Title Open Source Ophthalmic Data Sets Curated for R

Version 0.1.0

Description Open source data allows for reproducible research and helps advance our knowledge. The purpose of this package is to collate open source ophthalmic data sets curated for direct use. This is real life data of people with intravitreal injections with anti-vascular endothelial growth factor (anti-VEGF), due to age-related macular degeneration or diabetic macular edema. Associated publications of the data sets:

Fu et al. (2020) <doi:10.1001/jamaophthalmol.2020.5044>,
Fasler et al. (2019) <doi:10.1136/bmjopen-2018-027441>,
Arpa et al. (2020) <doi:10.1136/bjophthalmol-2020-317161>,
Kern et al. 2020, <doi:10.1038/s41433-020-1048-0>.

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URL https://github.com/tjebo/eyedata

BugReports https://github.com/tjebo/eyedata/issues

Encoding UTF-8

LazyData true

RoxygenNote 7.1.1

Depends R (>= 4.0)

Imports dplyr (>= 1.0.2)

NeedsCompilation no

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Description

To explore potential utility of survival analysis techniques for retrospective clinical practice visual outcomes in a retrospective cohort study with 12-year observation period.

Usage

data("amd")

Format

A data frame (tibble) with 6696 rows and 23 variables:

- **patID**: anonymised patient number
- **sex**: gender of patient (m = male, f = female)
- **age**: age at initiation of anti-VEGF therapy (50-59 years, 60-69 years, 70-79 years, 80 years and above)
- **avdays_induc**: arithmetical average of interval between injections in induction phase in days
- **ethnicity**: ethnicity of patient: asian (South East Asian), caucasian Afro Caribbean, Mixed, unknown_other
- **loaded**: induction phase was appropriately completed within 90 days (TRUE) or not (FALSE)
- **time**: days following initiation of anti-VEGF therapy
- **injgiven**: whether injection was given at appointment time or not (TRUE = injections given, FALSE = injection not given)
- **va**: visual acuity at time point in early treatment diabetic retinopathy study letter score
- **regimen**: anti-VEGF drug given is ranibizumab only or aflibercept only
- **pre2013**: anti-VEGF therapy initiated before October 2013 (TRUE) or after (FALSE) i.e. before or after the introduction of aflibercept, respectively
Study setting and design

Of 10,744 eyes with neovascular AMD receiving anti-VEGF therapy between October 2008 and February 2020, 7802 eyes met study criteria (treatment-naive, first-treated eyes starting anti-VEGF therapy). Institution: Moorfields Eye Clinic, London, UK Approval was granted by the Institutional Review Board of the hospital (ROAD17/031). The study complied with the Declaration of Helsinki.

Data source

All clinical information at Moorfields Eye Hospital is recorded within an electronic medical record (EMR) application (OpenEyes Foundation, London, UK). A SQL database (SQL Server Reporting Service, Microsoft Corporation, Richmond, USA) containing all the information from the EMR is in place and regular updates are performed overnight to keep the data warehouse up-to-date. VA is reported in ETDRS letter score (and categories for ETDRS < 1). The highest value (independent of measurement method) available at each visit was chosen.

Data

Types of data: de-identified human subjects data Information governance authorised Moorfields Eye Clinic 19/07/2018. age not provided as a continuous variable as in original analysis to facilitate de-identification.

Source

https://doi.org/10.5061/dryad.nvx0k6dqg

See Also

Associated publication: „Insights from survival analyses during 12 years of anti-vascular endothelial growth factor therapy for neovascular age-related macular degeneration” https://doi.org/10.1001/jamaophthalmol.2020.5044

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 amd2  
 Real life data of patients with neovascular AMD

Description

A dataset containing anonymized real life human subjects data on eyes with treatment naive neovascular age-related macular degeneration (AMD), which underwent intravitreal anti-VEGF therapy with ranibizumab and/or aflibercept.

Usage

data("amd2")
Format

A data frame (tibble) with 40764 rows and 7 variables:

- **patID**: Anonymized patient identifier
- **sex**: Sex of patient (m = male, f = female)
- **age0**: Age (years) at day of first appointment
- **eye**: Left or right eye of patient (r = right, l = left)
- **time**: Time in days after date of first appointment (0 = first appointment)
- **va**: Visual acuity in Early Treatment Diabetic Retinopathy Study letters
- **inj_no**: Current number of injection at appointment date

Details

The data was collected in Moorfields Eye Hospital, London, UK. (Information governance sign off Moorfields Eye Hospital 19/07/2018)

Data was accessed on the 25th May 2020

Missing values

There are two missing visual acuity entries in this data set. They result from data entry errors (ETDRS values above 100) in the original medical health records. Unfortunately, the correct VA value could not be retrieved and it was decided to assign missing values to those measurements.

Source

https://doi.org/10.5061/dryad.97r9289

See Also

Scientific article to which this data set was supplement: https://doi.org/10.1136/bmjopen-2018-027441

 amd3

Ten year neovascular AMD survival data

Description

Ten Year Survival Trends of Neovascular Age Related Macular Degeneration at First Presentation

Usage

data("amd3")
Format

A data frame (tibble) with 6696 rows and 23 variables:

- **patID** Anonymized patient identifier.
- **sex** Sex of patient (m = male, f = female)
- **time** time in days after date of first injection (0 = first injection or baseline)
- **eye** Left or right eye of patient (r = right, l = left)
- **ethnicity** Ethnicity of patient following the categories from the UK ETHNIC CATEGORY CODE 2001. (asian = Asian or Asian British, white = White, black = Black or Black British, other_unknown = Other ethnic group or unknown)
- **fellow** fellow eye requires anti-VEGF within observation period
- **ttofellow** time between baseline and fellow eye involvement in months
- **drug_switch** anti-VEGF drug agent is switched during observation period
- **ttodrugswitch** time between baseline and date of drug switch in months
- **ltfu** patient is lost to follow-up during observation period
- **ltfu_stable** of those lost to follow-up, are they stable prior to this timepoint
- **ltfu_outcome** reason for lost to follow-up (discharged, dna (did not attend last appointment), dead, transferred)
- **ttoltfu** time between baseline and loss to follow-up in months
- **stable10y** patient is stable at ten year timepoint
- **irf** presence of intraretinal fluid at last visit
- **srf** presence of subretinal fluid at last visit
- **disciform** presence of disciform scar at last visit (0 = no, 1a = type 1a, 1b = type 1b)
- **ga** presence of geographic atrophy at last visit
- **ga_foveal** presence of foveal geographic atrophy at last visit
- **ttoga** time to geographic atrophy development in months
- **injgiven** injection given (TRUE) or not given (FALSE)
- **va** visual acuity in Early Treatment Diabetic Retinopathy Study letters
- **crt** foveal thickness in microns

Central retinal thickness (CRT)

There are double CRT measurements for 1011 time points. This is because the data set curator provided both automatic measurement and manual measurement. Unfortunately, based on the information in the raw data, it is not possible to know which entry belongs to which. One possible way to adjust for this may be to create an average for both measurements. For one way to do this, see "Examples"

Missing values

There are 1388 missing visual acuity entries and 858 missing central retinal thickness entries.
Study setting and design

Retrospective cohort study of treatment-naïve, first-affected eyes with nAMD started on ranibizumab before January 1, 2009. Approval was granted by the Institutional Review Board of the hospital (ROAD17/031). The study complied with the Declaration of Helsinki.

Data source

All clinical information at Moorfields Eye Hospital is recorded within an electronic medical record (EMR) application (OpenEyes Foundation, London, UK). Information governance authorised Moorfields Eye Clinic 19/07/2018. Age not provided as a continuous variable as in original analysis to facilitate de-identification.

Source

https://doi.org/10.5061/dryad.9cnp5hqfm

See Also

Scientific article to which this data set was supplement: https://doi.org/10.1136/bjophthalmol-2020-317161

Examples

```r
library(dplyr)
amd3 %>%
group_by(patID, time) %>%
mutate(crt_av = mean(crt, na.rm = TRUE)) %>%
select(-crt) %>%
distinct()
```

amdoct

Real life OCT segmentation data of patients with AMD

Description

This CSV dataset (AMD_baseline_MEH_v1.csv) is associated with the paper Moraes et al Quantitative analysis of optical coherence tomography for neovascular age-related macular degeneration using deep learning. Ophthalmology. (2020). The dataset comprises anonymised metadata and OCT segmentation data of patients undergoing treatment for wet AMD at Moorfields Eye Hospital, London, United Kingdom. The dataset includes 2966 rows (2966 individual patients).

Usage

```r
data("amdoct")
```
Format

A data frame (tibble) with 2966 rows and 24 variables:

- **patID**  Anonymized patient identifier
- **sex**  Sex of patient (m = male, f = female)
- **ageStrat**  Stratified age (years) as factor. One missing value
- **ethnicity**  Ethnicity of patient following the categories from the UK ETHNIC CATEGORY CODE 2001. (asian = Asian or Asian British, white = White, black = Black or Black British, other_unknown = Other ethnic group or unknown)
- **eye**  Left or right eye of patient (r = right, l = left)
- **first_eye**  Is the eye the first injected eye
- **va**  Visual acuity in Early Treatment Diabetic Retinopathy Study (ETDRS) letters and categories hand motion (hm) or counting fingers (cf) for values <1. One value = 0 was not specified in the original data set.
- **inj**  Injection given (TRUE) or not
- **time**  Time in days from baseline, i.e. injection number 1
- **Neurosensory_volume_voxels**  Number of voxels segmented as the feature neurosensory retina (NSR)
- **RPE_volume_voxels**  Number of voxels segmented as the feature retinal pigment epithelium (RPE)
- **SRF_volume_voxels**  Number of voxels segmented as the feature subretinal fluid (SRF)
- **IRF_volume_voxels**  Number of voxels segmented as the feature intraretinal fluid (IRF)
- **SHRM_volume_voxels**  Number of voxels segmented as the feature subretinal hyperreflective material (SHRM)
- **Drusen_volume_voxels**  Number of voxels segmented as the feature drusen
- **sPED_volume_voxels**  Number of voxels segmented as the feature serous pigment epithelium detachment (sPED)
- **fvPED_volume_voxels**  Number of voxels segmented as the feature fibrovascular pigment epithelium detachment (fvPED)
- **HRF_volume_voxels**  Number of voxels segmented as the feature hyperreflective foci (HRF)
- **Neurosensory_thickness_um**  Thickness (measured in microns) of the segmented feature neurosensory retina (NSR)
- **IRF_thickness_um**  Thickness (measured in microns) of the segmented feature intraretinal fluid (IRF)
- **SRF_thickness_um**  Thickness (measured in microns) of the segmented feature subretinal fluid (SRF)
- **SHRM_thickness_um**  Thickness (measured in microns) of the segmented feature subretinal hyperreflective material (SHRM)
- **HRF_thickness_um**  Thickness (measured in microns) of the segmented feature hyperreflective foci (HRF)
- **CST_um**  Central subfield retinal thickness measured as the sum of NSR, IRF, SRF, SHRM, HRF thickness (measured in microns)
Missing values

There are 233 missing visual acuity entries in this data set. They result from data entry errors (ETDRS values above 100) and missing data in the original medical health records.

Study setting and design

Study setting and design: This study is a retrospective cohort study of patients that commenced anti-VEGF therapy for neovascular AMD between June 2012 and June 2017 at Moorfields Eye Hospital NHS Foundation Trust, London, UK. Approval was granted by the Institutional Review Board of the hospital (ROAD17/031). The study complied with the Declaration of Helsinki.

Data source

All clinical information at Moorfields Eye Hospital is recorded within an electronic medical record (EMR) application (OpenEyes Foundation, London, UK). A SQL database (SQL Server Reporting Service, Microsoft Corporation, Richmond, USA) containing all the information from the EMR is in place and regular updates are performed overnight to keep the data warehouse up-to-date. VA is reported in ETDRS letter score (and categories for ETDRS < 1). The highest value (independent of measurement method) available at each visit was chosen.

OCT

Each voxel equates to 2.60 x 11.72 x 47.24 microns in the A-scan, B-scan, and C-scan directions, respectively. All OCT data is captured using 3DOCT-2000 devices (Topcon Corp., Tokyo, Japan). All images comprise 512 x 885 x 128 voxels covering a volume of 6x6x2.3mm. In this dataset, only one image is available per eye. The image with the lowest segmented artifacts was chosen if multiple scans were available at the same visit.

Time variable

For first-treated eyes, time will generally equal 0. Where a scan at day 0 was not available, a scan up to 14 days prior to the first injection (i.e. up to -14 days), is used for analysis.

Segmentation

Segmentation data was output using a deep learning segmentation model described further in De Fauw et al. and Yim and Chopra et al.

Source

https://doi.org/10.5061/dryad.2rbnz57m4

See Also

Scientific article to which this data set was supplement: https://doi.org/10.1016/j.ophtha.2020.09.025
Real life data of patients with diabetic macular edema

Description

A dataset containing anonymized real life human subjects data on eyes with diabetic macular edema (DME), which underwent intravitreal anti-VEGF therapy with ranibizumab and/or aflibercept. Data was accessed on the 3rd July 2020.

Usage

data("dme")

Format

A data frame (tibble) with 40281 rows and 8 variables:

- **patID**: Anonymized patient identifier
- **sex**: Sex of patient (m = male, f = female)
- **ageStrat**: Age (years) at day of first appointment, stratified. Factor! Labels are constructed using "(a,b]" interval notation, e.g., "(60,70]" means x > 60 & x<= 70
- **ethnicity**: Ethnicity of patient following the categories from the UK ETHNIC CATEGORY CODE 2001. (asian = Asian or Asian British, white = White, black = Black or Black British, mixed = Mixed, other = Other ethnic group, unknown = Unknown)
- **eye**: Left or right eye of patient (r = right, l = left)
- **va**: Visual acuity in Early Treatment Diabetic Retinopathy Study (ETDRS) letters
- **time**: Time in days following baseline i.e. injection number 1
- **inj**: Injection given or not (TRUE = injection given, FALSE = no injection given)

Missing values

There are 18 missing visual acuity entries in this data set. They result from data entry errors (ETDRS values above 100) in the original medical health records. Unfortunately, the correct VA value could not be retrieved and it was decided to assign missing values to those measurements.

Study setting and design

Study setting and design: This study is a retrospective cohort study of diabetic patients treated for DMO by anti-VEGF at a tertiary referral center - Moorfields Eye Hospital NHS Foundation Trust, London, UK. Approval was granted by the Institutional Review Board of the hospital (ROAD17/031) - Audit registration was completed (MEH-233). The study complied with the Declaration of Helsinki and STROBE guidelines for the reporting of cohort studies.
Data source

All clinical information at Moorfields Eye Hospital is recorded within an electronic medical record (EMR) application (OpenEyes Foundation, London, UK). A SQL database (SQL Server Reporting Service, Microsoft Corporation, Richmond, USA) containing all the information from the EMR is in place and regular updates are performed overnight to keep the data warehouse up-to-date. VA is reported in ETDRS letter score. The highest value (independent of measurement method) available at each visit was chosen.

Participants

A data-warehouse query for patients that received one IVI for DMO (between March 2013 and October 2018) resulted in 3226 unique eyes from 2368 patients. Exclusion criteria were those that: (i) suffered from macular oedema secondary to other conditions than diabetes; (ii) under 18 years old; (iii) received fewer than 3 IVI; (iv) received bevacizumab, dexamethasone intravitreal implant, or fluocinolone acetonide intravitreal implant; leaving 2614 eyes of 1964 patients taken forward for analysis.

Source

https://doi.org/10.5061/dryad.pzgmsbcfw

See Also

Scientific article to which this data set was supplement: https://doi.org/10.1038/s41433-020-1048-0
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