# brought to you by 🌋 CORE

# **PREFACE**

In recent years, people have come to recognize that a healthy lifestyle can promote wellness and prevent illness and diseases, allowing them to enjoy long, high quality lives. People strive to maintain homeostasis and achieve their potential by meeting their individual physiological, safety, cultural and spiritual needs. The body's homeostatic balance is affected by diet. Consumption of massive amounts of sugar, salt, caffeine or fried foods can affect homeostatic balance.

Wellness is a state of health in which basic needs are being met and homeostasis maintained. Health problem can be any actual or potential concern or condition which must be resolved and prevented in order to maintain wellness. Unresolved problems will lead to the inability to meet basic needs and maintain homeostasis, eventually resulting in illness. Wellness technology can create healthy energetic environments that can optimize dynamic human system homeostasis targeting physical, mental, emotional and spiritual balances.

In an organization, wellness programs can be incorporated to improve the health and well-being of employees (and their families), in order to enhance organizational performance and reduce cost. Wellness programs typically address specific behaviours and health risk factors, such as poor nutrition, hypertension, coronary heart disease, obesity and smoking. These factors commonly lead to serious and expensive health problems and have negative impact on workplace productivity. While the return on investment (ROI) varies for each employer, studies in the United States have shown that for every USD1 an employer spends on wellness programs, employers can expect a USD3 to USD6 return on their investment. Wellness programs not only improve an individual's short-term and long-term health, but they also help curb absenteeism, improve productivity, and aid quicker return to work for employees on disability leave.









# ABSTRACTS FOR POSTER PRESENTATION

S2: BIOPROCESSING & AGRI-TECHNOLOGY

PS2-11

Production Of Erythromycin Antibiotic By Saccharoplyspora Erythraea Fermentation In Shake Flasks A **Bioreactor** 

Mohamud, M.A<sup>1</sup>, Abd Malek, R<sup>1</sup>, Mohamed, N.A<sup>2</sup>, Othman, Z<sup>1</sup>, Ramli, S<sup>1</sup>, Jalal, Y<sup>1</sup>, Aziz, R<sup>1</sup>, El Enshasy, H.A<sup>1</sup>, <sup>1</sup>Institute of Bioproduct Development (IBD), Universiti Teknologi Malaysia (UTM), Johor Bahru, Malaysia. <sup>2</sup>Natural and Microbial Products Department, National Research Centre, Dokki, Cairo, Egypt. <sup>3</sup>Bioprocess Development Department, City for Scientific Research and Technology Applications (CSAT), New B Al Arab, Alexandria, Egypt.

### Abstract

Recently success of erythromycin in antibiotic market over the other antibiotics was due to that erythromycin l high quality and it is cheap in price. Erythromycin received much attention because of the increasing application of its semi-synthetic modified derivatives to infection diseases, such as azithromycin, roxithromycin a clarithromycin. It is produced by the strain Saccharoplyspora erythraea (formerly known as Streptomyces erythrae In this research, the aims were to optimize medium components for high erythromycin antibiotic production by strain S. erythae via submerged fermentation using statistical technique known as response surface methodolo Glucose and yeast extract were found to have significant effect to erythromycin production using Placket-Burn experimental design for media screening. The Box-Benkhen experimental design was adopted for optimizat studied. Finally, the optimal concentration of glucose, yeast extract, sodium nitrate, dipotasium hydrod phosphate, sodium chloride and magnesium sulphate obtained using statistical media optimization approximately 45;8; 4; 2.5;1.0; 0.5 (g L-1), respectively. Result showed that the maximal erythromycin concentrat and CDW obtained in shake flasks of optimize medium were 412.5 mg L<sup>-1</sup> and 4.9 g L<sup>-1</sup>, respectively. Production erythromycin antibiotic reached 30.43% under the optimize medium. Furthermore, the batch culture using n medium formulation for erythromycin production was implemented using controlled and un-controlled conditions. Compared with the un-controlled pH bioreactor, the controlled bioreactor was increased erythromy concentration by 12.9 % up to 567.5 mg  $L^{-1}$ . This present work demonstrated that great potential production erythromycin antibiotic at industrial scale.

**Keywords:** *S. erythae*, Erythromycin, Medium optimization, Response surface methodology.

PS2-12

Probiotication Of Punica Granatum (Pomegranate) Juice By Lactobacillus Plantarum

Siti Marhaida Mustafa<sup>1</sup>, Lee Suan Chua\*, Hesham Ali El Enshasy<sup>1,2</sup>, Fadzilah Adibah Abdul Majid<sup>3</sup>, Roslin Abd Malek<sup>1</sup>

<sup>1</sup>Institute of Bioproduct Development (IBD), Universiti Teknologi Malaysia (UTM), Johor Bahru, Malaysia. <sup>2</sup>Department of Bioprocess Engineering, Faculty of Chemical Engineering, Universiti Teknologi Malaysia (UT Johor Bahru, Malaysia.

## Abstract

Fruit juice enriched with probiotics is increasingly accepted nowadays, mainly due to its health benefit for digest system. In particular, probioticated fruit juice is the good choice for those who are having lactose intoler problem from milk based drinks. In the present study, the whole fruit of *Punica granatum* (pomegranate) has be probioticated with Lactobacillus plantarum at different fermentation temperatures (22°C, 30°C and 35°C). growth rate of L. plantarum has been monitored based on the optical density and acidity of the broth culture at hours of time interval for 72 hours. The bacterial growth in the pomegranate juice was predicted by measur absorbance at 600 nm spectrophotometrically and pH value by a pH meter. There was an increasing trend in bacterial growth of L. plantarumincubated at 35°C compared to other temperatures at 22°C and 30°C. The resu also indicated there was no significant changes on pH during the fermentation as the bacterial strain was adaptation process with the new medium and conditions. Meanwhile, the antioxidant assay showed t probiotication of pomegranate juice by L. plantarum significantly increased the radical scavenging activity. pomegranate juice was shown to be a suitable substrate for L. plantarum cultivation at 35°C.

**Keywords:** Pomegranate juice; fermentation; *Lactobacillus plantarum*, antioxidant activity.