

MDA2016-08-02 - Adverse Events Data Dictionary

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Document Summary

Property	Value
Document Title	MDA2016-08-02 - Adverse Events: Data Dictionary
Date Created	06/29/2023
Sections	2
Entries	19
Document Filename	dictionary_adverse_events.rtf

MDA2016-08-02 - Adverse Events: Data Dictionary

Section 1: Identifiers

Class	Variable	Label	Description	Format Text
01. Principal	build_dt	Build Date		SAS Date
01. Principal	mrno	Participant ID		Char

Section 2: Adverse Events

Class	Variable	Label	Description	Format Text
01. Principal	ae_action	Change (If Any) In Agent Dose Due To Adverse Event		.M="Missing" .N="No Adverse Events Reported" 1="Agent Withdrawn" 2="Agent Dose Reduced" 3="Agent Dose Increased" 4="Agent Dose Not Changed" 5="Unknown" 6="Not Applicable" 7="Agent Interrupted"
01. Principal	ae_attribution	Likelihood Adverse Event Was Caused By Study Agent		.M="Missing" 1="Unrelated" 2="Unlikely" 3="Possible" 4="Probable" 5="Definite"
01. Principal	ae_comments	Comments		Char
01. Principal	ae_ctcae_term	CTCAE Term v4.0		Char
01. Principal	ae_dropped	Dropped Due To This Adverse Event?		.M="Missing" .N="N/A" 0="No" 1="Yes"
01. Principal	ae_end_dt	End Date Of Adverse Event		SAS Date .M="Missing"
01. Principal	ae_event	Adverse Event As Described Verbatim		Char
01. Principal	ae_grade	Severity Of Adverse Event		.M="Missing" 1="Mild Adverse Event" 2="Moderate Adverse Event" 3="Severe Adverse Event" 4="Life Threatening Adverse Event" 5="Death Related to Adverse Event"
01. Principal	ae_grade_description	Grade Description		Char
01. Principal	ae_interval_dt	Interval Date		SAS Date .M="Missing"
01. Principal	ae_meddra_v12	MedDRA v12.0 Code		Numeric .M="Missing" .N="No Adverse Events Reported" 99999999="No MedDRA Code Associated with CTCAE Term"
01. Principal	ae_onset_dt	Onset Date Of Adverse Event		SAS Date

Class	Variable	Label	Description	Format Text
01. Principal	ae_outcome	Outcome Of Adverse Event		.M="Missing" 1="Recovered/Resolved" 2="Recovering/Resolving" 3="Not recovering/Not Resolved" 4="Recovered/Resolved with sequelae" 5="Fatal" 6="Unknown"
01. Principal	ae_report_dt	Report Date Of Adverse Event		SAS Date
01. Principal	ae_sae	Serious Adverse Event?		.M="Missing" .N="N/A" 0="No" 1="Yes"
01. Principal	ae_system_organ	Toxicity Group		.M="Missing" .N="No Adverse Events Reported" 1="Blood and Lymphatic System Disorders" 2="Cardiac Disorders" 3="Congenital, Familial and Genetic Disorders" 4="Ear and Labyrinth Disorders" 5="Endocrine Disorders" 6="Eye Disorders" 7="Gastrointestinal Disorders" 8="General Disorders and Administration Site Conditions" 9="Hepatobiliary Disorders" 10="Immune System Disorders" 11="Infections and Infestations" 12="Injury, Poisoning and Procedural Complications" 13="Investigations" 14="Metabolism and Nutrition Disorders" 15="Musculoskeletal and Connective Tissue Disorders" 16="Neoplasms Benign, Malignant and Unspecified (Including Cysts and Polyps)" 17="Nervous System Disorders" 18="Pregnancy, Puerperium and Perinatal Conditions" 19="Psychiatric Disorders" 20="Renal and Urinary Disorders" 21="Reproductive System and Breast Disorders" 22="Respiratory, Thoracic and Mediastinal Disorders" 23="Skin and Subcutaneous Tissue Disorders" 24="Social Circumstances" 25="Surgical and Medical Procedures" 26="Vascular Disorders" 99="No Toxicity Group Associated With CTCAE Term"
01. Principal	ae_tac	Specify Treatment Assignment Code (TAC) At Time Of AE Onset		.M="Missing" 0="TAC0" 1="TAC1" 2="TAC2" 3="TAC3" 4="TAC4"

