Abstract

Biodiesel is a non-toxic and very low sulfur containing, renewable and biodegradable diesel fuel substitute with low volatility and high cetane number. It is derived from fresh or waste vegetable oils and animal fats transesterified with short chain alcohol such as ethanol or methanol in the presence of a catalyst. Several new types of carriers and technologies have been adopted in the recent past to improve the ability of a catalyst in transesterification. One of the new trends is nanoparticles-based immobilization of enzyme as a catalyst. The combination of the precise physical, chemical, optical and electrical properties of nanoparticles with the specific recognition site or catalytic properties has led them to appear in a myriad of novel nanomaterial application. Enzyme immobilized on nanoparticles showed a broader working temperature and pH range and thermal stability than the native enzyme. Enhancement in the reactivity of nanocatalysts is associated with their increased surface area, greater concentrations of highly reactive edge, unusual and stabilized lattice planes. The greater activity of nanomaterial immobilized biocatalyst affords operational simplicity, low energy consumption, and greater safety, in the process of transesterification. This review article highlights the issues including the exploration of the ability of nanomaterial to immobilize biocatalyst and factors that influence the activity of biocatalysts upon immobilization.

Key words: Biodiesel, Immobilization, nanomaterial, transesterification

Introduction

The issue of worldwide energy consumption has steadily increased, claiming higher budget for living, increasing transportation, industrial and petrochemicals. Fig. 1 shows that the majority of the consumed energy is provided by fuel in the world. Global energy consumption in 2010 rebounded strongly in all regions, driven by economic recovery. The world energy consumption growth reached 5.6% in 2010, the highest rate since 1973. After falling for two consecutive years in 2008 - 2009, global oil consumption grew by 2.7 million barrels per day (b/d), or 3.1%, to reach a record level of 87.4 million b/d. Although this was the largest percentage increase since 2004, the worldwide oil consumption growth rate started declining and remained the weakest among all fossil fuels till 2011 and grew slowly again in 2016 - the third consecutive year in which demand has grown by 1% or less - much weaker than the rates of growth world had become used to over the previous 10 years or so (Dudley 2017). As a result, global crude oil production increased by 2.8 million b/d in 2015, led by a 1 million b/d increase in U.S. production. The bulk of the rest of the world's oil production increase came from OPEC, which cumulatively boosted production by 1.6 million b/d over 2015. BP's definition of crude oil “includes crude oil, shale oil, oil sands and NGLs (natural gas liquids - the liquid content of natural gas where this is recovered separately).”
However, the petroleum is a limited source for fuel that is quickly becoming insufficient and more expensive. In addition, petroleum-based product is one of the main causes of emission of carbon dioxide (CO\textsubscript{2}) to the atmosphere. The transportation sector worldwide is almost dependent on petroleum-derived fuels (Fig. 1). The major causes of global CO\textsubscript{2} emissions are transport and industrial sector, which accounts for millions of barrels of oil consumption per day. As in 2015, the strength in oil demand was most pronounced in consumer-led fuels, such as gasoline. CO\textsubscript{2} emission goes up by 0.7%, being trivial in relation to concerns about energy security or climate change (EIA 2011). As a result, there has been a growing interest in alternative source.

Biodiesel is one of the energy sources as an alternative fuel for diesel engines. Biodiesel is defined as a mono alkyl ester, which is produced by transesterification of oil from renewable biological source like vegetable or animal fats. It is an alternative diesel fuel because of its environmental benefits such as being biodegradable, nontoxic and with low carbon dioxide emission profiles (Jegannathan et al. 2010).

In general, there are two prospective methods to produce biodiesel in industrial scale: chemical and enzyme catalyzed method. In chemical-catalyzed method, acid and alkali catalysts can be used (acidic or bases solution) for the production of biodiesel, but there are several drawbacks to this approach, including difficulties in the recovery of the glycerol and potassium and/or sodium salt, and the wastewater treatment problem (Jegannathan et al. 2010, Li et al. 2011). On the contrary, the enzymatic reaction by lipase is a clean technology due to its non-toxic and environment friendly nature. In addition, the process produces high purity grade product and enables easy separation from the by-product, glycerol (Winayawanuwattikun et al. 2008).

Lipase is a hydrolytic enzyme, which catalyzes a reversible reaction to hydrolyze triglycerides to free fatty acids and glycerol and the esterification of free fatty acids and alcohol to ester (e.g. biodiesel). So lipase is
used as industrial biocatalyst (Sangeetha et al. 2011). Much research is needed to overcome problems such as enzyme inhibition by methanol, exhaustion of enzyme activity and high cost of enzymes, which may contribute to the global effort on industrial implementation of the enzymatic production of biodiesel in the near future (Iso et al. 2001, Nordblad and Adlercreutz 2008, Fjerbaek et al. 2009). Immobilization improves the enzyme stability under the reaction conditions, and enhances enzyme activity, thus, makes the repeated use of the enzyme feasible, permits the use of enzyme for diverse applications and thus lowers the production cost (Bajaj et al. 2010, Tan et al. 2010, Sangeetha et al. 2011). Immobilization provides a better environment for enzyme to act and also offers better product recovery (Lee et al. 2006).

Lipase can be immobilized on different porous support materials. Immobilized enzymes are defined as biocatalysts restrained and localized into microenvironment to retain their catalytic properties. Immobilization usually can increase stability and makes the reuse of the enzyme preparation very simple (Twyman et al. 2005). Immobilization is studied using covalent bonding, cross-linking, entrapment, adsorption, and encapsulation. Selection of an immobilization strategy greatly influences the properties of biocatalyst (Iso et al. 2001, Yagiz et al. 2007, Meunier et al. 2010, Xie and Ma 2010).

Nanomaterials constitute novel and interesting matrices for enzyme immobilization. While their high surface to volume ratio is an obvious advantage, their Brownian motion can impact the behavior of enzymes immobilized on these matrices. Such immobilized enzyme systems have been used in both aqueous and low water media for bio catalysis and resolution of race mates (Xie and Ma 2010, Yiu and Keane 2012). This overview examines the behavior of enzymes immobilized on nanomaterials and discusses the results reported with such biocatalyst preparations.

**Biodiesel in general**

Biodiesel is mono alkyl esters of long chain fatty acids contained in the renewable natural resources, such as plant, oils and fats (Moser 2009). Biodiesel is obtained from the transesterification of triglycerides with alcohol using an acid or base catalyst. Biodiesel contains no nitrogen or aromatic component and has sulfur content less than 15 ppm. Biodiesel has efficiency as a fuel similar to diesel oil and at the same time it is non-toxic, biodegradable, and has a low emission value (Jegannathan et al. 2010). The use of biodiesel can reduce emissions of HCs, CO, PM, sulfates, polycyclic aromatic hydrocarbons and nitrated polycyclic aromatic hydrocarbons. Also NOx emissions increase with the concentration of biodiesel in the fuel (Balat and Balat 2010).

Generally, vegetable oil or animal fats are esters of saturated and unsaturated monocarboxylic acids with the trihydric alcohol glycerol. These esters are called triglycerides, which can react with alcohol in the presence of a catalyst, a process known as transesterification. The simplified form of its chemical reaction is presented in equation:
**Fig. 2.** Transesterification of Triglyceride.

Where, R in triacylglycerol is long-chain hydrocarbons, sometimes called fatty acid chains. Normally, there are five main types of chains in vegetable oils and animal oils: palmitic, stearic, oleic, linoleic, and linolenic. When a triglyceride is converted stepwise to diglyceride, monoglyceride, and glycerol - one mol of fatty acid is liberated at each step. Usually, methanol is the preferred alcohol for producing biodiesel because of its low-cost (Semwal et al. 2011).

Transesterification, also known as alcoholysis is the reaction of vegetable oil or fat with an alcohol to form esters and glycerol. Stoichiometrically to complete a transesterification reaction a 3:1 molar ratio of alcohol to triglycerides is needed. In practice, to have a maximum yield of esters, this ratio must be higher than the stoichiometric ratio (Leung et al. 2010). A catalyst is usually used to improve the reaction rate and yield. Because the reaction is reversible, excess alcohol is used to shift the equilibrium to the products side.

In transesterification, the alcohol is deprotonated with base to make stronger nucleophile. Commonly, ethanol or methanol is used. As can be seen, the reaction has no other inputs than the triglyceride and the alcohol. Transesterification consists of sequence of three consecutive reversible reactions. The first step is the conversion of triglyceride to diglyceride, followed by the conversion of diglyceride to monoglyceride, and finally monoglyceride to glycerol, yielding one ester molecule for each glyceride at each step. The basic mechanism of transesterification is shown in Fig. 3.

\[
\text{Triglyceride} + \text{ROH} \quad \rightarrow \quad \text{Diglyceride} + \text{R'}\text{COOR} \\
\text{Diglyceride} + \text{ROH} \quad \rightarrow \quad \text{Monoglyceride} + \text{R'}\text{COOR} \\
\text{Monoglyceride} + \text{ROH} \quad \rightarrow \quad \text{Glycerol} + \text{R'}\text{COOR}
\]

**Fig. 3:** General equations for transesterification on triglyceride.

The catalyst used for biodiesel production can be grouped as follows base-catalyzed, acid-catalyzed, lipase-catalyzed, heterogeneous catalyzed process.

**Homogeneous acid-base catalyzed transesterification**

The homogeneous acid-base catalysis is used if the feedstock contains high FFA. The feedstock first treated with \(\text{H}_2\text{SO}_4\) to reduce the level of FFA to below 1 wt%, followed by transesterification process catalyzed by homogeneous base catalysis. In this method, the yield of FAME is very high but the rate of FFA esterification
reaction is relatively slow. The drawback to this two-step process is even more pronounced due to the requirement of extra separation steps to remove the catalyst in both stages. Although the problem of catalyst's removal from the first stage can be avoided by using a catalyst from the second stage through the neutralization process, the use of extra base catalyst will add to cost of biodiesel production (Lam et al. 2010).

**Homogeneous base catalyzed transesterification**

The base catalyzed transesterification of vegetable oils proceeds faster than the acid-catalyzed reaction. Industrial processes usually favor base catalysts, such as alkali metal alkoxides and hydroxides as well as sodium potassium carbonates. The mechanism of the base-catalyzed transesterification is the reaction of the base with the alcohol, producing an alkoxide and the protonated catalyst (1). The nucleophilic attack of the alkoxide at the carbonyl group of the triglyceride generates a tetrahedral intermediate (2) from which the alkyl ester and the corresponding anion of the diglyceride are formed (3). The latter deprotonates the catalyst, thus regenerating the active species (4), which is now able to react with a second molecule of the alcohol, starting another catalytic cycle (Dias et al. 2008).

Alkali metal alkoxides (as CH$_3$ONa for the methanolysis) are the most active catalysts, since they give very high yields (>98%) in short reaction times, even if they are applied at low molar concentrations. Alkali metal hydroxides (KOH and NaOH) are cheaper than metal alkoxides, but less active. The presence of water gives rise to hydrolysis of some of the produced ester, with consequent soap formation and reduces the ester yield and considerably difficult to recover the glycerol due to the formation of emulsions. On the other hand, using potassium carbonate gives a high yield of fatty acyl alkyl ester and reduces the soap formation (Chanakya et al. 2013).

**Heterogeneous base-catalyzed transesterification**

Base catalysts have been developed for biodiesel production, such as basic zeolites, alkaline earth metal oxides and hydroxalicates. CaO had attracted much attention due to their relatively high basic strength, low solubility in methanol and can be synthesized from cheap sources like limestone and calcium hydroxide. The yield of FAME was up to 90% after 1 h reaction time at methanol reflux temperature and methanol to oil ratio 12:1 (Liu et al. 2008, Son and Kusakabe 2011, Kouzu and Hidaka 2012).

The other research, findings proved CaO as a potential solid catalyst in transesterifying triglycerides to methyl ester. CaO requires a thermal activation to remove CO$_2$ and moisture. The research reported that CaO acts as a heterogeneous catalyst in transesterifying the adsorbed waste palm oil on spent bleaching clay. Compared to the conventional catalysts (NaOH and KOH), the CaO-catalyzed reaction yielded much higher biodiesel (about 90%) from the waste cooking oil (6.6 - 6.8% FFA content) compared to only 46 and 61% yield using NaOH and KOH, respectively (Boey et al. 2011a,b).

The reaction mechanism for CaO-catalyzed transesterification is described as the methoxide ion that is attached to the catalyst surface attack the carbonyl carbon of the triglyceride molecule. This results in the formation of tetrahedral intermediate. Then the intermediate is rearranged to form a diglyceride anion and a mole of methyl ester. The charged-anion is then stabilized by a proton from the catalyst surface to form
diglyceride and at the same time regenerates the catalyst. The cycle continues until all three carbonyl centers of the triglyceride have been attacked by the methoxide ions to give one mole of glycerol and three moles of methyl esters (Boey et al. 2011c).

However, the yield of FAME dropped when waste cooking oil with high FFA was used under the same reaction condition. It is obvious that the basic sites of CaO were poisoned by strong adsorption of FFA on the surface catalyst. However, the catalytic activity of CaO can be regenerated if CaO is subjected to an activation treatment at 700°C in order to remove the main poisoning species (the carbonate groups) from the surface. Leaching of the catalyst was still observed in the transesterification reaction, although prior thermal treatment was employed (Lam et al. 2010).

**Heterogeneous acid-catalyzed transesterification**

Biodiesel research is focused on exploring new and sustainable solid acid catalyst for transesterification reaction. The advantages of using a solid catalyst are (1) they are insensitive to FFA content, (2) esterification and transesterification occur simultaneously, (3) eliminate the washing step of biodiesel, (4) easy separation of the catalyst from the reaction medium, resulting in lower product contamination level, (5) easy regeneration and recycling of catalyst, (6) reduced corrosion problem, even with the presence of acid species (Semwal et al. 2011).

The ideal solid acid catalyst for transesterification reaction should have characteristics such as an interconnected system of large pores, a moderate to high concentration of a strong acid site, and a hydrophobic surface. The overviews of various solid acid catalysts used (ZrO₂, TiO₂, SnO₂, zeolite, sulfonic ion-exchange resin, sulfonic modified mesostructure silica, sulfonated carbon-based catalyst, and heteropolyacids) in transesterification reaction that give the high conversion yield of FAME are discussed (Enweremadu and Mbarawa 2009, Petchmala et al. 2010).

**Lipase as biocatalyst in transesterification**

Lipase (EC.3.1.1.3, triacylglycerol acylhydrolases) are a group of enzymes, which can hydrolyze triacylglycerols at an oil-water interface to release fatty acids ester and glycerol. The substrates of lipase are triacylglycerols, which have very low solubility in water (Fig. 4). Lipase is present in microorganisms, plant and animals. Lipases catalyze a wide range of reactions, including hydrolysis, trans-esterification, alcoholysis, acidolysis, esterification and aminolysis (Joseph et al. 2008).

\[
\text{Triglyceride} \quad \xrightleftharpoons{\pm H_2O} \quad \text{Diglyceride} \quad + \quad \text{Fatty acid} \quad + \quad \xrightleftharpoons{\pm H_2O} \quad \text{Monoglyceride} \quad + \quad \text{Fatty acid} \quad + \quad \xrightleftharpoons{\pm H_2O} \quad \text{Fatty acid} \quad + \quad \text{Glycerol}
\]

*Fig. 4. Lipase catalyzed de-esterification reaction.*

Generally, bacterial lipases are glycoproteins but some extracellular bacterial lipases are lipoproteins. The production of extracellular lipase from bacteria is usually dependent on the carbon and nitrogen source, metal salt, chemical reagents, presence of lipids or oil, availability of oxygen, and temperature. The genera of bacteria are *Streptomyces* sp., *Achromobacter* sp., *Alcaligenes* sp., *Arthrobacter* sp., *Pseudomonas* sp. Other than bacteria, many researchers have exploited fungi as valuable sources of lipase due to the following properties: thermal stability, pH stability, substrate specificity and activity in organic solvents. The fungi that can produce commercial lipase are *Aspegillus niger*, *A. terrus*, *A. carneus*, *Candida cylindracea*, etc.

Lipase from bacteria and fungi are most commonly used for transesterification, and optimal parameters for the use of a specific lipase depend on the origin as well as the formulation of lipase. Based on the result of existing research, lipase from different sources has different properties suitable for the process. Thus, there has been a search for an ideal enzyme (Chen et al. 2009), with screened lipase producing from Candida sp. 99-125 from sewage water in north China has a very high conversion rate in lipase-catalyzed esterification and hydrolysis after several induced mutations. Sangeetha et al. (2011) isolated and screened a total of 360 strains of lipase producing bacteria, yeasts and fungi from the samples of oil-contaminated soil and waste water. Among all the screened microbes, the potential lipolytic bacterium, Staphylococcus warneri, unicellular yeast, Candida rugosa and filamentous fungus, Fusarium solani were selected because of their high specific activities. Table 1 enlists some recently screened lipase producing microorganisms with their substrate and optimum parameters.

Table 1. Different sources of microbial lipase.

<table>
<thead>
<tr>
<th>Type</th>
<th>Source</th>
<th>Substrate</th>
<th>Parameter Activity</th>
<th>Activity</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>Pseudomonas aeruginosa</td>
<td>Olive oil</td>
<td>T = 45°C, pH = 8, max activity = 0.760 U/ml</td>
<td>Mobarak-Qamsari (2011)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Burkholderia multivorans</td>
<td>Olive oil</td>
<td>T = 30°C, pH = 9, max activity = 33 U mg⁻¹ protein</td>
<td>Gupta et al. (2005)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Staphylococcus warneri</td>
<td>Palm oil</td>
<td>T = 40°C, pH = 8, max activity = 1.20 U/mg</td>
<td>Sangeetha et al. (2011)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Burkholderia cepacia</td>
<td>Palm oil</td>
<td>T = 45°C, pH = 7, max activity = 160 U/mg</td>
<td>Rathi et al. (2002)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Pseudomonas sp.</td>
<td>Groundnut oil</td>
<td>T = 37°C, pH = 7, max activity = 0.54 U/ml</td>
<td>Ghosh et al. (2005)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Pseudomonas aeruginosa</td>
<td>Olive oil</td>
<td>T = 35°C, pH = 9, max activity = 1098 U/ml</td>
<td>Bisht et al. (2012)</td>
<td></td>
</tr>
<tr>
<td>Fungi</td>
<td>Fusarium oxysporum</td>
<td>Olive oil</td>
<td>T = 50°C, pH = 8, max activity = 60 U</td>
<td>Prazeres et al. (2006)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Rhizopus oryzae</td>
<td>Palm oil</td>
<td>T = 55°C, pH = 8 - 8.5</td>
<td>Kharrat et al. (2011)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Fusarium Solani</td>
<td>Palm oil</td>
<td>T = 40°C, pH = 8, max activity = 1.48 U/ml</td>
<td>Sangeetha et al. (2011)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Geotrichum sp.</td>
<td>Waste cooking oil</td>
<td>T = 45 - 50°C, pH = 8</td>
<td>Kharrat et al. (2011)</td>
<td></td>
</tr>
<tr>
<td>Yeasts</td>
<td>Aspergillus fumigates</td>
<td>Olive oil</td>
<td>T = 30°C, pH = 8.5 - 10</td>
<td>Rajan et al. (2011)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Candida rugosa</td>
<td>Olive oil</td>
<td>T = 50°C, pH = 4 - 9, max activity = 430 U/g</td>
<td>Minovska et al. (2005)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Candida sp.</td>
<td>Waste cooking oil</td>
<td>T = 50°C, max activity = 222.5 U/mg</td>
<td>Chen et al. (2009)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Candida cylindracea</td>
<td>Palm oil mill effluent (POME)</td>
<td>T = 30°C, pH = 6.8, max activity = 4.02 U/ml</td>
<td>Salihu et al. (2011)</td>
<td></td>
</tr>
</tbody>
</table>
For the production of enough lipase the media can be optimized by using palm oil as an inducer and lipase activities for both hydrolytic and synthetic catalysis can be compared. Candida rugosa lipase, which exhibited the highest potential for catalyzing the biodiesel production, was further purified and immobilization on various hydrophobic support materials and was found to be the most promising for further development as a biocatalyst for biodiesel synthesis.

Lipase transesterification of triglycerides with an alcohol (alcoholysis) involves a two-step mechanism when looking at a single ester bond. The first step is the hydrolysis of the ester bond and release of the alcohol moiety followed by an esterification with the second substrate. The two steps are represented in equation (1) and (2) (Fjerbaek et al. 2009, Gog et al. 2012).

\[ E + Es \leftrightarrow E \cdot Es \leftrightarrow F \cdot Bp \leftrightarrow F + Bp \]  

(1)

Followed by

\[ F + As \leftrightarrow F \cdot As \leftrightarrow E \cdot Es_p \leftrightarrow E + Es_p \]  

(2)

Subscripts s and p indicate substrate and product, respectively. For biodiesel, \( As = \) alcohol substrate, \( Bp = \) product with an alcohol moiety (di or monoglyceride or glycerol), \( E = \) free enzyme, \( Es = \) ester substrate (triglycerides) \( Es_p = \) FAAE, \( F = \) fatty acid (Fjerbaek et al. 2009).

This mechanism conforms to a ping-pong bi bi mechanism as each product is released between additions of the substrate and is the widely accepted mechanism for alcoholysis of triglycerides, although simplifications such as Michaelis-Menten kinetics are applied when fitting to experimental result. An example of an initial rate equation for a ping pong bi bi mechanism can be seen in equation (3)

\[ v_i = \frac{V_{max} [TG][A]}{K_{m,TG} [A] (1 + \frac{K_{i,A}}{[TG]} + \frac{K_{m,A}}{[TG]} + [TG][A])} \]  

(3)

Where \( v_i = \) initial rate; \( V_{max} = \) kinetic constants; \( K_{m,TG} = \) concentration of triglycerides and acyl acceptor, respectively.

In order to have full image of the kinetics of enzymatic alcoholysis of triglycerides, other parameters must also be included such as lipase type, amount of reactant, mass transfer limitations, presence of organic solvent, formation and conversion of intermediates, the temperatures influence on enzyme deactivation or the equilibrium limitation for conversion. Thus, when trying to evaluate or determine the kinetics in such systems all these aspects become important (Gog et al. 2012). In general, lipases perform their catalytic activity in more gentle condition and with a variety of triglyceride substrate, including waste oils and fats with high level of FFA. Furthermore, biodiesel separation and purification is much easier, resulting in a more environmentally friendly process. Fig. 5 shows the transesterification process using lipase as catalyst.
Immobilized lipase in support material

Use of enzymes as industrial catalysts serve to be beneficial if the whole process is economical and the cost of any process involve the production of the biocatalyst also. Hence recovery of the catalyst for repeated use becomes necessary. Free enzymes are labile and vulnerable to degradation during the process of recovery of the used enzyme. Also, most lipases exhibit low stability and activity in organic media (Lee et al. 2006). The disadvantage could be overcome by the use of immobilization of enzyme. Immobilization improves the stability of the enzyme under the reaction conditions, enhances enzyme activity thus, makes the repeated use of the enzyme feasible, permits the use of enzyme for diverse applications and thus lowers the production cost (Sangeetha et al. 2011). Immobilization provides a better environment for enzyme to act and also offers better product recovery (Lee et al. 2006).

Enzyme immobilization methods are classified as chemical or physical. Chemical methods involve the formation of covalent bonds between functional groups on the enzyme. Chemical methods are sub classified as either non-polymerizing or cross-linking methods. Non-polymerizing methods involve the formation of both enzyme-support bonds only between enzyme and support, but not between individual enzyme molecules, while cross-linking methods allow the formation of both enzyme-support bonds as well as enzyme-enzyme cross-links (Mikkelsen and Corton 2004, Twyman 2005).

Physical immobilization methods do not involve covalent bond formation with the enzyme, so that the native composition of the enzyme remains unaltered. Physical immobilization methods are sub classified as adsorption, entrapment, or encapsulation methods. Adsorption of protein to the surface of a carrier is, in principle, reversible, but careful selection of the carrier material and the immobilization conditions can render desorption negligible. Entrapment of enzyme in a cross-linking polymer is accomplished by carrying out the polymerization reaction in the presence of enzyme; the enzyme becomes trapped in interstitial spaces in the polymer matrices (Winayanuwattikun et al. 2005, Yagiz et al. 2007, Meunier and Legge 2010). Encapsulation of enzymes results in regions of high enzyme concentration is being separated from the bulk solvent system by a semi-permeable membrane, through which substrate, but not enzyme, may diffuse (Li et al. 2011).
The new method of enzyme immobilization should be able to provide high enzyme loading (close to that of carrier-free enzyme), high retention of activity, and broad reactor configuration. The development of carrier with a predetermined chemical and physical nature, especially suitable geometric properties and binding chemistry, which can bind (or hold) enzyme directly under mild conditions and thus can be used in different reactor configurations. Lipase from different sources has been investigated for their transesterification activity on different support material in Table 2.

**Table 2.** Yield in transesterification reaction using various support materials and different lipases.

<table>
<thead>
<tr>
<th>Support material</th>
<th>Alcohol</th>
<th>Source of enzyme</th>
<th>Feedstock</th>
<th>Yield (%)</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Polyacrylonitrile (PAN) nanofibrous</td>
<td>Methanol</td>
<td><em>P. cepacia</em></td>
<td>Soybean Oil</td>
<td>90</td>
<td>Winayanuwattikun et al. (2008)</td>
</tr>
<tr>
<td>Textile cloth with co-fixing agents</td>
<td>Methanol</td>
<td>Candida’s lipase</td>
<td>Waste cooking Oil</td>
<td>91.08</td>
<td>Chen et al. (2009)</td>
</tr>
<tr>
<td>Hydro calcite</td>
<td>Methanol</td>
<td>Lipzyme-TL IM</td>
<td>Waste cooking Oil</td>
<td>92.8</td>
<td>Yagiz et al. (2007)</td>
</tr>
<tr>
<td>Microporous polymeric matrix (MPPM)</td>
<td>Methanol</td>
<td>Thermomyces lanuginosus</td>
<td>Sun flower Waste cooking oil</td>
<td>97.0  90.2</td>
<td>Dizge et al. (2009)</td>
</tr>
<tr>
<td>Sepabeads EC-OD</td>
<td>Methanol</td>
<td><em>C. rugosa</em></td>
<td>Palm oil</td>
<td>70.0</td>
<td>Sangeetha et al. (2011)</td>
</tr>
<tr>
<td>Chitosan-Glu</td>
<td>Ethanol</td>
<td><em>C. antarctica B</em></td>
<td>Oleic acid</td>
<td>75.0</td>
<td>Foresti and Ferreira (2007)</td>
</tr>
<tr>
<td>Polyurethane Foam</td>
<td>n-hexane</td>
<td><em>C. rugosa</em></td>
<td>Oleic acid</td>
<td>80.0</td>
<td>Awang et al. (2007)</td>
</tr>
<tr>
<td>Zeolite (delaminated zeolite-ITQ) Silicate-1(S-1)</td>
<td>Methanol</td>
<td><em>Rhizomucor miehei</em></td>
<td>Olive oil</td>
<td>92.0  99.0</td>
<td>MacAvio et al. (2007)</td>
</tr>
<tr>
<td>Phyllosilicate sol-gel matrix</td>
<td>Methanol</td>
<td><em>P. cepacia</em></td>
<td>Palm Menhaden Corn Grease</td>
<td>65.0  80.0  71.0  78.0  62.0  88.0  83.0  89.0</td>
<td>Winayanuwattikun et al. (2008)</td>
</tr>
<tr>
<td>Protein-coated micro-crystals (PCMCs)</td>
<td>Tert-butyl alcohol</td>
<td><em>Geotrichum sp. lipase</em></td>
<td>Waste cooking oil</td>
<td>64.0</td>
<td>Yan et al. (2011)</td>
</tr>
<tr>
<td>Ceramic beads</td>
<td>Methanol</td>
<td><em>P. cepacia</em></td>
<td>Waste cooking Oil</td>
<td>40.0</td>
<td>Al-Zuhair et al. (2009)</td>
</tr>
<tr>
<td>SiO₂-PVA</td>
<td>Ethanol</td>
<td><em>Burkholderia cepacia</em></td>
<td>Babassu oil Beef tallow Babassu oil Beef tallow</td>
<td>100.0  89.70  74.13  40.20</td>
<td>Da Rós et al. (2010)</td>
</tr>
<tr>
<td>Nb₂O₅</td>
<td>Ethanol</td>
<td><em>Burkholderia cepacia</em></td>
<td>Babassu oil Beef tallow Babassu oil Beef tallow</td>
<td>100.0  89.70  74.13  40.20</td>
<td>Da Rós et al. (2010)</td>
</tr>
</tbody>
</table>

**Nanoparticles as support material in immobilized lipase**

Currently, nanosized magnetic particles used widely in the immobilization of enzyme have received considerable attention. Based on the research of Lee et al. (2007), magnetic Fe₂O₄ nanoparticles treated
with (3-aminopropyl) triethoxysilane were used as immobilization material. The lipase from *T. lanuginosa* was covalently bound to the amino-functionalized magnetic nanoparticles by using glutaraldehyde as a coupling reagent with the activity recovery up to 70% and the enzyme binding efficiency of 84%. The optimal condition for immobilized lipase was dependent on the immobilization time, temperature, the concentration of glutaraldehyde, and the ratio of lipase to magnetic carrier.

Table 3 shows the nanoparticles used as support material in transesterification. Enzymes immobilized on support materials could catalyze the transesterification of vegetable oils with over 90% conversion to biodiesel being achieved.

**Table 3.** The nanoparticles used as support material in transesterification.

<table>
<thead>
<tr>
<th>Nanoparticle utilized</th>
<th>Material loaded</th>
<th>Feedstock</th>
<th>Parameter process</th>
<th>Yield</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fe₃O₄ nanoparticles</td>
<td>Lipase from <em>T. lanuginosa</em></td>
<td>Soybean oil</td>
<td>50°C, 30 h, (M:O = 1:1), 40% catalyst</td>
<td>90%</td>
<td>Xie and Ma (2009)</td>
</tr>
<tr>
<td>Fe₃O₄ magnetic nanoparticles</td>
<td><em>T. lanuginosa</em></td>
<td>Soybean oil</td>
<td>45°C, 8 h, (M:O = 1.5:1), 3% catalyst</td>
<td>94 %</td>
<td>Xie and Ma (2010)</td>
</tr>
<tr>
<td>Magnetic nanoparticles</td>
<td><em>Pseudomonas cepacia</em></td>
<td>Soybean oil</td>
<td>40°C, (M:O = 3:1), 72 h</td>
<td>93%</td>
<td>Mak et al. (2009)</td>
</tr>
<tr>
<td>Fe₃O₄ nanoparticles bio composite</td>
<td><em>Pseudomonas cepacia</em></td>
<td>Soybean oil</td>
<td>40°C, (0.2M M/H + 6:1:0.2), 24 h, 6.6g catalyst</td>
<td>&gt;99%</td>
<td>Wang et al. (2011)</td>
</tr>
<tr>
<td>Ferric silica nanocomposite</td>
<td>Bulkholderia sp.</td>
<td>Olive oil</td>
<td>40°C, (M:O = 4:1), 30 h, 11%</td>
<td>92%</td>
<td>Tran et al. (2012a)</td>
</tr>
<tr>
<td>Ferric silica nanocomposite</td>
<td>Bulkholderia sp.</td>
<td>Microalgal oil</td>
<td>40°C, (M:O = 61:75), 48 h, 1203.1 U g⁻¹</td>
<td>97.25%</td>
<td>Tran et al. (2012b)</td>
</tr>
</tbody>
</table>

Note: M = methanol, O = oil, DW = distilled water, H = n-hexane

Immobilization of lipase as a catalyst has a great potential for achieving the design and operation of enzymatic biodiesel production on the industrial scale. By using a packed-bed reactor system with lipase-Fe₃O₄ nanoparticles bio composite catalyst was successfully developed for biodiesel production. The nanoparticles bio composite showed elevated activity and stability in the four-packed-bed reactor with conversion of biodiesel was maintained at the high rate of over 88% for 192 h. The efficient reuse of the enzyme was realized via a simple and effective immobilization procedure that resulted in a high initial activity without inactivation or inhibitor. The packed-bed reactor system has a great potential for achieving the design and operation of enzymatic biodiesel production on the industrial scale (Wang et al. 2011).

In another study, porcine pancreas lipase was covalently immobilized on the surface of silica-coated magnetite nanoparticles. The diameter of silica-coated magnetite was about 17 nm, and the immobilization process did not change the phase of Fe₃O₄. The results showed that the covalent immobilization of lipase on support material improved the thermal, pH and storage stability. Moreover, kinetic study showed the activation of immobilized enzyme. The enzyme recovery represents the establishment of about 64% of residual activity after six cycles of washing (Ranjbakhsh et al. 2012).
The lipase-coated magnetic nanostructures were applied in a reactive extraction process that allowed separation of the products formed during transesterification. It is expected that reactive extraction can directly produce 77% ethyl oleic (biodiesel) using lower ethanol/triolein ratios compared to 35 - 40% purity in the normal stirred reactor with a higher ethanol/triolein ratio. This approach implies a novel and efficient location and use of lipase in column reactors for production of biodiesel (Dussan et al. 2010).

**Physicochemical properties of nanomaterial and immobilized lipase**

Immobilization is normally considered to be an important method to improve the stability of enzyme. The morphology of nanoparticles observed using SEM micrograph of coated magnetic particle showed agglomerations because the non-coated magnetic particles were not dispersed in an appropriate substance (Dussan et al. 2010). The pure magnetite was observed to be spherical with nano size (Xie and Wang 2012). The particles diameter is an important factor for support material. Smaller particles have larger surface-to-volume ratios and larger capacity to bind more substance on their surface and product will give less restriction for diffusion.

BET analysis for immobilized *Burkholderia* sp. shows that the surface area of pure magnetite, silica magnetite, silica-magnetite nanocomposite and alkyl grafted silica-magnetite nanocomposites has a different surface area. Silica-magnetite nanocomposite have a surface area (202 m²g⁻¹) and after grafting with the alkyl group at the surface of Fe₃O₄-SiO₂, the surface area decreasing (128 m²g⁻²). That would be due to the alkyl group may enter the Fe₃O₄-SiO₂ pores, thereby decreasing surface area and smaller pore size (Tran et al. 2012a).

The hydrolytic activity of free and immobilized lipase were measured at various temperatures (35 - 65 °C). The activity of bound and free lipase showed the highest activity at approximately 45°C. However, the immobilized lipase has higher stability than free lipase. The relative activity of immobilized lipase is 82% at 55 °C. The optimum hydrolytic activity was observed at pH 7.0 for both lipase. It indicated that immobilization did not change the activity of lipase. Immobilized lipase retained activity when the pH was higher than pH optimum. Immobilization method can improve the pH stability of the lipase, until pH 8.5 with the relative activity 60% (Xie and Ma 2010).

Furthermore, immobilized lipase can improve the storage stability and catalyst recycling than free lipase. The immobilized lipase conserved more than 64% of its activity after 21 days while free lipase only 47%. Also in reusing catalyst after 6 cycles, immobilized lipase retained 63% of its initial activity (Ranjbakhsh et al. 2012).

**Kinetic parameters**

Kinetic parameters of free and immobilized lipase were determined from Lineweaver-Burk plots. The measurement of Michaelis-Menten parameters also revealed a considerable improvement of the immobilized lipase. In a study (Ranjbakhsh et al. 2012), Kₘ value of immobilized lipase was lower than free lipase, which represents a higher affinity of immobilized lipase to substrates. Certainly with the nano size, magnetic nanoparticles could be imagined to offer lipase molecules a porous surface with a better orientation leading to higher affinity for substrate and more available sites. The result also demonstrated an increase in Vₘₐₓ due to immobilization of lipase. The improvement of Vₘₐₓ may also be due to more efficient conformation of
immobilized lipase with respect to free lipase. The improvement of kinetic parameters of immobilized lipase can be a good feature for possible industrial application.

Kinetic study showed a dependence of alkyl-grafted-Fe₃O₄-SiO₂-lipase on the substrate, the $V_{\text{max}}$ and $K_m$ values were estimated at 6251 U g⁻¹ (132.4 U mg⁻¹ protein) and 3.65 mM, respectively. The $K_m$ value of alkyl-grafted-Fe₃O₄-SiO₂-lipase is higher than free lipase but smaller than celite-lipase. The $V_{\text{max}}$ of alkyl-grafted-Fe₃O₄-SiO₂-lipase is smaller than free lipase and larger than celite-lipase. It indicated that the structure of enzymes could be rigidified on the surface of celite, thus the blocking of the active site of lipase would probably decrease lipase activity. However, with alkyl-grafted-Fe₃O₄-SiO₂-lipase, the binding of lipase on the surface of nanocomposite is multipoint hydrophobic interaction, which may cause the same phenomena as multipoint covalent bonding. The maximum reaction of alkyl-grafted-Fe₃O₄-SiO₂-lipase is much higher than celite lipase. This result indicates that hydrophobic interaction is the better approach for immobilization of the lipase (Tran et al. 2012a).

Conclusion

Nanocrystals and nanomaterials are serving as a novel supports for a catalyst in transesterification reaction. Currently, the use of nanoparticles has emerged as a versatile tool for generating excellent material for the catalyst due to their small size and large surface area and high catalytic activity. Nanoparticles strongly influence the mechanical properties in the material. The nano-magnetic biocatalyst also exhibits as a good catalytic property. Recently, immobilized lipase on nano-magnetic support showed high catalytic activity and advantages of easy separation and reuse. Moreover, nanocomposite grafted with a long alkyl group in order to create affinity for lipase bound on the surface of nanoparticles are shown to be a good matrix for lipase immobilization with high yield of biodiesel and high reusability. Support nanomaterials were prepared by co-precipitation method and also creating a functional group is one of the new trends in immobilization of lipase. Nanocatalysts is a potential material for transesterification reaction and for commercial applications.

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References


Trend In Enzyme Immobilization


