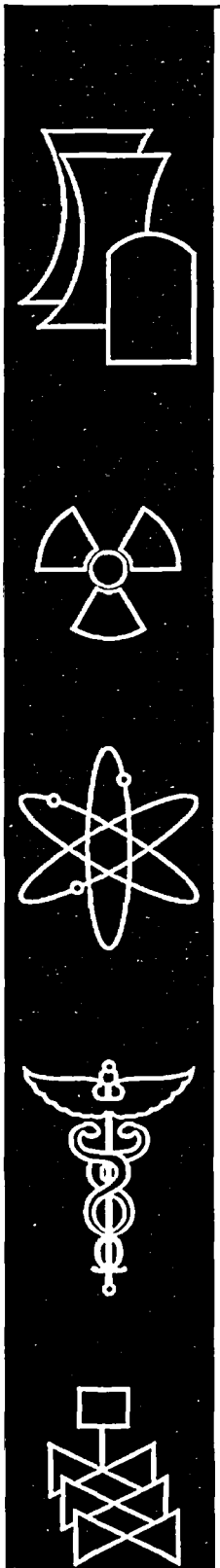
NUREG-1792



Good Practices for Implementing Human Reliability Analysis (HRA)

Draft Report for Comment

Science Applications International Corporation

Sandia National Laboratories

U.S. Nuclear Regulatory Commission
Office of Nuclear Regulatory Research
Washington, DC 20555-0001

AVAILABILITY OF REFERENCE MATERIALS IN NRC PUBLICATIONS

NRC Reference Material

As of November 1999, you may electronically access NUREG-series publications and other NRC records at NRC's Public Electronic Reading Room at http://www.nrc.gov/reading-rm.html.
Publicly released records include, to name a few, NUREG-series publications; *Federal Register* notices; applicant, licensee, and vendor documents and correspondence; NRC correspondence and internal memoranda; bulletins and information notices; inspection and investigative reports; licensee event reports; and Commission papers and their attachments.

NRC publications in the NUREG series, NRC regulations, and *Title 10, Energy*, in the Code of *Federal Regulations* may also be purchased from one of these two sources.

1. The Superintendent of Documents
   U.S. Government Printing Office
   Mail Stop SSOP
   Washington, DC 20402–0001
   Internet: bookstore.gpo.gov
   Telephone: 202-512-1800
   Fax: 202-512-2250
2. The National Technical Information Service
   Springfield, VA 22161–0002
   www.ntis.gov
   1–800–553–6847 or, locally, 703–605–6000

A single copy of each NRC draft report for comment is available free, to the extent of supply, upon written request as follows:

Address: Office of the Chief Information Officer,
Reproduction and Distribution
Services Section
U.S. Nuclear Regulatory Commission
Washington, DC 20555-0001
E-mail: DISTRIBUTION@nrc.gov
Facsimile: 301–415–2289

Some publications in the NUREG series that are posted at NRC's Web site address http://www.nrc.gov/reading-rm/doc-collections/nuregs are updated periodically and may differ from the last printed version. Although references to material found on a Web site bear the date the material was accessed, the material available on the date cited may subsequently be removed from the site.

Non-NRC Reference Material

Documents available from public and special technical libraries include all open literature items, such as books, journal articles, and transactions, *Federal Register* notices, Federal and State legislation, and congressional reports. Such documents as theses, dissertations, foreign reports and translations, and non-NRC conference proceedings may be purchased from their sponsoring organization.

Copies of industry codes and standards used in a substantive manner in the NRC regulatory process are maintained at—

The NRC Technical Library
Two White Flint North
11545 Rockville Pike
Rockville, MD 20852–2738

These standards are available in the library for reference use by the public. Codes and standards are usually copyrighted and may be purchased from the originating organization or, if they are American National Standards, from—

American National Standards Institute
11 West 42nd Street
New York, NY 10036–8002
www.ansi.org
212–642–4900

Legally binding regulatory requirements are stated only in laws; NRC regulations; licenses, including technical specifications; or orders, not in NUREG-series publications. The views expressed in contractor-prepared publications in this series are not necessarily those of the NRC.

The NUREG series comprises (1) technical and administrative reports and books prepared by the staff (NUREG–XXXX) or agency contractors (NUREG/CR–XXXX), (2) proceedings of conferences (NUREG/CP–XXXX), (3) reports resulting from international agreements (NUREG/IA–XXXX), (4) brochures (NUREG/BR–XXXX), and (5) compilations of legal decisions and orders of the Commission and Atomic and Safety Licensing Boards and of Directors' decisions under Section 2.206 of NRC's regulations (NUREG–0750).

NUREG-1792

Good Practices for Implementing Human Reliability Analysis (HRA)

Draft Report for Comment

Manuscript Completed: July 2004
Date Published: July 2004

Prepared by
A. Kolaczkowski/Science Applications International Corporation
J. Forester/Sandia National Laboratories
E. Lois/ U.S. Nuclear Regulatory Commission
S. Cooper/U.S. Nuclear Regulatory Commission

Science Applications International Corporation
405 Urban Street
Suite 400
Lakewood, CO 80220

Sandia National Laboratories
P.O. Box 5800
Albuquerque, NM 87185

E. Lois, NRC Project Manager

Prepared for
Division of Risk Analysis and Applications
Office of Nuclear Regulatory Research
U.S. Nuclear Regulatory Commission
Washington, DC 20555-0001

COMMENTS ON DRAFT REPORT

Any interested party may submit comments on this report for consideration by the NRC staff. Comments may be accompanied by additional relevant information or supporting data. Please specify the report number **(Draft NUREG-1792)**, in your comments, and send them-- by the end of the 90-day comment period specified in the *Federal Register* notice announcing the availability of this draft–to the following address:

Chief, Rules and Directives Branch,
Mail Stop T6-D59,
Office of Administration,
U.S. Nuclear Regulatory Commission,
Washington, DC 20555-0001.

You may also provide comments via the NRC's Web site:

http://www.nrc.gov/reading-rm/doc-collections/nuregs/docs4comment;

For any questions about the material in this report, please contact:

Erasmia Lois,
Mail Stop: T-10E50
U.S. Nuclear Regulatory Commission
Washington, DC 20555-0001
Phone: (301) 415-6560,
Email: exl1@nrc.gov or

or

Susan Cooper,
Mail Stop: T-10E50
U.S. Nuclear Regulatory Commission
Washington, DC 20555-0001
Phone: (301) 415-5183 or (302) 234-4423
Email: sec1@nrc.gov

ABSTRACT

The U.S. Nuclear Regulatory Commission is establishing "good practices" for performing human reliability Analysis (HRA) and reviewing HRAs to assess the quality of analyses. The HRA good practices are developed as part of the NRC's activities to address probabilistic risk assessment (PRA) quality issues and supports the implementation of Regulatory Guide (RG) 1.200 entitled: "An Approach for Determining the Technical Adequacy of Probabilistic Risk Assessment Results For Risk-Informed Activities."

The documented HRA good practices are of generic nature, that is, are not tied to any specific methods or tools that could be employed to perform an HRA. The good practices provide lower level guidance for implementing the RG 1.200 when performing a Level 1 and a limited Level 2 PRA for internal events (excluding fire) with the reactor at full power. Its elements are directly linked to RG 1.200 which reflects and endorses, with certain clarifications and substitutions, the American Society of Mechanical Engineers (ASME) Standard RA-S-2002, "Standard for Probabilistic Risk Assessment for Nuclear Power Plant Applications," and Revision A3 of the Nuclear Energy Institute (NEI) "Probabilistic Risk (PRA) Peer Review Process Guidance," (NEI-00-02).

TABLE OF CONTENTS

LIST OF TABLES

FOREWORD

This draft report documents good practices for implementing human reliability analysis (HRA). The report is being prepared as part of the NRC's activities to address probabilistic risk assessment (PRA) quality issues and supports the implementation of Regulatory Guide 1.200, which provides an acceptable approach for determining the technical adequacy of PRA results for risk informed regulatory decision making. The work was performed by the Office of Nuclear Regulatory Research (RES) with the support of Sandia National Laboratories.

The HRA good practices were developed on the basis of NRC and contractor experience on HRA method development (e.g., THERP and ATHEANA), performing HRAs (e.g., NUREG-1150) and reviewing HRAs, particularly the Individual Plant Examinations. The good practices are of generic nature, that is, are not tied to any specific methods or tools that could be employed to perform an HRA. Its elements are directly linked to elements of the American Society of Mechanical Engineers (ASME) PRA Standard for Nuclear Power Plant Applications, endorsed by Regulatory Guide 1.200. As such it provides lower level guidance for implementing the ASME PRA standard when performing a Level 1 and a limited Level 2 PRA for internal events (excluding fire) with the reactor at full power. The report provides a technical basis for performing an HRA or for formulating questions to evaluate the quality of an HRA. In this regard, it can be used to develop a regulatory guide and/or a standard review plan.

ACKNOWLEDGMENTS

We would like to thank Gareth Parry of the USNRC for several thorough reviews of this document at various stages of its development and for many important comments and suggestions. Similarly, we would like to thank Dennis Bley of Buttonwood Consulting, Inc., Bruce Hallbert of INEEL and James Bongarra, Andrew Kugler, Martin Stutske, Davis Lew, and Autumn Szabo of the USNRC for their reviews and comments.

Good Practices for Implementing Human Reliability Analysis (HRA)

1. INTRODUCTION

1.1 Background

In accordance with its policy statement [1] on the use of probabilistic risk assessment (PRA), during the last decade the Nuclear Regulatory Commission (NRC) has been increasingly using PRA technology in "all regulatory matters to the extent supported by the state of the art in PRA methods and data." Examples of risk informed initiatives are: undertaking risk-informed rulemaking activities such as risk-informing 10 CFR Part 50[2], generating a risk-informed framework for supporting licensee requests for changes to a plant's licensing basis (Regulatory Guide 1.174),[3] risk-informing the reactor oversight process, performing risk studies (e.g., for steam generator tube rupture (SGTR), and fire events), and evaluating the significance of events. In addition, the NRC is using PRA in the development of an infrastructure to license new reactors.

Given the increasing importance of the role of PRA in regulatory decision making, it is crucial that decision makers have confidence in the results produced by PRAs. To support this, the NRC has issued Regulatory Guide 1.200[4] that describes an acceptable approach for determining the technical adequacy of PRA results for risk informed activity. Regulatory Guide 1.200[4] reflects and endorses guidance provided by standards produced by societies and industry organizations. It currently addresses the American Society of Mechanical Engineers (ASME) Standard for Probabilistic Risk Assessment for Nuclear Power Plant Applications[5] which was developed for a full power, internal events (excluding fire) Level 1 PRA and a limited Level 2 PRA, and Nuclear Energy Institute's (NEI's) Probabilistic Risk Assessment Peer Review Process Guidance (NEI-00-02).[6]

The level of detail provided in Regulatory Guide 1.200,[4] the ASME Standard[5] and NEI-00-02[6] is at a high level, addressing what to do, but not how to do it. Consequently, there may be several approaches to address certain analytical elements, which though they may meet the guidance and standards, may do so by making different assumptions and approximations and, therefore, produce different results. This is particularly true of human reliability analysis (HRA) (see Section 1.2 for a discussion of HRA). Therefore, the guidance provided by these documents is not sufficient to address the detailed HRA quality issues that should be considered in regulatory decision making. For example, in Standard Review Plan 19[7], Section A.8, Modeling of Human Performance, the NRC staff is required to determine if "the modeling of human performance is appropriate." While Regulatory Guide 1.200[4], the ASME Standard[5] and NEI-00-02[6] can address whether the HRA addresses the right issues, they do not give detailed guidance on how they are addressed. Therefore, in order to support the review of human performance issues in the context of PRAs, the NRC is developing this guidance for performing and reviewing HRAs, as a document supporting Regulatory Guide 1.200[4]. The guidance is being developed in two phases. The first phase is the development of this "HRA Good Practices" document which has been prepared on the basis of the NRC experience and lessons learned from developing HRA methods (e.g., THERP,[8] SLIM,[9] and ATHEANA[10]), performing HRAs (e.g., NUREG-1150[11] studies), and reviewing HRAs (in particular the individual plant examinations [IPEs]). The second phase is a review and evaluation of existing HRA approaches for their capability to meet the good practices when employed to address different regulatory applications.

This volume describes what the NRC staff regards as good practices for a HRA as implemented within a broader PRA. The volume is written in the context of a risk assessment for commercial nuclear power plant (NPP) operations occurring nominally at full power. It is specifically intended for applications involving internal initiating events but should generally be appropriate for external initiating events. While written for full power applications, many of the good practices will also be applicable to low power and shutdown operations. Additionally, elements of this volume may be of benefit in examining human actions related to nuclear materials and safeguard types of applications.

As with any evolving technology, both PRA and the implementation of HRA within the PRA framework are continuing to improve. Hence, what is a good practice today may be somewhat inferior or outdated tomorrow. Much of what is in this volume will always constitute good practice; some of it may be subject to newer technology, methods, and tools. For this reason, this volume must be considered a snapshot of good practices in HRA circa 2004.

With the expectation that PRA will continue to be used in the commercial nuclear industry in assessing current operating risks, in estimating changes in risk as a result of temporary and permanent plant changes to existing plants, and as an adjunct to the design process of newer generation plants, it is important that HRA practitioners perform human reliability analyses in accordance with good practices and that reviewers recognize the implementation of good practices (or failure to do so) in these analyses.

It should be noted that failure to meet particular good practices will not necessarily render the analysis and its results invalid or unacceptable. As is discussed further below, for certain applications, it may not be necessary to thoroughly meet all of the good practices. With the good practices in mind, practitioners and reviewers will have to decide whether the application at hand can be performed adequately without completely meeting each good practice. Such judgments will require careful thinking about the goals of the analysis, the issues being addressed, and the importance of each good practice relative to those goals and issues. Knowing what the good practices are, however, at least provides a starting point for practitioners to decide what must be done and for reviewers to determine whether the analysis was done rigorously enough to address the issue.

1.2 HRA in the context of PRA

Human reliability analysis in the PRA context is that discipline that identifies and provides probabilities for the human failure events that can negatively impact normal or emergency plant operations. The human failure events (HFEs) modeled in PRAs that are associated with normal plant operation include: 1) events that leave equipment in an unrevealed, unavailable state, (e.g., miscalibration of a level sensor), 2) those that induce an initiating event (typically captured by the initiating event frequency), such as a human-caused loss of feedwater or 3) those modeled as human events contributing to an initiating event (e.g., for a total loss of service water, failing to backup the start of service water train B upon loss of train A). The human failure events modeled in PRAs associated with emergency plant operation include events that, if not performed, do not allow the desired function to be achieved, such as failing to initiate feed and bleed. Quantification of the probabilities of the human failure events is based on plant and accident specific conditions, where applicable, including any dependencies among actions and conditions.

This document provides HRA good practices that when implemented will result in determining the impacts of human actions as *realistically as necessary* in an assessment of risk. Note the emphasis on realistic as necessary rather than as realistic as possible. For example, depending on the purpose for which the PRA is to be used, a conservative treatment of human performance may be sufficient to address a PRA application; more realism may not be necessary and could be a waste of resources. However, a conservative approach may not be sufficient when used as the basis for not needing to further investigate the issue at hand. Such an approach could potentially constrain the capability of identifying weaknesses in plant operations and plant practices related to the particular human actions credited in the PRA.

Most of the HRA good practices should be addressed to some degree for any PRA application. However, recognizing that the document will be used to guide a wide variety of applications, it is not intended that all the practices be met for any specific PRA application; in fact, some may not be relevant or necessary. For example, a specific application may impact only post-initiator human events. In such a case, all the good practices associated with pre-initiator human events would not be relevant. As another example, rather than assessing the degree of dependencies among human actions as described in this document, it may be more efficient and acceptable for an application to simply assume complete dependence among the actions because the issue is only demonstrating that the resultant risk metric (e.g., core damage frequency) is no worse than some threshold value. In this case, using a conservative treatment for the human action dependencies is sufficient. As discussed in Section 1.1, practitioners and reviewers should determine the applicable good practices for the PRA application and perform or review the HRA accordingly.

In some cases, the necessity for meeting certain good practices may be a function of the importance of the human actions being modeled. Documents such as NUREG-1764,[12] "Guidance for the Review of Changes to Human Actions," provide guidance for categorizing human actions by measures of importance. Such a categorization scheme may help reviewers determine the degree of compliance with HRA good practices that is appropriate for each human action. However, as noted above, relative "importance" in the PRA should not be the only criterion for determining the needed level of analysis. Additional or other criteria may be needed because generic, non-detailed, or screening level analyses can fail to detect plant vulnerabilities and weaknesses in plant practices.

1.3 Purpose

This document serves as a reference guide of good practices in HRA. By "good practices," we mean those processes and individual analysis tasks and judgments that would be expected of a HRA (considering current knowledge and state-of-the-art) in order for the HRA results to sufficiently represent the anticipated operator performance when making risk-informed decisions. The document is principally focused on the process for performing HRA and does not, for instance, specifically address HRA data or details of specific quantification approaches. As such, it is written in a way that links the prescribed good practices to Regulatory Guide 1.200[4] and subsequent ties to requirements in the ASME Standard[5] and particularly the HRA section of that document. However, nearly all other sections of the ASME standard[5] also have some parallel requirements with regard to operator actions such as in the accident sequence analysis, success criteria, systems analysis, and large early release frequency (LERF) analysis sections. Appendix A provides a cross-reference table that links the good practices in this document to the corresponding sections in the ASME Standard,[5] Regulatory Guide

1.200,[4] and NEI-00-02,[6] as appropriate.

This document has at least two primary uses.

1. It provides guidance for performing HRAs (whether for the first time or when analyzing a change to current plant practices). It supports the implementation of Regulatory Guide 1.200[4] and focuses on the attributes of a good HRA regardless of the specific methods or tools that are used. The guidance is specifically for HRAs for reactor, full power, and internal events applications although most of the guidance should be useful for other applications (e.g., external events, other operating modes). It does not endorse nor is it meant to suggest that a specific method or tool be used since many exist, and all have strengths and limitations regarding their use and applicability. Nevertheless, the good practices come from those advocated in such sources as the ASME Standard[5], NEI -00-02[6], THERP[8], ASEP[13], SHARP1[14], SPAR-H Method[15], and ATHEANA[10], as well as the experiences of the authors and reviewers of this document.

2. It supports the review of HRAs in assessing the quality of the analyses. In this regard, the HRA good practices provided here should be useful in formulating questions about and evaluating the quality of a HRA. Its purpose is not to explicitly provide questions a reviewer should ask, but rather to provide the technical basis for developing questions and potentially a standard review plan.

2. OVERVIEW OF GOOD PRACTICES FOR HRA

2.1 Scope of the report

As stated in Section 1, the purpose of this report is to ensure consistency and quality in performing or reviewing HRAs. In order to assist in achieving consistency and quality in HRA practices, this document provides lower level guidance than that contained in Regulatory Guide 1.200[4], the ASME Standard[5] and NEI-00-02[6] PRA guidance. As such it is directed at specific HRA tasks or activities.

The performance of HRA typically involves several tasks or activities. Some of these tasks are dependent on the HRA method or quantification approach that is used. Because this HRA good practices document does not endorse or specify the use of specific HRA methods or quantification approaches, most of the guidance in this document is directed at the process for performing HRA. However, this document does provide some non-method-specific good practices with respect to HRA quantification.

The report is written assuming the HRA in question supports a Level 1 internal events PRA. It explicitly addresses good practices for both pre-initiator and post-initiator event analyses. It does not explicitly address human actions that cause or contribute to initiating events.

In current PRA practices, initiating event frequencies typically are developed by aggregating data from either equipment-induced or human-induced initiators. As a result, the contribution of human performance on the frequency of initiating events and, therefore, on plant risk, is not separately analyzed in most at-power PRAs today. However, the good practices documented here for pre- or post-initiating events would be applicable for an explicit treatment of human-induced failures. For example, fault trees developed for support system initiators could include HFEs depicting those actions that could lead to a human-induced support system failure. Such actions will have the characteristics of either pre- or post-initiating event HFEs. The techniques needed to analyze such HFEs are similar to those described here for either pre-initiating or post-initiating events.. Therefore the good practices needed to explicitly address human-induced initiators are covered by this document. (See for example, the high level requirement HLR-IE-C in the ASME Standard[5] and supporting requirements such as IE-C9 concerning the modeling of recovery actions in an initiator fault tree, and IE-C12 concerning procedural influences on the interfacing system loss of coolant accident (ISLOCA) frequency.) However, an explicit modeling of human-induced initiating events could potentially lead to important insights with respect to the overall significance of human performance on safety and enhance the predictability of PRA models. As the use of PRA for decision making is being expanded, the need to separately address human-induced initiators may emerge.

2.2 HRA good practices and the state-of-the-art in HRA

The HRA good practices given in this document were developed on the basis of past experience in performing and reviewing HRAs, including those used to support the IPEs, but also reflect perspectives regarding the impact of human performance gained from domestic and international HRA developmental efforts during the 1990s (e.g., ATHEANA[10], MERMOS[16] and CREAM[17]).

Consistent with the state-of-the-art in HRA, it is recommended that future HRA/PRAs attempt to identify and model not only errors of omission (EOOs) as is typically done, but also potentially important errors of commission (EOCs). Several existing sources provide useful guidance for identifying and treating EOCs in the context of PRAs, e.g., ATHEANA,[10] CESA,[18] Julius et al.,[19] and Wakefield.[20] This report provides guidance for identifying characteristics of situations that can facilitate EOCs, particularly in the context of risk-informed analyses being done to support plant changes. The good practices on EOCs would apply to any HRA method used for quantification.

2.3 Report organization

The good practices are presented in a PRA logical analysis approach. Chapter 3 provides overall HRA good practices. Chapters 4 and 5, for pre-initiating and post-initiating HFEs, respectively, address HRA good practices for identifying, screening, modeling, and quantifying HFEs. Chapter 6 discusses good practices for modeling EOCs, Chapter 7 provides good practices for documenting the analyses, and Chapter 8 provides the list of references. Appendix A presents a table of cross-references between the Good Practices in this document and the corresponding sections in the ASME Standard[5] and NEI 00-02[6]. It also notes where Regulatory Guide 1.200 has added clarification or qualification to the corresponding items in those documents. Appendix B provides guidance on the consideration of performance shaping factors (PSFs) for post-initiator HFEs .

While the document is written in a serial fashion, in practice, it is often desirable to perform or review a HRA in a more holistic manner and address multiple steps of the HRA process simultaneously to achieve greater resource efficiency.

2.4 Summary of HRA good practices

Table 2-1 below provides a summary of the good practices.

Table 2-1 Summary of Good Practices

Analysis Activity	Good Practice	Section
HRA team formation and techniques for a realistic analysis	GP 1 - Perform a Multi-Disciplinary, Integrated Analysis. The HRA assessment should involve a multi-disciplinary team that interacts with the rest of the PRA team in addressing each accident scenario at each stage of the analysis.	3.1.3.1
	GP 2 - Perform Field Observations and Discussions. In addition to the review of plant documents, the HRA should include walkdowns of relevant actions (particularly local actions), observations of simulator exercises, talk-throughs of accident scenarios and related actions with plant operators and trainers, and other field observations and discussions as needed.	3.1.3.2
Pre-Initiators: Identifying human actions that could leave equipment unavailable	GP 1 - Review Pre-Initiator Procedures, Actions, and Equipment. All routine (scheduled) test and maintenance as well as calibration procedures that affect equipment to be credited in the PRA should be identified and reviewed. Actions and equipment specified in the procedures should be examined to determine whether misalignment or miscalibration could occur and render the equipment unavailable or faulty. If so, the actions and equipment should be "considered for modeling" at least until initial screening is performed.	4.1.3.1
	GP 2 - Do Not Ignore Pre-Initiators. The identification process should identify pre-initiator human actions even if they may be potentially covered by the affected equipment failure data.	4.1.3.2
	GP 3 - Examine Other Operational Modes and Routine Actions Affecting Structures (if applicable). The identification process should address other operational modes and routine actions affecting barriers and other structures such as fire doors, block walls, drains, seismic restraints, etc. (if applicable for the analysis).	4.1.3.3
	GP 4 - Identify Actions Affecting Redundant and Multiple Diverse Equipment. The identification process needs to include possible pre-initiator actions *at least within each system* where redundant or multiple diverse equipment can be affected by (a) a single act, or (b) through a common failure with similar multiple acts.	4.1.3.4
Pre-Initiators: Screening human actions that do not need to be modeled	GP 1 - Screen Pre-Initiators with Acceptable Restoration Mechanisms or Aids. Pre-initiators with signals, signs, or checks that help ensure that the equipment will be reliably restored to its desired state can be screened from further analysis (with the exception noted in the next good practice below).	4.2.3.1
	GP 2 - Do Not Screen Actions Affecting Redundant and Multiple Diverse Equipment. In general, do not screen those pre-initiator actions that simultaneously affect multiple (redundant or diverse) equipment items.	4.2.3.2

Analysis Activity	Good Practice	Section
Pre-Initiators: Screening human actions that do not need to be modeled (continued).	GP 3 - Re-Evaluate the Screening Process For Special Applications. For a specific PRA application such as a plant change or analysis of a special issue, revisit the original PRA to ensure that the pre-initiator screening is still valid for the current application.	4.2.3.3
Pre-Initiators: Modeling specific human failure events (HFEs) corresponding to the unscreened human actions	GP 1 - Include HFEs for Unscreened Human Actions in the PRA Model. Define each specific pre-initiator HFE to be modeled in the PRA as a basic event that describes the human-induced failure mode and preferably locate it in the model such that it is linked closely to the unavailability of the affected component, train, system, or overall function (i.e., level of modeling) depending on the effect(s) of the HFE.	4.3.3.1
Pre-Initiators: Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs	GP 1 - Use Screening Values During the Initial Quantification of the HFEs. Use of screening HEPs is acceptable provided (a) it is clear that the individual values used are over-estimations of the probabilities if detailed assessments were to be performed and (b) dependencies among multiple HFEs appearing in an accident sequence are conservatively accounted for. Individual screening values should never be less than 1E-2 and the joint probability of multiple HEPs in a sequence should not be lower than 5E-3.	4.4.3.1
	GP 2 - Perform Detailed Assessments of Significant HFEs. To help understand the role of significant HFEs in plant safety and the factors influencing their likelihood, a detailed assessment (quantification) of at least the significant HFEs should be performed.	4.4.3.2
	GP 3 - Revisit the Use of Screening Values vs. Detailed Assessments For Special Applications. For a specific PRA application such as a plant change or analysis of a special issue, revisit the original PRA to ensure that the appropriate HFEs receive detailed assessments in the new analysis.	4.4.3.3
	GP 4 - Account for Plant and Activity Specific Performance Shaping Factors (PSFs) in the Detailed Assessments. Pre-initiator HEP assessments should account for the most relevant plant-specific and activity-specific performance-shaping factors (PSFs) in the analysis. Potentially important PSFs include written work plans, procedures, training, complexity and number of steps, reliance on memory, ergonomics, and the task environment.	4.4.3.4
	GP 5 - Apply Plant Specific Recovery Factors. To the extent the plant has "built-in" specific practices to recover any failures of pre-initiator human actions, they should be applied to the HEP evaluations for the HFEs. Multiple recoveries may be acceptable, but any dependencies among the initial failure and the recoveries, and among the recoveries themselves, must be considered. .Example recovery factors include testing, independent verification, scheduled checks, and compelling signals.	4.4.3.5

Analysis Activity	Good Practice	Section
Pre-Initiators: Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs (continued)	GP 6 - Account for Dependencies Among the HEPs in an Accident Sequence. Dependencies should be quantitatively accounted for by deriving HEPs so that they reflect commonalities and relationships among the HFEs. A main concern is human related "common -cause factors that would lead to similar errors occurring across similar systems. The impact of recovery factors should be included in evaluating dependencies.	4.4.3.6
	GP 7- Assess the Uncertainty in Mean HEP Values. Point estimates should be mean values for each HEP (excluding screening HEPs) and an assessment of the uncertainty in the mean values should be performed at least for the significant HEPs. Typical assessments of uncertainty involve propagating uncertainty distributions for the HEPs through the quantitative analysis of the entire PRA, performing sensitivity analyses that demonstrate the effects on the risk results for extreme estimates in the HEPs based on at least the expected uncertainty range about the mean value, or addressing through qualitative arguments. Aleatory and epistemic uncertainties should be addressed as necessary.	4.4.3.7
	GP 8 - Evaluate the Reasonableness of the HEPs Obtained Using Detailed Assessments. The pre-initiator HEPs (excluding the screening HEPs) should be reasonable from two standpoints: 1)first and foremost, relative to each other (i.e., the probabilistic ranking of the failures when compared one to another), and 2) in absolute terms (i.e., each HEP value), given the relative strengths of the positive and negative PSFs identified as being important and the presence or absence of recovery factors. Example evaluation techniques include consideration of actual plant history, comparisons with results of other analyses, and qualitative understanding of the actions and their contexts by experts.	4.4.3.8
Post-Initiators: Identifying post-initiator human actions (Note that the three GPs associated with this activity are to be performed in an iterative manner and a stringent order is not implied.)	GP 1 - Review Post-Initiator Related Procedures and Training Materials. Plant-specific emergency operating procedures (EOPs), abnormal operating procedures (AOPs), annunciator procedures, system operating procedures, and severe accident management guidelines (SAMGs) should be reviewed. Other relevant special procedures (e.g., fire emergency procedures) should also be reviewed as appropriate. Observations of simulator exercises and talk-throughs of accident scenarios and related actions with plant operators and trainers can support the identification of post-initiator human actions at this stage.	5.1.3.1

Analysis Activity	Good Practice	Section
Post-Initiators: Identifying post-initiator human actions (continued)	GP 2 - Review Functions and Associated Systems and Equipment to be Modeled in the PRA. The PRA team's plant and system knowledge should be used to identify critical functions and equipment needed and not needed for the given accident scenario. In addition, ways the equipment can functionally succeed and fail should be determined, along with ways the operators are 1) intended/required to interact with the equipment and 2) how they are to respond to equipment failure modes that can cause undesired conditions per the PRA. During the identification process, it is helpful to use action words such as actuate, initiate, isolate, terminate, control, change, etc. so that the desired actions are clear.	5.1.3.2
	GP 3 - Look for Certain Expected Types of Actions. The types of actions expected to be identified as post-initiator human actions include: necessary and desired actions to directly provide a critical function, backup actions to failed automatic responses, and anticipated procedure-guided or skill-of-the-craft recovery actions . Although the actions modeled will generally be error of omission, identification of errors of commission may sometimes be important.	5.1.3.3
Post-Initiators: Modeling specific human failure events (HFEs) corresponding to the human actions	GP 1 - Include HFEs for Needed Human Actions in the PRA Model. Define each specific post-initiator HFE to be modeled in the PRA as a basic event that describes the human-induced failure mode and preferably locate it in the model such that it is linked closely to the unavailability of the affected component, train, system, or overall function (i.e., level of modeling) depending on the effect(s) of the HFE. The nature of the action, the consequences if its failure, the nature of the sub-tasks involved, and the level of detail already in the model should be considered in deciding how to model the HFEs, e.g., where it is placed and whether it should be broken into separate HFEs. Dependencies must also be considered.	5.2.3.1
	GP 2 - Define the HFEs Such that they are Plant and Accident Sequence-Specific. Each of the modeled post-initiator HFEs should be defined such that they are plant and accident sequence-specific, and the basic events representing them are labeled uniquely. In order for the act to occur, the operator must diagnose the need to take the act and then execute the act. While many performance shaping factors are used to quantify the probability for failing to perform the act correctly (as discussed later under quantification), all of which should be evaluated based on plant and accident sequence-specifics, consideration of plant and accident specific timing information, along with procedural and training information is critical.	5.2.3.2

Analysis Activity	Good Practice	Section
Post-Initiators: Modeling specific human failure events (HFEs) corresponding to the human actions (continued)	GP 3 - Perform Talk-Throughs, Walkdowns, Field Observations, and Simulator Exercises (as necessary) to Support the Modeling of Specific HFEs. To fully understand the nature of the act(s) (e.g., who performs it, what is done, how long does it take, whether there are special tools needed, whether there are environmental issues or special physical needs, whether there is a preferred order of use of systems to perform a specific function, etc.) and help define the HFEs and their context, additional reviews, talk-throughs, walkdowns, field observations, and simulator exercises are performed (as discussed in Good Practice #2 under Section 3.1.3.2, with more about the benefits of these techniques presented in Appendix B). In addition, the results of these activities may add to the list of actions and/or help interpret how procedural actions should be defined based on how they are actually carried out.	5.2.3.3
Post-Initiators: Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs	GP 1 - Address Both Cognitive and Response Execution Failures. Whether using conservative or detailed estimations of the post-initiator HEPs, the evaluation should include both cognitive (i.e., thinking) as well as execution failures.	5.3.3.1
	GP 2 - Use Screening Values During the Initial Quantification of the Pots-Initiator HFEs. The use of conservative human error probability (HEP) estimates to screen unimportant HFEs is acceptable provided (a) it is clear that the individual values used are over-estimations of the probabilities if detailed assessments were to be performed AND (b) dependencies among multiple human failure events appearing in an accident sequence are conservatively accounted for. Individual screening values should never be less than 0.1 and the joint probability of multiple HEPs in a sequence should not be lower than 0.05.	5.3.3.2
	GP 3 - Perform Detailed Assessments of Significant Post-Initiator HFEs. To help understand the role of significant HFEs in plant safety and the factors influencing their likelihood, a detailed assessment (quantification) of at least the significant HFEs should be performed. Non-significant HFEs should also be assessed in detail if detection of <u>all</u> weaknesses in plant design or practices is desirable given the application.	5.3.3.3
	GP 4 - Revisit the Use of Post-Initiator Screening Values vs. Detailed Assessments For Special Applications. For a specific PRA application such as a plant change or analysis of a special issue, revisit the original PRA to ensure that the appropriate HFEs receive detailed assessments in the new analysis.	5.3.3.4

Analysis Activity	Good Practice	Section
Post-Initiators: Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs (continued)	GP 5 - Account for Plant and Activity Specific PSFs in the Detailed Assessments of Post-Initiator HEPs. Post-initiator HEP assessments should account for the most relevant plant-specific and activity-specific performance-shaping factors (PSFs). Potentially important main control room PSFs include (but are not limited to): procedures (and how the procedures are implemented), training, task complexity, workload, team dynamics, and scenario timing, Potentially important PSFs for local actions include (but are not limited to): procedures (and how the procedures are implemented), training, task complexity, workload (staff available for the action) , team dynamics, scenario timing, communication requirements, number of steps, reliance on memory, ergonomics, task environment, accessability, special fitness needs, and the need and location of special tools.	5.3.3.5
	GP 6- Account for Dependencies Among Post-Initiator HFEs. Dependencies among the post-initiator HEPs in an accident sequence should be quantitatively accounted for in the PRA model by virtue of the joint probability used for the HEPs. In analyzing for possible dependencies among the HFEs in an accident sequence, look for links among the acts including: the same crew member(s) is responsible for the acts, the actions take place relatively close in time in the sense that a crew "mindset" or interpretation of the situation might carryover from one event to the next, the procedures and cues used along with the plant conditions related to performing the acts are identical (or nearly so) or related, and the applicable steps in the procedures have few or no other steps in between the applicable steps, there are similar performance shaping factors for performing the acts, how the acts are performed is similar and they are performed in or near the same location, and the interpretation of the need for one action might bear on the crews decision regarding the need for another action. Once all the relationships are considered and the dependencies are included in the HEP values, the total combined probability of all the HFEs in the same accident sequence/cut set should not be below the ~0.0001 to 0.00001 range.	5.3.3.6
	GP 7 - Assess the Uncertainty in Mean HEP Values. Mean values for each HEP (excluding conservative HEPs) and an assessment of the uncertainty in the mean values are performed at least for the significant HEPs to the extent that these uncertainties need to be understood and addressed in order to make appropriate risk-related decisions . Typical assessments of uncertainty involve propagating uncertainty distributions for the HEPs through the quantitative analysis of the entire PRA, performing sensitivity analyses that demonstrate the effects on the risk results for extreme estimates in the HEPs based on at least the expected uncertainty range about the mean value, or addressing through qualitative arguments. Aleatory and epistemic uncertainties should be addressed.	5.3.3.7

Analysis Activity	Good Practice	Section
Post-Initiators: Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs (continued)	GP 8 - Evaluate the Reasonableness of the HEPs Obtained Using Detailed Assessments. The post-initiator HEPs (excluding the screening HEPs) should be reasonable from two standpoints: 1) first and foremost, relative to each other (i.e., the probabilistic ranking of the failures when compared one to another), and 2) in absolute terms (i.e., each HEP value), given the context and combination of positive and negative PSFs and their relative strengths. Example evaluation techniques include consideration of actual plant history, comparisons with results of other analyses, and qualitative understanding of the actions and their contexts by experts	5.3.3.8
Post-Initiators: Adding recovery actions to the PRA	GP 1 - Define Appropriate Recovery Actions. Based on the failed functions, systems, or components, identify recovery actions to be credited that are not already included in the PRA (e.g., aligning another backup system not already accounted for...) and that are appropriate to be tried by the crew to restore the failure. Examples of aspects to consider include the following: whether the cues will be clear and provided in time to indicate the need for a recovery action, the most logical recovery actions for the failure based on the cues that will be provided, that the recovery is not a repair action (e.g., the replacement of a motor on a valve so that it can be operated), whether sufficient time is available, whether sufficient crew resources exist to perform the recovery(ies), whether there is procedure guidance to perform the recovery(ies), whether the crew has trained on the recovery action(s) including the quality and frequency of the training, whether the equipment needed to perform the recovery(ies) is accessible and in a non-threatening environment (e.g., extreme radiation), and whether the equipment needed to perform the recovery(ies) is available in the context of other failures and the initiator for the sequence/cut set.	5.4.3.1
	GP 2 - Account for Dependencies. All the good practices provided above for post-initiator HFEs apply specifically to recovery actions as well. Particular attention should be paid to accounting for dependencies among the HFEs including the credited recovery actions.	5.4.3.2
	GP 3 - Quantify the Probability of Failing to Perform the Recovery(ies). Per the quantification good practices above, quantify the probability of failing to perform the recovery(ies) by: 1) using representative data that exists and deemed appropriate for the recovery event, or using the HRA method/tool(s) used for the other HFEs (i.e., using an analytical/modeling approach). If using data, ensure the data is applicable for the plant/sequence context or that the data is modified accordingly.	5.4.3.3

Analysis Activity	Good Practice	Section
Errors of Commission (EOCs)	GP 1- Address EOCs in Future HRAs/PRAs (Recommendation). Given the recent advances in the ability to address EOCs and the potential for regulatory requirements to make the need to address EOCs more important, it is recommended that future HRA/PRAs identify and model, to the extent necessary, potentially important EOCs. For any EOCs modeled, the guidance given in this document for either pre-initiators or post initiators are applicable for identifying, modeling, and quantifying EOCs. , That is, the same good practices apply whether the error is one of omission or commission.	6.1.3.1
	GP 2 - As a Minimum, Search for Conditions that May Make EOCs More Likely. The use of risk in any issue assessment should at least ensure that conditions that promote likely EOCs do not exist. For example, it should be ensured that such conditions have not been introduced by a plant change or modification, or that the plant is not more susceptible to EOCs under the unique set of conditions being examined.. When considering the potential for situations that may make EOCs somewhat likely, the premise of any evaluation should be that: 1) operators are performing in a rationale manner (e.g., no sabotage), and 2)the procedural and training guidance is being used by the crew based on the plant status inputs they are receiving. Using this premise, EOCs are considered to be largely the result of problems in the plant information/operating crew interface (wrong, inadequate information is present, or the information can be easily misinterpreted) or in the procedure-training/operating crew interface (procedures/training do not cover the actual plant situation very well because they provide ambiguous guidance, no guidance, or even unsafe guidance for the actual situation that may have evolved in a somewhat unexpected way). In either case, significant mismatches can occur between the scenario conditions and the crew's understanding of those conditions. Such mismatches should be searched for and their potential for leading to EOCs should be examined.	6.1.3.2

Analysis Activity	Good Practice	Section
HRA Documentation	GP 1 - Document the HRA. The HRA should be documented well enough to allow a knowledgeable reviewer to understand the analysis to the point that it can be at least approximately reproduced and the resulting conclusion reached, if the same methods, tools, data, key assumptions, and key judgments and justifications are used. Hence, the documentation should include the following, but only to the extent it is applicable for the application: the overall approach and disciplines involved in performing the HRA, including to what extent talk-throughs, walkdowns, field observations, and simulations were used; summary descriptions of the HRA methodologies, processes, and tools used; assumptions and judgments made in the HRA, their bases, and their impact on the results and conclusions; the PSFs considered (for at least each of the HFEs important to the risk decision to be made), the bases for their inclusion, and how they were used to quantify the HEPs, along with how dependencies among the HFEs and joint probabilities were quantified; the sources of data and related bases or justifications for the screening and conservative values, and the best estimate values, along with their uncertainties with related bases; the results of the HRA including a list of the important HFEs and their HEPs; and the conclusions of the HRA.	7.1.3.1

3. HRA TEAM FORMATION AND OVERALL GUIDANCE

If human actions are going to be included realistically in the PRA, the modeling of human interactions must consider each action evaluated in the context of a complete accident scenario or sequence of events. To do this, HRA has evolved from the days when PRA analysts provided the human events of interest to a HRA specialist who then assigned human error probabilities (HEPs) to the human events, sometimes in isolation. Such a process is no longer considered good practice. Understanding an accident sequence context is a complex, multi-faceted process. The interaction of plant hardware response and the response of plant operators must be investigated and modeled accordingly. Characteristics such as those in the following list need to be understood and reflected, as necessary, in the model of a specific human action or group of actions:

- plant behavior and conditions
- timing of events and the occurrence of human action cues
- the parameter indications used by the operators and changes in those parameters as the scenario proceeds
- the time available and locations necessary to take the human actions
- the equipment available for use by the operators based on the sequence
- the environmental conditions under which the decision to act must be made and the actual response must be performed
- the degree of training guidance and procedure applicability, among many other characteristics.

Much of the guidance in this document is aimed at good practices for understanding the context associated with each modeled human action, and how that context affects both the definition of human failure events and an assessment of their probabilities.

This emphasis on the need to adequately understand and address context in order to more realistically address human performance is based on advances in our understanding of the factors that can influence human performance. These advances come from recent reviews of operational events involving serious accidents (e.g., ATHEANA[10]) and from other international efforts and recent research in the cognitive sciences that together have provided a clearer picture of the ways in which various factors and situations can interact to influence the occurrence of inappropriate human actions (e.g., Reason[21], Woods[22], and Endsley[23]). Improvements have been made in how to address the broad range of potential influences on human performance, on both the identification of the human actions to be modeled in the PRA as well as what to consider during screening and detailed quantification of the actions. The guidance in this document provides good practices that reflect these improvements and ensures the proper treatment of context in performing a reasonably realistic HRA.

Identifying the appropriate participants for the HRA (i.e., forming the HRA team) and encouraging the use of certain techniques throughout the HRA process, both to ensure that the context associated

with each human action is properly understood and reflected in the PRA, is an activity discussed below with its associated good practices.

3.1 HRA team formation and techniques for a realistic analysis

3.1.1 OBJECTIVE: To provide a strong underlying basis to ensure the context for each human action is adequately determined. Properly determining the correct context associated with each modeled human action should lead to HRA results that are credible.

3.1.2 REGULATORY GUIDE 1.200 POSITION:

There is no specific guidance on team formation. However, because of past practices whereby HRA was often treated as an isolated task or for which much of the analysis was performed without, for instance, examining the locations involving the actions or obtaining operators and trainers perspectives on the specific scenarios, the following good practices are warranted to stress their importance. Further, while there are references to the use of talk-throughs and simulations in the regulatory guide and its endorsement of the ASME Standard[5], the need to broaden the use of such techniques is emphasized here to confirm or expand any judgement or assumption associated with understanding the context for a human action (e.g., not just to verify the time it takes to perform an action).

3.1.3 GOOD PRACTICES

3.1.3.1 Good Practice #1 - Perform a Multi-Disciplinary, Integrated Analysis:

The HRA should be an integral part of the PRA (not performed as an isolated task in the PRA process) whereby the inputs from the following types of disciplines are used together to define the PRA structure including which human events need to be modeled, how they are defined and modeled in the PRA, and the considerations used to quantify the associated HEPs:

- PRA modelers
- HRA practitioners
- Human factors specialists
  Thermal-hydraulic analysts
- Operations, training, and maintenance personnel
- Other disciplines (e.g., structural engineers, system engineers) as necessary (e.g., structural engineers if the timing of an action is dependent on when and how the containment might fail).

Each discipline provides a portion of the context knowledge. When the context is sufficiently understood, only then can human failure events be realistically modeled and quantified.

3.1.3.2 Good Practice #2 - Perform Field Observations and Discussions:

Besides the review of plant documents, the HRA is performed using the insights gained from the activities listed below . Performance of the activities will allow analysts to confirm judgments and

assumptions made from the document review and help them obtain a more well-informed understanding of the context for the various actions and scenarios. If these activities are not performed, there will be uncertainties associated with the context that could be important. Thus, such activities should only not be performed when a generic, non-detailed analysis is deemed adequate in the context of the issue being examined, e.g., the NRC's accident sequence precursor analyses. The activities include the following:

- Walkdowns of areas where decisions and actions are to take place (including timing of relevant actions)

- Talk-throughs of scenarios and actions of interest with plant operators, trainers, or maintenance staff

- Other field observations to help understand the equipment involved, the plant layout, and the general plant "environment"

- Simulator exercises (While it is realized that simulator exercises may not always be possible, it is good practice that at least a couple of the main, nominal scenarios for the issue under investigation be observed by the HRA team. In addition to allowing analysts to obtain scenario-specific information, it also allows them to see how plant crews perform as a team and implement their procedures. Moreover, observation of simulator exercises also provides a basis for discussions with operators and trainers about both the scenarios observed and those unobserved due to time or resource constraints.)

3.1.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES

Not meeting the above good practices may have little to no impact on the HRA, or it may have a significant impact. Generally, for human actions for which a very simple and straightforward context can reasonably be assumed, no impact is likely. For those actions for which a more complex context is likely (e.g., scenarios with multiple failures [equipment and/or human actions] or relatively unexpected scenarios), there could be a much greater impact (e.g., grossly under-estimating the failure probability for a human action). While there is no direct way to "measure" the effect of not performing the above good practices, the implication is that the context used to model and quantify a human action *may not* be correct or complete enough if the above team participants and confirmation techniques are not used at appropriate stages during the analysis. In other words, the modeling and quantification of the human action may not accurately reflect "the current design and operating practices" as discussed in Regulatory Guide 1.200.

The impact, if any, will be manifested via the other good practices such as not addressing all the appropriate performance shaping or recovery factors, or in results that are incorrect or do not make sense. If this occurs, the source of the insufficiency may be that the good practices in this section were not met (i.e., the team makeup and participation at various stages of the analysis was inadequate or the information gathering and confirmation techniques were not used, leading to incorrect judgements or assumptions about the context). For this reason, these good practices have been provided as an adjunct to all the other good practices in this document, since they provide an underlying basis for the success of the entire HRA process.

9. PRE-INITIATOR HRA

The NRC staff has stated their positions on the ASME Standard[5] in Appendix A of Regulatory Guide 1.200[4]. The standard separates its requirements into two broad classifications; those that address the modeling of failures of pre-initiator human actions and those that address the modeling of failures of post-initiator human actions. This section provides good practices for implementing the requirements for addressing pre-initiator human failure events in a PRA.

Pre-initiator human failure events are events that represent the impact of human failures committed prior to the initiation of an accident sequence (e.g., during test or maintenance or the use of calibration procedures). They are important to model because plant personnel can make the equipment needed to mitigate a particular accident sequence unavailable, thus reducing the overall capability to respond to the initiating event. Hence, depending on the issue being addressed, this impact may need to be included in a PRA if a detailed assessment of risk is required.

The following good practices are categorized under four major analysis activities for doing pre-initiator HRA. These analysis activities are:

1. Identifying activities that have the potential to result in pre-initiator human failures
2. Screening out the activities for which human failure events (HFEs) do not need to be modeled
3. Modeling specific HFEs corresponding to the unscreened activities
4. Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs.

4.1 Identifying potential pre-initiator HFEs

4.1.1 OBJECTIVE: To identify from routine plant actions, those pre-initiator human actions whose failure to perform correctly could result in the human-induced unavailability of PRA-modeled equipment that is credited (i.e., modeled, including the likelihood of either success or failure) in the PRA accident sequences. This is important because these actions represent other potential modes of unavailability of the credited equipment (besides the equipment simply failing to start or other failure modes in the PRA) that contribute to overall plant risk. Note that not all of the identified actions will be modeled since some may be screened from further analysis in the following analysis activity (screening). The following provides good practices for identifying potential pre-initiator human failures while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

4.1.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard calls for a systematic process to be used to identify routine activities that if not completed correctly, may impact the availability of equipment addressed in the PRA. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-A that address the need to consider test and maintenance activities, calibration activities, and actions that could affect multiple equipment. Regulatory Guide 1.200 states the NRC staff have no objections to the ASME Standard requirements covering this activity.

4.1.3 GOOD PRACTICES:

4.1.3.1 Good Practice #1: Review Pre-Initiator Procedures, Actions, and Equipment

The main goal of this good practice is to identify pre-initiator human actions with the potential to leave important equipment unavailable. Since the analysis of the actions will be an iterative process throughout the PRA, at this stage the HRA process should include reviews of the items listed below to the extent that is necessary to determine which actions should be included through the screening phase and to obtain the information necessary to apply the screening rules (Section 4.2). The sources of information can be returned to later if needed to support further analysis. The review should cover the following:

- All routine (scheduled) test and maintenance as well as calibration procedures that affect equipment to be credited in the PRA (for core damage frequency (CDF) and LERF) should be identified and reviewed.

- Actions specified in the above procedures that realign equipment outside their normal operation or standby status, or otherwise could detrimentally affect the functionality of credited equipment if not performed correctly (e.g., miscalibration) should be identified.

- "Affected" equipment that is routinely acted on and credited in the PRA. The equipment reviews should include:

  - the primary systems, structures, and components (SSCs) (e.g., emergency core cooling systems' components, containment cooling systems' components),

  - support systems (e.g., power, air, cooling water),

  - cascading effects among the equipment (e.g., if the realignment of an equipment item in one procedure such as an air-operated valve would implicitly require the subsequent realignment of another equipment item such as isolation of an air line that would then disable a portion of the air system), and

  - instrumentation (e.g., indicators, alarms, sensors, logic devices) and controls (e.g., hand switches) that (a) affect automatic operation of the above primary and support system equipment and/or (b) is the sole instrumentation relied upon (as opposed to multiple, redundant items) to credit post-initiator human actions to be included in the model (e.g., a single subcooling indication relied upon to meet an emergency core cooling termination criteria which if miscalibrated could induce failure of the appropriate post-initiator operator action).

4.1.3.2 Good Practice #2: Do Not Ignore Pre-Initiators

The identification process should identify pre-initiator human actions even if they may be potentially covered by the affected equipment failure data (see section 4.1.4 for additional information).

4.1.3.3 Good Practice #3: Examine Other Operational Modes and Routine Actions

If applicable and credited in the analysis, the identification process should address other operational modes and routine actions affecting barriers and other structures such as fire doors, block walls, drains, seismic restraints, etc.

4.1.3.4 Good Practice #4: Identify Actions Affecting Redundant and Multiple Diverse Equipment

The identification process needs to include possible pre-initiator actions *at least within each system* where redundant or multiple diverse equipment can be affected by (a) a single act (e.g., misalignment of a valve affecting multiple system trains or even multiple systems) or (b) through a common failure with similar multiple acts (e.g., mis-calibrating multiple sensors due to incorrect implementation of the same calibration procedure or use of the same mis-calibrated standard). For the latter case, the analyst should not duplicate that already covered under the common cause failure modeling of the equipment, but should include consideration of at least the following possible commonalities in deciding whether to include a particular action:

- same crew, same shift performing the actions (common "who" mechanism),
- common incorrect calibration source (common "what" mechanism),
- common incorrect tool, process, or procedure/training, or inadequate material (e.g., wrong grease) (common "what/how" mechanisms),
- proximity in time and/or space/location of similar multiple acts (common "when/where" mechanisms), and
- common cues (e.g., same indications, labels, alarms, procedures steps) (common "why" mechanism).

The more these commonalities exist concurrently with respect to a particular action being considered for initial inclusion in the model, the more the identification process should consider the act as a potentially important pre-initiator action to be included.

4.1.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to perform the above good practices could lead to an incomplete list of routine human activities that if not completed correctly, may impact the availability of equipment addressed in the PRA. Therefore, this could result in potentially missing (i.e., not modeling) a risk-significant pre-initiator human action. The following related observations should be considered.

- Missing or unnecessarily including a human action is often not a serious mistake (i.e., would not significantly affect the overall risk) unless the human action (a) can affect multiple equipment items, or (b) can affect a single equipment item with a high operating reliability. This is because with common nuclear plant practices and designs, typically those human

actions that could affect multiple trains of equipment tend to be the more significant pre-initiator human failures. Those affecting just one equipment item are usually not important unless the equipment item has a high operating reliability (e.g., failure to start or run is in the 0.0001 or lower probability range) and so the pre-initiator failure probability from failing the human action could be a significant contributor to the unavailability of the equipment.

- The possible human action failures associated with routine test and maintenance or calibration procedures should be included when they could affect critical instrumentation, diagnostic devices, or specific items like pushbuttons, etc. that have no redundancy or diverse means of function. While typically such situations do not exist in nuclear power plants, changes to the plant could conceivably and unintentionally create such a situation. Affecting the operator's ability to take the desired action is similar, functionally, to affecting the equipment item itself which is to be activated. Hence, the analysis approach should ensure that such situations, from a possible pre-initiator perspective, do not exist or if they do, they are addressed.

- In practice it is best to include pre-initiator human actions even if the associated failure already may be included in the failure data for the affected equipment item (e.g., in the failure-to-start data). This is because it is often hard to determine if the failure data bases include such human failures since data bases are typically insufficiently documented to know if the potential pre-initiator failure is already included. Generally, unless the failure can affect multiple equipment items, either missing the failure or double-counting the failure have small effects on the outcome of the PRA. Potential double-counting is the most conservative thing to do, and yet typically not a serious over-estimation of the failure's significance. In addition, including all identified pre-initiators gives analysts the opportunity to identify potentially problematic actions such as those with procedural or training problems, those that do not require appropriate checks, etc.

- If applicable, the possible failures associated with routine test and maintenance or calibration procedures should be included when they could affect equipment critical to external events such as fire barriers (e.g., opening a fire door and failing to restore it to a closed position), seismic restraints, floor drains and barriers, wind barriers, etc. While typically such situations do not exist in nuclear power plants since such equipment items often do not have routine test, maintenance, or calibration activities that would adversely affect their function, changes to the plant or plant practices, for instance, could conceivably and unintentionally create such a situation. To the extent the analysis assumes the functionality of these normally highly reliable devices, pre-initiator failures that could affect these devices could be potentially important. Hence, the analysis approach should ensure that such situations, from a possible pre-initiator perspective, do not exist or if they do, they are addressed.

- Considering the potential importance of acts that affect multiple equipment items, the identification process should search for acts that affect multiple equipment items *at least within a system* (e.g., auxiliary feedwater system, reactor core injection system) as this represents the current state of the art in PRA. A search across multiple systems (e.g., auxiliary feedwater and high pressure injection) is an expansion of the current state of the art and should not be expected except for those cases where the same instrumentation or equipment (e.g., pressure signals, same tank level equipment) activates or affects multiple systems.

4.2 Screening those activities for which HFEs do not need to be modeled

4.2.1 OBJECTIVE: To screen out those activities for which associated failures do not need to be analyzed because they should be probabilistically unimportant. The screening process, though largely qualitative, is based on the belief that certain design or operational practices make some pre-initiator failures sufficiently unlikely that they will not be risk significant failures and therefore do not need to be modeled. The following provides good practices for screening out pre-initiator human actions and associated human failures while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

4.2.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard addresses allowable screening of activities based on practices that limit the likelihood of errors in those activities. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-B that address screening rules or criteria, as well as the requirement to not screen actions that could affect multiple equipment items. Regulatory Guide 1.200 states the NRC staff have no objections to the ASME Standard requirements covering this activity.

4.2.3 GOOD PRACTICES:

4.2.3.1 Good Practice #1: Screen Pre-Initiators with Acceptable Restoration Mechanisms or Aids

A candidate pre-initiator action can be screened out (i.e., not to be modeled) if the nature of the associated action meets any one of the following criteria and the reason for screening is documented (see exception under Good Practice #2 below):

- the affected equipment will receive an automatic realignment signal and it can respond if demanded (i.e., the equipment will not have been disabled by the human actions), or
- there is a valid post-maintenance test/functional check (a test or functional check that has been shown to work consistently) after the original manipulation which will reveal misalignment or incorrect status (e.g., faulty position, improper calibration), or
- following the original action(s), there is an independent second verification of equipment status that uses a written checklist that will verify incorrect status, or
- there is a valid check (one that has been shown to work consistently), at least once per shift, of equipment status that will reveal misalignment or incorrect status, or
- there is a compelling signal (e.g., annunciator or indication) of improper equipment status or inoperability in the control room, it is checked at least shiftly or daily, and realignment can be easily accomplished, or
- other criteria as long as it can be demonstrated, using an acceptable model such as THERP[8] or ASEP[13] that the resulting human error probabilities would be low compared with the failure

probabilities (e.g., failure to open) of the equipment.

4.2.3.2 Good Practice #2: Do Not Screen Actions Affecting Redundant or Multiple Diverse Equipment

Do not screen those actions and possible pre-initiator failures that simultaneously affect multiple (redundant or diverse) equipment items (see Good Practice #4 under Section 4.1.3).

4.2.3.3 Good Practice #3: Re-Evaluate the Screening Process for Special Applications

For a specific PRA application and depending on the issue being addressed (e.g., examination of a specific procedure change), revisit the original PRA screening process to ensure issue-relevant human actions have not been deleted from the PRA prior to its use to assess the new issue.

4.2.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to perform the above good practices could lead to inappropriate screening out of human actions and therefore not including the actions in the PRA. This could result in potentially missing a risk-significant pre-initiator human action. The following related observations should be considered.

- Generally, screening out pre-initiator human failures (i.e., don't have to be modeled) is acceptable based on experience with past PRAs and the types of pre-initiator failures that are typically found to be unimportant. This is done to simplify the model and not expend resources addressing unimportant pre-initiator human actions. It should be clear that an appropriate level of investigation has been performed to ensure the above criteria have been met and if these or other criteria are used, their justification is documented for outside review. It is advisable to keep a record of all screened actions for later reference when performing specific applications (see Good Practice #3). When in doubt, it is recommended the pre-initiator action not be screened out, but the corresponding human failure modeled in the PRA for further analysis.

- Since pre-initiator human actions and related failures affecting multiple equipment items can sometimes be risk important, none of these should be screened out but should be modeled and examined in more detail in the PRA because of the potential consequences of the failure.

- There can be a tendency to use an existing PRA model to address issues such as changes to the plant, without spending the appropriate time to revisit some of the underlying assumptions and modeling choices made to create the original PRA. However, such a review should be done to see if these assumptions and choices still apply for the issue being addressed. Some pre-initiator human failures may not have been included in the original PRA (i.e., screened out) that in light of the new issue being addressed, should now be included in the model (i.e., could be important for addressing the issue). Hence, it is good practice to implement a process that determines whether some of the formerly screened out pre-initiator human failures should be added back in to the model in order to appropriately address the issue.

4.3 Modeling specific HFEs corresponding to the unscreened human actions

4.3.1 OBJECTIVE: To define how the specific pre-initiator HFE is to be modeled in the PRA to accurately represent the failure of each action identified and not screened out from the above analysis activities. The HFE needs to be linked to the affected equipment (single or multiple) and needs to appropriately define the failure mode of that equipment that makes the equipment unavailable. The following provides good practices for modeling pre-initiator human failure events while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

4.3.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard calls for the modeling of pre-initiator HFEs based on the impact of the mode of unavailability of the affected equipment item(s) in the PRA. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-C that address the modeling level of detail for each HFE and the modes of failure (as unavailabilities) to be considered. Regulatory Guide 1.200 states the NRC staff have no objections to the ASME Standard requirements covering this activity.

4.3.3 GOOD PRACTICES:

4.3.3.1 Good Practice #1: Include HFEs for the Unscreened Human Actions in the PRA Model

Define each specific pre-initiator HFE to be modeled in the PRA as a basic event that describes the human-induced failure mode and is located in the model such that it is linked to the unavailability of the affected component, train, system, or overall function (i.e., level of modeling) depending on the effect(s) of the HFE (e.g., a single valve will not close, a train will be inappropriately isolated, the automatic start signal for an entire system will be disabled). The following considerations, as a minimum, should be taken into account when defining the pre-initiator failure level properly in the PRA:

- whether the nature of the manipulation affects a whole train, system, etc., making it logical to define the HFE at the level of a whole train, system, etc,

- whether multiple individual acts affecting multiple pieces of equipment (e.g., different components) can be combined as a single pre-initiator HFE affecting a higher level of equipment resolution (e.g., the train containing the different components). This can be done as long as:

  - the acts and effects are related,

  - the factors affecting the quantification of the single HFE will not be significantly different than those that would have been relevant for the individual acts (e.g., the same performance shaping factors will be relevant [as is discussed later]) or the quantification result will be conservatively bounding compared to modeling and quantifying the individual acts separately, and

- there are no potential commonalities/dependencies with other pre-initiator acts elsewhere in the model so that potential common failures among similar individual acts might be missed (e.g., miscalibration of multiple signal channels), and

- the level of detail already modeled in the PRA (e.g., train, system) for failures of the associated equipment (but note that this is a less important factor than those above).

The failure modes (fail to close, fail to start, etc.) or modes of unavailability (valve left in wrong position, etc.) should be a direct result of considering the equipment affected and the effects of the human-induced failure (refer to all the Good Practices under Section 4.1.3), and stem from failure to restore equipment and/or otherwise correct the adverse effect (such as miscalibration) so that the equipment is again operable. The failure modes should clearly describe the HFE effect to ensure proper interpretation of the HFE in the model (e.g., only two of three redundant sensors need to be disabled to make the actuation signal unavailable, and not all three sensors have to be disabled).

As an aid to ensure appropriate modeling, it is a recommended practice (but not necessary) that the pre-initiator failure be placed in proximity, in the PRA model, to the equipment affected by the human failure. In this way, a quick comparison can be made between the equipment failure and the pre-initiator human failure to ensure they are consistent.

4.3.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to either (a) properly link the pre-initiator human failure to the affected equipment item(s), or (b) properly model the effect by the appropriate failure mode of the equipment item(s) will likely lead to misrepresentation of the unavailability of the equipment caused by a pre-initiator human failure in the PRA. Depending on the specific misrepresentation, the risk effect of the human failure could be over-emphasized (such as if the human failure is modeled as affecting more equipment items than it actually does or it is modeled as causing a failure mode that it does not cause), under-estimated (such as if the human failure is modeled as affecting less equipment items than it actually does or a failure mode that is caused by the human failure is missed in the model), or even missed entirely. This can result in inaccuracies in the PRA results and particularly incorrect assessments of the importance of pre-initiator human failures. The precise definition of the pre-initiator basic events and their placement in the model (from both a logic and failure mode standpoint) ultimately defines how the model addresses the effects of the human failures. This needs to be done accurately if the model is going to logically represent the real effects of each human failure and if the corresponding HFE is going to be correctly quantified (as discussed later).

4.4 Quantifying the corresponding HEPs for the specific HFEs

4.4.1 OBJECTIVE: To address how the human error probabilities (HEPs) for the modeled HFEs from the previous analysis activity are to be quantified. This section provides good practices guidance on an attribute or criteria level and does not endorse a specific tool or technique (although THERP[3] or its ASEP[4] simplification are among those often used). Ultimately, it is these probabilities along with the other equipment failure and post-initiator human error

probabilities as well as initiating event frequencies that are all combined to determine such risk metrics as CDF, LERF, ΔCDF, ΔLERF, etc. as addressed in Regulatory Guide 1.174[3]. The following provides good practices for quantifying pre-initiator human failure events while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

4.4.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard calls for a systematic process for assessing the pre-initiator HEPs that addresses plant-specific and activity-specific influences. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-D that address many factors associated with quantifying the HEPs. These include when screening vs. detailed estimates are appropriate, performance-shaping factors (PSFs) considered in the evaluations, treatment of recovery, consideration of dependencies among HFEs, uncertainty, and reasonableness of the HRA results. Regulatory Guide 1.200 states the NRC staff have no objections to the ASME Standard requirements covering this activity.

4.4.3 GOOD PRACTICES:

4.4.3.1 Good Practice #1: Use Screening Values During the Initial Quantification of the HFEs

The use of screening-level human error probability (HEP) estimates is usually desirable during the early stages of PRA development and quantification. This is acceptable (and almost necessary since not all the potential dependencies among human events can be anticipated) provided (a) it is clear that the individual values used are over-estimations of the probabilities that would be developed if detailed assessments were to be performed AND (b) dependencies among multiple human failure events appearing in an accident sequence are conservatively accounted for. These screening values should be set so as to make the PRA quantification process more efficient (by not having to perform detailed analysis on every HFE), but not so low that later detailed analysis would actually result in higher HEPs. The screening estimates should consider both the individual events and the potential for dependencies across multiple HFEs in a given accident sequence (scenario). To meet these conditions, it is recommended that (unless a more detailed assessment is performed of the individual or combination events to justify lower values):

- no individual pre-initiator HEP screening value should be lower than 0.01 (typical of highest pre-initiator values in PRAs), and

- multiple HFEs in the same sequence should not have a a joint probability value lower than 0.005 (accounts for a 0.5 high dependency factor) at this stage.

4.4.3.2 Good Practice #2: Perform Detailed Assessments of Significant HFEs

As needed for the issue being addressed to produce a more realistic assessment of risk, detailed assessments (not just screening estimates) should be performed for at least the significant HFE contributors (see Regulatory Guide 1.200[4], Table A-1 for a definition of the term "significant contributor"). The PRA analyst can define the significant contributors by using typical PRA criteria (not addressed here, but see NUREG-1764,[12] Section 2.3) such as importance measure thresholds as

well as other qualitative and quantitative considerations. While using screening-level values (supposedly purposely conservative) may, at first, seem to be a "safe" analysis process, it can have negative impacts. Screening values can focus the risk on inappropriate human actions or related accident sequences and equipment failures because of the intentionally high HEPs. Such incorrect conclusions need to be avoided by ensuring that a sufficient set of more realistic, detailed HEPs are included in the model.

4.4.3.3 Good Practice #3: Revisit the Use of Screening Values Versus Detailed Assessments for Special Applications

For a specific PRA application and depending on the issue being addressed (e.g., examination of a specific procedure change), revisit the use of screening vs. detail-assessed HEPs to ensure issue-relevant human actions have not been prematurely deleted from the PRA or there is an inappropriate use of screening vs. detailed values to properly assess the issue and the corresponding risk.

4.4.3.4 Good Practice #4: Account for Plant and Activity Specific Performance Shaping Factors (PSFs) in the Detailed Assessments

HEP assessments should account for the most relevant plant-specific and activity-specific PSFs in the analysis of each pre-initiator HFE. There is not one consensus list of appropriate contextual factors (e.g., plant conditions, PSFs, activity characteristics, etc.) to be considered in the evaluation of the pre-initiator HEPs. Additionally, for a specific action, which factors are most relevant may be different (e.g., perhaps one act is time-sensitive because it is done in a high radiation area while another is most affected by the complexity of steps with many opportunities to make undetected mistakes). It should be qualitatively apparent that the factors seemingly most relevant to the act (based on an understanding of the act [as might be derived from a sound task analysis]) have been considered in the corresponding HEP estimate.

Factors that are typically important to address because they tend to be variable and not always optimal based on typical nuclear plant practices, include:

- whether the activity relies on the use written work plans, job briefs, and related procedures (positive influences tending to lower the HEP) rather than verbal guidance and/or memory (more negative influences tending to raise the HEP), as well as the quality of the information (e.g., look for ambiguities, incompleteness, inconsistencies, etc. that are negative influences and thus tend to raise the HEP),

- whether the activity is complex (e.g., involves multiple and/or repetitive steps that are hard to track, requires coordination of multiple personnel, does not use appropriately human-factored checklists, involves several variables, requires that calculations be performed, requires sensitive adjustments), and

- what ergonomic issues (e.g., layout, available information [instruments, alarms, computer readouts, etc.], labeling, readability, physical demands) are relevant.

Note that information about such factors can be greatly enhanced by performing talk-throughs and

walkdowns of the actions and whenever possible, observing actual crews perform the actions (or at least sample actions).

Other factors that tend to be less important either because of typical nuclear plant practices or because the factors are typically less relevant include:

- skill level/experience/training of the crew (which is typically adequate in nuclear plants for the jobs each crew member is to perform),
- stress level (which is not usually relevant in pre-initiator failures unless special situations such as potential personal harm, the need for fast sequential responses, etc. play a role),
- environmental factors such as temperature, humidity, radiation, noise, lighting, etc., which are typically sufficiently benign (with the exception of special circumstances such as a high radiation environment that can lead to the desire to hurry the actions), and
- availability of time (which is not usually a strong factor in pre-initiator failures).

Nevertheless, the evaluation should ensure that the typical practice or "irrelevancy" is not compromised.

If the large majority of these factors affect the human performance negatively or if even just one or two have a strong negative influence, the HEP will tend to be higher (e.g., 0.01 to 0.1 or even higher, not accounting for recovery addressed under Good Practice #5 below). Conversely, mostly positive influences should yield lower HEPs (e.g., 0.001, with additional recovery factors still to be applied as addressed under Good Practice #5 below).

4.4.3.5 Good Practice #5: Apply Plant Specific Recovery Factors

Each HFE analyzed should be investigated for opportunities to recover the initial failure. Multiple recoveries may be acceptable, where appropriate. However, any dependencies among the initial failure and possible recovery actions, and among the recovery actions themselves, must be considered (see Good Practice #6 below). Typical pre-initiator recovery actions[1] or considerations include:

- post-maintenance or post-calibration tests that are required and performed by procedure,
- independent verification that uses a written check-off list which verifies component status following maintenance/testing/calibration and its practice has been verified by walk-throughs

[1] Note that the definition of a recovery action and its distinction from a repair action has been adopted from Regulatory Guide 1.200.[4] *Recovery action* is defined as: a PRA modeling term representing restoration of the function caused by a failed system, structure, or component (SSC), by bypassing the failure. Such a recovery can be modeled using HRA techniques regardless of the cause of the failure. *Repair* is defined as: a general term describing restoration of a failed SSC by correcting the failure and returning the failed SSC to operability. HRA techniques cannot be used since the method of repair is not known without knowing the specific causes.

and examination of plant experience,

- a separate check of component status that is made by the original performer at a later time and that uses a written check-off list,

- work shift or daily checks that are performed for component status and uses a written check-off list,

- compelling feedback (e.g., alarm) that supports detection and quick recovery of the original failure, or

- combinations of the above.

The more of these considerations that are applicable for a given pre-initiator HFE, the more the situation tends to increase the recovery potential (i.e., decrease the HEP). To the extent they are independent, each recovery can result in a multiplier (e.g., 0.1) on the original HEP estimate thereby reducing its overall value. However, the analyst should know that there is a logical limit to such reductions. For example, credit for two checks should not be given if there are dependencies (e.g., same or similar job aids and cues) between the checks done by the original performer and the independent verifier.

Basic HEPs for pre-initiator HFEs for nuclear plant applications (including recovery) are typically expected in the 0.01 (among the highest) to 0.0001 range. Any values below the 0.0001 to 0.00001 range should be considered suspect unless justified.

4.4.3.6 Good Practice #6: Account for Dependencies Among the HEPs in an Accident Sequence

Dependencies among the pre-initiator HFEs, and hence the corresponding HEPs in an accident sequence, should be quantitatively accounted for in the PRA model. This is particularly important so that combined probabilities are not inadvertently too optimistic, resulting in the inappropriate decrease in the risk significance of human actions and related accident sequences and equipment failures. In the extreme, this could result in the inappropriate screening out of accident sequences from the model because the combined probability of occurrence of the events making up an accident sequence drops below a threshold value used in the PRA to drop sequences from the final risk results.

To address these dependencies, usually a level or degree of dependence among the HFEs in an accident sequence is determined, at first qualitatively (e.g., low, high, complete), and then combined HEPs are assessed accordingly. Once the first HEP has been estimated, subsequent quantitative factors for dependent human failures or recoveries of the original failure are typically expected to be:

- 0.01 to 0.1 for low dependence
- 0.1 to 0.5 for high dependence
- >0.5 for very high or 1.0 for complete dependence

Note that specific tools/techniques may use somewhat different probabilities than provided here based on specific considerations.

In establishing the level of dependence, Good Practice #4 under Section 4.1.3 addresses typical commonalities that tend to make HEPs more dependent. If, for example, an HFE is not independent of another HFE, then once the first human failure occurs, there is an increased likelihood that a similar second or third, and so on, human failure will also occur. For example, failing to restore the lineup of one train of equipment after a test may increase the likelihood of failing to restore the second train of equipment after a similar test. Good Practice #5 just above addresses recovery characteristics that tend to break-up these commonalities because they "recover" any initial error, making the individual HFEs more independent. The more the types of commonalities addressed under Good Practice #4 under Section 4.1.3 exist and the less corresponding recoveries under Good Practice #5 above exist, the higher should be the assessed level of dependence among the HFEs. To the extent the converse is true, low or even no dependence should be assessed.

4.4.3.7 Good Practice #7: Assess the Uncertainty in Mean HEP Values

Point estimates should be mean values for each HEP (excluding screening HEPs) and an assessment of the uncertainty in the mean values should be performed at least for the significant HEPs to the extent that these uncertainties need to be understood and addressed in order to make appropriate risk-related decisions. Assessments of uncertainty are typically performed by:

- assigning uncertainty distributions for the HEPs and propagating them through the quantitative analysis of the entire PRA, such as by a Monte Carlo technique, and/or

- performing sensitivity analyses that demonstrate the effects on the risk results for extreme estimates in the HEPs based on at least the expected uncertainty range about the mean value.

Note that, in some cases, it may be sufficient to address the uncertainties by just qualitative arguments without the need to specifically quantify them (e.g., justifying why the HEP cannot be very uncertain or why a change in the HEP has little relevancy to the risk-related decision to be made).

In assessing the uncertainties, and particularly when assigning specific uncertainty distributions, the uncertainties should include (a) those epistemic uncertainties existing because of lack of knowledge of the true expected performance of the human for a given context and associated set of PSFs, and (b) consideration of the combined effect of the relevant aleatory (i.e., random) factors to the extent they are not specifically modeled in the PRA and to the extent that they could alter the context and PSFs for the HFE. For pre-initiator HFEs, there should be few or no aleatory factors worthy of consideration, since typically the procedure used, the environment experienced, etc. do not randomly change (at least significantly). But, for example, if different and significant crew experience levels are known to exist, it is random as to which crew will perform the pre-initiator act at any given time. In such a case, the mean should represent the average crew experience level and the uncertainty should reflect the uncertainty in the mean due to crew experience and any other relevant factors. Again, aleatory factors are typically not very relevant to pre-initiator HEPs and so typically are not important to address.

Whatever uncertainty distributions are used, the shape of the distributions (e.g., log-normal, beta, etc.) are typically unimportant to the overall risk results (i.e., the results are usually not sensitive to specific distributions). Further, typical uncertainties include values for the HEP that represent a factor of 10 to 100 between the lower bound value and the upper bound value that encompass the mean value.

However, it should be noted that some distributions, e.g., log-normal, can give probabilities greater than 1.0 for HEPs that are relatively high.

4.4.3.8 Good Practice #8: Evaluate the Reasonableness of the HEPs Obtained Using Detailed Assessments

The pre-initiator HEPs (excluding the screening HEPs) should be reasonable from two standpoints:

- first and foremost, relative to each other (i.e., the probabilistic ranking of the failures when compared one to another), and

- in absolute terms (i.e., each HEP value), given the relative strengths of the positive and negative PSFs identified as being important and the presence or absence of recovery factors.

Such reasonableness should be checked based on consideration of actual plant experience and history, against other evaluations (such as for similar acts at other plants), and the qualitative understanding of the actions and the relevant contexts and PSFs under which the acts are performed. It is suggested that a rank-ordered list of the pre-initiator HFEs by probability be used as an aid for checking reasonableness. For example, simple, procedure-guided, independently checked actions should have lower HEPs than complex, memorized, not checked actions, all other factors being the same. Similarly, with respect to the reasonableness of the absolute values, HFEs with many positive PSFs or several strong recovery factors could have values in the 0.0001 to 0.00001 range, while HFEs with negative PSFs dominating and no independent checks could be in the 0.1 to 0.01 range. Typical expectations of pre-initiator HEPs can be wide-spread (~0.01 to approximately 0.00001) and depend particularly on the relevant contextual factors, applicable recoveries, and proper consideration of dependencies as discussed under many of the Good Practices covered above.

4.4.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to quantify the pre-initiator HEPs as realistically as possible using the good practices articulated above (except for where higher screening estimates are used, purposely and appropriately), could result in improper HEPs and thus inaccuracies in the PRA results and particularly incorrect assessments of the importance of pre-initiator human failures. The risk effect of the human failure could be over-emphasized (such as if the human failure is estimated with too high [pessimistic] a HEP or a high screening estimate is used where a more realistic detailed estimate is appropriate), or under-estimated (such as if the human failure is estimated with too low [optimistic] a HEP or a dependency among failures is not accounted for resulting in too low a joint probability for multiple human failures). Besides these concerns about inaccuracies in the HEP quantification and thus whether the HEPs "make sense", as well as the resulting potential misinformation about the significant risk contributors if quantification is not done well, the following related observations are noted.

- Screening is a useful and most often, necessary part of HRA so as to avoid the expenditure of resources on unimportant human events and accident sequences. The above guidance is aimed at allowing a level of useful screening without inadvertently and inappropriately allowing the analytical phenomenon of, for instance, multiplying three human events in the same sequence each at a screening value of 0.01 to yield a 0.000001 combined probability, without checking

for dependencies among the human events. In such a case some human failure events and combinations of events, or even whole accident sequences, may inappropriately screen out of the PRA model entirely because the accident sequence frequency drops below a model threshold. Hence some of the significant individual or combination contributors may be missed. This is why the estimated HEP values both individually and for combined events should not be too low during the screening stage. Further, if these estimated screening values are left permanently assigned to some human failure events that should be assessed with more detail to obtain a more realistic assessment of risk (supposedly lowering the probability), the risk significance of these human failure events and related equipment failures are likely to be over-emphasized at the expense of improperly lessening the relative importance of other events and failures.

- It is important to be sure that dependencies among the various modeled HFEs including the associated recoveries, have been investigated (e.g., the same person as the originator of the action performing the recovery may be more prone to fail to detect the original failure than an independent checker). Treating HFEs and any corresponding recoveries as independent acts without checking for dependencies (thereby being able to multiply the individual HEPs) can inappropriately lessen the risk significance of those HFEs and related equipment failures in accident sequences. This can cause the inappropriate screening out of accident sequences because the sequences quantitatively drop below a model threshold value as discussed above under screening. Proper consideration of the dependencies among the human actions in the model is necessary to reach the best possible evaluation of both the relative and absolute importance of the human events and related accident sequence equipment failures.

- The use of mean values and addressing uncertainties are a part of the Regulatory Guide 1.174[3] guidance and to the extent addressed therein, the HRA quantification needs to be consistent with that guidance when making risk-informed decisions. The estimates should reflect, to the extent possible, the as-built and as-operated conditions as addressed in the plant- and activity-specific PSFs.

- There can be a tendency to use an existing PRA model to address issues such as changes to the plant, without spending the appropriate time to revisit some of the underlying assumptions and modeling choices made to create the original PRA. However, such a review should be done to see if these assumptions and choices still apply for the issue being addressed. For instance, some pre-initiator human failure events may be quantified in the original model using a set of screening estimates and detailed failure probabilities that may not be appropriate for the new issue being addressed. As an example, where high estimates as screening HEPs may have been acceptable for purposes of the original PRA, these supposedly conservative values may over-estimate the contribution of these human failure events for the issue being addressed. Further, the relative risk contribution of equipment and associated accident sequences with which the human failure events appear, may be artificially too high (and therefore other events too low) because of the screening values. Hence, it is good practice to revisit the use of screening and detailed human failure event probabilities in order to appropriately address the issue.

5. POST-INITIATOR HRA

The NRC staff has stated their positions on the ASME Standard[5] in Appendix A of Regulatory Guide 1.200[4]. The standard separates its requirements into two broad classifications; those that address the modeling of failures of pre-initiator human actions and those that address the modeling of failures of post-initiator human actions. This section provides good practices for implementing the requirements for addressing post-initiator human failure events (HFEs) in a PRA.

Post-initiator human failure events are events that represent the impact of human failures committed during actions performed in response to the initiation of an accident sequence (e.g., while following post-trip procedures or performing other recovery actions). They are important to model because humans can have a direct influence on the mitigation or exacerbation of undesired plant conditions after the initial plant upset. Hence, depending on the issue being addressed, this impact may need to be included in a PRA if a realistic assessment of risk is required.

The following good practices are categorized under four major analysis activities for doing post-initiator HRA. These analysis activities include:

1. Identifying potential post-initiator human failures
2. Modeling specific human failure events (HFEs) corresponding to the human actions
3. Quantifying the corresponding human error probabilities (HEPs) for the specific HFEs
4. Adding recovery actions to the PRA.

5.1 Identifying potential post-initiator human failures

5.1.1 OBJECTIVE: To identify the key human response actions that may need to be taken by the operators in response to a variety of possible accident sequences and that will therefore need to be modeled in the PRA. This is important since failures associated with these actions (e.g., failure to start standby liquid control, failure to initiate feed and bleed, failure to properly control steam generator feed flow, failure to align containment/suppression pool cooling) are represented in the PRA such that in combination with equipment failures, are expected to lead to core damage and/or large early releases. Such failures contribute to the overall risk and thus a systematic process needs to be followed to identify these response actions. The following provides good practices for identifying post-initiator human failures while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

5.1.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard calls for a systematic review to identify operator responses required for each of the accident sequences. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-E that address what to review as well as the types of actions to be included. Use of talk-throughs and simulator observations are also addressed as part of the supporting requirements (though this is best dealt with under the modeling and quantification analysis activities of the analysis and thus addressed more fully later in this report). Regulatory Guide 1.200 states the NRC staff have no objections to the ASME Standard requirements covering this activity.

5.1.3 GOOD PRACTICES:

5.1.3.1 Good Practice #1: Review Post-Initiator Related Procedures and Training Materials

Reviews of the following form the primary bases for identifying the post-initiator actions.

- Plant-specific emergency operating procedures (EOPs)
- Abnormal operating procedures (AOPs)
- Annunciator procedures
- System operating procedures.
- Severe accident management guidelines (SAMGs)
- Other special procedures (e.g., fire emergency procedures) as appropriate.

The review is done to identify ways operators (crews) are intended to interact with the plant equipment after an initiator. The ways they interact will be a function of the various conditions that can occur, as defined by the development of the PRA accident sequences and associated equipment unavailabilities and failure modes. Analysts should particularly note where operator actions that will directly influence the behavior of the system or affect critical functions are called out in these procedures and under what plant conditions and indications (cues) such actions are carried out. It will also be useful at this time to examine whether there are any potential accident conditions under which the procedures might not match the situation as well as would be desired, e.g., potentially ambiguous decision points or incorrect guidance provided under some conditions. Information about such potential vulnerabilities will be useful later during quantification and may help identify actions that need to be modeled.

While observation of simulator exercises and talk-throughs with crews about various accident scenarios are probably most important during the modeling phase, if time and resources allow, they may also be useful during the identification phase to help analysts understand the procedures and how they are implemented by the crews.

5.1.3.2 Good Practice #2: Review Functions and Associated Systems and Equipment to Be Modeled in the PRA

The PRA team's plant, system, and operations knowledge is important to the performance of this review and identification task. The process should accomplish the following:

- Identify the functions and associated systems and equipment to be modeled in the PRA.

- Identify whether the function is needed (e.g., injection) or undesired (e.g., stuck-open safety relief valve) for each scenario being addressed, recognizing that the need for a function may vary with different initiators and sequences.

- Identify the systems/equipment that can contribute to performing the function or cause the undesired condition (including structures and barriers such as fire door and floor drains, especially for external event analyses).

- Identify ways the equipment can functionally succeed (i.e., the success criteria) and fail.

Based on the above, the analysts together identify ways the operators are (a) intended/required to interact with the equipment credited to perform the functions modeled for the accident sequences modeled in the PRA and/or (b) to respond to equipment and failure modes that can cause undesired conditions per the PRA. During the identification process, it is helpful to use action words such as actuate, initiate, isolate, terminate, control, change, etc. so that the desired actions are clear. This process ensures that the logic models correctly model the impact of emergency operating procedures and other procedures, such as AOPs and annunciator response procedures, on the accident sequence development.

5.1.3.3 Good Practice #3: Look for Certain Expected Types of Actions

While the specific actions to be identified may be plant-specific, in general, the list below provides the types of actions that are expected to be identified. Note that actions that are "heroic" (e.g., they must enter an extreme high radiation environment in order to perform) or that are performed without any procedure guidance or are not trained on, should not be included or credited in the analysis. Exceptions may be justified, but this should not be normal practice.

The analysts should:

- Include necessary and desired/expected actions (e.g., initiate RHR, control vessel level, isolate a faulted steam generator, attempt to reclose a stuck-open relief valve).

- Include backup actions to failed automatic responses (e.g., manually start a diesel generator that should have auto started) but be sure the action can be credited to recover the auto failure mode.

- Include anticipated procedure-guided or skill-of-the-craft recovery actions (e.g., restore offsite power, align firewater backup) although these may best be defined later as the PRA quantification begins and important possible recovery actions become more apparent.

Consistent with present day state-of-the art, acts whose failure involve an error of omission (EOO) should be included when identifying post-initiator acts of concern. These involve failure to take the appropriate actions as called out in the procedures and/or trained on or expected as common practice. For example, failure to initiate feed and bleed or failure to start standby liquid control, are EOOs. Possible acts whose failure would involve an error of commission (EOC) have generally been beyond PRA practice, but some issues may require that the PRA/HRA address such failures. EOCs involve performing expected acts incorrectly or performing extraneous and detrimental acts such as shutting down safety injection when it is not appropriate. See Section 6 of this document for more information on the inclusion of EOCs and some related good practices.

Finally, it should be recognized that iterations as well as refinement and review of the PRA model construction may (and often do) provide additional opportunities to identify any potentially important human actions.

5.1.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to perform the above good practices could lead to an incomplete list of post-initiator human activities that if not completed correctly, may impact the outcome and frequencies of accident sequences addressed in the PRA. Therefore, this could result in potentially missing (i.e., not modeling) a risk-significant post-initiator human action that in turn, could make the model incomplete and/or inaccurate, potentially resulting in misinformation as to the risk significant plant features (including the important human actions).

Further, while not all the post-initiator actions will be important in the final assessment of risk, unlike the pre-initiator actions, it is difficult to predetermine (at this stage) a set of actions that do not have to be included as part of the identification process. Ways the operators interact with the plant equipment and affect the outcome of any accident sequence need to be assessed in order to determine their relative significance. Hence the good practices herein are aimed at ensuring potentially risk-significant post-initiator actions (based on the procedures as well as the ways the procedures are interpreted and carried out) are identified at this stage of the analysis.

5.2 Modeling specific HFEs corresponding to the human actions

5.2.1 OBJECTIVE: To define how each specific post-initiator HFE is to be modeled in the PRA to accurately represent the failure of each action identified. This involves (a) the modeling of the HFEs as human-induced unavailabilities of functions, systems, or components consistent with the level of detail in the PRA accident sequences and system models, (b) possible grouping of responses into one HFE, and (c) ensuring the modeling reflects certain plant-specific and accident sequence-specific considerations. The following provides good practices for modeling post-initiator human failure events while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

5.2.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard calls for the HFEs to be defined so that they represent the impact of not properly performing the required actions, consistent with the structure and level of detail of the accident sequences. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-F that address the modeling level of detail for each HFE and how to complete the definition of each HFE. Regulatory Guide 1.200 states the NRC staff have no objections to the ASME Standard requirements covering this activity.

5.2.3 GOOD PRACTICES:

5.2.3.1 Good Practice #1: Include HFEs for Needed Human Actions in the PRA Model

Define each specific post-initiator HFE to be modeled in the PRA as a basic event that describes the human failure of not properly performing the required actions and locate the HFE in the model such that it is linked to the unavailability of the affected component, train, system, or overall function depending on the effect(s) of the HFE (e.g., failure to manually depressurize using the safety relief valves, failure to manually scram, failure to align the backup train of service water). The following

considerations should be used to define the post-initiator failure level properly in the PRA:

- whether the action is performed on a train, system, or component level, making it logical to define the HFE at the level of a whole train, system, or component

- whether the consequences of the failure and what would be affected by the failure is at the component, train, system, multiple systems, or entire function level

- whether multiple individual acts/responses such as at a system or component level (e.g., starting high pressure injection and then subsequently opening a power-operated pressurizer relief valve) can be combined as a single post-initiator HFE affecting a higher level of equipment resolution such as at a system or a function level (e.g., initiating feed and bleed). This could be done as long as:

  - the acts and effects are related,

  - the factors affecting the quantification of the single HFE will not be significantly different than those that would have been relevant for the individual acts (e.g., the same performance shaping factors will be relevant) or the quantification result will be conservatively bounding compared to modeling and quantifying the individual acts separately, and

  - there are no potential commonalities/dependencies with other post-initiator acts elsewhere in the model so that potential common failures among similar individual acts might be missed (see the discussion presented below)

- the level of detail already modeled in the PRA (e.g., train, system) for failures of the associated equipment (but note that this is a less important factor than those above)

As an example of how human responses may be grouped and modeled as one or more HFEs, consider the case in a boiling water reactor (BWR) of a desired response to control reactivity in an anticipated transient without scram scenario. Failure to control reactivity could be defined as one HFE, or as several HFEs based on the subtasks involving inhibiting the automatic depressurization system, lowering reactor water level, and initiating the standby liquid control system.

For situations such as the above example, usually it is best to model separate HFEs if failure to perform the subtasks:

- have different effects,

- may individually be impacted by very different performance shaping factors (e.g., in-control room actions vs. local actions in a high steam environment area, a subtask performed early in the scenario vs. another subtask performed much later in the scenario), or

- involves an action that has a dependency with some other action to be modeled in the PRA (e.g., failure to trip two reactor coolant pumps followed by subsequent failure to trip the remaining reactor coolant pumps when conditions warrant).

An alternative is to model them all as one HFE and model the bounding consequence (such as the failure to control reactivity example cited above) as long as the most limiting performance shaping factors are used (e.g., the shortest time that any of the subtasks must be performed, the most complex of the subtasks, etc.) and any subtask dependencies with other HFEs are identified, treated in the model, and properly quantified.

The failure mode defined by the HFE should be consistent with both the human failures and the equipment affected by these human failures (refer to the Good Practices under Section 5.1.3). The failures should sufficiently describe the HFE and its effect to ensure proper interpretation of the HFE in the model (e.g., fail to initiate feed and bleed within 5 minutes of the reactor pressure achieving 2400 psig).

As an aid to ensure appropriate modeling, it is recommended practice that the post-initiator failure be placed in proximity, in the PRA model, to the component, train, system, or function affected by the human failure. In this way, a quick examination of the model can reveal the modeled effect of the human failure.

5.2.3.2 Good Practice #2: Define the HFEs Such that they are Plant and Accident Sequence-Specific

Each of the modeled post-initiator HFEs should be defined such that they are plant- and accident sequence-specific, and the basic events representing them are labeled uniquely.

In order for the act to occur, the operator must diagnose the need to take the act and then execute the act. While many performance shaping factors are used to quantify the probability for failing to perform the act correctly (as discussed later under quantification), all of which should be evaluated based on plant and accident sequence-specifics, the following requirements are particularly germane to a basic understanding of the HFE and should be met to complete the definition of each HFE:

- To the extent possible, the time by which the act needs to be performed (e.g., fail to initiate feed and bleed by 2 minutes after primary pressure reaches 2400 psig), and the time necessary to diagnose the need for and to perform the act should be based on plant and accident sequence-specific timing and nature of the complexity and/or subtasks involved in implementing the act. In other words, timing information should not be based on another plant's analysis or a general analysis for the "average" plant, since the number and nature of the specific manipulations could be different, the plant thermal hydraulic response could be different, the location for local actions may require different travel times, and some sequences require a fast response while others may require a much quicker response for the same act, etc.

- Similar to the above, the availability and timing of plant and accident sequence-specific cues (i.e., indications, alarms, visual observations, etc. and when they will be manifested) should be used as this timing information can differ plant-to-plant and sequences-to sequence and will affect the likelihood and timing of diagnosing the need for the action.

- Plant-specific procedure and training guidance should be used based on the reviews under the Good Practices in Section 5.1.3.

- Where the act is performed (e.g., in the control room, locally in the auxiliary building) should be noted.

5.2.3.3 Good Practice #3: Perform Talk-Throughs, Walkdowns, Field Observations, and Simulator Exercises (as necessary) to Support the Modeling of Specific HFEs

To fully understand the nature of the act(s) (e.g., who performs it, what is done, how long does it take, whether there are special tools needed, whether there are environmental issues or special physical needs, whether there is a preferred order of use of systems to perform a specific function, etc.) and help define the HFEs and their context, additional reviews, talk-throughs, walkdowns, field observations, and simulator exercises are performed (as discussed in Good Practice #2 under Section 3.1.3.2, with more about the benefits of these techniques presented in Appendix B). In addition, the results of these activities may add to the list of actions and/or help interpret how procedural actions should be defined based on how they are actually carried out. Analysts should:

- Review training material and where possible, perform talk-throughs or walkdowns of the actions with operations or training staff to ensure consistency with training policies and teachings and to identify likely operator response tendencies for various conditions that may not be evident in the procedures For example, operators may cite a reluctance to restart reactor coolant pumps in spite of the procedure direction based on their training and perceived adverse effects, or they may have a preference to use condensate as a BWR injection source before using lower pressure emergency core cooling system. These added "interpretations" of the procedures can help complete and/or clarify the identified actions and ensure that later modeling and quantification of the actions will reflect the "as-operated" plant.

- Observe simulator exercises of accident scenarios since these can provide valuable insights with regard to how the actions are actually carried out, by whom, and particularly how procedure steps are interpreted by plant crews, especially where the procedure is ambiguous or leaves room for flexibility in the crew response. For example, through simulation it may be observed that a "single action" in the procedure (e.g., align recirculation) is actually carried out by a series of numerous and sequential individual actions (e.g., involving the use of many handswitches in a certain sequence). Again these observed "interpretations" of the procedures can help complete and/or clarify the identified actions and ensure that later modeling and quantification of the actions will reflect the "as-operated" plant.

5.2.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to perform the above good practices could lead to improper modeling of the HFE and thus a misrepresentation of its effects on the plant equipment and sequence outcome. Depending on the degree of failure to follow the above good practices, the risk effect of failing to take the proper action could be inappropriately over-emphasized (such as improperly linking the action to more equipment than is actually affected or using too conservative thermal-hydraulic information for the relevant timing associated with the action), under-estimated (such as if combining many actions into too broad a single HFE without properly bounding the consequences of the failure or not accounting for some plant-specific and accident sequence-specific negative PSFs as confirmed by simulations), or even missed entirely (such as not reviewing all appropriate procedures). This can result in inaccuracies in

the PRA results and particularly incorrect assessments of the importance of post-initiator human failures. The precise definition of the post-initiator basic events and their placement in the model (from both a logic and failure mode standpoint) ultimately define how the model addresses the effects of the human failures. This needs to be done accurately if the model is going to logically represent the real effects of each human failure and if the corresponding HFE is going to be correctly quantified (as discussed later). This accuracy is best obtained if plant-specific and accident sequence-specific information is used. Nevertheless, the following observation is noted.

- Not using plant/accident sequence-specific thermal hydraulic information for timing may or may not be critical based on the relevancy and thus appropriateness of the non-specific (i.e., "general") timing information that is used. It is better to use plant and accident-specific information, though it is recognized that in some areas (e.g., containment response for LERF), from a practical standpoint, modified "general" information may be all that is readily available. Further, as long as the timing considerations used are reasonable and accurate to within the resolution of the HRA quantification tool to be used, differences between plant and accident-specific versus more "general" timing considerations may not be a significant issue. Analysts should ensure that if non-specific timing information is used, the timing is reasonable for the plant and accident sequence being analyzed.

5.3 Quantifying the corresponding HEPs for the specific post-initiator HFEs

5.3.1 OBJECTIVE: To address how the human error probabilities (HEPs) for the modeled HFEs from the previous analysis activity are to be quantified. This section provides good practices on an attribute or criteria level and does not endorse a specific tool or technique. Ultimately, it is these probabilities along with the other equipment failure and pre-initiator human error probabilities as well as initiating event frequencies that are all combined to determine such risk metrics as CDF, LERF, ΔCDF, ΔLERF, etc. as addressed in Regulatory Guide 1.174[3]. The following provides good practices for quantifying post-initiator HEPs while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

5.3.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard requires that a well-defined and self-consistent process be used to quantify the post-initiator HEPs. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-G that address many factors associated with quantifying the HEPs. These include when conservative vs. detailed estimates are appropriate, consideration of cognitive and execution failures, performance shaping factors considered in the evaluations, consideration of dependencies among HFEs, uncertainty, and reasonableness of the HRA results. Regulatory Guide 1.200 states the NRC staff has just one clarification to the ASME Standard requirements covering this activity and that is to ensure that the availability of staff resources to carry-out all the desired actions is considered. This has been addressed among the PSFs covered in the good practices.

5.3.3 GOOD PRACTICES:

5.3.3.1 Good Practice #1: Address Both Cognitive and Response Execution Failures

Whether using conservative or detailed estimation of the post-initiator HEPs, the evaluation should include both cognitive (i.e., "thinking") as well as execution failures. For example, incorrectly interpreting a cue or not seeing a cue and thus not performing the act can be one mode of failure. Or, the operator can intend to take the act based on the proper and recognized cues but still otherwise fail to take the act or perform it correctly. Both need to be part of the HEP evaluations. However, the qualitative HRA analysis may indicate that one of these failure modes predominates the other in such a way that the effect of only one failure mode needs to be quantified, but this should be justified.

5.3.3.2 Good Practice #2: Use Screening Values During the Initial Quantification of the Post-Initiator HFEs

The use of conservative human error probability (HEP) estimates (screening values) is usually desirable during the early stages of PRA development and quantification. This is acceptable (and almost necessary since not all the potential dependencies among human events can be anticipated) provided (a) it is clear that the individual values used are over-estimations of the probabilities that would be developed if detailed assessments were to be performed AND (b) dependencies among multiple human failure events appearing in an accident sequence are conservatively accounted for. These screening values should be set so as to make the PRA quantification process more efficient (by not having to perform detailed analysis on every HFE), but not so low that later detailed analysis would actually result in higher HEPs. The screening estimates should consider both the individual events and the potential for dependencies across multiple HFEs in a given accident sequence (scenario). To meet these conditions, it is recommended that:

- no conservative HEP value assigned to an individual post-initiator HFE should be lower than the worse case anticipated detailed value and generally not lower than 0.1 (which is typical of high post-initiator values in PRAs, but 0.5 is also often appropriate and sometimes used), and

- multiple HFEs in the same sequence should not have a joint probability value lower than the worse case anticipated detailed joint probability value and generally not lower than 0.05 (accounts for a 0.5 high dependency factor) at this stage.

5.3.3.3 Good Practice #3: Perform Detailed Assessments of Significant Pots-Initiator HFEs

As needed for the issue being addressed to produce a realistic assessment of risk, detailed assessments of the significant HFE contributors should be performed (see Regulatory Guide 1.200[4], Table A-1 for a definition of the term "significant contributor"). The PRA analyst can define the significant contributors by use of typical PRA criteria (not addressed here, but see NUREG-1764,[12] Section 2.3) such as importance measure thresholds as well as other qualitative and quantitative considerations. Conservative estimates, or screening values should not be used in place of detailed analysis. While the use of conservative values may, at first, seem to be a "safe" analysis process, it can have negative impacts. More conservative values can focus the risk on the wrong human actions or related accident sequences and equipment failures because of the intentionally high HEPs. Such incorrect conclusions need to be avoided by ensuring a sufficient set of more realistic, detailed HEPs are included in the

model. In fact, non-significant HFEs should also be assessed in detail if detection of all weaknesses in plant design or practices is desirable given the application.

5.3.3.4 Good Practice #4: Revisit the Use of Post-Initiator Screening Values versus Detailed Assessments for Special Applications

For a specific PRA application and depending on the issue being addressed (e.g., examination of a specific procedure change), revisit the use of conservative (screening) versus detailed- assessed HEPs to ensure issue-relevant human actions have not been prematurely deleted from the PRA or there is an inappropriate use of conservative vs. detailed values to properly assess the issue and the corresponding risk.

5.3.3.5 Good Practice #5: Account for Plant and Activity Specific PSFs in the Detailed Assessments of Post-Initiator HEPs

As "good practice," the following table of PSFs (Table 5-1) for both in-control room and local (ex-control room) actions should be treated in the evaluation of each HEP per the table guidance. The guidance should fit most cases, but it should be recognized that for specific actions, some of the factors may not apply. Also, there may be HFEs and contexts for which some PSFs are so important, the others do not matter (e.g., time available is so short, the act almost assuredly cannot be done regardless of the other factors). Further, if a specific situation warrants treatment of unique factors that are not, and cannot be addressed by the list of factors in the table, identification of other PSFs should complement the list. Consideration of the impact of the factors on the HEPs should be as plant- and accident sequence-specific as necessary to address the issue. In addition, such influences should be confirmed, where useful, by such techniques as talk-throughs, walkdowns, field observations, simulations, and examination of past events in order to be realistic. Appendix B provides more specific guidance and discussion of the PSFs presented in the table.

It should be noted that there are aleatory aspects of some or all of the PSFs listed below and not all of these aleatory influences will need to be considered for all applications. For example, for a license amendment, the more "systematic" PSFs (e.g. procedures, training) or the more systematic aspects of the PSFs (e.g., for the team/crew dynamics PSF, the plant practice for the division of control room responsibilities would tend to be systematic) may be the more important ones. On the other hand, when performing assessments of specific events, some of the more aleatory factors or aspects (e.g., crew aggressiveness, failed instrumentation, etc.) may be directly relevant. Nevertheless, for any application, there may be aleatory factors that could be very important for given scenarios and analysts should at least consider whether various aleatory influences could be important enough in the context of the event to be considered directly. For example, if the timing of the scenario is such that only aggressive crews will be able to reach the relevant step in the procedure soon enough, and it is determined that the plant operating crews vary significantly in terms of their aggressiveness in implementing the procedures, then direct modeling of this variable may be appropriate. At a minimum, such aleatory factors should be considered in determining the mean HEP value and in consideration of the uncertainty.

The analysis should ensure that the factors seemingly most relevant to the HFE (either as positive or negative influences) and having the most impact on the HEP, have been considered quantitatively. Furthermore, the quality of the HFE evaluations is improved when the impacts of relevant factors have

been determined from collected information (e.g., talkthroughs, walkdowns, field observations, simulations) rather than simple judgments.

Table 5-1 Post-Initiator PSFs To Be Considered for Both Control Room and Local (Ex-Control Room) Actions

Post-Initiator PSFs To Consider (Relative to Each Scenario and HFE)	Conditions When Particularly Relevant	
	Control Room Actions	Local Actions
Applicability and suitability of training and experience	Always	Always
Suitability of relevant procedures and administrative controls	Always	Always
Availability and clarity of instrumentation (cues to take actions as well as confirm expected plant response)	Always	When time is short
Time available and time required to complete the act, including the impact of concurrent and competing activities	Always	Always
Complexity of required diagnosis and response. In addition to the usual aspects of complexity, the need for special sequencing, organization, and coordination can also be contributors to complexity	Unfamiliar Situation	Unfamiliar Situation
Workload, time pressure, stress	Crew is aware of time constraints	Crew is aware of time constraints
Team/crew dynamics and crew characteristics (degree of independence among individuals, operator attitudes - biases - rules, use of status checks, approach for implementing procedures, e.g., aggressive vs. slow and methodical). Note. Observation of simulator exercises and discussions with operating crews and trainers are particularly important to obtaining this type of information. Understand that organizational attitudes and rules may bear on aspects of crew behavior and should be considered when possible.	Always	When the timing and the appropriateness of the directions from the control room (CR) is critical or could be affected by the scenario, or when the subsequent carrying out of the local [ex-CR action(s)] involves teamwork.
Available staffing and resources	If typical CR staff is expected to be decreased or impacted so that others must perform more than their typical tasks. (Not usually an issue, but a fire scenario might be an exception).	Particularly when many or complex actions need to occur concurrently or in a short time, and staffing needs may be stretched. Time of day may be a contributing factor to the availability of staff.

Post-Initiator PSFs To Consider (Relative to Each Scenario and HFE)	Conditions When Particularly Relevant	
	Control Room Actions	Local Actions
Human-system interface	If could be problematic, or not easily accessed or used (not usually an issue but consider, for instance, the need to use backboards, deal with common workarounds)	If could be problematic (e.g., poor labeling) or not easily accessed or used
Environment in which the act needs to be performed	Potentially adverse or threatening situations such as fire, flood, seismic, loss of ventilation (not usually an issue)	Potentially adverse situations such as high radiation, high temperature, high humidity, smoke, toxic gas, noise, poor lighting, weather, flooding, seismic
Accessability and operability of equipment to be manipulated	If operability could be problematic, or not easily accessed or used, such as the need to use backboards, or when indications/controls could be affected by the initiating event or other failures (e.g., loss of DC)	If operability could be problematic (e.g., rarely used), or not easily accessed, such as when the equipment could be affected by the initiating event (e.g., fire, flood, loss of power)
The need for special tools (keys, ladders, hoses, clothing such as to enter a radiation area...)	Not usually an issue but consider, for instance, when accessability of keys for keylock switches could be important	For situations where other than simple switch or similar type operations are necessary, or when needed to be able to access the equipment
Communications (strategy and coordination) as well as whether one can be easily heard	Not usually an issue - simply ensure that communication strategy allows crisp direction and proper feedback; otherwise only in special situations such as needing to communicate while wearing a self-contained breathing apparatus (SCBA)	For situations where communication among crew members (locally and/or with CR) are likely to be needed and there could be a threat such as too much noise, failure of the communication equipment, availability and location issues associated with the communication equipment.

Post-Initiator PSFs To Consider (Relative to Each Scenario and HFE)	Conditions When Particularly Relevant	
	Control Room Actions	Local Actions
Special fitness needs	Typically not an issue	For special situations expected to involve the use of heavy or awkward tools/equipment, carrying hoses, climbing, etc.
Consideration of 'realistic' accident sequence diversions and deviations (e.g., extraneous alarms, failed instruments, outside discussions, sequence evolution not exactly like that trained on..) Note. This item is essentially addressing aleatory factors that could have important affects on performance. While analysts may choose to explicitly address these factors only when a more detailed investigation of the scenario is warranted or when they are explicitly part of the question being asked, it should be recognized that in some cases they could have strong effects. If they are not addressed explicitly in the analysis, it is suggested that their potential impacts be considered in addressing the uncertainty associated with the HEP values.	When reasonable variations in plant conditions for the scenario (even if of low probability) have the potential to confuse operating crews.	Rarely important, but to the extent that these aspects could affect the timing, specific directions, or successful performance of the local action(s), they should be considered

5.3.3.6 Good Practice #6: Account for Dependencies Among Post-Initiator HFEs

Dependencies among the post-initiator HFEs and hence the corresponding HEPs in an accident sequence should be quantitatively accounted for in the PRA model by virtue of the joint probability used for the HEPs. This is to account for the evaluation of each sequence holistically, considering the performance of the operators throughout the sequence response and recognizing that early operator successes or failures can influence later operator judgments and subsequent actions. This is particularly important so that combined probabilities that are overly optimistic are not inadvertently assigned, potentially resulting in the inappropriate decrease in the risk significance of human actions and related accident sequences and equipment failures. In the extreme, this could result in the inappropriate screening out of accident sequences from the model because the combined probability of occurrence of the events making up an accident sequence drops below a threshold value used in the PRA to drop sequences from the final risk results.

In analyzing for possible dependencies among the HFEs in an accident sequence, look for links among the acts represented in different HFEs including:

- the same crew member(s) is responsible for the acts,
- the actions take place relatively close in time such that a crew "mindset" or interpretation of the situation might carryover from one event to the next,
- the procedures and cues used along with the plant conditions related to performing the acts are

identical (or nearly so) or related, and the applicable steps in the procedures have few or no other steps in between the applicable steps,

- similar PSFs apply to acts,

- how the acts are performed is similar and they are performed in or near the same location, and

- there is reason to believe that the interpretation of the need for one action might bear on the crews decision regarding another action, i.e., the basis for one decision in a scenario may influence another decision later in the scenario.

The more the above commonalities and similarities exist, the greater the potential for dependence among the HFEs (i.e., if the first act is not performed correctly, there is a higher likelihood the second, third... act(s) will also not be performed correctly; and vice versa if the act(s) are successful). For example, if nearly all or all of the above characteristics exist, very high or complete dependence should generally be assumed. If only one or two of the above characteristics exist, then analysts will need to evaluate the likely strength of their effect and the degree of dependence that should be assumed and addressed in quantification.

The resulting joint probability of the HEPs in an accident sequence should be such that it is in line with the above characteristics and the following guidance, unless justified otherwise:

- The total combined probability of all the HFEs in the same accident sequence/cut set should not be less than a justified value. It is suggested that the value not be below ~0.00001 since it is typically hard to defend that other dependent failure modes that are not usually treated (e.g., random events such as even a heart attack) cannot occur. Depending on the independent HFE values, the combined probability may need to be higher.

- To the extent the joint HEPs are looked at separately, but a previous human action in the sequence has failed, then:

  - A factor of 3-10 higher than what would have been the independent HEP value for the subsequent act(s) exists for low to moderate dependence

  - 0.1 up to 0.5 is the resulting probability value used for the subsequent HEP(s) for high dependence

  - $\geq$0.5 exists for the subsequent HEP(s) for very high or 1.0 for complete dependence.

5.3.3.7 Good Practice #7: Assess the Uncertainty in Mean HEP Values

Mean values for each HEP (excluding conservative HEPs) and an assessment of the uncertainty in the mean values should be performed at least for the significant HEPs to the extent that these uncertainties need to be understood and addressed in order to make appropriate risk-related decisions. Assessments of uncertainty are typically performed by:

- assigning uncertainty distributions for the HEPs and propagating them through the quantitative

analysis of the entire PRA, such as by a Monte Carlo technique, and/or

- performing sensitivity analyses that demonstrate the effects on the risk results for extreme estimates in the HEPs based on at least the expected uncertainty range about the mean value.

Note, in some cases, it may be sufficient to address the uncertainties by just qualitative arguments without the need to specifically quantify them (e.g., justifying why the HEP cannot be very uncertain or why a change in the HEP has little relevancy to the risk-related decision to be made).

Generally, however, a more thorough treatment of uncertainty is needed. While the guidance provided below describes good practice for treating uncertainty, in most cases it will probably go beyond what has been done in even recent PRA applications. Nevertheless, analysts should consider such approaches to the extent necessary to ensure that their application is as realistic as necessary. In other words, they need to examine the question being addressed and ask whether a more thorough treatment of uncertainty would be beneficial.

Thus, it is recommended that in assessing the uncertainties and particularly when assigning specific uncertainty distributions, the uncertainties should include:

- those epistemic uncertainties existing because of lack of knowledge of the true expected performance of the human for a given context and associated set of performance shaping factors (i.e., those factors for which we do not have sufficient knowledge or understanding as to the "correct" HEP, such as how time of day affects the bio-rhythm and hence, performance of operators), and

- consideration of the combined effect of the relevant aleatory (i.e., random) factors to the extent they are not specifically modeled in the PRA and to the extent that they could significantly alter the context and PSF evaluations for the HFE, and thereby the overall HEP estimate.

Concerning the latter, it is best to specifically model the aleatory factors in the PRA (i.e., those factors that are random and could significantly affect operator performance, for example, whether or not other nuisance alarms or equipment failures may co-exist with the more important failures in the sequence, whether a critical equipment failure occurs early in the sequence or late in the sequence, etc.). However, this is often impractical and is typically not done as it would make the PRA model excessively large and unwieldy. Thus in assigning the mean HEP and uncertainty distribution, analysts should reflect an additional contribution from random factors associated with the plant condition or overall action context. This can be done by considering the relevant aleatory (i.e., random) factors, their likelihoods of occurrence, and their effects on the HEP estimate.

For example, suppose for an accident sequence(s) it is judged that the human performance will be significantly affected by the number of "nuisance and extraneous failures," as opposed to when no or few nuisance/extraneous failures exist (and yet these two plant "states" are not explicitly defined by the PRA model). Further, based on the analyst considering how the HEP is affected, a value of P_0 would be estimated for when no or few nuisance/extraneous failures exist and a value of P_1 would be estimated for when many do exist, and the difference between P_0 and P_1 is significant (e.g., factor of 10). It is also judged that many nuisance/extraneous failures will occur about 50% of the time based on past experience. The resulting combined mean HEP value is $0.5P_0 + 0.5P_1$ considering this random

factor. The overall uncertainty about the combined mean HEP value should reflect the weighted epistemic uncertainties in P_0 and P_1 (such as by a convolution approach, via an approximation, or other techniques). While it is not expected that such a detailed evaluation be done for every random situation or for every HEP, the mean and uncertainty estimates for the most significant HEPs should account for any such perceived important aleatory factors that have not otherwise been accounted for (i.e., if the factors, considering their likelihoods and effects on the HEP, are anticipated to have a significant impact on the resulting overall HEP).

Whatever uncertainty distributions are used, the shape of the distributions (log-normal, beta, etc.) are typically unimportant to the overall risk results (i.e., the PRA results are usually not sensitive to specific distributions). Further, typical uncertainties include values for the HEP that represent a factor of 10 to 100 or even more between the lower bound value and the upper bound value that encompass the mean value. However, it should be noted that some distributions, e.g., log-normal, can give probabilities greater than 1.0 for HEPs that are relatively high.

5.3.3.8 Good Practice #8: Evaluate the Reasonableness of the HEPs Assigned Using Detailed Assessments

The HEPs for post-initiator HFEs (excluding the conservative HEPs) should be reasonable from two standpoints:

- first and foremost, relative to each other (i.e., the probabilistic ranking of the failures when compared one to another), and

- in absolute terms (i.e., each HEP value), given the context and combination of positive and negative PSFs and their relative strengths .

This reasonableness should be checked based on consideration of actual plant experience and history, against other evaluations (such as for similar acts at other plants), and the qualitative understanding of the actions and the relevant contexts and performance shaping factors under which the acts are performed.

It is suggested that a rank-ordered list of the post-initiator HFEs by probability be used as an aid for checking reasonableness. As part of such a list, it is particularly worthwhile to compare "like" HFEs for different sequences such as failure to manually depressurize in a BWR when all high pressure injection is lost during a small LOCA as compared to the same action but during a simple transient. For example, simple, procedure-guided actions with easily recognized cues and plenty of time to perform the actions, should have lower HEPs than complex, memorized, short time available type actions, all other factors being the same. Typical expectations of most post-initiator HEPs are in the 0.1 to 0.0001 range and depend particularly on the relevant contextual factors and proper consideration of dependencies as discussed under many of the Good Practices covered above. Helpful checks include:

- For a HFE, are there one or two dominant PSFs that exist or is the cumulative effect of the relevant PSFs such that they are either so negative or so positive that a 'sanity check' would

suggest a high HEP (e.g., 0.1) or a low HEP (e.g., 0.0001), respectively? For example, if the procedures and training fit the scenario well, the cues will be clear, there is plenty of time available, and the actions are simple and familiar, then an HEP of 0.001 to 0.0001 would be reasonable. Alternatively, if there is a reasonable expectation that the scenario could be confusing, it is not frequently trained on, there may be some hesitancy associated with taking the action, and only an aggressive crew would be likely to diagnose and complete the action in the time available, then an HEP 0.5 or even 1.0 would not be unreasonable. Accordingly, this very high or low probability HFE should be one of the higher or lower probability HFEs relative to the other HFEs in the model.

- Are there seemingly balanced combinations of both positive and negative factors, or are there weak to neutral factor effects? If so, this is likely to lead to in-between values for the HEPs (e.g., ~0.01) placing these HFEs (relative to others) 'in the middle'.

- Do the individual HEPs and the relative ranking of the HFEs seem consistent with actual or simulated experience? For example, if it is known that operators 'have trouble with' a specific act(s) in simulations or practiced events, and yet the assigned HEP is very low (e.g., 0.001 or lower), this may be a reason to question and revisit the assigned HEP.

- Do other similar plant and action analyses support the HEP evaluation? Recognize, however, that there may be valid reasons why differences may exist and thus this check is not likely to be as helpful as the others above.

5.3.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to quantify the post-initiator HEPs as realistically as possible using the good practices articulated above (except for where higher screening estimates are used, purposely and appropriately), could result in improper HEPs and thus inaccuracies in the PRA results and particularly incorrect assessments of the importance of pre-initiator human failures. The risk effect of the human failure could be over-emphasized (such as if the human failure is estimated with too high [pessimistic] a HEP or a high screening estimate is used where a more realistic detailed estimate is appropriate), or under-estimated (such as if the human failure is estimated with too low [optimistic] a HEP or a dependency among failures is not accounted for resulting in too low a joint probability for multiple human failures). Besides these concerns about inaccuracies in the HEP quantification and thus whether the HEPs "make sense," as well as the resulting potential misinformation about the significant risk contributors if quantification is not done well, the following related observations are noted.

- Use of conservative values is a useful and most often necessary part of HRA so as to avoid the expenditure of resources on unimportant human events and accident sequences. The above guidance is aimed at allowing some conservative values without inadvertently and inappropriately allowing the analytical phenomenon of, for instance, multiplying four human events in the same sequence each at a conservative estimate of 0.1 to yield a 0.0001 combined probability, without checking for dependencies among the human events. In such a case some human failure events and combinations of events, or even whole accident sequences, may

inappropriately screen out of the PRA model entirely because the accident sequence frequency drops below a model threshold. Hence some of the significant individual or combination contributors may be missed. This is why the conservative estimates both individually and for combined events should not be too low. Further, if conservative values are left permanently assigned to some human failure events that should be assessed with more detail to obtain a more realistic assessment of risk (supposedly lowering the probability), the risk significance of these human failure events and related equipment failures are likely to be over-emphasized at the expense of improperly lessening the relative importance of other events and failures.

- It is important to be sure that dependencies among the various modeled HFEs including those with conservative values, have been investigated. Treating HFEs, whether with conservative values or based on more detailed analysis, as independent acts without checking for dependencies (thereby being able to multiply the individual HEPs) can inappropriately lessen the risk significance of those HFEs and related equipment failures in accident sequences. This can cause the inappropriate dropping out of accident sequences because the sequences quantitatively drop below a model threshold value as discussed above under screening. Proper consideration of the dependencies among the human actions in the model is necessary to reach the best possible evaluation of both the relative and absolute importance of the human events and related accident sequence equipment failures.

- The use of mean values and addressing uncertainties are a part of the Regulatory Guide 1.174[3] guidance and to the extent addressed therein, the HRA quantification needs to be consistent with that guidance when making risk-informed decisions. The estimates should reflect, to the extent possible, the as-built and as-operated conditions as addressed in the plant- and activity-specific PSFs.

- There can be a tendency to use an existing PRA model to address issues such as changes to the plant, without spending the appropriate time to revisit some of the underlying assumptions and modeling choices made to create the original PRA. However, such a review should be done to see if these assumptions and choices still apply for the issue being addressed. For instance, some post-initiator human failure events may be quantified in the original model using conservative estimates and detailed failure probabilities that may not be appropriate for the new issue being addressed. As an example, where high conservative HEPs may have been acceptable for purposes of the original PRA, these may over-estimate the contribution of these human failure events for the issue being addressed. Further, the relative risk contribution of equipment and associated accident sequences with which the human failure events appear, may be artificially too high (and therefore other events too low) because of the conservative values. Hence it is good practice to revisit the use of conservative estimates and detailed human failure event probabilities in order to appropriately address the issue.

5.4 Adding recovery actions to the PRA

5.4.1 OBJECTIVE: To address what recovery actions can be credited in the post-initiator HRA and the requirements that should be met before crediting recovery actions. Adding recovery actions is common practice in PRA and accounts for other reasonable actions the operators might take

to avoid severe core damage and/or a large early release that are not already specifically modeled. For example, in the PRA modeling of an accident sequence involving loss of all injection, it would be logical and common to credit the operators attempting to locally align an independent firewater system for injection. The failure to successfully perform such actions would subsequently be added to the accident sequence model thereby crediting the actions and further lowering the overall accident sequence frequency because it takes additional failures of these actions before the core is actually damaged. The following provides good practices for crediting post-initiator recovery actions while implementing Regulatory Guide 1.200[4] and the related ASME Standard[5] requirements.

5.4.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard requires that recovery actions be modeled only if it has been demonstrated that the actions are plausible and feasible for those sequences to which they are applied. There are multiple supporting requirements in the Standard under high-level requirement HLR-HR-H that address what recovery actions can be credited as well as the need to consider dependencies among the HFEs and any recovery actions that are credited. Regulatory Guide 1.200 states the NRC staff has just one clarification to the ASME Standard requirements covering this activity and that is with regard to factors that need to be considered when accounting for dependencies between the recovery HFE and other HFEs in the sequence. This has been addressed in the good practices.

5.4.3 GOOD PRACTICES:

5.4.3.1 Good Practice #1: Define Appropriate Recovery Actions

Based on the failed functions, systems, or components, identify recovery actions (see Footnote #1 for the specific definition being used and the distinction between recovery actions and repair actions) to be credited that are not already included in the PRA (e.g., aligning another backup system not already accounted for) and that are appropriate to be tried by the crew to restore the failure. The following should be considered in defining appropriate recovery actions:

- the failure to be recovered,
- whether the cues will be clear and provided in time to indicate the need for a recovery action(s), and the failure that needs to be recovered,
- the most logical recovery action(s) for the failure, based on the cues that will be provided,
- the recovery is not a repair action (e.g., the replacement of a motor on a valve so that it can be operated),
- whether sufficient time is available for the recovery action(s) to be diagnosed and implemented to avoid the undesired outcome,

- whether sufficient crew resources exist to perform the recovery(ies),
- whether there is procedure guidance to perform the recovery(ies),
- whether the crew has trained on the recovery action(s) including the quality and frequency of the training,
- whether the equipment needed to perform the recovery(ies) is accessible and in a non-threatening environment (e.g., extreme radiation), and
- whether the equipment needed to perform the recovery(ies) is available in the context of other failures and the initiator for the sequence/cut set.

In addressing the above issues and assessing which recovery action, or actions, to credit in the PRA, for post-initiator HFEs all the good practices provided earlier in Sections 5.1, 5.2, and 5.3 apply (i.e., the failure to recover is just another HFE like all the other post-initiator HFEs). In general, no recovery should be credited where any of the above considerations are not met (e.g., there is not sufficient time, there are no cues that there is a problem, there are not sufficient resources, there is no procedure or training). It may be possible to justify exceptions in unique situations, such as a procedure is not needed because the recovery is a skill-of-the-craft, non-complex, and easily performed; or the specific failure mode of the equipment is known for the sequence (this is usually not the case at the typical level of detail in a PRA) and so "repair" of the failure can be credited because it can be easily and quickly diagnosed and implemented. Any exceptions should be documented as to the appropriateness of the recovery action.

When considering multiple recoveries (i.e., how many recoveries to be credited in one accident sequence/cut set), the above considerations apply to all the recoveries. The analyst should also consider that one recovery may be tried (perhaps even multiple times) and then the second recovery may be tried but with even less time and resources available because of the attempts on the first recovery. Hence the failure probability of the second recovery should be based on more pessimistic characteristics (e.g., less time available, less resources) than if such a possibility is not considered.

5.4.3.2 Good Practice #2: Account for Dependencies

As stated above, all the good practices provided earlier in Sections 5.1, 5.2, and 5.3 apply. From these good practices, particular attention should be paid to accounting for dependencies among the HFEs including the credited recovery actions. More specifically, dependencies should be assessed:

- among multiple recoveries in the accident sequence/cut set being evaluated, and
- between each recovery and the other HFEs in the sequence/cut set being evaluated.

As part of this effort, the analyst should give proper consideration to the difficulties people often have in overcoming an initial mind-set despite new evidence. For a real world example, consider how long the pressure-operated relief valve (PORV) path remained open in the Three Mile Island accident despite new cues of the problem, different personnel arriving, etc. For this and similar reasons, the

assessing of no dependence needs to be adequately justified to ensure the quantified credit for the recovery action(s) is not unduly optimistic.

5.4.3.3 Good Practice #3: Quantify the Probability of Failing to Perform the Recovery(ies)

Quantify the probability of failing to perform the recovery(ies) by:

- using representative data that can be shown to be appropriate for the recovery event(s) (e.g., using data that exists for typical times to recover offsite power), or
- using the HRA method/tool(s) used for the other HFEs (i.e., using an analytical/modeling approach).

In performing the quantification, one should ensure that all the good practices under Section 5.3 are followed (for each individual recovery as well as for multiple/joint recovery credit). In addition, if using data, ensure the data is applicable for the plant/sequence context or that the data is modified accordingly. For example, a plant may use available experience data for the probability of failing to align a firewater system for injection but the experience data is based on designs for which all the actions can be taken from the main control room whereas for this plant, the actions have to be performed locally.

5.4.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS:

Failure to follow the above good practices is likely to lead to recovery credit that is applied too optimistically; that is, the failure to recover is assigned too low a probability. Hence an under-estimate of the failure to recover is applied to the PRA accident sequence/cut set, making the affected sequence/cut set artificially too low in risk significance. This can subsequently affect the ranking of the important sequences, equipment failures, and human actions, potentially leading to false conclusions as to the significant risk contributors.

6. ERRORS OF COMMISSION (EOCs)

Explicit modeling of errors of commission[a] has generally been beyond current PRA practice and is not explicitly addressed in Regulatory Guide 1.200[4] or ASME Standard[5] HRA requirements. Many practitioners believe that the number of potential operator acts that could result in an EOC is unmanageably large. That is, they believe that there are a large number of potential acts that would be adverse to safe shutdown (e.g., fail or make unavailable equipment/functions relevant to mitigating the scenario, or otherwise exacerbate the scenario such as by opening a PORV and causing an unwanted loss of coolant accident), even for what may appear to be justifiable reasons. Errors of

[a] Error of Commission (EOC): A *human failure event* resulting from an overt, unsafe action, that, when taken, leads to a change in plant configuration with the consequence of a degraded plant state. Examples include terminating running safety-injection pumps, closing valves, and blocking automatic initiation signals.

omission[b] are typically modeled in PRAs because they are easily defined and limited by the requirements of operating procedures (especially emergency operating procedures). At best, PRAs have handled EOCs implicitly (e.g., as part of a base HEP) without a systematic or adequate search for this type of error.

However, recent work in the area (e.g., ATHEANA,[10] CESA,[18] Julius et al.,[19] and Wakefield[20]) has made advances in the ability to identify EOCs without the need to perform an exhaustive search. One of the lessons learned from the development and application of ATHEANA[10] is that the effort needed to identify EOCs can be substantially reduced by focusing the search on identifying systematic vulnerabilities in plant operations associated with plant critical functions.

6.1 Including and modeling EOCs

6.1.1 OBJECTIVE: To describe the conditions under which EOCs should be considered for inclusion in the HRA modeling.

6.1.2 REGULATORY GUIDE 1.200 POSITION:

There is no specific treatment of EOCs.

6.1.3 GOOD PRACTICES

6.1.3.1 Good Practice #1: Address EOCs in Future HRAs/PRAS (Recommendation)

Given the recent advances in the ability to address EOCs and the potential for regulatory requirements to make the need to address EOCs more important, it is recommended that future HRA/PRAs attempt to identify and model potentially important EOCs. Sources available to support this process are listed above. It is suggested that multiple sources be consulted to assist this process.

To the extent any EOCs are modeled, all the guidance in this document has been written with both types of errors in mind; that is, all the same good practices apply whether the error is one of omission or commission. However, as a point of emphasis, in assessing the probability of identified potential EOCs, it is particularly important to consider the role that plant indications will play in supporting a crew's ability to detect and recover from EOCs. At least for many EOCs, there may be immediate changes in plant conditions that would alert them to such recovery actions.

6.1.3.2 Good Practice #2: As a Minimum, Search for Conditions That May Make EOCs More Likely

Even if the recommended first good practice above is not performed, the use of risk in any issue assessment should at least ensure that conditions that promote likely EOCs do not exist. For example,

---

[b] Error of Omission (EOO): A *human failure event* resulting from a failure to take a required action, that leads to an unchanged or inappropriately changed plant configuration with the consequence of a degraded plant state. Examples include failures to initiate standby liquid control system, start auxiliary feedwater equipment, and block automatic depressurization system signals.

it should be ensured that such conditions have not been introduced by a plant change or modification, or that the plant is not more susceptible to EOCs under the unique set of conditions being examined, (e.g., pressurized thermal shock scenarios). When considering the potential for situations that may make EOCs somewhat likely, the premise of any evaluation should be that:

- operators are performing in a rationale manner (e.g., no sabotage), and
- procedural and training guidance used by the crew will be selected on the basis of the plant status inputs they are receiving.

Using this premise, EOCs are typically the result of problems in the plant information/operating crew interface (e.g., wrong, inadequate information is present, or the information can be easily misinterpreted) or in the procedure-training/operating crew interface (e.g., procedures/training do not cover the actual plant situation very well because they provide ambiguous guidance, no guidance, or even unsafe guidance for the actual situation that may have evolved in a somewhat unexpected way). In either case, significant mismatches can occur between the scenario conditions (i.e., context) and the crew's understanding of those conditions (see ATHEANA[10] for more detailed discussion on the role of mismatches in facilitating EOCs).

With a focus on analysts and on reviewers reviewing potential applications of current PRAs, the following is offered as guidance in this area to aid in ensuring EOC-prone conditions do not exist or have not been introduced as part of a plant change. Hence, a review of a plant change should look for situations where one or more of the following characteristics are introduced as a result of the change and thus should be corrected if possible.

- To address the potential impacts of mismatches between scenario context and plant information/interface, an analysis/review should at least look for those acts that operators may take that (a) would fail or otherwise make unavailable a PRA function or system, or (b) would reduce the accident mitigating redundancy available, or (c) would exacerbate an accident challenge, because the change has caused such an action to be performed on the basis of just one primary input/indication for which there is no redundant means to verify the true plant status. Such a situation identifies a vulnerable case where EOCs may likely be performed based on just one erroneous (failed, spurious, etc.) input such as an alarm, indicator, or verbal cue of an observed condition.

  In identifying such cases, one should keep in mind that multiple indications may use the same faulty input (e.g., subcooling margin indication and primary system indication may use the same pressure transmitter(s); multiple reactor vessel level indications may rely on the same power supply) and therefore a single fault may actually affect multiple inputs observable to the operator. Depending on the how the failure affects the indications (fail high, low, mid-scale, etc.), the failure may not be "obvious" and a EOC-prone situation may exist that may need to be rectified.

- To identify potential problems with the procedure-training interface, an analysis/review should at least look for those acts that operators may take that (a) would fail a PRA function or system, or (b) would reduce the accident mitigating redundancy available, or (c) would

exacerbate an accident challenge, because the change has caused the procedure (including entry conditions) and/or training guidance:

- to become ambiguous/unclear (e.g., vague criteria as to when to abandon the main control room),
- to introduce a repetitive situation in the response steps where a way to proceed out of the procedure and/or the specific repetitive steps is not evident (e.g., at the end of a series of steps, the procedure calls for a return to a previous step with no clear indication as to how the operators ultimately get out of the repetitive loop of steps),
- to place the operators in dilemma conditions without some guidance/criteria as to how to "solve" the dilemma (e.g., being vague as to whether or not to shutdown a diesel with a cooling malfunction when all other ac power is unavailable),
- to require the operators to rely on memory, especially for complex or multi-step tasks, or
- to require the operators to perform calculations or make other manual adjustments, especially in time-sensitive situations.

The above identify vulnerable cases where EOCs may likely be performed because the procedures and/or training do not adequately cover accident situations that may be faced by the operator or rely on techniques (require memory or adjustments) that may be difficult to perform properly, especially when in a dynamic response situation. In these cases, mismatches between the actual event response that is required and the procedure/training guidance can become magnified making conditions potentially more prone to EOCs.

Note that additional discussions on situations that can facilitate the occurrence of EOCs is provided in ATHEANA,[10] CESA,[18] and Julius et al.[19]

6.1.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS

One concern with failing to adequately address EOCs is that reviews of operational events (e.g., ATHEANA[10]) have shown that they are often important contributors in serious accidents. Moreover, when they do occur, it is often because the operating crew has become confused because of some unexpected characteristics of the context and there is a strong "basis" for taking the inappropriate action. Thus, not including EOCs will likely cause the overall risk profile to be optimistic since these additional sources of risk will have been neglected. Even though contexts that can lead to EOCs may be relatively rare, when they do occur there may be a high probability that the EOCs will follow. From a safety standpoint and in terms of identifying potential plant vulnerabilities, it becomes important to at least perform the 2nd good practice above (if the 1st good practice cannot or is not performed) to at least qualitatively identify those plant conditions that could lead crews to make errors of commission, and address them if necessary.

7. HRA DOCUMENTATION

The ASME Standard[5] provides a set of requirements for documenting a human reliability analysis (HRA) in a manner that facilitates PRA applications, upgrades, and peer review. Specific requirements are provided. The following provides good practice for documenting an HRA building on those requirements.

7.1 Documenting the HRA

7.1.1 OBJECTIVE: To provide the requirements for documenting a HRA. It is recognized that these requirements may be met at various degrees of detail depending on the application; but that most likely, all the requirements should be addressed.

7.1.2 REGULATORY GUIDE 1.200 POSITION:

The ASME Standard provides a list of requirements for documenting a HRA under high-level requirement HLR-HR-I, all of which are included in the good practice below. Regulatory Guide 1.200 states the NRC staff has just one clarification; that this section of the standard is written for impact on CDF and not LERF (seemingly implying that the LERF documentation is covered elsewhere in the standard). Additionally, the reader is referred to Section 1.2.6 of the regulatory guide for further guidance on documentation, in general. That section focuses on the need to ensure traceability and defensibility of the work.

7.1.3.1 Good Practice #1: Document the HRA

The level of detail to be addressed in the documentation depends on the PRA application and the issue being addressed as well as the objectives, scope, and level of detail of the analysis. Whatever documentation is provided, the test for adequate documentation should be: Can a knowledgeable reviewer understand the analysis to the point that it can be at least approximately reproduced and the resulting conclusion reached if the same methods, tools, data, key assumptions, and key judgments and justifications are used? Hence, the documentation should include the following, but only to the extent it is applicable for the application:

- the overall approach and disciplines involved in performing the HRA including to what extent talk-throughs, walkdowns, field observations, and simulations were used,
- summary descriptions of the HRA methodologies, processes, and tools used to:
  - identify the pre-and post-initiator human actions,
  - screen pre-initiators from modeling,
  - model the specific HFEs, including decisions about level of detail and the grouping of individual failures into higher order HFEs,
  - quantify the HEPs with particular attention to the extent to which plant and accident sequence-specific information was used, as well as how dependencies were identified and treated,

- assumptions and judgments made in the HRA, their bases, and their impact on the results and conclusions (generic or on a HFE-specific basis, as appropriate),

- for at least each of the HFEs important to the risk decision to be made, the PSFs considered, the bases for their inclusion, and how they were used to quantify the HEPs, along with how dependencies among the HFEs and joint probabilities were quantified,

- the sources of data and related bases or justifications for:

  - the screening and conservative values,
  - the best estimate values and their uncertainties with related bases,

- the results of the HRA including a list of the important HFEs and their HEPs, and

- conclusions of the HRA.

7.1.4 POSSIBLE IMPACTS OF NOT PERFORMING GOOD PRACTICES AND ADDITIONAL REMARKS

Failure to address the topic areas addressed above in the HRA documentation is likely to mean that an outside reviewer or subsequent user of the HRA (peer reviewer, NRC regulator, a new analyst not involved in the original work, etc.) will have insufficient information to be able to independently understand the analysis. In particular, it may be difficult to decide if the good practices have largely been met and whether the HRA results and conclusions are appropriate and defensible. This could be cause for not accepting a proposed plant change or for misuse of the results in other risk-informed decisions (e.g., Maintenance Rule applications).

8. REFERENCES

[1] *Use of Probabilistic Risk Assessment Methods in Nuclear Activities: Final Policy Statement*, Federal Register, Vol 60, p. 42622 (60FR 42622), US Nuclear Regulatory Commission, August 16, 1995.

[2] *Code of Federal Regulations 10, Parts 1 to 50*, Office of the Federal Register National Archives and Records Admission, Revised as of January 1, 2001.

[3] *An Approach for Using Probabilistic Risk Assessment in Risk-Informed Decisions on Plant-Specific Changes to the Licensing Basis*, Regulatory Guide 1.174, Rev. 1, US Nuclear Regulatory Commission, November 2002.

[4] *An Approach for Determining The Technical Adequacy of Probabilistic Risk Assessment Results For Risk- Informed Activities*, Draft Regulatory Guide 1.200, U.S. Nuclear Regulatory Commission, February 2004.

[5] *Standard for Probabilistic Risk Assessment for Nuclear Power Plant Applications*, ASME RA-S-2002, American Society of Mechanical Engineers, April 5, 2002.

[6] *Probabilistic Risk Assessment Peer Review Process Guidance*, NEI 00-02 Revision A3, Nuclear Energy Institute, March 2000

[7] *Use of Probabilistic Risk Assessment in Plant-Specific, Risk-Informed Decisionmaking: General Guidance*, NUREG-0800, Chapter 19, Rev. 1, US Nuclear Regulatory Commission, November 2002.

[8] A.D. Swain and H.E. Guttmann, *Handbook of Human Reliability Analysis with Emphasis on Nuclear Power Plant Applications - Final Report*, NUREG/CR-1278, SAND80-0200, Sandia National Laboratories, August 1983.

[9] D.E. Embrey, et al., *SLIM-MAUD: An Approach to Assessing Human Error Probabilities Using Structured Expert Judgment (Vols. I & II)*, NUREG/CR-3518, Brookhaven National Laboratory, Upton, NY, 1984.

[10] *Technical Basis and Implementation Guidelines for A Technique for Human Event Analysis (ATHEANA)*, NUREG-1624, Rev. 1, US Nuclear Regulatory Commission, May 2000.

[11] *Severe Accident Risks: An Assessment for Five U.S. Nuclear Power Plants*, NUREG-1150, US Nuclear Regulatory Commission, December 1990

[12] *Guidance for the Review of Changes to Human Actions*, NUREG-1764, US Nuclear Regulatory Commission, December, 2002.

[13] A.D. Swain, *Accident Sequence Evaluation Program Human Reliability Analysis Procedure*, NUREG/CR-4772, SAND86-1996, Sandia National Laboratories, February 1987.

[14] *SHARP1 - A Revised Systematic Human Action Reliability Procedure,* EPRI NP-7183-SL, Electric Power Research Institute, December 1990.

[15] *SPAR-H Method,* NUREG/CR- later, INEEL/EXT-02-10307, Idaho National Engineering and Environmental Laboratory, DRAFT, November 2002.

[16] C. Bieder, P. Le-Bot, E. Desmares, J-L Bonnet, F. Cara, "MERMOS: EDF's new advanced HRA method," in *Probabilistic Safety Assessment and Management (PSAM 4),* A. Mosleh and R.A. Bari (eds), Springer-Verlag, New York, 1998.

[17] Hollnagel, E., *Cognitive Reliability and Error Analysis Method (CREAM).* York: Elsevier Science, New York, 1988.

[18] B. Reer, V.N. Dang, and S. Hirschberg, "The CESA Method and Its Application in a Plant - Specific Pilot Study on Errors of Commission," *Reliability Engineering & System Safety, 83:* 187-205, 2004.

[19] J. Julius, E. Jorgenson, G.W. Parry, and A.M. Mosleh, "A Procedure for the Analysis of Error of Commission in a Probabilistic Safety Assessment of a Nuclear Power Plant at Full Power," *Reliability Engineering and System Safety, 50:* 189-201, 1995.

[20] D.J. Wakefield, "Application of the Human Cognitive Reliability Model and Confusion Matrix Approach in a Probabilistic Risk Assessment," *Reliability Engineering and System Safety, 22:* 295-312, 1988.

[21] J. Reason, *Human Error,* Cambridge, England: Cambridge University Press, 1990 and *Managing the Risks of Organizational Accidents,* Aldershot, UK: Ashgate, 1997.

[22] D.D. Woods et. al., *Behind Human Error: Cognitive Systems, Computers, and Hindsight,* Crew System Ergonomics Information Analysis Center (CSERIAC), The Ohio State University, Wright-Patterson Air Force Base, Columbus, OH, December 1994.

[23] M.R. Endsley, Towards a Theory of Situation Awareness in Dynamic Systems, *Human Factors, 37,* 65-84, 1995.

APPENDIX A

Summary of HRA Good Practices Audit

An audit of HRA good practices was conducted against the Supporting Requirements (SRs) contained in Section 4.5.5, Human Reliability Analysis (HR), of ASME RA-Sa-2003 and Table HR, Human Reliability Analysis (HRA) Modeling Related Grades - Element HR, Appendix B of NEI 00-02. Other SRs and technical elements (TEs) relevant to HRA practice may be contained in other sections of these references; however, these were not identified or considered by the reviewer for this draft. *(However, reviews of the other sections are planned for a later draft).*

The audit indicated that draft HRA good practices existed for all SRs in ASME RA-Sa-2003, Section 4.5.5 and all TEs in NEI 00-02, Table HR, with the exception of HR-29 pertaining to the need for an independent review of the HRA results. Note that the empty cells under the RG 1.200 column indicates that the NRC had no objection to the SRs in ASME RA-Sa-2003, Section 4.5.5 or the TEs in NEI 00-02, Table HR.

Table A-1
Summary of HRA Good Practices Audit Against ASME RA-Sa-2003 and NEI 00-02

Location	Good Practice	Description	ASME RA-Sa-2003	NEI 00-02	RG 1.200	Notes
HRA team formation and techniques for a realistic analysis						
Section 3.1.3.1	1	Perform a Multi-Disciplinary, Integrated Analysis	not addressed	not addressed		
Section 3.1.3.2	2	Perform Field Observations and Discussions	HR-E3	HR-10 HR-14 HR-20		
Identifying potential pre-initiator human failure events (HFEs)						
Section 4.1.3.1	1	Review Pre-Initiator Procedures, Actions, and Equipment	HR-A1 HR-A2 HR-D3	HR-4 HR-5	clarification to NEI 00-02	self-assessment needs to confirm and document that the factors listed in ASME HR-D3 were considered in the pre-initiator HEPs
Section 4.1.3.2	2	Identify All Pre-Initiator Human Actions (Do Not Ignore Pre-Initiators)	HR-C2 HR-C3	HR-5 HR-7 HR-27		
Section 4.1.3.3	3	Examine Other Operational Modes and Routine Actions	not addressed	not addressed		

Table A-1 Summary of HRA Good Practices Audit Against ASME RA-Sa-2003 and NEI 00-02						
Location	Good Practice	Description	ASME RA-Sa-2003	NEI 00-02	RG 1.200	Notes
Section 4.1.3.4	4	Identify Actions Affecting Redundant and Multiple Diverse Equipment	HR-A3 HR-B2	HR-5 HR-6 HR-7 HR-26		
Screening those pre-initiator activities for which HFEs do not need to be modeled						
Section 4.2.3.1	1	Screen Pre-Initiators with Acceptable Restoration Mechanisms or Aids	HR-B1	HR-5 HR-6		
Section 4.2.3.2	2	Do Not Screen Actions Affecting Redundant and Multiple Diverse Equipment	HR-A3 HR-B2	HR-5		
Section 4.2.3.3	3	Re-Evaluate the Screening Process For Special Applications	Section 3	not addressed		Section 3 of the ASME standard provides detailed discussion about the risk assessment application process
Modeling specific HFEs corresponding to the unscreened pre-initiator human actions						
Section 4.3.3.1	1	Include HFEs for Unscreened Human Actions in the PRA Model.	HR-C1 HR-C2 HR-C3	HR-5 HR-7 HR-27		
Quantifying the corresponding human error probabilities (HEPs) for the specific pre-initiator HFEs						
Section 4.4.3.1	1	Use Screening Values During the Initial Quantification of the HFEs	HR-B1 HR-D1 HR-D2	HR-5 HR-6 HR-13		
Section 4.4.3.2	2	Detailed assessments of at least the significant HFE contributors should be performed	HR-B1 HR-D1 HR-D2	HR-5 HR-6		
Section 4.4.3.3	3	Revisit the use of screening vs. detailed-assessed HEPs for specific PRA applications	Section 3	not addressed		Section 3 of the ASME Standard provides detailed discussion about the risk assessment application process

Table A-1 Summary of HRA Good Practices Audit Against ASME RA-Sa-2003 and NEI 00-02						
Location	Good Practice	Description	ASME RA-Sa-2003	NEI 00-02	RG 1.200	Notes
Section 4.4.3.4	4	Account for Plant and Activity Specific Performance Shaping Factors (PSFs) in the Detailed Assessments	HR-G3 HR-G4 HR-G5	TBD		
Section 4.4.3.5	5	Apply Plant Specific Recovery Factors	HR-D4	not addressed	clarification to NEI 00-02	NEI 00-02 does not address use of expert judgement
Section 4.4.3.6	6	Account for Dependencies Among the HEPs in an Accident Sequence	HR-D5	HR-26 HR-27		
Section 4.4.3.7	7	Assess the Uncertainty in Mean HEP Values	HR-D6	not addressed		
Section 4.4.3.8	8	Evaluate the Reasonableness of the HEPs Obtained Using Detailed Assessments	HR-D7	not addressed		
Identifying potential post-initiator human failures						
Section 5.1.3.1	1	Review Post-Initiator Related Procedures and Training Materials (and plant simulator training	HR-E1(a) HR-E3 HR-E4	HR-9 HR-10 HR-14 HR-16 HR-20		
Section 5.1.3.2	2	Review Functions and Associated Systems and Equipment to be Modeled in the PRA	HR-E1(b)	HR-9 HR-10 HR-16		
Section 5.1.3.3	3	Look for Certain Expected Types of Actions (general types of post-initiator actions to be considered)	HR-E2	HR-8 HR-9 HR-10 HR-21 HR-22 HR-23 HR-25		
Modeling specific HFEs corresponding to the post-initiator human actions						
Section 5.2.3.1	1	Include HFEs for Needed Human Actions in the PRA Model	HR-F1	HR-16		

Table A-1
Summary of HRA Good Practices Audit Against ASME RA-Sa-2003 and NEI 00-02

Location	Good Practice	Description	ASME RA-Sa-2003	NEI 00-02	RG 1.200	Notes
Section 5.2.3.2	2	Define the HFEs Such that they are Plant and Accident Sequence-Specific	HR-F2	HR-11 HR-16 HR-17 HR-19 HR-20		
5.2.3.2	3	Perform Talk-Throughs, Walkdowns, Field Observations, and Simulator Exercises (as necessary) to Support the Modeling of Specific HFEs	HR-E3 HR-E4 HR-G5	TBD		
Quantifying the corresponding HEPs for the specific post-initiator HFEs						
Section 5.3.3.1	1	Address Both Cognitive and Response Execution Failures	HR-G2	HR-2 HR-11	qualification to NEI 00-02	self-assessment needs to document if cognitive and execution errors are included
Section 5.3.3.2	2	Use Screening Values During the Initial Quantification of the Pots-Initiator HFEs	HR-G1	HR-13 HR-15 HR-17 HR-18		
Section 5.3.3.3	3	Perform Detailed Assessments of Significant Post-Initiator HFEs.	HR-G1	HR-15 HR-17 HR-18		
Section 5.3.3.4	4	Revisit the Use of Post-Initiator Screening Values vs. Detailed Assessments For Special PRA	Section 3	not addressed		Section 3 of the ASME Standard provides detailed discussion about the risk assessment application process
Section 5.3.3.5	5	Account for Plant and Activity Specific PSFs in the Detailed Assessments of Post-Initiator HEPs	HR-G3 HR-G4 HR-G5	HR-16 HR-17 HR-18 HR-19 HR-20		
Section 5.3.3.6	6	Account for Dependencies Among Post-Initiator HFEs	HR-G7 HR-G8	HR-26 HR-27	clarification to ASME HR-G7	RG 1.200 added a requirement to justify multiple recovery actions

Table A-1 Summary of HRA Good Practices Audit Against ASME RA-Sa-2003 and NEI 00-02						
Location	Good Practice	Description	ASME RA-Sa-2003	NEI 00-02	RG 1.200	Notes
Section 5.3.3.7	7	Assess the Uncertainty in Mean HEP Values. Mean values for each HEP	HR-G9	not addressed		
Section 5.3.3.8	8	Evaluate the Reasonableness of the HEPs Obtained Using Detailed Assessments	HR-G6	HR-12		
Adding recovery actions to the PRA						
Section 5.4.3.1	1	Define Appropriate Recovery Actions	HR-H1 HR-H2	HR-21 HR-22 HR-23 HR-24		
Section 5.4.3.2	2	Account for Dependencies (all earlier guidance applies for recovery actions; particular attention is needed for dependencies)	HR-H3	HR-26	clarification to ASME HR-H3	RG 1.200 added a requirement to justify multiple recovery actions
Section 5.4.3.3	3	Quantify the Probability of Failing to Perform the Recovery(ies).	qunatifca-tion not explicitly addressed	TBD		
Including and modeling errors of commission (EOCs)						
Section 6.1.3.1	1	Address EOCs in Future HRAs/PRAs (Recommendation).	not addressed	not addressed		
Section 6.1.3.2	2	As a Minimum, Search for Conditions that May Make EOCs More Likely.	not addressed	not addressed		
Documenting the HRA						
Section 7.1.3.1	1	HRA documentation	HR-I1	HR-1 HR-28 HR-30	clarification to ASME HR-I1	RG 1.200 maintains that the ASME SRs pertaining to HRA "are written for CDF and not LERF"

APPENDIX B

Guidance on Consideration of Performance Shaping Factors for Post-Initiator HFEs

The following provides more detail on the performance shaping factors (PSFs) presented in Section 5.3.3.5, including some key characteristics to consider when assessing the influence of these performance shaping factors on the failure probability for a human failure event (HFE). Included are important interactions among the factors that should also be examined when assessing the holistic impact of the performance shaping factors on operator performance. These factors need to be assessed on a plant-specific and accident sequence-specific basis, considering the relevant context and the act to be performed.

It is important to re-iterate that this Appendix is written for the specific purpose of addressing post-initiator HFEs in a risk assessment for commercial nuclear power plant operations occurring nominally at full power, and for internal initiating events. However, much of it is considered useful to other modes of operation and for other industry applications such as safety assessments of chemical plants, space mission risk assessments, and others. Similarly, much of it is considered applicable for external initiating events, but it should be used with the additional context of such events in mind (e.g., shaking during a seismic event). Additionally, portions of this Appendix may be of benefit in examining human actions related to nuclear materials and safeguard types of applications.

Specific HRA methods and tools used by the industry may define and "measure" these performance shaping factors somewhat differently than described here. That is, they may use a different explicit set of performance shaping factors that 'roll-up' many of the factors listed below into the definitions of their specific factors (e.g., stress, workload). Nevertheless, these summaries are provided as one means with which to assess that the specific HRA method/tool has been used such that the characteristics described here have indeed been accounted for in the evaluation of post-initiator human error probabilities (HEPs).

While quantitative guidance is not provided (specific quantification depends on the method/tool that is used), the following should be useful in determining qualitatively whether a performance-shaping factor is a weak/strong positive, neutral (or not applicable), or negative influence, regardless of the method/tool. The method/tool being used should provide definitions and guidance for assessing PSFs qualitatively (e.g., "good', "adequate", "poor"), along with a way to interpret the results into a quantified HEP, that can be used in conjunction with this information.

The performance shaping factors are addressed below.

Applicability and suitability of training/experience. For both in-control room and local actions, this is an important factor in assessing operator performance. For the most part, in nuclear plants, operators can be considered "trained at some minimum level" to perform their desired tasks.

However, from a HRA perspective, the degree of familiarity with the type of sequences modeled in the PRA and the actions to be performed, can provide a negative or positive influence that should be used to assess the likelihood of operator success. In cases where the type of PRA sequence being

examined or the actions to be taken are not periodically addressed in training (such as covered in classroom sessions or simulated every one to two years or even more often) or the actions are not performed as part of their normal experience or on-job duties, this factor should be treated as a negative influence. The converse would result in a positive influence on overall operator performance.

One should also attempt to identify systematic training biases that may affect operator performance either positively or negatively. For example, training guidance in a pressurized water reactor (PWR) may provide a reluctance to use "feed and bleed" in a situation where steam generator feed is expected to be recovered. Other biases may suggest operators are allowed to take certain actions before the procedural steps calling for those actions are reached, if the operators are sure the actions are needed. Such training "biases" could cause hesitation and result in higher HEPs for the desired actions, as in the first case above, or as in the case of not waiting to take obvious actions, they may be a positive influence.

It is incumbent on the analyst to ensure that training and/or experience is relevant to the PRA sequence situation and desired actions. The more it can be argued that the training is current, "is like the real event," is varied enough to represent differences in the way the event can evolve, and proficiency is demonstrated on a periodic basis, the more positive this factor. If there is little or no training/experience or there are potentially negative training biases for the PRA sequence being examined, this factor should be considered to have a negative influence.

Suitability of relevant procedures and administrative controls. For both in-control room and local actions, this is an important factor in assessing operator performance. Similar to training, for the most part, procedures exist to cover many types of sequences and operator actions.

However, from a HRA perspective, the degree the procedures clearly and unambiguously address the types of sequences modeled in the PRA and the actions to be performed, dictates whether they are a negative or positive influence on operator performance. Where procedures have characteristics like those listed below related to the desired actions for the sequences of interest, this factor should be considered a negative influence:

- ambiguous/unclear/non-detailed steps for the desired actions in the context of the sequence of interest
- situations can exist where the operators are likely to have trouble identifying a way to proceed forward through the procedure
- there is a requirement to rely on considerable memory
- operators must perform calculations or make other manual adjustments especially in time-sensitive situations
- there is no procedure or the procedure is likely to not be available, especially when taking local actions "in the heat of the scenario" and it cannot be argued that the desired task is simple and a "skill of the craft" or that it is an automatic/memorized activity that is trained on and for

which there is routine experience

Otherwise, this PSF should be considered as adequate or even a positive influence.

Talk-throughs with operations and training staff (in the context of the scenario being examined) can be helpful in uncovering 'difficulties' or 'ease' in using the relevant procedures, considering the associated training that the operators receive and the way the operators interpret the use of the procedures.

Availability and clarity of instrumentation (cues to take actions as well as confirm expected plant response). For both in-control room and local actions, this is an important factor since operators, other than for immediate and memorized response actions, take actions based on diagnostic indications and look for expected plant responses to dictate follow-on actions. For in-control room situations, typical nuclear plant control rooms have sufficient redundancy and diversity for most important plant parameters. For this reason, most HRA methods inherently assume that adequate instrumentation typically exists. Nevertheless, this should be verified looking for the following characteristics that could make this a negative performance-shaping factor, particularly in situations where there is little redundancy in the instrumentation associated with the act(s) of interest:

- the key instrumentation associated with an act is adversely affected by the initiating event or subsequent equipment failure (e.g., loss of DC power causing loss of some indications, spurious or failed as a result of a hot short from a fire)

- the key instrumentation is not readily available and may not be typically scanned such as on an obscure back panel

- the instrumentation could be misunderstood or may be ambiguous because it is not a direct indication of the equipment status (e.g., PORV position is really the position of the solenoid valve and not the PORV itself)

- the instrumentation is operating under conditions for which it is not appropriate (e.g., calibrated for normal power conditions as opposed to shutdown conditions)

- there are so many simultaneous changing indications and alarms or the indication is so subtle, particularly when the time to act is short, it may be difficult to "see and pick out" the important cue in time (e.g., a changing open-close light for a valve without a concurrent alarm or other indication, finding one alarm light among hundreds).

The above also applies to local actions outside the control room, recognizing that in some situations, less instrumentation may exist (e.g., only one division of instrumentation and limited device actuators on the remote shutdown panel). However, on the positive side, local action indications often can include actual/physical observation of the equipment (e.g., pump is running, valve stem shows it is closed) that compensates for any lack of other indicators or alarms.

It is incumbent on the analyst to ensure that adequate instrumentation is available and clear so that the

operators will know the status of the plant and when certain actions need to be taken. If this is demonstrated, then this PSF would be positive. Task analysis will often facilitate determining whether the instrumentation is adequate.

Time available and time required to complete the act, including the impact of concurrent and competing activities. This can be an important influence for both in-control room and local actions since clearly, if there is not enough or barely enough time to act, the estimated HEP is expected to be quite high. Conversely, if the time available far exceeds the time required and there are not multiple competing tasks, the estimated HEP is not expected to be strongly influenced by this factor.

It is important that the time available and the time needed to perform the act be considered *in concert with* many of the other performance shaping factors and the demands of the sequence. This is because the thermal-hydraulic inputs (e.g., time to steam generator dryout, time to start uncovering the core), while important, are not the only influences. (Note, it is best if the thermal-hydraulic influences are derived from plant-specific or similar analyses rather than simple judgments).

The time to perform the act, in particular, is a function of the number of available staff, the clarity and repetitiveness of the cues that the act needs to be performed, the human system interface (HSI-discussed later), the complexity involved (discussed later), the need to get special tools or clothing (discussed later), consideration of diversions and other concurrent requirements (discussed later), where in the procedures the steps for the act of interest are called out, crew characteristics such as whether the crews are generally aggressive or slow and methodical in getting through the procedural steps (discussed later), and other potential 'time sinks'.

Clearly there is judgment involved, but as described here, it is not as simple as watching an operator perform an act in ideal conditions with a stop watch to determine the time required to perform the act. Only when the sequence context is considered holistically with the interfacing performance shaping factors that have been mentioned here, can more meaningful "times" be estimated. Hence, especially for complex acts and/or situations, walkdowns and simulations can be helpful in ensuring overly optimistic "times" have not been estimated. Whatever HRA method/tool is used, determination of these times should include the considerations provided here.

Complexity of the required diagnosis and response, the need for special sequencing, and the familiarity of the situation. This factor attempts to measure the overall complexity involved for the situation at hand and for the act itself (e.g., many steps have to be performed by the same operator in rapid succession vs. one simple skill-of-the craft action). Many of the other performance shaping factors bear on the overall complexity, such as the need to decipher numerous indications and alarms, the presence of many and complicated steps in a procedure, poor HSI, etc. Nevertheless, this factor should also capture 'measures' such as the ambiguity associated with assessing the situation or in executing the task, the degree of mental effort or knowledge involved, whether it is a multi-variable or single variable associated task, whether special sequencing or coordination is required in order for the act to be successful (especially if it involves multiple persons in different locations), whether the activity may require very sensitive and careful manipulations by the operator, etc. The more these "measures" describe an overall complex situation, this performance shaping factor should be found to be a negative influence. To the extent these "measures" suggest a simple, straightforward, unambiguous process (or

one that the crew or individual is very familiar with and skilled at performing), this factor should be found to be nominal or even ideal (i.e., positive influence).

Workload, Time Pressure, and Stress. Although these factors are often associated with complexity and can certainly contribute to perceived complexity, the emphasis here is on the amount of work a crew or individual has to accomplish in the time available (e.g., task load), along with their overall sense of being pressured and/or threatened in some way with respect to what they are trying to accomplish (e.g., see Swain and Guttmann[8] for a more detailed definition and discussion of stress and workload). To the extent crews or individuals expect to be under high workload, time pressure, and stress, it is generally thought to have a negative impact on performance (particularly if the task being performed is considered complex). However, the impact of these factors should be carefully considered in the context of the scenario and the other PSFs thought to be relevant. For example, if the scenario is familiar, the procedures and training for the scenario are very good, and the rate at which the crews normally implement their procedures will allow them to achieve their goal on time, then analysts might decide that even relatively high expected levels of workload and stress will not have a significant impact on performance. Although these factors may be difficult to measure, analyst's should demonstrate a careful evaluation of the their potential influence in the scenario being examined, before deciding on the strength of their effect.

Team/crew dynamics and crew characteristics (degree of independence among individuals, operator attitudes - biases - rules, use of status checks, approach for implementing procedures, e.g., aggressive vs. slow and methodical crew). This is a "catch-all" type of factor which can be important, especially to in-control room actions where the early responses to an event occur and the overall strategy for dealing with the event develops. In particular, the way the procedures are written and what is (or is not) emphasized in training (which may be related to an organizational influence), can cause systematic and nearly homogeneous biases and attitudes in most or all the crews that can affect overall crew performance. A review of this factor should include looking for such characteristics as:

- Are independent actions encouraged or discouraged among crew members (allowing independent actions may shorten response time but could cause inappropriate actions going unnoticed until much later in the scenario)?

- Are there common biases or 'informal rules'? For example, is there a reluctance to do certain acts, is there an overall philosophy to protect equipment or run it to destruction if necessary, or are there informal rules regarding the way procedural steps are interpreted.

- Are periodic status checks performed (or not) by most crews so that everyone has a chance to 'get on the same page' and allow for checking on what has been performed to ensure that the desired activities have taken place? In general, are there good communication strategies used to help ensure that everyone stays informed?

- Is the overall approach of most crews to aggressively respond to the event, including taking allowed shortcuts through the procedural steps (which will shorten response times), or are typical responses slow and methodical (we trust the procedures type of attitude), thereby tending to slow down response times but making it less likely to make mistakes.

In general, deciding whether the crew characteristics have a positive or negative effect will be contingent on the scenario being examined. For example, a particular bias may be very positive for some scenarios, but not for others.

Observing simulations and using talk-throughs and walkdowns can provide valuable insights into the overall crew response dynamics, attitudes, and the typical times it takes them to get through various procedure steps and deal with unexpected failures or distractions. This knowledge can be a key input into the HEP evaluation including determining the typical time to respond (see that factor above).

Available staffing/resources. For in-control room actions, this is generally not an important consideration (i.e., not a particularly positive or negative factor) since plants are supposed to maintain an assigned minimum crew with the appropriate qualified staff available in or very near the control room.

However, for ex-control room local actions, this can be an important consideration particularly dependent on (a) the number and locations of the necessary actions, (b) the overall complexity of the actions that must be taken, and (c) the time available to take the actions and the time required to perform the actions (see above for more on these related factors). For instance, where the number of actions are few and complexity is low and time available is high, one or two personnel available to perform the local actions may be more than enough and thus the available staffing can be considered to be adequate or even a positive factor. On the other hand, where the number of actions and their complexity is high, and with little time, perhaps three or more staff may be necessary. Additionally, the time of the day the initiating event occurs may be a factor since typically, night and "back" shifts have fewer people available than the day shift. If there are significant differences in the staffing levels depending on the time of day, it is advisable to either treat the staffing level in a HEP evaluation as the minimum available depending on the shift, or probabilistically account for these aleatory differences more explicitly in the PRA model

It is incumbent on the analyst to demonstrate that the available staffing is sufficient to perform the desired actions and/or assess the HEP(s) accordingly.

Human-system interface (HSI). This is generally not an important factor relative to main control room actions since, given the many control room design reviews and improvements and the daily familiarity of the control room boards and layout, problematic HSIs have been taken care of or are easily worked around by the operating crew. Of course, any known very poor HSI should be considered as a negative influence for an applicable action even in the control room. For example, if common workarounds are known to exist that may negatively influence a desired act, this should be accounted for in the HEP evaluation. Furthermore, it is possible that some unique situations may render certain HSIs less appropriate and for such sequences, the relevant interfaces should be examined.

However, since local actions may involve more varied (and not particularly "human-factored") layouts and require operators to take actions in much less familiar surroundings and situations, any problematic HSIs can be an important negative factor on operator success. For instance, if to reach a valve to open it manually requires the operator to climb over pipes and turn the valve with a tool while in a laid out position, or in-field labeling of equipment is generally in poor condition and could

lengthen the time to find the equipment, etc., such 'less ideal' HSIs could mean this is a negative performance-shaping factor. Otherwise, if a review reveals no such problematic interfaces for the act(s) of interest, this influence can be considered adequate, or even positive if the interface helps ensure the appropriate response in some way.

Walkdowns and field or simulator observations can be useful tools in discovering problems (if any exist) in the HSI for the actions of interest. Sometimes, discussions with the operators will reveal their own concerns about issues in this area.

Environment in which the act needs to be performed. Except for relatively rare situations, this factor is not particularly relevant to in-control room actions given the habitability control of such rooms and the rare challenges to that habitability (e.g., control room fire, loss of control room ventilation, less lighting as a result of station blackout). However, for local actions, this could be an important influence on the operator performance. Radiation, lighting, temperature, humidity, noise level, smoke, toxic gas, even weather for outside activities (e.g., having to go on a potential snow-covered roof to reach the atmospheric dump valve isolation valve), etc., can be varied and far less than ideal. Hence any HEP assessment should ensure that the influence of the environment where the act(s) needs to take place is accounted for as a performance-shaping factor. This factor may be non-problematic (adequate) or a negative influence (even to the point of not being able to perform the act).

Accessability and operability of the equipment to be manipulated. As with the environment factor, this factor is not particularly relevant most of the time to in-control room actions except for special circumstances such as loss of operability of indications or controls as a result of the initiator or equipment failures (e.g., loss of DC). However, for local actions, accessability and the operability of the equipment to be manipulated may not always be ensured, and needs to be assessed in the context with such influences as the environment, the need to use special equipment (discussed later), and HSI. Hence any HEP assessment should ensure that this factor, for where the act(s) needs to take place, is accounted for as a performance-shaping factor. This factor may be non-problematic (adequate) or a negative influence (even to the point of not being able to perform the act).

The need for special tools (keys, ladders, hoses, clothing such as to enter a radiation area). As for the environment and accessability factors, this factor is not particularly relevant to in-control room actions with the common exception of needing keys to manipulate certain control board switches or similar controls (e.g., key for explosive valves for standby liquid control injection in a BWR). However, for local actions, such needs may be more commonplace and necessary in order to successfully perform the desired act. If such equipment is needed, it should be ensured that the equipment is readily available, its location is readily known, and it is either easy to use or periodic training is provided, in order for this factor to be considered to be adequate. Otherwise, this factor should be considered to have a negative influence on the operator performance, perhaps even to the point of making the failure of the desired action very high.

Communications (strategy and coordination) as well as whether one can be easily heard). For in-control room actions, this factor is not particularly relevant although there should be verification that the strategy for communicating in the control room is one that tends to ensure that directives are not easily misunderstood (e.g., it is required that the board operator repeat the act to be performed and then

wait for confirmation before taking the act). Generally, it is expected that this will not be problematic; but any potential problems in this area (such as having to talk with special air packs and masks on in the control room in a minor fire) should be accounted for if they exist.

For local actions, this factor may be much more important because of the possible less than ideal environment or situation. It should be assured that the initiating event (e.g., loss of power, fire, seismic) or subsequent equipment faults are not likely to negatively affect the ability for operators to communicate as necessary to perform the desired act(s). For instance, having to set up the equipment and talk over significant background noise and possibly having to repeat oneself many times should be a consideration - even if just as a possible 'time sink' for the time to perform the act. Additionally, there should be training on the use of the communication equipment, its location should be readily known, and its operability periodically demonstrated and shown to be in good working condition. Depending on the status of these characteristics, this factor may be non-problematic (adequate) or a negative influence (even to the point of not being able to perform the act).

Special fitness needs: While typically not an issue for in-control room actions, this could be an important factor for a few local actions depending on the specific activity involved. Having to climb up or over equipment to reach a device, needing to move and connect hoses, using an especially heavy or awkward tool, are examples of where this factor could have some influence on the operator performance. In particular, the response time for an action may be increased for successful performance of the act. Physically demanding (or not) activities should be considered in the evaluation of any HEP where it is appropriate to do so. Talk-throughs or field observations of the activities involved can help determine whether such issues are relevant to a particular HFE.

Consideration of 'realistic' accident sequence diversions and deviations (e.g., extraneous alarms, outside discussions, or the sequence evolution is not exactly like that trained on). Particularly for in-control room actions where the early responses to an event occur and the overall strategy for dealing with the event develops, this can be an important factor to be considered. Through simulations, training, and the way the procedures are written, operators 'build up' some sense of expectations as to how various types of sequences are likely to proceed; even to the extent of recognizing alarm and indication patterns and what actions will likely be appropriate. To the extent the actual sequence may not be 'just like in the simulator,' such as involving other unimportant or spurious alarms, the need for outside discussions with other staff or even offsite entities such as a fire department, differences in the timing of the failed events, and behavior of critical parameters, etc., all can add to the potential diversions and distractions that may delay response timing or in the extreme, even confuse the operators as to the appropriate actions to take.

Hence, the 'signature' of the PRA accident sequence and the potential acts of interest should be examined against the expectations of the operators to determine if there is a considerable potential for such distractions and deviations. Observing simulations and talking with the operators can help in discovering such possibilities. This could impact the HEP mean value estimate as well as the uncertainty in the HEP, which may be important to assessing the potential risk or in establishing the limits for doing sensitivity studies with the HEP.

NRC FORM 335 (2-89) NRCM 1102, 3201, 3202

U.S. NUCLEAR REGULATORY COMMISSION

BIBLIOGRAPHIC DATA SHEET

(See instructions on the reverse)

1. REPORT NUMBER (Assigned by NRC, Add Vol., Supp., Rev., and Addendum Numbers, if any.)

NUREG-1792

2. TITLE AND SUBTITLE

Good Practices for Implementing Human Reliability Analysis (HRA)
Draft Report for Comment

3. DATE REPORT PUBLISHED

MONTH	YEAR
July	2004

4. FIN OR GRANT NUMBER

5. AUTHOR(S)

Alan Kolaczkowski, Science Applications International Corp.
John Forester, Sandia National Laboratories
Erasmia Lois, Nuclear Regulatory Commission
Susan Cooper, Nuclear Regulatory Commission

6. TYPE OF REPORT

Technical

7. PERIOD COVERED *(Inclusive Dates)*

n/a

8. PERFORMING ORGANIZATION - NAME AND ADDRESS *(If NRC, provide Division, Office or Region, U.S. Nuclear Regulatory Commission, and mailing address; if contractor, provide name and mailing address.)*

Division of Risk Analysis and Applications
Office of Nuclear Regulatory Resrach
U.S. Nuclear Regulatory Commission
Washington, DC, 20555-0001

Sandia National Laboratories
P.O. Box 5800, MS0748
Albuquerque, NM 87185

Science Applications International Co.
405, Uban St., Suite 400
Lakewood, Co, 80220

9. SPONSORING ORGANIZATION - NAME AND ADDRESS *(If NRC, type "Same as above"; if contractor, provide NRC Division, Office or Region, U.S. Nuclear Regulatory Commission, and mailing address.)*

As above

10. SUPPLEMENTARY NOTES

E. Lois, NRC Project Manager

11. ABSTRACT *(200 words or less)*

The U.S. Nuclear Regulatory Commission is establishing "good practices" for performing human reliability analysis (HRA) and reviewing HRAs to assess the quality of analyses. The HRA good practices are developed as part of the NRC's activities for addressing probabilistic risk assessment (PRA) quality issues and supports the implementation of Regulatory Guide (RG) 1.200 entitled: "An Approach for Determining the Technical Adequacy of Probabilistic Risk Assessment Results For Risk-Informed Activities."

The documented HRA good practices are of generic nature, that is, are not tied to any specific methods or tools that could be employed to perform an HRA. The report provides low level guidance for implementing the RG 1.200 when performing a Level 1 and a limited Level 2 PRA for internal events (excluding fire) with the reactor at full power. Its elements are directly linked to RG 1.200 which reflects and endorses, with certain clarifications and substitutions, the American Society of Mechanical Engineers (ASME) Standard RA-S-2002, "Standard for Probabilistic Risk Assessment for Nuclear Power Plant Applications," and Revision A3 of the Nuclear Energy Institute (NEI) "Probabilistic Risk (PRA) Peer Review Process Guidance," (NEI-00-02).

12. KEY WORDS/DESCRIPTORS *(List words or phrases that will assist researchers in locating the report.)*

Human Reliability Analysis
HRA
Probabilistic Risk Assessment
PRA
good practices
HRA guidance
HRA review guidance

13. AVAILABILITY STATEMENT

unlimited

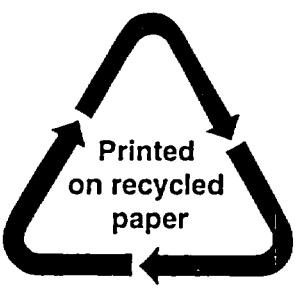
14. SECURITY CLASSIFICATION

(This Page)
unclassified

(This Report)
unclassified

15. NUMBER OF PAGES

16. PRICE



Federal Recycling Program

UNITED STATES
NUCLEAR REGULATORY COMMISSION
WASHINGTON, DC 20555-0001

OFFICIAL BUSINESS