

Borillo, GC and Tadano, YS and Godoi, AFL and Santana, SSM and Weronka, FM and Penteado Neto, RA and Rempel, D and Yamamoto, CI and Potgieter, SS and Potgieter, JH and Godoi, RHM (2015)*Effectiveness of Selective Catalytic Reduction (SCR) systems on reducing gaseous emissions from an engine using Diesel and Biodiesel Blends.* Environmental Science and Technology, 49 (5). pp. 3246-3251. ISSN 0013-936X

Downloaded from: http://e-space.mmu.ac.uk/620398/

Publisher: American Chemical Society

DOI: https://doi.org/10.1021/es505701r

Please cite the published version

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1	Effectiveness of Selective Catalytic Reduction (SCR) systems on
2	reducing gaseous emissions from an engine using Diesel and Biodiesel
3	Blends
4	
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32	ABSTRACT

33 There is an urgent and pressing need to further understand petroleum-based emission 34 control systems. To date, a limited number of emission studies have reported on the 35 effects on automotive emissions when vehicles equipped with Selective Catalytic 36 Reduction (SCR) systems run on a mixture of regular petroleum-based and biodiesel. 37 The aim of this investigation was to quantify organic and inorganic gas emissions from 38 a four-cylinder diesel engine equipped with urea-SCR system. Using a bench 39 dynamometer, the emissions from the following mixtures were evaluated using an **40** FTIR spectrometer: low sulphur diesel (LSD), ultra-low sulphur diesel (ULSD) and a 41 blend of 20 % soybean biodiesel and 80% ULSD (B20). Our results confirmed that the 42 use of the SCR system yields statistically significant (p<0.05) lower NO_x emissions in 43 comparison to all the studied fuels. The LSD and ULSD fuels also significantly reduced 44 emissions of compounds with high photochemical ozone creation potential, such as 45 formaldehyde. However, the SCR system produced significantly (p<0.05) higher 46 emissions of N₂O comparing the used fuels. In the case of LSD, the NH₃ emissions 47 were elevated and in the case of ULSD and B20 fuels, the non-methane hydrocarbon **48** (NMHC) and total hydrocarbon (HCD) emissions were significantly higher.

49

50 Keywords: Selective Catalytic Reduction (SCR); biodiesel; hydrocarbons; diesel;
51 emissions; gaseous pollutants.

52



60 based emission control systems. Global pressure to meet emission standards lead to the

61 development and use of new engine technologies and as of late also for the use of new62 fuels and fuel blends, such as ultra-low sulphur diesel and biodiesel blends.

63 Emissions depend on a variety of factors, such as engine technology, maintenance and emission control technology,¹ as well as the type and quality of the 64 employed fuel. Besides the greenhouse gas pollutants with global warming potential, it 65 66 is widely known that engine exhaust systems produce also organic gases that have an 67 impact on photochemical ozone and other secondary pollutants' formation. Among 68 such different gases emitted by petroleum-based systems, nitrogen oxides (NO_x) are 69 one of the major threats to the environment and therefore its emission in diesel engines has been widely investigated.²⁻⁵ NO_x suppression strategies consist of combustion 70 controls, such as Selective Catalytic Reduction (SCR) systems, using a urea solution as 71 reducing agent, a well-established technique of stationary diesel engines.⁶⁻⁸ Biodiesel 72 73 seems to be a promising alternative, as it can be used in diesel engines without major modifications,⁹ reducing qualitative and quantitatively several pollutant emissions.¹⁰⁻¹⁴ 74 75 The use of biofuels and fuel blends, in combination with exhaust aftertreatment systems 76 as a means of mitigating emissions, are promising and therefore the topic of this 77 investigation.

78 New standard guidelines are being established worldwide concerning heavy-79 duty diesel engine emissions, aiming mostly at the simultaneous reduction of particles 80 and NO_x (Euro V and Euro VI regulations in Europe and 40 Code of Federal Regulations 86.007-11).¹⁵ In Brazil, the ruling legislation is equivalent to the Euro V 81 emission standards and it was established on January 1st, 2012, as a result of the seventh 82 83 stage of the Program to Control Vehicular Air Pollution (PROCONVE, in Portuguese). 84 In order to achieve the Brazilian air quality guidelines, the sulphur content of diesel fuels was reduced and new aftertreatment systems have been implemented, with the 85 urea-SCR (Selective Catalytic Reduction) system being mostly utilized.^{4,5,16} 86

87 To date, a limited number of emission studies have reported on the effects of
88 biodiesel additions to regular petroleum-based diesel on emissions from vehicles
89 equipped with Selective Catalytic Reduction (SCR) systems.

90 In order to fill the gap, the aim of this investigation was to quantify organic and
91 inorganic gas emissions (gas- and particle-phase) from a four-cylinder diesel engine
92 equipped with an urea-SCR system using Diesel or Biodiesel blends.

93

2. Materials and methods

In this study, we used an engine dynamometer following the European Steady
Cycle (ESC) testing cycle, in agreement with the Directive 1999/96/EC of the European
Parliament and the Directive of the December 13th, 1999 Council,¹⁷ which establishes
engine and dynamometer settings, and also NO_x and other pollutants emission limits.
The dynamometer used in this study has a power output of 440 kW at 6000 rpm and a
torque of 2334 Nm. The engine employed is in accordance with the Euro V standards,
using an urea-SCR after-treatment system. Table 1 specifies the engine details.

103

104

Table 1. Engine specifications, BR- model 2012.					
Specifications					
Emission	Euro V "Heavy Duty"/Proconve P7				
Configuration	4 cylinders, inline				
Displacement	4,8 liters				
Bore x Stroke	105 x 137 mm				
Combustion System	Direct injection				
Injection System	Common Rail Electronic				
Aspiration	TGV Intercooler				
Power Output	187hp (139,7kW) 2,200rpm				
Peak Torque	720Nm (73kgf.m) 1,200 ~ 1,600rpm				
Weight (dry)	426 kg				
Aftertreatment SCR					
Dimensions (H x L x W)	900 x 975 x 826 mm				

105

106 The emission data were sampled in the laboratory of vehicular emissions of the 107 Federal University of Parana –Curitiba/Brazil, employing an engine dynamometer driving cycle using LSD (Low Sulphur Diesel - 50 ppm sulphur content), ULSD (Ultra 108 109 Low Sulphur Diesel - 10 ppm sulphur content) and B20 (soybean biodiesel blended 110 (20%) with ULSD). The main difference between LSD and ULSD is their sulphur 111 content, which may affect SO₂ and particulate emissions. However, the cetane number 112 also differs and is considered a key fuel property comprising NMHC and CO emissions.9,18 113

114 Table 2 shows the quality parameters of the reference diesel fuels and the biodiesel 115 blend used in this research. The Standard Test Methods established by ASTM were 116 followed. The main properties having an influence on exhaust emissions are sulphur 117 content and cetane number, as will be discussed in the results section.

118

Tuble 2. Tuble Troperties of ESD and OESD aleser and D20 biodieser.				
Property	LSD	ULSD	B20	
Sulphur, mg/kg	24	4	6	
Cetane number	49.2	53.8	51	
Glow point (°C)	58.5	44.5	70.5	
Viscosity at 40°C (mm ² /s)	2.6	3.0	3.15	
Specific mass at 20°C (kg/m ³)	835.2	830.5	848.1	

119 Table 2. Fuel Properties of LSD and ULSD diesel and B20 biodiesel.

120

121 The gas emission data were obtained by a SESAM i60 FT, a Fourier Transform 122 InfraRed (FTIR) multi-component measurement system from AVL. Table 3 presents 123 some important technical characteristics of the FTIR analysis. The FTIR was calibrated 124 to detect specific hydrocarbons (HC), nitrogen compounds (NO, NO₂, N₂O and NH₃) 125 and other pollutants. It also calculates NO_x, total (HCD) and non-methane hydrocarbons (NMHC) concentrations. The HCD is the sum of all hydrocarbons that 126 127 FTIR can analyse using a method for diesel fuel (HCD = CH₄, C_2H_2 , C_2H_4 , C_2H_6 , C_3H_6 , 128 $C_{3}H_{8}$, $C_{4}H_{6}$, nC_{8} and AHC-aromatic hydrocarbons). The HCD expresses the total 129 hydrocarbons (HC) for diesel emission analysis. The NMHC comprises the HCD 130 concentration, except for the methane fraction.

131

132 Table 3. FTIR settings.

FTIR Spectrometer Data	
Sampling Rate	1 scan per second (1 Hz)
Data Rate	All measured gas components at 1 Hz
Spectral Resolution	0.5 cm^{-1}
Measurement Cell	Gas cell heated to 191 °C (375.8 °F)
Response Time	t ₁₀ to t ₉₀ within 1 s
Sample Flow Rate	10 l/min per stream
Detector Cooling	Liquid nitrogen, 50 ml/h
Zero/Purge Gas	Nitrogen / Synthetic Air, 0.6 – 1.5 l/min
Compressed Air	5 – 6 bar and max. 100 l/min per FTIR stream

3. Results and discussions

- **136** 3.1 Nitrogen Compounds
- 137

138 Analysis of Variance (ANOVA), normal probability plot of residuals and Bartlett's 139 test of homogeneity of variances were applied to the studied compounds. The statistical 140 analysis were performed using R software.¹⁹ A preliminary analysis showed that the 141 residuals have a normal distribution and a parametric behaviour. The Bartlett's test 142 presented, for almost all samples, p-values less than the significance level of 0.05, 143 confirming the homogeneity of sample variances. In conclusion, the analysis of 144 variance results are valid, except for C_2H_2 and C_2H_6 .

145 According to the analysis of variance results the means differ due to fuel and after-146 treatment system choice. To analyse the interactions between fuel and after-treatment 147 system, we applied the Tukey significant difference test. Differences between mean 148 values at a level of p < 0.05 (95% confidence level) were considered statistically 149 significant.²⁰

Our results, presented in Table 4, have shown that, for all studied fuels the use of
the SCR system presented statistically significant different means of nitrogen oxides
(NO_x), nitrogen monoxide (NO) and nitrogen dioxide (NO₂) emissions, compared to
results when the SCR system was not used. Quantitatively, the use of the SCR system
decreased NO_x, NO and NO₂ concentrations.

According to Chin et al.¹, some biodiesel blends may reduce emissions of regulated
pollutants, such as PM, CO, NMHC and CO₂. However, it usually increases fuel
consumption and NO_x emissions.

Only NO₂ emission means showed statistically significant differences between
LSD and ULSD fuels when the engine was not equipped with the SCR system.
However this trend was not observed between the ULSD and the B20 fuels. In contrast,
the use of different fuels statistically affected NO_x, NO and NO₂ emission means when
the engine was equipped with the SCR system, where the highest emissions were
observed for the ULSD and B20 fuels.

164 According to Chin et al.¹ and Agarwal and Das^{21} , a NO_x emission increase due to 165 biodiesel blend fuels use, is a result of some fuel properties, such as viscosity, and also 166 is a result of the advance in injection timing, temperature rise and abundance of oxygen 167 available in the combustion chamber.^{1,21} Viscosity interfere in the fuel nebulization generating different sizes of droplets in the combustion chamber. The burning
efficiency is higher with small droplets, due to a lower viscosity, leading a lower NO_x
emission.

171Despite the fact that the WHO^{22} has reported that sulphur content of fuels can172increase NO_x emissions, as it reduces catalyst efficiency, our results showed similar173concentrations to all tested fuels (scenarios without SCR system), although higher174concentrations using ULSD in comparison to LSD with the use of SCR system were175observed.

176

177 Table 4. Average and standard deviation of exhaust emissions for nitrogen compounds

	Low Sulfur Diesel		Ultra Low Sulfur Diesel		Biodiesel B20	
Pollutant	SCR off (±SD)	SCR on (±SD)	SCR off (±SD)	SCR on (±SD)	SCR off (±SD)	SCR on (±SD)
NO _x	7.55 ± 0.04	0.52 ± 0.02	7.66 ± 0.07	2.4 ± 0.8	7.6 ± 0.2	1.6 ± 0.4
NO	4.89 ± 0.02	0.34 ± 0.01	4.84 ± 0.03	1.5 ± 0.5	4.8 ± 0.1	0.98 ± 0.24
NO_2	0.06 ± 0.01	< M.D.C.	0.26 ± 0.04	0.15 ± 0.04	0.31 ± 0.07	0.06 ± 0.01
NH ₃	0.004 ± 0.002	0.07 ± 0.02	0.002 ± 0.001	0.007 ± 0.003	$\begin{array}{c} 0.0008 \pm \\ 0.0007 \end{array}$	0.006 ± 0.001
N_2O	0.0133 ± 0.0001	0.0434 ± 0.0003	0.0127 ± 0.0005	0.044 ± 0.004	0.013 ± 0.001	0.061 ± 0.008

178 (g/kWh) using SCR system on and off.

179 NO_x - Nitrogen Oxides, NO- Nitrogen Monoxide, NO₂- Nitrogen Dioxide, NH₃- Ammonia, N₂O 180 Nitrous Oxide.

181 * MDC (Minimal Detectable Concentration) is the detection limit of each gas component, determined as two times the standard deviation σ of zero gas measurement over 60 seconds.

183 Inferior to MDC: NO_2 – Nitrogen dioxide (MDC = 0,011 g/kWh).

184 185

186 While designed to reduce NO_x emissions, the SCR system may increase other
187 pollutants' emissions. As demonstrated in our study, the SCR system satisfies its
188 purpose of reducing NO_x emissions. However, it brings forth new problems, such as
189 higher emissions of N₂O, NH₃ and some hydrocarbons.

190 Table 4 shows an increase in ammonia emissions due to SCR system use. The only

191	increase considered	statistically	significant	(p<0.05)) was for LSD.
		2	<u> </u>	VI .	

192 On the other hand, while the engine was equipped with the SCR system, there is a

193 statistically significant difference between NH₃ emission means from LSD to B20 and

194 from LSD to ULSD. The NH₃ emission means for ULSD and B20 could not be

195 considered significantly different at a 95% confidence level.

Koebel et al.⁶ reported that the SCR system uses continuous urea injections
(ammonia content) to neutralize NO_x emissions, which may lead to an excess of urea,

called ammonia slip. It is therefore not unreasonable to assume that the ammonia slipmay be responsible for the higher NH₃ emissions observed.

When the injected urea solution fails to be completely decomposed below 200°C, it can produce ammonium nitrate (NH4NO3), cyanuric acid ((HNCO)3), and other compounds as sub-products.⁸ As a consequence, ammonia and ammonium salts have a relevant impact on the ecosystem, accounting for the modification of the atmosphere global radioactive balance, the reduction of atmospheric visibility, the acidification and eutrophication of the environment.²³

As has been reported by European Environment Agency²⁴, road transport contributes only 2% of total ammonia (NH₃) emissions, though it is a significant source from a local perspective in urban areas. Many studies²⁵⁻²⁹ reported that an increase in NH₃ emission has occurred due to introduction of vehicles equipped with catalytic converters and adoption of urea-SCR system.

The main source of anthropogenic N₂O is agriculture,³⁰ but some concern has
arisen due to new diesel exhaust after-treatment systems being responsible for N₂O
production, for example, the chemical reactions in urea-SCR system.³¹

In our experiment, the use of the SCR system increased N₂O concentrations for all studied fuels. With 95% confidence level, these increases can be considered statistically significant, with the highest increase observed for the B20 biodiesel blend (about 361%) and the lowest for the ULSD (about 83%). These results can be explained by the undesirable processes that may occur in the SCR systems, including several competitive, non selective reactions with oxygen that can produce secondary emission.³¹

221 While the engine was equipped with the SCR system, a statistically significant 222 increase of N_2O emission due to B20 biodiesel use was verified, in comparison with 223 ULSD and LSD fuels (p<0.05).

- **224** 3.2 Hydrocarbons
- 225

226 The FTIR equipment is also able to detect the non-methane hydrocarbons (NMHC)227 and hydrocarbons of diesel (HCD). The results are shown in Table 5.

228

229 Table 5. Average exhaust emissions for hydrocarbons compounds (g/kWh).

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Low Sulfur Diesel Ultra Low Sulfur Diesel Biodiesel B20
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Pollutant	SCR off (±SD)	SCR on (±SD)	SCR off (±SD)	SCR on (±SD)	SCR off (±SD)	SCR on (±SD)
NMHC	0.1888 ± 0.0002	0.1857 ± 0.0004	0.135 ± 0.003	0.159 ± 0.003	0.136 ± 0.007	0.164 ± 0.006
HCD	0.1917 ± 0.0004	0.1878 ± 0.0004	0.137 ± 0.003	0.161 ± 0.003	0.137 ± 0.007	0.166 ± 0.006
C_3H_6	0.0233 ± 0.0009	0.0236 ± 0.0002	0.012 ± 0.002	0.006 ± 0.001	0.0138 ± 0.0004	0.013 ± 0.003
C_2H_2	0.0142 ± 0.0003	0.0120 ± 0.0003	0.0125 ± 0.0008	0.0122 ± 0.0004	0.0104 ± 0.0006	0.0124 ± 0.0008
C_2H_6	0.0653 ± 0.0006	0.0673 ± 0.0007	0.064 ± 0.002	0.089 ± 0.003	0.068 ± 0.004	0.087 ± 0.002
C_3H_8	0.030 ± 0.001	0.0169 ± 0.0007	0.0276± 0.002	0.0281 ± 0.0008	0.0168 ± 0.0007	0.025 ± 0.005
CH ₄	0.0028 ± 0.0003	0.00213 ± 0.00003	0.0021 ± 0.0002	0.0023 ± 0.0001	0.00165 ± 0.00007	0.0022 ± 0.0004
НСНО	0.0285 ± 0.0007	0.0063 ± 0.0005	0.011 ± 0.002	0.0037 ± 0.0002	0.010 ± 0.004	0.006 ± 0.002
nC8	0.056 ± 0.001	0.0659 ± 0.0002	0.0204 ± 0.0005	0.024 ± 0.002	0.027 ± 0.002	0.027 ± 0.004

NMHC- Non-Methane Hydrocarbons, HCD- Hydrocarbons of Diesel, C3H6-Propylene, C2H2-230231 Acetylene, C₂H₆- Ethane, C₃H₈-Propane, CH₄ - Methane, HCHO- Formaldehyde and nC8- N-Octane. 232 233 Inferior to MDC: C_2H_4 - Ethene (MDC = 0,0173 g/kWh), C_4H_6 - 1, 3 Butadiene (MDC = 0,0666 g/kWh) 234 and AHC- Aromatic hydrocarbon (MDC = 0.0134 g/kWh). 235 236 The NMHC emission means were statistically different between LSD and ULSD 237 for both situations, SCR-on and SCR-off, showing a reduction of 30% for SCR off and 238 15% for SCR on. The influence of the SCR system in NMHC emissions means was 239 statistically significant only for ULSD and B20. The means increased by nearly 20% 240 using ULSD and B20 (p<0.05). Diesel hydrocarbons emissions (HCD) showed a similar trend to that observed for NMHC emissions described previously. 241

Fuels with a smaller cetane number has a higher ignition delay time, which "along
with the combustion of a partially premixed charge results in excessive emissions from
incomplete combustion, specifically total hydrocarbons (THC) and CO".¹⁸

Regarding recent changes on fuel properties, such as lower sulphur content in diesel and the use of biodiesel blends, considering measures of each hydrocarbon to engine not equipped with SCR system, the use of ULSD showed statistically significant difference on means in comparison to LSD to all hydrocarbons, with exception of ethane and acetylene (analysis of variance invalid). However, the only hydrocarbons showing significant differences on means (p<0.05) from ULSD to B20 were propane and n-octane, with decrease of propane and increase of n-octane.

252 Statistical treatment of data indicates that formaldehyde emissions were
253 significantly (p<0.05) lower (78%) with LSD and (59%) with ULSD due to SCR system
254 use. It also indicates that n-octane emissions were significantly (p<0.05) higher (18%)
255 with LSD due to SCR system use.

Besides the toxicity of some organic compounds like BTEX and HPA's, well
known as potential carcinogenic compounds, Atkinson³² pointed out that a variety of
hydrocarbons may lead to ozone production in low latitudes, through their reaction to
OH radicals in the presence of NO_x and SO₂.

260 The ground-level ozone is a well-known atmospheric pollutant, which can cause 261 several deleterious impacts on the environment and human health. In high 262 concentrations, the tropospheric O₃ can interfere with photosynthesis and the growing of some plant species.^{33,22} The latest European directive 2002/3/CE recommends that 263 at least 30 NMHCs (saturated, unsaturated or aromatic) should be measured.³⁴ As far 264 as ozone formation due to high NMHCs and SO_x emissions are concerned, the critical 265 266 situation in our study was that of LSD, which presented elevated NMHC and SO₂ emissions. 267

In this context, it is widely known that organic compounds participated in the
formation of secondary pollutants that may contribute to some of the undesirable
environmental effects associated with photochemical smog episodes.

Essentially, each compound has a different contribution due to the amount emitted and some properties that affect the secondary pollutants production during photochemical reactions. Some of these compounds are said to be more reactive than others. Consequently, the most reactive organic compounds should be addressed towards a strategy to reduce ozone and PAN (Peroxyacetylnitrate) exposure levels.³⁵

A ranking of most reactive organic compounds, based on ozone formation under 276 specific atmospheric conditions has been developed, the so-called reactivity scale. 277 Derwent et al.³⁵ created a reactivity scale for Northwestern Europe. They estimated the 278 279 Photochemical Ozone Creation Potentials (POCPs) and Photochemical PAN Creation 280 Potentials (PPCPs) for 120 organic compounds and their sensitivity to NO_x emissions taking ethylene (POCP = 100) and propylene (PPCP = 100), respectively, as the 281 282 reference compound. Table 6 presents the values calculated by Derwent et al. (1998).³⁵ 283 Table 6. Photochemical Ozone Creation Potential POCP and Photochemical PAN

284

Creation Potential

Organic Compounds	POCP	PPCP
Propylene	112.3	100
Formaldehyde	51.9	14.8
N-octane	45.3	42.9

Propane	17.6	13.7
Ethane	12.3	17.3
Acetylene	8.5	2.2
Methane	0.6	0.9
D		

Source: Derwent et al.³

287 Relating the results of Table 6 with our study, n-octane POCP is only 13% lower
288 than formaldehyde's one, while its PPCP is 65% higher than the formaldehyde one.
289 With regards to ozone and PAN formation, LSD fuel presented the higher
290 concentrations for the compounds with the higher POCP and PPCP values: propylene,
291 formaldehyde and n-octane.

292 Considering only the LSD fuel, it was statistically verified (p<0.05) an increase in
293 n-octane emission and a decrease in formaldehyde when the SCR system was used.
294 These results indicate a beneficial effect in ozone photochemical creation, as the
295 formaldehyde POCP is higher than n-octane one. In addition, as reported by WHO²²,
296 formaldehyde was classified as a carcinogenic compound.

The SCR system combined with ULSD or B20 has increased alkanes emissions,
however their POCP and PPCP are lower than those of formaldehyde, propylene and
n-octane. Therefore, the ULSD and B20 fuels are, apparently, a better alternative than
LSD, considering the hydrocarbons emissions and their photochemical potentials.

Recently Derwent et al.³⁶ developed a similar study applying the same models to
 create an activity scale for different emission sources of organic compounds. They
 indicated road transport-exhaust as the major contributor to POCP levels. Furthermore,
 Derwent et al.³⁷ made the same conclusion for secondary organic aerosol formation
 from organic compounds.

The POCP and PPCP analysis applied in our study is interesting since the
combination of megacities, atmospheric conditions and significant emissions of ozone
and PAN precursors can favour photochemical reactions in smog systems, creating
serious pollution episodes.

Regarding the use of the SCR system scenarios, the results are of similar magnitude
for all tested fuels. However, when the engine was not equipped with the SCR system,
the LSD showed higher emissions, with differences over 60% in comparison to ULSD,
with little difference between ULSD and B20.

Open literature describes decreases in aldehyde emissions from some biodiesel
fuels, in comparison to diesel.³⁸⁻⁴⁰ However, specifically with regard to formaldehyde,
some researchers observed an increase or no alteration in its emission.^{41-43,9} Tan et al.⁴⁴
showed an increase of formaldehyde emissions mainly for pure biodiesel fuel in
comparison to diesel, and showed little difference between diesel and B20 blend.

319 Taken together, this study showed that the emissions of NO and NO₂ while the 320 engine was equipped with the SCR system using the ESC cycle were lower and 321 statistically significant (p<0.05). However, the use of the SCR system produced 322 significantly increased concentrations of: N₂O for all studied fuels; NH₃ just for LSD; 323 and non-methane hydrocarbons (NMHC) and hydrocarbons of diesel (HCD) for ULSD 324 and B20. On the other hand, the use of SCR system significantly (p<0.05) supressed 325 formaldehyde emissions for LSD and ULSD fuels, having a beneficial impact since it 326 has a huge POCP and PPCP and is considered as a carcinogenic compound.

327 Soybean biodiesel blend used, in combination with the SCR system, can
328 successfully reduce harmful pollutant emissions such as NO_x, however, increases the
329 HCD production.

330

331332

4. Acknowledgement

333

This work was supported financially by the National Council for Scientific andTechnological Development (CNPq).

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- **337 5.** References
- 338

319 (1) Chin, J.; Batterman, S. A.; Northrop, W. F.; Bohac, S. V.; Assanis, D. N. Gaseous
340 and Particulate Emissions from Diesel Engines at Idle and under Load: Comparison of
341 Biodiesel Blend and Ultralow Sulfur Diesel Fuels. *Energy & Fuels.* 2012, 26, 6737–6748;
342 DOI 10.1021/ef300421h.
343 (2) U.S. EPA United States Environmental Protection Agency Technical Bulletin;

343 (2) U.S. EPA. United States Environmental Protection Agency Technical Bulletin:
344 Nitrogen oxides (NO_x) why and how they are controlled. Clean Air Technology Center
345 (MD-12), 57pp, 1999.

346 (3) Alkemade, U.; Schumann, B. Engines and exhaust after treatment systems for future automotive applications. *Solid State Ionics.* 2006, 177, 2291–2296; DOI 10.1016/j.ssi.2006.05.051

349 (4) Furfori, S.; Russo, N.; Fino, D.; Saracco, G.; Specchia, V. NO SCR reduction by hydrogen generated in line on perovskite-type catalysts for automotive diesel exhaust gas

- 351
 treatment.
 Chemical
 Engineering
 Science.
 2010,
 65,
 120–127;
 DOI

 352
 10.1016/j.ces.2009.01.065.
 10.1016/j.ces.2009.01.065.
 10.1016/j.ces.2009.01.065.
 10.1016/j.ces.2009.01.065.
- 353 (5) Miquel, P.; Granger, P.; Jagtap, N.; Umbarkar, S.; Dongare, M.; Dujardin, C. NO
 354 reduction under diesel exhaust conditions over Au/Al₂O₃ prepared by deposition355 precipitation method. *Journal of Molecular Catalysis A: Chemical.* 2010, 322, 90-97; DOI 10.1016/j.molcata.2010.02.024.
- 357 (6) Koebel, M.; Elsener, M.; Kleemann, M. Urea-SCR: a promising technique to reduce
 358 NO_x emissions from automotive diesel engines. *Catalysis Today*. 2000, 59, 335–345; DOI 10.1016/S0920-5861(00)00299-6.
- 360 (7) Hu, Y.; Griffiths, K.; Norton, P.R. Surface science studies of selective catalytic
 361 reduction of NO: Progress in the last ten years. *Surface Science*. 2009, 603, 1740–1750;
 362 DOI 10.1016/j.susc.2008.09.051.
- 363 (8) Jiang, L.; Ge, Y.; Shah, A. N.; He, C.; Liu, Z. Unregulated emissions from a diesel
 364 engine equipped with vanadium-based urea-SCR catalyst. *Journal of Environmental*365 *Sciences.* 2010, 22, 575–581; DOI 10.1016/S1001-0742(09)60148-0.
- 366 (9) Turrio-Baldassarri, L.; Battistelli, C. L.; Conti, L.; Crebelli, R.; De Berardis, B.;
 367 Iamiceli, A. L.; Gambino, M.; Iannaccone, S. Emission comparison of urban bus engine
 368 fueled with diesel oil and "biodiesel" blend. *The Science of the Total Environment.* 2004,
 369 327, 147–62; DOI 10.1016/j.scitotenv.2003.10.033.
- (10) Lobo, F. A.; Goveia, D.; Oliveira, A. P.; Romão, L. P. C.; Fraceto, L. F.; Dias Filho,
 N. L.; Rosa, A. H. Development of a method to determine Ni and Cd in biodiesel by graphite
 furnace atomic absorption spectrometry. *Fuel.* 2011, 90, 142-146; DOI
 10.1016/j.fuel.2010.09.009.
- 374 (11) Karavalakis, G.; Bakeas, E.; Stournas, S. Influence of oxidized biodiesel blends on
 375 regulated and unregulated emissions from a diesel passenger car. *Environmental Science*376 and Technology. 2010, 44, 5306-5312; DOI 10.1021/es100831j.
- 377 (12) Bakeas, E.; Karavalakis, G.; Stournas, S. Biodiesel emissions profile in modern diesel
 378 vehicles. Part 1: Effect of biodiesel origin on the criteria emissions. *The Science of the Total*379 *Environment.* 2011, 409, 1670–1676; DOI 10.1016/j.scitotenv.2011.01.024.
- 380 (13) Bermúdez, V.; Lujan, J. M.; Pla, B.; Linares, W. G. Comparative study of regulated
 381 and unregulated gaseous emissions during NEDC in a light-duty diesel engine fuelled with
 382 Fischer Tropsch and biodiesel fuels. *Biomass and Bioenergy*. 2011, 35, 789–798; DOI 10.1016/j.biombioe.2010.10.034.
- 384 (14) Demirbas, A. Biodiesel from oilgae, biofixation of carbon dioxide by microalgae: A
 385 solution to pollution problems. *Applied Energy.* 2011, 88, 3541–3547; DOI 10.1016/j.apenergy.2010.12.050.
- 387 (15) U.S. EPA. United States Environmental Protection Agency, 2013. Website
 388 http://www.epa.gov/otaq/standards/heavy-duty/hdci-exhaust.htm.
- 389 (16) Koebel, M.; Madia, G.; Elsener, M. Selective catalytic reduction of NO and NO₂ at low temperatures. *Catalysis Today.* 2002, 73, 239–247; DOI 10.1016/S0920-391 5861(02)00006-8.
- 392 (17) EC (European Commission). European Commission Report from the Commission to
 393 the European Parliament and the Council: Quality of petrol and diesel fuel used for road
 394 transport in the European Union: Sixth annual report (Reporting year 2007), 15pp.
- 395 (18) Lilik, G. K.; Boehman, L. Advanced Diesel Combustion of a High Cetane Number
 396 Fuel with Low Hydrocarbon and Carbon Monoxide Emissions. *Energy & Fuels.* 2011, 25, 1444–1456; DOI 10.1021/ef101653h.
- 398 (19) R Core Team. 2014. R: A language and environment for statistical computing. R
 399 Foundation for Statistical Computing, Vienna, Austria. Website http://www.R-project.org/.
- 400 (20) Montgomery, D. C. Design and Analysis of Experiments. 5th Edition. John Willey &
 401 Sons, INC. 699 pp, 2011.
- 402 (21) Agarwal, A. K.; Das, L. M. Biodiesel development and characterization for use as a
 403 fuel in compression ignition engines. *Journal of Engineering for Gas Turbines and Power*.
- **404 2001**, 123, 440-447; DOI 10.1115/1.1364522.

- 405 (22) WHO. World Health Organization. Health effects of transport-related air pollution.
 406 Copenhagen: WHO Regional Office for Europe, 205p, 2005.
- 407 (23) Reche, C.; Viana, M.; Pandolfi, M.; Alastuey, A.; Moreno, T.; Amato, F.; Ripoll,
 408 A.; Querol, X. Urban NH₃ levels and sources in a Mediterranean environment. *Atmospheric*409 *Environment.* 2012, 57, 153–164; DOI 10.1016/j.atmosenv.2012.04.021.
- 410 (24) EEA (European Environment Agency). European Union Emission Inventory Report
 411 1990 e 2009 under the UNECE convention on long-range transboundary air pollution. EEA
- 411 1990 e 2009 under the UNECE convention on long-range transboundary air pollution. EEA
 412 Technical Report 9, 2011. Website http://www.eea.europa.eu/publications/eu-emissioninventory-report-lrtap.
- 414 (25) Sutton, M. A.; Dragosits, U.; Tang, Y. S.; Fowler, D. Ammonia emissions from non415 agricultural sources in the UK. *Atmospheric Environment.* 2000, 34, 855–869; DOI
 416 10.1016/S1352-2310(99)00362-3.
- 417 (26) Perrino, C.; Catrambone, M.; Menno, A. Di; Bucchianico, D.; Allegrini, I. Gaseous
 418 ammonia in the urban area of Rome, Italy and its relationship with traffic emissions.
 419 Atmospheric Environment. 2002, 36, 5385–5394; DOI 10.1016/S1352-2310(02)00469-7.
- 420 (27) Cape, J. N.; Tang, Y. S.; van Dijk, N.; Love, L.; Sutton, M.; Palmer, S. C. F.
 421 Concentrations of ammonia and nitrogen dioxide at roadside verges, and their contribution
 422 to nitrogen deposition. *Environmental Pollution*. 2004, 132, 469–478; DOI 10.1016/j.envpol.2004.05.009.
- 424 (28) Burgard, D. A.; Bishop, G.; Stedman, D. H. Remote sensing of ammonia and sulfur
 425 dioxide from on-road light duty vehicles. *Environmental Science & Technology*. 2006, 40,
 426 7018–22; DOI 10.1021/es061161r.
- 427 (29) Kean, A. J.; Littlejohn, D.; Ban-Weiss, G. A.; Harley, R. A.; Kirchstetter, T. W.;
 428 Lunden, M. M. Trends in on-road vehicle emissions of ammonia. *Atmospheric Environment.*429 2009, 43, 1565–1570; DOI <u>10.1016/j.atmosenv.2008.09.085</u>.
- 430 (30) Wang, Z.; Zheng, H.; Luo, Y.; Deng, X.; Herbert, S.; Xing, B. Characterization and
 431 influence of biochars on nitrous oxide emission from agricultural soil. *Environmental*432 *Pollution.* 2013, 174, 289-296; DOI 10.1016/j.envpol.2012.12.003.
- 433 (31) Majewski, W. A. 2005. Selective catalytic reduction. Website
 434 http://www.dieselnet.com/tech/cat_scr.php.
- 435 (32) Atkinson, R. Atmospheric chemistry of VOCs and NO_x. *Atmospheric Environment*.
 436 2000, 34, 2063-2101; DOI 10.1016/S1352-2310(99)00460-4.
- 437 (33) Susaya, J.; Kim, K.; Shon, Z. Demonstration of long-term increases in tropospheric
 438 O₃ levels: Causes and potential impacts. *Atmospheric Environment.* 2013, 92 (11), 1520–
 439 1528; DOI 10.1016/j.chemosphere.2013.04.01.
- 440 (34) Arsene, C.; Bougiatioti, A.; Mihalopoulos, N. Sources and variability of non-methane
 441 hydrocarbons in the eastern Mediterranean. *Global Nest Journal.* 2009, 11 (3), 333-340;
 442 DOI 10.1007/978-1-4020-6429-6 14.
- 443 (35) Derwent, R. G.; Jenkin, M. E.; Saunders, S. M.; Pilling, M. J. Photochemical ozone
 444 creation potentials for organic compounds in northwest Europe calculated with a master
 445 chemical mechanism. *Atmospheric Environment.* 1998, 32, 2429–2441; DOI
 446 10.1016/S1352-2310(98)00053-3.
- 447 (36) Derwent, R. G.; Jenkin, M. E.; Passant, N. R.; Pilling, M. J. Photochemical ozone
 448 creation potentials (POCPs) for different emission sources of organic compounds under
 449 European conditions estimated with a Master Chemical Mechanism. *Atmospheric*450 *Environment.* 2007, 41, 2570–2579; DOI 10.1016/j.atmosenv.2006.11.019.
- 451 (37) Derwent, R. G.; Jenkin, M. E.; Utembe, S. R.; Shallcross, D. E.; Murrells, T. P.;
 452 Passant, N. R. Secondary organic aerosol formation from a large number of reactive man453 made organic compounds. *Science of the Total Environment.* 2010, 408, 3374–3381; DOI
 454 10.1016/j.scitotenv.2010.04.013.
- 455 (38) Sharp, C. A.; Howell, S. A.; Jobe, J. Effect of Biodiesel Fuels on Transient Emissions
 456 from Modern Diesel Engines, Part II, Unregulated Emissions and Chemical
 457 Characterization. *SAE Technical.* 2000, 01-1968, DOI: 10.4271/2000-01-1968.

- 458 (39) Peng, C.; Yang, H.; Lan, C.; Chien, S. Effects of the biodiesel blend fuel on aldehyde
 459 emissions from diesel engine exhaust. *Atmospheric Environment.* 2008, 42, 906–915; DOI 10.1016/j.atmosenv.2007.10.016.
- 461 (40) Ratcliff, M. A.; Dane, A. J.; Williams, A; Ireland, J.; Luecke, J.; McCormick, R. L.;
 462 Voorhees, K. J. Diesel Particle Filter and Fuel Effects on Heavy-Duty Diesel Engine
 463 Emissions. *Environmental Science & Technology.* 2010, 44, 8343–8349; DOI 10.1021/es1008032.
- 465 (41) Corkwell, K. C.; Jackson, M. M.; Daly, D. T. Review of Exhaust Emissions of
 466 Compression Ignition Engines Operating on E Diesel Fuel Blends. *SAE Technical.* 2003,
 467 01-3283; DOI 10.4271/2003-01-3283.
- 468 (42) He, B. Q.; Wang, J. X.; Yan, X. G.; Tian, X.; Chen, H. Study on Combustion and
 469 Emission Characteristics of Diesel Engines Using Ethanol Blended Diesel Fuels. *SAE*470 *Technical.* 2003, 01-0762; DOI 10.4271/2003-01-0762.
- 471 (43) Correa, S. M.; Arbilla, G. Carbonyl emissions in diesel and biodiesel exhaust.
 472 *Atmospheric Environment.* 2008, 42 (4), 769-775; DOI 10.1016/j.atmosenv.2007.09.073.
- 473 (44) Tan, P. Q.; Hu, Z. Y.; Lou, D. M. FTIR detection of unregulated emissions from a
- **474** diesel engine with biodiesel fuel. *Spectroscopy and Spectral Analysis.* **2012**, 32 (2), 360– **475** 363; DOI 10.3964/j.issn.1000-0593(2012)02-0360-04