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## EMULSIFICATION OF WASTE COOKING OILS AND FATTY ACID DISTILLATES AS DIESEL ENGINE FUEL

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### KEYWORDS

Emulsified biofuels, Waste cooking oils, Fatty acid distillates

### ABSTRACT

The scope of this paper is to analyze the possibilities and feasibilities, as well as the main experimental results reported about the emulsification method applied to waste cooking oils and fatty acid distillates as diesel engine fuels, compared with other commonly used methods. These waste products are obtained in the refining oil industry, food industry and service sector from the frying process, mainly. However, they are a little used as feedstocks to produce biofuels and constitute a potential source of contamination. From the review of the state of arts, significant decreases in exhaust emissions of nitrogen oxides, cylinder pressure, as well as increases of the ignition delay, brake specific fuel consumption, hydrocarbon, smoke opacity, carbon monoxide, particulate matters to emulsified waste cooking oils and fatty acid distillates compared with diesel fuel are reported. In some experiments the emulsified waste cooking oils achieved better performance than neat fatty acid distillates, neat waste cooking oils and their derivatives methyl esters.

### INTRODUCTION

Since fossil fuels increase greenhouse gas emissions and cause global warming, the use of alternative resources like biofuels are more pronounced everyday (Ozbay et al., 2008). For this reason, recently much attention has been paid to the development of alternative fuels in order to meet the emission standards and to reduce the dependency on fossil fuel (Kannan and Anand, 2011; Melo et al., 2014), as well as to counteract the recent changes in fossil fuels' prices and their influence on the energy worldwide scenario. In this context, taking into account the biodiversity and the “food vs. fuel” debate in mind (Singh and Singh, 2011), special attention has been paid to feedstocks such as non-edible vegetable oils and waste products (e.g. fatty acid distillates and waste cooking oils).

Although previous researches (Al-Widyan et al., 2002; Dorado et al., 2003; Cetinkaya and Karaosmanoglu, 2005; Pugazhvadivu and Jeyachandran, 2005; Felizardo et al., 2006; Sudhir et al., 2007; Chhetri et al., 2008; Lapueta et al., 2008; Ozsezen et al., 2008; Giraçol et al., 2011; Kalam et al., 2011; Kartina and Suhaila, 2011; Murlidharan and Vasudevan, 2011; Galle et al., 2012; Elshaib et al., 2014; Ferrer and Piloto, 2014; Hirkude et al., 2014; Melo et al., 2014; Piloto et al., 2014; Piloto et al., 2014) using waste cooking oils (WCOs) and fatty acid distillates (FADs) have demonstrated that their use as feedstocks to produce biofuels is possible, their use is not common yet and represent a potential source of contamination. However, with their uses it is possible to reduce their effects on environment due to the dumping on rivers and seas. In addition, their exploitations to produce biofuels and energy might improve the efficiency in the refining oil, food and service industries (restaurants, hotels and food factories), adding values to these byproducts and contributing to the concept of reaching zero-waste.

On the other hand, WCOs and FADs are potential alternatives for diesel engines due to their very similar properties to diesel fuel. However, the direct use of WCOs and FADs might bring several problems in engine performance and emissions. These problems are mainly associated to a lower heating value and cetane number of WCOs compared to diesel fuel. Also, properties as kinematic viscosity and surface tension have a significant influence on the injection process as well as the process of mixture formation in diesel engines (Kruczynski, 2013; Melo et al., 2014).

From the above reasons exposed and to obtain a more engine-friendly fuel, is necessary to change the biofuels properties applying different methods such as: preheating, blending, dual fuel operation, transesterification, cracking/pyrolysis or emulsification (Melo et al., 2014). Among these methods, the emulsification techniques applied to WCOs and FADs

have not been studied thoroughly. The scope of this paper is to analyze the possibilities and feasibilities, though the main experimental results reported about emulsification method applied to WCOs and FADs compared with other commonly used methods.

## WASTE PRODUCTS AS BIOFUEL'S FEEDSTOCKS

One of the most important challenges for the industrial sector dedicated to refining oil, service and food industry is to find solutions to use, reduce or eliminate large quantities of waste products generated from themselves. The waste cooking oils and fatty acid distillates are available around the world and generally have a low commercial value and a little use. The FADs from the refining oil industry are obtained, as is in Fig.1 showed.

Despite the WCOs are used for soap and animal feed productions, in agreement with Chhetri (Chhetri et al., 2008) part of they are discharged into the environment. Also, it is important to take into account that the use of WCO in the production of animal feeds in few countries is prohibited and this has resulted in the availability of surplus quantity WCO (Pugazhvadivu and Jeyachandran, 2005). In addition, the reuse of cooking oils might cause serious difficulties on human health. Due to the high temperatures, carcinogens as benzopyrene are released.

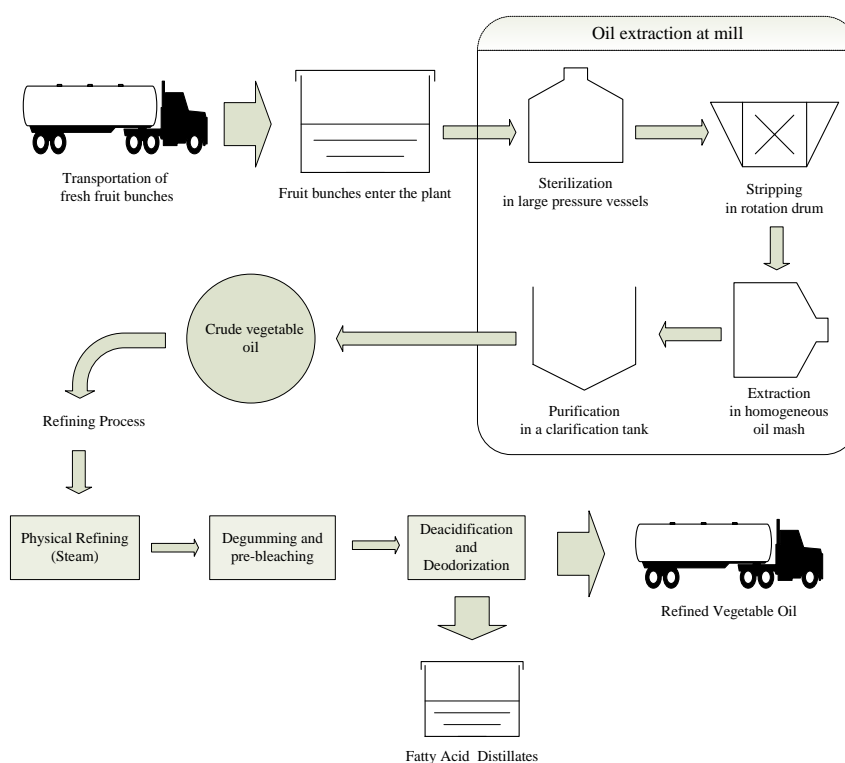


Fig 1: Full processing flow chart for a general vegetable oil physical refining process (Melo et al., 2014; Piloto et al., 2014; Piloto et al., 2014)

According to the composition, WCOs and FADs are suitable for the production of biofuels (see Table 1) increasing the efficiency to the above industries mentioned; contributing to the concept of reaching zero-waste concerning the utilization of by-products generated in the oil refineries (Piloto et al., 2013), reducing the environmental degradation (Pugazhvadivu and Jeyachandran, 2005). In agreement, researchers as Kartina and Suhaila (Kartina and Suhaila, 2011) refer that WCO is the cheapest source and can reduce problems on waste oil disposal whereas FAD is a byproduct from oil refining, therefore can be a readily available feedstock to produce engine biofuels.

However, the direct use of these waste products as engine fuel might affect the engine assessment and components. In order to obtain an economic and environmentally-friendly engine fuel from renewable feedstocks such as vegetable oils and animal fats, it is necessary to change the biofuels' properties (e.g.: viscosity, surface tension, free fatty acid, etc.). For this reason, different methods have been used, such as preheating, blending, dual fuel operation, transesterification, cracking/pyrolysis and emulsification. Among these methods, Demirbas et al. (Demirbas and Fatih, 2010) and Meher (Meher et al., 2006) specified that the transesterification is the most promising solution to the high viscosity problem and is an interesting method to produce a cleaner and environmentally safe fuel.

Transesterification of triglycerides produces biodiesel and proceeds through a reaction with alcohols in the presence of a catalyst and producing glycerol as a co-product (Attaphong, 2012). Biodiesel also has drawbacks, including cold weather limitations due to a relatively higher cloud point and pour point, and might increase emissions of nitrogen oxides (NO<sub>x</sub>) (Attaphong, 2012). Atmanli (Atmanli et al., 2013) and Agarwal (Agarwal and Rajamanoharan, 2009) pointed out that the transesterification process is a relatively expensive chemical process since it involves the use of chemicals, catalysts and a heating process. Also, depending on the quality of the feedstock (free fatty acids, glyceride and moisture content), different steps are necessary.

The free fatty acids (FFA) is hydrolysis/oxidation by-products of oil due to cooking and storage, and monoglyceride and diglycerides are degrade products of oil (Chai et al., 2014). The free fatty acids and glyceride content in fatty acid distillates and waste cooking oils from different feedstocks are shown in Table 2. Due to the high free fatty acids (FFAs) content of WCO and FAD, these sources cannot be converted directly to biodiesel via alkaline transesterification (Kartina and Suhaila, 2011). Although different methods to decrease FFA and glyceride content are reported (Chongkhong et al., 2009; Kombe et al., 2013; Talebian-Kiakalaieh et al., 2013) in order to improve the transesterification efficiency, economically the biodiesel productions are not profitable.

Table 1: Physical-chemical properties of WCO and FAD (Yoshimoto et al., 1999; Kerihuel et al., 2005; Pugazhvadivu and Jeyachandran, 2005; Nanthagopal and Subbarao, 2009; Atmanli et al., 2013; García et al., 2013; Qi et al., 2013; Gandón et al., 2014; Senthil and Jaikumar, 2014)

Properties	WCO	FAD (from Soybean)	Diesel Fuel
Viscosity ( <i>cSt</i> )	33,40-43,36	38,10	3.9-4.6
Density ( <i>g/cm<sup>3</sup></i> )	0,88-0,925	0,923	0.829-0.84
Flash Point ( <i>°C</i> )	210-302	232	67.5-78
Moisture Content (%)	0,2433-0.0693	0,3869	NS
LHV ( <i>MJ/Kg</i> )	36.47-39.60	NS	42.39-43.38
Cetane number	33.4-37	NS	45-60.5
C ( <i>w/w%</i> )	76.8	NS	84-87
H ( <i>w/w%</i> )	11.6	NS	16-33
O ( <i>w/w%</i> )	10.6	NS	0
<b>Fatty acid composition (% wt)</b>			
Myristic	NS	0,106-0,108	---
Palmitic	NS	12,45-12,81	---
Palmitoleic	NS	0,108-0,166	---
Stearic	NS	3,79-3,89	---
Oleic	NS	23,33-23,36	---
Linoleic	NS	51,67-51,94	---
Linolenic	NS	6,34-6,39	---

NS: Not specified

Table 2: Free fatty acids and glyceride content in WCO and FAD from different feedstocks

Type	Feedstocks	FFA ( <i>wt %</i> )	Glycerides ( <i>wt %</i> )	Ref.
FAD	Cotton	85.0	NS	(Keskin et al., 2008)
	Palm	70.0-93.0	20-30	(Chongkhong et al., 2009; Budiman et al., 2012)
	Hazelnut	45-50	NS	(Usta et al., 2005)
	Soybean	30.1-45.4	13.0-23.3	(Hirota et al., 2003; Dumont and Narine, 2008; Gunawan et al., 2008)
	Rapeseed	48.8	32.9	(Liu and Wang 2009)
WCO	NS	5-37.96	54.4-96.2	(Wang et al., 2006; Chhetri et al., 2008; Patil et al., 2012; Wang et al., 2012; Kombe et al., 2013; Chai et al., 2014)

FFA: free fatty acids, FAD: fatty acid distillate, WCO: waste cooking oil, NS: Not specified

## EMULSIFICATION METHOD APPLIED TO WASTE COOKING OILS AND FATTY ACID DISTILLATES

An alternative to the transesterification process may be the use of the emulsification method. Emulsification is the process of dispersing one liquid in a second immiscible liquid using a third substance known as emulsifier. Through this process, a dispersed system is obtained containing small droplets of water suspended in WCO or FAD. Emulsification is a simple process and might need no modification of the original engine design (Senthil et al., 2006).

Emulsions are interesting as fuel in diesel engines due to their simultaneous reduction of smoke and NO<sub>x</sub> emissions using vegetable oils/diesel as fuel (Yoshimoto et al., 1999; Lin and Wang, 2004; Senthil et al., 2006). Additionally, due to the microexplosion phenomenon, it is also possible to recover in some proportion the decrease in combustion efficiency.

Different researches about the formulation of emulsified biofuels using WCO or FAD reported the use of surfactants of the Span series and mainly co-surfactants such as ethanol. However, it is possible to use other surfactants with a hydrophilic-lipophilic balance (HLB) between 4-6 (Debnath et al., 2013), as well as short-chain alcohols as co-surfactants with the aim to increase the water amount and/or to further improve the stability of the emulsified biofuels. The characteristics of some surfactants and co-surfactants used as emulsifier agents are shown in Table 3.

Table 3: Surfactants and co-surfactants used in emulsified biofuels' formulation

Chemical name	Chemical formula	HLB	FP (°C)	CN	Density (g/mL)	Ref.
Sorbitan monopalmitate (Span 40)	C <sub>22</sub> H <sub>42</sub> O <sub>6</sub>	6.7	NS	NS	NS	(Chow and Ho, 1996; Muñoz et al., 2007)
Sorbitan monostearate (Span 60)	C <sub>24</sub> H <sub>46</sub> O <sub>6</sub>	4.7	NS	NS	NS	(Muñoz et al., 2007)
Sorbitan monooleate (Span 80)	C <sub>24</sub> H <sub>44</sub> O <sub>6</sub>	4.3	NS	NS	0.99	(Kerihuel et al., 2005; Kannan and Gounder, 2011; Attaphong and Sabatini, 2013)
Sorbitan sesquiolate (Span 83)	C <sub>66</sub> H <sub>130</sub> O <sub>18</sub>	3.7	NS	NS	0.95	(Kerihuel et al., 2005; Mura et al., 2012; Attaphong and Sabatini, 2013)
Ethanol	C <sub>2</sub> H <sub>5</sub> OH	NS	10.2-13.5	5-8	0.788	(Qi et al., 2010; Tangka, 2011)
Bioethanol	C <sub>2</sub> H <sub>5</sub> OH	NS	13	NS	0.79	(Barroso et al., 2010)

FP: Flash Point, CN: Cetane number, NS: Not specified

Researchers such as Morais et al. (Morais et al., 2006), Porras (Porras et al., 2008) and Bhimani (Bhimani et al., 2013) point out that a mixture of hydrophilic and hydrophobic surfactants yields a more stable emulsion. For this reason, in order to obtain a mixture of surfactants with HLB number according to the interval previously recommended, a mathematical equation given by Mollet (Mollet and Grubenmann, 2001) and Bhimani (Bhimani et al., 2013) can be used. Through this equation, the mass percentage (%) of the surfactants involved in the mixture can be obtained.

$$\% \text{Surfactant A} = [100 \cdot (X - \text{HLB}_B)] \cdot (\text{HLB}_A - \text{HLB}_B)^{-1} \quad (1)$$

$$\% \text{Surfactant B} = 100 - \% \text{Surfactant A} \quad (2)$$

With:

X: HLB required

HLB<sub>A</sub>: hydrophilic-lipophilic balance of the surfactant A

HLB<sub>B</sub>: hydrophilic-lipophilic balance of the surfactant B

An accurate methodology in order to formulate emulsified biofuels is difficult to establish because there are different feedstocks and different methods of preparation such as mechanical stirrer or membrane emulsification. Among these methods, the use of dispersion has been the most applied. Different researchers (Waltra, 1993; Khumar, 1996; Kerihuel et al., 2006) noticed that the higher rotational speed leads to the formation of smaller water droplets.

As was previously mentioned, the emulsification techniques applied to WCOs and FADs have not been studied thoroughly. However, researchers as Yoshimoto, Mubarak, Nanthagopal, Subbarao, Senthil and Jaikumar (Yoshimoto et al., 1999; Nanthagopal and Subbarao, 2009; Mubarak and Senthil, 2012; Senthil and Jaikumar, 2014) formulated emulsified biofuels using waste cooking oil (WCO). Yoshimoto (Yoshimoto et al., 1999) conducted an investigation about the emulsified WCOs, which were discarded from restaurants and households. The emulsified WCO were prepared with different percentage of water (10-40%), 1% of surfactant (CRS-75) and a mixture in equal proportions of WCO and diesel fuel as continuous phase. The kinematic viscosity of emulsified WCO formulated increased exponentially with increasing of water content. According to stabilities test, emulsified WCO showed good stability, also it was better than emulsified WCOME.

Mubarak and Senthil (Mubarak and Senthil, 2012) prepared emulsified biofuels using WCO, ethanol, water and Span 80 as surfactant. The emulsions were prepared varying the amount of neat waste cooking oil, water and the ratio of surfactant/co-surfactant (Span 80/ethanol) in the system. These emulsified biofuels were prepared stirring vigorously. From the stability test, it was found that the mixture of 70% of WCO, 15% of water, 10% of ethanol and 5% of Span 80 was stable for two weeks (Mubarak and Senthil, 2012). The physical-chemical properties of emulsified biofuels formulated in both researches were not reported.

Senthil and Jaikumar (Senthil and Jaikumar, 2014) obtained their emulsions with specified amount of neat WCO, water, ethanol and Span 80 surfactant. From the stability test, it was found that the mixture of 70% of WCO, 15% of water, 10% of ethanol and 5% of surfactant by volume was stable for two weeks (Senthil and Jaikumar, 2014). On the other hand, Nanthagopal and Subbarao (Nanthagopal and Subbarao, 2009), using high-speed stirrer prepared their emulsions with diesel fuel-WCO mixture (equal quantities), different water amount (10, 20 and 30%) and surfactant. The physical-chemical properties of the emulsions in both studies were not reported.

On the other hand, emulsified FADs from soybean were formulated by Melo (Melo et al., 2014). The emulsified FADs were prepared as ternary systems using residual FAD, methanol anhydrous and the moisture content into FAD (see Table 1), take into account the less as possible stabilization time during the experiments. For this reason, a factorial design 2<sup>3</sup> was developed (selected factors analyzed: temperature, methanol percentage and intensity of agitation). The microemulsions formation was detected as the formation of one phase of a very clear, transparent and totally stable liquid system. The dynamic viscosity of emulsified FAD is 6.1 mPa·s higher than the value found for the reference diesel fuel. The difference between neat FAD and emulsified FAD was 20.0 mPa·s. The density measurements did not evidence significant differences among fuels.

## **PERFORMANCE AND EXHAUST EMISSIONS ASSESSMENT OF DIESEL ENGINES FUELLED WITH BIOFUELS FROM WCO AND FAD**

The effects of the emulsified fuels on the engine performance are different from one study to another (Armas et al., 2005). The results obtained depend, mainly, on the engine operation mode (Castro et al., 1997; Samec et al., 2002), type and tuning of the injection system, and finally, on the optimized combustion chamber configuration (Armas et al., 2005). The physical-chemical properties of the emulsified fuel also play a significant role. A summary of the some experimental results reported about the use of WCO, FAD and their derivatives compared with diesel fuel are shown in table 4 and 5.

The WCO and FAD have a lower heating value, cetane number and poor volatility than diesel fuel (see Table 1) and their properties play an important influence on the premixed combustion phase. For this reason, differences in engine performance and exhaust emissions are expected, e.g. higher ignition delays for emulsified biofuels compared with diesel fuel and neat WCO, as was reported by Senthil, Jaikumar and Mubarak (Mubarak and Senthil, 2012; Senthil and Jaikumar, 2014). Generally, also the water content into emulsified biofuels delays the ignition.

Moreover, as a consequence of the lower heating value of the emulsified WCO and FAD, an increase of the specific fuel consumption was reported. In some experiments, the emulsified WCO achieved lower HC, CO and smoke opacity than neat WCO (Mubarak and Senthil, 2012) and diesel fuel (Nanthagopal and Subbarao, 2009). This behavior might be attributed to improvements on spray formation and the oxygen content into biofuels, despite adverse phenomena such as low temperature bulk quenching of the oxidation reactions, etc.

From the nitrogen oxide emissions reported (see Table 4 and Table 5), the use of emulsification method decreases the NO<sub>x</sub> emissions compared with diesel fuel, neat WCO, neat FAD and their derivatives. This topic is one of the most important advantages to use the emulsification. An explanation to these results is the thermal effect of the water on the combustion temperature. Nevertheless, formation of NO<sub>x</sub> is quite complex because numerous intermediate species exist (Imtanan, Varman et al. 2014). Similar results have been reported by different researchers (Crookes et al., 1995; Singh et al., 2010; Qi et al., 2013; Qi et al., 2013) using other feedstocks.

Finally, increases of the exhaust gas temperatures (Pugazhavadivu and Jeyachandran, 2005; Narayana et al., 2008; Hirkude et al., 2014) to WCOs and their derivatives, compared with diesel fuel might be associated to slow combustion, the amount increases of fuel burned to meet the power requirement and the presence of the oxygen in the chemical

structure of these biofuels. Also, an important role plays in the case of WCOME the higher latent heat of vaporization of methanol, in concordance with Senthil et al. (Senthil, 2003).

Table 4: Performance and exhausts emissions assessment of diesel engines fuelled different biofuels from fad compared with diesel fuel

Type	Engine	Performance, emissions and engine components wear	Ref.
Neat FAD (preheated at 70°C)  Different blends (10,15, 25, 50%) of FAD in diesel fuel	Petter 1-cylinder DI	Slight differences in ignition delay ↓ Cylinder peak pressure ↑ BSFC ↓ NOx ↑ CO and HC	(Ferrer and Piloto, 2014; Melo, Piloto et al., 2014; Melo et al., 2014)
Emulsified FAD	Petter 1-cylinder DI	↑ BSFC ↓ NOx	(Melo et al., 2014)
Neat FAD (preheated at 110°C)	6-cylinder DI	↑ Dark deposits on the piston crown, the rings, the combustion chamber and the injector. Heavy erosion produced by particles in the fuel facilitates the start of microcracks, producing fatigue loads and the failure of fuel injectors	(Galle et al., 2012; Piloto et al., 2014)

DI: Direct injection, FAD: fatty acid distillates, BSFC: brake specific fuel consumption, HC: unburned hydrocarbons, PM: particulate matter, CO: carbon monoxide, NOx: nitrogen oxides

Table 5: Performance and exhausts emissions assessment of diesel engines fuelled different biofuels from wco compared with diesel fuel

Type	Engine	Performance and emissions	Ref.
Neat WCOME and two blends (70 and 30%) of WCOME in diesel fuel	Deutz 2-cylinders DI	↓ Cylinder pressure, Heat release rate and Ignition delay ↓ CO, HC ↑ NOx	(Elshaib et al., 2014)
Different blends (10, 20, 30 and 50%) of WCOME in diesel fuel	1-cylinder DI	↑ BSFC and Cylinder peak pressure ↓ BTE, CO and HC ↑ NOx	(El-Kassaby and Nemit-allah, 2013)
Emulsified WCOME and emulsified WCO-Diesel fuel in equal proportion (both emulsion with 10, 20, 30 and 40% of water)	1-cylinder DI	↑ Kinematic viscosity with the increase of water amount ↓ NOx and Smoke density without worsening BSFC with water to fuel volume ratios	(Yoshimoto et al., 1999)
Neat WCOME and blends (90, 70 and 50%) in diesel fuel	Kirloskar 1-cylinder DI	↓ BTE with the increases of WCOME in the blends ↑ BSFC with the increases of WCOME in the blends ↑ Exhaust gas temperature and Smoke opacity with the increases of WCOME in the blends	(Hirkude et al., 2014)
Neat WCO (preheated at 30°C, 75°C and 135°C)	Kirloskar 1-cylinder DI	↑ BSEC, Exhaust gas temperature, CO and smoke density ↓ BTE and NOx	(Pugazhvadivu and Jeyachandran, 2005)
Neat WCOME and blends of (10, 30, 50 and 70%) in diesel fuel	Kirloskar 1-cylinder DI	↑ BSEC ↓ BTE and NOx ↑ Smoke density	(Subramaniam et al., 2013)

Neat WCOME	Petter 1-cylinder DI	↑ BSFC ↓ NOx and CO	(Gandón et al., 2014)
Different blends (25, 50 and 75% ) of WCOME in diesel fuel	TTF 8000s 4-cylinder DI	↓ Smoke and HC ↑ CO and NOx	(Arslan, 2011)
Neat WCOME and blends of (20, 40, 50 and 80%) in diesel fuel	Kirloskar 1-cylinder DI	↓ Rate of heat release and BTE ↑ Cylinder peak pressure and BSFC ↓ Ignition delay, CO and HC ↑ NOx and exhaust gas temperature	(Narayana et al., 2008)
Neat WCO (preheated at 70°C) and a blend of 15% of WCO in diesel fuel	Petter 1-cylinder DI	↓ Cylinder peak pressure ↑ Ignition delay at two experimental points (3.14KW and 4.63KW) ↑ BSFC, CO and HC ↓ NOx	(Ferrer 2013; Gandón et al., 2014; Melo et al., 2014)
Neat WCO and Emulsified WCO (70% WCO, 15% water, 10% ethanol and 5% Span 80)	Kirloskar 1-cylinder DI	↓ Cylinder pressure. However, in three experimental points the emulsified WCO was higher than neat WCO ↑ BSEC and Ignition delay ↓ BTE and NOx ↑ HC, CO and Smoke opacity. In some experimental points the emulsified WCO was lower than neat WCO	(Mubarak and Senthil, 2012; Senthil and Jaikumar, 2014)
A blend of WCO in diesel fuel (equal quantities)  WCO-diesel fuel emulsions with different water contents (10, 20 and 30%)	DI	↓ BSFC, except to emulsified WCO-diesel fuel with 30% of water ↑ BTE ↓ CO and NOx ↑ PM and Smoke intensity	(Nanthagopal and Subbarao, 2009)

DI: Direct injection, WCOME: waste cooking oil methyl ester, BTE: brake thermal efficiency, BSFC: brake specific fuel consumption, BSEC: brake specific energy consumption, HC: unburned hydrocarbons, PM: particulate matter, CO: carbon monoxide, NOx: nitrogen oxides

## CONCLUSIONS

The review of state of the art developed here is an approach to emulsified waste cooking oils and fatty acid distillates with the aim of assemble experimental results reported in the scientific literature and enhancing the knowledge about this topic (formulation, characterization, engine performance and emissions). The investigations developed about the use of WCO and FAD as diesel engine fuel shown the transesterification as the method more commonly applied; in spite of the necessary step to remove the higher free fatty acid, glyceride and moisture content into FAD and WCO, as well as economic and environmental facilities that brings the emulsification method over other.

Studies focused on the analysis of formulation, stability and optimization of the FAD and WCO emulsification have been a little developed. In addition, the physical-chemical properties of emulsified biofuels formulated were not reported.

According to the engine performance and exhaust emissions, it is possible through the emulsification to use the FAD and WCO as engine biofuels. Differences between physical-chemical properties of FAD, WCO and diesel fuel are responsible of variations of the specific fuel consumption, ignition delay and exhaust emissions reported. Even the contribution of oxygen in the chemical structure of the biofuels, the water amount and the microexplosion process played their role. However, although the engine performance and exhaust emissions behavior depend on the physical-chemical properties of the emulsified biofuels, engine type and operation conditions but the exact relationship need to be further clarified.

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