

PARAMETER ESTIMATION OF MICROBIAL MODELS USING HYBRID  
OPTIMIZATION METHODS

AFNIZANFAIZAL BIN ABDULLAH

UNIVERSITI TEKNOLOGI MALAYSIA

PARAMETER ESTIMATION OF MICROBIAL MODELS USING HYBRID  
OPTIMIZATION METHODS

AFNIZANFAIZAL BIN ABDULLAH

A thesis submitted in fulfilment of the  
requirements for the award of the degree of  
Doctor of Philosophy (Computer Science)

Faculty of Computing  
Universiti Teknologi Malaysia

JUNE 2013

To my beloved wife, children, family, and friends...

## ACKNOWLEDGEMENT

I would like to express my sincere thanks to Prof. Dr. Safa'ai Deris, my supervisor, for his encouragement, guidance and advices. He has corrected all my mistakes patiently and taught me everything that I need not only for the successfulness of my study but also for my life and career. His continual support and inspiration have led me to explore the area of Bioinformatics and made this research possible. My sincere appreciation also goes to Assoc. Prof. Dr. Siti Zaiton Mohd Hashim, Assoc. Prof. Dr. Mohd Saberi Mohamad, my colleagues in Artificial Intelligence and Bioinformatics Research Group (AIBIG) and all of my friends for their continuous help and support. Finally, I would like to thank my beloved wife, Aishah Yusoff, for her everlasting patience and motivation which help me through my difficult time. Thank you all.

## ABSTRACT

Development of biological models is essential as it represents and predicts complex processes within microbial cells. These models are formed by mathematical formulations that depend heavily on a set of parameters whose accuracy is often influenced by noisy and incomplete experimental data. This study is aimed to design and develop new optimization methods that can effectively estimate these parameters by iteratively fitting the model outputs to the experimental data. To achieve this goal, two new hybrid optimization methods based on the Firefly Algorithm (FA) method are proposed. Firstly, a method using evolutionary operations from Differential Evolution (DE) method was developed to improve the estimation accuracy of the parameters. Then, a second method using Chemical Reaction Optimization (CRO) method was proposed to surmount the convergence speed problem during parameter estimation. The effectiveness of the proposed methods was evaluated using synthetic transcriptional oscillator and extracellular protease production models. Computational experiments showed that these methods were able to estimate plausible parameters which produced model outputs that closely fitted in the experimental data. Statistical validation confirmed that these methods are competent at estimating the identifiable parameters. These findings are crucial to ensure that the estimated parameters can generate predictive and sensitive model outputs. In conclusion, this study has presented new hybrid optimization methods, capable of estimating the model parameters effectively whilst taking into account noisy and incomplete experimental data.

## ABSTRAK

Pembangunan model biologi adalah penting kerana ia mewakili dan meramalkan proses-proses kompleks di dalam sel-sel mikrob. Model-model ini dibentuk dengan ungkapan matematik yang sangat bergantung kepada satu set parameter yang ketepatannya sering dipengaruhi oleh data eksperimen yang hingar dan tidak lengkap. Kajian ini bertujuan untuk merekabentuk dan membangunkan kaedah-kaedah pengoptimuman yang mampu menganggarkan parameter-parameter ini dengan memadankan output model kepada data eksperimen secara berlelaran. Untuk mencapai tujuan ini, dua kaedah pengoptimuman hibrid berasaskan Algoritma Kunang-Kunang telah dicadangkan. Pertama, kaedah yang menggunakan operasi-operasi evolusi daripada kaedah Evolusi Perbezaan telah dibangunkan untuk memperbaiki ketepatan padanan parameter-parameter. Kemudian, kaedah kedua menggunakan kaedah Pengoptimuman Tindakbalas Kimia telah dicadangkan untuk mengatasi masalah kepantasan penumpuan semasa menganggarkan parameter. Keberkesanan kaedah-kaedah yang dicadangkan telah dinilai menggunakan model pengayun transkripsi sintetik dan model pengeluaran protease luar sel. Eksperimen-eksperimen komputeran menunjukkan kaedah-kaedah ini mampu menganggarkan parameter-parameter yang dapat menghasilkan model output yang hampir padan dengan data eksperimen. Pengesahsahihan statistik mengesahkan bahawa kaedah-kaedah ini adalah memuaskan dalam menganggarkan parameter-parameter yang dapat dikenalpasti. Penemuan-penemuan ini adalah penting untuk memastikan parameter-parameter yang dianggarkan dapat menjana model output yang boleh diramal dan peka. Kesimpulannya, kajian ini telah mempersembahkan kaedah-kaedah pengoptimuman hibrid yang baru, yang mampu menganggarkan parameter-parameter model secara efektif dengan mengambilkira data eksperimen yang hingar dan tidak lengkap.

## TABLE OF CONTENTS

CHAPTER	TITLE	PAGE
	<b>DECLARATION</b>	ii
	<b>DEDICATION</b>	iii
	<b>ACKNOWLEDGEMENTS</b>	iv
	<b>ABSTRACT</b>	v
	<b>ABSTRAK</b>	vi
	<b>TABLE OF CONTENTS</b>	vii
	<b>LIST OF TABLES</b>	xi
	<b>LIST OF FIGURES</b>	xii
	<b>LIST OF ABBREVIATION</b>	xiv
	<b>LIST OF APPENDICES</b>	xvi
<b>1</b>	<b>INTRODUCTION</b>	1
	1.1 Problem Background	1
	1.2 Problem Statement	4
	1.3 Research Goal and Objectives	6
	1.4 Research Scope and Significance	6
	1.5 Thesis Outline	7
	1.6 Summary	8
<b>2</b>	<b>LITERATURE REVIEW</b>	9
	2.1 Introduction	9
	2.2 Overview of the Bacterial Cell	11
	2.3 Bacterial Extracellular Protease Production	15
	2.4 Biological Modeling	16
	2.5 Parameter Estimation	18

2.6	Solving Parameter Estimation Using Optimization Methods	22
2.6.1	Local Optimization Methods	22
2.6.2	Global Optimization Methods	23
2.6.3	Hybrid Optimization Methods	26
2.7	Comparative Analysis on the Optimization Methods	29
2.8	Non-Identifiability Validation in Parameter Estimation	34
2.8.1	Structural Non-Identifiability	34
2.8.2	Practical Non-Identifiability	36
2.9	Research Trends and Directions	38
2.10	Summary	40
<b>3</b>	<b>RESEARCH METHODOLOGY</b>	<b>41</b>
3.1	Introduction	41
3.2	Research Framework	42
3.3	Model and Experimental Data	45
3.3.1	Synthetic Transcriptional Oscillators	47
3.3.2	Extracellular Protease Production	48
3.4	Parameter Estimation Problem Formulation	51
3.5	Non-Identifiability Analysis	53
3.6	Performance Measurements for Proposed Methods	54
3.7	Summary	55
<b>4</b>	<b>A COMPARATIVE ANALYSIS OF EXISTING OPTIMIZATION METHODS FOR PARAMETER ESTIMATION IN BIOLOGICAL MODELS</b>	<b>56</b>
4.1	Introduction	56
4.2	Existing Optimization Methods	57
4.2.1	Nelder-Mead	57
4.2.2	Levenberg-Marquardt	58



4.2.3	Particle Swarm Optimization (PSO)	59
4.2.4	Differential Evolution (DE)	59
4.2.5	Firefly Algorithm (FA)	61
4.2.6	Chemical Reaction Optimization (CRO)	62
4.2.7	Particle Swarm Evolutionary Optimization (PSEO)	65
4.2.8	Differential Evolutionary Bee Colony Optimization (DEBCO)	67
4.3	Experimental Results	69
4.3.1	Synthetic Transcriptional Oscillators	69
4.3.2	Extracellular Protease Production	79
4.4	Discussion	88
4.5	Summary	90
<b>5</b>	<b>A NEW HYBRID OPTIMIZATION METHOD FOR BIOLOGICAL MODEL PARAMETER ESTIMATION</b>	<b>92</b>
5.1	Introduction	92
5.2	An Evolutionary Firefly Algorithm	93
5.3	Experimental Results	97
5.3.1	Synthetic Transcriptional Oscillators	97
5.3.2	Extracellular Protease Production	105
5.4	Discussion	114
5.5	Summary	116
<b>6</b>	<b>AN IMPROVED HYBRID OPTIMIZATION METHOD FOR PARAMETER ESTIMATION</b>	<b>117</b>
6.1	Introduction	117
6.2	An Improved Swarm-Based Optimization Method	118
6.2.1	Solution Representation and Initialization	118
6.2.2	Solution Discrimination	119

6.2.3	Swarm Evolutionary Improvement	119
6.2.4	Weak Solution Randomization	120
6.3	Experimental Results	124
6.3.1	Synthetic Transcriptional Oscillators	124
6.3.2	Extracellular Protease Production	133
6.4	Discussion	143
6.5	Summary	145
<b>7</b>	<b>CONCLUSION AND FUTURE WORK</b>	<b>146</b>
7.1	Conclusion	146
7.2	Future Work	149
	<b>REFERENCES</b>	<b>151</b>
	Appendices A-D	164-206

## LIST OF TABLES

<b>TABLE NO.</b>	<b>TITLE</b>	<b>PAGE</b>
2.1	Comparative analysis of the optimization methods	32
3.1	Parameter values of synthetic transcriptional oscillators model	48
3.2	Parameter values of extracellular protease production model	50
4.1	Parameter tunings for the evaluated methods using trial-and-error approach	71
4.2	Average best fitness and standard deviation of the evaluated methods	72
4.3	Average computation times for the synthetic transcriptional oscillators model. Average time taken for 500 iterations in 100 independent runs	73
4.4	Variance points and intervals for non-identifiability analysis	76
4.5	Parameter tunings for the evaluated methods using trial-and-error approach	80
4.6	Average best fitness and standard deviation of the evaluated methods	81
4.7	Average computation times for the synthetic transcriptional oscillators model. Average time taken for 500 iterations in 100 independent runs	83
4.8	Variance points and intervals for non-identifiability analysis	84
5.1	Parameter tunings for the evaluated methods	98
5.2	Average best fitness and standard deviation of the	99

	evaluated methods	
5.3	Average computation times for the synthetic transcriptional oscillators model. Average time taken for 500 iterations in 100 independent runs	100
5.4	Variance points and intervals for non-identifiability analysis	102
5.5	Parameter tunings for evaluated methods	106
5.6	Average best fitness and standard deviation of the evaluated methods	107
5.7	Average computation times for the extracellular protease production model. Average time taken for 1000 iterations in 100 independent runs	109
5.8	Variance points and intervals for non-identifiability analysis	110
6.1	Parameter tunings for the evaluated methods	125
6.2	Average best fitness and standard deviation of the evaluated methods	126
6.3	Average computation times for the synthetic transcriptional oscillators model. Average time taken for 500 iterations in 100 independent runs	127
6.4	Variance points and intervals for non-identifiability analysis	130
6.5	Parameter tunings for the evaluated methods	134
6.6	Average best fitness and standard deviation of the evaluated methods	135
6.7	Average computation times for the extracellular protease production model. Average time taken for 1000 iterations in 100 independent runs	137
6.8	Variance points and intervals for non-identifiability analysis	139

## LIST OF FIGURES

FIGURE NO.	TITLE	PAGE
2.1	General structure of the bacterial cell	12
2.2	Gram-positive and Gram-negative bacteria	13
2.3	Bacterial extracellular protease production (Veening <i>et al.</i> , 2008)	15
2.4	The use of optimization method for solving parameter estimation problem	21
2.5	Contours by a pair of parameters, $p_1$ and $p_2$ .	35
3.1	Research framework	44
3.2	Simulation and experimental data preparation	46
3.3	Framework of solving parameter estimation problem using optimization methods	52
4.1	Fitness convergence of the evaluated methods	74
4.2	Data fit of model outputs with the experimental data	77
4.3	Fitness convergence of the evaluated methods	82
4.4	Data fit of model outputs with the experimental data	85
5.1	Pseudocode for NEFA method	95
5.2	NEFA method for parameter estimation	96
5.3	Fitness convergence of the evaluated methods	101
5.4	Data fit of model outputs with the experimental data	103
5.5	Fitness convergence of the evaluated methods	108
5.6	Data fit of model outputs with the experimental data	111
6.1	Pseudocode for S-CRO method	122
6.2	S-CRO method for parameter estimation	123
6.3	Fitness convergence of the evaluated methods	128

6.4	Data fit of model outputs with the experimental data	131
6.5	Fitness convergence of the evaluated methods	136
6.6	Data fit of model outputs with the experimental data	140

**LIST OF ABBREVIATION**

ABC	-	Artificial Bee Colony
ACO	-	Ant Colony Optimization
AIC	-	Aikake Information Criterion
AIS	-	Artificial Immune System
AMP	-	Adenosine Monophosphate
AS	-	Atorvastatin
ASL	-	Lipophilic Lactone
ASLpOH	-	Para-Hydroxy-Atorvastatin Lactone Acid
ATP	-	Adenosine Triphosphate
BA	-	Bee Algorithm
BFGS	-	Broyden–Fletcher–Goldfarb–Shanno
CPU	-	Central Processing Unit
CRO	-	Chemical Reaction Optimization
CSA	-	Clonal Selection Algorithm
DE	-	Differential Evolution
DEBCO	-	Differential Evolution Bee Colony Optimization
DNA	-	Deoxyribonucleic Acid
EA	-	Evolutionary Algorithm
ES	-	Evolution Strategy
EKF	-	Extended Kalman Filter
FA	-	Firefly Algorithm
FIM	-	Fisher’s Information Matrix
FGF	-	Fibroblast Growth Factor
GA	-	Genetic Algorithms
HIV	-	Human Immunodeficiency Virus
KEGG	-	Kyoto Encyclopedia Of Genes And Genomes
LPS	-	Lipopolysaccharide

mRNA	-	Messenger Ribonucleic Acid
NEFA	-	New Evolutionary Firefly Algorithm
NFL	-	No Free Lunch
NLP	-	Nonlinear Programming
ODEs	-	Ordinary Differential Equations
OS	-	Operating System
PCR	-	Polymerase Chain Reaction
PSM	-	Presomitic Mesoderm
PSEO	-	Particle Swarm Evolutionary Optimization
PSO	-	Particle Swarm Optimization
RA	-	Retinoic Acid
RNA	-	Ribonucleic Acid
SBFA	-	Selective Breeding Firefly Algorithm
SBML	-	Systems Biology Markup Language
S-CRO	-	Swarm-Based Chemical Reaction Optimization
SQP	-	Sequential Quadratic Programming
SRES	-	Stochastic Ranking Evolutionary Strategy
SS	-	Scatter Search



**LIST OF APPENDICES**

<b>APPENDIX</b>	<b>TITLE</b>	<b>PAGE</b>
A	List of Publications	164
B	Publication 1: An Evolutionary Firefly Algorithm for the Estimation of Nonlinear Biological Model Parameters	166
C	Publication 2: An Improved Swarm Optimization for Parameter Estimation and Biological Model Selection	183
D	Publication 3: A New Hybrid Bee Evolution Algorithm for Parameter Estimation in Biological Model	200

# CHAPTER 1

## INTRODUCTION

### 1.1 Problem Background

Computational systems biology is aimed to elucidate complex behaviours of biochemical reactions within cells through computational approaches (Sun *et al.*, 2012; Lages *et al.*, 2012). This field of research is important to acquire better understanding of how these reactions work as a system (Karr *et al.*, 2012; Isalan, 2012). In general, these reactions can be represented by using computational models. These models are constructed based on mathematical formulations such as ordinary differential equations (ODEs) to quantify the changes of specific biochemical concentrations over a sample of observation time. The development of these models commonly involves two major stages, namely network structure identification and parameter estimation (Tashkova *et al.*, 2011; Chou and Voit, 2009). On the network structure identification stage, the structures of the models are formed by modelling experts. This often requires prior knowledge of the reaction networks. On the other hand, the parameter estimation stage is performed to determine parameter values in the constructed models. These parameters are usually approximated based on the available experimental data obtained from the high-throughput experiments. This is a challenging task because the biochemical reactions in the systems are highly nonlinear and the experimental data are frequently noisy and incomplete (Tashkova *et al.*, 2011; Sun *et al.*, 2012; Lages *et al.*, 2012).

The purpose of the parameter estimation stage is to equip the models with a plausible set of parameters. These parameters are used to produce model outputs

that are consistent with the experimental data. Thus, these parameters are crucial to signify physical properties of the systems, such as kinetic constants and reaction rates. However, these parameters are difficult to be extracted from the high-throughput experiments. In most cases, these parameters are estimated using nonlinear programming (NLP) methods (Balsa-Canto *et al.*, 2012; Sun *et al.*, 2012).

The parameter estimation task is considered as an optimization problem, in which optimal model parameters are repeatedly determined by the difference between the model outputs and the corresponding experimental data is minimized. Generally, there are two major approaches for this task: gradient and stochastic searching strategies (Chou and Voit, 2009). The gradient searching strategy usually utilizes local search algorithms to find the parameters based on the initial guesses of the state measurements. However, due to the complexity of the biological systems, it is difficult to determine these initial guesses as the values are often unknown. Moreover, the nonlinearity of the models may lead the searching to least substantial parameters (Balsa-Canto *et al.*, 2011). To overcome this limitation, stochastic searching strategy is applied. This strategy employs global optimization methods that initiate the searching processes with a set of randomly selected model parameters. Nevertheless, a major drawback of this strategy is the high computational cost. Furthermore, the strategy sometimes has difficulty to converge the searching to the local optimum solutions (Sun *et al.*, 2012).

To surmount these bottlenecks, many researches have considered the use of hybrid optimization methods. In these methods, the convergence performance of the stochastic searching strategy is improved by incorporating other optimization algorithms (Rodriguez-Fernandez *et al.*, 2008; Ashyraliyev *et al.*, 2009). In recent years, the hybrid optimization methods have shown potential achievements in estimating parameters of the biological models (Lages *et al.*, 2012; Sun *et al.*, 2012; Tashkova *et al.*, 2011). Moreover, Evolutionary Algorithms (EAs) such as Differential Evolution (DE) (Storn and Price, 1997), Clonal Selection Algorithm (CSA) (De Castro and Von Zuben, 2002), and Chemical Reaction Optimization Algorithm (CRO) (Lam and Li, 2010) have presented promising capabilities in handling measurement noise and incompleteness of the experimental data (Sun *et al.*, 2012; Abdullah *et al.*, 2011). This is due to the fact that these algorithms employ random recombination searching approaches that utilize neighbouring

vectors within the population of solutions. Therefore, incorporating EAs may facilitate improvements of stochastic searching strategy, especially in terms of robustness over noisy and incomplete experimental data during the estimation process (Tashkova *et al.*, 2011).

Besides handling measurement noise and incompleteness of the experimental data, another challenge may also arise in the parameter estimation task, namely parameter non-identifiability. The parameter non-identifiability occurred when the estimated parameters fail to produce distinctive model outputs. There are two types of parameter non-identifiability, namely structural and practical non-identifiability (Balsa-Canto and Banga, 2010). The structural non-identifiability generally occurred due to limitations in the model structure, in which can be solved by modifying the models until the model outputs are consistent with the experimental data (Bandara *et al.*, 2009). On the other hand, practical non-identifiability problem is much complicated as it often occurs due to the quality and amount of the experimental data (Chis *et al.*, 2011).

Providing sufficient experimental data and constraints may be useful to solve this problem. This may allow the parameter estimation task to find unique parameters based on the experimental data (Balsa-Canto and Banga, 2010). Finding identifiable parameters is useful for model selection. Model selection is generally defined as the capability to choose plausible models based on the given experimental data. Therefore, unique parameters that are estimated by the optimization methods may produce distinctive model outputs, in which will facilitate to differentiation of models (Miao *et al.*, 2009; Lillacci and Khammash, 2010). The model selection permits further analyses of the model structures according to the available experimental data, especially for finding new pathways to improve certain biochemical productions.

## 1.2 Problem Statement

Computational systems biology has become an increasingly important research area in the recent years (Sun *et al.*, 2012; Lages *et al.*, 2012). This field of research is aimed to gain better understanding of how complex biological process response as a system within living cells. This is often facilitated using computational models (Sun *et al.*, 2012; Tashkova *et al.*, 2011; Chou and Voit, 2009). These models commonly contain a set of parameters that represent the physiological properties of the systems. Generally, obtaining these parameters is a challenging task. In recent years, many optimization methods have been proposed to estimate these parameters by fitting the model outputs with the corresponding experimental data. This is usually performed by minimizing the difference between these two data. However, the available experimental data are usually incomplete and has measurement noise. Thus, designing and developing robust optimization methods are crucial to ensure the accuracy of the estimation. Moreover, the estimated parameters are sometime non-identifiable, which thwart the possibility of finding plausible parameters that may produce informative model outputs. As the reliable parameters are difficult to be attained, this may lead to further difficulty in selecting feasible models based on the given experimental data.

Currently, there is an increasing number of nonlinear optimization methods proposed to estimate the parameters in the biological models (Sun *et al.*, 2012; Balsa-Canto *et al.*, 2012; Tashkova *et al.*, 2011). The aim of these methods is to find the optimal parameter set which may produce the model outputs that closely fit the corresponding experimental data. Conventionally, derivative-based optimization methods are utilized, including maximum likelihood (Lloyd-Smith, 2007) and gradient descent (Ashyraliyev *et al.*, 2008) methods. More recently, a local optimization method, namely Extended Kalman Filter (EFK) (Costa, 1994) method, is employed (Sun *et al.*, 2008). Lillacci and Khammash (2010) introduced an improved EFK method that incorporates the continuous model outputs and the experimental measurements to estimate the parameters using state space searching approach. Additionally, Zheng and co-workers (2012) proposed inequality constraints to improve the estimation by the EFK method. However, both improved methods commonly require the use of model refinement phases to avoid the searching processes from being trapped into suboptimal solutions. Furthermore, these methods need to consider the limitations of the EFK method that heavily rely

on a good set of initial values for both states and parameters in the models (Sun *et al.*, 2008).

In contrast, several previous works have presented prospective achievements by using metaheuristics methods (Balsa-Canto *et al.*, 2012). Methods such as Particle Swarm Optimization (PSO) (Kennedy and Eberhart, 1995) and Genetic Algorithm (GA) (Goldberg, 1988) were also used to estimate the parameters in biological systems, which showed promising results (Besozzi *et al.*, 2008; Tutkun, 2009). More recently, evolutionary-based metaheuristics methods have received the remarkable attentions (Sun *et al.*, 2012; Tashkova *et al.*, 2011; Buhry *et al.*, 2011). Generally, these methods utilize evolutionary operations such as crossover, mutation, and selection operations to exploit the information of the solutions in the population. Tashkova and co-workers (2011) suggested that the use of DE method is more practical compared to the existing meta-heuristic methods. However, it was also presented that the method may use a substantial amount of computational cost to obtain the best solution (Sun *et al.*, 2012; Abdullah *et al.*, 2011). Moreover, there is no guarantee that these methods will converge to the global optimum solutions (Balsa-Canto *et al.*, 2012). These generally lead to the use of hybrid optimization methods that combine several searching techniques of different metaheuristics methods to overcome these limitations.

Therefore, the problem of this research can be formulated as follows: given the noisy and incomplete experimental data, it is a challenging task to design and develop an effective hybrid optimization method that robustly estimate the model parameters within an acceptable amount of computational time. The proposed method also needs to consider the non-identifiability of the estimated parameters.

### 1.3 Research Goal and Objectives

The goal of this research is to propose a new hybrid optimization method for estimating model parameters based on noisy and incomplete experimental data. In order to achieve this goal, the following objectives are required to be met:

1. To design and develop a new hybrid optimization method that can handle noisy and incomplete experimental data during the parameter estimation;
2. To evaluate the effectiveness of the proposed method in dealing with practically non-identifiable parameters;

### 1.4 Research Scopes and Significance

In this research, the metabolic systems of well-studied bacteria are used to evaluate the effectiveness of the proposed optimization methods. The systems, which are formed by a series of biochemical reactions, are used to observe the concentration changes of certain biochemical compounds in specific biological processes. The models used for the parameter estimation problem and non-identifiability analysis is obtained from Biomodels database (Le Novere *et al.*, 2006). The models are in the form of Systems Biology Markup Language (SBML) file format. The file contains the information of involved metabolites, reaction rates, parameters and the initial concentration volume used in the high-throughput experiments. The model is simulated using general purpose modelling software, COPASI (Hoops *et al.*, 2006). The experimental data for this model is generated *in silico*. This is performed by adding the Gaussian noise into the model outputs to simulate the measurement noise (Lillacci and Khammash, 2010).

The significance of the research is addressed as follows. Firstly, the design and development of new hybrid optimization methods is valuable in term of the computational contribution. The methods utilize the advantages of the evolutionary operations employed by DE and CRO methods to enhance the searching capability of the Firefly Algorithm (FA) method (Yang, 2009) and reduce computational time

significantly. In addition, the methods are capable to handle noisy and incomplete experimental data during the parameter estimation process. Secondly, the outcomes of this research can benefit the systems biology community. This is due to the contribution of the new approach to parameter estimation and non-identifiability analysis. As the optimization methods are robust to measurement noise and incompleteness, this provides effective tools to implement the methods for diverse parameter estimation problems of other biological models.

## 1.5 Thesis Outline

The organization of the thesis is outlined as follows:

- Chapter 1: This chapter provides the introduction of the research, which encompasses research background, problem statement, goal, objectives, scope and significance of the study.
- Chapter 2: This chapter provides the literature review of the research. The chapter starts with the overview of the bacterial cell. Then, the biological model development is described. This leads to the use of optimization methods for parameter estimation and discussion of related issues on the problem.
- Chapter 3: This chapter provides the research methodology. This chapter presents the research operational framework, description of the data used and an overview of the evaluation measurement.
- Chapter 4: This chapter presents an empirical analysis of the existing optimization methods, ranging from three categories: local, global, and hybrid optimization methods. The methods used in this analysis are Nelder-Mead, Levenberg-Marquardt, PSO, DE, FA, CRO, and two recently proposed hybrid optimization methods, namely Particle Swarm Evolutionary Optimization (PSEO) (Abdullah *et al.*, 2013a) and Differential Evolutionary Bee Colony Optimization (DEBCO) (Abdullah *et al.*, 2013b) methods. The effectiveness of this method is evaluated using two biological models: synthetic transcriptional oscillators (Kim and Winfree, 2011), and an extracellular protease production (Veening *et al.*, 2008) models.



- Chapter 5: In this chapter, a new hybrid optimization method based on FA and CRO methods is proposed. The method is validated for parameter estimation accuracy and its capability on handling non-identifiable parameters.
- Chapter 6: In this chapter, an improved hybrid optimization method based on FA and CRO methods is proposed. This method is aimed to overcome limitations of the method proposed in the previous chapter.
- Chapter 7: This chapter discusses the contribution of the works and future plans to fulfil the research objectives.

## **1.6 Summary**

In this chapter, the introduction of the research is presented. Firstly, the background of the research is discussed. This includes the designing of models for synthetic biology, current parameter estimation methods and challenges of parameter estimation of biological models. Then, the problem statement of the research is addressed. Next, the research goal and objectives are described. Later, the research scopes and the significance of the study are discussed. In the next chapter, the literature review of this research is presented.

## REFERENCES

- Abdullah A., Safaai D. and Anwar S. Hybrid Evolutionary Clonal Selection for Parameter Estimation of Biological Model. *International Journal of Computer Applications in Engineering Sciences*, 2011, 1(3): 313-319.
- Abdullah A., Deris S., Mohamad M.S. and Hashim S.Z.M. A New Particle Swarm Evolutionary Optimization for Parameter Estimation of Biological Models. *International Journal of Computer Information Systems and Industrial Management Applications*, 2013a, 5(1): 571-580.
- Abdullah A., Deris S. and Mohamad M.S. A New Hybrid Bee Evolution Algorithm for Parameter Estimation in Biological Model. *ICIC Express Letter Part B: Application*, 2013b, 4(1): 1-6.
- Apgar J.F., Toettcher J.E., Endy D., White F.M. and Tidor B. Stimulus design for model selection and validation in cell signaling. *PLoS Computational Biology*, 2008, 4: e30.
- Arias-Montano A., Coello Coello C. and Mezura-Montes E. Evolutionary Algorithms Applied to Multi-Objective Aerodynamic Shape Optimization. *Computational Optimization, Methods and Algorithms*, 2011: 211-240.
- Ashyraliyev M., Fomekong-Nanfack Y., Kaandorp J.A. and Blom J.G. Systems Biology: Parameter Estimation for Biochemical Models. *FEBS Journal*, 2009, 276(4): 886-902.
- Ashyraliyev M., Jaeger J. and Blom J.G. Parameter Estimation and Determinability Analysis Applied to *Drosophila* Gap Gene Circuits. *BMC Systems Biology*, 2008, 2(1): 83-102.
- Assareh E., Behrang M.A., Ghalambaz M., Noghrehabadi A.R. and Ghanbarzadeh A. A New Approach to Solve Blasius Equation using Parameter Identification of Nonlinear Functions based on the Bees Algorithm (BA). *World Academy of Science, Engineering and Technology*, 2011, 73: 1119-1121.

- Balsa-Canto E. and Banga J.R. Computational procedures of model identification, in Choi S. (eds.) *Systems Biology for Signaling Networks*: Springer. 111-137; 2010.
- Balsa-Canto E., Banga J.R., Egea J.A., Fernandez-Villaverde A. and Hijas-Liste G.M. Global Optimization in Systems Biology: Stochastic Methods and their Applications. *Advances in Systems Biology*, 2012, 736(4): 409-424.
- Balsa-Canto E., Banga J.R. and Garcia M.R. Dynamic Model Building Using Optimal Identification Strategies, with Applications in Bioprocess Engineering. *Process Systems Engineering*, 2011: 441-467.
- Balsa-Canto E., Alonso A.A. and Banga J.R. Computational Procedures for Optimal Experimental Design in Biological Systems. *IET Systems Biology*, 2008a, 2(4): 163-172.
- Balsa-Canto E., Peifer M., Banga J., Timmer J. and Fleck C. Hybrid Optimization Method with General Switching Strategy for Parameter Estimation. *BMC Systems Biology*, 2008b, 2(1): 26-35.
- Bandara S., Schlöder J., Eils R., Bock H. and Meyer T. Optimal Experimental Design for Parameter Estimation of a Cell Signaling Model. *PLoS Computational Biology*, 2009, 5(11): 1-12.
- Barr R. S., Golden B. L., Kelly J., Steward W. and Resende M. Guidelines for Designing and Reporting on Computational Experiments with Heuristic Methods. *Proceedings of International Conference on Metaheuristics for Optimization*, 2001: 1-17.
- Basu B. and Mahanti G.K. FireFly and Artificial Bees Colony Algorithm for Synthesis of Scanned and Broadside Linear Array Antenna. *Progress in Electromagnetics Research*, 2011, 32: 169-190.
- Batchelor E., Loewer A., Lahav G. The ups and downs of p53: understanding protein dynamics in single cells. *Nature Review Cancer*, 2009, 9: 371-377.
- Benson D.A., Karsch-Mizrachi I., Lipman D.J., Ostell J. and Wheeler D.L. GenBank. *Nucleic Acids Research*, 2008, 36(suppl. 1): D25.
- Besozzi D., Cazzaniga P., Mauri G., Pescini D. and Vanneschi L. A comparison of Genetic Algorithms and Particle Swarm Optimization for parameter estimation in stochastic biochemical systems. *Evolutionary Computation, Machine Learning and Data Mining in Bioinformatics*, 2009: 116-127.

- Bonilla J., Diehl M., Logist F., De Moor B. and Van Impe J. An Automatic Initialization Procedure in Parameter Estimation Problems with Parameter-Affine Dynamic Models. *Computers and Chemical Engineering*, 2010, 34(6): 953-964.
- Borchers D.L. and Efford M.G. Spatially Explicit Maximum Likelihood Methods for Capture-Recapture Studies, *Biometrics*, 2008, 64(2): 377-385.
- Bronte V. and Zanovello P. Regulation of immune responses by L-arginine metabolism. *Nature Review Immunology*, 2005, 5: 641–654.
- Bucher J., Riedmaier S., Schnabel A., Marcus K., Vacun G., Weiss T.S., Thasler W.E., Nüssler A.K., Zanger U.M. and Reuss M. A Systems Biology Approach to Dynamic Modelling and Inter-Subject Variability of Statin Pharmacokinetics in Human Hepatocytes. *BMC Systems Biology*, 2011, 5: 66-85.
- Buhry L., Grassia F., Giremus A., Grivel E., Renaud S. and Saighi S. Automated Parameter Estimation of the Hodgkin-Huxley Model Using the Differential Evolution Algorithm: Application to Neuromimetic Analog Integrated Circuits. *Neural Computation*, 2011: 1-27.
- Bujara M., Schümperli M., Billerbeck S., Heinemann M., Panke S. Exploiting Cell Free Systems: Implementation and De-Bugging of a System Of Biotransformations. *Biotechnology and Bioengineering*, 2010, 106: 376–389.
- Burgess C., O’Connell-Motherway M., Sybesma W., Hugenholtz J. and van Sinderen D. Riboflavin Production in *Lactococcus lactis*: Potential for *in-situ* Production of Vitamin-enriched Foods. *Applied and Environmental Microbiology*, 2004, 70(10): 5769-5777.
- Capistrán M.A., Moreles M.A. and Lara B. Parameter Estimation of Some Epidemic Models: The Case of Recurrent Epidemics Caused by Respiratory Syncytial Virus. *Bulletin of Mathematical Biology*, 2009, 71(8): 1890-1901.
- Casey F.P., Baird D., Feng Q., Gutenkunst R.N., Waterfall J.J., *et al.* Optimal Experimental Design in an Epidermal Growth Factor Receptor Signalling and Down-Regulation Model. *IET Systems Biology*, 2007, 1: 190–202.
- Chen Y.T. and Wang F.S. Determination of Kinetic Parameters for Enzymatic Cellulose Hydrolysis using Hybrid Differential Evolution. *International Journal of Chemical Reactor Engineering*, 2012, 9(1): 1-16.
- Chen D.B. and Zhao C.X. Particle Swarm Optimization with Adaptive Population Size and its Application. *Applied Soft Computing*, 2009, 9(1): 39-48.

- Chin S.V. and Chappell M.J. Structural Identifiability and indistinguishability Analyses of the Minimal Model and a Euglycemic Hyperinsulinemic Clamp model for glucose-insulin dynamics. *Computer Methods and Programs in Biomedicine*, 2010, 104(2): 120-134.
- Chou I. and Voit E.O. Recent Developments in Parameter Estimation and Structure Identification of Biochemical and Genomic Systems. *Mathematical Bioscience*, 2009, 219(2): 57-83.
- Chis O.T., Banga J.R. and Balsa-Canto E. Structural Identifiability of Systems Biology Models: A Critical Comparison of Methods. *PLoS One*, 2011, 6(11): e27755.
- Costa P.J. Adaptive Model Architecture and Extended Kalman-Bucy filters. *IEEE Transaction on Aerospace Electron System*, 1994, 30: 525-533.
- Das S., Abraham A. and Konar A. Particle Swarm Optimization and Differential Evolution Algorithms: Technical Analysis, Applications and Hybridization Perspective. *Studies in Computational Intelligence*, 2008, 116: 1-38.
- De Castro L.N. and Von Zuben F.J. Learning and Optimization using the Clonal Selection Principle. *IEEE Transactions on Evolutionary Computation*, 2002, 6(3): 239-251.
- Dennis J. E., and Woods D. J. Optimization on Microcomputers: The Nelder-Mead Simplex Algorithm. *New Computing Environments: Microcomputers in Large-Scale Computing*, 1987, 116-122.
- Dräger A., Kronfeld M., Ziller M., Supper J., Planatscher H., Magnus J., Oldiges M., Kohlbacher O. and Zell A. Modeling Metabolic Networks in *C. Glutamicum*: A Comparison of Rate Laws in Combination with Various Parameter Optimization Strategies. *BMC Systems Biology*, 2009, 3(1): 5-29.
- Dorigo M. and Di Caro G. Ant Colony Optimization: A New Meta-Heuristic. *Proceedings of the Congress on Evolutionary Computation*, 1999, 2, DOI: 10.1109/CEC.1999.782657
- Dubrulle J., McGrew M.J. and Pourquié O. FGF Signaling Controls Somite Boundary Position and Regulates Segmentation Clock Control of Spatiotemporal Hox Gene Activation. *Cell*, 2001, 106: 219-232.
- Durot M., Bourguignon P.Y. and Schachter V. Genome-Scale Models of Bacterial Metabolism: Reconstruction and Applications. *FEMS Microbiology Review*, 2008, 33: 164-190.

- Egea J.A., Marti R. and Banga J.R. An Evolutionary Method for Complex-Process Optimization. *Computers and Operations Research*, 2010, 37(2): 315-324.
- Evans N.D., Chapman M.J., Chappell M.J. and Godfrey K.R. Identifiability of uncontrolled nonlinear rational systems. *Automatica*, 2002, 38(10): 1799-1805.
- Fomekong-Nanfack Y., Kaandorp J.A. and Bloom J. (2007), Efficient parameter estimation for spatio-temporal models of pattern formation: case study of *Drosophila melanogaster*, *Bioinformatics*, 23, 3356-3363.
- Freeman M. and Gurdon J.B. Regulatory Principles of Developmental Signaling. *Annual Review on Cell Developmental Biology*, 2002, 18: 515-539.
- Gao X.Z., Wang X. and Ovaska S.J. Fusion of Clonal Selection Algorithm and Differential Evolution Method in Training Cascade-Correlation Neural Network, *Neurocomputing*, 2009, 72(10-12): 2483-2490.
- Glover F, Laguna M, Marti R. Fundamentals of Scatter Search and Path Relinking. *Control and Cybernetics*, 2000, 39(3): 653-684
- Goldberg D.E. Genetic Algorithms in Search, Optimization, and Machine Learning: Addison-Wesley Professional. 1989.
- Goldbeter A., Gonze D. and Pourquie O. Sharp Developmental Thresholds Defined Through Bistability by Antagonistic Gradients of Retinoic Acid and FGF Signaling. *Developmental Dynamics*, 2007, 236(6): 1495-1508.
- Goroehowski T.E., Matyjaszkiewicz A., Todd T., Oak N., Kowalska K., Reid S., Tsaneva-Atanasova K.T., Savery N.J., Grierson C.S. and di Bernardo M. BSim: An Agent-Based Tool for Modeling Bacterial Populations in Systems and Synthetic Biology. *PLoS One*, 2012, 7(8): e42790.
- Guo T. and Kang L.S. A New Evolutionary Algorithm for Function Optimization. *Wuhan University Journal of Natural Sciences*, 1999, 4: 409-414.
- Gutenkunst R.N., Waterfall J.J., Casey F.P., Brown K.S., Myers C.R. and Sethna J.P. Universally Sloppy Parameter Sensitivities in Systems Biology Models. *PLoS Computational Biology*, 2007, 3(10): e189.
- Harris S.L., Levine A.J. The p53 Pathway: Positive and Negative Feedback Loops. *Oncogenetics*, 2005, 24(17): 2899-2908.
- Hattersley J.G., Pérez-Velázquez J., Chappell M.J., Bearup D., Roper D., Dowson C., Bugg T. and Evans N.D. Indistinguishability and Identifiability of Kinetic Models for the Murc Reaction in *Peptidoglycan* Biosynthesis. *Computer Methods and Programs in Biomedicine*, 2011, 22(1): 109-116.

- Ho W.H. and Chan A.L.F. Hybrid Taguchi-Differential Evolution Algorithm for Parameter Estimation of Differential Equation Models with Application to HIV Dynamics. *Mathematical Problems in Engineering*, 2011, DOI:10.1155/2011/514756.
- Hoops S., Sahle S., Gauges R., Lee C., Pahle J., Simus N., Singhal M., Xu L., Mendes P. and Kummer U. COPASI - A Complex Pathway Simulator. *Bioinformatics*, 2006, 22(24): 3067-3074.
- Hughenoltz J., Sybesma W., Nierop Groot M., Wisselink W., Ladero V., Burgess K., van Sinderen D., Piard J.C., Eggink G., Smid E.J., Savoy G., Sesma F., Jansen T., Hols P. and Kleerebezem M. Metabolic Engineering of Lactic Acid Bacteria for the Production of Nutraceuticals. *Antonie Van Leeuwenhoek*, 2002, 82(1): 217-235
- Hunziker A., Jensen M.H. and Sandeep K. Stress-Specific Response of the p53-Mdm2 Feedback Loop. *BMC Systems Biology*, 2010, 4: 94-102.
- Iacca G., Neri F. and Mininno E. Noise analysis compact differential evolution. *International Journal of Systems Science*, 2011, DOI:10.1080/00207721.2011.598964.
- Isalan M. A cell in a computer. *Nature*, 2012, 488: 40-41.
- Ishige T., Honda K. and Shimizu S. Whole Organism Biocatalysis. *Current Opinion in Chemical Biology*, 2005, 9(2): 174-180.
- Jin Y. and Branke J. Evolutionary Optimization in Uncertain Environments-A Survey, *IEEE Transactions on Evolutionary Computation*, 2005, 9(3): 303-317.
- Jin Y., Olhofer M. and Sendhoff B. A Framework for Evolutionary Optimization with Approximate Fitness Functions. *IEEE Transactions on Evolutionary Computation*, 2002, 6(5): 481-494.
- Jiménez-Hornero J.E., Santos-Dueñas I.M. and García-García I. Optimization of Biotechnological Processes. The Acetic Acid Fermentation. Part II: Practical Identifiability Analysis and Parameter Estimation. *Biochemical Engineering Journal*, 2009, 45(1): 7-21.
- Kanehisa M. and Goto S. KEGG: Kyoto Encyclopedia of Genes and Genomes. *Nucleic Acids Research*, 2000, 28(1): 27.
- Karr J.R., Sanghvi J.C., Macklin D.N., Gutschow M.V., Jacobs J.M., Bolival B., Assad-Garcia N., Glass J.I. and Covert M.W. A Whole-Cell Computational Model Predicts Phenotype from Genotype, *Cell*, 2012, 150(2): 389-401.

- Karaboga D. and Basturk B. A Powerful and Efficient Algorithm for Numerical Function Optimization: Artificial Bee Colony (ABC) Algorithm, *Journal of Global Optimization*, 2007, 39(3): 459-471.
- Katare S., Bhan A., Caruthers J.M., Delgass W.N., and Venkatasubramanian V. A Hybrid Genetic Algorithm for Efficient Parameter Estimation of Large Kinetic Models, *Computational Chemical Engineering*, 2004, 28(12): 2569-2581.
- Kennedy J. and Eberhart R.C. Particle Swarm Optimization, *Proceeding of IEEE International Conference on Neural Network*, 1995, 4: 1942-1948.
- Kephart J.O. A Biologically Inspired Immune System for Computers, *Artificial Life IV: Proceedings of the Fourth International Workshop on the Synthesis and Simulation of Living Systems*, 1994: 130-139.
- Kim K.A., Spencer S., Albeck J., Burke J., Sorger P., Gaudet S. and Kim D.H. Systematic Calibration of a Cell Signaling Network Model. *BMC Bioinformatics*, 2010, 11(1): 202-216.
- Kim J. and Winfree E. Synthetic *In Vitro* Transcriptional Oscillators. *Molecular and Systems Biology*. 2011, 7(1): 1-15.
- Kitano H. Computational Systems Biology. *Nature*, 2002, 420(12): 206-210.
- Kramer J.A., Sagartz J.E. and Morris D.L. The Application of Discovery Toxicology and Pathology Towards the Design of Safer Pharmaceutical Lead Candidates. *National Review on Drug Discover*, 2007, 6: 636-649.
- Lages N.F., Cordeiro C., Sousa Silva M., Ponces Freire A., Ferreira A.E.N. Optimization of Time-Course Experiments for Kinetic Model Discrimination. *PLoS One*, 2012, 7(3): e32749.
- Lam A.Y.S. and Li V.O.K. Chemical-Reaction-Inspired Metaheuristic for Optimization. *IEEE Transaction on Evolutionary Computation*, 2010, 14(3): 381-399.
- Le Novere N., Bornstein B., Broicher A., Courtot M., Donizelli M. *et al.* BioModels Database: a free, centralized database of curated, published, quantitative kinetic models of biochemical and cellular systems. *Nucleic Acids Research*, 2006, 34(suppl. 1): D689.
- Levine A.J., Hu W., Feng Z. The p53 Pathway: What Questions Remain to be Explored? *Cell Death Differentiation*, 2006, 13(6): 1027-1036.



- Liang H., Miao H. and Wu H. Estimation of Constant and Time-Varying Dynamic Parameters of HIV Infection in a Nonlinear Differential Equation Model. *The Annals of Applied Statistics*, 2010, 4(1): 460-483.
- Li L., Yang Y., Peng H. and Wang X. Parameters Identification of Chaotic Systems via Chaotic Ant Swarm. *Chaos, Solitons and Fractals*, 2006, 28(5): 1204-1211.
- Lillacci G. and Khammash M. Parameter estimation and model selection in computational biology. *PLoS Computational Biology*, 2010, 6(3): e1000696.
- Lillacci G. and Khammash M. A Distribution-Matching Method for Parameter Estimation and Model Selection in Computational Biology. *International Journal of Robust and Nonlinear Control*, 2012, 22: 1065-1081.
- Liu B., Zhang J., Tan P.Y., Hsu D., Blom A.M., Leong B., Sethi S., Ho, B., Ding J.L. and Thiagarajan P.S. A Computational and Experimental Study of the Regulatory Mechanisms of the Complement System. *PLoS Computational Biology*, 2011, 7(1): e1001059.
- Liu F. and Burrage K. Novel Techniques in Parameter Estimation for Fractional Dynamical Models Arising from Biological Systems. *Computers and Mathematics with Applications*, 2011, 62(3): 822-833.
- Lloyd-Smith, J.O. Maximum Likelihood Estimation of the Negative Binomial Dispersion Parameter for Highly Overdispersed Data, with Applications to Infectious Diseases. *PLoS One*, 2007, 2(2): e180.
- Lukasik S. and Zak S. Firefly Algorithm for Continuous Constrained Optimization Tasks. *Computational Collective Intelligence, Semantic Web, Social Networks and Multiagent Systems*, 2009: 97-106.
- Mente C., Prade I., Bruschi L., Breier G. and Deutsch A. Parameter Estimation with a Novel Gradient-Based Optimization Method for Biological Lattice-Gas Cellular Automaton Models. *Journal of Mathematical Biology*, 2010, 63(1): 173-200.
- Miao H., Dykes C., Demeter L.M. and Wu H. Differential Equation Modeling of HIV Viral Fitness Experiments: Model Identification, Model Selection, and Multimodel Inference. *Biometrics*, 2009, 65(1): 292-300.
- Miro A., Pozo C., Guillen-Gosalbez G., Egea J.A. and Jiménez L. Deterministic Global Optimization Algorithm based on Outer Approximation for the Parameter Estimation of Nonlinear Dynamic Biological Systems. *BMC Bioinformatics*, 2012, 13: 90.

- Modares H., Alfi A. and Naghibi Sistani M.B. Parameter Estimation of Bilinear Systems based on An Adaptive Particle Swarm Optimization. *Engineering Applications of Artificial Intelligence*, 2010, 23(7): 1105-1111.
- Moles C.G., Mendes P. and Banga J.R. Parameter Estimation in Biochemical Pathways: A Comparison of Global Optimization Methods, *Genome Research*, 2003, 13: 2467-2474
- Montanez R., Rodriguez-Caso C., Sanchez-Jimenez F., Medina M.A. *In Silico* Analysis of Arginine Catabolism as a Source of Nitric Oxide or Polyamines in Endothelial Cells. *Amino Acids*, 2008, 34(2): 223-229
- Moré J. J. The Levenberg-Marquardt algorithm: implementation and theory. In Numerical analysis. Springer Berlin Heidelberg. 105-116: 1978.
- Nelder J.A. and Mead R. A Simplex Method for Function Minimization. *The Computer Journal*, 1965, 7(4): 308-313.
- Noman N. And Iba H. Accelerating Differential Evolution using an Adaptive Local Search. *IEEE Transaction on Evolutionary Computation*, 2008, 12(1): 107-125.
- Papp B., Teusink B. and Notebaart A. A Critical View of Metabolic Network Adaptations. *HFSP Journal*, 2009, 3(1): 24-35.
- Pham D.T., Ghanbarzadeh A., Koc E., Otri S., Rahim S. and Zaidi M. Bees Algorithm, *Technical Note*, Manufacturing Engineering Centre, Cardiff University: 2005.
- Poovathingal S.K. and Gunawan R. Global Parameter Estimation Methods for Stochastic Biochemical Systems. *BMC Bioinformatics*, 2010, 11: 414-426.
- Proctor C.J. and Gray D.A. Explaining Oscillations and Variability in the p53-Mdm2 System, *BMC Systems Biology*, 2008, 2(1): 75-95.
- Prügel-Bennett A. Benefits of a Population: Five Mechanisms that Advantage Population-Based Algorithms. *IEEE Transactions on Evolutionary Computation*, 2010, 14(4), 500-517.
- Pourquié O. The Segmentation Clock: Converting Embryonic Time into Spatial Pattern, *Science*, 2003, 301: 328–330.
- Qin A. K., Huang V. L., and Suganthan P. N. Differential Evolution Algorithm with Strategy Adaptation for Global Numerical Optimization. *IEEE Transactions on Evolutionary Computation*, 2009. 13(2): 398-417.
- Quaiser T., Dittrich A., Schaper F. and Mönnigmann M. A Simple Work Flow for Biologically Inspired Model Reduction – Application to Early JAK-STAT Signaling. *BMC Systems Biology*, 2011, 5: 30-43.

- Raue A., Kreutz C., Maiwald T., Bachmann J., Schilling M., Klingmüller U. and Timmer J. Structural and Practical Identifiability Analysis of Partially Observed Dynamical Models by Exploiting the Profile Likelihood, *Bioinformatics*, 2009, 25(15): 1923-1929.
- Raue A., Kreutz C., Maiwald T., Klingmüller U. and Timmer J. Addressing Parameter Identifiability by Model-Based Experimentation. *IET Systems Biology*, 2011, 5(2): 120-130.
- Rechenberg I. Evolution Strategy. *Computational Intelligence: Imitating Life*, 1994: 147-159.
- Rodriguez-Fernandez M., Mendes P. and Banga J.R. A Hybrid Approach for Efficient and Robust Parameter Estimation in Biochemical Pathways, *Biosystems*, 2008, 83(2-3): 248-265.
- Rodriguez-Fernandez M., Egea J.A. and Banga J.R. Novel Metaheuristic for Parameter Estimation in Nonlinear Dynamic Biological Systems, *BMC Bioinformatics*, 2006, 7(1): 483-501.
- Roeva O. Genetic Algorithms for a Parameter Estimation of a Fermentation Process Model: A Comparison, *Bioautomation*, 2005, 3: 19-28.
- Runarsson T.P. and Yao X. Stochastic Ranking for Constrained Evolutionary Optimization. *IEEE Transactions on Evolutionary Computation*, 2000, 4(3): 284-294.
- Salton M.R.J. and Kim K.S. Structure, in Baron S. (Ed.) *Medical Microbiology 4th Edition*, University of Texas Medical Branch, 1996: Available online: <http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=mmed&part=A289&rendertype=figure&id=A294>.
- Schlatter R., Conzelmann H., Gilles E., Sawodny O. and Sauter T. Analysis of an Apoptotic Core Model Focused on Experimental Design using Artificial Data, *IET System Biology*, 2009, 3(4): 255–265.
- Sims G.K. Nitrogen Starvation Promotes Biodegradation of N-heterocyclic Compounds in Soil, *Soil Biology and Biochemistry*, 2006, 38(8): 2478-2480.
- Slezak D.F., Suarez C., Cecchi G.A., Marshall G. and Stolovitzky G. When the Optimal is Not the Best: Parameter Estimation in Complex Biological Models. *PLoS One*, 2010, 5(10): e13283.
- Stiert B., Raue A., Timmer J. and Kreutz C. Experimental Design for Parameter Estimation of Gene Regulatory Networks. *PLoS ONE*, 2012, 7(7): e40052.

- Stephan R., Katrin M., Gabriele V., Thomas W., Wolfgang T., Andreas N., Ulrich Z., and Matthias R. A Systems Biology Approach to Dynamic Modeling and Inter-Subject Variability of Statin Pharmacokinetics in Human Hepatocytes. *BMC Systems Biology*, 2011, 5: 66-85.
- Storn R. and Price K. Differential evolution - A Simple and Efficient Heuristic for Global Optimization Over Continuous Spaces. *Journal of Global Optimization*, 1997, 11, 341–359.
- Sun J., Garibaldi J. and Hodgman C. Parameter estimation using meta-heuristics in systems biology: a comprehensive review. *IEEE/ACM Transactions on Computational Biology and Bioinformatics*, 2012, 9(1): 185-202.
- Sun X., Jin L. and Xiong M. Extended Kalman Filter for Estimation of Parameters in Nonlinear State-Space Models of Biochemical Networks. *PLoS One*, 2008, 3(11): e3758.
- Surovtsova I., Simus N., Hübner K., Sahle S. and Kummer U. Simplification of Biochemical Models: A General Approach Based on the Analysis of the Impact of Individual Species and Reactions on the Systems Dynamics. *BMC Systems Biology*, 2012, 6: 14.
- Syberfeldt A., Ng A., John R.I. and Moore P. Evolutionary Optimisation of Noisy Multi-Objective Problems using Confidence-Based Dynamic Resampling. *European Journal of Operational Research*, 2010, 204(3): 533-544.
- Sybesma W., Starrenburg M., Kleerebezem M., Mierau I., De Vos W.M. and Hugenholtz J. Increased Production of Folate by Metabolic Engineering of *Lactococcus lactis*. *Applied and Environmental Microbiology*, 2003, 69(6): 3069-3076.
- Szekely D., Vandenberg J.I., Dokos S. and Hill A.P. An Improved Curvilinear Gradient Method for Parameter Optimization in Complex Biological Models. *Medical and Biological Engineering and Computing*, 2011, 49(3): 289-296.
- Tabata T., and Takei Y. Morphogens, their Identification and Regulation. *Development*, 2004, 131: 703–712.
- Takahashi K., Arjunan S.N. and Tomita M. Space in Systems Biology of Signaling Pathways-Towards Intracellular Molecular Crowding *In Silico*, *FEBS Letter*, 2005, 579(8): 1783-8.
- Tanaka U., Ogata Y. and Stoyan D. Parameter Estimation and Model Selection for Neyman-Scott Point Processes, *Biometrical Journal*, 2008, 50(1): 43-57.

- Tang Y. and Guan X. Parameter Estimation for Time-Delay Chaotic System by Particle Swarm Optimization. *Chaos, Solitons and Fractals*, 2009, 40(3): 1391-1398.
- Tashkova K., Korosec P., Silc J., Todorovski L. and Dzeroski S. Parameter Estimation with Bio-Inspired Meta-Heuristic Optimization: Modeling the Dynamics of Endocytosis. *BMC Systems Biology*, 2011, 5(1): 159.
- Tian T. and Song J. Mathematical Modelling of the MAP Kinase Pathway Using Proteomic Datasets. *PLoS One*, 2012, 7(8): e42230.
- Turner T.E., Schnell S. and Burrage K. Stochastic Approaches for Modelling *In Vivo* Reactions. *Computational Biology and Chemistry*, 2004, 28: 165-178.
- Tutkun N. Parameter Estimation in Mathematical Models using the Real Coded Genetic Algorithms. *Expert Systems with Applications*, 2009, 36(2): 3342-3345.
- Veening J.W., Igoshin O.A., Eijlander R.T., Nijland R., Hamoen L.W. and Kuipers O.P. Transient Heterogeneity in Extracellular Protease Production by *Bacillus subtilis*. *Molecular and Systems Biology*, 2008, 4(1): 184-209.
- Villarreal M.R. Average Prokaryote Cell, Wikimedia Commons: 2008. Available at: [http://en.wikipedia.org/wiki/File:Average\\_prokaryote\\_cell-\\_en.svg](http://en.wikipedia.org/wiki/File:Average_prokaryote_cell-_en.svg) (Accessed June 2011)
- Vilas C., Balsa-Canto E., García S.G., Banga J.R. and Alonso A.A. Dynamic Optimization of Distributed Biological Systems using Robust and Efficient Numerical Techniques. *BMC Systems Biology*, 2012, 6: 79.
- Wolpert D.H. and Macready W.G. No Free Lunch Theorems for Optimization. *IEEE Transactions on Evolutionary Computation*, 1997, 1(1): 67-82.
- Yang X.S. Firefly Algorithms for Multimodal Optimization. *Stochastic Algorithms: Foundations and Applications*, 2009, 5792: 169-178.
- Yates J.W.T., Evans N.D. and Chappell M.J. Structural Identifiability Analysis via Symmetries of Differential Equations. *Automatica*, 2009, 45(11): 2585-2591.
- Yuting Z. and Ganesh S. Mathematical Modeling: Bridging the Gap Between Concept and Realization in Synthetic Biology, *Journal of Biomedicine and Biotechnology*, 2010: DOI: 10.1155/2010/541609.
- Zeng N., Wang Z., Li Y., Du M. and Liu X. A Hybrid EKF and Switching PSO Algorithm for Joint State and Parameter Estimation of Lateral Flow Immunoassay Models. *IEEE/ACM Transaction on Computational Biology and Bioinformatics*, 2012, 9(2): 321-329.

Zhang W. and Zou X. The Data-based Mathematical Modeling and Parameter Identification in JAK-STAT Signaling Pathway by using a Hybrid Evolutionary Algorithm, *Advances in Computation and Intelligence*, 2011: 516-522.

optimization method on RSM based regression models. Karabulut and Karakoc [8] investigated the machinability of silicon carbide and aluminum alloy-based metal matrix composite during milling. operation. He used three different feature selection methods including the ACO based metaheuristics. Zuperl and Cus [29] proposed an approach of using ANFIS to represent the manufacturer's objective function and an Ant Colony Optimization algorithm (ACO) to obtain the optimal objective value. Besides these, Liao [12] presented hybrid differential evolution and harmony search algorithms for. optimizing fourteen engineering design problems selected from different engineering fields. These models usually depend on several parameters and initial conditions. If these parameters are unknown, results from simulation studies can be misleading. Such a scenario can be avoided by fitting the model to experimental data before analyzing the system. This involves parameter estimation which is usually performed by minimizing a cost function which quantifies the difference between model predictions and measurements. Results: In this work we propose a new hybrid global method, based on the combination of an evolutionary search strategy with a local multiple-shooting approach, which offers a reliable and efficient alternative for the solution of large scale parameter estimation problems.