DICOM Correction Item

Correction Number	CP 747
Log Summary: Add clinical trial series identifier and description	
Type of Modification Addition	Name of Standard PS 3.3, 3.6 2007
Rationale for Correction When the clinical trial attributes were added, identifiers and descriptors were added for subjects, sites and time points, but not the series. There is a requirement to provide an additional identifier at the Series level to allow for the insertion of a trial-specific series identifier and description when it is desirable to retain the original manufacturer's values of Series Description, Series Number and Protocol Name.	
Sections of documents affected PS 3.3 C.7.3.2	
Correction Wording:	

PS 3.3: Add attributes to the Clinical Trial Series Module:

C.7.3.2 Clinical Trial Series Module

Table C.7-5b contains attributes that identify a Series in the context of a clinical trial.

Table C.7-5b
CLINICAL TRIAL SERIES MODULE ATTRIBUTES

Attribute Name	Tag	Type	Attribute Description
Clinical Trial Coordinating Center Name	(0012,0060)	2	The name of the institution that is responsible for coordinating the medical imaging data for the clinical trial. See C.7.3.2.1.1.
Clinical Trial Series ID	**(0012,xxx1)**	**3**	**An identifier of the series in the context of a clinical trial. See C.7.3.2.1.2.**
Clinical Trial Series Description	**(0012,xxx2)**	**3**	**A description of the series in the context of a clinical trial. See C.7.3.2.1.2.**

C.7.3.2.1 Clinical Trial Series Attribute Descriptions

C.7.3.2.1.1 Clinical Trial Coordinating Center Name

The Clinical Trial Coordinating Center Name (0012,0060) identifies the institution responsible for coordinating the collection of images and associated data for subjects enrolled in the clinical trial.

C.7.3.2.1.2 Clinical Trial Series Identifier and Description

The Clinical Trial Series ID (0012,xxx1) and Clinical Trial Series Description (0012,xxx2) attributes can be used to identify and describe a Series within the context of a clinical trial without requiring the replacement of the values in the Series Number (0020,0011) and Series Description (0008,103E) attributes in the General Series Module, whose manufacturer or user provided values may be relevant and important to retain.

PS 3.6: Add attributes:

(0012,0010)	Clinical Trial Sponsor Name	LO	1
(0012,0020)	Clinical Trial Protocol ID	LO	1
(0012,0021)	Clinical Trial Protocol Name	LO	1
(0012,0030)	Clinical Trial Site ID	LO	1
(0012,0031)	Clinical Trial Site Name	LO	1
(0012,0040)	Clinical Trial Subject ID	LO	1
(0012,0042)	Clinical Trial Subject Reading ID	LO	1
(0012,0050)	Clinical Trial Time Point ID	LO	1
(0012,0051)	Clinical Trial Time Point Description	ST	1
(0012,0060)	Clinical Trial Coordinating Center Name	LO	1
(0012,0062)	Patient Identify Removed	CS	1
(0012,0063)	De-identification Method	LO	1-n
(0012,0064)	De-identification Method Code Sequence	SQ	1
(0012,xxx1)	**Clinical Trial Series ID**	**LO**	**1**
(0012,xxx2)	**Clinical Trial Series Description**	**LO**	**1**