

# Optimization of Environmental Conditions for remediation of contaminated water with oil using Plackett-Burman Model

## Abstract

Plackett-Burman is beneficial design not only in determining the significant variables of bioremediation, but also in optimization of these variables. In this study Plackett-Burman (PB) experimental model had been applied to assess the significant of some nutritional and environmental condition affecting oily wastewater bioremediation by *Aspergillus niger*. Eleven variables through twelve assays were planned, namely: Temperature, pH, sucrose,  $\text{KH}_2\text{PO}_4$ ,  $\text{NaCl}$ ,  $\text{MgSO}_4$ ,  $\text{Na}_2\text{HPO}_4$ ,  $\text{NH}_4\text{Cl}$ ,  $\text{NaNO}_3$ , urea and spore suspension to explain their effects on oily wastewater removal. The degradation process was enhanced based on oil and grease experiment. pH, sucrose,  $\text{KH}_2\text{PO}_4$ ,  $\text{NaCl}$ ,  $\text{Na}_2\text{HPO}_4$  and urea were recognized as the positive factors that stimulate the degradation. On the other hand, other variables affected negatively on oily wastewater removal. The regression coefficient  $R^2$  (0.99) ensure the adequate integrity of the model. Plackett-Burman Model was used to optimize the method of oily wastewater bioremediation by fungal isolate. It was showed that sucrose(A),  $\text{NaCl}$  (B) and  $\text{KH}_2\text{PO}_4$ (C) give high removal of oily wastewater when approaching to +1, on the other hand, the temperature (D) give high removal of oily wastewater when approaching to -1. B had positive effects on oily wastewater degradation, whereas C, D had negative effects. The factor with confidence level above 95% is considered as significant parameter. It was clear that variable B was the chief factor, while variables A, C, D, with levels below 95%, were considered insignificant.

**Keywords:** Plackett- Burman design, Bioremediation, *Aspergillus niger*, optimization

## 1. Introduction

Industrial wastewater includes some major pollutants that affect negatively on public health as petroleum hydrocarbon, heavy metals and dyes. Diesel oil that polluted wastewater has been considered as one of the most concerned pollution sources. Oily wastewater produced from different sources such as crude oil production, automotive garage, oil refinery, petrochemical industry, metal processing, lubricant and car washing. These sources serve as the major contributor to the environmental problems especially in soil and water. Wastewater contains some toxic constituents such as petroleum hydrocarbons, phenols, polyaromatic

33 hydrocarbons which are inhibitory to animal and plant growth and also are mutagenic and  
34 carcinogen to human being [1]. Hydrocarbons have an effect on aquatic life, coastal  
35 environment and habitats as coral reefs and the marine organisms in addition to causing  
36 damage and death to agricultural crops [2]. As concern for effect of hydrocarbon  
37 contaminated wastewater continues to grow, so are the different technologies that continue to  
38 emerge to remediate contaminated site. One of these technologies is bioremediation.  
39 Bioremediation of oily wastewater is treatment technology that use of microorganisms or  
40 their enzymes to reduce the concentration or toxicity of hydrocarbon contaminants into less  
41 toxic forms [3]. Microorganisms as fungi and bacteria may be indigenous to contaminated  
42 area or they are isolated from other area and brought to the hydrocarbon contaminated area  
43 [4]. According to Atlas [5], the bioremediation process can be occurred at site, less expensive, less  
44 required energy and can be used with other physical or chemical treatment method.

45 Fungi have several advantages over bacteria. Due to their hyphal growth mode they can form  
46 mycelial networks, which they can use to transport water, nutrients and electron acceptors  
47 within mycelia. Unlike bacteria they can also grow though air-filled pores and penetrate soil  
48 aggregates. Furthermore, because these produced fungal enzymes have low specificity they  
49 are able to degrade a wide range of organic compounds and mixtures of various chemicals  
50 [6].

51 The capability of microorganisms to degrade contaminants and growth of cells are strongly  
52 affected by nutritional and ecological factors such as carbon sources, nitrogen sources,  
53 inorganic salts, temperature, and pH. These experiments may take several times and consume  
54 large amounts of chemicals for achieving optimization. Therefore, it is essential to design  
55 suitable process for maximizing the removal efficiency of diesel oil by *Aspergillus niger*.  
56 Plackett-Burman design provides a fast and effective way to identify the important factors  
57 among a large number of variables, thereby, saving time and maintaining convincing  
58 information on each parameter [7]. In the present study, a Plackett-Burman model has been  
59 employed to determine the weighty factors and optimize degradation. The main aim of the  
60 present study is to determine the optimum concentrations of some nutritional and  
61 environmental parameters affecting diesel oil removal from industrial wastewater using  
62 *Aspergillus niger*.

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## 2. Materials and Methods

### 2.1 Media preparation

The enrichment procedure as described by Nwachukwu [8] was used in the estimation of hydrocarbon utilizes. A minimal salt broth containing 2.0g of Na<sub>2</sub>HPO<sub>4</sub>, 0.17g of K<sub>2</sub>SO<sub>4</sub>, 4.0g of NH<sub>4</sub>NO<sub>3</sub>, 0.53g of KH<sub>2</sub>PO<sub>4</sub>, 0.10g of MgSO<sub>4</sub> · 7H<sub>2</sub>O and 5.0g of agar – agar dissolved in 1000 ml of distilled water was prepared. The solution was sterilized by autoclaving. Diesel oily wastewater is added as main carbon source in concentration of 0.5% v/v.

### 2.2 Isolation and identification of microorganisms

Plates were incubated at room temperature (29° – 31°C) for 15 days. The fungus used in this study was isolated from minimal salt broth medium. It was purified and identified morphologically as *Aspergillus niger*.

### 2.3 Plackett-Burman experiment

Every variable was prepared from stock solution to obtain accurate results. The different factors were prepared in three levels: -1 for low concentration, 0 for medium concentration and +1 for high concentration, depending on Plackett- Burman modeling design [9]. A control experiment was made for every assay in this design as the same manner of experiment without spore suspension. Table (1) illustrates the factors under investigation and their levels that used in this model. Oily wastewater concentration was kept constant in all trails at the level of 0.5%.

### 2.4 Analytical determination of oil hydrocarbons

Oil concentration was determined gravimetrically using petroleum ether extraction method in acidified medium according to standard methods for examination of water and wastewater [10]. Petroleum biodegradation % based on oil and Grease content. It was calculated according to the following equation:

$$\text{Petroleum degradation \%} = \frac{\text{control} - \text{fungal}}{\text{control}} * 100 [1]$$

Where *control*= amount of oil and grease without spore suspension. *Fungal*= amount of oil and grease with spore suspension.

### 2.5 Plackett-Burma experimental design

For screening purpose, various medium components have been evaluated using Plackett–Burman (PB) statistical design [9]. The different factors were prepared in two levels: -1 for low level, 0 for medium level and +1 for high level. Table (1) illustrates the factors under investigation as well as levels of each factor used in the experimental design. The nitrogen compounds were prepared in equimolar bases to give 0.2 M nitrogen for higher concentration (+1) and the compounds that containing carbon and phosphorus were prepared to give 0.04 M

phosphorus for the higher level assays (+1). Petroleum oil concentration was kept constant in all assays at the level of 0.5%. The PB experimental design is based on the first order model (2):

$$Y = a_0 + \sum_{i=1}^n a_i x_i \quad (1)$$

Where Y is the response (productivity or specific activity),  $a_0$  is the model intercept and  $a_i$  is the variable estimates. This model describes no interaction among factors and is used to evaluate the important factors that influence petroleum oil bioremediation and fungal growth. Eleven variables were screened in twelve experiments; each variable being either medium constituent or environmental variable. Variables with high confidence levels are considered significant on their effect on petroleum bioremediation.

## 2.6 Screening of significant medium variables by PB design:

The PB design was applied to obtain the estimates of the different culture determinants for petroleum removal by *Aspergillus niger*. A polynomial model for oily wastewater removal% was developed by using the estimated coefficients (coded units) and given in Eq. (2).

$$Y = 56.66 + 0.61A + 3.96B - 9.43C - 1.66D + 25.81E - 5.09F - 6.40G - 3.03H + 3.22J - 2.93K - 2.5L \quad (2)$$

## 2.7 Optimization of significant nutritional and environmental parameters

Based on growth of *Aspergillus niger* in the preliminary experiment, four variables (Temperature,  $\text{KH}_2\text{PO}_4$ , sucrose and NaCl) were selected as the various nutritional and environmental parameters for Plackett-Burman design in this study. The concentrations for the different variables were selected according to some preliminary experiments and given in Eq. (3).

$$Y = 57.4 + 1.41A + 6.15B - 0.58C - 0.15D + 5.68AB + 1.72AC + 12.37AD - 0.16BC - 9.03BD - 4.95CD - 4.89ABD - 3.58ACD - 6.61BCD \quad (3)$$

## 3. Results and Discussion

Eleven variables were chosen in PB design that resembles the most important nutritional and environmental affecting oily wastewater bioremediation using fungal isolate Table (1). Twelve assays were made as stated by PB design and the response, oily wastewater removal was obtained as given in Table (2). The data listed in Table (2) showed a wide variation in oily wastewater degradation, from 7.6 to 96.3, in the 12 assays. The variation suggested that process optimization was important for improving the bioremediation efficiency. Figure (1)

shows that pH, sucrose,  $\text{KH}_2\text{PO}_4$ , **NaCl**,  $\text{Na}_2\text{HPO}_4$  and urea were recognized as the positive factors that stimulate the degradation. On the other hand, other variables affected negatively on diesel oily removal. Figure (2) shows the ranking of variable evaluations in a Pareto chart. The Pareto chart displays the magnitude of each factor estimate (independent on its contribution, either positive or negative) and is a appropriate technique to show the results of a PB model [11]. The highest positive significant variable is  $\text{KH}_2\text{PO}_4$ , while spore suspension is the highest negative significant variable. **Abd El Hamid [6]** shows that spore suspension had a highest contribution while pH had a lowest contribution for the growth of *Pencillium sp.* **Hegazy et al. [12]** showed that sodium dihydrogen phosphate, temperature, sodium chloride inhibited the growth of oil bioremediation. Figure (3) shows the model validation, a comparison was held between estimated and predicted results. The linearity of correlation is an evidence of the excellent agreement between experimental and predicted data. According to positive effects of selected variables on oily wastewater bioremediation, optimization of selected variables was chosen Table (3). **Zhou et al. [13]** stated that inoculum concentration was vital element for the xenobiotic compounds degradation. A similar result was observed by **Ghanem et al. [14]** for the chloroxylenol degradation by *Aspergillus niger*. Nitrogen sources like yeast extract has been proved to support a quick growth of cells and metabolites biosynthesis, as well as extracellular enzymes production.

Low concentration of ammonium sulfate leads to the high degradation of naphthalene in validated tests. The kinetics of ammonium uptake is first order and its specific uptake rate and microbial growth increase as the ammonium ion concentration increases. However, growth and ammonium uptake level reach a maximum and then decrease with the increase in aqueous ammonium ion concentration related to some possible inhibition mechanism [15]. Table (4) shows that B value had significant as positive effects on oily wastewater degradation, whereas C, D had negative effects. The variable with confidence level above 95% is considered as significant parameter. It was clear that variable B was the significant factor, while variables A, C, D, with confidence levels below 95%, were reflected an insignificant value. The statistical significance of the model was checked by F-test and the results were presented in Table (4). The  $R^2$  values (multiple correlation coefficients) closer to 1 denoted high agreement between the experimental and predicted responses and indicate that the mathematical model is very reliable in the present study. The coefficient of variation (CV) indicated the degree of accuracy in the comparison of experiments. A lower reliability of the experiment is usually indicated by high value of CV; in the present case the low value of CV (3.40) indicated that experiments conducted were precise and reliable. ANOVA

analysis for oily wastewater degradation by *Aspergillus niger* showed that the regression model was significant and the lack of fit was insignificant. The better the model would explain the variability between the experimental and the model predicted values. The graphical representations of the regression model, named the response surface plots and their corresponding contour plots were obtained by Design-Expert software and are obtainable in figure (4). Here, each response surface plot represented the effect of two independent variables, holding the other factors at zero levels. The shape of the corresponding contour plot showed whether or not the mutual interaction among the independent factors is significant figure (5). It was showed that A, B and C give high removal of oily wastewater when approaching to +1, on the other hand D give high removal of oily wastewater when approaching to -1. Zahed *et al.* [16] suggests that increasing phosphorus and nitrogen concentration can increase n-alkane removal. The optimized predicted removal was obtained at a phosphorus concentration of 20 mg/L at approximately 24 days. Due to dominating interaction effects of time and phosphorus, higher, levels of these variables increase biodegradation up to 23 days. Optimum levels of nutrient are both economically and ecologically important: high nutrient concentrations may cause eutrophication and harmful algal blooms (HABs) in the aquatic ecosystems [17]. In this study, result implies that it can be a good degrader in presence of significant nutritional and environmental parameter with in a very short period of time figure (6). The effect of the interaction of selected parameters on oily wastewater bioremediation by *Aspergillus niger* was investigated by plotting the response surface curves against any two independent variables while keeping the other independent variable at constant level. They provided data about the interaction between two factors and allowed an easy interpretation of the results and prediction of the best standards.

#### 4. Conclusion

This present study showed the ability of *Aspergillus niger* in degrading oily wastewater under different conditions. Optimum conditions that achieved in this research encourage using of this isolate in the remediation of high-strength of oily wastewater discharges. This study showed that the response surface methodology was suitable design to enhance the culture conditions for obtaining the maximum bioremediation of oily wastewater. This method may be modified for more removal of toxic environmental pollutants in different industrial aspects.

197 Table (1): List of different variables under study and their coded levels.

Factor	Factor Name	unit	Low Level (-1)	Medium Level (0)	High Level (+1)
A	Sucrose	g/L	0.1	2	4
B	MgSO <sub>4</sub>		0.1	0.5	0.9
C	NaCl		0.1	0.4	0.8
D	KH <sub>2</sub> PO <sub>4</sub>		0.2	0.6	1.2
E	pH		3	5	7
F	Temperature	°C	18	20	30
G	Urea	g/L	0.1	0.5	0.8
H	NH <sub>4</sub> Cl		0.1	0.4	0.8
I	Spore suspension	%	0.2	0.3	0.53
J	Na <sub>2</sub> HPO <sub>4</sub>	g/L	0.2	0.4	0.8
K	NaNO <sub>3</sub>		0.1	0.3	0.9

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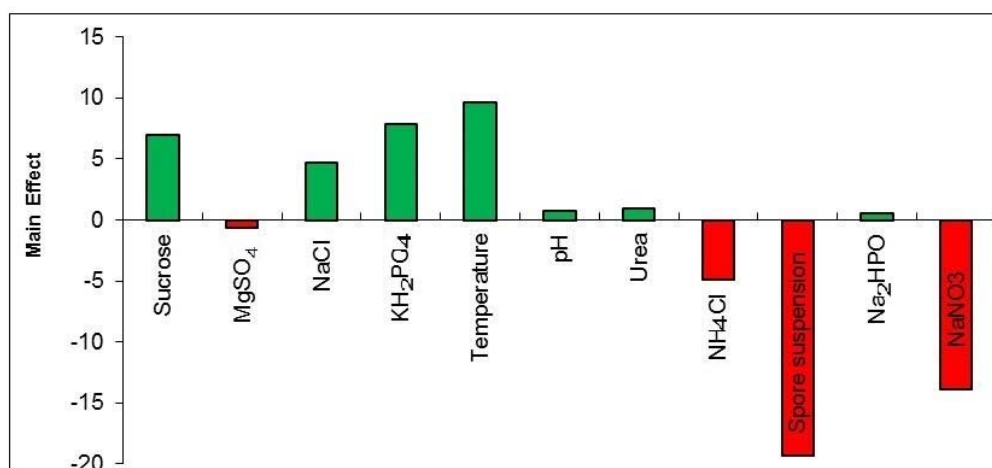
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200 **Table (2):** PB experimental design for evaluating the effect of different nutritional and  
 201 environmental categories on oil bioremediation.

Assay	Sucrose	MgSO <sub>4</sub>	NaCl	KH <sub>2</sub> PO <sub>4</sub>	Temperature	pH	Urea	NH <sub>4</sub> Cl	Spore suspension	Na <sub>2</sub> HPO <sub>4</sub>	NaNO <sub>3</sub>	Oil removal
1	+1	-1	1	-1	-1	-1	1	1	1	-1	1	7.69
2	+1	1	-1	+1	-1	-1	-1	1	+1	1	-1	39.69
3	-1	1	+1	-1	1	-1	-1	-1	+1	+1	+1	20.52
4	+1	-1	+1	+1	-1	1	-1	-1	-1	+1	1	72.64
5	+1	+1	-1	1	1	-1	1	-1	-1	-1	+1	80.51
6	+1	1	1	-1	+1	1	-1	+1	-1	-1	-1	91.8
7	-1	1	1	+1	-1	+1	+1	-1	+1	-1	-1	47.29

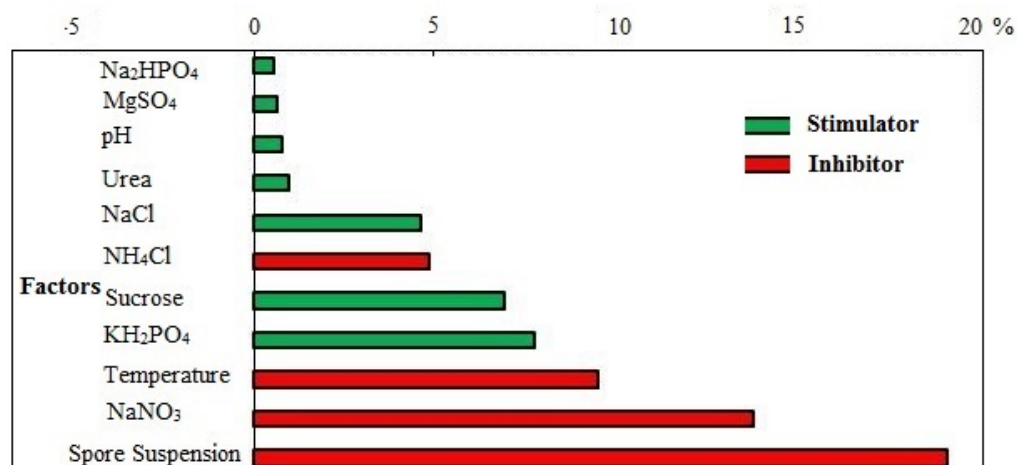
8	-1	-1	+1	1	+1	-1	+1	+1	-1	+1	-1	96.32
9	-1	-1	-1	+1	+1	1	-1	+1	+1	-1	1	18.88
10	+1	-1	-1	-1	1	1	+1	-1	1	1	-1	57.85
11	-1	+1	-1	-1	-1	+1	+1	1	-1	1	+1	24.38
12	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	58.82
13	0	0	0	0	0	0	0	0	0	0	0	58.82

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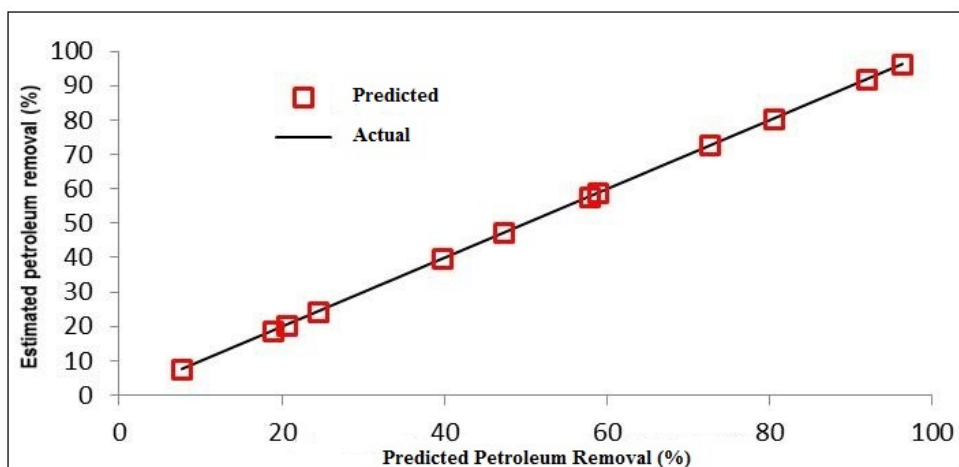
204 **Figure (1)** Effects of different culture conditions on diesel oil removal %.



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206 **Figure (2)** Pareto plot for PB parameter estimates.





**Figure (3):** The relationship between predicted and actual value.

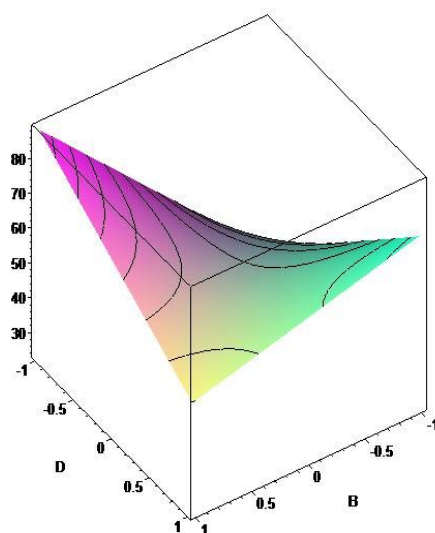
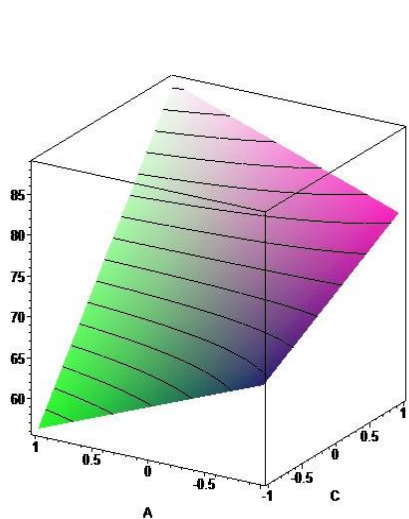
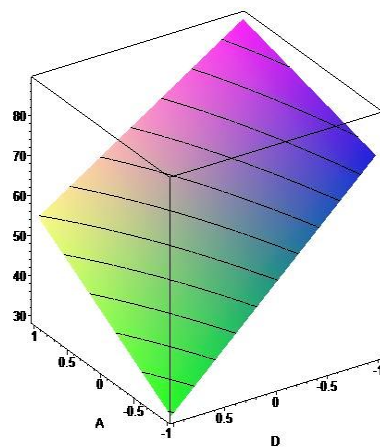
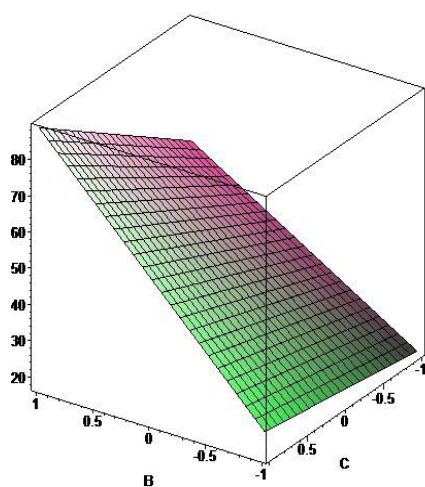
**Table (3):** Plackett-Burman design for optimization of selected variables.

RUN	Sucrose	NaCl	KH <sub>2</sub> PO <sub>4</sub>	Temperature
	A	B	C	D
1	-1	-1	-1	-1
2	-1	1	-1	1
3	1	1	1	1
4	1	1	-1	-1
5	1	1	-1	1
6	-1	1	1	-1
7	-1	-1	-1	1
8	-1	1	1	1
9	1	-1	1	-1
10	1	-1	-1	1
11	-1	-1	1	-1
12	1	1	1	-1
13	1	-1	1	1
14	-1	1	-1	-1
15	1	-1	-1	-1
16	-1	-1	1	1

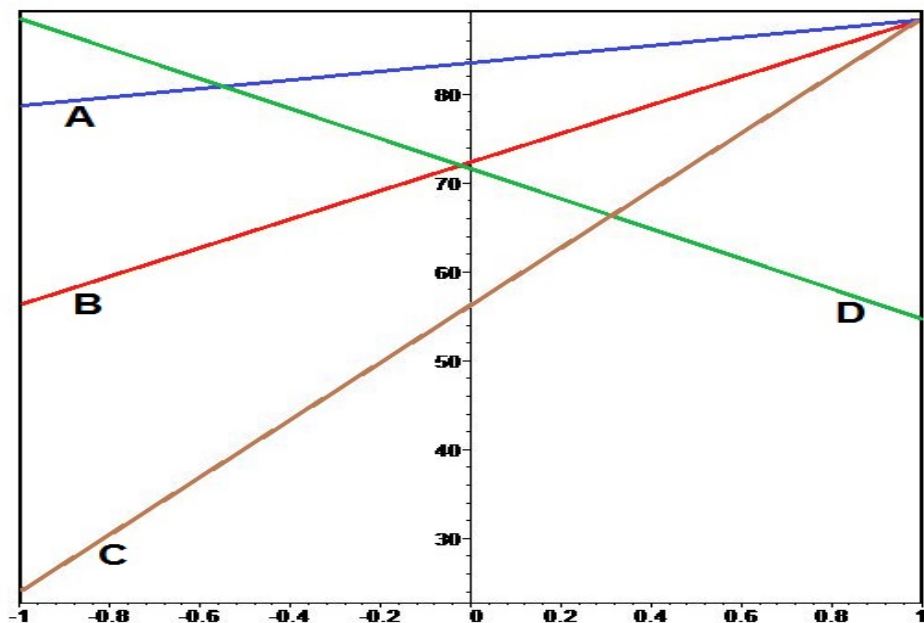
216 **Table (4):** Effects of the variables and statistical analysis of the Plackett-Burman design.

Source	Sum of Squares	df	Square	F- Value	p-value Prob > F
Model	6639.01	13.00	510.69	134.44	0.01
A-A	31.71	1.00	31.71	8.35	0.10
B-B	605.08	1.00	605.08	159.29	0.01
C-C	5.44	1.00	5.44	1.43	0.35
D-D	0.34	1.00	0.34	0.09	0.79
AB	515.33	1.00	515.33	135.66	0.01
AC	47.36	1.00	47.36	12.47	0.07
AD	2449.67	1.00	2449.67	644.89	0.00
BC	0.39	1.00	0.39	0.10	0.78
BD	1305.39	1.00	1305.39	343.65	0.00
CD	392.28	1.00	392.28	103.27	0.01
ABD	382.08	1.00	382.08	100.59	0.01
ACD	205.47	1.00	205.47	54.09	0.02
BCD	698.49	1.00	698.49	183.88	0.01
Residual	7.60	2.00	3.80		

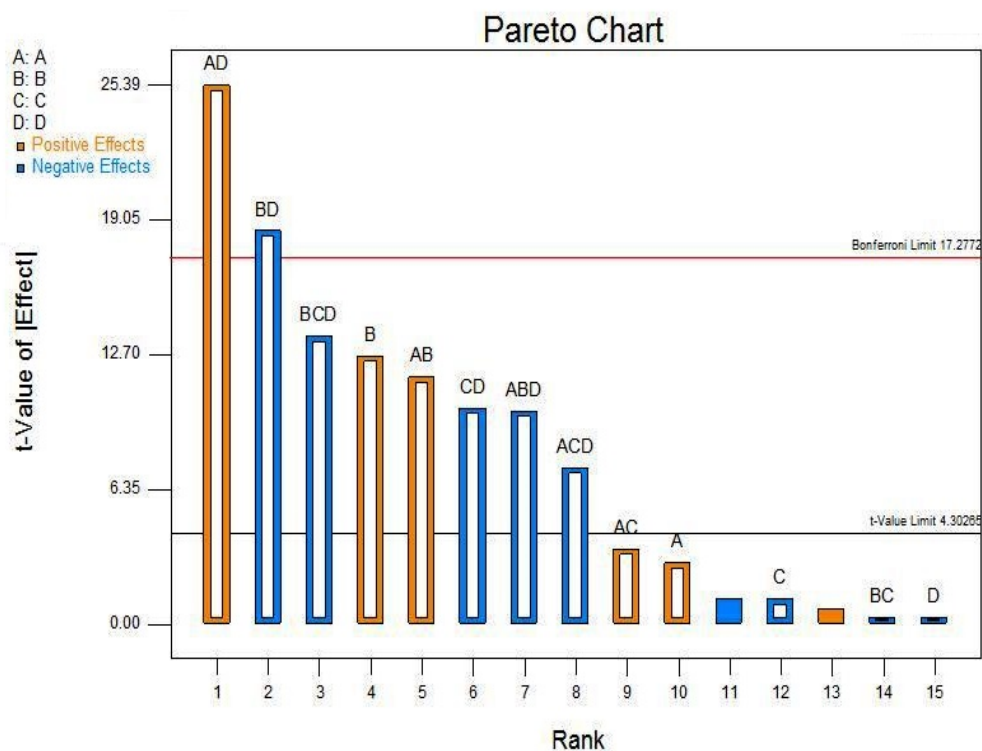
217 C.V. % -3.40;  $R^2$ :0.9268; adjusted  $R^2$ : 0.9914



**Figure (4):**Three-dimensional response surface plots and two-dimensional contour plots for degradation by *Aspergillus niger* showing variable interactions of: (A) Sucrose; (B) sodium chloride; (C)  $\text{KH}_2\text{PO}_4$  and (D) Temperature.



**Figure (5):** positive and negative effects of selected variables on oily wastewater removal.



**Figure (6):** Relationship between t-value of effect and various parameters.

231     **5. References**

- 232     **[1] Kanluen, R. and Amer, S. I. (2000):** A new treatment successfully removes contaminants  
233             from oily wastewater generated by aircraft maintenance operations. Environmental  
234             Protection. Aquachem Incorporation.
- 235     **[2] Al-Jumaily, E. and Al- wahab, N. (2012):** Nutritional requirement of *Enterobactercloacae*  
236             for biodegradation of hydrocarbons. Global. *J. Biol. Sci. Biotechnol.*,1: 65-70.
- 237     **[3] Boboye, B., Olukunle, O. F. and Adetuyi, F. C. (2010):** Degradative activity of bacteria  
238             isolated from hydrocarbon polluted site in Ilaje, Ondo State Nigeria. African Journal of  
239             Microbiology Research. 4(23): 2484-2491.
- 240     **[4] Vidali, M. (2005):** Bioremediation: An overview. Pure Applied Chemical. 7: 1162-1172.
- 241     **[5] Atlas, R. M. (1991):** Microbial hydrocarbon degradation-bioremediation of oil spills. J.  
242             Chem. Technol. Biotechnol., 52: 149-156.
- 243     **[6] Abd El- Hamid, H. T. (2015):** Detection and remediation of hydrocarbon wastes along the  
244             coastline from Damietta to Port Said. Ph.D. thesis, Faculty of Sci. Damietta, Univ.  
245             Egypt. pp.4.
- 246     **[7] Abdel-Fattah, Y. R.; Saeed, H. M.; Gohar, Y. M. and El-Baz, M. A. (2005):** Improved  
247             production of *Pseudomonas aeruginosa* uricase by optimization of process parameters  
248             through statistical experimental designs. *Process Biochemistry*, 40(5): 1707– 1714.
- 249     **[8] Nwachukwu, S. C. U. (2000):** Enhanced rehabilitation of tropical aquatic environment  
250             polluted with crude petroleum using *Candida utilis*. J. Environ. & Biol.,21: 241 – 250.
- 251     **[9] Plackett, R. L. and Burman, J. P. (1946):** The design of optimum multifactorial  
252             experiments. Biometrika, 33: 305-325.
- 253     **[10] American Public Health Association (APHA) (1998):** Standard methods for  
254             examination of water and wastewater, 20th edn., Washington D.C.
- 255     **[11] Strobel, R. J. and Sullivan, G. R. (1999):** Experimental design for improvement  
256             offermentations. In: Manual of industrial microbiology and biotechnology (Demain,  
257             A.L, Davies, J. E., eds.): 80-93. Washington: ASM Press.
- 258     **[12] Hegazy, T. A.; Ibrahim, M. S. and Abd El Hamid, H. T. (2015):** Placket Burman  
259             Design of Environmental and Nutritional Parameters for Petroleum Bioremediation by  
260             Pencillium Chrsogenum. *Scientfic Journal of Damietta Faculty of Science* 5: 2, 40-44.

- 261 [13] **Zhou, J.; Yu, X., Ding, C.; Wang, Z.; Zhou, Q.; Pao, H. and Cai, W. (2011):**  
262 Optimization of phenol degradation by *Candida tropicalis* Z-04 using Plackett-Burman  
263 design and response surface methodology. *Journal of Environmental Science and*  
264 *Health*, **231**, 22- 30.
- 265 [14] **Ghanem, K. M.; Al-Fassi, F. A. and Al-Hazmi, N. M. (2012):** Optimization of  
266 chloroxylenol degradation by *Aspergillus niger* using Plackett-Burman design and  
267 response surface methodology. *African Journal of Biotechnology*, **84**, 15040-15048.
- 268 [15] **Annur, M. S. M.; Tan, I. K. P.; Ibrahim, S. and Ramachandran, K. B. (2006):**  
269 Ammonium uptake and growth kinetics of *Pseudomonas putida* PGA1. Asia Pacific  
270 Journal of Molecular Biology and, Biotechnology, v. 14 (1), 1-10
- 271 [16] **Zahed, M. A.; Aziz, H. A.; Isa, M. H. and Mohajeri, L. (2010):** Enhancement  
272 Biodegradation of n-alkanes from Crude Oil Contaminated Seawater. *Int. J. Environ.*  
273 *Res.*, 4(4):655-664.
- 274 [17] **Tam, N. F. Y.; Wong, Y. S. and Wong, M. H. (2009):** Novel technology in pollutant  
275 removal at source and bioremediation. *Manage.* **52**, 368-373.  
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