## EFFECTS OF SALT CONCENTRATION ON VAPOR-LIQUID EQUILIBRIUM (VLE) OF AZEOTROPIC MIXTURE IN ULTRASONIC DISTILLATION SYSTEM

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A thesis submitted as fulfillment of the award of the degree of Master of Engineering (Chemical)

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Specially dedicated to

my beloved father, Razali bin Ismail, my beloved mother, Rujomah bte Jamil

and those people who have guided and inspired me throughout my journey of education nt my jourr.

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#### ABSTRACT

Methanol and methyl acetate form an azeotrope in their mixtures at 34.78 mole % of methanol. It is difficult and may be impossible to separate azeotropic mixture using conventional distillation column. This phenomenon gives big challenges in the chemical industry in order to solve it. In this study, potassium chloride was added in ultrasonic distillation system to study the combination effect of salt and ultrasonic wave on methanol-methyl acetate mixture. The salt was added with different range of concentration to select the optimum concentration that can eliminate the azeotropic point. Ultrasonic wave with frequency of 25 kHz and intensity of 200 W/A.cm<sup>2</sup> were used. The studies on the effect of different salt concentration at 0 wt%, 5 wt%, 10 wt% and 15 wt% to VLE of binary mixtures were done at that frequency and intensity to obtain the best salt concentration. The results obtained show that, as the salt concentration increased in the liquid phase, the equilibrium line shifts upwards and in the same time, the azeotropic point also move upward. The salt concentration used in this work give the results in the following order 15 wt% >10 wt% > 5 wt% > 0 wt% where the azeotropes point form at 70 mole %, 54 mole %, 48 mole % and 38 mole % of methanol accordingly. As the result, the best concentration of the potassium chloride for the methanol-methyl acetate separation in this project was at 15 wt% of concentration. These results show that the combination of ultrasonic and salt as a separating agent gave positive results and have a potential to be apply for industry in the future.

#### ABSTRAK

Campuran metanol dan metil acetat membentuk azeotrop pada titik 34.78 mol % dari metanol. Sebatian azeotrop ini merupakan campuran yang agak sukar dipisahkan dan mungkin tidak boleh dipisahkan oleh penyulingan biasa. Hal ini membuatkan industri kimia pada masa kini menghadapi cabaran yang agak besar untuk mengatasi masalah ini. Kajian ini memperkenalkan kaedah baru dalam percubaan untuk memisahkan campuran azeotrop ini, dimana garam digunakan sebagai agen pemisahan dalam sistem penyulingan ultrabunyi. Garam yang digunakan adalah kalium klorida, dimana garam ini di campurkan ke dalam sebatian metanol-metil acetat, dalam julat kepekatan yang berbeza agar nilai optimum kepekatan garam dapat diperoleh. Frekuensi gelombang ultrabunyi yang digunakan adalah 25 kHz, manakala keamatan ultrabunyi yang dibekalkan adalah pada 200 W/A.cm<sup>2</sup>. Julat kepekatan garam yang digunakan adalah sebanyak 0 wt%, 5 wt%, 10 wt% dan 15 wt%. Keputusan yang diperoleh menunjukkan kesan yang agak baik apabila lengkungan keseimbangan metanol-metil acetat berganjak menjauhi garisan 45°C dan seterusnya melonjakkan nilai titik azeotrop. Hasil yang diperoleh adalah mengikut urutan 15 wt% >10 wt% > 5 wt% > 0 wt% dimana titik azeotrop yang diperoleh adalah pada 70 mol %, 54 mol %, 48 mol % dan 38 mol % dari metanol mengikut urutan. Kesimpulannya, kepekatan garam kalium klorida yang terbaik dalam projek ini adalah pada kepekatan 15 wt %. Keputusan ini menunjukkan kaedah yang diperkenalkan ini memberi hasil yang positif. Ini memungkinkan kaedah ini untuk diteruskan kajiannya, seterusnya diaplikasikan dalam industri pada masa hadapan.



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## LIST OF SYMBOLS AND ABBREVIATION

VLE	-	Vapor-Liquid Equilibrium
m	-	Meter
°C	-	Degree Celsius
ml	-	Mill liter
g	-	Gram
KCl	-	Potassium Chloride
%	-	Percentage

### **CHAPTER I**

#### INRODUCTION

### 1.1 Introduction



Distillation is the most widely applied separation technology and will continue as important process in the future because there is simply no industrially viable alternative around. Eventhough this technique confronted challenges from other technologies, it still improves from time to time and moves to a higher level of sophistication. Nevertheless, there are still many technical barriers faced by distillation. Not all liquid mixture can be separated by ordinary fractional distillation. The separation becomes difficult and expensive when the components of the system have low relative volatilities ( $1.00 < \alpha < 1.05$ ). This is because a large numbers of trays are required and, usually, a high reflux ratio as well. A different problem occurs if the system forms azeotropes, where the azeotropic composition limits the separation (Cheresources.com website, 2008).

Azeotrope means literally that the vapor boiling from a liquid has the same composition as the liquid. The azeotropic mixture depends upon the degree of non-ideality of a mixture and the difference in boiling points between the two pure components (Kim et al., 1997).

It has been known that the separation of components by simple distillation is impossible if the mixture exhibits an azeotropic at a specified temperature and pressure. For a mixture that has an azeotrope, the separation process relies on the addition of specially chosen chemicals to eliminate the azeotropes entirely. Salts are one of the mass-separating agents that can be introduced in distillation system in order to solve this problem (Banat et al., 1997).

A salt dissolved in a mixture of volatile components may affect the activities of the components through the formation of complexes (Yao et al., 1999). The salt will dissociates into ions in the liquid mixture and alters the relative volatilities to make the separation becomes possible (Banat et al., 1997).

Normally, even little salt may bring an appreciable effect on the relative volatility where this is called the effect of preferential salvation. The use of a salt instead of another separating agent in distillation gives several advantages including lower energy consumption. It also gives high purity of the overhead products because salts are non-volatile and hence do not evaporate or condense during distillation process (Banat et al., 1997). Salts also has lower toxicity lever comparing with other liquid separation agents such as benzene.

The previous research has verified the feasibility of using ultrasonic wave to enhance the separation of binary mixtures in distillation column and overcoming the challenges of azeotropic separation. In this project, salt was used to be tested in ultrasonic distillation system. The salt was introduced in the flask with different concentration. Once the optimum salt concentration was achieved, further analysis can be done by calculate its relative volatility.

This research was about to see the effect of salts in ultrasonic distillation system in order to enhance the separation process. The selected binary mixture was methanol and methyl acetate, while the salt used was potassium chloride (KCl).

#### 1.2 **Problem statement**

Distillation is the common separating method to separate components in liquid mixtures. However, this process may be complicated by the formation of azeotropes due to non idealities in the mixture. These azeotropes can make a given separation impossible by conventional distillation processes. Industries always looking forward to obtain the solution for this problem.

Salt is one of the separating agents in distillation process for separating closeboiling or azeotropes systems that cannot easily be purified using ordinary distillation. Salt has been proved for eliminating the azeotropes entirely. In this work, vapor-liquid equilibrium (VLE) studies were conducted to determine the optimum concentration of selected salt in ultrasonic distillation system.

#### 1.3 Objective

NKU TUN AMINA The main objective of this research is to study the separation of azeotropic mixture by using salt on VLE in ultrasonic distillation system.

#### 1.4 Scope of Research

The scope of this research was to identify the appropriate salt concentration to be applied in ultrasonic distillation system.

### **CHAPTER II**

### LITERATURE REVIEW

### 2.1 Distillation

Distillation is a method of separating mixtures where a liquid or vapor mixture of two or more substances is separated into its component fractions of desired purity, by the application and removal of heat. The separation of mixture is based on differences in their volatilities in a boiling liquid mixture (Fair, 2000). Distillation is the most common separation technique and it consumes enormous amounts of energy, both in terms of cooling and heating requirements. It can contribute to more than 50% of plant operating costs. The best way to reduce operating costs of existing unit is to improve their efficiency and operation via process optimization and control (Distillation website, 2008).

Distillation exists either in batch or continuous mode. In batch distillation, the composition of the source material, the vapors of the distilling compounds and the distillate change during the distillation. In batch distillation, a still is charged (supplied) with a batch of feed mixture, which is then separated into its component fractions which are collected sequentially from most volatile to less volatile, with the bottoms (remaining least or non-volatile fraction) removed at the end. The still can then be recharged and the process repeated. In continuous distillation, the source materials, vapors, and distillate are kept at a constant composition by carefully adding the source material and removing fractions from both vapor and liquid in the system. This results in a better control of the separation process (Wikipedia website, 2008).

#### 2.2 Vapour-Liquid-Equilibrium (VLE)

#### Introduction 2.2.1



However, when automation expertise has been successfully created in the laboratory, the goal should be to implement data acquisition programs and automation software to increase the measurement output of the experimental devices. It is thereby possible to decrease the cost of one individual measurement point substantially. Suitable methods for determination of VLE vary. In some cases several methods can be applied, but in the most difficult cases measurements are almost

impossible. The selection of methods and apparatuses depend on the physical properties of the system studied such as vapor pressure, component stability, material compatibility, measurement accuracy and safety. The properties determined specifically for binary vapor liquid equilibrium systems are temperature, pressure and the compositions of the constituent phases (Ussi-Kyyny, 2004).

The determination of composition is the most complex task. The devices needed are often expensive and there is no universal analytical device that is suitable for all components. Gas chromatography is used most often for the determination of the composition of phases. Other methods for composition determination, although seldom applied in VLE measurements, include mass spectrometry, various spectroscopic methods, and density and refractive measurement (Ussi-Kyyny, 2004).

### 2.2.2 Vapour-Liquid-Equilibrium (VLE) Curves

Constant pressure VLE data is obtained from boiling point diagrams. Figure 2.1 shows the plot that often presented for VLE data of binary mixtures. The VLE plot expresses the bubble-point and the dew-point of a binary mixture at constant pressure. The curved line is called the equilibrium line and describes the compositions of the liquid and vapor in equilibrium at some fixed pressure. This particular VLE plot shows a binary mixture that has a uniform vapor-liquid equilibrium that is relatively easy to separate (Distillation website, 2008).



Figure 2.1: VLE graph of binary mixtures

## 2.3 Azeotropic Formation in Binary Mixtures



The most intriguing VLE curves are generated by azeotropic systems. An azeotrope is a liquid mixture which when vaporised, produces the same composition as the liquid. Figure 2.2 shows two different azeotropic systems, one with a minimum boiling point and one with a maximum boiling point. In both plots, the equilibrium curves cross the diagonal lines, and this are azeotropic points where the azeotropes occur. In other words azeotropic systems give rise to VLE plots where the equilibrium curves crosses the diagonals (Distillation website, 2008).



Figure 2.2: Two different azeotropic systems (homogenous azeotrope)

Note the shapes of the respective equilibrium lines in relation to the diagonal lines that bisect the VLE plots. Both plots are however, obtained from homogenous azeotropic systems. Homogenous azeotrope is an azeotrope that contains one liquid phase in contact with vapour. A homogenous azeotrope cannot be separated by conventional distillation. However, vacumn distillation may be used as the lower pressures can shift the azeotropic point. Figure 2.3 shows the VLE curve that also generated by an azeotropic system, in this case a heterogenous azeotrope. Heterogenous azeotropes can be identified by the flat portion on the equilibrium diagram. They may be separated in 2 distillation columns since these substances usually form two liquid phases with widely differing compositions. The phases may be separated using settling tanks under appropriate conditions (Distillation website, 2008).



Figure 2.3: Heterogenous azeotrope

#### Physical Phenomenon Leading to Azeotropy 2.3.1

The tendency of mixture to form azeotrope depends on two factors, which are: (i) the difference in the pure component boiling points, and (ii) the degree of non-ideality (Kim and Simmrock, 1997). The closer the boiling points of the pure components and the less-ideal mixture, the greater the likelihood of an azeotrope The mixtures that have wide differences in their components boiling point may not exhibit an azeotrope even though they form a nonideal mixture.

Most mixtures of organic compounds form nonideal systems. The presence of some specific groups, particularly polar groups such as oxygen, nitrogen, chlorine, and fluorine often results in the formation of azeotropes. The nonideality of mixture depends largely on intermolecular forces of attraction among the mixture AKAAN TUNKU TUN AMINA components such as dispersion forces, dipole-dipole interactions, dipole-induced dipole interactions, and hydrogen bonding (Swietoslawski, 1963).

#### Ultrasound 2.4

#### 2.4.1**About Ultrasound**

Ultrasonic waves are mechanical pressure waves formed by actuating the ultrasonic transducers with high frequency, high voltage current generated by electronic oscillators. A typical industrial high power generator produces ultrasonic frequencies ranging from 20 - 120 kHz. Typical transducers are normally immersed in the liquid. The generated ultrasonic waves propagate perpendicularly to the resonating surface (Awad, 1996).

The waves interact with liquid media to generate cavitation implosions. High intensity ultrasonic waves create micro vapor bubbles in the liquid medium, which grow to maximum sizes proportional to the applied ultrasonic frequency and then implode, releasing their energies. The higher the frequency, the smaller the cavitation size. The high intensity ultrasonics can also grow cavities to a maximum in the course of a single cycle (Awad, 1996).

#### 2.4.2 Cavitation

Cavitation is the formation and collapse of either gas or vapor bubbles in a liquid subjected to pressure changes. The formation of cavities in liquids is analogous to tensile failure in solids. When the "tensile strength" of a liquid is exceeded, cavities form. Actual values of these "strengths" are much lower than theoretical values, as a result of imperfections (gas pockets) in the liquid which serve as nuclei for cavitation. These nuclei grow, through net diffusion of dissolved gas from the liquid to the nuclei, to form cavitational bubbles. When a high enough pressure amplitude (cavitation threshold) is reached, the nucleus becomes unstable and rapidly grows into a mostly vapor-filled bubble or transient cavity (Busnaina et al., 1994).

#### 2.5 Salts

An additional substance may add to shift the azeotropic point to a more favourable position (Distillation website, 2008). Salt is one of the additional substances that can be added to enhance the separation process. The salt acts as a separating agent by raising the relative volatility of the mixture and by breaking any azeotropes that may otherwise form.

The addition of a salt, instead of a liquid, as a separating agent in distillation provides several advantages including lower energy consumption and high purity of the overhead products because salts are nonvolatile and hence do not evaporate or condense during distillation. The salt effect is believed to be a complex function of salt and solvent interaction and self-interaction among all other components. In most cases, but not necessarily always, the molecules of the more polar solvent, in which the salt is more soluble, are preferentially attracted by the electrostatic field of the salt ions and thereby the molecules of the less polar solvent are liberated to the vapor phase (salted-out) (Banat et al., 1997). The list below show the general advantages of using salt in distillation column:

- Allows continuous operation because of the high efficiency and the low waste of solvent.
- A high purity product can be obtained.
- The relative volatility of mixture is increased, makes the separation processes easier.
- Improves the solvent performance. Compared with normal extractive distillation, the quantity of the solvent to recycle is reduced to its fourth or fifth, the number of theoretical stages required can be reduced to its third, as well as energy consumption.

### **CHAPTER III**

#### METHODOLOGY

### 3.1 Introduction

This chapter discussed about the research methodology that applied in this study. A set of distillation apparatus equipped with ultrasonic wave generating equipment was set up as the experimental rig to obtain the vapor-liquid equilibrium data. Only one type of binary mixture was studied in this research. The ultrasonic wave generating equipments, which were ultrasonic transducers and ultrasonic generators, were supplied by Crest Ultrasonic (M) Sdn. Bhd. based in Penang.

#### REFERENCES

Awad, S. B. (1996). "Ultrasonic Cavitations and Precision Cleaning". The Magazine of Critical Cleaning Technology. 12-17.

Banat, F., Al-Asheh, S. and Simandl, J. (2003). "Vapor-Liquid Equilibria of
Propionic Acid-Water System in The Presence of Different Types of Inorganic Salts:
Effect of Temperature and Salt Concentration". Chemical Engineering and
Processing. 42:917-924.

Busnaina, A. A., Gale, G. W. and Kashkoush, I. I. (1994). "Ultrasonic and Megasonic Theory and Experimentation". The Magazine of Critical Cleaning Technology. 13-19.

Cai, J., Yang, J., Du, Y., Fan, L., Qiu, Y., Li, J. and Kennedy, J., F. (2005). "Enzymatic Preparation of Chitosan From Waste Aspargillus Niger Mucelium of Citric Acid Production Plant." Carbohydrate Polymers. 1-7.

Chemicalland21.com website. http://chemicalland21.com/industrialchem/solalc/METHYL%20ACETATE.htm. Accessed on 10 October 2008.

Cheresources.com website. http://www.cheresources.com.my. Accessed on 9 August 2008.

Crest Ultrasonic Product Bulletin (2004). Tanks, Generators, Immersible Transducer and Push Pull. Penang, Malaysia: Crest Ultrasonics.

Distillation website. http://lorien.ncl.ac.uk/ming/distil/distiltyp.htm. Accesed on 9 August 2008.

Fair, J.R. (2000). Distillation: The Engineering Handbook. Boca Raton: CRC Press LLC.

Geankoplis, C.J. (1993). Transport Processes and Unit Operations. 3<sup>rd</sup> ed. Englewood Cliffs, N. J.: Prentice-Hall.

Kim, Y. J. and Simmrock, K. H. (1997). "Azeopert: An Expert System for The Prediction of Azeotrope Formation-I. Binary Azeotropes." Pergamon. 21:93-111.

Modul Amali Makmal Unit Operasi 2 (2004). Keseimbangan Wap-Cecair. Skudai, Johor: Chemical Engineering Department, UTM. Unpublished.

Shinskey, F.G. (1984). Distillation Control for Productivity and Energy Consevation. New York: McGrawHill Book Company.

Swietoslawski, W. (1963). Azeotropy and Polyazeotropy. Oxford: Pergamon Press. Thornycroft, J. and Barnaby, S. (1985). Proc. Inst. Civ. Engin. 122: 51.

The Suslick Research Group website. http://www.scs.uiuc.edu/~suslick/britannica.html. Accessed on 14 March 2008.

Ussi-Kyyny, P. (2004). "Vapour Liquid Equilibrium Measurements For Process Design". Chemical Engineering Report Series.

Wikipedia website. http://en.wikipedia.org/wiki/distillation. Accessed on 14 March 2008.

Yao, J., Li, H. and Han, S. (1999). "Vapor-Liquid Equilibrium Data for Methanol-Water-NaCl at 45 °C." Fluid Phase Equilibria. 163:253-260.