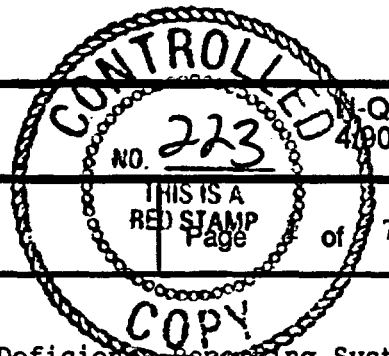


INTERIM CHANGE NOTICE



QA-023

ICN Number:

1

Effective Date:

10/19/90

Applies to:

Number OMP-16-03 Rev. 1 Title Standard Deficiency Reporting System

REQUIRED CHANGE(S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

Entire procedure

Change Project Office QA, Project Office organization, and Project Office personnel to Project Office QA staff.

Entire procedure

Change Project Office PQM, Project Quality Manager (PQM), PQM, and QA Division Manager to DOE Director, QA/designee.

Para. 2.0

Change the third sentence to read:

The deficiencies may be identified by Yucca Mountain Project Office (Project Office) staff personnel during the performance . . .

Para. 4.2

Add the following:

The Project Office QA Staff is also responsible for tracking all SDRs initiated via a Standard Deficiency Report Log, assuring committed corrective actions have been properly implemented and reporting the status of SDRs.

Para. 4.4

Delete entirely.

Para. 5.1.1.1.2

Change Branch Chief to Division Director and delete "or Department Manager" in the last sentence.

Para. 5.1.1.1.4

Change Branch Chief to Division Director and delete "or Department Manager."

Para. 5.1.1.2

Change QA Administrative Assistant to Project Office QA Staff.

Para. 5.2.1.1

Delete "responsible QA Manager," from the fourth sentence.

Para. 5.2.2

Delete the phrase "obtained from the Project Office QA Administrative Assistant" from the first sentence.

APPROVALS

Division Director

Director, QA

Project Manager

Date

N/A

Date

10/18/90

Date

10/19/90

9011010204 901026
PDR WASTE
WM-11 PDC

**INTERIM CHANGE NOTICE
CONTINUATION PAGE**

N-QA-023
4/90

ICN Number: 1	Applies to: QMP-16-03	Effective Date: 10/19/90	Page 2 of 7
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REQUIRED CHANGE (S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

Para. 5.2.2 (con't)	Combine the second and the last sentences to read "Once the draft SDR is finalized, the responsible Quality Assurance Engineer (QAE) or Lead Auditor signs and dates the SDR in block 11 and obtains the approval of the DOE QA Director/designee."
Para. 5.2.3	Change the first sentence to read "If the DOE QA Director/designee does not approve . . ."
Para. 5.3.1	Delete the second paragraph entirely.
Para. 5.3.2	Change QA Verification Department to Project Office QA Staff.
Para. 5.3.2.1	Change QA Verification Department to Project Office QA Staff.
Para. 5.4.2	Delete the words "the respective QA Division Manager and" from the second sentence.
Para. 5.4.5.2	Change to read "Block 8 shall cite the QARD as the violated requirement."
Para. 5.5.1.1	Delete "the respective QA Division Manager and" from the second sentence.
Para. 5.5.2.1	Delete "the responsible QA Division Manager and."
Para. 5.6	Delete "the appropriate QA Division Manager;" change signatures to signature in the first sentence. Change the second sentence to read "A notation of closure shall be made on the SDR log."
Para. 5.8.1	Change QA Verification Department Manager to Project Office QA Staff in the second sentence.
Figure 1 (Instructions)	Change Branch Chief/Department Manager to DOE Division Director in Block 1. (See revised Deficiency Evaluation Report (DER) Format Sheet and Completion Instructions. on page 4 of 7).

**INTERIM CHANGE NOTICE
CONTINUATION PAGE**

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4/90

ICN Number: 1	Applies to: QMP-16-03	Effective Date: 10/19/90	Page 3 of 7
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REQUIRED CHANGE (S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

Figure 2
(Instructions)

Delete "from the QA Administrative Assistant" from the instructions for Block 4.

Delete Block 12 from the instructions and renumber accordingly.

Delete "and responsible Division Manager" from Block 18.

(See revised Yucca Mountain Project Office Standard Deficiency Report Format Sheet and Completion Instructions on page 5 of 7.)

Figure 4
(Instructions)

Delete "from QA Administrative Assistant" from the instructions for Block 1.

Change Branch Manager to DOE Director, Quality Assurance Department in Block 10.

(See revised Revised Yucca Mountain Project Office Observation Form and Completion Instructions on page 6 of 7).

Figure 5
(Instructions)

Delete the block for the Signature/Date of the QA Division Manager.

(See revised SDR Severity Level Checklist on page 7 of 7).

INTERIM CHANGE NOTICE CONTINUATION PAGE

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4/90

ICN Number: 1	Applies to: QMP-16-03	Effective Date: 10/19/90	Page 4 of 7
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REQUIRED CHANGE (S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

YUCCA MOUNTAIN PROJECT Deficiency Evaluation Report		N-QA-013 9/90
1	To _____	
	Initiator _____	Date _____
	DOE Division Director _____	Date _____
2	Requirement	
3	Deficiency	
4	Evaluated By _____ Date _____	
	Surveillance Report No. _____	
5	Approved By _____ Date _____	
	DOE Director, Quality Assurance	
	SDR No. Issued _____	
6	Disapproved By _____ Date _____	
	DOE Director, Quality Assurance	
	Reason	

Figure 1. Revised Deficiency Evaluation Report (DER) Format Sheet.

INTERIM CHANGE NOTICE CONTINUATION PAGE

N-QA-023
4/90

ICN Number: 1	Applies to: QMP-16-03	Effective Date: 10/19/90	Page 5 of 7
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REQUIRED CHANGE (S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

YMPO STANDARD DEFICIENCY REPORT			N-QA-038 8/90
Completed by Originating QA Organization	1 Date		2 Severity Level <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3
	3 Discovered During		4 SDR No. _____
	3a Identified By		Rev. _____
	5 Organization	6 Person(s) Contacted	
	7 Response Due Date is 20 Working Days from Date of Transmittal		
Completed by Organization in Block 5	8 Requirement (Audit Checklist Reference, if Applicable)		
	9 Deficiency		
	10 Recommended Action(s): <input type="checkbox"/> Remedial <input type="checkbox"/> Investigative <input type="checkbox"/> Corrective		
	11 QAE/Lead Auditor/Date		12 DOE Director, Quality Assurance/Date
	13 Remedial/Investigative Action(s)		14 Effective Date _____
Comp. by Org. QA Org.	15 Cause of the Condition & Corrective Action to Prevent Recurrence		16 Effective Date _____
	17 Signature/Date		
	18 Response Accepted	QAE/Lead Auditor/Date	DOE Director, Quality Assurance/Date
	19 Corrective Action Verif. Satisfactory	QAE/Lead Auditor/Date	DOE Director, Quality Assurance/Date
	20 Remarks		
21 QA CLOSURE		QAE/Lead Auditor/Date	DOE Director, Quality Assurance/Date

Figure 2. Revised Yucca Mountain Project Office Standard Deficiency Report Format Sheet.

INTERIM CHANGE NOTICE CONTINUATION PAGE

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4/90

ICN Number: 1	Applies to: QMP-16-03	Effective Date: 10/19/90	Page 6 of 7
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REQUIRED CHANGE (S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

YUCCA MOUNTAIN PROJECT OFFICE YMPO OBSERVATION NO. _____		N-QA-012 8/90
Completed by Originating Organization	2 Noted During:	3 Identified By:
	4 Date:	
	5 Organization:	6 Person(s) Contacted:
	7 Response Due Date is 20 Days from Date of Transmittal	
Completed by Respondee	8 Discussion:	
	9 QAE/Lead Auditor Date	10 DOE Director, Quality Assurance Date
Completed by QA Org.	11 Response:	
	12 Signature: Date:	
Completed by QA Org.	13 Response Receipt Acceptable <input type="checkbox"/>	
	Initiator Date	QA/Lead Auditor Date
14 Remarks:		
		Page ____ of ____

Figure 4. Revised Yucca Mountain Project Office Observation Form.

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4/90

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REQUIRED CHANGE (S): (Minor ☐ Yes ☒ No)

PARAGRAPH

CHANGE TO

SDR SEVERITY LEVEL CHECKLIST		N-QA-037 8/90																								
<p>I. ASSIGN A SEVERITY LEVEL OF 1 IF ONE OR MORE OF THE FOLLOWING IS TRUE.</p> <table style="width: 100%;"> <thead> <tr> <th></th> <th style="text-align: center;">Yes</th> <th style="text-align: center;">No</th> </tr> </thead> <tbody> <tr> <td>1. Did the deficiency result in significant damage to natural barriers, structures, systems, or components that will require extensive evaluation, extensive redesign, or extensive repair in order to assure public health and safety?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>2. Does the deficiency involve loss of essential data or information needed for licensing?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>3. Does the deficiency constitute a significant deficiency in design, construction, testing, or performance assessment that were detected subsequent to formal quality verification and acceptance?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>4. Does the deficiency constitute a significant deficiency in design as approved for construction such that the design deviates extensively from design criteria and bases?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>5. Does the deficiency constitute a significant deviation from performance objectives or specifications that will require extensive evaluation, extensive redesign, or extensive repair to establish the adequacy of a natural barrier, structure, system, or component to meet design criteria and bases?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>6. Does the deficiency constitute a significant error detected in a computer program after it has been released for use?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>7. Does the deficiency constitute a significant breakdown in a participant's QA program and/or repetitive, programmatic and hardware deficiencies for which previous corrective action has not been reasonably prompt or effective?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> </tbody> </table>				Yes	No	1. Did the deficiency result in significant damage to natural barriers, structures, systems, or components that will require extensive evaluation, extensive redesign, or extensive repair in order to assure public health and safety?	—	—	2. Does the deficiency involve loss of essential data or information needed for licensing?	—	—	3. Does the deficiency constitute a significant deficiency in design, construction, testing, or performance assessment that were detected subsequent to formal quality verification and acceptance?	—	—	4. Does the deficiency constitute a significant deficiency in design as approved for construction such that the design deviates extensively from design criteria and bases?	—	—	5. Does the deficiency constitute a significant deviation from performance objectives or specifications that will require extensive evaluation, extensive redesign, or extensive repair to establish the adequacy of a natural barrier, structure, system, or component to meet design criteria and bases?	—	—	6. Does the deficiency constitute a significant error detected in a computer program after it has been released for use?	—	—	7. Does the deficiency constitute a significant breakdown in a participant's QA program and/or repetitive, programmatic and hardware deficiencies for which previous corrective action has not been reasonably prompt or effective?	—	—
	Yes	No																								
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<p>II. ASSIGN A SEVERITY LEVEL OF 2 IF THE ANSWERS TO ALL QUESTIONS IN PART I ARE NO AND ONE OR MORE OF THE FOLLOWING IS TRUE:</p> <table style="width: 100%;"> <thead> <tr> <th></th> <th style="text-align: center;">Yes</th> <th style="text-align: center;">No</th> </tr> </thead> <tbody> <tr> <td>1. Could failure to correct deficiency have a potentially adverse impact on the health or safety of operations personnel?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>2. Does the deficiency constitute operating outside the scope of the quality program or approved quality procedures where both remedial and corrective actions are required?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> <tr> <td>3. Does the deficiency constitute a repetitive hardware deficiency for which no previous corrective action measures exist?</td> <td style="text-align: center;">—</td> <td style="text-align: center;">—</td> </tr> </tbody> </table>				Yes	No	1. Could failure to correct deficiency have a potentially adverse impact on the health or safety of operations personnel?	—	—	2. Does the deficiency constitute operating outside the scope of the quality program or approved quality procedures where both remedial and corrective actions are required?	—	—	3. Does the deficiency constitute a repetitive hardware deficiency for which no previous corrective action measures exist?	—	—												
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3. Does the deficiency constitute a repetitive hardware deficiency for which no previous corrective action measures exist?	—	—																								
<p>III. ASSIGN A SEVERITY LEVEL OF 3 IF THE ANSWERS TO ALL QUESTIONS TO PARTS I AND II ARE NO.</p>																										
QAE/Lead Auditor		DOE Director, Quality Assurance																								
Signature/Date		Signature/Date																								

Figure 5. Revised SDR Severity Level Checklist.