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Multi-objective Optimization of Biochemical System Production Using an Improve Newton Competitive Differential Evolution Method

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Abstract— In this paper, an improved method of multi-objective optimization for biochemical system production is presented and discussed in detail. The optimization process of biochemical system production become hard and difficult when involved a large biochemical system that contains many components. In addition, the multi-objective problem also needs to be considered. Due to that, this study proposed and improved a method that comprises with Newton method, differential evolution algorithm (DE) and competitive co-evolutionary algorithm(ComCA). The aim of the proposed method is to maximize the production and simultaneously minimize the total amount of chemical concentrations involves. The operation of the proposed method starts with Newton method by dealing with biochemical system production as a nonlinear equations system. Then DE and ComCA are used to represent the variables in nonlinear equation system and tune the variables in order to find the best solution. The used of DE is to maximize the proposed method is evaluated using two benchmark biochemical systems, and the experimental results show that the proposed method performs well compared to other works.

Keywords- Newton method; differential evolution algorithm; competitive co-evolutionary algorithm; biochemical system

I. INTRODUCTION

Solving a nonlinear equation system requires the process of finding all solutions for each equation in the system. It is hard task because the equations are usually а nondeterministic polynomial equations. In addition, nonlinear equations system has a complex structure due to the number equations and variables, which makes it difficult to solve [1], [2]. Nowadays, many applications use nonlinear equations system such as in the chemistry domain [3], [4]. One example application in the chemistry domain that uses nonlinear equations system is multi-objective optimization of the production in a biochemical system. In the multiobjective optimization of the production in a biochemical system, many researchers use nonlinear equations system to represent the chemical reactions. This can be achieved by using the knowledge of biotechnology where the chemical

reactions can be represented by a mathematical model, known as ordinary differential equations (ODEs) model.

In the optimization process, the ODEs model is used to model the biochemical system. The optimization of biochemical system production can be considered as solving nonlinear equations system because the optimization process always constrained by steady state condition. This situation makes all the ODEs model equal to 0 thus lead to the operation of solving the nonlinear equations system. Currently, there are many methods used in solving a nonlinear equation system, for example, Newton method [5], [6], Secant method [7], [8], and Bisection method [9], [10]. Among these methods, it was found that Newton method is the suitable method to be used in solving a nonlinear equation system. This is due to advantages offered by Newton method which is the convergence speed of Newton method [11]-[13] and the simplicity in using Newton method [2], [11]. Due to that, this study use Newton method in solving nonlinear equations system.

In order to improve the biochemical system production, the variable value in nonlinear equations system needs to be altered and tuned. The tuning process of all variables in nonlinear equations system is performed to find the suitable value that able to produce the best result. The tuning is a process of changing the value of all variables in nonlinear equations system. The process becomes hard and difficult when involves a large and complex structure of biochemical system where a large and complex structure of biochemical system make nonlinear equations system become more complex. The large and complex biochemical system involves many chemical reactions and involves many interactions between them. Generally, a large biochemical system comprises with many variables where these variables represent the components and the interaction in the biochemical system [14]. Due to the involvement of many variables, the optimization process becomes a problem where it makes the optimization process become hard and complicated. Due to that, there is need an automated approach in tuning all variables in nonlinear equations system [15]. There are many methods available in automated tuning approaches such as genetic algorithm (GA), differential evolution (DE) algorithm, Particle Swarm Optimization (PSO) algorithm and Artificial Bee Colony (ABC) algorithm. Based on comparative studies, used DE is more suitable compared to others because DE is more robust [16] and only involved a few parameters control [17], [18]. In this work, the DE is used to represent the variables in nonlinear equations system in chromosome form. Then, the chromosome is tuned in order to search the best solution that can produce the best result in the optimization process.

In the multi-objective optimization of the production in a biochemical system, two objectives need to be considered. The first objective is to maximize the production while the second objective is to minimize the total amount of chemical concentrations involves. Besides the multi-objective optimization, several constraints are involved which contribute to the difficulty in optimization process [3]. The first constraint is a steady-state constraint. The purpose of this constraint is to ensure a continuous optimal operation of a biochemical system. The second constraint is the chemical reaction constraint. This constraint makes the biochemical concentration remains within a specific range to maintain the survival of the cell. Because the multi-objective optimization of the production in a biochemical system involves two objectives, the optimization process become hard. The two objectives make the evaluation process of the solution that represents by DE become hard. Moreover, it is timeconsuming to evaluate the solution. Like other methods used in tuning approach, DE also suffers from long computational times because of their evolution/iteration process. Therefore, there is a method to embodied into DE to improve its performance. Apply competitive coevolutionary algorithm (ComCA) is a good choice. This is because of the nature of ComCA where ComCA decomposed a problem into multi sub-problems. The used of ComCA is used in handling the multi-objective problem where ComCA breaks the multiobjective into sub-objective by dividing the population of the chromosome into two part. The first population is for improving the biochemical objective while the second is for minimizing the total amount of chemical concentrations involves.

In this study, an improved method that knowns as Newton Competitive DE (NComDE) are presented. The NComDE combine Newton method, DE, and ComCA. The Newton method is used in solving the nonlinear equations system, DE for tuning the variable in nonlinear equations system while ComCA for dealing with multi-objective optimization. In order to access the performance of the NComDE, two biochemical system are chosen which are ethanol production in Saccharomyces cerevisiae (S. cerevisiae) pathway and the optimization of tryptophan (trp) biosynthesis in Escherichia coli (E. Coli) pathway. In the following section, the methods that were used in this study are discussed in detail. It starts with the DE algorithm then followed by the ComCA. Then the proposed method is presented in detail. This is followed by a discussion of case studies before result and discussion are discussed and finally followed by a conclusion.

II. MATERIAL AND METHODS

A. Differential Evolution Algorithm

DE is one of the population-based algorithm used to solve many optimization problems. The advantages of using DE include for its ease of use, speed, a simple structure where it used the real number and its efficiency a robustness over many problems [19]–[22]. DE operates by maintaining a population of candidate solutions known as chromosome for an optimization problem. There are three operations in DE including mutation, crossover, and selection. The NComDE which combine Newton method, DE and ComCA began with a population of m candidate solutions and defined as follows:

$$X_{m} = \{ \mathbf{x}_{m1}, \mathbf{x}_{m2}, \dots, \mathbf{x}_{mn} \}$$
(1)

where *m* is the number of chromosomes while *n* is denoted as the number of variable in nonlinear equations system. The variables in nonlinear equations system are defined in a specific range which is x_{mn}^{l} for upper range while x_{mn}^{u} for the lower range are as follows:

$$x_{mn} = \{x_{mn}^{l} + rand(x_{mn}^{u} - x_{mn}^{l})\}$$
(2)

where rand is random number in the range of 0 and 1 [23]. For expends the search/optimization space, DE use mutation operation where it creates a trivial chromosome as follows:

$$V_m = X_{best(g)} + F(X_{rand1(g)} - X_{rand2(g)})$$
(3)

where $X_{best(g)}$ is the current best chromosome, *F* is scaling factor while $X_{rand1(g)}$ and $X_{rand2(g)}$ are randomly chosen chromosome in the current generation, *g*. Then, the trivial chromosome with a parent is mixed together to produce Y_m number of their offspring in crossover operation. The crossover operation used the following rule to produce offspring:

$$Y_{m(g)} = \begin{cases} V_{m(g)} \text{ if } rand_2 < CR \\ X_{m(g)} \text{ Otherwise} \end{cases}$$
(4)

where *CR* is the rate of crossover. The *rand*₂ is the random number between 0 and 1 [23]. After the crossover operation is performed, then the $Y_{m(g)}$ is compared with its parent. This is intended to determine which one has good quality and survive in the next generation using rule as follows:

$$X_{m(g+1)} = \begin{cases} Y_{m(g)} \text{ if } f(Y_{m(g)}) \le f(X_{m(g)}) \\ X_{m(g)} \text{ if Otherwise} \end{cases}$$
(5)

B. Competitive Coevolutionary Algorithm

The concept of the coevolutionary algorithm (CA) was proposed by Darwin in work entitled On the Origin of species [24]. The main objective of CA was to enhance the performance of the evolutionary algorithm method in solving the complex optimization problem especially in solving the multi-objective problem. The CA can be viewed as a collection if EAs where CA can simplify the complex problem into multiple sub-components of problem. By doing this, the sub-components will evolve separately to solve the optimization problem.

The ComCA is one type of CA where it refers to the competition of sub-components. It happens when the fitness of sub-component is in competition with another sub-component. This lead to an arms race where the race tends to produce a new strategy in order make the solution to be improved. The fitness evaluation process (arms race) can be performed when a representative from every sub-components was chosen and being evaluated based on the optimization problem. Then, the winner from the evaluation process is chosen to go to the next step in solving the optimization problem.

C. An Improved Method of Newton Competitive Differential Evolution Algorithm

In this section, the proposed method that combines Newton method, ComCA and DE that known as NComDE is presented. In the NComDE, the Newton method is used in solving the nonlinear equations system, while the ComCA and DE is used for multi-objective problem where ComCA for minimizing the total amount of chemical is concentrations involves by populate the chromosome into two separate population and DE for maximize the biochemical system production by tuning the variable in nonlinear equations system. In NComDE, two populations of chromosomes are used to represent the variables in nonlinear equations system. The first population focus on maximizing the production while the second population is for minimizing the total amount of chemical concentrations involves. The number of chromosome in both populations is same. The flowchart of NComDE is given in Fig. 1. Next, are the steps involved in the NComDE.

Step 1: Generation of the initial population. This step is about the first generation of the chromosome in two separate population. The generation process is perform using equation 1. The first population is generated to improve the production while the second population is to minimize the total amount of chemical concentrations involves. The chromosomes in both population are in binary format. The representation of the variables in nonlinear equation system in chromosome form is given in Fig. 2.

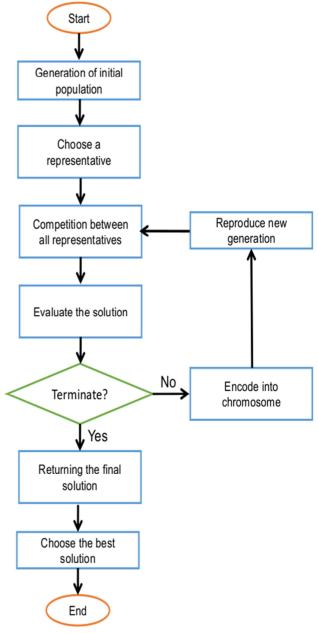


Fig. 1 The flowchart of NComDE

1) Step 2: Choose a representative. This step about the choosing process of a representative from all population. The selection process of representation is based on their fitness where it refers to the objective of each population. For the first population, the chromosome that has higher fitness value (denoted as F_1) is chosen as a representative due to this population is about improve the production. The F_1 is the fitness value where it comes from the production rate. For the second population, the chromosome with lowest fitness value (denoted as F_2) is selected as this population is aiming to minimize the total amount of chemical concentrations involves. The F_2 is the fitness value where it comes from the total amount of chemical involves.

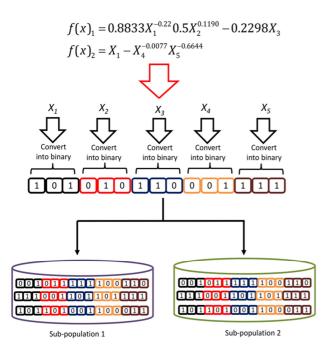


Fig. 2 The representation of the variables in chromosome form

2) Step 3: Competition between all representatives. This step is about the selection of a winner between all population. In this step, all representatives compete with each other, and the winner is chosen based on the representative that has higher fitness function (denoted as F_3). The F_3 is formulated in equation 6. Then, the representative that has higher of F_3 is chose as a solution.

$$F_3 = \left|\frac{F_1 - F_2}{2}\right| \times 100$$
 (6)

3) Step 4: Evaluate the solution. This step is about the evaluation process of the representative that chosen from the previous step. The evaluation process starts with the decoding process of the solution into variables in nonlinear equations system. The Newton method is used to in solving the nonlinear equations system. At this stage, termination conditions occur, which is whether the steady-state constraint and chemical reaction constraint are fulfilled or not. If the solution meets the termination conditions, if move forward to Step 7, otherwise it enters the next step.

4) Step 5: Encode into the chromosome. In this stage, the variables in nonlinear equations system are encoded back into chromosome form after being tested by Newton method. Then, the chromosome goes back into its population for reproduction process.

5) Step 6: Reproduce new generation. In this step, the selection, mutation, and crossover operations are applied to all populations to produce the new generation.

6) Step 7: Returning the final solution. In this step, the best set of solutions (variables in the nonlinear equations system) that obtained from the evolution process is given.

In order to show the performance of the proposed method, two benchmark case studies were used, namely the optimization of ethanol production in *S. cerevisiae* pathway and the optimization of *trp* biosynthesis in *E. Coli*. To run the experiment, a Java program was developed that was based on jMetal [25] and JAMA version 1.0.3 were used. The jMetal can be downloaded from http://jmetal.sourceforge.net/ index.html while JAMA can be obtained from http://math.nist.gov/javanumerics/jama/.

A. Case Study 1: Optimization of Ethanol Production in Saccharomyces Cerevisiae Pathway

In *S. cerevisiae* pathway, the method that proposed in this study is used to optimize the ethanol production. The detail description of this pathway is discussed in the work that proposed in [26]. The ODEs model that represents this pathway is given by the equation 7 as follows:

$$\frac{dX_1}{dt} = V_{in} - V_{HK}$$

$$\frac{dX_2}{dt} = V_{HK} - V_{PFK} - V_{Carb}$$

$$\frac{dX_3}{dt} = V_{PFK} - V_{GAPD} - 0.5V_{Gro}$$

$$\frac{dX_4}{dt} = 2V_{GAPD} - V_{PK}$$

$$\frac{dX_5}{dt} = 2V_{GAPD} + V_{PK} - V_{HK} - V_{Carb} - V_{PFK} - V_{ATPase}$$
(7)

All fluxes (denoted by V) has the following rates when it is in steady state condition:

$$V_{in} = 0.8122 X_{2}^{-0.2344} Y_{1}$$

$$V_{HK} = 2.8632 X_{1}^{0.7464} X_{5}^{0.0243} Y_{2}$$

$$V_{PFK} = 0.5232 X_{2}^{0.7318} X_{5}^{-0.3941} Y_{3}$$

$$V_{Carb} = 8.904 \times 10^{-4} X_{2}^{8.6107} Y_{6}$$

$$V_{GAPD} = 7.6092 \times 10^{-2} X_{3}^{0.6159} X_{5}^{0.1308} Y_{4}$$

$$V_{Gro} = 9.272 \times 10^{-2} X_{3}^{0.05} X_{4}^{0.533} X_{5}^{-0.0822} Y_{7}$$

$$V_{PK} = 9.471 \times 10^{-2} X_{3}^{0.05} X_{4}^{0.533} X_{5}^{-0.0822} Y_{5}$$

$$V_{ATPase} = X_{5} X_{8}$$
(8)

The ethanol production that was tried to be improved in this pathway is given by the flux V_{PK} thus it becomes the F_I . Therefore, the first objective (first population) of the multi-objective problem in this pathway is as follows:

$$max F_1 = V_{PK} \tag{9}$$

For the second objective, F_2 (second population), the proposed method is trying to minimize the total amount of chemical concentrations involves and it can be formulated as follows:

$$\min F_2 = \sum_{j=1}^{5} X_j + \sum_{j=6}^{6} Y_j$$
(10)

In the steady state constraint, all the ODEs model are force to be equals to 0 [3], [4]. This make equation 7 become as follows:

$$V_{in} - V_{HK} = 0$$

$$V_{HK} - V_{PFK} - V_{Carb} = 0$$

$$V_{PFK} - V_{GAPD} - 0.5V_{Gro} = 0$$

$$2V_{GAPD} - V_{PK} = 0$$

$$2V_{GAPD} + V_{PK} - V_{HK} - V_{Carb} - V_{PFK} - V_{ATPase} = 0$$
(11)

Meanwhile, the chemical reaction constraint in this pathway is divided into two categories, metabolites constraint and enzymes constraint. For metabolites constraint, all metabolites were set in range 0.8 to 1.2 and given in equation 12, while enzymes constraint were in the range 0-50 and given in equation 12 [3], [4].

$$\begin{split} X_{j}^{0.8} &\leq X_{j} \leq X_{j}^{1.2} \qquad j = 1, 2, 3, 4, 5 \\ Y_{j}^{0} &\leq Y_{j} \leq Y_{j}^{50} \qquad j = 1, 2, 3, 4, 5, 8 \end{split} \tag{12}$$

B. Case Study 2: Optimization of Tryptophan Production in Escherichia Coli Pathway

For case study 2, the proposed method was tried to optimize the end product of *E.coli* which is *trp*. Detail description of this pathway can be further read in works that performed by Xiu and co-workers [27]. The ODEs model of this pathway is given by equation 13.

$$\frac{dX_1}{dt} = V_{11} - V_{12}$$

$$\frac{dX_2}{dt} = V_{21} - V_{22}$$

$$\frac{dX_3}{dt} = V_{31} - V_{32} - V_{33} - V_{34}$$
(13)

At the steady state condition, the rate of all reactions (denoted by *V*) has the following value:

$$V_{11} = 0.6403 X_{3}^{-5.87 \times 10^{-4}} X_{5}^{-0.8332}$$

$$V_{12} = 1.0233 X_{1} X_{4}^{0.0035} X_{11}^{0.9965}$$

$$V_{21} = X_{1}$$

$$V_{22} = 1.4854 X_{2} X_{4}^{-0.1349} X_{12}^{0.8651}$$

$$V_{31} = 0.5534 X_{2} X_{3}^{-0.5573} X_{6}^{0.5573}$$

$$V_{32} = X_{3} X_{4}$$

$$V_{33} = 0.9942 X_{3}^{7.0426 \times 10^{-4}} X_{7}$$

$$V_{34} = 0.8925 X_{3}^{3.5 \times 10^{-6}} X_{4}^{0.9760} X_{8} X_{9}^{-0.0240} X_{10}^{-3.5 \times 10^{-6}}$$
(14)

The *trp* production was given by reaction V_{34} and become the F_1 . Therefore, the first objective (first population) of the multi-objective problem in case study two is given as follows:

$$max F = V_{34} \tag{15}$$

For the second objective, F_2 (second population), the proposed method is trying to minimize the total amount of chemical concentrations involves and it can be formulated as follows:

$$\min F_2 = \sum_{j=1}^{6} X_j + X_8 \tag{16}$$

In the steady-state constraint, all the ODEs model are a force to be equal to 0 [28]. This makes equation 13 become as follows:

$$V_{11} - V_{12} = 0$$

$$V_{21} - V_{22} = 0$$

$$V_{31} - V_{32} - V_{33} - V_{34} = 0$$
(17)

For the chemical reaction constraint of this pathway, only several involves in the fine-tuning process which was X_1 to X_6 and X_8 , while the other components used a fixed value. The chemical reactions that involve were given as follows:

$$X_{j}^{0.8} \leq X_{j} \leq X_{j}^{1.2} \qquad j = 1, 2, 3$$

$$0 \leq X_{4} \leq 0.00624$$

$$4 \leq X_{5} \leq 10$$

$$500 \leq X_{6} \leq 5000$$

$$X_{7} = 0.0022X_{5}$$

$$0 \leq X_{8} \leq 1000$$

$$X_{9} = 7.5$$

$$X_{10} = 0.005$$

$$X_{11} = 0.9$$

$$X_{12} = 0.02$$

$$X_{13} = 0$$

(18)

III. RESULTS AND DISCUSSION

In accessing the performance of the proposed method, several experiments were performed. The Newton method use fixed parameters setting which are the number of iterations was fixed to 100, and the tolerance was set to 10^{-6} . For the DE parameters, Table 1 list the DE parameters setting in obtaining the best result.

TABLE I PARAMETERS SETTING IN PRODUCING THE BEST RESULT

Parameter	S. cerevisiae pathway	<i>E.coli</i> pathway	
No of chromosomes	250	200	
No of generations	300	300	
Scaling factor	0.8	0.7	
Crossover rate	0.2	0.2	

Table 2 gives the result obtained by the NComDE when the NComDE uses on the *S. cerevisiae* pathway. The best result obtained by the NComDE is 52.93 compared to its steady-state value. This means that the NComDE is able to improve the ethanol production. Meanwhile, the NComDE able to reduce the total amount of chemical concentrations involves where the value is 294.8 compared to its steadystate value. In addition, it can be found that the performance of the NComDE outperforms the result from other works in terms of improving the ethanol production and at the same time reduce the total amount of chemical concentrations involves.

For the *E.coli* pathway, the best result produced by the NComDE is 3.99 compared to its steady state value where it shows the improvement of *trp* production. For the total amount of chemical concentrations involves, the NComDE

able to reduce it to 6015.01. For the detail result, Table 3 lists the results including the comparison with previous works. Like the previous pathway, the performance of the NComDE outperforms the other works and confirm that the proposed method able to simultaneously improve the production and reduce the total amount of chemical concentrations involves.

TABLE II
THE RESULT OBTAINED BY NCOMDE IN S. CEREVISIAE PATHWAY

	Best	Work	Work	Work	Work
	Solution	by [29]	by [4]	by [30]	by [15]
X_{I}	0.9590	1.14	1.102	1.11	1.11
X_2	0.8087	1.05	1.046	1.03	1.03
X_3	0.8479	1.15	1.141	1.13	1.19
X_4	1.1524	1.17	1.171	1.18	1.17
X_5	0.9033	1.12	1.113	1.14	0.91
Y_1	49.9170	49.97	50	49.99	49.73
Y_2	44.8006	44.77	45.953	45.83	45.81
Y_3	49.6112	49.89	50	49.92	48.89
Y_4	47.2629	47.26	47.772	47.97	48.13
Y_5	48.7968	48	48.366	48.30	47.85
Y_8	49.7482	49.75	50	49.79	48.98
F_{I}	52.97	52.0843	52.5118	52.57	52.91
F_2	294.80	295.28	297.664	297.384	294.80

TABLE III THE RESULT OBTAINED BY NCOMDE IN E.COLI PATHWAY

	Best	Work	Work	Work	Work
	Solution	by [31]	by [28]	by [4]	by [30]
X_{I}	1.06	1.19	1.2	1.2	1.11
X_2	1.11	1.15	1.15	1.12	1.114
X_3	0.8	0.8	0.8	0.8	0.8
X_4	0.0054	0.0041	0.004	0.0054	0.0054
X_5	4.50	4	4	4.011	4.75
X_6	5000	5000	5000	5000	5000
X_8	1000	1000	1000	1000	1000
F_{I}	3.99	3.06	3.06	3.95	3.98
F_2	6015.01	6016.38	6016.38	6016.57	6016.22

In addition, the NComDE is compared with the method that not apply the ComCA (Newton DE). The purpose in using ComCA is to reduce the total amount of chemical concentrations involves. Several experiments were conducted using parameters setting in Table 1, Fig. 3 and Fig. 4 give the comparison of NComDE and Newton DE in a line graph. From the graph, it can be observed that all NComDE results are lower compared to Newton DE thus confirm that using the ComCA able to reduce the total amount of chemical concentrations involves.

Besides that, the distribution result of ethanol production and *trp* production was given by the box plot in Fig.5 and Fig. 6. The results of these figures were collected from 100 independent runs and use parameters setting form Table 1. From both figures, it can be observed that the performance of NComDE outperformed the performance of Newton DE where the distribution of results produced by NComDE was not too wide compared to the results produced by Newton DE. This point out that the performance of NComDE was consistent. It can be suggested that NComDE could produce better results if the experiment runs several times.

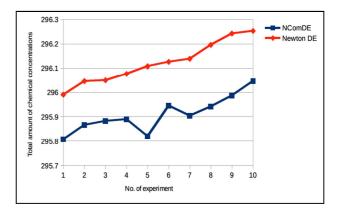


Fig. 3 The comparison of the total amount of chemical concentrations between NComDE and Newton DE in S. cerevisiae pathway

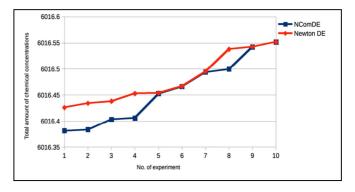


Fig. 4 The comparison of the total amount of chemical concentrations between NComDE and Newton DE in E.coli pathway

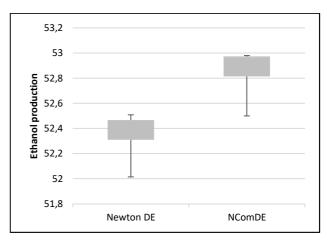


Fig. 5 The box plot of ethanol production

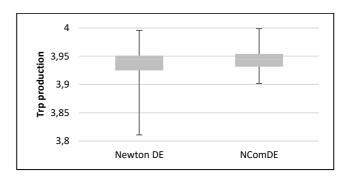


Fig. 6 The box plot of trp production

IV. CONCLUSION

As a conclusion, NComDE has a higher tendency in improving the biochemical system's production. In future development, this research could be enhanced by referring to various other works available such as [32]-[42].

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