TMSS/100 8/30/91 REVISION NO.:	2	[x]STA	& MANAGEMENT SUPPORT SE NDARD PRACTICE PROCED N PROCEDURE [] WORK	URE	WBS: <u>12914</u> QA: <u>QA</u>				
Title RECORDS MANAGEMENT: RECORD SOURCE IMPLEMENTATION									
Procedure Numbe SP 1.36	r Revision Number 7	ICN Number N/A	Check as [] Complete Revision [] N/ Appropriate [X] Revision Bars	(Rev. 0) Effective Date 3/2/92	Page 1 Of 24				
I have read, unders	tood, and complied with S	P1.1, Iha ny Rev	piect Manager (SPs) Date Date 2-28-52 ave read, understood, and complied with SP 1.1, v ICN # in accomplishing my ponsibilities in this procedure.	QA Manager (SPs(QPs/VAs) // Left B / D / MM I have read, understood, and co Rev ICN # in acc responsibilities in this procedure	mpiled with SP 1.1, complishing my				
1.0	PURPOSE			· · · · · · · · · · · · · · · · · · ·					
This procedure describes the methods for identifying and preparing Yucca Mountain Site Characterization Project (YMP) records for submission to the Las Vegas Local Records Center (LRC) operated by the Management & Operations Contractor. (Hereinafter, the "Las Vegas LRC" shall be referred to as "LRC.)									
, 2.0	SCOPE								
2.1	maintain, or	retrieve	s to T&MSS personnel who cro records while performing wo gram Description (QAPD).						
2.2		ate, sub	pplies to all T&MSS particip mit, maintain, or retrieve p work.						
2.3	generate, sub	mit, mai	s to subcontractors and othe ntain, or retrieve records a ty Assurance Program Descrip	while performing w					
3.0	REFERENCES AN	D DEFINI	TIONS						
3.1	REFERENCES								
	3.1.1 Record	ls Manage	ment Policies and Requirement	nts, DOE/RW-0194					
	3.1.2 Record	ls Manage	ment Plan, YMP/CC-0016, Rev	. 3					
			ons and Lettering From the Entation Practices, ANSI Y 14		y and				
I	3.1.4 Projec	t Glossa	ry, YMP-89-15						
			Management Support Services ption (QAPD), SAIC-90/8002	Quality Assurance					
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:	3.1.6	Contro 5635.1		ssified I	Ocuments and	Information,	DOE Order			
3	3.1.7	Record	s Disposi	ition, DO	DE 1324.2A					
3.2 1	EFINI	TIONS								
:	8.2.1	The definitions of Standard Terms may be found in the T&MSS QAPD and the Project Glossary referenced in 3.1.3.								
-	only and o form organ docum indiv legit sourc			Authentication - Records shall be considered valid QA records only if stamped, initialed, signed, or otherwise authenticated and dated by authorized personnel. Authentication may take the form of a statement by the responsible individual or organization. Handwritten signatures are not required if the document is clearly identified as a statement by the reporting individual or organization. The records may be originals or legible copies. When a QA record is authenticated by a record source it means that the record is validated as accurate, legible, and completed appropriate to the work accomplished.						
. 3	suffic by the copies data a facili			bual Storage - Two copies of records are stored in areas that are sufficiently remote from each other that they cannot be destroyed by the same disaster. The order, format or condition of those copies does not have to be exact as long as the information or lata are the same. The Valley Bank Center Tower and the garage facilities are considered sufficiently remote from each other that they can provide dual storage facilities for records.						
3	be det		Legibility - Is a characteristic of a record that enables it to be read or the contents deciphered. If all the contents can be determined, either directly or through deduction, then the record shall be considered legible.					n be		
3	3.2.5	as sta	ted in th	ne Projec	d which meets t Glossary, 1 information	but does not		cord		
3	3.2.6				ersized record ter than 14 in		s that have	a		
:	3.2.7	alpha- record is the	numeric o s package three-di	code used identif igit alpl	fier - A reco to identify fier is written a code used to QRP for Qual	records pack en as "X. to identify t	ages. The X.X", where he type of	F. 11		

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TECHNICAL & MANAGEMENT SUPPORT SERVICES [x] STANDARD PRACTICE PROCEDURE [] ORGANIZATION PROCEDURE [] WORK INSTRUCTION (CONTINUATION PAGE)

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for Records Turnover Package [non-quality related materials]) and "X" is the Work Breakdown Structure (WBS) number assigned to the record package that corresponds to the activity.

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QA: ____A

- 3.2.8 Records Package Segment A records package segment is a record or a group of records that shall be included as part of a record package.
- 3.2.9 Records Package Tracking Number A records package tracking number is a number used by the LRC and the record source(s) to track and control records package segment collection and the building of records packages in the LRC. This number becomes obsolete upon completion of the records package, and is not to be entered into the records system.
- 3.2.10 Record Source A record source, as used in this procedure, is an individual working under the T&MSS QAPD, including subcontractors or other contractors, who is responsible for generating YMP records, or receiving YMP records from an entity within or from outside the YMP.
- 3.2.11 Special Process Records Special process records are records that cannot be microfilmed, but can be duplicated. These records (e.g., magnetic tapes, film, etc.) may be duplicated and stored in dual storage or a one-hour fire-rated safe.

4.0 BACKGROUND

This procedure applies to record sources. It provides step by step instructions that will enable record sources to meet the requirements that apply to creating, maintaining, protecting, submitting, and retrieving YMP records. In addition to the responsibilities of record sources, Assistant Project Managers (APMs), or their designees, have responsibilities assigned in this procedure.

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SP 1.36	7	N/A	Appropriate [X] Revision Bars 3/2/92 Page 4 24
5.0 PH	ROCEDURE		
5.1 II	DENTIFICATIO	N AND PR	EPARATION OF RECORDS
5.1.1	Identificat	ion of R	ecords
RESPONS	SIBILITY		ACTION
APM (or des	5 signee)	5.1.1.1	Ensure, by review, that design specifications, procurement documents, task plans, study plans, test procedures, implementing procedures, instructions, or other documents directing the conduct of quality related activities identify the records and/or records packages to be generated, supplied or maintained.
			NOTE: For examples of documents that are not records refer to Exhibit 1.
	5	5.1.1.2	Identify a record source responsible for collecting, authenticating (if applicable), and submitting the records and records packages for activities within a specific area of responsibility.
Record	Source 5	.1.1.3	Complete steps 5.1.1.4 and 5.1.1.5 if documents to be generated are to be part of a records package; otherwise go to step 5.1.1.6.
	5	5.1.1.4	If the LRC is to maintain the records package, contact the LRC to obtain a records package tracking number and provide the LRC with the following information.
			a. A records package title,
			b. A records package identifier,
			c. A record source name and organization, and
			d. A quality-affecting designation (i.e., QA or QA: NA).

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	TMSS/100 8/30/91	TE		& MANAGEMENT SUPPORT SERVICES WBS: 12914		
	REVISION NO.: 2			NDARD PRACTICE PROCEDURE		
	l	[]ORG <i>P</i>	ANIZATION	N PROCEDURE [] WORK INSTRUCTION		
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	RESPONS	SIBILITY _	<u> </u>	ACTION -		
	Record	Source 5	5.1.1.5	If the record source is to maintain the records package until complete, then ensure the following:		
				a. Store non-QA records or documents that will become QA records so that damage will not occur due to excessive (1) light, stacking, and electromagnetic fields (in the case of magnetic tapes, store in accordance with manufacturer's specifications) and (2) temperatures and humidity. Also prevent loss of documents by maintaining access controls.		
				NOTE: If at anytime records, or documents destined to become records, cannot be properly stored and protected by the record source, submit those records to the LRC in advance of records package completion.		
				b. Completed QA, records when not in actual use by record sources, shall be stored in a one-hour fire-rated container bearing a U.L. label or certified by a person competent in the field of fire protection, or dual storage shall be provided.		
		·		NOTE: When providing dual storage, records must be protected and stored in separate locations as a safeguard against both copies being destroyed in the same fire, flood, or similar event.		
		5	5.1.1.6	Generate or receive documents that will become records. Provide adequate care, as identified in Step 5.1.1.5.a and Step 5.1.1.5.b.		
	ľ			NOTE: When materials received from outside entities become records or part of a record package, they shall be prepared and submitted to the LRC in accordance with this procedure.		

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Procedure N SP 1		Revision Number 7	ICN Number N/A	Check as [] Complete Revision [] N/A (Rev. 0) Effective Date Appropriate [X] Revision Bars 3/2/92 Page 01
		Preparation	of Hard	Copy (Paper) Records
<u>R</u> J	ESPONS	IBILITY		ACTION
Re	ecord	Source 5	.1.2.1	Prepare documents generated or received that will become records according to the following requirements for completeness and legibility.
				NOTE: Step 5.1.2.1 applies to all record types. For additional requirements for specific types of records (e.g., magnetic media, microform, and one-of-a-kind) go to Section 5.1.3.
		5	.1.2.2	Complete the following for all final technical and scientific reports:
				a. Contact the LRC and obtain a pre-assigned accession number.
				NOTE: Journal Articles, bulletins, or professional papers generated as a result of YMP activities must be submitted to the LRC for processing, but are not required to receive a pre-assigned accession number.
				b. Place the pre-assigned accession number on the inside back cover or within the acknowledgment section.
				c. The designation "readily available" may be printed in lieu of the accession number. Contact the LRC to obtain the accession numbers for cited references which have already been submitted to the RIS, or if unsure if the reference has been submitted previously.
				d. Indicate an accession number (placed within the citation) for all cited references except for references that are readily available to the general public (i.e. dictionaries, codes and regulations, etc.).

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RESPON	SIBILITY	<u></u>	ACTION
Record	Source !	5.2.2.2	 Submit cited references that have not been previously submitted in order to obtain an accession number.
	:	5.1.2.3	Clearly mark DRAFT on the first page of draft documents.
÷	ļ	5.1.2.4	Place the correct WBS number to at least the third level, and quality-affecting designation of QA or QA:NA in the upper right portion of the first page of the record.
			NOTE: Identify the WBS to highest level possible to identify the activity and place the decimal in the appropriate place.
•	5	5.1.2.5	Ensure corrections to documents that will become records are authenticated by an authorized individual or organization as follows:
			a. Line through the incorrect information with black ink.
			b. Place the correct information (if applicable) in close proximity of the crossed out information.
			c. Initial and date the correction. Insertions and write overs are considered corrections.
			d. Documents being reviewed with suggested changes, line-outs, deletions and writing in margins are not considered corrections and do not have to be dated and initialed.
	5	5.1.2.6	On the Transmittal/Receipt Acknowledgment, Form TMSS/010 (Exhibit 2) and the table of contents (Exhibit 3), the titles of the document must concisely identify and describe the contents of the document in order to enable future identification and timely retrieval.

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	Procedure Number SP 1.36	Revision Number 7	ICN Number N/A	er Check as [] Complete Revision [] N/A (Rev. 0) Effective Date Appropriate [] Revision Bars 3/2/92 Page 0 2	
		SIBILITY		ACTION	<u> </u>
	· <u>····································</u>		5.1.2.7		
				attachments or enclosures.	
				NOTE: If an attachment or enclosure is not included because it is not record material or will be later submitted as part of a record package, state this where the enclosures or attachments are listed. If	
				attachment has been previously submitted and processed, identify an accession number or a document identification number.	
		5	5.1.2.8	Verify that no portions of a page are missing due to tearing or folding of record edges, and that no information is obliterated.	
/			5.1.2.9	If the original hardcopy is not available for processing, the copy submitted for processing shall emulate the original to the degree possible. The copy image must be aligned properly on the page.	
		5	5.1.2.10	Colored paper shall not be submitted as a recording medium. The record shall be copied to a first generation copy; use "auto contrast" button on copier if available.	
	·			NOTE: The only exception to this requirement is oversize records that are of a color that can be filmed on a 35-mm planetary camera for aperture card production. These exceptions require special handling and require approval by the CRF manager prior to submittal for microfilming.	
		5	5.1.2.11	Ensure that the documents are legible, including photo reductions. If a record is not legible, then one of the following actions is required:	
1				NOTE: For records that are to be microfilmed, the microfilm image becomes the official program record, therefore the microfilm image must be legible. To produce a legible microfilm image, the hard copy record must have a sufficient contrast	
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	RESPONS	SIBILITY	<u>.</u>	ACTION
	Record	Source	5.1.2.1	NOTE: between the data on the record and the background media. To assist in determining sufficient contrast, copy the record at normal settings and if the copy is legible, the record will meet the legibility requirements.
				a. Determine if information can be deduced through the context in which the information is used or through reference numbers such as form numbers and purchase order numbers that can lead to the discovery of the information from within the same record or record package.
·				b. Regenerate or obtain a legible copy.
				c. Correct the record by enhancing or transcribing the illegible information. The enhancement or transcription is considered a correction which must be processed in accordance with procedures which provide for appropriate review or approval by the originating organization. The correction shall include the date and the identification (signature or initials) of the person authorized to issue such corrections.
				d. If the record cannot be corrected or regenerated, documentation must be provided stating the impact of the lost information, due to being illegible, on future, in-process, or completed work. This documentation must be signed and dated by the individual responsible for the record and approved by the record sources supervisor. This documentation must accompany the deficient record to the LRC. Use of a "Best Available Copy" stamp is inappropriate.
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/	TMSS/100 8/30/91 REVISION NO.: 2		[_x] STA	& MANAGEMENT SUPPORT SERVICES NDARD PRACTICE PROCEDURE N PROCEDURE [] WORK INSTRUCTION (CONTINUATION PAGE)
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	RESPONS	SIBILITY		ACTION -
	Record	Source 5	5.1.2.12	Record data on records and drawings in black ink against a light background.
				NOTE: Information recorded on certain data records may be accepted in other than black ink. Such uses shall be handled on a case-by-case basis and approved in advance. If appropriate, blanket approval may be obtained from the YMPO.
		5	5.1.2.13	Complete data to be recorded on drawings in accordance with American National Standards Institute (ANSI) Y 14.2.
		5	5.1.2.14	Create blackline drawings as opposed to blueprint or sepia copies.
		. 5	5.1.2.15	Use only stamps or other marks that do not intersect and obliterate the text, and do not write extraneous (unrelated) information on the documents.
		5	5.1.2.16	Fill in all applicable blanks on the documents, including signature, or enter NA, unless the record clearly states that given a certain response only a portion of the record needs to be completed or the individual responsible for the record or record package states that having reviewed the record it is determined that all the blanks are intentional. This statement must be signed and dated by the individual. Forms that go through a phased approach in completing should only be submitted when all parts have been completed (e.g., QA deficiency reports, document review sheets, etc.).
				NOTE: As an alternative to writing N/A in each blank space, N/A may be placed in the first space and an arrow drawn down through all following spaces, which are also N/A. The arrow shall stop where information other than N/A is applicable.
		5	5.1.2.17	Avoid folding oversized documents (i.e., minimum dimension greater than 14").
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	SP 1.36	<u> </u>	N/A	
		Preparation	n of Magne	
	RESPON	SIBILITY		ACTION
	Record	Source S	5.1.3.1	Convert all magnetic media (excluding video and audio tapes) to 9 track magnetic tape reels that are new or recertified and have been rewound under controlled tension.
				NOTE: Media must be tested and certified no more than 6 months before using them to record information designed for permanent retention. Generate two copies for submittal to LRC.
	I	5	5.1.3.2	Media must be written in ASCII, with all extraneous control characters removed from the data (except record length indicators for variable length records or marks designating a datum word, field, block, or file), and blocked no higher than 30,000 bytes per block, at 800, 1600, or 6250 bpi. Labeled externally to include the following:
				a. The name of the organizational unit responsible for the data.
				b. File title(s)
				c. Dates of creation and coverage
				d. The recording density
				e. Types of internal labels
				f. If applicable, data set name(s), volume serial number, number of tracks, character code/software dependency, record length, block size, and reel sequence (if the file is part of a multi-reel set).
				g. Verified as error-free.
		5	5.1.3.3	Machine-readable files, which have been designated for preservation and maintained on a direct access storage device, must comply with the requirements of this procedure.

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RESPONS	IBILITY		ACTION
Record	Source	5.1.3.4	Temporary records shall not be submitted on the same magnetic tape as permanent records.
		5.1.3.5	Documentation adequate for servicing and interpreting machine readable records that have been designated for preservation shall be transferred along with the machine-readable file. This documentation shall include, but not be limited to, the following:
			a. Narrative description of the file(s)
			b. Physical file characteristics
			c. Recording mode information, including the coding structure (code books)
-			NOTE: Where it has been necessary to strip data of extraneous control characters, the code book specifications defining the data elements and their values must match the new format of the data.
			d. Recording system information
			e. A record layout that should break down the file by fields. Each field shall have a name, size, starting position, and a description of the form of the data (alphabetic, zoned decimal, packed decimal, or numeric).
			NOTE: Floppy disks and microfiche are not an acceptable storage media for records.
5.1.4	Submittal	of Oversi:	red and Special Process Records
Record	Source	5.1.4.1	Submit two copies of oversized records rolled with a Special Instruction Form, TMSS/009/1 (Exhibit 4) to the LRC.
1		5.1.4.2	Insert a copy of Form TMSS/009/1 in the record package where records would have been placed.
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RESPON	SIBILITY		ACTION
Record	Source 5	.1.4.3	Submit two copies of electronic records, film records, and other special process records as soon as they become inactive or whenever they cannot be maintained properly in accordance with this procedure.
	5	.1.4.4	Submit one-of-a-kind records when no longer in use.
	. 5	.1.4.5	Records shall be sent unbound or loose-leaf when possible.
5.1.5	Authenticat	ion and 1	Preparation of Records for Submittal
Record	Source 5	.1.5.1	Authenticate QA records if all requirements listed above have been met.
		.1.5.2	Determine if the record is part of a records package. If yes, then go to Step 5.1.1.4. Otherwise, go to step 5.2.
	5	.1.5.3	Determine if the record package is being maintained at the LRC or by the record source.
			a. If at the LRC, then go to Step 5.1.1.4.
			b. If by the record source, then include the record within the package, and continue to store in accordance with Step 5.1.1.5.b, and go to Step 5.1.1.5.
1	5	.1.5.4	Determine if the record(s) or record package(s) are identified as privileged records. If yes, then go to step 5.2.1.
	5	.1.5.5	If the record package is complete, then prepare a table of contents which contains the following (at a minimum):
			a. Record Package Identifier (upper right-hand corner) (See Section 3.2.7 of this procedure.)

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RESPONS	SIBILITY		ACTION						
Record	Source 5	.1.5.5	b. QA Designation (QA or QA:N/A) (upper right-hand corner directly under the Record Package Identifier)						
			c. Descriptive Title						
			d. Individual listing of the contents						
			NOTE: The table of contents is to be the first item listed.						
			e. Individual page count						
			f. Total page count.						
			g. Signature (and date) of the person preparing the table of contents indicating authentication (QA record/record package) or submittal (non-QA record/record package) in accordance with this procedure.						
			NOTE: See Exhibit 3 for a sample table of contents.						
5.2 RI	CORD SUBMIT	TAL							
the LRO Yucca N	with the e	xception e. These	pared under this procedure shall be submitted to of records created or initially received at the e records shall be submitted to the Yucca Mountain						
Record	Source 5	.2.1	Identify all privileged records and ensure that they are clearly labeled as such.						
	5	.2.2	Submit privileged records on a separate form TMSS/010 and identify them as "Privileged" on the form.						
	5	.2.3	Submit individual non-QA records and record packages with a completed Form TMSS/010 no later than 10 working days after the date of completion or receipt.						

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RESPONSIBILITY ACTION Record Source 5.2.4 Submit QA records and record packages with a completed Form TMSS/010 no later than 10 workin days after authentication. 5.2.5 Determine if any additional segments will be submitted for records packages being compiled b the LRC. NOTE: Record package segments may be submitted for records packages being compiled b the LRC with a completed Form TMSS/010 anytime during creation of the record package. 5.2.6 If the package is complete, notify the LRC and review the segments to ensure all applicable requirements of this procedure have been met. sign and date the table of contents and have th table of contents authenticated. 5.3 RECORD DISCREPANCIES Record Source 5.3.1 After submittal of records to the LRC, resolve discrepancies stated on the LRC Record Rejection Notice. NOTE: NOTE: Notify the LRC if the discrepancy(ies) cannot be resolved within the allowable time. 5.3.2 Notify the LRC of any errors found in previousl processed records or record packages. Submit th corrected, modified, or supplemental records, referencing the accession number of the previou processed record or record package, to the LRC.	s: <u>12914</u> A: <u>O</u> A	QA:	OCEDURE	EMENT SUPPO RACTICE PRO DURE [] W UATION PAGE	PR	NDARD N PROC	[_x] STA	. –	00 DN NO.: 2	TMSS/10 8/30/91 REVISIO				
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	5.4 RI	ETRIEVAL OF	RECORDS	
	RESPON	SIBILITY		ACTION
	Record	Source 5	5.4.1	Contact the LRC (e.g., via telephone, in person, electronic mail, interoffice mail) to retrieve a record(s) or to complete a Records Request Form (available in the LRC) and submit it to the LRC.
				a. Provide adequate information that will identify the record(s) {e.g., record date, author, organization, title, subject, etc.}.
				b. Provide a specific date or time that the record(s) is required.
,		5	5.4.2	Provide appropriate authorization or identification for retrieval of privileged records (i.e., training records, qualification records).
				NOTE: To obtain access to privileged records, authorization must be approved by Director of Office of Quality Assurance (OQA) or OCRWM, unless otherwise directed.
		5	5.4.3	Obtain assistance from the LRC to use the microfilm reader/printer to retrieve the record from microfilm if the request volume is large.
				NOTE: There are many factors that will determine the timely retrieval of records. These include, but are not limited to, the following: volume, priority, location of storage, specific information provided, and record location.
	6.0 RI	ECORDS		
	No reco	ords are ger	nerated a	as a result of this procedure.
	Record	Source f	6.1	Submit the signed documentation along with deficient record(s) when a record cannot be corrected or generated.

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	7.0 E	EXHIBITS AND	FORMS REF	ERENCED	IN THIS PROCED	URE		
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TECHNICAL & MANAGEMENT SUPPORT SERVICES [x] STANDARD PRACTICE PROCEDURE [] ORGANIZATION PROCEDURE [] WORK INSTRUCTION (CONTINUATION PAGE)

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EXHIBIT 1

LIST OF MATERIALS NOT PROCESSED INTO THE RECORDS SYSTEM

1.0 PURPOSE AND APPLICABILITY

This exhibit is a list of some non-processed materials. This list is a guideline to be used to determine whether materials should be processed into the records system. Contact the LRC to resolve differences involving interpretation of this guidance.

1.1 LIST OF NON-PROCESSED MATERIALS

The following materials shall not be captured in the records system and may be disposed of without special authority, except when the procedure governing the activity specifically requires the submission of this material.

Such material includes, but is not limited to, the following:

- Correspondence that is circulated or transmitted for information purposes only, and other materials on which no documented action is taken or required. Such materials should be identified as "Information Copy" (or designated information copy through a buckslip/routing slip).
- Correspondence and other materials documenting fringe activities such as employee welfare activities or charitable fund drives. Other materials of short-term value, that, after action has been completed, have neither programmatic nor informational value, such as requests for publications and communications on hotel reservations.
- 3. Tickler, follow-up, suspense, or reading file copies of records; duplicate copies of all records maintained in the same file; and extra copies of printed or processed material, official copies of which have been retained for record purposes.
- 4. Superseded manuals or other directives maintained outside the issuing office.

5. Routing slips.

.6. Electronic mail.

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	EXHIBIT 1
LIS	T OF MATERIALS NOT PROCESSED INTO THE RECORDS SYSTEM (Continued)
7.	Working papers, such as personal notes, reminders, or handwritten drafts.
8.	Transmittals sheets, which do not require action, unless used to transmit materials for action.
9.	Blank forms.
10.	Initial stenographic notes after the transcription is available.
11.	Processed or published material received from other activities or offices, which requires no action and is not needed for documentary purposes (the originating office or activity is required to maintain record copies).
12.	Catalogs, trade journals, and other publications or papers that are received from government agencies, commercial firms, or private institutions, which require no action and are not part of a case (activity or Project task) upon which action is taken.
13.	Reproduction materials such as stencils and offset masters.
14.	Physical exhibits, artifacts, and material lacking documentary value.
15.	Telecopies (facsimiles). If telecopies (facsimiles) of signed documents are sent, the original of the signed document(s), including draft documents, must be forwarded immediately through the mail system.
16.	Pre-award information and documents (e.g., information on a procurement prior to contract award, Source Evaluation Board materials, proposal information), except as required as a QA record.
17.	Personnel records, except as required as QA records (e.g., qualification and training records).
18.	Business sensitive (financial or commercial) information.

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