Package 'RLumModel'

February 24, 2016

```
Type Package
Title Modelling Ordinary Differential Equations Leading to
     Luminescence
Version 0.1.1
Date 2016-02-24
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Description A collection of functions to simulate luminescence signals in the
     mineral quartz based on published models.
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License GPL-3
Depends R (>= 3.2.3), utils, Luminescence (>= 0.5.1)
Imports deSolve (>= 1.12), methods, Rcpp
Collate RLumModel-package.R RcppExports.R calc_signal.R
     calc_concentrations.R create_DRT.sequence.R
     create_SAR.sequence.R extract_pars.R
     model_LuminescenceSignals.R read_SEQ2R.R set_ODE.R
     set_ODE_LM_OSL.R set_pars.R simulate_CW_OSL.R simulate_LM_OSL.R
     simulate RF.R simulate TL.R simulate heating.R
     simulate_illumination.R simulate_irradiation.R simulate_pause.R
     translate_sequence.R
RoxygenNote 5.0.1
Suggests testthat, R.rsp
VignetteBuilder R.rsp
LinkingTo Rcpp, RcppArmadillo
NeedsCompilation yes
```

R topics documented:

RLumi	Model-package	Мос	lellii	ıg (Or	dii	nai	ry .	Dί	ffe	ere	nt	ia	l E	Eqi	ıai	tio	ns	Le	eaa	lin	gi	to	Lı	ım	in	esc	cer	ıсе	?
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Author(s)

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Project source code repository

https://github.com/R-Lum

Related projects

```
http://www.r-luminescence.de
http://cran.r-project.org/package=Luminescence
```

http://shiny.r-luminescence.de

http://cran.r-project.org/package=RLumShiny

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Acknowledgement

The work of Johannes Friedrich is gratefully supported by the DFG in framework of the project 'Modelling quartz luminescence signal dynamics relevant for dating and dosimetry' (SCHM 305114-1)

ExampleData.ModelOutput

Example data (TL curve) simulated from Bailey (2001, fig. 1)

Description

Example data (TL curve) simulated from Bailey (2001, fig. 1)

Format

A RLum. Analysis object containing one TL curve as RLum. Data. Curve.

Note

```
This example has only one record (TL). The used sequence was sequence <- list(IRR = c(temp = 20, dose = 10, DoseRate = 1), TL = c(temp_begin = 20, temp_end = 400, heating_rate = 5))
```

Source

model_LuminescenceSignals()

References

Bailey, R.M., 2001. Towards a general kinetic model for optically and thermally stimulated luminescence of quartz. Radiation Measurements 33, 17-45.

Examples

```
data(ExampleData.ModelOutput)
TL_curve <- get_RLum(model.output, recordType = "TL$", drop = FALSE)
##plot TL curve
plot_RLum(TL_curve)

TL_concentrations <- get_RLum(model.output, recordType = "(TL)", drop = FALSE)
plot_RLum(TL_concentrations)</pre>
```

model_LuminescenceSignals

Model Luminescence Signals

Description

This function models luminescence signals for quartz based on published physical models. It is possible to simulate TL, (CW-) OSL, RF measurements in a arbitrary sequence. This sequence is definded as a list of certain abrivations. Furthermore it is possible to load a sequence direct from the Riso Sequence Editor. The output is an RLum. Analysisobject and so the plots are done by the plot_RLum. Analysis function. If a SAR sequence is simulated the plot output can be disabled and SAR analyse functions can be used.

Usage

```
model_LuminescenceSignals(model, sequence, lab.dose_rate = 1,
    simulate_sample_history = FALSE, plot = TRUE, verbose = TRUE,
    show.structure = FALSE, ...)
```

Arguments

model character (**required**): set model to be used. Available models are: "Bai-

ley 2001", "Bailey 2002", "Bailey 2004", "Pagonis 2007", "Pagonis 2008"

sequence list (required): set sequence to model as list or as *.seq file from the Riso

sequence editor. To simulate SAR measurements there is an extra option to set

the sequence list (cf. details).

lab.dose_rate numeric (with default): laboratory dose rate in XXX Gy/s for calculating sec-

onds into Gray in the *.seq file.

simulate_sample_history

logical (with default): FALSE (with default): simulation begins at laboratory conditions, TRUE: simulations begins at crystallization (all levels 0) process

plot logical (with default): Enables or disables plot output

verbose logical (with default): Verbose mode on/off

show.structure logical (with default): Shows the structure of the result. Recommended to show record.id to analyse concentrations.

... further arguments and graphical parameters passed to plot.default. See details for further information.

Details

Defining a sequence

Arguments	Description	Sub-arguments
TL	thermally stimulated luminescence	'temp begin', 'temp end', 'heating rate'
OSL	optically stimulated luminescence	'temp', 'duration', 'optical_power'
ILL	illumination	'temp', 'duration', 'optical_power'
LM_OSL	linear modulated OSL	'temp', 'duration', optional: 'start_power', 'end_power'
RL/RF	radioluminescence	'temp', 'dose', 'dose_rate'
IRR	irradiation	'temp', 'dose', 'dose_rate'
CH	cutheat	'temp', optional: 'duration', 'heating_rate'
PH	preheat	'temp', 'duration' optional: 'heating_rate'
PAUSE	pause	'temp', 'duration'

Note: 100 % illumination power equates to 20 mW/cm^2

Defining a SAR-sequence

Abrivation	Description	examples
RegDose	Dose points of the regenerative cycles	c(0, 80, 140, 260, 320, 0, 80)
TestDose	Test dose for the SAR cycles	50
PH	Temperature of the preheat	240
CH	Temperature of the cutheat	200
OSL_temp	Temperature of OSL read out	125
OSL_duration	Duration of OSL read out	default: 40
Irr_temp	Temperature of irradiation	default: 20
PH_duration	Duration of the preheat	default: 10
dose_rate	Dose rate of the laboratory irradiation source	default: 1
optical_power	Percentage of the full illumination power	default: 90
Irr_2recover	Dose to be recovered in a dose-recovery-test	20

Value

This function returns an RLum. Analysis object with all TL, (LM-) OSL and RF/RL steps in the sequence. Every entry is an RLum. Data. Curve object and can be plotted, analysed etc. with further RLum-functions.

Function version

0.1.0

Author(s)

Johannes Friedrich, University of Bayreuth (Germany), Sebastian Kreutzer, IRAMAT-CRP2A, Universite Bordeaux Montaigne (France)

References

Bailey, R.M., 2001. Towards a general kinetic model for optically and thermally stimulated luminescence of quartz. Radiation Measurements 33, 17-45.

Bailey, R.M., 2002. Simulations of variability in the luminescence characteristics of natural quartz and its implications for estimates of absorbed dose. Radiation Protection Dosimetry 100, 33-38.

Bailey, R.M., 2004. Paper I-simulation of dose absorption in quartz over geological timescales and it simplications for the precision and accuracy of optical dating. Radiation Measurements 38, 299-310.

Pagonis, V., Chen, R., Wintle, A.G., 2007: Modelling thermal transfer in optically stimulated luminescence of quartz. Journal of Physics D: Applied Physics 40, 998-1006.

Pagonis, V., Wintle, A.G., Chen, R., Wang, X.L., 2008. A theoretical model for a new dating protocol for quartz based on thermally transferred OSL (TT-OSL). Radiation Measurements 43, 704-708.

Soetaert, K., Cash, J., Mazzia, F., 2012. Solving differential equations in R. Springer Science & Business Media.

See Also

```
plot, RLum, read_SEQ2R
```

Examples

```
##=========##
## Example 1: Simulate sample history of Bailey2001
## (cf. Bailey, 2001, Fig. 1)
##============##

##set sequence with the following steps
## (1) Irradiation at 20 deg. C with a dose of 10 Gy and a dose rate of 1 Gy/s
## (2) TL from 20-400 deg. C with a rate of 5 K/s

sequence <-
list(
    IRR = c(20, 10, 1),
    TL = c(20, 400, 5)
)

##model sequence
model.output <- model_LuminescenceSignals(
    sequence = sequence,
    model = "Bailey2001"
)</pre>
```

```
##get all TL concentrations
TL_conc <- get_RLum(model.output, recordType = "(TL)", drop = FALSE)</pre>
plot_RLum(TL_conc)
##plot 110 deg. C trap concentration
plot_RLum(TL_conc, recordType = "conc. level 1")
##============================##
## Example 2: compare different optical powers of stimulation light
# call function "model_LuminescenceSignals", model = "Bailey2004"
# and simulate_sample_history = FALSE (default),
# because the sample history is not part of the sequence
# the optical_power of the LED is varied and then compared.
optical_power <- seq(from = 0, to = 100, by = 20)
model.output <- lapply(optical_power, function(x){</pre>
 sequence <- list(IRR = c(20, 50, 1),
                PH = c(220, 10, 5),
                OSL = c(125, 50, x)
                )
 data <- model_LuminescenceSignals(</pre>
         sequence = sequence,
          model = "Bailey2004",
          plot = FALSE,
          verbose = FALSE
return(get_RLum(data, recordType = "OSL$", drop = FALSE))
})
##combine output curves
model.output.merged <- merge_RLum(model.output)</pre>
##plot
plot_RLum(
object = model.output.merged,
xlab = "Illumination time [s]",
ylab = "OSL signal [a.u.]",
main = "OSL signal dependency on optical power of stimulation light",
legend.text = paste("Optical power density", 20*optical_power/100, "mW/cm^2"),
combine = TRUE)
## Not run:
##============================##
```

```
## Example 3: Simulate Thermal-Activation-Characteristics (TAC)
##-----##
##set temperature
act.temp <- seq(from = 80, to = 600, by = 20)
##loop over temperature
model.output <- vapply(X = act.temp, FUN = function(x) {</pre>
##set sequence, note: sequence includes sample history
  sequence <- list(</pre>
    IRR = c(20, 1, 1e-11),
    IRR = c(20, 10, 1),
    PH = c(x, 1),
    IRR = c(20, 0.1, 1),
    TL = c(20, 150, 5)
)
##run simulation
  temp <- model_LuminescenceSignals(</pre>
    sequence = sequence,
    model = "Pagonis2007",
    simulate_sample_history = TRUE,
    plot = FALSE,
    verbose = FALSE
    ## "TL$" for exact matching TL and not (TL)
  TL_curve <- get_RLum(temp, recordType = "TL$")</pre>
  ##return max value in TL curve
  return(max(get_RLum(TL_curve)[,2]))
}, FUN. VALUE = 1)
##plot resutls
plot(
  act.temp[-(1:3)],
  model.output[-(1:3)],
  type = "b",
  xlab = "Temperature [\u00B0C]",
  ylab = "TL [a.u.]"
##-----##
## Example 4: Simulate SAR sequence
##-----##
##set SAR sequence with the following steps
## (1) RegDose: set regenerative dose [Gy] as vector
## (2) TestDose: set test dose [Gy]
## (3) PH: set preheat temperature in deg. C
## (4) CH: Set cutheat temperature in deg. C
## (5) OSL_temp: set OSL reading temperature in deg. C
## (6) OSL_duration: set OSL reading duration in s
sequence <- list(</pre>
```

```
RegDose = c(0,10,20,50,90,0,10),
TestDose = 5,
PH = 240,
CH = 200,
OSL_{temp} = 125,
OSL_duration = 70)
# call function "model_LuminescenceSignals", set sequence = sequence,
# model = "Pagonis2007" (palaeodose = 20 Gy) and simulate_sample_history = FALSE (default),
# because the sample history is not part of the sequence
model.output <- model_LuminescenceSignals(</pre>
sequence = sequence,
model = "Pagonis2007",
plot = FALSE
)
# in environment is a new object "model.output" with the results of
# every step of the given sequence.
# Plots are done at OSL and TL steps and the growth curve
# call "analyse_SAR.CWOSL" from RLum package
results <- analyse_SAR.CWOSL(model.output,</pre>
                           signal.integral.min = 1,
                           signal.integral.max = 15,
                           background.integral.min = 601,
                           background.integral.max = 701,
                           fit.method = "EXP",
                           dose.points = c(0,10,20,50,90,0,10))
##============================##
## Example 5: generate sequence from *.seq file and run SAR simulation
##============================##
# load example *.SEQ file and construct a sequence.
# call function "model_LuminescenceSignals", load created sequence for sequence,
# set model = "Bailey2002" (palaeodose = 10 Gy)
# and simulate_sample_history = FALSE (default),
# because the sample history is not part of the sequence
path <- system.file("extdata", "example_SAR_cycle.SEQ", package="RLumModel")</pre>
sequence <- read_SEQ2R(file = path)</pre>
model.output <- model_LuminescenceSignals(</pre>
 sequence = sequence,
 model = "Bailey2001",
 plot = FALSE
)
```

```
## call RLum package function "analyse_SAR.CWOSL" to analyse the simulated SAR cycle
results <- analyse_SAR.CWOSL(model.output,</pre>
                            signal.integral.min = 1,
                            signal.integral.max = 10,
                            background.integral.min = 301,
                            background.integral.max = 401,
                            dose.points = c(0,8,14,26,32,0,8),
                            fit.method = "EXP")
print(get_RLum(results))
## Example 6: Simulate sequence at labour without sample history
##===========================##
##set sequence with the following steps
## (1) Irraditation at 20 deg. C with a dose of 100 Gy and a dose rate of 1 Gy/s
## (2) Preheat to 200 deg. C and hold for 10 s
## (3) LM-OSL at 125 deg. C. for 100 s
## (4) Cutheat at 200 dec. C.
\#\# (5) Irraditation at 20 deg. C with a dose of 10 Gy and a dose rate of 1 Gy/s
## (6) Pause at 200 de. C. for 100 s
## (7) OSL at 125 deg. C for 100 s with 90 % optical power
## (8) Pause at 200 deg. C for 100 s
## (9) TL from 20-400 deg. C with a heat rate of 5 K/s
## (10) Radiofluorescence at 20 deg. C with a dose of 200 Gy and a dose rate of 0.01 Gy/s
sequence <-
list(
  IRR = c(20, 100, 1),
  PH = c(200, 10),
  LM_{OSL} = c(125, 100),
  CH = c(200),
  IRR = c(20, 10, 1),
  PAUSE = c(200, 100),
  OSL = c(125, 100, 90),
  PAUSE = c(200, 100),
  TL = c(20, 400, 5),
  RF = c(20, 200, 0.01)
)
# call function "model_LuminescenceSignals", set sequence = sequence,
# model = "Pagonis2008" (palaeodose = 200 Gy) and simulate_sample_history = FALSE (default),
# because the sample history is not part of the sequence
model.output <- model_LuminescenceSignals(</pre>
   sequence = sequence,
  model = "Pagonis2008"
  )
```

read_SEQ2R

```
## End(Not run)
```

read_SEQ2R Parse a Risoe SEQ-file to a sequence neccessary for luminescence	for simulating quartz
---	-----------------------

Description

A SEQ-file created by the Risoe Sequence Editor can be imported to simulate the sequence written in the sequence editor.

Usage

```
read_SEQ2R(file, lab.dose_rate = 1, txtProgressBar = TRUE)
```

Arguments

file character (required): a *.seq file created by the Risoe Sequence Editor

lab.dose_rate character (with default): set the dose rate of the radiation source in the labo-

ratory [Gy/s]. Default: 1 Gy/s

txtProgressBar logical (with default): enables or disables the txtProgressBar for a visuell con-

trol of the progress. Default: txtProgressBar = TRUE

Details

Supported versions

Suppored and tested: version 4.36.

Value

This function returns a list with the parsed *.seq file and the required steps for model_LuminescenceSignals.

Function version

0.1.0

Author(s)

Johannes Friedrich, University of Bayreuth (Germany),

References

Riso: Sequence Editor User Manual. Available at: http://www.nutech.dtu.dk/english/-/media/Andre_Universitetsenheder/Nutech/Produkter%20og%20services/Dosimetri/radiation_measurement_instruments/tl_osl_reader/Manuals/SequenceEditor.ashx?la=da

read_SEQ2R

See Also

```
{\tt model\_LuminescenceSignals, readLines}
```

Examples

```
##search "example_SAR_cycle.SEQ" in "extdata" in package "RLumModel"
path <- system.file("extdata", "example_SAR_cycle.SEQ", package="RLumModel")
sequence <- read_SEQ2R(file = path)</pre>
```

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