Feedstocks
Most homebrewed biodiesel is made from waste vegetable oil (WVO) collected from restaurants or from raw pressed vegetable oil. The WVO often contains water and high levels of free fatty acids as well as solids and other contaminants. The solids can be filtered out, but the water must be removed by heating the oil to above 212°F for an extended period (several hours) to drive the water from the oil. WVO typically contains from 2 to 5% free fatty acids.

If the free fatty acid (FFA) level is above this amount, it is difficult to produce biodiesel without an acid esterification process first that most homebrewers do not have the capability to implement. The FFA level is determined by a titration process. A full description of this process is provided in TechNote #33.

Raw vegetable oils that are produced by pressing oilseeds such as canola, mustard, or soybeans can be converted to biodiesel. The FFA level of these raw oils is usually 0.5% or less, which is low enough not to pose a problem. However, the oils may contain gum compounds that can create sludge deposits in the processing equipment and can make it more difficult to separate the glycerin at the end of the reaction. The easiest way for homebrewers to reduce the gum level in their oil is to let the oil sit in a tank for 2 to 3 weeks before converting it to biodiesel.

The oil will undergo natural degumming and reach a low enough gum level that conversion to biodiesel is not a problem. If some gums remain in the oil, they will generally be removed with the glycerin or during the wash process.

The common recipe for biodiesel production is to combine 100 pounds of oil with 22 pounds of methanol and 0.5 to 1.0 pounds of sodium or potassium hydroxide. Additional sodium or potassium hydroxide is added to neutralize the free fatty acids if present, based on the titration described above. Generally, the catalyst is dissolved in the alcohol and then the mixture is added to the oil while stirring.

This mixture is heated to about 140°F and stirred for one to two hours. A frequent mistake of homebrewers is to add too little methanol or not allow sufficient time. Reports of as little as 13 to 15% methanol (compared with the 22% recommended here) are heard frequently. This is unlikely to produce a good quality biodiesel. Although it can be difficult for homebrewers to confirm that their product meets the specifications of the ASTM standard, this is a goal that we recommend they strive to achieve. In addition to
adding enough alcohol, it is important for homebrewers to provide enough agitation, enough temperature, and enough time for the reaction to reach completion. We recommend that the reaction be allowed 1-2 hours at 140°F, 2-4 hours at 105°F, or 4-8 hours at 70°F. The oil, alcohol, and catalyst should be stirred vigorously throughout the reaction period.

One approach that is usually successful is to do the reaction in two steps. This involves adding 80% of the methanol and catalyst in the first reaction step, separating and removing any glycerin that has formed, and then adding the remaining 20% of the methanol and catalyst for a second reaction.

At the end of the reaction period, the biodiesel should be allowed to settle for several hours. Overnight is even better. A clearly defined line should exist between the biodiesel layer on top and the darker glycerin layer on the bottom. The glycerin can be drawn off leaving biodiesel that is ready for washing. A common observation by homebrewers is that instead of a sharp separation between the biodiesel and glycerin, they end up with a semi-solid, gelatinous mixture that cannot be used as fuel. This is an indication that the oil contained excessive free fatty acids and/or water.

The glycerin by-product is usually a disposal problem for small producers. It may contain up to 40% methanol so it is flammable, and releases methanol to the atmosphere. The methanol is an air pollutant and a potential ground water contaminant if the glycerin is spread on the ground. Methanol is also toxic to human beings and animals. A possible answer is for several homebrewers to combine their glycerin so there is enough to justify the expense of a methanol recovery system. A cooperative of small producers would be a good way to do this. After the methanol has been removed, the glycerin can be used as a composting accelerant, fed to animals, or further refined for other uses. A good way to dispose of glycerin is to add it to an anaerobic digester that produces methane gas from animal waste.

Washing consists of adding an amount of water equal to about 50 to 100% of the biodiesel volume to the biodiesel and gently agitating the mixture. Then, the water is allowed to settle so it can be separated. The process is repeated until the water is clear when it is removed. A good quality reaction can usually be washed in 3 to 4 steps.

Disposal of the wash water is a similar problem to glycerin disposal for small producers. It can probably be put into a municipal sewer system although before doing this, you should visit them and explain what you intend to do.

After the biodiesel has been washed, it may be cloudy due to small water droplets suspended in the fuel. If the fuel is allowed to sit for several days, these will settle out. Or, if the fuel is heated to 160-210°F, it will become clear. At this point, the fuel should be ready to use, either as a pure fuel, or in blends with petroleum diesel fuel.

Some homebrewers have argued that washing is optional. We always recommend washing biodiesel because of the presence of soaps, residual free glycerin, catalyst, and methanol that remain in the biodiesel. These are easily removed with water washing and a very pure product will result.

Safety
While the oils and fats used to make biodiesel and the biodiesel itself are non-toxic and not very flammable, the methanol and catalyst present several safety concerns. Methanol is very toxic and while ingestion is particularly dangerous, long term inhalation or skin exposure should be avoided. All processing work with methanol should be done in covered containers in well-ventilated areas, preferably out-of-doors. Methanol vapors can be ignited by sparks or exposed flames so use explosion-proof motors for agitation. If these are not available, recirculating air pumps or hydraulic motors can also be used. Methanol burns with an invisible flame so serious burns are possible before you even know there is a fire. Explosions have occurred when tanks that have held methanol are exposed to welding arcs and cutting torches. All used tanks should be filled with water before cutting or welding.

Sodium and potassium hydroxide are strong bases that can cause burns to exposed skin and potentially life-threatening injury if their dust is inhaled. Full chemical protective clothing and gloves along with a respirator are recommended when handling these dangerous materials. Always add the catalyst slowly to the alcohol so that any heat that is generated has a chance to dissipate.

Both sodium and potassium hydroxide will absorb moisture from the air so they should be kept in sealed containers until they are to be used. The alcohol-catalyst mixture should also be protected from contact with air until it is added to the oil.