

<u>ชัญวลัย จิ<mark>1</mark></u>ชีพิ,นิรัพญเฆ่งค์ ดัง ¹ณุวิทยา ปั้นสูวรรณ^{1,3}

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งานวิจัยนี้ เป็นการที่กษาการผลิตไบโอพีเซลจากไขมันในน้ำเสียของโรงงานอุดสาหกรรมผลิตและแปรรูปอาหารทะเล ที่มีปริมาณกรดไขมันอิสระประกอบอยู่สูง ถึง 70-85 เปอร์เซ็นค์ โดยใช้กระบวนการผลิตไบโอพีเซลแบบสองนั้นดอน ขั้นดอน แรกใช้กระบวนการเอสเทอริฟิเกชั่นโดยใช้กรดชัลฟีวริก เป็นด้วยงปฏิกิริยาระหว่างกรด ใพมันอิสระ กับเมทานอล จนปริมาณกรด ใพมันอิสระมีก่าด่ากว่า 2 เปอร์เซ็นด์ และขั้นดอนที่สองคือกระบวนการทรานส์เอสเทอริฟิเกชั่น โดยนำผลิตภัณฑ์ จากขั้นคอนแรกมาทำปฏิกิริยากับเมทานอลค่อ โดยใช้ไพแทสเซียมไฮครอกไซค์ เป็นตัวเร่งปฏิกิริยา และได้ผลิตภัณฑ์เป็นเมทิลเอสเทอร์ จากการทดลองพบว่า ขั้นตอนแรกของกระบวนการเอสเทอริพิเคชั่น ใช้อัตราส่วนโดยไมลของ เมทานอลค่อ ไขมันเท่ากับ 10: 1 ความเข้มข้นของกรดข้อฟัววิกเท่ากับ 3 เปอร์เซ็มด์โดยเทียบกับน้ำหนักไขมัน และเวลาในการทำปฏิกิริยาเท่ากับ 240นาที ส่วนในขั้นตอนที่ 2 ใช้กระบวนการทรานส์เอสเทอริพิเคชั่น โดยใช้ไทแทสเซียมไฮครอกไซด์เป็นด้วเร่ง ปฏิธีริยาที่ความเข้มข้น 1.5 เปอร์เซ็นต์โดยน้ำหนัก อัพราส่วนโดยโมลของเมทานอลด่อผลิตภัณฑ์ที่ได้จากขั้นดอนที่ 1 เท่ากับ 6::1 ระขะเวลาในการทำปฏิธีริยาเท่ากับ 30 นาที สามารถสังเคราะห์ใบไอดีเซลได้บริสุทธิ์ สูงสุด 67.0 เปอร์เซ็นต์ ไบโอดีเซลที่ได้เมื่อทดสอบคุณสมบัติดามมาตรฐานไบโอดีเซลชุมชนพบว่าผ่านเกณท์ดามมาตรฐาน ASTM ที่กำหนดไว้

Abstract:

ADSTRACE: This research was to study the waste fat in waste water from sea food processing plants as the raw material for biodiesel production. Due to high free fatty acid content(about 70-85%), two-step process reaction was required. The first step was esterification using sulfuric acid as a catalyst to reduce free fatty acid from 70-85 percent to lower than 2 percent. In the second step, transesterification, potassium hydroxide was applied as a catalyst in the reaction between the product from the first step and methanol. The optimum conditions for the esterification step were 10:1 molar ratio of methanol to oil, 3 percent of sulfuric acid by weight on fat and 240 minutes of reaction time. The optimum conditions for the transesterification step were 1.5 percent by weight of catalyst, 6:1 molar ratio of methanol to the product from the esterification step and 30 minutes of reaction time. The application were 1.5 percent by weight of catalyst, 6:1 molar ratio of methanol to the product from the esterification step and 30 minutes of reaction time. The maximum purity of biodiesel was 67.0 percent. The quality of biodiesel passed the standard community biodiesel follow ASTM standard.

Introduction: Biodiesel is a renewable fuel. It is produced from high quality food grade vegetable oil which makes biodiesel more expensive than diesel fuel. Therefore, a low cost feedstock is needed. The waste fat from a grease tap of a waste water treatment plant for marine production plants is considered as an available feedstock to produce biodiesel in this study. Due to high free fatty acid (FFA) content in waste fat, alkali is used as a catalyst in transesterification to produce biodiesel, which gives low biodiesel yield because FFA reacts with alkali to form soap, resulting emulsification and separation problems. To resolve these problems, acid catalyst has been as been as perteratment step. Sulfuric acid is the most commonly used catalyst. The optimization of pretreatment reaction was investigated with sulfuric acid dosage to reduce FFA content to below 2%. Alcohol molar ratio and reaction gives a retreatment step. Sulfuric acid is reaction is carried out with an alkali catalyst to investigate the optimum dosage of potassium hydroxide, methanol molar ratio and reaction time. Biodiesel quality will be evaluated according to ASTM standard for community biodiesel.



Conclusion

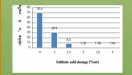
Conclusion : Waste fat from waste water of marine products processing plant can be a possible feedstock for production of community biodiesel. Acid esterification followed by alkaline transesterification was necessary as the waste fat has been found to possess high free fatty acid content of 70.14%. Molar ratio of alcohol to oil of 10:1, 3% w/w of sulfuric acid, reaction time of 4 hr. at 60°C and 500 rpm of stirrer speed were optimum conditions to bring down the free fatty acid to below 2 %. The optimum conditions of alkaline transesterification process were 1.5% of potassium hydroxide as catalyst, molar ratio of methanol to the product from the esterification step of 6:1, reaction time of 30 min. at 60°C temperature and 500 rpm of stirrer speed. The final product bears viscosity of 5.81 est at 40°C, acid value of 0.8 mg KOH/g, flash point 164°C and water and sediment 0.05% which conform to ASTM standard of community biodiesel

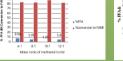
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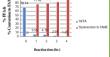
caracteristic and J.V. Gerpen. 2001. A pilot plant to produce biodiesel from high free fatty acid feedstocks. ASAE an ternational Meeting presented as paper no.016049 erpen, J.V. 2005. Biodiesel processing and production. Fuel Processing Technology. 86:1097-1107

Results and Discussion :

The initial content of FFA of the waste fat in this study was 70.14%. Esterification process ,the FFA content decreases with increasing sulfuric acid dosage ,as shown in Figure 1. The reaction was controlled at a constant stirring speed of 500 rpm., a constant temperature of 60 °C, a reaction time of 4 hr. and molar ratio of methanol to fat 10:1. The optimum dosage of sulfuric acid was found at 3.0 % www. Different molar ratios, 6:1, 8:1, 10:1 and 12:1 were tried for acid sprimm uosge of summa also way found also way way indicate model matter model matters of other than the fore the determinant setterminant of the setterminant and the setterminant model in the setterminant and the setterminant %conversion as shown in Figure 2. When varied the reaction time, the optimum reaction time for the waste fat in this study was 4 hr.The FFA was decreased to 1.25% and get maximum %conversion to FAME at 88.44% as shown in Figure 3.The summarized result of optimum conditions of the esterification step for the waste fat in this study were 3.0% ww of sulfuric acid, 10:1 of molar ration and 4 hr. of .reaction time. This condition can decrease %FFA from 70.14% to 1.25% and get maximum %conversion to FAME at 88.44%.



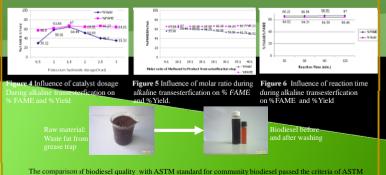




Figuer 1 Influence of catalyst dosage during acid esterfication on FFA(%) Figure 2 Influence of molar ratio of alcohol to fatty acids during acid esterification on %FFA and %conversion toFAME

Figure 3 Influence of reaction tin during acid esterification on %FFA and %conversion to FAME

Alkaline catalyzed transesterification process was further studied by using a pretreated product from the fication step that had an FFA content of 1.25%. Potassium hydroxide was used as catalyst at different dosages range 3.0%, molar ratios of methanol to the product from the esterification step of 6:1, 10:1, 15:1, 20:1, 25:1, 35:1 and 40:1. different reaction time of 30-120 minutes. The result showed that catalyst dosage of 1.5% w/w gave maximum FAME as shown in Figure 4. The further experiment with molar ratio as shown in Figure 5. It show that the molar was not a significant effect on %FAME (2.6% difference). The effect of different reaction time of 30-120 min, was not a significant effect on %FAME as shown in Figure 6. The summary optimum conditions of transesterification ess were 1.5 % of catalyst dosage, 6:1 of molar ratio of methanol to the product from the esterification step, reaction of 30 min at temperature of 60°C and the reaction stirring speed of 500 rpm. The color of biodiesel turned from dark to neolor after transesterification step.



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Table 1 Properties of biodiesel from waste fat of marine products processing plant compared to ASTM standard munity biodiesel.

Properties	Unit	Biodiesel from waste f	at ASTM criteria
Kinematic viscosity (40°	C) mm ² s ⁻² (cst)	5.81	1.9-8.0
Acid number	mgKOHg-1	0.80	<0.8
Flash point (closed cup)	°C	164	<120
Water and sediment	%volume	0.05	< 0.2
Total glycerine	%wt	0.08 <1.5	0