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**Assessing combustion and emission performance of direct use of SVO in a diesel engine  
by oxygen enrichment of intake air method**

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20 **Abstract**

21 This work investigated the effect of the oxygen enrichment in the intake air of diesel engines  
22 on the combustion and emissions performance using Rape Seed Oil (RSO) as a fuel. The  
23 purpose of the paper is to investigate the potential of oxygen enrichment in the intake air  
24 method to restrain the deterioration of particulate emissions of the RSO due to its high  
25 viscosity so as to explore the possibility of direct use of SVO (Straight Vegetable Oil) in  
26 diesel engines, which can reduce CO<sub>2</sub> emissions and save cost. The combustion parameters  
27 such as ignition delay, heat release rate, in-cylinder peak temperature and pressure were  
28 determined. Engine out particulate and gaseous emissions of the RSO were measured at  
29 oxygen concentrations from 21% (by volume) (no enrichment) to 24% (by volume) and  
30 compared to diesel results. The enrichment of the intake air with oxygen decreased the  
31 ignition delay and premixed combustion duration, and increased the in-cylinder peak pressure  
32 and temperature. The particulate, CO and hydrocarbon emissions were significantly reduced  
33 while the NO<sub>x</sub> emissions increased as the oxygen enrichment rate increased. 22% oxygen  
34 enrichment rate was suggested to achieve lower than diesel particulate emissions with the  
35 lowest NO<sub>x</sub> penalty. Increased NO<sub>x</sub> could be controlled by other methods. The results show  
36 that the oxygen enrichment in intake air method enabled direct combustion of SVO in diesel  
37 engines with reduced particulate, hydrocarbon and CO emissions.

38 **Key words**

39 Biofuel, Straight vegetable oil, Exhaust emission, Diesel engine, Oxygen enrichment

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## 44 1. INTRODUCTION

45       Increasing environmental concerns and security of energy supply have accelerated the  
46 research and development of alternative fuels for transport and power generation  
47 applications. Biodiesel, as a most popular biofuel in transport, is made from  
48 transesterification of vegetable oils or animal fats. The transesterification process consumes  
49 the energy and produces CO<sub>2</sub> and increases the cost as well as rejects about 10% of the  
50 feedstock as waste glycerine. Straight vegetable oils (SVO) and waste vegetable oils or  
51 waste cooking oils (WVO or WCO) can be used as fuels in diesel engines with a better  
52 carbon footprint compared to FAME (Fatty Acid Methyl Ester). The use of waste cooking  
53 oils and SVOs is potentially cheaper than using petroleum diesel or edible oil biodiesel [1].  
54 However, performance problems have been reported [2, 3]. These unprocessed (non-  
55 esterified) biofuels are not commonly used due to their high viscosity and low volatility,  
56 leading to poor atomisation and thus high particulate emissions as well as other problems  
57 such as handling problems and engine deposits and injector coking [4]. This study has  
58 investigated the impact of oxygen enrichment in the air inlet of the engine on the combustion  
59 and emissions characteristics of rape seed oil burned in a diesel engine. The aim is to explore  
60 the possibility of using oxygen enrichment method to overcome the adverse effect on engines  
61 and environment caused by the high viscosity of SVO so as to provide essential information  
62 to enable further evaluation of oxygen enrichment method as a way to utilise SVO (including  
63 waste cooking oils) directly in diesel engines without causing adverse impact. The primary  
64 potential application is aimed at stationary applications such as power generation.

65           The concept of the oxygen enrichment of intake air as a mean of improving combustion  
66 and reducing emissions in diesel engines has been investigated for a long time [5-14]. These  
67 researches focused on the air intake enrichment of conventional petroleum diesel and  
68 compared with addition of oxygenates in the petroleum diesel. Enrichment of the air intake of  
69 naturally aspirated diesel engines can be an effective measure to reduce emission and  
70 improve thermal efficiency. Previous publications on the subject suggest that the amount of  
71 O<sub>2</sub> addition is a trade off between NO<sub>x</sub> increase and reduction of PM (Particulate Matter),  
72 HC, CO and smoke emissions. Watson and Rigby [5] conducted experiments on a Caterpillar  
73 V-8 engine with oxygen enrichment up to 30 % (volume fraction and so as hereinafter). They  
74 found smoke emissions to decrease gradually with O<sub>2</sub> increase to around 10 H.S.U.  
75 (Hartridge Smoke Unit) at all engine speeds compared to 80 H.S.U. for 21% O<sub>2</sub> (no oxygen  
76 enrichment). NO<sub>x</sub> emissions gradually increased with increasing oxygen enrichment to about  
77 20% at 30% oxygen, acceptable levels were found at 24 % oxygen. Particulate emissions  
78 were reduced up to 80 % at full load conditions. Desai and Watson [7] further investigated  
79 impact of different grades of diesel fuels at various O<sub>2</sub> enrichment rates for emission  
80 characteristics and performance. They found that smoke and particulate emissions  
81 substantially decreased due to oxygen enrichment for all fuels. Ignition Delay was found to  
82 decline at higher oxygen levels which in combination with injection timing will maintain  
83 NO<sub>x</sub> levels within acceptable levels while large benefits in PM and smoke reduction can be  
84 achieved. Donahue and Foster [8] conducted oxygen enrichment tests on a Cummins N 14  
85 single cylinder diesel engine with oxygen rates of 21, 22 and 23 %. They found an increase to  
86 22 % oxygen showed drastic reductions of particulate matter emissions while only  
87 moderately increasing NO<sub>x</sub>, the exact magnitude depending on injection timing. A further  
88 increase to 23 % only showed marginal decrease in PM but large unfavourable increase in  
89 NO<sub>x</sub>. Increased O<sub>2</sub> intake was found to reduce ignition delay and increase peak cylinder

90 pressure for all load conditions. The increase in NO<sub>x</sub> with higher O<sub>2</sub> rates was attributed to  
91 increased temperatures and the PM reduction to increased fuel pyrolysis and soot oxidation.  
92 Rakopoulos et al [10] did an extensive literature review on previous research of the subject.  
93 Their conclusions are consistent with above mentioned findings; the shorter ignition delay  
94 was attributed to the accelerated pre-ignition reactions. Poola and Sekar [14] investigated the  
95 impact of oxygen enrichment of combustion air on the operational and emissions properties  
96 using a locomotive research diesel engine (two cylinder, EMD 567B). They found that the  
97 NO<sub>x</sub> could be reduced simultaneously along with PM emissions when the concentration of  
98 intake air oxygen, fuelling rate, and injection timing were optimized. Their results showed  
99 that particulates were reduced by approximately 60% and NO<sub>x</sub> emissions were reduced by  
100 15–20% with the optimal operating strategy. Higher gross power, lower peak cylinder  
101 pressures, and lower brake specific fuel consumption were also observed. A recent study by  
102 Perez and Boehman [15] studied the effect of air inlet oxygen enrichment on engine  
103 performance at simulated high altitude conditions using a small aero diesel engine. One of  
104 their key findings was that the oxygen enrichment decreased brake-specific fuel consumption  
105 (BSFC) by 40% due to increased fuel conversion efficiency. The emission was not measured  
106 as it was not the objectives of the research. Table 1 below summarized the key elements of  
107 major publications related to experimental results of oxygen enrichment of air intake for  
108 diesel engines. Besides the experimental work, there are some theoretical researches on air  
109 intake oxygen enrichment. Zannis et al [16, 17] modelled the effect of air intake oxygen  
110 enrichment method on combustion and emissions and found similar results such as the  
111 shorter ignition delay, higher in-cylinder pressure, lower PM and higher NO<sub>x</sub> emissions with  
112 oxygen enrichment.

113

114 Table 1. Review of previous key O<sub>2</sub> enrichment in air intake experiments on diesels

Author and year of publication	O <sub>2</sub> -Enrichment Rates	Recommended Rate	Conclusions
Watson and Rigby;1990 [5]	21-30	24	NOx control difficult but possible by retarded injection. Large reductions in PM and smoke possible.
Virk et al;1993 [6]	21-27	22.5	NOx levels were kept at the same value as the unenriched-air, while particulates and smoke emissions were reduced by 30%.
Desai and Watson;1997 [7]	21, 25, 27	25	NOx levels acceptable by injection timing control for low smoke and PM
Donahue and Foster; 2000 [8]	21, 22, 23	22	22 % only small NOx increase but large PM. 23 % only small decrease in PM but large increase in PM.
Poola and Sekar; 2003 [14]	21,23,25,27	23	At optimum oxygen-enrichment rate (~23%) and injection timing, PM emissions were reduced by ~60%, NOx emissions were reduced by ~14%, gross power was increased by ~16%, and BSFC was reduced by ~2% Such wide benefits demonstrate the potential of the oxygen-enrichment technique.
Perez and Boehman; 2009~ 2010 [12] [15]	21, 22, 23	22-23	A modest decrease in BSFC by oxygen enrichment. PM emissions were reduced while NOx deterioration was minimized when combined with EGR.

115

116 **2. EXPERIMENTAL**

117 **2.1 Test engine and fuels**

118 A Perkins Phaser 180Ti 6 cylinder, 6 L (0.006 m<sup>3</sup>), DI (Direct Injection), turbo-charged inter-  
 119 cooled, heavy duty diesel engine was used, fitted with an oxidation catalyst. The  
 120 specifications of the engine are shown in Table 2. The engine was of Euro 2 emissions

121 standards compliance. The engine was running at 47kW (50% of its maximum load at this  
 122 speed) and 157 rad s<sup>-1</sup> constant conditions.

123 Table 2. Engine specifications

Type	Perkins Phaser 180Ti
Displacement (L)	6.0
Cylinder No.	6
Compression ratio	17.5 : 1
Maximum power rating	134kW at 272 rad s <sup>-1</sup>
Aspiration	Turbo-charged and Inter-cooled
EGR	No
Fuel injection	Pump-line-nozzle
Injector hole diameter, (mm)	0.2
The number of holes	6
Injection timing, °BTDC	11
BMEP at 47kW & 157 rad s <sup>-1</sup> (KPa)	628

124

125 A food grade virgin Rapeseed Oil (RSO), Agri green extended life cooking oil and produced  
 126 by Agri Energy Food Service co, was used as the testing fuel. The oil was produced by deep  
 127 degumming process and the key fuels properties were listed in table 3.

128 Table 3. Selected physical properties of the test fuel RSO

Properties	Values
Colour	15Yellow



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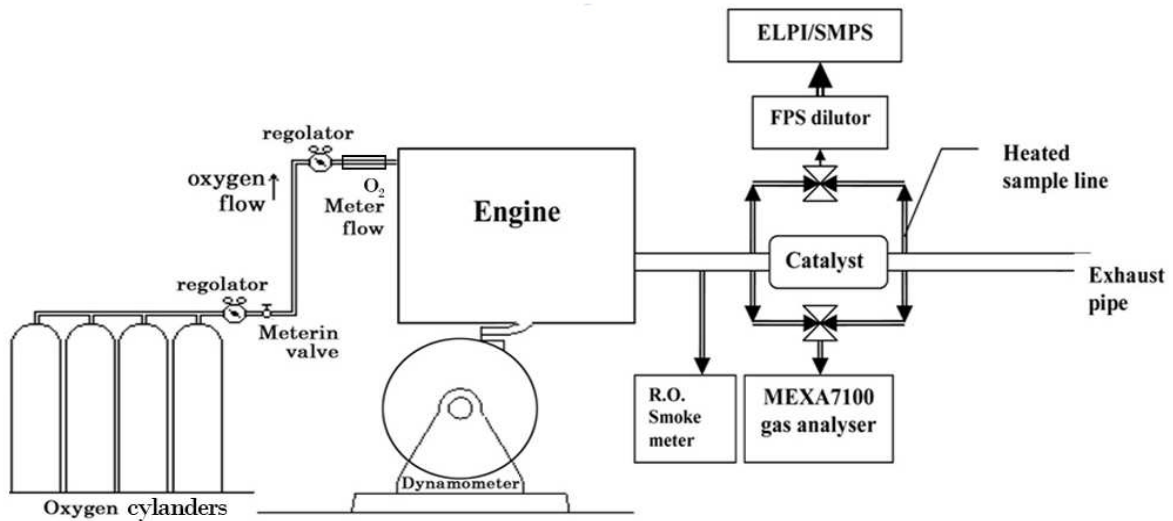
	/1.5 red
Free fatty acids (mass fraction %) max	0.15
Protein (mass fraction %)	0
Insoluble impurities (mass fraction %) max	0
Density @15°C (kg m <sup>-3</sup> )	922
Kinematic viscosity @40°C (mm <sup>2</sup> s <sup>-1</sup> )	35.61
Flash point (°C)	320
Oxygen (mass fraction %)	17.5
Calorific value (MJ kg <sup>-1</sup> )	39

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129

### 130 **2.3 Oxygen enrichment layout**

131 The oxygen was supplied by four 46.6 L (0.0466 m<sup>3</sup>) capacity cylinders, passing through a  
132 pressure regulator and a volumetric flow meter and transported to a valve. The oxygen was  
133 then passed through a mass flow meter with an accuracy of 1%, which was connected to a  
134 laptop for the control and measurement of oxygen flow rate. The influence of surrounding  
135 temperatures on the oxygen flow rate was minimal as all the pipes and meters were housed  
136 indoors and the temperature was maintained at 15±2°C. The metered oxygen was injected  
137 into the air stream by inserting a probe downstream of the intercooler and one meter upstream  
138 of air inlet manifold of the engine. A dedicated sample probe was inserted in the engine air  
139 inlet manifold and connected to a Servomex paramagnetic oxygen analyser to measure the  
140 oxygen concentration after mixing with an accuracy of ±0.1% of oxygen concentration. The  
141 readings were used to monitor oxygen level and adjust oxygen flow rate if necessary. Figure  
142 1 shows the schematic view of the experimental set up.



143

144 Figure 1. Schematic view of oxygen enrichment test set up

145 **2.4 In-cylinder pressure and crank angle measurement**

146 One of the cylinders on the test engine was fitted with a piezo-electric pressure transducer for  
 147 measuring in-cylinder pressure, which was connected to a data acquisition system that could  
 148 take around 100, 000 pressure readings per second. The Top Dead Centre (TDC) position was  
 149 measured using an optical sensor and indentation on the flywheel at TDC. The time based  
 150 pressure results were then converted into P- $\theta$  (pressure-crank angle) diagram. At least 50  
 151 cycles were converted for each condition. An average P- $\theta$  diagram was then obtained. The  
 152 heat release rate was thus calculated. The ignition delay and premixed combustion duration  
 153 (PCD) was then determined. The variations were determined and indicated by the error bars  
 154 (absolute error) in the figure 7. This method has also been used by other researchers [18-20]  
 155 and proved to be reliable. The ignition delay was defined as from the beginning of fuel  
 156 injection at 11 ° BTDC (Before Top Dead Centre) until the start of premixed combustion.  
 157 The PCD was defined between the end of ignition delay and the start of stratified combustion.  
 158 The in-cylinder peak temperature was computed based on peak pressure data.

## 159 **2.5 Emissions sampling systems**

160 The particulate and gaseous exhaust samples were taken at 1.3 m from the turbocharger  
161 upstream of the catalyst and at 2.05 m downstream of the catalyst respectively, as shown in  
162 figure 1. A constant temperature sampling (CTS) technique was used with a 6.3 mm (0.0063  
163 m) stainless steel pipe inserted with a curved bend onto the centre line of the exhaust pipe.  
164 The sample for particulate collection was passed through a heated sample line to a heated  
165 filter in an oven at 50°C. The SAE smoke meter (R.O.smoke meter manufactured by Richard  
166 Oliver Ltd.) was used, which was modified to include separate temperature control of the  
167 filter paper block and the oven. There was sufficient sample residence in the oven to cool the  
168 sample to 50°C. Whatman 0.055 m diameter GF/F glass fibre filters were used with a 0.029  
169 m spot size. The sampling flow rate was kept at 5 L min<sup>-1</sup> and at least 0.001 g of particulate  
170 was collected, weighed to 0.000010 g resolutions and hence the minimum accuracy of the  
171 filter weights was ±2%.

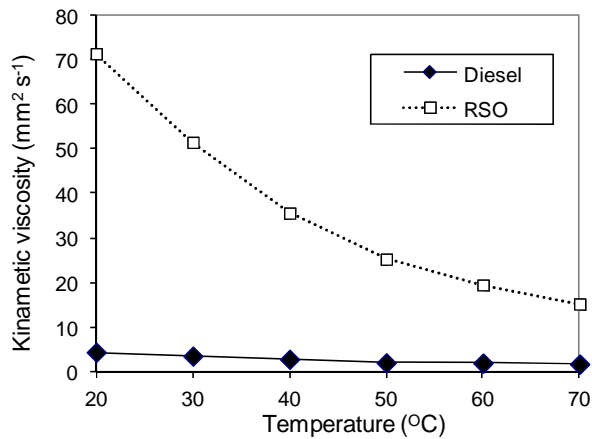
172 Gas samples were taken from the same sample position in the exhaust as the particulates and  
173 transported via a 6 m long 6.3 mm diameter electrically heated Teflon tube to a Horiba  
174 MEXA-7100D on-line gas analysis system. The sample line temperature was kept at 190°C.  
175 The exhaust gas analysis system contained a heated FID (Flame Ionization Detector) analyzer  
176 for total hydrocarbon analysis, a Chemiluminescence analyzer for NO and NO<sub>2</sub> analysis, and  
177 two NDIR (Non-Dispersive Infrared) analyzers for CO and CO<sub>2</sub> analysis. The air/fuel ratio  
178 was calculated from the gas analysis results based on the carbon balance principle, taking into  
179 account the oxygen content of the biofuels. This was used to convert the measured emissions  
180 data from volumetric to mass emissions. The exhaust gas samples were measured every 5  
181 seconds and at least 300 readings were taken on each testing condition.

182

### 183 3. RESULTS AND DISCUSSION

#### 184 3.1 Fuel viscosity

185 One of the major problems for using RSO in diesel engines is its much high viscosity  
186 compared to diesel fuel. The kinematic viscosity of the RSO was measured as a function of  
187 temperature along with the standard diesel fuel as shown in figure 2. The RSO had a much  
188 higher viscosity than the diesel fuel and its viscosity was much more variable with  
189 temperature. It was observed during the experiment that the temperature of the RSO was  
190 raised from room temperature in the fuel tank to ~55 °C before entering the fuel injector due  
191 to the pumping heat and the heat from the engine block. However, the difference in viscosity  
192 between the RSO and diesel at the fuel injection point was still significant. The high viscosity  
193 of the RSO would deteriorate the fuel spray and atomization and affect mixing of fuel with  
194 air and therefore adversely affect combustion and emissions [21]. The cone angle of fuel  
195 spray decreases, and the size and penetration of fuel droplets increase as the viscosity of fuels  
196 increases. This could cause direct contact of fuels with the combustion chamber wall, the  
197 piston surface and rings, resulting in carbon deposits formation. The high viscosity could  
198 also lead to higher pressure accumulations in the fuel injectors and fuel lines and cause early  
199 fuel injections. Another adverse effect of the high viscosity of the RSO is the dilution of the  
200 engine lubricating oil.



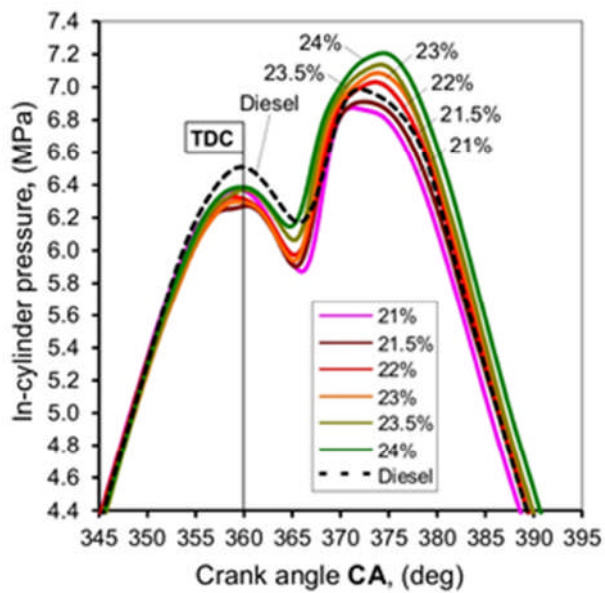
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202 Figure 2. Fuels kinematic viscosity as a function of temperature

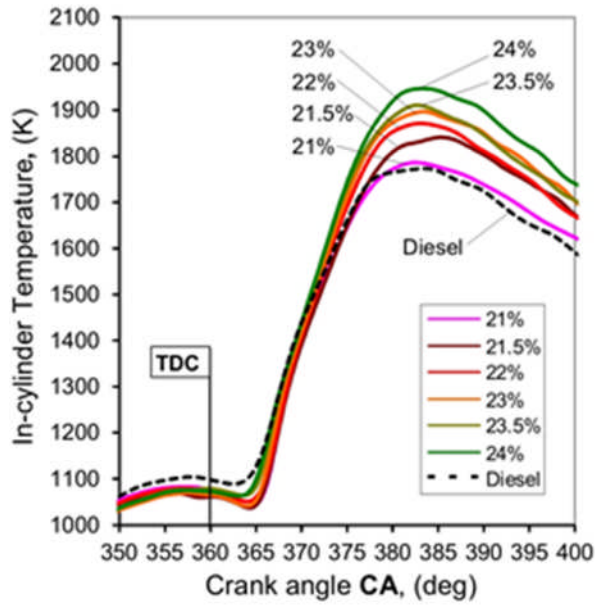
203 **3.2 Combustion parameters - ignition delay, premixed combustion, peak in-cylinder**  
 204 **pressure and temperature, heat release and indicated efficiency**

205 Figure 3 shows the pressure-crank angle (p-θ) diagram of the RSO at four different oxygen  
 206 enrichment rates/levels. The fuel was injected at 11° BTDC. The ignition occurred at  
 207 approximately 5° ATDC. The results show a clear trend towards shorter ignition delays as the  
 208 oxygen rate increased. The peak pressure was increased from ~6.8 MPa to 7.2 MPa, ~6%  
 209 increase, as the oxygen rate increased from 21% to 24%. Figure 4 shows computed in-  
 210 cylinder gas temperatures for different oxygen rates using the equation of state for ideal gas.  
 211 Obviously, this was the mean gas temperatures inside the cylinder. Though the emissions in  
 212 diesel engine combustion are dominated by local temperature but the mean in-cylinder  
 213 temperature is strongly connected with the local temperature in the IC engines. The peak  
 214 temperature was increased from ~1780 to ~1950 K when the oxygen rate was increased from  
 215 21% to 24%, about 9% increase. Compared to diesel, the RSO showed the lower in-cylinder  
 216 pressure at the TDC. This could be due to two reasons. The first possible reason is that the  
 217 typical value of the specific heat capacity of rape seed oil is 1.91 kJ kg<sup>-1</sup>K<sup>-1</sup> while diesel is 1.8  
 218 kJ kg<sup>-1</sup>K<sup>-1</sup>. The higher specific heat of RSO leads to more heat absorption and heat transfer

219 from hot air to the RSO. The second possible reason is due to the advance of fuel injection.  
 220 Several researches have reported an inadvertent advance in fuel-injection timing using  
 221 biodiesel. It is caused by a higher bulk modulus comparing to diesel fuel [22-24]. The second  
 222 peak of in-cylinder pressure is higher for diesel fuel (compared to 21% oxygen RSO) because  
 223 the ignition delay of RSO is longer (fuel injection and evaporation of RSO takes more time)  
 224 than diesel. The higher pressure by oxygen enrichment for the RSO at around 370~375 °  
 225 crank angle showed more intensive diffusion combustion, reflected by the higher peak in-  
 226 cylinder temperature. Figure 5 shows the peak in-cylinder temperature as a function of  
 227 oxygen enrichment rate for the RSO. A good linear correlation is observed.

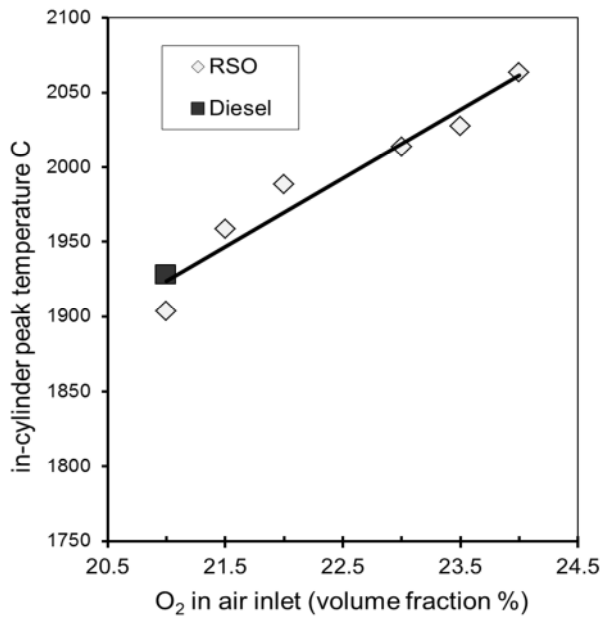


228  
 229 Figure 3. In-cylinder pressure variation as a function of crank angle (CA) for different  
 230 oxygen rates



231

232 Figure 4. In-cylinder temperature variation as a function of crank angle (CA) for different  
 233 oxygen rates

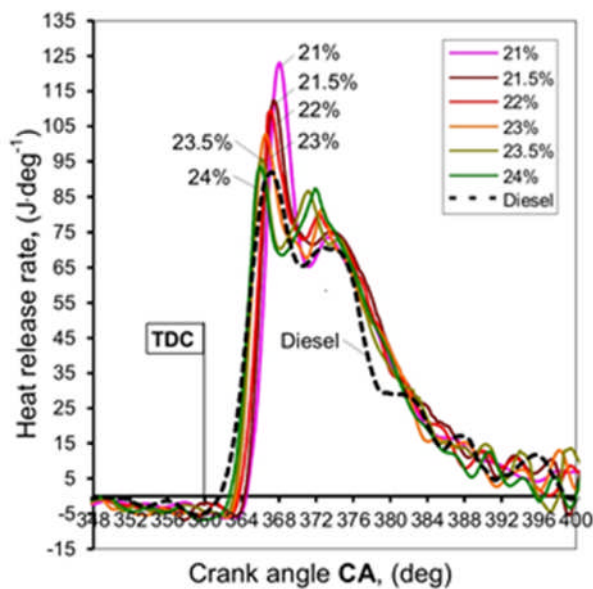


234

235 Figure 5. In-cylinder peak temperature as a function of oxygen rate

236 Figure 6 shows the heat release rate for different oxygen rates. The negative values before  
 237 ignition reflected the heat absorption by the RSO due to its evaporation, which was  $\sim 5 \text{ J deg}^{-1}$

238 <sup>1</sup>. The results show that the peak heat release rate was reduced as the oxygen rate increased.  
 239 This was due to that the shorter ignition delay at the higher oxygen rates reduced the fractions  
 240 of fuels burned in the premixed combustion phase. In contrast to the reduction of premixed  
 241 combustion phase, the diffusion combustion phase was extended at the higher oxygen rates,  
 242 along with a higher diffusion peak heat release (the second heat release peak) as shown in  
 243 figure 6. It was the higher heat release rate in the diffusion phase that caused higher mean in-  
 244 cylinder peak temperatures in figure 4, which then resulted in higher NOx emissions.

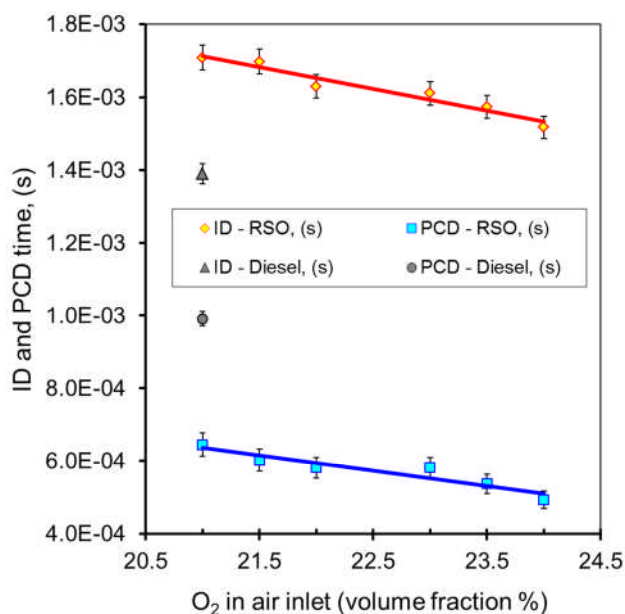


245  
 246 Figure 6. Heat release traces for different oxygen rates

247  
 248 The trend of ignition delay and PCD was plotted against the oxygen rates with error bars as  
 249 shown in figure 7. The error bars indicated the absolute error of the measurements. Both the  
 250 ignition delay and PCD show a significant decrease with increasing oxygen rates. The  
 251 reduction of the ignition delay was 1.7 ° at 24 % oxygen compared to oxygen level of 21%.  
 252 The shorter ignition delay resulted in shorter PCDs and longer diffusion combustion periods



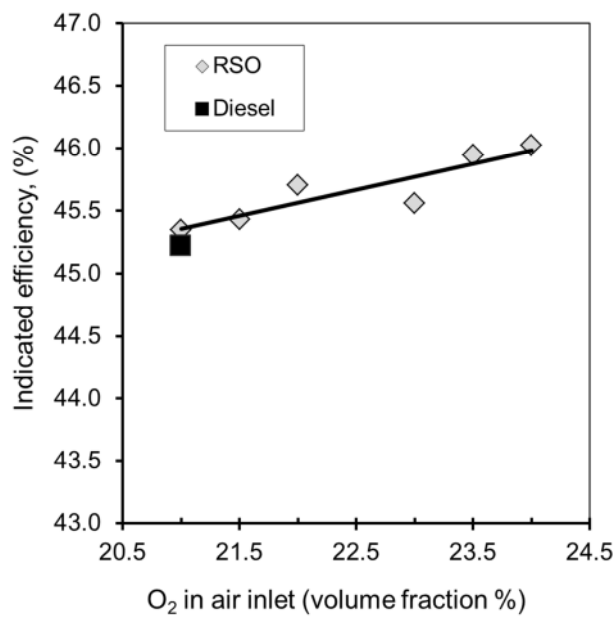
253 and thus higher flame temperatures and consequently higher NO<sub>x</sub>. The oxygen enrichment  
 254 caused the air fuel mixture during the PCD to be leaner, supporting higher NO<sub>x</sub> formation  
 255 through higher local temperatures in the PCD and, hence, in the NO<sub>x</sub>-forming diffusion  
 256 flame. The CO and hydrocarbon formation was reduced as a result of higher in-cylinder  
 257 temperatures and the air fuel ratio being leaner and further away from stoichiometric with the  
 258 increased oxygen rate. The additional oxygen could also result in late cycle oxidation of CO  
 259 and hydrocarbons further reducing these emissions. The ignition delay was gradually reduced  
 260 because the air fuel mixture will ignite at a lower temperature with more oxygen present.  
 261 Compared to diesel's ignition delay with the RSO without oxygen enrichment in figure 7, it  
 262 shows that the RSO had approximately 20% longer ignition delay than the diesel. This is  
 263 related to a lower cetane number of the RSO (~20% lower). Another possible reason is due to  
 264 the slower evaporation of the RSO. The results in figure 7 also show that the diesel had a  
 265 longer premixed combustion phase. This is due to that the amount of diesel injected during  
 266 the ignition delay period was greater than the viscous RSO. In other words, the low cetane  
 267 number of the RSO extended its ignition delay and the high viscosity of the RSO shortened  
 268 its premixed combustion period due to less amount of fuel being injected.



269

270 Figure 7. Changes in ID and PCD for different oxygen rates

271 Figure 8 shows the indicated engine efficiency as a function of oxygen rate. As expected,  
272 oxygen enrichment increased indicated engine efficiency. The maximum improvement of  
273 indicated efficiency is about 0.6% at 24% oxygen rate, which could be translated to about 1.2  
274 % fuel saving due to improved combustion efficiency by oxygen enrichment.



275  
276 Figure 8. Indicated efficiency as a function of oxygen rate

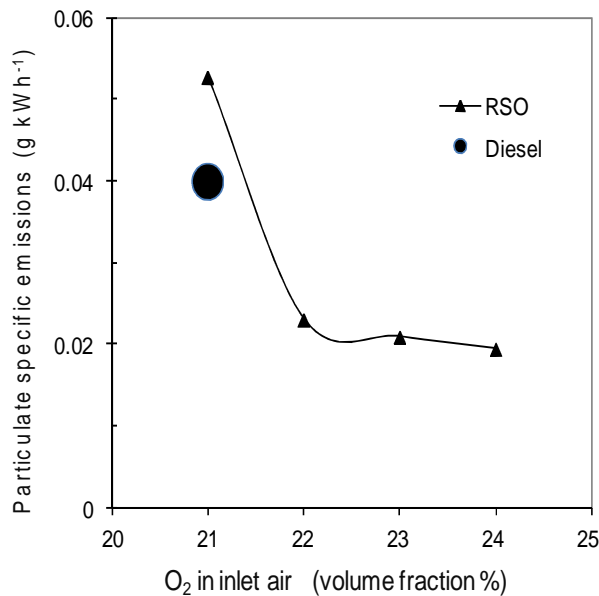
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### 278 3.3 Exhaust emissions

#### 279 3.3.1. Particulate emissions

280 The particulate matter (PM) was collected on the filter papers and mass emissions were  
281 determined. Figure 9 shows the particulate specific emissions ( $\text{g kWh}^{-1}$ ) as a function of  
282 oxygen rate. The PM specific emissions from the standard diesel were also plotted as a  
283 reference. Compared to diesel, RSO PM emissions (without oxygen enrichment) were

284 increased by ~30% due to its high viscosity and low volatility. The oxygen enrichment could  
285 effectively reduce the PM emissions from RSO. In fact, PM emissions from oxygen  
286 enrichment rate of 22% onwards were significantly lower than that of diesel fuel. The  
287 reductions in the PM mass of the RSO were the most significant between 21% and 22% of  
288 oxygen levels and became much slower afterwards. The higher in-cylinder gas temperature  
289 and extra oxygen helped the oxidation of the soot, resulting in less PM emissions. The EU  
290 emission regulation for EURO 2 heavy duty diesel after 1998 production is  $0.15 \text{ g kWh}^{-1}$ ,  
291 which is based on ECE R-49 test cycle with 13 steady state modes. The direct comparison of  
292 this research results with the EU regulation is not appropriate but the results in figure 9 could  
293 give an indication of PM emission levels relative to the regulation on a particular mode,  
294 i.e.50% of load at low speed mode, which was the operation condition of the test in this  
295 research.



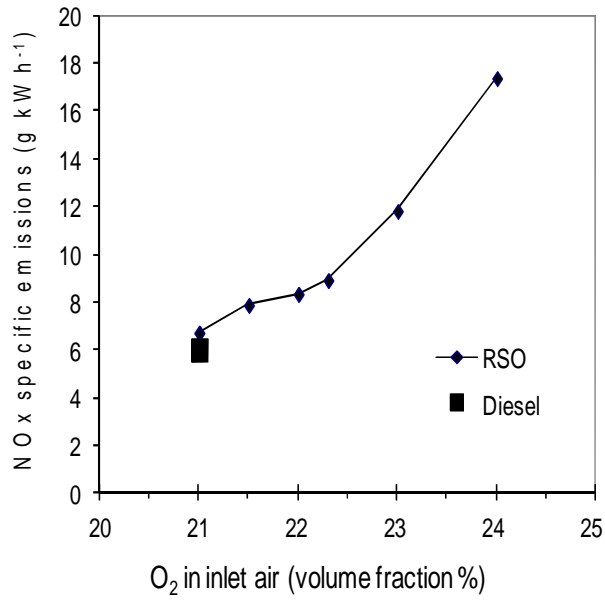
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297 Figure 9. PM specific emissions as a function of oxygen rate

298

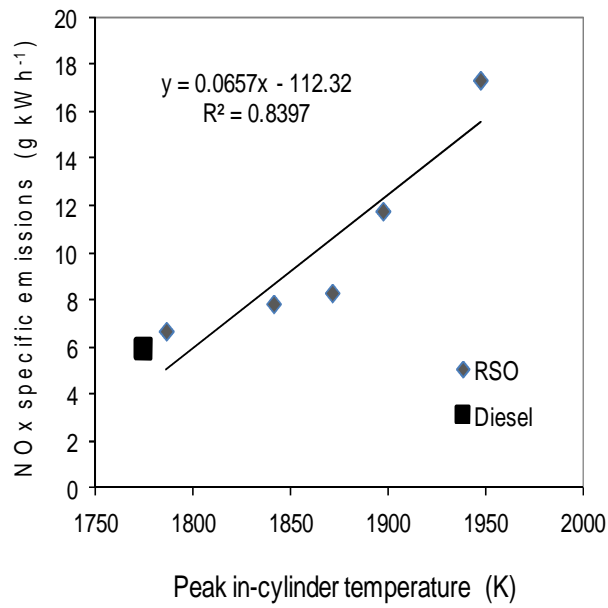
### 299 3.3.2. Nitrogen oxides (NO<sub>x</sub>) emissions

300 The EU NO<sub>x</sub> emission standard for this type of engine is 7 g kWh<sup>-1</sup> based on ECE R-49  
301 cycle. The condition the engine was tested represented one of the ECE R-49 mode, 50% of  
302 maximum power at low engine speed. The NO<sub>x</sub> level (6 g kWh<sup>-1</sup>) from the diesel fuel was  
303 already close to the standard. Figures 10 and 11 show the specific NO<sub>x</sub> emissions as a  
304 function of oxygen rate and peak in-cylinder temperature respectively. As expected, the NO<sub>x</sub>  
305 emissions increased significantly as the oxygen enrichment rate rose due to three major NO<sub>x</sub>  
306 formation contributors: the higher flame temperature, the higher local oxygen concentration  
307 and the longer diffusion combustion phase. A good linear correlation between NO<sub>x</sub> emissions  
308 and oxygen rate was observed. The NO<sub>x</sub> emissions were almost trebled to 17 g kWh<sup>-1</sup> at the  
309 highest oxygen rate (24%). The adverse effect on the NO<sub>x</sub> by oxygen enrichment could be  
310 minimised or even eliminated by changing fuel injection strategy - late fuel injection and  
311 adjusting fuel injection rate, as illustrated by Poola and Sekar [14]. The NO<sub>x</sub> could also be  
312 reduced by the use of EGR. Essentially, EGR and oxygen enrichment in intake air are two  
313 opposite methods serving two different purposes. The former is for NO<sub>x</sub> reduction while the  
314 latter is for PM reduction. A common practice is the combination of EGR with the other  
315 oxygen enrichment methods such as oxygenated fuels. However, these are out of the scope of  
316 this research. The NO<sub>x</sub> could also be treated with exhaust aftertreatment technologies such as  
317 SCR (Selective Catalytic Reduction) using urea. But a cost/benefit analysis is needed to  
318 assess the benefit of particulate reduction and penalty of NO<sub>x</sub> increase and associated cost of  
319 aftertreatment.



320

321 Figure 10. NO<sub>x</sub> specific emissions as a function of oxygen rate



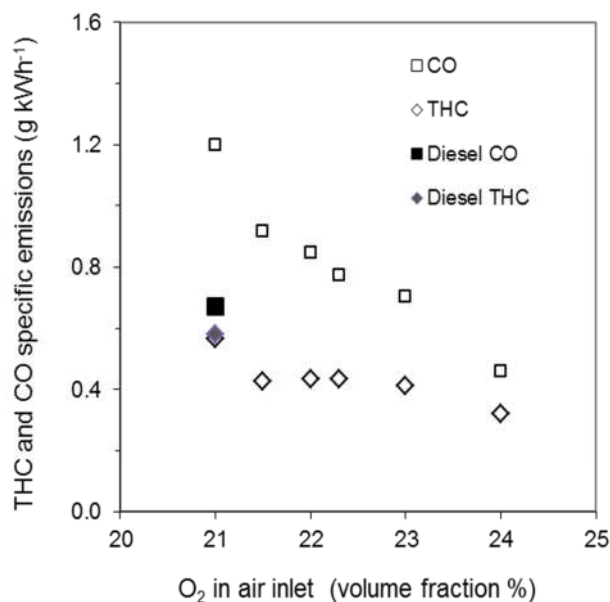
322

323 Figure 11. NO<sub>x</sub> specific emissions as a function of peak in-cylinder temperature

324

325 **3.3.3. CO and THC emissions**

326 The total hydrocarbon (THC) and CO specific emissions are presented in figure 12 as a  
 327 function of oxygen rate, along with the reference data of diesel results. Both the THC and CO  
 328 emissions are clearly showing strong tendencies of reductions as the oxygen enrichment  
 329 increased due to the oxidation effect taking place with additional oxygen. By comparison of  
 330 CO emissions between the diesel and the RSO at 21% oxygen level, it was suggested that the  
 331 poor atomization and mixing of the RSO occurred, indicated by much higher CO emissions  
 332 from the RSO. Surprisingly, there was no deterioration of hydrocarbon emissions from the  
 333 RSO compared to diesel. This could be attributed to the fuel born oxygen in the RSO assisted  
 334 the oxidation of the fuel.



335

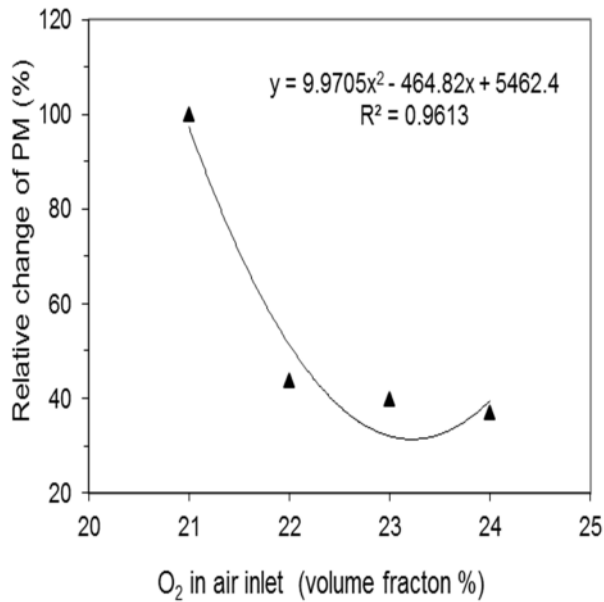
336 Figure 12. Specific emissions of THC and CO as a function of oxygen rate

337

### 338 3.4 Normalised emissions.

339 The PM and gaseous specific emissions were normalised to the values at oxygen  
 340 concentration of 21% (no oxygen enrichment) and presented in figures 13-15 in order to

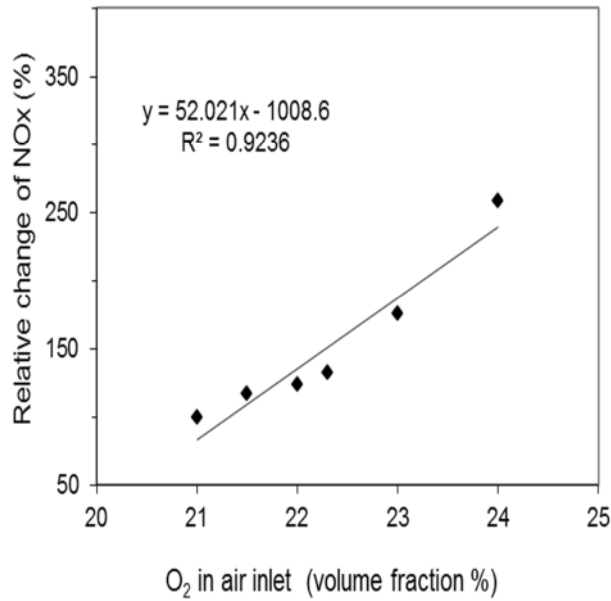
341 show the relative changes of emissions as a function of oxygen enrichment rate. The PM had  
342 a large decrease at oxygen concentration 22%, a reduction of ~60% compared to no oxygen  
343 enrichment. The reductions were much slower afterwards to an almost flat level.



344

345 Figure 13. Normalised particulate specific emissions as a function of oxygen rate

346 The normalised NO<sub>x</sub> specific emissions in figure 14 demonstrated a very good positive linear  
347 correlation with oxygen enrichment rate. The NO<sub>x</sub> emissions were increased by 150% at  
348 oxygen concentration of 24%, an average increase of ~50% for every 1% of increment of  
349 oxygen enrichment.



350

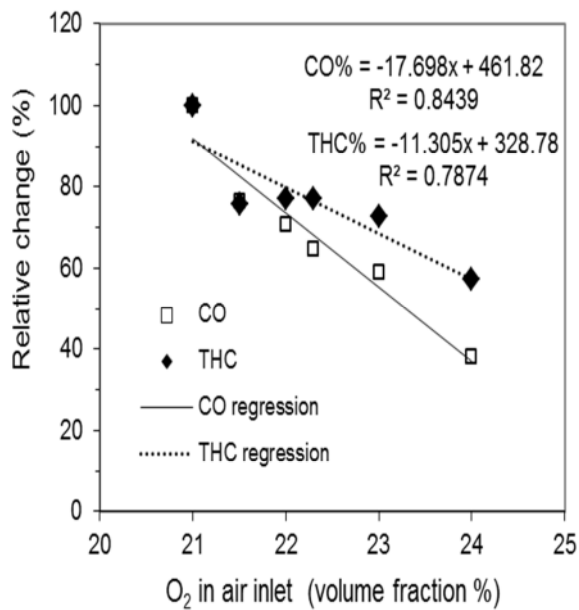
351 Figure 14. Normalised NOx specific emissions as a function of oxygen rate

352 Both the THC and CO specific emissions showed negative correlations with oxygen

353 enrichment rate. The CO emissions showed a better linear correlation than the THC

354 emissions. The CO and THC emissions were reduced by approximately 20% and 15% per

355 1% oxygen enrichment rate respectively.



356

357 Figure 15. Normalised THC and CO specific emissions as a function of oxygen rate



358 **4. CONCLUSION**

359 Direct use of pure SVO as a fuel in diesel engines without transesterification can offer greater  
360 CO<sub>2</sub> savings and lower cost compared to conventional biodiesels (FAME). However, high  
361 viscosity, low volatility of SVOs could lead to deteriorated combustion and emissions. The  
362 oxygen enrichment in the air inlet of the engine method as a way to improve the SVO's  
363 combustion so as to reduce emissions was investigated in this work. The air inlet oxygen  
364 levels tested were from 21% (no enrichment) to 24%. The results show:

365 1. The oxygen enrichment in the engine air inlet resulted in the shorter ignition delay and  
366 premixed combustion duration and longer diffusion combustion phase. The shorter ignition  
367 delay reduced the peak heat release rate at the premixed combustion phase but increased the  
368 peak heat release rate at the diffusion combustion phase. Overall the in-cylinder peak  
369 pressure and temperature were increased by 6% and 9% respectively when the oxygen rate  
370 was increased from 21% to 24%. This resulted in an improvement in engine efficiency  
371 measured by indicated efficiency.

372 2. Without oxygen enrichment, the RSO produced ~30% higher particulate emissions than  
373 the diesel at the same engine operating condition. However, the particulate emissions could  
374 be reduced to below the diesel level with oxygen enrichment rate of 22%. The reductions in  
375 particulate emissions were not linear with the oxygen enrichment rate. There was a large  
376 reduction in particulate emissions (~60%) between 21% (no enrichment) and 22% of oxygen  
377 level. The reductions were in a much less scale with the further oxygen enrichment.

378 3. NO<sub>x</sub> emissions showed a good positive linear correlation with oxygen enrichment rates  
379 and were increased on average by 50% for every 1% of oxygen enrichment rate. The high in-  
380 cylinder temperature and pressure are accounted for the higher engine out NO<sub>x</sub> emissions.

381 4. Without oxygen enrichment, the significantly higher CO emissions of the RSO, compared  
382 to diesel, indicated a poor atomisation and mixing of the RSO with air. The oxygen  
383 enrichment could reduce the CO emissions to a compatible level to diesel. The fuel born  
384 oxygen in the RSO helped the RSO to produce similar levels of hydrocarbon emissions to  
385 diesel. The oxygen enrichment overall reduced engine out CO and THC emissions.

386 5. The optimised oxygen enrichment rate should be 21~22%, where a significant reduction in  
387 particulate can be achieved with the lowest increase in NOx emissions.

388 6. The findings from this work demonstrated that adverse effects (deteriorated combustion  
389 and high emissions) of direct use of SVO in diesel engines can be solved and overcome by  
390 the oxygen enrichment method. This enables industrial diesel applications such as electric  
391 power generations to use the pure SVO directly to achieve greater CO<sub>2</sub> reductions and  
392 reduced fuel cost compared to biodiesel.

### 393 **NOTES**

394 The authors declare no competing financial interest.

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### 399 **NOMENCLATURE**

400 ATDC: After Top Dead Centre

401 BSFC: Brake Specific Fuel Consumption

- 402 BTDC: Before Top Dead Centre
- 403 CO: Carbon Monoxide
- 404 DI: Direct Injection
- 405 EGR: Exhaust Gas Recirculation
- 406 FAME (Fatty Acid Methyl Ester)
- 407 H.S.U.: Hartridge Smoke Unit
- 408 NOx: Nitrogen Oxides
- 409 PCD: Premixed Combustion Duration
- 410 PM: Particulate Matter
- 411 RSO: Rape Seed Oil
- 412 SCR: Selective Catalytic Reduction
- 413 SVO: Straight Vegetable oil
- 414 TDC: Top Dead Centre
- 415 THC: Total Hydrocarbon

416

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