# $\operatorname{secuTrialR}$ - a walkthrough

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# Introduction

"If I had just five minutes to chop down a tree I would spend the first two and a half minutes sharpening my axe."

— An anonymous woodsman

This R package provides functions for handling data from the clinical data management system (CDMS) secuTrial. The most important components are related to reading data exports from secuTrial into R. In brief, the package aims to enable swift execution of repetitive tasks in order to allow spending more time on the unique aspects of a dataset. It is developed and maintained by the Swiss Clinical Trial Organisation (SCTO).

This vignette will teach you how to use the **secuTrialR** package and you will likely learn quite a bit about secuTrial exports in general along the way.

# Installing

Let's get started by installing the package straight from GitHub and then loading it. For this you will need to have devtools installed. We are planning to add secuTrialR to CRAN but we are not there yet.

```
# install
devtools::install_github("SwissClinicalTrialOrganisation/secuTrialR")

# load silently
suppressMessages(library(secuTrialR))
# show secuTrialR version
installed.packages()["secuTrialR", "Version"]
#> [1] "0.8.3"
```

#### The CTU05 dataset

Before we continue with the functionalities let's briefly talk about the test data which is delivered as a part of the package. We refer to it as the CTU05 (clinical trial unit project 05) data. This dataset has been fabricated for demonstration purposes only and is not real clinical data. Principally it is made up of eight forms. These are called "surgeries", "baseline", "outcome", "treatment", "allmedi", "studyterminat", "ae" and "sae". You will see these names again later when the data has been read into R. The project setup includes most data types implementable in secuTrial. It is, however, not exhaustive. Since the data is delivered with the installation of the secuTrialR package we can point to it via the system.file() function.

If you work on your own datasets you can specify a path as a regular character string without using system.file().

# secuTrial export options

Prior to reading your data into R you need to export it with the secuTrial ExportSearchTool. We suggest exporting non-rectangular, zipped, English data with reference values stored in a separate table including Add-IDs, centre information, form status, project setup and without duplicated meta data. Furthermore, it is important to use "CSV format"/"CSV format for MS Excel" and suggested to select UTF-8 encoding. Most of these options are truly optional and reading your data should work even with differences from the above specifications.

# Reading a secuTrial data export into R

There is one principle function to read your data (i.e. read\_secuTrial()). Below you can see it in action with the CTU05 dataset.

```
ctu05_data <- read_secuTrial(data_dir = ctu05_data_location)
#> Read export successfully.
```

If the "Read export successfully." message appears your data was correctly read.

# The secuTrialdata object

If you inspect the class() of ctu05\_data you will find that it is a secuTrialdata object.

```
class(ctu05_data)
#> [1] "secuTrialdata"
```

Really this is only a list containing all the information from your secuTrial export.

```
typeof(ctu05_data)
#> [1] "list"
```

#### The data tables in the secuTrialdata object

We have implemented a custom variation of the print() function for secuTrialdata objects.

```
print(ctu05 data)
#> secuTrial data imported from:
#> /home/wrightp/R/x86 64-pc-linux-qnu-library/3.6/secuTrialR/extdata/sT exports/
#> export_options/s_export_CSV-xls_CTU05_20191003-144349_all_info.zip
#>
              table nrow ncol meta
                                          original_name
#>
                     4 12 TRUE
                                                 vp.xls
                           2 TRUE
#>
                       8
                                               upfs.xls
               vpfs
                            8 TRUE
                       8
#>
                 fs
                                                 fs.xls
#>
                      29
                            8 TRUE
                                                 qs.xls
                 qs
                      85
                           9 TRUE
#>
                 is
                                                 is.xls
#>
                ctr
                      3
                           3 TRUE
                                                ctr.xls
#>
                      11
                           15 TRUE
                                                 cn.xls
                 cn
#>
                       0
                           9 TRUE
                                               atcn.xls
               atcn
                       0
                           16 TRUE
#>
              atcvp
                                              atcup.xls
#>
                       0
                           12 TRUE
                                                qac.xls
                qac
#>
                           10 TRUE
                                                cts.xls
                cts
```

```
#>
                              10
                                 TRUE
                                                     miv.xls
                 miv
#>
                             15 FALSE
               atmiv
                         0
                                                   atmiv.xls
                            107 FALSE
#>
            baseline
                        17
                                               baseline.xls
#>
          atbaseline
                         0
                             79 FALSE
                                             atbaseline.xls
#>
             outcome
                        13
                             48 FALSE
                                                outcome.xls
#>
           atoutcome
                         5
                             47 FALSE
                                              atoutcome.xls
#>
           treatment
                        11
                             44 FALSE
                                              treatment.xls
#>
        attreatment
                         0
                             45 FALSE
                                            attreatment.xls
#>
                        17
                             47 FALSE
                                                allmedi.xls
             allmedi
                         2
#>
           atallmedi
                             47 FALSE
                                              atallmedi.xls
                                          studyterminat.xls
#>
      studyterminat
                        10
                             41 FALSE
#>
    atstudyterminat
                         1
                             39 FALSE atstudyterminat.xls
#>
                              66 FALSE
                         1
                                                      ae.xls
#>
                         0
                             58 FALSE
                                                    atae.xls
                atae
#>
                         2
                             64 FALSE
                                                     sae.xls
                 sae
#>
               atsae
                         0
                             57 FALSE
                                                  atsae.xls
#>
                        18
                              11 FALSE
                                             esurgeries.xls
          esurgeries
#>
                         0
                               9 FALSE
       atesurgeries
                                           atesurgeries.xls
#>
                       205
                               3
                                  TRUE
                                                      cl.xls
                  cl
                             20 FALSE
#>
               atae1
                         0
                                                  atae1.xls
```

It shows you where the export archive of your secuTrialdata object is located, tells you which tables (i.e. table) it contains, what the source files (i.e. original\_name) are and specifies each table's dimensions (i.e. ncol, nrow).

By now you have possibly realized that all the forms specified earlier (i.e. "surgeries", "baseline", "outcome", "treatment", "allmedi", "studyterminat", "ae" and "sae") are present, but also that there are many tables which you do not recognize.

The majority of the unrecognizable tables are tagged as TRUE in the meta column. This means that they are structural data tables. Their names and data structures are fixed in secuTrial exports. In the following we will briefly explain which information the most relevant meta tables contain.

- vp visitplan definition
- vpfs visitplan form linkage
- fs forms information
- qs questions
- is items i.e. variable definitions
- ctr centre information
- cn casenodes i.e. table of entered study participants
- cl information how the data in the variables is coded

Furthermore, there is a set of tables whose names start with "at". These are audit trail tables and only relevant if you need to investigate changes in the data. Last but not least you may have also realized that the "surgeries" table is called esurgeries. This is because it is a so-called repetition form. Repetition forms are labelled with a leading "e" and are implemented as subforms in other forms. In this case, esurgeries is a subform in baseline and the linkage is defined by the mnpdocid column in both tables. If this sounds cryptic to you we suggest you talk so someone who has implemented a database in secuTrial and let them explain it with a specific example. It is pretty straight forward when you look at a concrete implementation.

# Accessing the tables and values

Since the secuTrialdata object is a list and the data tables within this list are data.frames you can simply access the tables using \$. Let's say you would like to have a look at the placebo to verum ratio in your treatment data or what types of other medication were entered in allmedi.

```
table(ctu05_data$treatment$rando_treatment)
#>
#> Placebo Verum
#> 8 3
table(ctu05_data$allmedi$med_product)
#>
#> Amoxicillin Doxycycline Importal
#> 2 1 3
```

#### Data transformations

During the loading process, coded categorical data is transformed. For example the gender variable in the baseline form is categorical. The raw data is accessible via gender and the transformed version of the data is added during the reading process and accessible via gender.factor. Thus, data is not overwritten but added with the .factor extension. If there are issues during factorization a warning() will inform you of this.

```
# raw gender data
ctu05_data$baseline$gender
#> [1] 1 NA NA 2 1 2 1 NA NA 1 2 NA NA 1 2 2 NA
#> attr(,"label")
#> [1] "Gender"
# transformed gender data
ctu05_data$baseline$gender.factor
#> [1] male
              <NA>
                     <NA> female male female male
                                                       <NA>
                                                              <NA>
                                                                     male
#> [11] female <NA>
                     <NA> male female female <NA>
#> attr(,"label")
#> [1] Gender
#> Levels: male female
# raw more meds
ctu05_data$allmedi$no_more_meds
#> [1] 1 1 0 1 1 1 0 0 0 1 0 0 1 1 1 1 0
#> attr(,"label")
#> [1] "No further medication"
# transformed more meds
ctu05_data$allmedi$no_more_meds.factor
#> [1] yes yes no yes yes yes no no yes no no yes yes yes yes no
#> attr(,"label")
#> [1] No further medication
#> Levels: no yes
```

Note that descriptive labels have also been automatically added to the data.

```
label(ctu05_data$allmedi$no_more_meds.factor)
#> [1] "No further medication"
label(ctu05_data$baseline$gender.factor)
#> [1] "Gender"
label(ctu05_data$esurgeries$surgery_organ.factor)
#> [1] "Organ"
```

Datetime data is also transformed and similarly to the factorization process the names are concatenated with .date or .datetime.

```
# raw
ctu05_data$baseline$visit_date
#> [1] 20190401 20190402 20190403 20190402 20190403 20190404 20190405
#> [8] 20190406 20190407 20190411 20190412 20190413 20190414 20190413
#> [15] 20190414 20190415 20180501
#> attr(,"label")
#> [1] "Date of visit"
# processed
ctu05_data$baseline$visit_date.date
#> [1] "2019-04-01" "2019-04-02" "2019-04-03" "2019-04-02" "2019-04-03"
#> [6] "2019-04-04" "2019-04-05" "2019-04-06" "2019-04-07" "2019-04-11"
#> [11] "2019-04-12" "2019-04-13" "2019-04-14" "2019-04-13" "2019-04-14"
#> [16] "2019-04-15" "2018-05-01"
# raw only head
head(ctu05_data$baseline$hiv_date)
#> [1] 201903052356
                                           NA
                                                        NA
                                                                      NA
#> [6]
# processed only head
head(ctu05_data$baseline$hiv_date.datetime)
#> [1] "2019-03-05 23:56:00 CET" NA
#> [3] NA
                                 NA
#> [5] NA
                                 NA
# classes
class(ctu05_data$baseline$visit_date)
#> [1] "integer"
class(ctu05_data$baseline$visit_date.date)
#> [1] "Date"
class(ctu05_data$baseline$hiv_date)
#> [1] "numeric"
class(ctu05_data$baseline$hiv_date.datetime)
#> [1] "POSIXct" "POSIXt"
```

Depending on the setup, incomplete dates can be valid entries in a secuTrial database. Thus they will also occasionally appear in your exports. The datetime conversion does not work in these cases and NAs are created. If this happens, <code>secuTrialR</code> will warn you accordingly and you should have a closer look into the affected datetime variables and whether you would like to perform so-called date imputation.

# **Export options**

The secuTrialdata object also contains information on the export options.

```
ctu05_data$export_options
#> secuTrial version: 5.5.1.10
#> Time of export on server: 03.10.2019 - 14:43:49 (CEST)
#> Project version: (20.06.2019 - 11:22:04 (CEST))
#> Exported with short names
#> File extension: xls
#> Seperator: ' '
#> 32 files exported
     including 13 metadata elements
#> Reference values exported - factorize possible
#> Metadata elements:
#>
                type exportname available
#>
               forms
                             fs
                                      TRUE
#>
           case nodes
                                      TRUE
                             cn
#>
             centres
                             ctr
                                      TRUE
#>
               items
                            is
                                      TRUE
#>
                                      TRUE
           questions
                             qs
             queries
                                      TRUE
                             qac
#>
           visitplan
                                      TRUE
                             vp
#>
      visitplanforms
                            vpfs
                                      TRUE
#>
         at case nodes
                                      TRUE
                           atcn
#>
    at case visitplans
                           atcvp
                                      TRUE
#>
            comments
                                      TRUE
                             cts
#>
                 miv
                             miv
                                      TRUE
#>
                                      TRUE
                  cl
                              cl
```

export\_options itself is a list. If you are interested in more information than is printed you can also access it. Let's assume you would like to know the project\_name and encoding.

```
ctu05_data$export_options$project_name
#> [1] "secuTrialR example CDMA"
ctu05_data$export_options$encoding
#> [1] "UTF-8"
```

Much more information is stored in the elements of export\_options. The names of the elements should be descriptive enough to infer the contents.

```
names(ctu05_data$export_options)
#> [1] "sep"
                                                    "datetime_format"
                              "date_format"
#> [4] "date_format_meta"
                              "na.strings"
                                                    "short names"
#> [7] "is zip"
                              "is rectangular"
                                                    "audit trail"
#> [10] "column_names"
                              "lang_not_supported" "dict_items"
#> [13] "refvals separate"
                              "add id"
                                                    "lab id"
#> [16] "meta_names"
                                                    "duplicate\_meta"
                              "meta\_available"
#> [19] "all_files"
                                                    "data_names"
                              "data\_files"
#> [22] "file_end"
                              "extension"
                                                    "data\_dir"
#> [25] "secuTrial_version" "project_version"
                                                    "project_name"
#> [28] "format_info"
                              "time\_of\_export"
                                                    "encoding"
#> [31] "form_status"
                              "centre_info"
                                                    "proj_setup"
#> [34] "factorized"
                              "dated"
                                                    "labelled"
```

# Generic functions for secuTrialdata objects

Now that you understand the **secuTrialdata** object we will show you some generic functions you can use on objects of this class.

# Show the study participants

First off you may be interested in a table of participants.

```
get_participants(ctu05_data)
#>
                    mnpaid mnpctrid
                                                           mnpctrname
      mnppid
#> 1
        1204 RPACK-CBE-001
                                462
                                               Charité Berlin (RPACK)
#> 2
        1205 RPACK-CBE-002
                                462
                                               Charité Berlin (RPACK)
#> 3
       1206 RPACK-CBE-003
                                462
                                               Charité Berlin (RPACK)
                                               Charité Berlin (RPACK)
#> 4
       1207 RPACK-CBE-004
                                462
#> 5
       1208 RPACK-CBE-005
                                462
                                               Charité Berlin (RPACK)
#> 6
       1209 RPACK-INS-011
                                461
                                             Inselspital Bern (RPACK)
#> 7
       1210 RPACK-INS-012
                                461
                                             Inselspital Bern (RPACK)
#> 8
        1211 RPACK-INS-013
                                461
                                             Inselspital Bern (RPACK)
#> 9
        1212 RPACK-INS-014
                                461
                                             Inselspital Bern (RPACK)
#> 10
       1213 RPACK-INS-015
                                461
                                             Inselspital Bern (RPACK)
#> 11
        1214 RPACK-USB-123
                                441 Universitätsspital Basel (RPACK)
```

#### Recruitment over time

You can return recruitment per centre and year.

```
annual_recruitment(ctu05_data)
#>
                                Center Total 2018 2019
#> 1
                                   All
                                           11
                                                 1
                                                     10
#> 2
               Charité Berlin (RPACK)
                                            5
                                                      5
#> 3
             Inselspital Bern (RPACK)
                                            5
                                                 0
                                                      5
#> 4 Universitätsspital Basel (RPACK)
```

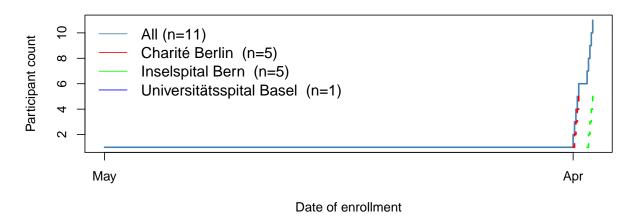
Since the centre names often have a systematic addition (e.g. (RPACK)) we have enabled the option to remove certain parts of the centre descriptions via regular expressions (i.e. rm\_regex argument). In this case the regular expression removes trailing parentheses and everything they enclose.

```
annual_recruitment(ctu05_data, rm_regex = "\\(.*\\)$")
                       Center Total 2018 2019
#> 1
                           All
                                 11
                                        1
                                            10
#> 2
               Charité Berlin
                                  5
                                        0
                                             5
#> 3
                                   5
                                        0
                                             5
             Inselspital Bern
#> 4 Universitätsspital Basel
                                   1
                                        1
```

It is also possible to plot the recruitment over time.

```
plot_recruitment(ctu05_data, cex = 1.2, rm_regex = "\\(.*\\)$")
```

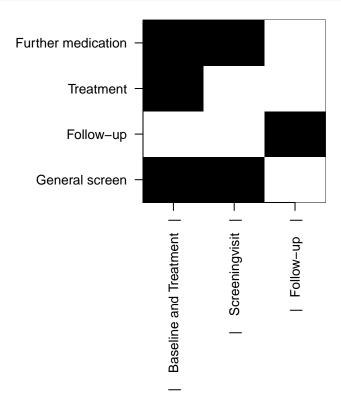
# Recruitment over time



# Visit plan visualization

secuTrialR can provide a depiction of the visit structure, although only where the visit plan is fixed.

```
vs <- visit_structure(ctu05_data)
plot(vs)</pre>
```



# Completeness of forms

If you are not sure about how complete the data in your export is, it may be useful to get a quick overview of how well the forms have been filled. The below table shows both absolute and relative numbers for a few forms.

```
fss <- form_status_summary(ctu05_data)</pre>
tail(fss, n = 5)
#>
          form_name partly_filled completely_filled empty with_warnings
#> 6
                                  3
                                                             0
                                                                            0
            baseline
                                                     14
#> 7
                                  1
                                                     12
                                                             0
                                                                            0
             outcome
#> 8
                                  0
                                                      2
                                                             0
                                                                            0
                 sae
                                  0
                                                             0
                                                                            0
#> 9
      study terminat
                                                     10
#> 10
           treatment
                                  0
                                                     11
                                                             0
                                                                            0
      with_errors partly_filled.percent completely_filled.percent
#>
                               0.17647059
#> 6
                 0
                                                             0.8235294
#> 7
                 0
                               0.07692308
                                                             0.9230769
#> 8
                 0
                               0.00000000
                                                             1.0000000
#> 9
                 0
                               0.00000000
                                                             1.0000000
#> 10
                 0
                               0.00000000
                                                             1.0000000
#>
      empty.percent with_warnings.percent with_errors.percent form_count
#> 6
                   0
                                                                            17
#> 7
                   0
                                           0
                                                                 0
                                                                            13
#> 8
                   0
                                           0
                                                                 0
                                                                             2
                   0
                                           0
#> 9
                                                                 0
                                                                            10
                   0
                                           0
#> 10
                                                                 0
                                                                            11
```

Please note that a form is only complete if all required fields have been filled. Thus, a whole study may have 99% completeness on variable basis while showing 0% completeness on form basis. It is currently not technically possible to assess completeness on variable basis in a generic way. Hence, high completeness on form basis implies high completeness on variable basis but **NOT** vice versa.

For a more participant id centred statistic you can perform the following.

```
fsc <- form_status_counts(ctu05_data)</pre>
# show the top
head(fsc)
            pat_id form_name completely_filled partly_filled empty
                                                3
#> 1 RPACK-CBE-001 baseline
                                                               0
                                                                     0
#> 2 RPACK-CBE-002
                     baseline
                                                1
                                                               0
                                                                     0
#> 3 RPACK-CBE-003
                     baseline
                                                1
                                                               0
#> 4 RPACK-CBE-004
                     baseline
                                                1
                                                                     0
                                                               2
                                                                     0
#> 5 RPACK-CBE-005
                     baseline
                                                1
#> 6 RPACK-INS-011
                                                                     0
                    baseline
#>
     with_warnings with_errors
                  0
#> 1
                               0
                               0
#> 2
                  0
#> 3
                  0
                               0
                  0
                               0
#> 4
#> 5
                               0
                  0
#> 6
```

# Form linkage

Linkages amongst forms can be explored with the links\_secuTrial() function. This relies on the igraph package to create a network. It is possible to interact with the network, e.g. move nodes around in order to read the labels better. The device ID is returned to the console, but can be ignored. Forms are plotted in deep yellow, variables in light blue.

```
links_secuTrial(ctu05_data)
```

The output can not be shown within this vignette but you should give it a try. Please note that the linkage plot is likely most useful **without** the audit trail data in the export.

#### Sampling random participants

During study monitoring it is common practice to check random participants from a study database. These participants should be retrieved in a reproducible fashion. The below function allows this for a loaded secuTrial data export.

```
# retrieve at least 25 percent of participants recorded after March 18th 2019
# from the centres "Inselspital Bern" and "Charité Berlin"
return_random_participants(ctu05_data,
                           percent = 0.25,
                           seed = 1337,
                           date = "2019-03-18",
                           centres = c("Inselspital Bern (RPACK)",
                                        "Charité Berlin (RPACK)"))
#> $participants
#>
            mnpaid
                                     centre mnpvisstartdate
#> 2 RPACK-INS-012 Inselspital Bern (RPACK)
                                                 2019-04-12
#> 4 RPACK-INS-014 Inselspital Bern (RPACK)
                                                 2019-04-14
#> 5 RPACK-CBE-005
                     Charité Berlin (RPACK)
                                                 2019-04-05
#> 3 RPACK-CBE-003
                     Charité Berlin (RPACK)
                                                 2019-04-03
#>
#> $rng_config
#> [1] "Mersenne-Twister" "Inversion"
                                              "Rejection"
```

Please note that earlier R versions may return different results because there is a different rng\_config. For this reason we have added the rng\_config to the output.

#### Retrieve score variables

Generally, it is advisable to recalculate score variables before data analysis. A rule of thumb: The more complex a score is and the more data from different forms is necessary for its calculation the more likely its value should be recalculated. The below function will allow you to detect which variables this concerns.

# Finding changes/differences in exports

In ongoing studies it is possible that changes to the secuTrial data entry interface (i.e. the electronic case report forms) are made. Sometimes these changes may call for adjustments in analysis code. It is considered good practice to run diff\_secuTrial() on the last export and the current export of a project to at least make yourself aware of potential changes in the setup. If there are differences, the results of this function should be interpreted as a first indicator since they may not cover all alterations. Information is returned on new forms and variables. A detailed list of changes can be produced in the secuTrial FormBuilder with "Compare project setup".

```
# load second export from the same project
export_location <- system.file("extdata", "sT_exports", "longnames",</pre>
                                 "s export CSV-xls CTU05 long ref miss en utf8.zip",
                                package = "secuTrialR")
# read all export data
ctu05_data_new <- read_secuTrial_raw(data_dir = export_location)</pre>
# show diff
diff_secuTrial(ctu05_data, ctu05_data_new)
#> $new_forms
#> character(0)
#>
#> $new_variables
#> character(0)
#>
#> $removed_forms
#> character(0)
#>
#> $removed_variables
#> character(0)
```

This list contains only empty entries. Thus, you can conclude that nothing has changed.

## Conversion to SPSS, STATA, SAS

Given that you are working with R it is unlikely that you need such conversions for yourself. However, collaborators may ask for data which is readily importable into SPSS, STATA or SAS. For this you can use write\_secuTrial().

```
# retrieve path to a temporary directory
tdir <- tempdir()
# write spss
write_secuTrial(ctu05_data, format = "sav", path = tdir)</pre>
```

Since this has not been heavily tested or used there may be issues and you might prefer doing this manually with the haven package.

#### Subsetting secuTrialdata

In some cases it may be useful to subset your secuTrialdata object. For example if you have cohort data and would like to supply a subset of the data for a retrospective study. We have implemented this option with subset\_secuTrial(). It will truncate your secuTrialdata object and return a new secuTrialdata object which is a subset of the original data. It is possible to subset by including or excluding specific participant ids or centres.

```
# initialize some subset identifiers
participants <- c("RPACK-INS-011", "RPACK-INS-014", "RPACK-INS-015")
centres <- c("Inselspital Bern (RPACK)", "Universitätsspital Basel (RPACK)")

# exclude Bern and Basel
ctu05_data_berlin <- subset_secuTrial(ctu05_data, centre = centres, exclude = TRUE)
# exclude Berlin
ctu05_data_bern_basel <- subset_secuTrial(ctu05_data, centre = centres)
# keep only subset of participants
ctu05_data_pids <- subset_secuTrial(ctu05_data, participant = participants)

class(ctu05_data_berlin)
#> [1] "secuTrialdata"
class(ctu05_data_bern_basel)
#> [1] "secuTrialdata"
class(ctu05_data_pids)
#> [1] "secuTrialdata"
```

If you subset based on centres all traces of deleted centres will be removed. If you remove based on participant ids all traces of deleted participants will be removed.

```
# only Berlin remains
ctu05_data_berlin$ctr
#>
   mnpctrid
                        mnpctrname mnpcname
         462 Charité Berlin (RPACK)
#> 1
# all centres remain eventhough all three ids are from Bern
ctu05_data_pids$ctr
#> mnpctrid
                                 mnpctrname mnpcname
#> 1
                    Charité Berlin (RPACK)
       462
        461 Inselspital Bern (RPACK)
#> 2
                                                  NA
       441 Universitätsspital Basel (RPACK)
#> 3
                                                  NA
```

Since the truncated object's class remains unchanged (i.e. secuTrialdata) you can still use the generic functions on it. Let's say you would only like to look at the recruitment plot for Bern alone.

```
# keep only Bern
ctu05_data_bern <- subset_secuTrial(ctu05_data, centre = "Inselspital Bern (RPACK)")
# plot
plot_recruitment(ctu05_data_bern)</pre>
```

#### Recruitment over time



... or Bern and Berlin.

#### Recruitment over time



# The as.data.frame function

This vignette has been working with the secuTrialdata object as a list. For some users, working with a list can be tiresome so secuTrialR provides an as.data.frame() method to save the data.frames in the secuTrialdata object to an environment of your choice.

Let's have a look at the state of your globalenv() before running as.data.frame()...

```
ls(globalenv())
#> [1] "centres"
                                  "ctu05_data"
#>
   [3] "ctu05_data_berlin"
                                  "ctu05\_data\_bern"
    [5] "ctu05 data bern basel"
                                  "ctu05 data bern berlin"
   [7] "ctu05_data_location"
                                  "ctu05\_data\_new"
#> [9] "ctu05_data_pids"
                                  "export_location"
#> [11] "fsc"
                                  "fss"
#> [13] "participants"
                                  "vs"
# add files to global environment
as.data.frame(ctu05_data)
```

... and afterwards.

```
ls(globalenv())
#> [1] "ae"
                                  "allmedi"
#>
   [3] "atae"
                                  "atae1"
   [5] "atallmedi"
                                  "atbaseline"
  [7] "atesurgeries"
                                  "atmiv"
#> [9] "atoutcome"
                                  "atsae"
#> [11] "atstudyterminat"
                                  "attreatment"
#> [13] "baseline"
                                  "centres"
#> [15] "ctu05 data"
                                  "ctu05\_data\_berlin"
#> [17] "ctu05_data_bern"
                                  "ctu05_data_bern_basel"
#> [19] "ctu05_data_bern_berlin" "ctu05_data_location"
#> [21] "ctu05_data_new"
                                  "ctu05\_data\_pids"
#> [23] "esurgeries"
                                  "export_location"
#> [25] "fsc"
                                  "fss"
#> [27] "outcome"
                                  "participants"
#> [29] "sae"
                                  "studyterminat"
#> [31] "treatment"
```

# A note on mnp\* variables

There is a plethora of variables in the tables of secuTrial exports whose names start with mnp. These are meta variables which are e.g. important to logically link the different tables. Explaining them all is beyond the scope of this vignette. For detailed explanations please refer to the secuTrial user manual.

#### Disclaimer

The descriptions of the export data and the logic within are our understanding of them and come with no warranty. For in depth details please refer to the original secuTrial manuals.

# The sessionInfo() of the vignette

```
sessionInfo()
#> R version 3.6.1 (2019-07-05)
#> Platform: x86_64-pc-linux-qnu (64-bit)
#> Running under: Ubuntu 18.04.3 LTS
#> Matrix products: default
#> BLAS: /usr/lib/x86_64-linux-gnu/blas/libblas.so.3.7.1
#> LAPACK: /usr/lib/x86_64-linux-gnu/lapack/liblapack.so.3.7.1
#> locale:
#> [1] LC_CTYPE=C.UTF-8
                         LC NUMERIC=C
                                             LC_TIME=C.UTF-8
\#> [4] LC\_COLLATE=C.UTF-8 LC\_MONETARY=C.UTF-8 LC\_MESSAGES=C.UTF-8 \#> [7] LC\_PAPER=C.UTF-8 LC\_NAME=C LC\_ADDRESS=C
#> [10] LC TELEPHONE=C
                         LC MEASUREMENT=C.UTF-8 LC IDENTIFICATION=C
#>
#> attached base packages:
#> [1] tcltk
            stats graphics grDevices utils datasets methods
#> [8] base
#>
#> other attached packages:
#> [1] secuTrialR_0.8.3 lubridate_1.7.4 dplyr_0.8.3
                                                  tidyr_1.0.0
#>
#> loaded via a namespace (and not attached):
#> [1] Rcpp_1.0.3 knitr_1.25 magrittr_1.5 hms_0.5.2
#> [5] tidyselect_0.2.5 R6_2.4.0
                                                 tools 3.6.1
                                   rlang_0.4.1
#> [25] cellranger_1.1.0 forcats_0.4.0 backports_1.1.5 tufte_0.5
#> [29] pkqconfiq_2.0.3
```