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Analysis the Efficiency and Productivity of Indonesian Pharmaceutical Public Companies Using Data Envelopment Analysis

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As one of the biggest medicines market in the South East Asia, the pharmaceutical industry in Indonesia has a huge potential market. However, the majority supply of raw materials has been imported. Besides, regulations of the Health Ministry and the Trade Ministry have still hampered most players in Indonesia pharmaceutical industry. Therefore, this study used Data Envelopment Analysis (DEA) models to analyze efficiency and productivity change in the Indonesian pharmaceutical industry between 2006 and 2011, listed in the Indonesia Stock Exchange and also supported by applying efficiency financial ratio. This study finds that the decision for the most relatively efficient company is different using DEA compared to efficiency financial ratios, yet DEA has better measurement of efficiency. It is proven by one of State-owned Enterprises has been evaluated underperformed by the financial ratio analysis, unexpectedly is efficient using the DEA approach. This study has also proposed and tested a hypothesis on the average efficiency to check if the domestic and foreign pharmaceutical companies differ in their efficiency but the result implies that there is no significant statistical difference among them. This study indicates that firms having dominant contribution in selling over-the-counter medicines are more efficient than selling ethical medicines. Lastly, technological change contribution has more influence to productivity change instead of pure technical efficiency change in Indonesia pharmaceutical companies.

Keywords: *efficiency, productivity, pharmaceutical companies, Data Envelopment Analysis (DEA), financial ratio*

Introduction

Indonesia is one of the biggest potential markets for pharmaceutical products in the South East Asian region. Indonesia accounts for 37% out of the 650 million populations living in the region¹. Most of the market share for pharmaceutical products, up to 70%, is still held by local pharmaceutical companies with a total value of around USD2.4 billion². Profit growth opportunities are also promising due to

the population size and the rising annual population growth. However, these growth opportunities may be offset by various challenges.

Indonesia imports more than 95% of raw materials from India, China and Europe³. This situation creates vulnerability to the Indonesian pharmaceutical industry, since the cost of raw materials becomes highly dependent to the currency value of IDR compared to the currency of the supplier's country. Other challenges come from local regulations considered as constrain-

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¹ <http://www.businessnews.co.id/ekonomi-bisnis/industri>

² <http://www.businessreview.co.id/bisnis-investasi>

³ <http://www.businessnews.co.id/ekonomi-bisnis/industri-farmasi>

ing to foreign pharmaceutical players under the International Pharmaceutical Manufacturers Group⁴, especially the Ministry of Health Regulation (*Permenkes*) No.1010/2008 concerning Medicine Registration. The regulation implies that foreign pharmaceutical companies without local manufacturing plant in Indonesia may not import pharmaceutical products to Indonesia. On the other hand, 95% raw materials for Indonesian pharmaceutical industry are still imported. Moreover, foreign owned companies usually import their raw materials from their home country and then formulated the final product in the host country (Saranga and Phani, 2008). The regulation was further strengthened by revisions of the definition of pharmaceutical industry in the Decree of Health Minister (*SK Menkes*) No 245/1990 which stated that pharmaceutical companies operating in Indonesia at minimum must own packaging facilities located in Indonesia, reducing the requirement from owning a production facility to only a packaging facility. Other regulation comes from the Ministry of Trade Regulation (*Permendag*) No.45/2009 regulating Importer Identifying number and limitation of foreign ownership up to 75% stated in the regulation of Negative Investment List. These regulations become a disincentive for foreign investors to invest their funds and the development of foreign owned pharmaceutical company in Indonesia.

Despite pharmaceutical industry in Indonesia facing many challenges, in fact its market share is still very attractive and promising. Thus, to improve competitiveness in order to expand market share, pharmaceutical companies are required to be more productive and efficient in running the company operations.

Based on the problems stated above, this paper aims to investigate the efficiency and productivity of publicly owned pharmaceutical company in Indonesia. Several researchers have studied efficiency and productivity of pharmaceutical industries, most of which are conducted in Europe, Japan and India (Fare et al., 1995; Gonzales and Gascon, 2004; Danzon et al., 2005; Hashimoto et al., 2008; Saranga and Phani, 2008; Mazumdar and Rajeev, 2009;

and Pannu et al., 2011). Conversely, the efficiency and productivity of pharmaceutical industries in the South East Asian region -such as Indonesia- are rarely studied. Therefore, this study tries to expand the body of research conducted previously in Europe, Japan and India, by conducting the study in Indonesia. This study also compares relative efficiency measured with DEA with efficiency calculated using efficiency financial ratios. Furthermore, this study tests whether difference of means exists between the efficiencies of local and foreign owned pharmaceutical companies, as well as identifying changes in productivities and the dominant determining factors.

Review of Related Studies

Most of the previous studies regarding the pharmaceutical industries are motivated by various constraining regulations in relation of the pharmaceutical industries themselves (Gombola and Ketz, 1983; Lothgren and Tambour, 1998; Barnum et.al., 2001; Gonzales and Gascon, 2004; Saranga and Phani, 2008; Mazumdar and Rajeev, 2009; Xu and Wang, 2009; Lin et al, 2011; Pannu et al., 2011; Maricica and Georgeta, 2012; Patel and Pande, 2012; Delen, Kuzey and Uyar, 2013). Saranga and Phani (2008) discovered that positive relations exist between internal efficiency and corporate growth in the dynamic environment of Indian pharmaceutical industries. The dynamic environment was influenced by the changes in regulation regarding foreign ownership of pharmaceutical companies from a maximum of 40% to 74% and also of Intellectual Property Protection (IPI). This finding is further strengthened by the study of Pannu et al. (2011) which discovered that a pharmaceutical corporation with higher efficiency enjoys higher compounded average growth rate (CAGR) of their sales.

Mazumdar and Rajeev (2009) studied factors influencing the efficiency of pharmaceutical corporations, such as corporate size, existence of R&D unit, target market segment and existing products. The study concludes that large pharmaceutical corporation has greater

⁴ <http://www.businessreview.co.id/bisnis-investasi>

efficiency due to the profits from invested technological innovations. They also find that the sector experienced rapid technological progress in recent years. Leading to a rise in the inefficiency for this sector, the most firms have failed to appropriate the benefits of the technological change. Their other finding mentioned that in the context of the Indian pharmaceutical industry, an extremely effective strategy for firms' benefit from efficiency and technological gains can be implemented by merging vertically with downstream raw material.

Conversely, the existence of R&D unit was found to have no significant effect to corporate efficiency. However, firms that invested in R&D profited from technological innovation. This suggests that the few frontier firms profited most from their R&D related outlays, but this investment enlarged the distance between the inefficient and efficient firms. Large firms that invested more in R&D have profited from technological growth. Thus, firms can expand their production possibilities, enabling themselves to realize higher margins by increasing the scale of operation backed by sufficient R&D activity. This finding supports earlier study by Hashimoto and Hanaeda (2008) in Japanese pharmaceutical companies which assessed that the existence of R&D unit is meant to increase company value more rather than increase corporate efficiency. The firms have continued to increase R&D expenditure every year although R&D efficiency has not improved. Possibly, firms might have found another meaning of R&D expense than R&D itself. However, it is certain that there has been the shortage of firms' R&D efficiency evaluation.

Alternatively, Pannu et al. (2011) proved that the existence of R&D unit can increase efficiency. Pannu found that the efficiency and productivity change leaders and laggards over 10 year period. Pannu also proved that the efficiency of local companies is lower than the average efficiency of MNCs between 1998 and 2007. However, there is no difference between multinational and local companies in the average productivity change between 1998 and 2007. According to this paper, close to 70% of local companies have average efficiency scores

less than the worst efficiency score among MNCs. This basically shows that the input–output balance among multinational companies is better than that of Indian companies. They also noticed that the inefficient companies are lower than the CAGR of efficient companies. Also, they have analysed the effect of firm size and found that compared to small and micro firms, larger firms posted a higher CAGR, efficiency and relative efficiency. However, smaller firms led the larger firms in productivity change and technical change. Their finding shows that, though it is declining for both the groups after 2004 for the ten year period, efficiency of innovative firms was higher than non-innovative firms on a year to year basis. While neither global nor domestic market orientation influences efficiency, local regulation significantly influences them. Finally, corporations which produce bulk of medicines and formulation medicine were found to have greater efficiency so that integrated production strategy would lead to greater efficiency in pharmaceutical companies. This finding supports earlier finding by Saranga and Phani (2008). They have seen that the internal efficiency ratings are not dictated by size of a company. Excepting for a few instances where companies which were in the mode of expansion, the methodology establish the fact that there is a direct relationship between internal efficiencies and higher growth rates. Those instances may not have achieved full efficiencies. Internal efficiencies have been established by the local companies in bulk medicine and formulation businesses, which have helped them to arise so far, will also play a principle role in grabbing the new opportunities of product patent era, to survive and arise in future.

Research on productivity of pharmaceutical companies was pioneered by Fare et al. (1995) by studying pharmaceutical companies in Sweden. Fare et al. (1995) classified changes in total productivity factors into technical efficiency change, technological change and quality change. The study discovered that differences exist between productivity measures when quality was entered as an attribute. On average, pharmaceutical companies experience an in-

crease on productivity, technological changes and efficiency changes when quality was entered in the equation.

Proposing a modification of the DEA network model by Fare and Grosskopf (1996), Lothgren and Tambour (1998) allow a representation of both production and consumption technologies in a unified framework. The inputs directed to customer activities, together with the production quality attributes is presumed to give customer satisfaction, here represented by assessments of quality from barometer surveys of customer satisfaction. The results obtained from the network model and non-network productivity model are similar in some acknowledgement. In both models the estimated productivity change is positive. However, the network model indicates a lower estimated productivity progress than the non-network model. The progress in productivity is indicated by both the efficiency and technical change components in the Malmquist index decomposition. The accounts of technical change component for most of the progress, although this is more clear in the non-network model than in the network model. However, there are some differences regarding which pharmacies are experiencing progress and regress in productivity in the different models. Taking customer satisfaction into account, the network model proposed by Lothgren and Tambour (1991) was constructed with the purpose to estimate efficiency and productivity for the sample of pharmacies. However, it should be noted that the proposed model can be applied to arbitrary industries as long as the data include information on customer satisfaction, production quality attributes and firm-specific input allocation to customer and production oriented activities. It means that the proposed model is not limited to the specific pharmacy application.

Gonzales and Gascon (2004) examined the productivity of pharmaceutical laboratories after the *deficit cero* policy in Spain which caused many pharmaceutical companies to reduce their profit margin. They discovered that pure technical efficiency change and scale efficiency change are the two dominant factors influencing productivity growth. Similar finding was

obtained in the Indian pharmaceutical industry under the amended Medicines and Cosmetic Act (Mazumdar and Rajeev, 2009). It also found that technological innovation applied by the Indian companies does not increase productivity growth.

Focusing the study on performance evaluation of pharmacy retail stores, Patel and Pande (2012) found that retail stores are operating at average efficiency levels and there is a lot of improvements scope. The stores performing inefficiently need to focus upon their footfalls, sales, operating expenses and store size. Experiencing increasing returns to scale is a well planned strategy that must be practiced by the inefficient stores. The most suitable strategy for the stores experiencing decreasing returns to scale, however, is to lower down their scale of operations. Patel and Pande conclude that the study provides a framework for performance evaluation of pharmacy retail stores along with identification of efficient drivers of performance

Suggesting DEA model in hospital pharmacies, Barnum et al. (2001) developed a study of integrated procedures that can both mitigate the problems and take advantage of the frequent and detailed information now available. *First*, they suggest a set of compatible and comprehensive measure of distributional and clinical output, some of which are only available because of the computerized information system. As a result, all of the key distributional and clinical outputs are covered. *Second*, they suggest DEA models that can separately summarize the efficiency of the pharmacies' clinical and distributional activities; with a third model that appraises the pharmacies' comprehensive efficiency as a whole. *Third*, it suggests a methodology for developing statistical control charts through the use of statistical PDA models, so pharmacy managers can quickly and accurately differentiate when conditions are indeed out of control and when the reports merely reflect random variations. They believe that the collective use of these suggestions could provide significant assistance to hospital pharmacies seeking to increase and maintain their efficiency.

In the proposed prediction scheme, the efficiency of corporation is obtained through DEA.

In that study, Xu and Wang (2009) explained on their paper. They propose a prediction approach using efficiency of corporation as a predictor. The efficiency and some selected financial ratios are then used as predictors in prediction methods. The numerical results of application to corporations in Shanghai stock exchange validate the efficacy of the proposed prediction scheme.

Lin, Liang and Chen (2011) study a set of financial features that includes 21 commonly used financial ratios that are expected to achieve a more accurate prediction of corporate financial distress than a model based exclusively on scholars' survey results from experts. In a similar case, identifying and using a large and feature rich dataset was studied by Delen, Kuzey and Uyar (2013). After a long search they identified FINNET, a company providing variety of financial data, software, and Web-based analysis tools to their members. They covered the time period of 2005 to 2011 for the final dataset of financial ratios. They find that profitability ratios can predict financial distress well. These ratios indicate the potential ability of a company in case of controlling costs and expenses. The more successfully the firm can control its costs and expenses, and by improving its performance (represented as ROE and ROA), the higher these ratios. Further, the Leverage and Debt Ratios are also found to be able to predict financial distress as well.

Either classic or modern bankruptcy prediction model, are principally based on financial ratios. The study was done by Maricica and Georgeta (2012). They argue about the importance of the information quality provided by financial ratios in bankruptcy prediction because it impacts on the accuracy of the bankruptcy prediction. Surely this can be used to measure how efficiently a resource that can be used by the company, which in this study is the financial resources.

Methodology

Efficiency measures were calculated using non parametric approach with *Data Envelopment Analysis* (DEA) method introduced by

Charnes et al. (1978). Efficiency measures using DEA involves a minimum of one output and three inputs (Pannu et al., 2011), in which sales is the only output entered while the inputs entered are 1) Cost of Material, 2) Cost of Manpower, and 3) Capital Cost. The last input, capital cost, was calculated from the total cost of sales and production minus the cost of raw material and worker's compensation.

In calculating efficiency using panel data, efficiency measurement was conducted using *DEA-like linear programs* while the *Malmquist Total Factor Productivity (TFP) Index* was employed to measure *productivity change* as well as to decompose *productivity change* into *technological change* and *technical efficiency change*.

Caves et al. (1982) showed that productivity changes can be assessed by Malmquist indices, defined in terms of ratios of distance functions. Following Fare et al. (1995) an input-based Malmquist productivity index for the P-Node can be defined (and decomposed) as:

$$M_i^{P,t,t+1}(\cdot) = (E_i^{P,t,t+1})(TC_i^{P,t,t+1}) \quad 1)$$

$$M_i^{P,t,t+1}(\cdot) = \frac{D_i^{P,t+1}(x^{P,t+1}, y^{t+1})}{D_i^{P,t}(x^{P,t}, y^t)} \times \left[\frac{D_i^{P,t+1}(x^{P,t}, y^t)}{D_i^{P,t+1}(x^{P,t+1}, y^{t+1})} \right]^{1/2} \quad 2)$$

where $D_i^{P,t}(x^{P,t+1}, y^{t+1})$ is the distance function which measures the maximum proportional change in outputs required to make (x^{t+1}, y^{t+1}) feasible in relation to the technology at time t .

Productivity improvement is signaled by a Malmquist index less than one and a negative change in productivity is signaled by an index greater than one. The term outside the brackets, E_i , measures change in efficiency and can be interpreted as a "catching up" (to the production frontier) effect. The TC_i term measures change in technology in terms of shifts in the production frontier.

The *Malmquist Productivity change index* was specified based on Fare et al. (1994) as follows:

$$m_0(y_{t+1}, x_{t+1}, y_t, x_t) = \left[\frac{d_0^t(x_{t+1}, y_{t+1})}{d_0^t(x_t, y_t)} \times \frac{d_0^{t+1}(x_{t+1}, y_{t+1})}{d_0^{t+1}(x_t, y_t)} \right]^{1/2} \quad 2.1)$$

The value (x_{t+1}, y_{t+1}) is the value of productivity relative to (x_t, y_t) . The value of productivity greater than one indicates the existence of positive TFP growth from period t to period $t+1$. This index was derived from four *linear programming functions*, as follows:

$$\begin{aligned} & \max_{\phi, \lambda} \phi, \\ \text{st} \quad & -\phi + Y\lambda \geq 0, \\ & x_t - X\lambda \geq 0, \\ & N1'\lambda = 1 \\ & \lambda \geq 0 \end{aligned} \quad (2.2)$$

Function 2.2 above is the function of *Variable Return to Scale* (VRS). VRS is the assumptions used on the basis that no company in the pharmaceutical industry is currently operating at optimal level due to imperfect competition, one of which is caused by the government regulation. While the other three *linear programming functions* are:

$$\begin{aligned} & [d_0^{t+1}(x_{t+1}, y_{t+1})]^{-1} = \max_{\phi, \lambda} \phi, \\ \text{st} \quad & -\phi y_{i,t+1} + Y_{t+1}\lambda \geq 0, \\ & x_{i,t+1} - X_{t+1}\lambda \geq 0, \\ & \lambda \geq 0 \end{aligned} \quad (2.3)$$

$$\begin{aligned} & [d_0^t(x_{t+1}, y_{t+1})]^{-1} = \max_{\phi, \lambda} \phi, \\ \text{st} \quad & -\phi y_{i,t+1} + Y_t\lambda \geq 0, \\ & x_{i,t+1} - X_t\lambda \geq 0, \\ & \lambda \geq 0 \end{aligned} \quad (2.4)$$

$$\begin{aligned} & [d_0^{t+1}(x_t, y_t)]^{-1} = \max_{\phi, \lambda} \phi, \\ \text{st} \quad & -\phi y_{it} + Y_{t+1}\lambda \geq 0, \\ & x_{it} - X_{t+1}\lambda \geq 0, \\ & \lambda \geq 0 \end{aligned} \quad (2.5)$$

Whereas θ is a scale, λ is the vector $N \times 1$ at the condition of *constant return to scale*, y_i is the *output* and x_i is the *input*, Y is the *matrix output* for all companies in the industry, X is the *matrix input* for all companies in the industry while $N1\lambda = 1$ is the *convexity constraint* ensuring that an inefficient company is selected as benchmark for similar sized companies.

Additionally, other than efficiency measures using DEA, this study also used financial ratios represented by *cash efficiency ratio* (*cash ratio*, *quick ratio* and *current ratio*) and *asset efficiency ratio* (*Inventory turnover*, *net working capital turnover*, *fixed asset turn over* and *total asset turnover*).

Ross (2008) explained the formula for all the financial ratios in this study. Xu and Wang (2009) using *quick ratio*, *cash ratio*, *inventory turnover* and *total asset turnover* as the predictors in prediction methods for financial failure prediction. Maricia and Georgeta (2012) using *current ratio* and *quick ratio* to analyze the business failure risk in their study. Delen et. al. (2013) using Liquidity ratios (*cash ratio*, *quick ratio*, *current ratio*) and turnover ratios (*inventory turnover*, *net working capital turnover*, *fixed asset turnover* and *total asset turnover*) for measuring firm performance with a decision tree approach. The ability of a company to pay a short-term debt can be evaluated by liquidity ratios, whereas long-term solvency ratios used to investigate the risk level of an investment in the firm could be for creditors. Turnover ratios used to measure the success level of the company, so that it generates revenues through utilizing assets, collecting receivables, and selling its inventories. According to the previous related studies, this paper used the following efficiency ratios:

$$\text{Current ratio} = \frac{\text{current assets}}{\text{current liabilities}} \quad (3)$$

$$\text{Quick ratio} = \frac{\text{current asset} - \text{inventory}}{\text{current liabilities}} \quad (4)$$

$$\text{Inventory turnover} = \frac{\text{Cost of Goods Sold}}{\text{Inventory}} \quad (5)$$

$$\text{Net working capital turnover} = \frac{\text{Sales}}{\text{Net Working Capital}} \quad (6)$$

$$\text{Net working capital} = \text{current assets} - \text{current liabilities} \quad (7)$$

$$\text{Fixed asset turnover} = \frac{\text{Sales}}{\text{Net Fixed Assets}} \quad (8)$$

$$\text{Total asset turnover} = \frac{\text{Sales}}{\text{Total Assets}} \quad (9)$$

$$\text{Cash Ratio} = \frac{\text{Cash}}{\text{Current Liabilities}} \quad (10)$$

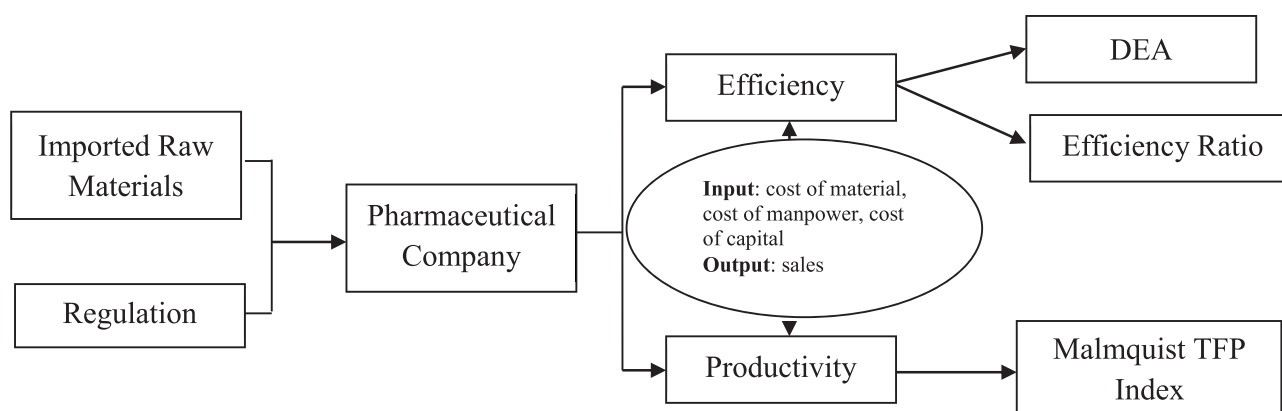


Figure 1. Research framework

Table 1. Descriptive Statistics Analysis (in USD*)

Description	Mean	Std Dev	Minimum	Maximum	Lower Quartile	Middle Quartile	Upper Quartile
Sales	275,485	339,723	6,731	1,196,093	39,387	104,329	382,000
Cost of Material	183,357	243,417	2,315	1,031,575	18,796	81,542	250,116
Cost of Manpower	21,417	26,031	645	116,609	3,853	10,686	31,988
Cost of Capital	41,390	61,706	1,918	266,579	4,741	18,051	39,087

Source: Output Stata 11 (Processed)

*Exchange rate: Rp 9,113/USD published by Bank Central of Indonesia per December 30th, 2011

Data and Design

This study is an *explanatory analysis* which aims to explain the value of relative efficiency and productivity changes produced from three inputs to a single output. The data employed in this study are *balanced panel data* covering the entire pharmaceutical companies listed in the Indonesian Stock Exchange period 2006 – 2011. During that period, there are twelve pharmaceutical company listed with various type of ownership. However, only eleven companies are included in the sample due to one company operates mainly in the supply of medical equipments and also incomplete data. All data was obtained through Thomson Reuters Knowledge and DataStream. Research framework employed in the study is shown in Figure 1.

Empirical Result

Statistic Descriptive

The publicly listed pharmaceutical companies included as sample for this study can be categorized based on ownership into three

groups: 1) two state owned/majority companies, 2) three privately owned domestic companies and 3) six foreign owned companies. Inputs and *output values* for each companies used in the DEA calculation are shown in Table 1.

Descriptive statistics showed that the largest expenditure for Indonesian pharmaceutical companies is the *cost of materials* with an average of USD183,357.18 covering raw materials, manufacturing and warehousing. The second is the *cost of capital* with an average of USD 41,390 covering fixed assets expenditures, machinery installation and intangible assets. While the *cost of manpower* is the least expenditure allocated with an average of USD21,417.46 covering both direct and indirect labor costs. Thus, the proportion between the total *cost of material*, *cost of capital* and *cost of manpower* are 9:2:1 respectively. The size of expenditures of *material* is mostly due to the majority of raw materials for Indonesian pharmaceuticals which must be imported from various countries such as India, China and European countries. Thus the material cost is influenced heavily by the low value of Indonesian currency relative to the country of the suppliers.

Table 2. Average Value of the Pharmaceutical Company Efficiency during 2006 – 2011 Period Using DEA

Company Name	Efficiency (VRS)					
	2006	2007	2008	2009	2010	2011
Darya-Varia Laboratoria Tbk	0.096	0.091	0.092	0.104	0.091	0.108
Enseval Putra Megatrading Tbk	0.219	0.182	1	0.182	0.182	0.226
Indofarma Tbk	1	1	1	0.333	1	1
Kimia Farma (Perseroan) Tbk	0.786	0.571	0.571	0.414	0.571	0.583
Kalbe Farma Tbk	1	1	0.455	0.537	0.455	0.455
Merck Tbk	0.610	0.545	0.699	0.565	0.545	0.592
Pyridam Farma Tbk	0.943	0.636	0.651	1	0.636	1
Schering Plough Indonesia Tbk	0.727	0.727	1	0.727	0.727	0.938
Millennium Pharmacon International Tbk	0.818	0.818	0.971	1	0.818	1
Taisho Pharmaceutical Indonesia Tbk	0.909	0.909	0.909	0.909	0.909	0.909
Tempo Scan Pasific Tbk	1	1	1	1	1	1

Source: *Output Deap* (Processed)

Companies Efficiency

Table 2 illustrates the relative efficiency value of Pharmaceutical Companies listed in the Indonesian Stock Exchange during the 2006-2011 periods using the DEA calculation. Efficiency value of 1 indicated the most efficient corporation.

Efficiency measurement using DEA showed that on average, the five most efficient companies are Tempo Scan Pasific Tbk with efficiency value of 1,00, followed by Taisho Pharmaceutical Indonesia Tbk (0.909), Millennium Pharmacon International Tbk (0.904), Indofarma Tbk (0.889) and Pyridam Farma Tbk (0.811). Conversely, the two least efficient pharmaceuticals companies in Indonesia are Darya-Varia Laboratoria Tbk (0.097) and Enseval Putra Megatrading Tbk (0.332). Based on the ownership, three out of five most efficient companies are foreign owned companies, one is a privately owned domestic company and one is a state owned/majority company.

Tempo Scan Pasific Tbk and Taisho Pharmaceutical Indonesia Tbk as the two top most efficient companies have different ownership structure, with Tempo Scan Pasific Tbk as domestic privately owned and Taisho Pharmaceutical Indonesia Tbk as foreign owned company. However, both are producers focusing on making and selling Over the Counter (OTC) Medications.

Tempo Scan Pasific Tbk sales of OTC medications contribute to up to 90% of total sales,

while prescription only contributes 10% (*Ethical*) of total sales. Based on the 2011 data, the market share of Tempo Scan Pasific Tbk in the OTC segment reached up to 45.1% of total OTC sales in Indonesia⁵. Similar focus was also adopted by Taisho Pharmaceutical Indonesia Tbk whose OTC sales contributed up to 78% of total sales⁶. Good target market and strong focus on OTC sales has made this company to have the best efficiency in the Indonesia pharmaceutical industry. Thus, it can be said that stronger focus on OTC medicines might increase efficiency of Pharmaceutical companies in Indonesia.

This factor may be attributed to the greater size of annual total sales for OTC medicines as well as the tendency of Indonesia consumer to self prescribes their medication. Based on the data by the Indonesian Central Biro of Statistics (BPS) in 2010, majority people in Indonesia (68.71%) prefer to self medicate themselves, by choosing their medicine without doctor's prescriptions. Additionally, 27.58% of Indonesians prefer self medication by using factory made traditional medicine recognized and registered by the Indonesian Health Ministry. This means, only 3.71% Indonesian prefer to doctor's prescriptions when choosing medications and the rest prefers using self prescribed over the counter medicine sold openly in retail outlets. Such Indonesian demographic characteristics become the main reason Tempo Scan Pasific Tbk and Taisho Pharmaceutical Indonesia Tbk focusing on the production and sales of OTC

⁵ <http://www.indonesiainfinance today.com>

⁶ <http://www.indonesiainfincetoday.com/read/24520>

Table 3. Average Value of the Pharmaceutical Company Efficiency during 2006 – 2011 Period Using Efficiency Financial Ratios (in times)

Company Name	Average Cash Efficiency				Average Asset Efficiency		
	Cash Ratio	Quick Ratio	Current Ratio	Inventory Turnover	NWC Turnover	Total Asset Turnover	Fixed Asset Turnover
Darya-Varia Laboratoria Tbk	1.70	3.60	4.30	3.057	1.813	1.016	4.883
Enseval Putra Megatrading Tbk	0.25	1.11	1.92	7.050	7.104	2.883	23.397
Indofarma Tbk	0.35	1.10	1.46	5.127	5.671	1.390	11.458
Kimia Farma (Perseroan) Tbk	0.49	1.43	2.25	5.612	5.128	1.833	6.709
Kalbe Farma Tbk	1.42	2.73	4.06	2.900	2.561	1.372	6.069
Merck Tbk	2.46	4.59	6.36	3.399	2.573	1.702	11.977
Pyridam Farma Tbk	0.13	1.12	2.03	2.187	6.336	1.144	2.074
Schering Plough Indonesia Tbk	0.09	0.63	1.32	1.596	-8.515	1.161	6.340
Millennium Pharmacon International Tbk	0.13	0.85	1.41	7.966	11.940	3.194	101.071
Taisho Pharmaceutical Indonesia Tbk	1.87	3.67	4.31	4.154	2.145	1.126	4.724
Tempo Scan Pasific Tbk	1.89	2.81	3.70	4.623	2.503	1.270	5.739
Industry Average for Indonesian Pharmaceutical Industry	0.98	2.15	3.01	4.330	3.570	1.640	16.770

Source: Thomson-Reuters Knowledgege (Processed)

medicines enjoy greater efficiency relative to other pharmaceutical companies with less focus on OTC medicine.

However, such demographic characteristic may come as a shock since according to the 2010 Census by the Indonesian Central Biro of Statistics; up to 31,023,400 people (16.56%) of the total population are considered below the poverty line. While the Indonesian government through the Ministry of Health provides free health care and medical treatment in form of the Health Insurance (*ASKES*), the Public Health Insurance (*JAMKESMAS*) and the Local Public Health Insurance (*JAMKESDA*) for 40% of the poorest Indonesians to enable their access to better health care (Holloway, 2011). Ideally, with the lower portion of poor people and better access to health care, people would use that opportunity to go to health care facility and consume medication as prescribed by the doctor instead of doing self medication.

Therefore, there are two possible explanations on why Indonesians still prefers self-medication than doctor's prescriptions. First, government health care aid might not reach the real segment, thus many of the poorer population can not access health care aids given in medical facilities due to their financial limitation. This condition forces them to consume OTC medication sold in retail outlets rather than prescribed medicines. Second, the mind set of many Indonesians may really prefer OTC medicines, thus avoiding or ignoring the doctor's prescriptions

when they are ill.

On the other hand, even though both Tempo Scan Pasific Tbk and Taisho Pharmaceutical Indonesia Tbk have the greatest relative efficiency based on DEA calculation, their financial efficiency ratios are not better than those with lesser relative efficiency. Tempo Scan Pasific Tbk and Taisho Pharmaceutical Indonesia Tbk has *cash efficiency ratio* above the industry average, while Millennium Pharmacon International Tbk has the lowest *cash efficiency ratio* in the industry (See Table 3). Lower *cash efficiency ratio* means greater ability of the company to manage their cash and current assets. On the other hand, Millennium Pharmacon International Tbk has the highest *asset efficiency ratio* in the industry, while Tempo Scan Pasific Tbk and Taisho Pharmaceutical Indonesia Tbk have below average ratios. *Asset efficiency ratio* implies that higher ratios means higher efficiency in utilizing their assets.

Millennium Pharmacon International Tbk has *asset turnover* ratio of 101.071, well above the industry average of 16.77 as shown in Table 3. This is due to the core business of the company which focuses on the distributions of pharmaceutical products. As a distributor, Millennium Pharmacon International Tbk does not need fixed assets as large as the manufacturers which mass produce pharmaceutical products. Therefore, the company obtained far greater *fixed asset turnover* compared to others in the industry.

Differences in the order of the most to the least efficient companies between DEA and the *efficiency financial ratio* may be attributed to the number of factors entered. In order to measure efficiency, DEA used multiple inputs to assess one or more outputs; while *financial ratio* can only use one input to assess one output in a given time. This limitation of *efficiency financial ratios* to describe the overall company activities in measuring efficiency may lead to a bias in measurements (Saranga and Phani, 2008).

One interesting finding is the efficiency of Indofarma Tbk, which obtain the best relative efficiency together with Tempo Scan Pasific Tbk in five out of the six periods employed in the study. Indofarma Tbk is one pharmaceutical company which per Desember 2011, was majority owned by the state (80.1%), while 19.3% of the shares was owned by public and 0.6% owned by the employees. This is contrary to the well known fact that this company has low profitability. There are several possible explanations to this finding. First, that is efficiency measurement using DEA assumed that sales, and not profitability to be the output of pharmaceutical companies. Thus, even though Indofarma Tbk may have low profitability, this is not due to low sales volume. As known, Indofarma Tbk as Majority State Owned-company has greater authorization to distribute pharmaceutical products than foreign owned or domestically owned private companies. As the largest producer of generic medicines in Indonesia, Indofarma Tbk distributes its production to various government owned service centers in the capital as well as in the provinces. Even though the margin for generic medicines has been set to a minimum by the government, the large volume of sales in generic medicines has kept Indofarma Tbk efficient.

Second, the efficiency gained by Indofarma Tbk may have been obtained due to tangible assets supporting the production capacity. Historical record showed Indofarma Tbk as the first pharmaceutical company founded by the Dutch-Indian colonial government. At that time, the colonial government invested large capacity machinery to mass produce medicines,

thus gave Indofarma Tbk a head start as the prime supplier of medicine to Indonesians. This head start in production capacity enabled lower and more efficient fixed costs for the company due to *process innovation*, which is conducting manufacturing process at the lowest possible cost by maximizing the *economies of scale* (Belcher and Nail, 2000).

However, when measured using *efficiency financial ratio*, Indofarma Tbk has a slightly lower *total asset turnover* than Kimia Farma (Perseroan) Tbk, even though Indofarma Tbk still has better overall *cash* and *asset efficiency*. Kimia Farma (Perseroan) Tbk is another state owned/majority pharmaceutical company included in the sample. Even though they are both state controlled, Kimia Farma (Perseroan) Tbk has greater total sales due to more vertically integrated operation while Indofarma Tbk focuses more on the production of generic medicines with lower price per unit. This shows that, despite larger the sales volume, Indofarma Tbk has lower ratio of sales to total asset compared to Kimia Farma (Perseroan) Tbk due the low cost nature of generic medicines it produces.

During 2009, Indofarma Tbk experiences a sharp drop in efficiency from 1.00 in 2008 to only 0.33 in 2009. This drop in efficiency is due to the increase of raw material price imported mostly from India. The price increased due to the global financial crisis which led to the devaluation of Rupiah. The problem thus compounded due to the inability of Indofarma Tbk to cross subsidize the production of generic medicines, thus forcing the management to postpone the import of raw material and reducing the production of least profitable products. This decision resulted in the decrease of sales which in turn reduces efficiency. This condition does not affect Indofarma Tbk alone, but also influences other pharmaceutical companies, such as Pyridam Farma Tbk, Kalbe Farma Tbk, Schering Plough Indonesia Tbk, Kimia Farma (Perseroan) Tbk and Merck Tbk, as shown in their lower efficiency during the 2008-2009 periods.

Even though the global financial crisis in 2009 has affected the efficiencies of pharmaceutical companies in Indonesia across the

board, according to DEA calculation Darya-Varia Laboratoria Tbk and Enseval Putra Megatrading Tbk was shown to be the least efficient on average, relative to other pharmaceutical companies operating in Indonesia. Darya-Varia Laboratoria Tbk is a foreign owned pharmaceutical company which focuses on the prescribed medicine segment, contributing up to 51% of their total sales compared to 36% contribution of *consumer health* products. This focus on prescribed medicine is in total opposite to the strategy of Tempo Scan Pasific Tbk and Taisho Pharmaceutical Indonesia Tbk which are shown as the most efficient companies of publicly listed pharmaceutical companies in Indonesia. This difference in product and consumer target may be the cause of differences in efficiency between the companies.

Not only when calculated using DEA, the efficiency of Darya-Varia Laboratoria Tbk was also low when calculated using financial ratios. On average, Darya-Varia Laboratoria Tbk has greater cash efficiency ratios (See Table 3), which means that Darya-Varia Laboratoria Tbk reserved more current assets to meet its short term liabilities, which led to less efficient utilization of cash. Conversely, its *asset efficiency* ratio was constantly above the industry average, implying that the company's asset utilization was relatively efficient.

Enseval Putra Megatrading Tbk is a domestic private owned pharmaceutical company. In contrast to other pharmaceutical companies which experiences sharp decrease of efficiency in 2008 due to the global financial crisis. Enseval Putra Megatrading Tbk experienced the best efficiency relative to other Indonesian pharmaceutical companies, despite having lower efficiency during other periods. The calculation using DEA was in opposite to the calculation using financial ratio. On average, the *cash efficiency* ratio of Enseval Putra Megatrading Tbk are lower than industry average while on average the *asset efficiency* ratio was consistently higher than the industry average. This showed that according to the financial ratios, Enseval Putra Megatrading Tbk was efficient in managing its cash and assets.

Domestic vs Foreign Efficiency Companies

Based on differences in the efficiency between groups of domestic and foreign owned pharmaceutical companies, this study tests the significance of such differences. This idea stems from the regulations regarding pharmaceutical companies in Indonesia considered to be constricting to foreign pharmaceutical players united in the International Pharmaceutical Manufacturers Group⁷.

Despite the government effort to relax the regulation regarding foreign owned pharmaceutical companies, existing regulation are still thought to be strong disincentive for foreign investments in the pharmaceutical industry. Such disincentive, if true, would constrict the activities of foreign owned pharmaceutical companies proven by differences in the size of investment activities between domestic and foreign owned pharmaceutical companies.

Based on Table 4, the top three companies with the largest investment activities are Kalbe Farma Tbk, Tempo Scan Pasific Tbk and Enseval Putra Megatrading Tbk which were all domestic owned companies. On the other hand, two companies with the least investment activities are Millennium Pharmacon International Tbk and Pyridam Farma Tbk which were both foreign owned companies. Also, *t-test* in Table 5 shows there are significant differences in investment activities between domestic against foreign owned company at confidence level 95%.

Further testing was also conducted to compare the differences between the efficiency of domestic and foreign owned companies using independent sample *t-test* with confidence level of 95%. Whereas, the hypothesis for this test is:
 Ho: There is no difference between the average efficiency of domestic and foreign-owned pharmaceutical companies in Indonesia.
 Ha: There is a difference between the average efficiency of domestic and foreign-owned pharmaceutical companies in Indonesia.

Based on the test result shown in Table 6, the null hypothesis can not be rejected. This means that there is no difference between the

⁷ <http://www.businessreview.co.id/bisnis-investasi-886.html>

Table 4. Investment Activities of Indonesian Pharmaceutical Companies during 2006 – 2011 Period (in USD*)

Company	2011	2010	2009	2008	2007	2006
Kalbe Farma Tbk	69,236,695	25,472,841	70,826,292	20,435,093	4,638,538	28,543,070
Tempo Scan Pasific Tbk	19,846,264	9,327,774	11,118,622	3,929,222	8,408,318	6,555,360
Enseval Putra Megatrading Tbk	17,153,736	15,213,541	10,251,070	5,543,729	5,584,769	-20,331,943
Kimia Farma (Perseroan) Tbk	5,968,287	1,462,197	8,105,125	2,915,725	3,094,700	2,525,623
Darya-Varia Laboratoria Tbk	3,751,344	3,491,166	2,182,377	1,749,259	3,637,551	1,037,529
Taisho Pharmaceutical Indonesia Tbk	1,428,948	2,422,583	461,648	1,537,803	2,274,333	855,810
Schering Plough Indonesia Tbk	1,171,513	428,618	627,236	446,944	878,525	1,305,278
Indofarma Tbk	378,909	496,214	2,074,289	1,113,355	811,588	20,191
Pyridam Farma Tbk	229,123	299,901	205,201	206,408	513,113	150,664
Millennium Pharmacon International Tbk	116,098	205,531	682,761	1,079,008	325,908	67,047
Merck Tbk	-9,811,588	2,336,772	1,829,913	1,412,488	868,649	549,325

Source: Annual Cash Flow - Reuters Knowledge (2011)

*Selling rate: Rp 9.113/USD published by Bank Central of Indonesia per December 30th, 2011

Table 5. Descriptive Statistics Independent Sample t-test between Indonesian Domestic and Foreign Owned Pharmaceutical Companies in Investing Activities

Ownership	Obs.	Investment	
		Mean	Variance
Domestic	30	24.79	17.05
Foreign Owned	36	109.63	30.51
<i>t-statistics</i>			-24.272
<i>p-value</i>			0.0192

Source: Output Stata (Processed)

Table 6. Descriptive Statistics Independent Sample t-test between Indonesian Domestic and Foreign Owned Pharmaceutical Companies

Ownership	n	Average Efficiency	
		Mean	Variance
Domestic	5	0.690733	0.117688
Foreign Owned	6	0.686917	0.126936
<i>t-statistics</i>			-0.022
<i>p-value</i>			0.9829

Source: Output Stata (Processed)

average efficiency of domestic and foreign-owned pharmaceutical companies in Indonesia. This result is in contrast to the finding of Pannu et al. (2009), which stated that partially, foreign-owned companies has lower efficiencies than domestic-owned companies due to lesser size of investments. This outcome led to earlier hypothesis that foreign-owned companies may not be able to achieve optimum *economies of scale* due to regulation issues. The current finding in this study, however, showed that on average there are no differences between the efficiencies of domestic and foreign-owned pharmaceutical companies in Indonesia.

The absence of differences between the mean efficiency of domestic and foreign owned group of companies refute the opinion that ex-

isting government regulation, such as ownership proportion and importing licence policies, may constrain the operation of foreign owned companies disproportionately. Foreign owned pharmaceutical companies such as Taisho Pharmaceutical Indonesia Tbk, Millennium Pharmacon International Tbk and Pyridam Farma Tbk can operate efficiently and compete with various domestic companies, both private and state controlled. This result supports the hypothesis that market and product selection influences efficiency. Companies which sold more *over the counter* medicines has greater efficiency than companies which sold more prescribed medicines, irrespective of their ownership.

The absence of differences between foreign and domestic companies does not refute the

Table 7. Average Efficiency, Technological Change, Efficiency Change and Total Factor Productivity Change

Year	Output Efficiency	Technological Change	Pure Technical Efficiency Change	Total Factor Productivity Change
2006	0.737	--	--	--
2007	0.680	0.862	0.908	0.766
2008	0.759	0.600	1.166	0.625
2009	0.616	1.429	0.767	1.256
2010	0.630	0.550	1.041	0.572
2011	0.710	0.653	1.135	0.741

Source: Output Deap (Processed)

view that Indonesian pharmaceutical industry may be considered as a highly regulated industry in comparison to other pharmaceutical industries in South East Asia. The strict regulation may force foreign owned companies to abide the regulations by allocating their investments to build manufacturies and purchase heavy machineries as well as operational licence in Indonesia. The large initial cost may reduce efficiency across the board and lengthen the break even point to new entrants, compared to existing players. On the other hand, the regulation for domestic companies may have been set more relaxed in a way that the market nearly becomes perfect competition, which enables players to enter and exit the market freely. This condition forces existing players to adopt price penetration strategy to be able to compete and sell more products. This strategy in turn caused domestic players to be relatively less efficient in generating revenues than it should be (Mazumdar dan Rajeev, 2009). Both sides of the regulation experiences different challenges which created inefficiencies, thus domestic and foreign owned pharmaceutical companies in Indonesia exhibit relatively similar level of efficiency regardless of ownership and regulations.

Productivity Changes

It is shown in the second column of Table 6 that the average output efficiency of pharmaceutical companies in Indonesia is 0.737. This meant that on average pharmaceutical companies in Indonesia can increase their output by 26% without adding any additional input. The fluctuating average output efficiency between the 2006 to 2011 period indicates two possible outcomes. First, an outward shift may occur across the board when the average efficiency

increases, which meant an increase of internal efficiency from each decision making unit (DMU) below the *frontier*. Second, an outward shift may occur only to DMU at the frontier, when the average efficiency decreases, which may indicate technological or process innovation which increases the gap efficiency between the leader and the rest of the industry. Thus, internally this also meant decrease in relative efficiency across the board to players below the *frontier* (Mazumdar dan Rajeev, 2009).

The third column in Table 7 shows technological change of greater than one means that there may be technological applications which increase efficiency significantly, relative to the previous year. Data showed that the Indonesian pharmaceutical industry only experienced one increase (progress) during the 2006-2011 periods, which is during 2009 with an increase of Technological Change of 43%. This increase in Technology may occur when new entrants utilize superior technology or existing players invest in a new technology which influences the frontier and existing DMU below the frontier (Mazumdar dan Rajeev, 2009). However, more accurate explanation would be that the global financial crisis in 2009 forced existing players to upgrade their technology to cope with the fluctuation of raw material prices due to currency fluctuations, by using less input to produce more or the same amount of output.

The fourth column of Table 7 shows *Efficiency Change* which explains the changes in efficiency between periods. *Efficiency change* of greater than one showed an increase in efficiency from the previous year, while a value of less than one showed a decrease in efficiency from the previous year. During 2007 and 2009, Indonesian pharmaceutical industry experienced decrease in average efficiency of 9% and

23% when compared to 2006 and 2008, respectively. While during year 2008, 2010 and 2011, there were increasing in the average efficiency from earlier year, of 16%, 4.1% and 13.5% respectively.

Compared to the *technological change*, the level of *efficiency change* tends to be more fluctuative, even though the overall *technological change* was decreasing. This showed that companies in the Indonesian pharmaceutical industry could increase their efficiency without increasing application of technologies. However, in overall the Total Factor Productivity (TFP) shown in the fifth column of Table 7 showed TFP decreased during the decrease in Technological change and increased during the application of newer technology. In other words, TFP are more affected by technological changes rather than changes in internal efficiency. This finding may be contradictive to the findings of Gonzales and Gascon (2004) studying the Spanish pharmaceutical industry along with Mazumdar and Rajeev (2009) studying the Indian pharmaceutical companies. Both study found that TFP was affected more by changes in the internal efficiency rather than technological changes. This difference showed that the Indonesian pharmaceutical companies can enjoy the benefits of technological innovations in producing pharmaceutical products due to an increase of efficiency.

In summary, the findings in this paper suggest the importance of using a more accurate measure based on an input-output framework, such as DEA, to evaluate firm efficiency instead of relying on efficiency measures based on financial ratios. Different with previous studies, such as Pannu et al. (2011), this paper does not find significant difference between the efficiency of foreign owned and locally owned pharmaceutical companies. One possible explanation about this finding is the stricter regulation in the Indonesian pharmaceutical industry which hinders the benefits of economies of scale and transfer of technology. However, whether and how regulation may impact firm efficiency in different countries, especially in the pharmaceutical industry, may need further research.

Conclusion

This study concludes that the five topmost efficient pharmaceutical companies in Indonesia are Tempo Scan Pasific Tbk, Taisho Pharmaceutical Indonesia Tbk, Millennium Pharmacon International Tbk, Indofarma Tbk and Pyridam Farma Tbk. Calculating efficiency using DEA are more accurate than using efficiency financial ratios due to its capability in capturing all aspect of company activities rather than a partial view as used in the ratios. It is also in line with research Xu and Wang (2009) which states that, in the proposed prediction scheme, the efficiency of corporation is more efficient obtained through DEA rather than using efficiency financial ratio analysis.

Further, this study finds no evidence that foreign owned pharmaceutical companies are more efficient than local companies. Differences in efficiency are more due to selection of market and product segment rather than ownership. Companies with greater sales contribution of over the counter medicine are generally more efficient than companies which focus on prescribed medicines. This study also showed that the *Total Factor Productivity* (TFP) of Indonesian pharmaceutical industry is more influenced by technological changes than changes in internal efficiency. This finding is in opposite to the findings in India and Spain in which changes in TFP is influenced more by changes in internal efficiency rather than technological changes.

The findings of this study must be seen in the context of several limitations. This study only used pharmaceutical companies listed in the Indonesian stock exchange which only accounts to 9.24% of the total pharmaceutical companies operating in Indonesia. This study measures difference of efficiency based on ownership, but not based on productivity due to insufficient data on production volume and the utilization of machinery and manpower by each company. Even so, this study employed a valid and robust research method which have been tried and tested in previous researches, therefore the results are accountable.

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