LMU	LUDWIG- MAXIMILIANS- UNIVERSITÄT MÜNCHEN	
-----	---	--

RUHR UNIVERSITÄT BOCHUM

# SimThyr 4.0

# Handbook and Reference



Johannes W. Dietrich Documentation version 4.0.3 Last Change: 2022-04-23, 12:54:31



#### Preface

Thyroid hormones act on virtually any cell of the vertebrate organism. They play a central role in metabolic coordination and differentiation. Robust control of thyroid homeostasis is therefore essential for survival, differentiation and reproduction [1].

In 1936, only seven years after the discovery of the thyrotropic hormone TSH by Aron in the year 1929, Karl Fellinger suggested a negative feedback effect of thyroid hormones on thyrotropic cells of the pituitary [2]. W. T. Salter probably independently postulated the existence of a control loop linking the pituitary and thyroid gland in 1940. His idea was inspired by the then recent description of the two gonadotropic feedback control loops. Only a few years later, Astwood and Hoskins could reveal both the existence and the pathophysiological relevance of this thyrotropic feedback control system. In 1968, Panda and Turner succeeded in a quantitative description of the interdependence of thyroxine and TSH levels, but the first mathematical models of the feedback loop had already been developed up to 12 years earlier [3-6].

Over the years, cybernetic models of thyroid homeostasis were increasingly improved, a success that was enabled by improved mathematical modelling methodology as well as growing empirical data. As a consequence of this continued evolution some of the newest models are also suitable for medical decision-making [1, 7-11]. However, due to nonlinear interactions, none of these cybernetic models can simultaneously describe static and dynamic behaviour. This gap may be filled with computer simulations that additionally facilitate an intuitive insight into the dynamics of thyrotropic feedback.

One of the more popular simulation programs for the pituitary thyroid feedback control is SimThyr. Based on a parametrically isomorphic model of the overall system [1, 12-15] SimThyr looks back on more than 25 years of continued development and a plethora of publications cited in PubMed and Web of Science [11-16]. Today, SimThyr is a user-friendly "virtual thyroid" program for modern desktop and server operating systems. Applications of SimThyr cover research, including development of hypotheses, and education of students in biology and medicine, nurses and patients.

Software, source code and additional documentation of SimThyr are available from http://simthyr.sf.net.

# Contents

Preface	2
Introduction	5
System Requirements	5
macOS / Mac OS X	5
Windows	5
Mac OS Classic	5
Linux	5
Installation	5
Uninstalling SimThyr	6
Source Code	6
SimThyr Guided Tour	7
Starting SimThyr	7
First simulation	7
Experiment 1	8
Experiment 2 1	1
User Interface1	3
Toolbar 1	3
Main Menu 1	4
Scenarios1	15
Sensitivity Analysis1	6
Preferences1	8
Supported File Formats1	9

Appendix	
References	
Contact	

### Introduction

SimThyr is a continuous simulator for pituitary-thyroid homeostasis. Pre-compiled Software is available for macOS (formerly referred to as Mac OS X or OS X), Mac OS Classic, Windows and ReactOS, but it can also be compiled for popular Linux distributions and several Unix derivatives. SimThyr was developed with Lazarus / Free Pascal (macOS, Linux, Windows and ReactOS) or THINK Pascal (Mac OS Classic), respectively. Source code is provided for all supported platforms and may also be used in order to compile SimThyr for additional platforms.

This document describes the standard version of SimThyr for graphical user interfaces (GUI). Please refer to the handbook and reference information for SimThyr CLI for usage of the version for command line interfaces (CLI).

# System Requirements

#### macOS / Mac OS X

SimThyr 4.0 for Mac supports Mac OS X 10.4 (Tiger) or newer on systems with either Intel or PPC processors. Currently ARM processors (Apple Silicon) are supported via the Rosetta 2 engine.

#### Windows

SimThyr 4.0 for Windows has been developed for Windows XP and was successfully tested also on Windows 2000, Windows 7, Windows 10 and ReactOS.

#### Mac OS Classic

Mac OS Classic (6.0.7 or newer) is supported by the legacy version SimThyr 2.0 only.

#### Linux

In order to use SimThyr on Linux please download the source code and compile it for your processor and operating system with Lazarus / Free Pascal.

#### Installation

For installing the program on Mac computers, simply drag the application file from the disk image (or BinHex package for legacy SimThyr 2.0) to the desired location on your file system. SimThyr for Mac doesn't come with and doesn't require an installer.

The recommended way to install SimThyr on Windows-based computers is to use the provided installer. For special requirements, e.g. for installation on computers without administrative rights, a portable application is available, too. In order to install the portable version simply drag the application file from the archive to the desired location on your file system. You may also start portable SimThyr directly from a USB stick, a CD and even a web server (depending on the settings of your browser). This may have advantages e.g. for CBT labs and other computer resource centres.

On Linux, please download the source code and compile it for your processor and operating system with Lazarus / Free Pascal.

# Uninstalling SimThyr

Before uninstalling SimThyr make sure that the application is not running. Then follow the directions of your operating system to delete the application file (on macOS this can be easily done by dragging the file representation to the trash can icon).

As required you may also want to delete scenarios and preferences files.

# Source Code

Source code is available from http://sourceforge.net/projects/simthyr. For compiling SimThyr from source please follow the directions given there.

#### SimThyr Guided Tour

#### Starting SimThyr

Start SimThyr by double-clicking its icon. After a short while a dialog box asking for simulation options is presented:



#### First simulation

For a first simulation run simply hit the **Start** button on the dialog box or press the **Enter** key on your keyboard. This will start a simulation of one day under standard conditions.



The Log window shows simulated values of time and hormone levels. In the plot window results are presented graphically as time series. The small window titled "Prediction" displays predicted results from analytically solving the equations describing the feedback loop in a parametrically isomorphic model. Small deviations from the depicted results above may ensue from simulated random effects.

#### Experiment 1

The first experiment simulates the development of subclinical (latent) hypothyroidism.

In order to do this, first click the stop and reset button in the toolbar:



This will delete all simulated values and set the time to zero. Then, start a new simulation over 14 days. Press the start button:



and enter 14 days in the simulation settings dialog:

W	Basic settings	for next simulation run:	SimThyr 4.0.0.836
Ð	Simulated time		
Duration:	14	Days 💙	
t.c.	Optional test si	gnal:	
	Off	200 μg TRH 50% T	BG rise
Starts at:	0	Hours	

After clicking "Start" you will obtain the result of 14 days in normal conditions that may be similar but not necessarily identical as depicted below.



Now, let's develop subclinical hypothyroidism. In order to achieve this, click the start button again *without* hitting the stop button first. This will cause the subsequent simulation results to be appended to the previous results.

In the simulation settings dialog don't click the start button. Rather click on the button titled **Structure parameters**... in the middle of the lower part. This will open a new dialog enabling you to modify structure parameters of the overall feedback system. Here, change the thyroid's secretory capacity (GT) to 1.2 pmol/sec, i.e. you should enter 1.2E-12 into the field depicted GT.

			Sec.				<b>→</b> (	Ô.⊷⊓	
AlphaR:	0.4	/I	7	GR:	1	mol/s		Ŧ -	]
BetaR:	0.0023105	/s	2	DR:	1E-10	mol/l			
AlphaS:	0.4	/I				_			
BetaS:	0.00023	/s		GH:	471.85	mU/s	SS:	100	l/ml
AlphaS2:	2.6E5	/I		DH:	4.7E-8	mol/l	DS:	50	mU/
BetaS2:	140	/s		LS:	1.6879E6	l/mol			
AlphaT:	0.1	/I		GT:	1.2E-12	mol/s			
BetaT:	1.1E-6	/s	1	DT:	2.75	mU/I			
Alpha31:	0.026	/1		GD1:	2.8E-8	mol/s	Disso	ciation c	onstants:
Beta31:	8E-6	/s		KM1	5E-7	mol/l	K30:	2E9	l/mo
							K31:	2E9	I/mo
Alpha32:	1.3E5	/I	100	GD2	4.3E-15	mol/s	K41:	2E10	l/mo
Beta32:	0.00083	/s	~	KM2	1E-9	mol/l	K42:	2E8	l/mo
Time con	stants:								Options
Tau0 R:	1800	s	t.c.					ſ	Chandrad
Tau0 S:	120	s		Tau0 '	T: 300	s		L	Standard
Tau0 S2:	3240	s		Tau0	3z: 3600	s			Cancel
First orde	r time consta	nts a	nd assoc	iated h	alf-lifes			E	011

After clicking the **OK** button the new value for GT is stored. Then change the simulated time to 21 days to allow for enough settling time. After clicking **Start** you should receive results that are similar to the following:



Obviously, FT4 falls into the low-normal range while TSH oscillates between normal and elevated levels.

In the structure parameters window you may change virtually every constant structure parameter of thyroid homeostasis including volumes of distribution, time constants and dissociation constants. Useful starting points for experimentation may be the sum activity of peripheral deiodinases (GD) and the secretory capacity of thyrotropic anterior pituitary cells (GH).

#### Experiment 2

This simulates the effects of modifications in plasma proteins, e.g. in pregnancy or on anticonceptive medication.

First, reconstruct standard conditions by clicking **Standard** in the structure parameters window and confirm with **OK**. Then, click the stop and reset button in the toolbar in order to delete all entries and start with time point 0.

After clicking the start button you will get the already well-known simulation settings dialog. Here click on the radio box titled 50% TBG rise in the lower part of the window.

Ð	Simulated time:
- Duration:	30 Days 💌
t,	Optional test signal:
Starts at:	Off 0200 µg TRH ● 50% TBG rise 7 Days ▼

To allow for the necessary settling times enter 30 days for total simulation and 7 days as beginning of the effect.



The results should be similar as depicted above, showing a decrease of FT4 levels in the beginning of the effect, than a short increase in TSH levels and a consecutive slight TSH decrease as FT4 rises in the achievement of a new equilibrium point.

You may change the combo boxes in the plot window in order to study the effects on other hormones of the feedback loop.

#### User Interface

# Toolbar

Most functions of SimThyr are accessible from the toolbar. If you move the mouse cursor to a button and leave it there for a second without clicking, a small tool tip is displayed explaining its function.



Parameters button: Modify structure parameters of feedback control here.

IPS button: Displays information processing structure.

Settings button: Set global application preferences here.



#### Main Menu

Additional options are available from the main menu. It will look slightly different on Mac OS, macOS (Mac OS X), Windows and Linux. However, the principal functions are identical.

🗯 SimThyr File Edit Simulation Window Help

SimThyr menu: Application-wide options, e.g. Preferences and Services (macOS only)

File menu: File- and scenario-wide options, e.g. for opening and saving files.

Edit menu: Standardised editing options like cut, copy and paste.

Simulation menu: Options for starting, stopping and suspending simulations as well as for changing structure parameters and for editing model annotations.

Window menu: Select background or hidden windows here.

Help menu: Help and additional information.

#### Scenarios

Simulation scenarios, e.g. containing a set of structure parameters in certain thyroid disorders, may be stored as XML files and reused. This may be a valuable option for research use, talks or educational purposes. Examples for useful scenarios are sets of structure parameters in certain thyroid disorders and influencing factors in some conditions like pregnancy or critical illness.

Saved scenarios are also an excellent medium to make *in-silico* experiments better reproducible. An increasing number of journals requires submission of open scientific data as a supplement to submitted manuscripts. SimThyr supports this endeavour by delivering the foundations of a simulation experiment in form of an XML file, which is readable by both humans and machines. In order to make the files compliant to MIRIAM and MIASE statements, the required information may be entered in the model annotations window.



Save a scenario by selecting the entry Save from the File menu and make sure that the format Scenario (XML) is selected in the save dialog.

You can open and reuse an existing scenario by selecting **Open Scenario**... from the File menu.

Scenario files are cross-compatible for all subversions of SimThyr 3.x and 4.x on all supported platforms. Despite the addition of additional information in SimThyr 4.0 the file format is backwards-compatible to SimThyr 3.x. Scenario XML files created with the GUI version of SimThyr can also be read by the CLI version.

#### Sensitivity Analysis

In its essence, cybernetic modelling maps a system's structure onto its behaviour. This applies to both mathematical descriptions and simulation. A more methodical implementation of this approach is sensitivity analysis, a very powerful method in systems science.

Sensitivity analysis systematically varies a structure parameter and investigates resulting behavioural variables. This helps to study the influence of certain marginal conditions on relevant output variables and to define factors that are critical for definite wanted (or unwanted) results. Additionally, this method allows assessing how the uncertainty of behavioural variables can be apportioned to different sources of uncertainty in model input.



Two-way sensitivity analysis



One-way sensitivity analysis, two-way sensitivity analysis, tornado plots and nullcline plots (equilibrium diagrams) may be accessed from the Simulation menu. Usage should be self-explanatory. In any case of classical sensitivity analysis select independent structure parameter(s) and dependent behavioural parameter(s) (i.e. hormone levels). In null isocline plots select two hormones, the interaction of which you want to study. The resulting plots may be modified with respect to colour and (for one-way and two-way sensitivity analysis) for lower and upper boundaries of structure parameters.

Form of sensitivity analysis	Kind of relation	Comments
One-way	Multivariate (one independent	Maps selected hormone
	and <i>n</i> dependent variables)	concentrations to one structure
		parameter of feedback loop.
Two-way	Bivariable (two independent	Maps a selected hormone
	and one dependent variables)	concentration to two structure
		parameters of feedback loop.
Tornado plot	Multivariable ( <i>n</i> independent	Maps a selected hormone
	and one dependent variables)	concentration to fixed changes
		of an arbitrary number of
		structure parameters.
Nullcline plot	Bivariable and bivariate	Maps the interaction of two
		hormone concentrations to
		visualise the evolution of an
		equilibrium point. Also referred
		to as interaction plot, null
		isocline plot or zero growth
		isocline plot

Table 1: Forms of sensitivity analysis provided by SimThyr

#### Preferences

Global application preferences allow for setting units of measurement and number formats for all your simulations.

Units of m	easureme	nt:					
TSH	m 岁	U/			~	r.	Example: 1 mU/I
TT4	μ 🞽 g	~	1	d	~	Ľ,	Example: 10 µg/dl
FT4	μ 🞽 g	~	7	d	~	r.	Example: 0.001 µg/dl
ттз	n 🞽 g	~	1	m	~	I.	Example: 1 ng/ml
FT3	n 🎽 g	~	1	m	~	I.	Example: 0.003 ng/ml
Formats:							
Numbers	###,###.	00##					123,456.789
Time	"d"D hh:r	n:ss					
							Cancel OK

The flexible modular menu structure enables you to compose virtually every imaginable measurement unit. Your entry is parsed and automatically converted to correct conversion factors.

Number formats are also very flexible. You may specify Free Pascal-style formatstrings to set the number of digits and type of notation:

**0**: Digit place holder. The digit of the corresponding place in the number will replace 0, otherwise a missing digit is filled with 0.

#: Digit place holder. A digit in the value will replace the corresponding #, a missing digit is replaced by a space.

.: Location of decimal separator

,: Location of thousands separator

E+, e+, E- and e-: Scientific notation. # signs after E+ denote the number of places.

; Separates positive, negative and zero numbers, respectively.

Examples:

###,###.00##:	1,234.50
###### <b>.</b> 0:	1234.5
#.###E+:	1.235E+5

Date formats may be specified with format strings, too:

D: day dd: day with leading zero h: hour n: minute s: second

Examples:

"d"D hh:nn:ss:	d1 03:31:40
hh:nn "h":	03:31 h

#### Supported File Formats

SimThyr supports export of simulation results in various formats including tabdelimited, CSV and DIF files. Both tab delimited and CSV are simple table formats, where columns (i.e. different simulated parameters) are separated by tabulator or comma signs and points in time are delimited by carriage return or linefeed symbols (depending on platform conventions). DIF (data interchange format) is a standard for a neutral representation of contents that facilitates the transfer of data among various applications. All of the mentioned formats are supported by a wide range of database, statistics and spreadsheet applications.

Plots may be saved in multiple varieties of image formats including bitmap (BMP), pixmap (XPM), portable pixmap (PPM), joint picture expert group (JPEG) format, tagged image file format (TIFF), portable network graphic (PNG) and, with some restrictions, scalable vector graphic (SVG).

Scenarios and preferences are saved as XML files.

# Appendix

#### References

- 1. Dietrich JW: Der Hypophysen-Schilddrüsen-Regelkreis. Entwicklung und klinische Anwendung eines nichtlinearen Modells, vol. 2. Berlin: Logos-Verlag; 2002.
- Fellinger K: Klinische und experimentelle Untersuchungen über das Verhalten und die Bedeutung des thyreotropen Hormons im Blute. Wien Arch inn Med 1936, 29:375-406.
- 3. Danziger L, Elmergreen GL: **The thyroid pituitary homeostatic mechanism**. *Bulletin of Mathematical Bology* 1956, **18**(1):1-13.
- 4. DiStefano JJ, 3rd, Stear EB: Neuroendocrine control of thyroid secretion in living systems: a feedback control system model. *Bull Math Biophys* 1968, **30**(1):3-26.
- 5. Norwich KH, Reiter R: Homeostatic control of thyroxin concentration expressed by a set of linear differential equations. *Bull Math Biophys* 1965, **27**(2):133-144.
- 6. Roston S: Mathematical representation of some endocrinological systems. Bulletin of Mathematical Bology 1959, **21**(3):271-282.
- 7. Degon M, Chipkin SR, Hollot CV, Zoeller RT, Chait Y: A computational model of the human thyroid. *Math Biosci* 2008, **212**(1):22-53.
- 8. Eisenberg M, Samuels M, DiStefano JJ, 3rd: Extensions, validation, and clinical applications of a feedback control system simulator of the hypothalamo-pituitary-thyroid axis. *Thyroid* 2008, **18**(10):1071-1085.
- 9. Leow MK: A mathematical model of pituitary--thyroid interaction to provide an insight into the nature of the thyrotropin--thyroid hormone relationship. *J Theor Biol* 2007, **248**(2):275-287.
- 10. Li G, Liu B, Liu Y: A dynamical model of the pulsatile secretion of the hypothalamo-pituitary-thyroid axis. *Biosystems* 1995, **35**(1):83-92.
- 11. Dietrich JW, Landgrafe-Mende G, Wiora E, Chatzitomaris A, Klein HH, Midgley JE, Hoermann R: Calculated Parameters of Thyroid Homeostasis: Emerging Tools for Differential Diagnosis and Clinical Research. Front Endocrinol (Lausanne) 2016, 7:57.
- 12. Dietrich JW, Landgrafe G, Fotiadou EH: **TSH and Thyrotropic Agonists: Key Actors** in **Thyroid Homeostasis**. *Journal of thyroid research* 2012, **2012**:351864.
- 13. Dietrich JW, Tesche A, Pickardt CR, Mitzdorf U: Thyrotropic Feedback Control: Evidence for an Additional Ultrashort Feedback Loop from Fractal Analysis. Cybernetics and Systems 2004, 35(4):315-331.
- 14. Hoermann R, Midgley JE, Larisch R, Dietrich JW: Is pituitary TSH an adequate measure of thyroid hormone-controlled homoeostasis during thyroxine treatment? *European journal of endocrinology / European Federation of Endocrine Societies* 2013, 168(2):271-280.
- 15. Midgley JE, Hoermann R, Larisch R, Dietrich JW: Physiological states and functional relation between thyrotropin and free thyroxine in thyroid health and disease: in vivo and in silico data suggest a hierarchical model. *Journal of clinical pathology* 2013, 66(4):335-342.
- 16. Hoermann R, Midgley JE, Larisch R, Dietrich JW: Integration of Peripheral and Glandular Regulation of Triiodothyronine Production by Thyrotropin in Untreated and Thyroxine-Treated Subjects. *Horm Metab Res* 2015, **47**(9):674-680.

# Contact

PD Dr. med. Johannes W. Dietrich<sup>1, 2, 3, 4</sup>

<sup>1</sup>Sektion Diabetologie, Endokrinologie und Stoffwechsel, St. Josef-Hospital, Ruhr University of Bochum, Gudrunstr. 56, D-44791 Bochum, NRW, Germany

<sup>2</sup>Diabeteszentrum Bochum/Hattingen, Klinik Blankenstein, Im Vogelsang 5–11, D-45527 Hattingen, NRW, Germany

<sup>3</sup>Zentrum für Seltene Endokrine Erkrankungen (ZSE), Centrum für Seltene Erkrankungen Ruhr (CeSER), Alexandrinenstr. 5, D-44791 Bochum, NRW, Germany

<sup>4</sup>Zentrum für Diabetestechnologie (ZDT), Katholisches Klinikum Bochum, Im Vogelsang 5–11, D-45527 Hattingen, NRW, Germany

# http://simthyr.sf.net