

Package ‘pharmaverseadam’

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Type Package

Title ADaM Test Data for the 'Pharmaverse' Family of Packages

Version 1.0.0

Description A set of Analysis Data Model (ADaM) datasets constructed using the Study Data Tabulation Model (SDTM) datasets contained in the 'pharmaversesdtm' package and the template scripts from the 'admiral' family of packages. ADaM dataset specifications are described in: CDISC Analysis Data Model Team (2021) <https://www.cdisc.org/system/files/members/standard/foundational/ADaMIG_v1.3.pdf>.

License Apache License (>= 2)

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adae

Adverse Events Analysis

Description

An example of ADAE ADaM dataset

Usage

adae

Format

A data frame with 101 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
AESEQ Sequence Number
AESPID Sponsor-Defined Identifier
AETERM Reported Term for the Adverse Event
AELLT Lowest Level Term
AELLTCD Lowest Level Term Code
AEDECOD Dictionary-Derived Term
AEPTCD Preferred Term Code
AEHLT High Level Term
AEHLTCD High Level Term Code
AEHLGT High Level Group Term
AEHLGTCD High Level Group Term Code
AEBODSYS Body System or Organ Class
AEBDSYCD Body System or Organ Class Code
AESOC Primary System Organ Class
AESOCCD Primary System Organ Class Code
AESEV Severity/Intensity
AESER Serious Event
AEACN Action Taken with Study Treatment
AEREL Causality
AEOUT Outcome of Adverse Event
AESCAN Involves Cancer
AESCONG Congenital Anomaly or Birth Defect
AESDISAB Persist or Signif Disability/Incapacity
AESDTH Results in Death
AESHOSP Requires or Prolongs Hospitalization
AESLIFE Is Life Threatening
AESOD Occurred with Overdose
AEDTC Date/Time of Collection
AESTDTC Start Date/Time of Adverse Event
AEENDTC End Date/Time of Adverse Event
AESTDY Study Day of Start of Adverse Event
AEENDY Study Day of End of Adverse Event

TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
DTHDT Date of Death
EOSDT End of Study Date
ASTDTM Analysis Start Date/Time
ASTDTF Analysis Start Date Imputation Flag
ASTTMF Analysis Start Time Imputation Flag
AENDTM Analysis End Date/Time
AENDTF Analysis End Date Imputation Flag
AENTMF Analysis End Time Imputation Flag
ASTDT Analysis Start Date
AENDT Analysis End Date
ASTDY Analysis Start Relative Day
AENDY Analysis End Relative Day
ADURN Analysis Duration (N)
ADURU Analysis Duration Units
LDOSEDTM End Date/Time of Last Dose
ASEV Analysis Severity/Intensity
AREL Analysis Causality
TRTEMFL Treatment Emergent Analysis Flag
ASEVN Analysis Severity/Intensity (N)
AOCCIFL 1st Max Sev./Int. Occurrence Flag
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity

ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adae.R).

References

None

Examples

```
data("adae")
```

adbcva_ophtha

*Best Corrected Visual Acuity Analysis***Description**

An example of ADBCVA ADaM dataset for Ophthalmology studies

Usage

adbcva_ophtha

Format

A data frame with 116 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
OESEQ Sequence Number
OECAT Category for Ophthalmic Test or Exam
OESCAT Subcategory for Ophthalmic Test or Exam
OEDTC Date/Time of Collection
VISIT Visit Name
VISITNUM Visit Number
VISITDY Planned Study Day of Visit
OESTRESN Numeric Result/Finding in Standard Units
OESTRESC Character Result/Finding in Std Format
OEORRES Result or Finding in Original Units
OETEST Name of Ophthalmic Test or Exam
OETESTCD Short Name of Ophthalmic Test or Exam
OETSTDTL Ophthalmic Test or Exam Detail
OELAT Laterality
OELOC Location Used for the Measurement
OEDY Study Day of Visit/Collection/Exam
OEMETHOD Method of Test or Examination
OEORRESU Original Units
OESTRESU Standard Units
OESTAT Completion Status
OETPT Planned Time Point Name
OETPTNUM Planned Time Point Number

TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
STUDYEYE Study Eye Location
AVAL Analysis Value
AVALU Analysis Value Unit
DTYPE Derivation Type
AFEYE Affected Eye
PARAM Parameter
PARAMCD Parameter Code
AVALC Analysis Value (C)
ADT Analysis Date
ADY Analysis Relative Day
ATPTN Analysis Timepoint (N)
ATPT Analysis Timepoint
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
BASETYPE Baseline Type
ONTRTFL On Treatment Record Flag
ABLFL Baseline Record Flag
ANL01FL Analysis Flag 01
ANL02FL Analysis Flag 02
WORS01FL Worst Post Baseline Obs
BASE Baseline Value
BASEC Baseline Value (C)
CHG Change from Baseline
PCHG Percent Change from Baseline
ASEQ Analysis Sequence Number
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death

DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country/Region
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSST End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag
CRIT1 Analysis Criterion 1

CRIT1FL Criterion 1 Evaluation Result Flag
CRIT2 Analysis Criterion 2
CRIT2FL Criterion 2 Evaluation Result Flag
CRIT3 Analysis Criterion 3
CRIT3FL Criterion 3 Evaluation Result Flag
CRIT4 Analysis Criterion 4
CRIT4FL Criterion 4 Evaluation Result Flag
CRIT5 Analysis Criterion 5
CRIT5FL Criterion 5 Evaluation Result Flag
CRIT6 Analysis Criterion 6
CRIT6FL Criterion 6 Evaluation Result Flag
CRIT7 Analysis Criterion 7
CRIT7FL Criterion 7 Evaluation Result Flag
CRIT8 Analysis Criterion 8
CRIT8FL Criterion 8 Evaluation Result Flag
AVALCA1N Analysis Value Category 1 (N)
AVALCAT1 Analysis Value Category 1

Source

Generated from admiralophtha package (template ad_adbcva.R).

References

None

Examples

```
data("adbcva_ophtha")
```

adce_vaccine

Clinical Events Analysis for Vaccine

Description

An example of ADCE dataset for Vaccine studies

Usage

```
adce_vaccine
```

Format

A data frame with 56 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
CESEQ Sequence Number
CELNKID Link ID
CELNKGRP Link Group ID
CETERM Reported Term for the Clinical Event
CEDECOD Dictionary-Derived Term
CELAT Laterality
CELOC Location of Event
CECAT Category for the Clinical Event
CESCAT Subcategory for the Clinical Event
CEPRESP Clinical Event Pre-specified
CEOCCUR Clinical Event Occurrence
CESEV Severity/Intensity
CEREL Causality
CEOUT Outcome of Event
EPOCH Epoch
CEDTC Date/Time of Event Collection
CESTDTC Start Date/Time of Clinical Event
CEENDTC End Date/Time of Clinical Event
CEDUR Duration of Clinical Event
CETPT Planned Time Point Name
CETPTNUM Planned Time Point Number
CETPTREF Time Point Reference
CERFTDTC Date/Time of Reference Time Point
CEEVINTX Evaluation Interval Text
CESTAT Completion Status
CEREASND Reason Clinical Event Not Collected
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
ASTDT Analysis Start Date
AENDT Analysis End Date
ASTDY Analysis Start Relative Day
AENDY Analysis End Relative Day

APERIOD Period
APERSDT Period Start Date
APEREDT Period End Date
APERSTDY Analysis Sub-period Start Relative Day
AREL Analysis Causality
ASEV Analysis Severity/Intensity
ASEVN Analysis Severity/Intensity (N)
AOCC01FL Event Occurrence Flag
ASEQ Analysis Sequence Number
ADURN Analysis Duration (N)
ADURU Analysis Duration Units
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
AGE Age
AGEU Age Units
SEX Sex
RACE Race
COUNTRY Country
ETHNIC Ethnicity
SITEID Study Site Identifier
SUBJID Subject Identifier for the Study

Source

Generated from admiralvaccine package (template ad_adce.R).

References

None

Examples

```
data("adce_vaccine")
```

adcm

Concomitant Medications Analysis

Description

An example of ADCM ADaM dataset

Usage

adcm

Format

A data frame with 90 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
CMSEQ Sequence Number
CMSPID Sponsor-Defined Identifier
CMTRT Reported Name of Drug, Med, or Therapy
CMDECOD Standardized Medication Name
CMINDC Indication
CMCLAS Medication Class
CMDOSE Dose per Administration
CMDOSU Dose Units
CMDOSFRQ Dosing Frequency per Interval
CMROUTE Route of Administration
VISITNUM Visit Number
VISIT Visit Name
VISITDY Planned Study Day of Visit
CMDTC Date/Time of Collection
CMSTDTC Start Date/Time of Medication
CMENDTC End Date/Time of Medication
CMSTDY Study Day of Start of Medication
CMENDY Study Day of End of Medication
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
DTHDT Date of Death
EOSDT End of Study Date

TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
ASTDTM Analysis Start Date/Time
ASTDTF Analysis Start Date Imputation Flag
ASTTMF Analysis Start Time Imputation Flag
AENDTM Analysis End Date/Time
AENDTF Analysis End Date Imputation Flag
AENTMF Analysis End Time Imputation Flag
ASTDT Analysis Start Date
AENDT Analysis End Date
ASTDY Analysis Start Relative Day
AENDY Analysis End Relative Day
ADURN Analysis Duration (N)
ADURU Analysis Duration Units
ONTRTFL On Treatment Record Flag
PREFL Pre-treatment Flag
FUPFL Follow-up Flag
ANL01FL Analysis Flag 01
AOCCPFL 1st Occurrence of Preferred Term Flag
APHASE Phase
APHASEN Description of Phase N
TRTP Planned Treatment
TRTA Actual Treatment
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race

ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Treatment End Datetime Imput Flag
TRTDURD Total Treatment Duration (Days)
SCRFDI Screen Failure Date
EOSST End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adcm.R).

References

None

Examples

```
data("adcm")
```

adeg

Electrocardiogram Tests Analysis

Description

An example of ADEG ADaM dataset

Usage

adeg

Format

A data frame with 105 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
EGSEQ Sequence Number
EGTESTCD ECG Test or Examination Short Name
EGTEST ECG Test or Examination Name
EGORRES Result or Finding in Original Units
EGORRESU Original Units
EGSTRESC Character Result/Finding in Std Format
EGSTRESN Numeric Result/Finding in Standard Units
EGSTRESU Standard Units
EGSTAT Completion Status
EGLOC Lead Location Used for Measurement
EGBLFL Baseline Flag
VISITNUM Visit Number
VISIT Visit Name
VISITDY Planned Study Day of Visit
EGDTC Date/Time of ECG
EGDY Study Day of ECG
EGTPT Planned Time Point Name
EGTPTNUM Planned Time Point Number
EGELTM Planned Elapsed Time from Time Point Ref
EGTPTREF Time Point Reference
temp undocumented field
TRTSDT Date of First Exposure to Treatment

TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
ADTM Analysis Datetime
ATMF Analysis Time Imputation Flag
ADY Analysis Relative Day
PARAMCD Parameter Code
AVAL Analysis Value
AVALC Analysis Value (C)
ADT Analysis Date
ATPTN Analysis Timepoint (N)
ATPT Analysis Timepoint
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
DTYPE Derivation Type
ONTRTFLL On Treatment Record Flag
ANRLO Analysis Normal Range Lower Limit
ANRHI Analysis Normal Range Upper Limit
ANRIND Analysis Reference Range Indicator
BASETYPE Baseline Type
ABLFL Baseline Record Flag
BASE Baseline Value
BASEC Baseline Value (C)
BNRIND Baseline Reference Range Indicator
CHG Change from Baseline
PCHG Percent Change from Baseline
ANL01FL Analysis Flag 01
TRTP Planned Treatment
TRTA Actual Treatment
ASEQ Analysis Sequence Number
AVALCA1N Analysis Value Category 1 (N)
AVALCAT1 Analysis Value Category 1
CHGCAT1N Change from Baseline Category 1 (N)
CHGCAT1 Change from Baseline Category 1
PARAM Parameter
PARAMN Parameter (N)
SUBJID Subject Identifier for the Study

RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag

RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adeq.R).

References

None

Examples

```
data("adex")
```

adex

Exposure Analysis

Description

An example of ADEX ADaM dataset

Usage

```
adex
```

Format

A data frame with 88 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
EXSEQ Sequence Number
EXTRT Name of Treatment
EXDOSE Dose
EXDOSU Dose Units
EXDOSFRM Dose Form
EXDOSFRQ Dosing Frequency per Interval
EXROUTE Route of Administration

VISITNUM Visit Number
VISIT Visit Name
VISITDY Planned Study Day of Visit
EXSTDTC Start Date/Time of Treatment
EXENDTC End Date/Time of Treatment
EXSTDY Study Day of Start of Treatment
EXENDY Study Day of End of Treatment
EXADJ Reason for Dose Adjustment
EXPLDOS Planned Dose
TRTSDT Date of First Exposure to Treatment
TRTSDTM Datetime of First Exposure to Treatment
TRTEDTM Datetime of Last Exposure to Treatment
ASTDTM Analysis Start Datetime
ASTDTF Analysis Start Date Imputation Flag
ASTTMF Analysis Start Time Imputation Flag
AENDTM Analysis End Datetime
AENDTF Analysis End Date Imputation Flag
AENTMF Analysis End Time Imputation Flag
ASTDY Analysis Start Relative Day
AENDY Analysis End Relative Day
EXDURD Duration of Treatment (Days)
ASTDT Analysis Start Date
AENDT Analysis End Date
DOSEO Dose O
PDOSEO PDose O
PARAMCD Parameter Code
AVAL Analysis Value
AVALC Analysis Value (C)
PARCAT1 Parameter Category 1
PARAM Parameter
PARAMN Parameter (N)
AVALCAT1 Analysis Value Category 1
ASEQ Analysis Sequence Number
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment

RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
TRTSTMF Time of First Exposure Imput. Flag
TRTETMF Time of Last Exposure Imput. Flag
TRTEDT Date of Last Exposure to Treatment
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adex.R).

References

None

Examples

```
data("adex")
```

adface_vaccine

Findings About Clinical Events Analysis

Description

An example of ADFACE for Vaccine studies

Usage

```
adface_vaccine
```

Format

A data frame with 60 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
SUBJID Subject Identifier for the Study
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
SAFFL Safety Population Flag
ARM Description of Planned Arm
ARMCD Planned Arm Code
ACTARM Description of Actual Arm
ACTARMCD Actual Arm Code
TRTSDT Date of First Exposure to Treatment
TRTSDTM Datetime of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRTEDTM Datetime of Last Exposure to Treatment

FATEST Findings About Test Name
FALNKID Link ID
FALNKGRP Link Group ID
FATESTCD Findings About Test Short Name
PARAMCD Parameter Code
PARAM Parameter
PARAMN Parameter (N)
FAOBJ Object of the Observation
PARCAT1 Parameter Category 1
PARCAT2 Parameter Category 2
AVALC Analysis Value (C)
AVAL Analysis Value
FASTAT Completion Status
FAREASND Reason Not Performed
FAEVAL Evaluator
EPOCH Epoch
ADT Analysis Date
ADTM Analysis Datetime
FAEVINTX Evaluation Interval Text
ADY Analysis Relative Day
ATPT Analysis Timepoint
ATPTN Analysis Timepoint (N)
ATPTREF Analysis Timepoint Reference
EXDOSE Dose
EXTRT Name of Treatment
EXSTDTC Start Date/Time of Treatment
EXENDTC End Date/Time of Treatment
TRTA Actual Treatment
TRTP Planned Treatment
APERIOD Period
APERSDT Period Start Date
APEREDT Period End Date
FAORRES Result or Finding in Original Units
TRT01P Planned Treatment for Period 01
TRT02P Planned Treatment for Period 02
TRT01A Actual Treatment for Period 01
TRT02A Actual Treatment for Period 02

VAX01DT Vaccination Date 01
VAX02DT Vaccination Date 02
EVENTFL Event Value Flag
EVENTDFL Day Event Value Flag
ANL01FL Analysis Flag 01
ANL02FL Analysis Flag 02

Source

Generated from admiralvaccine package (template ad_adface.R).

References

None

Examples

```
data("adface_vaccine")
```

adis_vaccine

Immunogenicity Specimen Assessments

Description

An example of ADIS dataset for Vaccine studies

Usage

```
adis_vaccine
```

Format

A data frame with 102 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
ISSEQ Sequence Number
ISTESTCD Immunogenicity Test/Exam Short Name
ISTEST Immunogenicity Test or Examination Name
ISCAT Category for Immunogenicity Test
ISORRES Results or Findings in Original Units
ISORRESU Original Units
ISSTRESC Character Result/Finding in Std Format

ISSTRESN Numeric Results/Findings in Std. Units
ISSTRESU Standard Units
ISSTAT Completion Status
ISREASND Reason Not Done
ISNAM Vendor Name
ISSPEC Specimen Type
ISMETHOD Method of Test or Examination
ISBLFL Baseline Flag
ISLLOQ Lower Limit of Quantitation
VISITNUM Visit Number
EPOCH Epoch
ISDTC Date/Time of Collection
ISDY Study Day of Visit/Collection/Exam
ISULOQ Upper Limit of Quantitation
LOD Limit of Detection
AVISITN Analysis Visit (N)
AVISIT Analysis Visit
ATPTN Analysis Timepoint (N)
ATPT Analysis Timepoint
ATPTREF Analysis Timepoint Reference
ADT Analysis Date
RFSTDTC Subject Reference Start Date/Time
PPROTFL Per-Protocol Population Flag
ADY Analysis Relative Day
PARAMCD Parameter Code
PARAM Parameter
PARAMN Parameter (N)
PARCAT1 Parameter Category 1
CUTOFF02 First Cutoff Value
CUTOFF03 Second Cutoff Value
AVAL Analysis Value
AVALU Analysis Value Unit
SERCAT1 Pre-vaccination seropositivity status
SERCAT1N Pre-vaccination sero status (n)
DTYPE Derivation Type
BASETYPE Baseline Type
BASE Baseline Value

ABLFL Baseline Record Flag
BASECAT1 Baseline Category 1
CHG Change from Baseline
R2BASE Ratio to Baseline
CRIT1FL Criterion 1 Evaluation Result Flag
CRIT1FN Criterion 1 Evaluation Result Flag (N)
CRIT1 Analysis Criterion 1
APERIOD Period
APERSDT Period Start Date
APEREDT Period End Date
TRTA Actual Treatment
TRTP Planned Treatment
PPSRFL Per-Protocol Record-Level Flag
SUBJID Subject Identifier for the Study
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
INVID Investigator Identifier
INVNAM Investigator Name
BRTHDTC Date/Time of Birth
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection

TRT01P Planned Treatment for Period 01
TRT02P Planned Treatment for Period 02
TRT01A Actual Treatment for Period 01
TRT02A Actual Treatment for Period 02
TRTSDTM Datetime of First Exposure to Treatment
TRTEDTM Datetime of Last Exposure to Treatment
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
AGEGR1 Pooled Age Group 1
REGION1 Geographic Region 1
VAX01DT Vaccination Date 01
VAX02DT Vaccination Date 02
AP01SDT Period 01 Start Date
AP01EDT Period 01 End Date
AP02SDT Period 02 Start Date
AP02EDT Period 02 End Date

Source

Generated from admiralvaccine package (template ad_adis.R).

References

None

Examples

```
data("adis_vaccine")
```

adlb

Laboratory Analysis

Description

An example of ADLB ADaM dataset

Usage

adlb

Format

A data frame with 111 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
LBSEQ Sequence Number
LBTESTCD Lab Test or Examination Short Name
LBTEST Lab Test or Examination Name
LBCAT Category for Lab Test
LBORRES Result or Finding in Original Units
LBORRESU Original Units
LBORNULO Reference Range Lower Limit in Orig Unit
LBORNURHI Reference Range Upper Limit in Orig Unit
LBSTRESC Character Result/Finding in Std Format
LBSTRESN Numeric Result/Finding in Standard Units
LBSTRESU Standard Units
LBSTNRLO Reference Range Lower Limit-Std Units
LBSTNRHI Reference Range Upper Limit-Std Units
LBNRIND Reference Range Indicator
LBBLFL Baseline Flag
VISITNUM Visit Number
VISIT Visit Name
VISITDY Planned Study Day of Visit
LBDTC Date/Time of Specimen Collection
LBDY Study Day of Specimen Collection
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
ADT Analysis Date
ADY Analysis Relative Day
PARAMCD Parameter Code
PARAM Parameter
PARAMN Parameter (N)
PARCAT1 Parameter Category 1
AVAL Analysis Value
AVALC Analysis Value (C)

ANRLO Analysis Normal Range Lower Limit
ANRHI Analysis Normal Range Upper Limit
DTYPE Derivation Type
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
ONTRTFL On Treatment Record Flag
ANRIND Analysis Reference Range Indicator
BASETYPE Baseline Type
ABLFL Baseline Record Flag
BASE Baseline Value
BASEC Baseline Value (C)
BNRIND Baseline Reference Range Indicator
CHG Change from Baseline
PCHG Percent Change from Baseline
ATOXSCL Analysis Toxicity Description Low
ATOXSCH Analysis Toxicity Description High
ATOXGRL Analysis Toxicity Grade Low
ATOXGRH Analysis Toxicity Grade High
ATOXGR Analysis Toxicity Grade
BTOXGRL Baseline Toxicity Grade Low
BTOXGRH Baseline Toxicity Grade High
BTOXGR Baseline Toxicity Grade
R2BASE Ratio to Baseline
R2ANRLO Ratio of Analysis Val compared to ANRLO
R2ANRHI Ratio of Analysis Val compared to ANRHI
SHIFT1 Shift from Baseline to Analysis Value
SHIFT2 Shift from Baseline to Overall Grade
ANL01FL Analysis Flag 01
LVOTFL Last Value On Treatment Record Flag
TRTP Planned Treatment
TRTA Actual Treatment
ASEQ Analysis Sequence Number
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment

RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDI Screen Failure Date
EOSDT End of Study Date
EOSST End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adlb.R).

References

None

Examples

```
data("adlb")
```

adlbhy

Analysis of Lab Hy's Law

Description

A template of the ADLBHY dataset

Usage

```
adlbhy
```

Format

A data frame with 14 columns:

STUDYID Study Identifier

USUBJID Unique Subject Identifier

TRT01A Actual Treatment for Period 01

PARAMCD Parameter Code

LBSEQ Sequence Number

ADT Analysis Date

AVISIT Analysis Visit

ADY Analysis Relative Day

AVAL Analysis Value

ANRHI Analysis Normal Range Upper Limit

CRIT1 Analysis Criterion 1

CRIT1FL Criterion 1 Evaluation Result Flag

AVALC Analysis Value (C)

PARAM Parameter

Source

Generated from admiral package (template ad_adlbhy.R).

References

None

Examples

```
data("adlbhy")
```

admh

*Medical History Analysis***Description**

A template of ADMH ADaM dataset

Usage

admh

Format

A data frame with 110 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
MHSEQ Sequence Number
MHSPID Sponsor-Defined Identifier
MHTERM Reported Term for the Medical History
MHLLT Lowest Level Term
MHDECOD Dictionary-Derived Term
MHHLT High Level Term
MHHLGT High Level Group Term
MHCAT Category for Medical History
MHBODSYS Body System or Organ Class
MHSEV Severity/Intensity
VISITNUM Visit Number
VISIT Visit Name
VISITDY Planned Study Day of Visit
MHDTC Date/Time of History Collection
MHSTDTC Start Date/Time of Medical History Event
MHDY Study Day of History Collection

MHENDTC End Date/Time of Medical History Event
MHPRESP Medical History Event Pre-Specified
MHOCCUR Medical History Occurrence
MHSTRTPT Start Relative to Reference Time Point
MHENRTPT End Relative to Reference Time Point
MHSTTPT Start Reference Time Point
MHENTPT End Reference Time Point
MHENRF End Relative to Reference Period
MHSTAT Completion Status
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
DTHDT Date of Death
EOSDT End of Study Date
ASTDT Analysis Start Date
AENDT Analysis End Date
ASTDY Analysis Start Relative Day
AENDY Analysis End Relative Day
ADT Analysis Date
ADY Analysis Relative Day
SMQ02NAM SMQ 02 Name
SMQ02CD SMQ 02 Code
SMQ02SC SMQ 02 Scope
SMQ02SCN SMQ 02 Scope (N)
SMQ03NAM SMQ 03 Name
SMQ03CD SMQ 03 Code
SMQ03SC SMQ 03 Scope
SMQ03SCN SMQ 03 Scope (N)
SMQ05NAM SMQ 05 Name
SMQ05CD SMQ 05 Code
SMQ05SC SMQ 05 Scope
SMQ05SCN SMQ 05 Scope (N)
CQ01NAM Customized Query 01 Name
CQ04NAM Customized Query 04 Name
CQ04CD Customized Query 04 Code
AHIST Response of Med Hx (past or current)

AOCCFL 1st Occurrence within Subject Flag
AOCCSFL 1st Occurrence of SOC Flag
AOCCPFL 1st Occurrence of Preferred Term Flag
AOCPFL 1st Occur w/in Trt Prd FL
AOCPSFL 1st Occur of SOC w/in Trt Prd FL
AOCPPFL 1st Occur of PT w/in Trt Prd FL
ANL01FL Analysis Flag 01
TRTP Planned Treatment
TRTA Actual Treatment
APHASE Phase
APHASEN Description of Phase N
MHTERMN Medical History Term (N)
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Input. Flag
TRTEDTM Datetime of Last Exposure to Treatment

TRTETMF Treatment End Datetime Input Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_admh.R).

References

None

Examples

```
data("admh")
```

adoe_ophtha

Exam Analysis for Ophthalmology

Description

An example of ADOE dataset for Ophthalmology studies

Usage

```
adoe_ophtha
```

Format

A data frame with 98 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
OESEQ Sequence Number
OECAT Category for Ophthalmic Test or Exam
OESCAT Subcategory for Ophthalmic Test or Exam
OEDTC Date/Time of Collection
VISIT Visit Name
VISITNUM Visit Number
VISITDY Planned Study Day of Visit
OESTRESN Numeric Result/Finding in Standard Units
OESTRESC Character Result/Finding in Std Format
OEORRES Result or Finding in Original Units
OETEST Name of Ophthalmic Test or Exam
OETESTCD Short Name of Ophthalmic Test or Exam
OETSTDTL Ophthalmic Test or Exam Detail
OELAT Laterality
OELOC Location Used for the Measurement
OEDY Study Day of Visit/Collection/Exam
OEMETHOD Method of Test or Examination
OEORRESU Original Units
OESTRESU Standard Units
OESTAT Completion Status
OETPT Planned Time Point Name
OETPTNUM Planned Time Point Number
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
STUDYEYE Study Eye Location
AVAL Analysis Value
AVALC Analysis Value (C)
AVALU Analysis Value Unit
DTYPE Derivation Type
AFEYE Affected Eye

PARAM Parameter
PARAMCD Parameter Code
ADT Analysis Date
ADY Analysis Relative Day
ATPTN Analysis Timepoint (N)
ATPT Analysis Timepoint
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
BASETYPE Baseline Type
ONTRTFL On Treatment Record Flag
ABLFL Baseline Record Flag
ANL01FL Analysis Flag 01
ANL02FL Analysis Flag 02
WORS01FL Worst Post Baseline Obs
BASE Baseline Value
BASEC Baseline Value (C)
CHG Change from Baseline
PCHG Percent Change from Baseline
ASEQ Analysis Sequence Number
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code

ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiralophtha package (template ad_adoe.R).

References

None

Examples

```
data("adoe_ophtha")
```

adpc

Pharmacokinetic Concentrations

Description

An example of ADPC dataset

Usage

adpc

Format

A data frame with 123 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
NFRLT Nom. Rel. Time from Analyte First Dose
PCTESTCD Pharmacokinetic Test Short Name
PCTEST Pharmacokinetic Test Name
PCORRES Result or Finding in Original Units
PCORRESU Original Units
PCSTRESC Character Result/Finding in Std Format
PCSTRESN Numeric Result/Finding in Standard Units
PCSTRESU Standard Units
PCNAM Vendor Name
PCSPEC Specimen Material Type
PCLLOQ Lower Limit of Quantitation
VISIT Visit Name
VISITNUM Visit Number
PCDTC Date/Time of Specimen Collection
PCDY Actual Study Day of Specimen Collection
PCTPT Planned Time Point Name
PCTPTNUM Planned Time Point Number
TRTSDT Date of First Exposure to Treatment
TRTSDTM Datetime of First Exposure to Treatment
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
ADTM Analysis Datetime
ATMF Analysis Time Imputation Flag

ADT Analysis Date
ATM Analysis Time
ADY Analysis Relative Day
FANLDTM First Datetime of Dose for Analyte
AVISITN Analysis Visit (N)
AVISIT Analysis Visit
ASTDT Analysis Start Date
ASTDTM Analysis Start Datetime
AENDT Analysis End Date
AENDTM Analysis End Datetime
ASTTM Analysis Start Time
AENTM Analysis End Time
AFRLT Act. Rel. Time from Analyte First Dose
ARRLT Actual Rel. Time from Ref. Dose
PCRFTDTM Reference Datetime of Dose for Analyte
FANLDT First Date of Dose for Analyte
FANLTM First Time of Dose for Analyte
PCRFTDT Reference Date of Dose for Analyte
PCRFTTM Reference Time of Dose for Analyte
NRRLT Nominal Rel. Time from Ref. Dose
PARCAT1 Parameter Category 1
ATPTN Analysis Timepoint (N)
ATPT Analysis Timepoint
ATPTREF Analysis Timepoint Reference
ABLFL Baseline Record Flag
BASETYPE Baseline Type
DOSEA Actual Treatment Dose
DOSEP Planned Treatment Dose
DOSEU Treatment Dose Units
FRLTU Rel. Time from First Dose Unit
RRLTU Rel. Time from Ref. Dose Unit
PARAMCD Parameter Code
ALLOQ Analysis Lower Limit of Quantitation
AVAL Analysis Value
AVALU Analysis Value Unit
AVALCAT1 Analysis Value Category 1
SRCDOM Source Data

SRCVAR Source Variable
SRCSEQ Source Sequence Number
DTYPE Derivation Type
MRRLT Modified Rel. Time from Ref. Dose
ANL01FL Analysis Flag 01
ANL02FL Analysis Flag 02
BASE Baseline Value
CHG Change from Baseline
ASEQ Analysis Sequence Number
PARAM Parameter
PARAMN Parameter (N)
HTBL Numeric Result/Finding in Standard Units
HTBLU Standard Units
WTBL Numeric Result/Finding in Standard Units
WTBLU Standard Units
BMIBL Baseline Body Mass Index (kg/m²)
BMIBLU BMI at Baseline (Unit)
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country

DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTEDT Date of Last Exposure to Treatment
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Under 30 Group
DTHA30FL Over 30 Group
DTHB30FL Over 30 plus 30 days Group

Source

Generated from admiral package (template ad_adpc.R).

References

None

Examples

```
data("adpc")
```

adpp

Pharmacokinetic Parameters

Description

An example of ADPP ADaM dataset

Usage

adpp

Format

A data frame with 75 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
PPTESTCD Parameter Short Name
PPTEST Parameter Name
PPCAT Parameter Category
PPORRES Result or Finding in Original Units
PPORRESU Original Units
PPSTRESU Standard Units
PPSPEC Specimen Material Type
PPRFDTC Date/Time of Reference Point
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
DTHDT Date of Death
EOSDT End of Study Date
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
ADT Analysis Date
ADY Analysis Relative Day
PARAMCD Parameter Code
PARCAT1 Parameter Category
AVAL Numeric Result/Finding in Standard Units
AVALU Standard Units
SRCDOM Domain Abbreviation
SRCVAR Source Variable
SRCSEQ Sequence Number

AVISITN Analysis Visit (N)
AVISIT Analysis Visit
VISITNUM Visit Number
VISIT Visit Name
TRTP Planned Treatment
TRTA Actual Treatment
AVALCA1N Analysis Value Category 1 (N)
AVALCAT1 Analysis Value Category 1
SUBJID Subject Identifier for the Study
RFSTDTTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Input. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Input. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSSTT End of Study Status

FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adpp.R).

References

None

Examples

```
data("adpp")
```

adppk

Population Pharmacokinetic

Description

A template of ADPPK ADaM dataset

Usage

```
adppk
```

Format

A data frame with 61 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
EVID Event ID
NFRLT Nom. Rel. Time from Analyte First Dose
AFRLT Act. Rel. Time from Analyte First Dose
APRLT Actual Rel Time from Previous Dose
NPRLT Nominal Rel Time from Previous Dose
DOSEA Actual Treatment Dose
DOSEP Planned Treatment Dose
PARAMCD Parameter Code
ALLOQ Analysis Lower Limit of Quantitation
CMT Compartment
BLQFL Below Lower Limit of Quant Flag
BLQFN Below Lower Limit of Quant Flag (N)
AMT Actual Amount of Dose Received (unit)
DV Dependent Variable Result
AVAL Analysis Value
DVL Log DV
MDV Missing Dependent Variable Result
AVALU Analysis Value Unit
UDTC Date/Time
II Dosing Interval (unit)
SS Steady State
ASEQ Analysis Sequence Number
PARAM Parameter
PARAMN Parameter (N)
PROJID Project Identifier
PROJIDN Project Identifier (N)
STUDYIDN Study Identifier (N)
SITEID Study Site Identifier
SITEIDN Study Site Identifier (N)
USUBJIDN Unique Subject Identifier (N)
SUBJID Subject Identifier for the Study
SUBJIDN Subject Identifier for the Study (N)
AGE Age

SEX Sex
SEXN Sex (N)
COHORT Cohort Subject Enrolled Into
COHORTC Description of Planned Arm
ROUTE Route of Administration
ROUTEN Route of Administration (N)
RACE Race
RACEN Race (N)
ETHNIC Ethnicity
ETHNICN Ethnicity (N)
FORM Drug Formulation
FORMN Drug Formulation (N)
COUNTRY Country
COUNTRYN Country (N)
COUNTRYL Country Name
HTBL Numeric Result/Finding in Standard Units
WTBL Numeric Result/Finding in Standard Units
ALTBL Numeric Result/Finding in Standard Units
ASTBL Numeric Result/Finding in Standard Units
TBILBL Numeric Result/Finding in Standard Units
CREATBL Numeric Result/Finding in Standard Units
BMIBL Baseline Body Mass Index (kg/m²)
BSABL Numeric Result/Finding in Standard Units
CRCLBL Baseline Creatinine Clearance
EGFRBL Age
RECSEQ Record Sequence

Source

Generated from admiral package (template ad_adppk.R).

References

None

Examples

```
data("adppk")
```

adrs_onco

Tumor Response Analysis

Description

An example of ADRS dataset for Oncology studies

Usage

adrs_onco

Format

A data frame with 79 columns:

DOMAIN Domain Abbreviation
STUDYID Study Identifier
USUBJID Unique Subject Identifier
VISITNUM Visit Number
VISIT Visit Name
RSTESTCD Assessment Short Name
RSTEST Assessment Name
RSORRES Result or Finding in Original Units
RSSTRESC Character Result/Finding in Std Format
RSEVAL Evaluator
RSEVALID Evaluator Identifier
RSACPTFL Accepted Record Flag
RSDDTC Date/Time of Assessment
RSSEQ Sequence Number
RANDDT Date of Randomization
PARAMCD Parameter Code
PARAM Parameter
PARCAT1 Parameter Category 1
PARCAT2 Parameter Category 2
PARCAT3 Parameter Category 3
ADT Analysis Date
ADTF Analysis Date Imputation Flag
AVISIT Analysis Visit
AVALC Analysis Value (C)
AVAL Analysis Value

ANL01FL Analysis Flag 01
ANL02FL Analysis Flag 02
ASEQ Analysis Sequence Number
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Input. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Input. Flag
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status

FRVDT Final Retrieval Visit Date
DTHDT Date of Death
DTHDTF Date of Death Imputation Flag
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
DTHCAUS Cause of Death
DTHDOM Domain for Date of Death Collection
DTHCGR1 Cause of Death Reason 1
LSTALVDT Date Last Known Alive
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
AGEGR1 Pooled Age Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiralonco package (template ad_adrs.R).

References

None

Examples

```
data("adrs_onco")
```

adsl

Subject Level Analysis

Description

An example of ADSL ADaM dataset

Usage

```
adsl
```

Format

A data frame with 54 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Input. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Input. Flag
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date

EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHDTF Date of Death Imputation Flag
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
DTHCAUS undocumented field
DTHDOM undocumented field
DTHCGR1 undocumented field
LSTALVDT Date Last Known Alive
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
AGEGR1 Pooled Age Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adsl.R).

References

None

Examples

```
data("adsl")
```

adsl_vaccine

Subject Level Analysis for Vaccine

Description

An example of ADSL ADaM dataset for Vaccine studies

Usage

adsl_vaccine

Format

A data frame with 46 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
INVID Investigator Identifier
INVNAM Investigator Name
BRTHDTC Date/Time of Birth
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country/Region

DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRT01P Planned Treatment for Period 01
TRT02P Planned Treatment for Period 02
TRT01A Actual Treatment for Period 01
TRT02A Actual Treatment for Period 02
TRTSDTM Datetime of First Exposure to Treatment
TRTEDTM Datetime of Last Exposure to Treatment
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
SAFFL Safety Population Flag
PPROTFL Per-Protocol Population Flag
RACEGR1 Pooled Race Group 1
AGEGR1 Pooled Age Group 1
REGION1 Geographic Region 1
VAX01DT Vaccination Date 01
VAX02DT Vaccination Date 02
AP01SDT Period 01 Start Date
AP01EDT Period 01 End Date
AP02SDT Period 02 Start Date
AP02EDT Period 02 End Date

Source

Generated from admiralvaccine package (template ad_adsl.R).

References

None

Examples

```
data("adsl_vaccine")
```

adtr_onco

*Tumor Results Analysis for Oncology***Description**

An example of ADTR ADaM dataset for Oncology studies

Usage

adtr_onco

Format

A data frame with 99 columns:

DOMAIN Domain Abbreviation
STUDYID Study Identifier
USUBJID Unique Subject Identifier
TRGRPID Group ID
TRLNKID Link ID
TRTESTCD Tumor/Lesion Assessment Short Name
TRTEST Tumor/Lesion Assessment Test Name
TRORRES Result or Finding in Original Units
TRORRESU Original Units
TRSTRESC Character Result/Finding in Std Format
TRSTRESN Numeric Result/Finding in Standard Units
TRSTRESU Standard Units
VISITNUM Visit Number
VISIT Visit Name
TREVAL Evaluator
TREVALID Evaluator Identifier
TRACPTFL Accepted Record Flag
TRDTC Date/Time of Tumor/Lesion Measurement
TRSEQ Sequence Number
RANDDT Date of Randomization
TULOC Location of the Tumor/Lesion
TULOCGR1 Tumor Site Group 1
LSEXP Lesion IDs Expected
LSASS Lesion IDs Assessed
ADT Analysis Date

ADTF Analysis Date Imputation Flag
ADY Analysis Relative Day
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
PARAMCD Parameter Code
PARAM Parameter
PARCAT1 Parameter Category 1
PARCAT2 Parameter Category 2
PARCAT3 Parameter Category 3
AVAL Analysis Value
ANL01FL Analysis Flag 01
ABLFL Baseline Record Flag
BASE Baseline Value
NADIR NADIR
CHG Change from Baseline
PCHG Percent Change from Baseline
CHGNAD Change from NADIR
PCHGNAD Percent Change from NADIR
PDFL Pharmacodynamic Analysis Set Flag
ANL02FL Analysis Flag 02
ANL03FL Analysis Flag 03
ANL04FL Analysis Flag 04
ASEQ Analysis Sequence Number
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race

ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRT01P Planned Treatment for Period 01
TRT01A Actual Treatment for Period 01
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
DTHDT Date of Death
DTHDTF Date of Death Imputation Flag
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
DTHCAUS Cause of Death
DTHDOM Domain for Date of Death Collection
DTHCGR1 Cause of Death Reason 1
LSTALVDT Date Last Known Alive
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
AGEGR1 Pooled Age Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiralonco package (template ad_adtr.R).

References

None

Examples

```
data("adtr_onco")
```

adtte_onco	<i>Time to Event Analysis for Oncology</i>
------------	--

Description

An example of ADTTE ADaM dataset for Oncology studies

Usage

```
adtte_onco
```

Format

A data frame with 20 columns:

STUDYID Study Identifier
USUBJID Unique Subject Identifier
ADT Analysis Date
EVNTDESC Event or Censoring Description
SRCDOM Source Data
SRCVAR Source Variable
SRCSEQ Source Sequence Number
CNSR Censor
CNSDTDSC Censor Date Description
STARTDT Time-to-Event Origin Date for Subject
PARAMCD Parameter Code
PARAM Parameter
AVAL Analysis Value
ASEQ Analysis Sequence Number
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
AGE Age
SEX Sex

Source

Generated from admiralonco package (template ad_adtte.R).

References

None

Examples

```
data("adtte_onco")
```

 advfq_ophtha

Visual Function Questionnaire Analysis

Description

An example of ADVFQ ADaM dataset for Ophthalmology studies

Usage

```
advfq_ophtha
```

Format

A data frame with 85 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
QSSEQ Sequence Number
QSTESTCD Question Short Name
QSTEST Question Name
QSCAT Category of Question
QSSCAT Subcategory for Question
QSORRES Finding in Original Units
QSORRESU Original Units
QSSTRESC Character Result/Finding in Std Format
QSSTRESN Numeric Finding in Standard Units
QSSTRESU Standard Units
QSBFL Baseline Flag
QSDRVFL Derived Flag
VISITNUM Visit Number
VISIT Visit Name

VISITDY Planned Study Day of Visit
QSDTC Date/Time of Finding
QSDY Study Day of Finding
TRTSDT Date of First Exposure to Treatment
TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
ADT Analysis Date
ADY Analysis Relative Day
PARAMCD Parameter Code
AVAL Analysis Value
AVALC Analysis Value (C)
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
ONTRTFL On Treatment Record Flag
ABLFL Baseline Record Flag
BASE Baseline Value
CHG Change from Baseline
PCHG Percent Change from Baseline
ANL01FL Analysis Flag 01
ASEQ Analysis Sequence Number
PARAM Parameter
PARCAT1 Parameter Category 1
PARCAT2 Parameter Category 2
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment
RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex

RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Input. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Input. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiralophtha package (template ad_advfq.R).

References

None

Examples

```
data("advfq_ophtha")
```

 advs

Vital Signs Analysis

Description

An example of ADVS ADaM dataset

Usage

advs

Format

A data frame with 101 columns:

STUDYID Study Identifier
DOMAIN Domain Abbreviation
USUBJID Unique Subject Identifier
VSSEQ Sequence Number
VSTESTCD Vital Signs Test Short Name
VSTEST Vital Signs Test Name
VSPOS Vital Signs Position of Subject
VSORRES Result or Finding in Original Units
VSORRESU Original Units
VSSTRESC Character Result/Finding in Std Format
VSSTRESN Numeric Result/Finding in Standard Units
VSSTRESU Standard Units
VSSTAT Completion Status
VSLOC Location of Vital Signs Measurement
VSBLFL Baseline Flag
VISITNUM Visit Number
VISIT Visit Name
VISITDY Planned Study Day of Visit
VSDTC Date/Time of Measurements
VSDY Study Day of Vital Signs
VSTPT Planned Time Point Name
VSTPTNUM Planned Time Point Number
VSELTM Planned Elapsed Time from Time Point Ref
VSTPTREF Time Point Reference
TRTSDT Date of First Exposure to Treatment

TRTEDT Date of Last Exposure to Treatment
TRT01A Actual Treatment for Period 01
TRT01P Planned Treatment for Period 01
ADT Analysis Date
ADY Analysis Relative Day
PARAMCD Parameter Code
AVAL Analysis Value
ATPTN Analysis Timepoint (N)
ATPT Analysis Timepoint
AVISIT Analysis Visit
AVISITN Analysis Visit (N)
DTYPE Derivation Type
ONTRTFL On Treatment Record Flag
ANRLO Analysis Normal Range Lower Limit
ANRHI Analysis Normal Range Upper Limit
A1LO Analysis Range 1 Lower Limit
A1HI Analysis Range 1 Upper Limit
ANRIND Analysis Reference Range Indicator
BASETYPE Baseline Type
ABLFL Baseline Record Flag
BASE Baseline Value
BNRIND Baseline Reference Range Indicator
CHG Change from Baseline
PCHG Percent Change from Baseline
ANL01FL Analysis Flag 01
TRTP Planned Treatment
TRTA Actual Treatment
ASEQ Analysis Sequence Number
AVALCA1N Analysis Value Category 1 (N)
AVALCAT1 Analysis Value Category 1
PARAM Parameter
PARAMN Parameter (N)
SUBJID Subject Identifier for the Study
RFSTDTC Subject Reference Start Date/Time
RFENDTC Subject Reference End Date/Time
RFXSTDTC Date/Time of First Study Treatment
RFXENDTC Date/Time of Last Study Treatment

RFICDTC Date/Time of Informed Consent
RFPENDTC Date/Time of End of Participation
DTHDTC Date/Time of Death
DTHFL Subject Death Flag
SITEID Study Site Identifier
AGE Age
AGEU Age Units
SEX Sex
RACE Race
ETHNIC Ethnicity
ARMCD Planned Arm Code
ARM Description of Planned Arm
ACTARMCD Actual Arm Code
ACTARM Description of Actual Arm
COUNTRY Country
DMDTC Date/Time of Collection
DMDY Study Day of Collection
TRTSDTM Datetime of First Exposure to Treatment
TRTSTMF Time of First Exposure Imput. Flag
TRTEDTM Datetime of Last Exposure to Treatment
TRTETMF Time of Last Exposure Imput. Flag
TRTDURD Total Treatment Duration (Days)
SCRFDT Screen Failure Date
EOSDT End of Study Date
EOSSTT End of Study Status
FRVDT Final Retrieval Visit Date
RANDDT Date of Randomization
DTHDT Date of Death
DTHADY Relative Day of Death
LDDTHELD Elapsed Days from Last Dose to Death
LSTALVDT Date Last Known Alive
AGEGR1 Pooled Age Group 1
SAFFL Safety Population Flag
RACEGR1 Pooled Race Group 1
REGION1 Geographic Region 1
LDDTHGR1 Last Dose to Death - Days Elapsed Grp 1
DTH30FL Death Within 30 Days of Last Trt Flag
DTHA30FL Death After 30 Days from Last Trt Flag
DTHB30FL Death Within 30 Days of First Trt Flag

Source

Generated from admiral package (template ad_adv.R).

References

None

Examples

```
data("adv")
```


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