Reduction of Low Temperature Engine Pollutants by Understanding the Exhaust Species Interactions in a Diesel Oxidation Catalyst

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10

11 **Abstract**

12 The interactions between exhaust gas species and their effect (promotion or inhibition) on the light-off

13 and activity of a diesel oxidation catalyst (DOC) for the removal of pollutants are studied, using actual

14 engine exhaust gases from the combustion of diesel, alternative fuels (rapeseed methyl ester and gas-

15 to-liquid fuel) and diesel/propane dual fuel combustion. The activity of the catalyst was recorded

16 during a heating temperature ramp where carbon monoxide (CO) and hydrocarbon (HC) light-off

17 curves were obtained. From the catalyst activity tests, it was found that the presence of species

18 including CO, medium-heavy HC, alkenes, alkanes and NO_x and their concentration influence the

19 catalyst ability to reduce CO and total HC emissions before release to the atmosphere. CO could

20 inhibit itself and other species oxidation (e.g. light and medium-heavy hydrocarbons) while suffering

21 from competitive adsorption with NO. Hydrocarbon species were also found to inhibit their own

22 oxidation as well as CO through adsorption competition. On the other hand, NO_2 was found to

23 promote low temperature HC oxidation through its partial reduction, forming NO. The understanding

24 of these exhaust species interactions within the DOC could aid the design of an efficient

25 aftertreatment system for the removal of diesel exhaust pollutants.

26
27 **Keywords**
28 DOC; CO/HC light-off; alternative fuels; exhaust species interactions
29

Introduction

Recently, the International Agency for Research on Cancer (IARC) changed the classification of diesel engine exhaust from *possibly carcinogenic to human* (Group 2A) in 1988 to *carcinogenic to human* (Group 1) in 2012 [1]. This classification justifies the ever tightening emission legislations imposed on diesel cars affecting the air pollution. The trend for this legislation is now to focus more on cold start and low to medium load emissions which can be produced in the urban environment when low vehicle speed, stop/start and idle phases are frequent [2]. It is also in these urban environments that humans are more exposed to vehicle exhaust gas. Several strategies have been developed in order to reduce the quantity of pollutants emitted to the atmosphere and among them is the use of the diesel oxidation catalyst (DOC) which oxidises CO and HC to CO_2 and water before release to the atmosphere. Another potential role of the DOC is to oxidise NO to NO_2 for use in passive regeneration of the diesel particulate filter (DPF) downstream of the DOC in the engine exhaust [3] or to enhance NO_x reduction in the selective catalytic reduction (SCR) process [4]. The limiting factor for the DOC efficiency is its light-off temperature, i.e. the temperature below which the oxidation is kinetically limited [5], which can reduce its efficiency, especially during urban operating conditions.

Other parameters also affecting the catalyst efficiency, apart from the temperature, are the exhaust gas residence time within the catalyst [6], the oxygen availability, the choice of catalytic components and loadings used to promote the oxidation reactions [7] and the ageing state of the coating and catalyst substrate caused by the thermal and chemical environment. Finally, the presence and concentrations of some exhaust species and their interactions with each other can also affect the catalyst behaviour, promoting or inhibiting oxidation, especially at low temperatures. Therefore, at an engine operating condition where the exhaust temperature is limited, pollutant removal could still take place, by encouraging promoting effects and limiting inhibition of DOC activity, which can be achieved with a clear understanding of these species interactions.

These interactions between exhaust components and their capacity to be oxidised have been researched and modelled using synthetic mixtures of gases to represent engine exhaust gas in order to understand the kinetics and oxidation mechanisms within the oxidation catalyst [7-13]. The majority of these studies focus on tracing the behaviour of an individual or a limited number of exhaust gas species in a synthetic gas mixture with a known composition. However, a single exhaust species can have different oxidation behaviour depending on whether it is studied on its own or as part of a mixture [13]. To our understanding the number of studies using actual engine exhaust gas to examine the interactions between different exhaust components is limited and focused upon exhaust gases from conventional fuel combustion [14-16]. In this study, a single cylinder engine is fuelled with diesel, alternative fuels and advanced combustion modes [17] to produce a variety of exhaust gases. The use

65 of genuine exhaust gas allows a valuable comparison with vehicle emissions met on the road and

66 targeted by the emission legislations for air pollution. The efficiency of a DOC fed with these

67 different exhaust gas compositions is studied in order to gain a wider knowledge of the diversity and

68 priority among the various interactions existing between exhaust pollutant species (CO, hydrocarbon

69 species and NO_x) within the DOC. A better understanding of these interactions and their effect

70 (inhibition/promotion) on CO and HC oxidation can eventually support the design of the

71 aftertreatment system to enhance the removal of pollutants, most especially at low temperatures.

72 **<u>Methodology and Experimental Setup</u>**

73 **Methodology**

74 In order to obtain a diverse spectrum of exhaust gas compositions, a single cylinder diesel engine was

75 operated with different diesel fuels (diesel, Rapeseed Methyl Ester (RME), Gas-to-Liquid (GTL)) as

76 well as with propane/diesel in a dual fuel combustion mode. Exhaust gas recirculation (EGR) was also

77 used in most of the cases, as it is a well known engine calibration strategy currently used to reduce

78 NO_x. During the experiment, the engine was run at a steady state point (40% load at 1500 rpm). In the

79 case of the dual fuel combustion experiment, diesel was used as the pilot fuel and 0.2% or 0.5% of

80 gaseous propane (based on the volume of intake air replacement) was injected in the intake manifold.

81 These percentages were chosen in order to limit the usual loss in volumetric efficiency caused by the

82 replacement of a portion of intake air by a gaseous fuel while maintaining combustion stability.

83 The concentrations of the engine exhaust species used to study the DOC activity are shown in Figure

84 1 a) and b) and were recorded under the following operating conditions:

85 - **Diesel without EGR** (Ultra Low Sulfur Diesel) used as a reference fuel produces the lowest
86   level of CO and greatest level of NO.

87 - **Diesel with EGR** shows a significant reduction of NO_x emissions, while CO and THC
88   concentrations are higher compared to diesel without EGR.

89 - **RME with EGR** produces a similar level of CO compared to diesel without EGR with a
90   lower THC concentration. It is assumed that the exhaust content of aromatic hydrocarbons is
91   lower (RME does not contain aromatics) than those emitted using conventional diesel but the
92   hydrocarbons produced are on average heavier than for diesel [18,19].

93 - **GTL with EGR** produces a higher level of CO and a level of THC similar to the baseline.
94   However, it is assumed that these hydrocarbons contain less aromatics than those emitted
95   using conventional diesel, similarly to RME, as GTL does not contain aromatics. This
96   condition also produces the lowest NO_x emission compared to the other studied exhaust
97   gases.

98   • **Dual fuel combustion** shows a rather different exhaust gas composition compared to single
99   fuel combustion, with a dramatic increase in CO and THC exhaust concentration. It also
100   produces higher concentrations of light hydrocarbon species and especially light alkanes
101   (propane) and alkenes (ethylene and propylene). Medium-heavy hydrocarbon concentration is
102   similar to single fuel combustions, as those hydrocarbons are produced from the combustion
103   of diesel fuel. The use of propane also produces a lower level of NO but increases the level of
104   NO_2 compared to single fuel combustions.

105   For diesel fuel combustion, heavy hydrocarbons represent around 80% of the total hydrocarbon
106   concentration found in the engine exhaust. Diesel/propane dual fuel combustion produces a more
107   varied spectrum of hydrocarbon species as can be seen in Figure 1 b). Therefore, dual fuel combustion
108   exhaust gases are used to study individual hydrocarbon species interactions while for the other fuels,
109   only medium-heavy hydrocarbon oxidation is studied.

110   The exhaust system was fitted with a diesel oxidation catalyst and CO, HC species and NO_x
111   conversion efficiencies were monitored during a heating temperature ramp. For every tests, the
112   oxidation catalyst was subjected to a heating temperature ramp of around 2°C/min and an exhaust gas
113   space velocity of 35,000/h. Engine out exhaust concentrations were measured at the beginning and at
114   the end of each experiment in order to compare any changes in the exhaust gas composition during the
115   test. Throughout the experiment, the evolution of the DOC outlet exhaust gas concentration was
116   continuously recorded to calculate the conversion efficiency based on the inlet concentration.

117   The use of EGR in an engine decreases the oxygen content of the exhaust [20]. To eliminate the effect
118   of this reduced oxygen availability on the catalyst activity, oxygen was injected upstream of the DOC
119   when EGR was used, in order to maintain a similar level of oxygen at the inlet of the catalyst, for all
120   the tests.

121   **Experimental Setup**

122   The engine used in this study is a single cylinder, direct injection, diesel engine [17]. The test rig is
123   composed of a DC motor-generator dynamometer coupled to a load cell used to load and motor the
124   engine. Other standard engine test rig instruments are used to monitor intake air flow, temperatures
125   and pressures (air, oil, inlet manifold and exhaust system). The EGR system is externally cooled and
126   its flow is controlled by a valve. The EGR level is determined volumetrically as the percentage
127   reduction in volume flow rate of inlet air at a fixed engine operating point. The fuels used in the study
128   were supplied by Shell Global Solutions UK and their properties can be found in [17].

129   A schematic of the experimental setup used in this study is shown in the Supporting Information
130   section (Figure S1). A MultiGas 2030, FTIR spectrometry based analyser was used for the
131   measurement of gaseous emissions (CO, HC and NO_x). A temperature controlled line was used to

132 direct a portion of the exhaust gas towards the catalyst placed in a furnace whose temperature was
133 externally controlled. The sampling line temperature was maintained at 150°C to limit the
134 condensation of hydrocarbons and water that could affect the measurements. The exhaust gas oxygen
135 content was measured using an AVL DiGas analyser fitted with an electrochemical oxygen sensor.
136 The measurement determined the quantity of oxygen to be added to the sample flow throughout the
137 experiment, to equal the 15% oxygen concentration found in the engine exhaust gas when no EGR
138 was used. The diesel oxidation catalyst used in this study is a 120 g/ft^3 Platinum/Palladium (weight
139 ratio 1:1) with alumina and zeolite washcoat (2.6 g/in^3 loading) coated on a cordierite honeycomb
140 monolith (25.4 mm x 91.4 mm) of 400 cells per in^2 and 4.3 mil wall thickness.

141 **<u>Results and Discussions</u>**

142 **CO Oxidation**

143 The start of CO oxidation in the DOC is dependent on its concentration in the exhaust gas (Figure 2).
144 As CO concentration increases (except for diesel combustion without EGR), its light-off is delayed
145 towards higher temperatures. CO can strongly adsorb onto active sites at low temperatures and cover
146 the majority of the catalyst surface [7,8,14]. This, in return, limits oxygen access to the catalytic sites,
147 preventing the start of CO oxidation which is directly released to the atmosphere. This CO self
148 inhibition is reduced under lower concentration of CO, as can be seen in the case of RME exhaust gas.

149 The presence of hydrocarbons can also affect CO oxidation by competing for the same active sites.
150 This inhibition can be altered as some of the hydrocarbons are adsorbed onto the zeolites coated on
151 the catalyst, limiting the quantity of tailpipe hydrocarbons emitted at low temperature. The lower HC
152 concentration recorded in the exhaust gas from RME combustion (Figure 1 a)) can also reduce the
153 potential inhibition from CO and HC competition and allows earlier CO light-off. In dual fuel
154 combustion, at 130°C-140°C, both CO and light alkenes (Figures 2 and 3) show some limited
155 conversion (up to 15% conversion) which could highlight mutual inhibition as, similarly to CO, light
156 alkenes strongly adsorb on active sites, due to their double bonds [9,21]. The greater light alkene
157 concentration recorded in dual fuel combustion (Figure 1 b)) would therefore affect more CO light-
158 off, as can be noticed in Figure 2. It is unlikely that propane has any influence on CO light-off as,
159 when its oxidation starts, CO is already fully oxidised. Moreover, light alkanes adsorb poorly on
160 metallic surfaces [11] and therefore are not supposed to compete to a large extent with respect to CO
161 strong adsorption.

162 NO$_x$ species can also compete with CO for adsorption and potentially inhibit CO oxidation [8,10]. It
163 can be noticed that the exhaust gases containing the lowest NO concentrations are the ones showing
164 an earlier CO light-off (RME, GTL and Diesel combustion with EGR). On the opposite, the exhaust
165 gas from the combustion of Diesel without EGR contains the highest level of NO and the lowest level

of CO while showing a rather delayed start in CO oxidation. CO inhibition from NO is especially visible for diesel without EGR exhaust gas as CO presents some limited conversion between 50°C and 110°C (5% oxidation) before the reaction rate increases (Figure 2). This shows that even if CO oxidation starts at similar temperatures for diesel without EGR exhaust gas compared to the other exhaust gases, it remains limited to just a few percentage of conversion for some time before increasing again. The comparison of CO oxidation from diesel combustion with and without EGR confirms that a higher level of NO (diesel without EGR) seems to be more detrimental on CO light-off than a higher level of CO and HC (with zeolites present) for these ranges of concentrations. As can be noticed from the NO_x catalyst outlet concentration during the experiment (Figure 4), all exhaust gases show an increase of NO at low temperature from NO_2 partial reduction with hydrocarbons, except for diesel without EGR exhaust gas (NO remains constant before decreasing). It has been observed [14,22,23] that NO and NO_2 can easily adsorb onto Pt and Pd active sites and efficiently dissociate, producing a high coverage of adsorbed nitrogen species and oxygen atoms on the active sites and limiting other species adsorption (e.g. CO).

The competitive adsorption at low temperatures between CO, light alkene and NO_x species only happens once CO oxidation starts, as before that, CO was strongly covering most of the active sites. Once the temperature allows CO oxidation to start, its oxidation products desorb, freeing some active sites on which CO, as well as NO and alkenes try to adsorb [10]. This adsorption competition eventually reduces CO access to the active sites and limits its oxidation (5-10% conversion efficiency) over a certain range of temperatures until the other adsorbed species react and desorb, freeing the active sites for more CO to adsorb again, allowing the oxidation rate to increase.

While CO oxidation in the DOC is kinetically limited at low temperatures when exhaust gases from the combustion of RME, GTL and diesel with EGR are used, for the exhaust gas from the combustion of diesel without EGR and dual fuel combustion, the limiting factor is CO access to the active sites, as previously mentioned. Thus, once CO overcomes the inhibitions and adsorbs on the active sites at higher temperatures, its oxidation shows greater reaction rates as most of the catalytic sites were already kinetically active for CO oxidation. Another reason for the sharp increase in CO conversion efficiency noticed in dual fuel combustion exhaust gas is that CO oxidation reaction is exothermic. As CO concentration in dual fuel combustion is much higher than in the other exhaust gases, it involves a greater heat release during the reaction. This increases the catalyst local temperature once the oxidation starts and especially increases the catalytic site activity which would translate into greater reaction rate. As a result, the maximum CO conversion efficiency in dual fuel combustion is reached at similar temperatures to that of the exhaust gases from the other operating conditions, even though the light-off was considerably later.

200 Figure 5 summarises the required temperatures for the different exhaust gases to reach 10%, 50% and
201 100% CO conversion efficiency in the DOC. The graph shows that the exhaust gas with an early start
202 of oxidation is not necessarily the one that will reach its maximum conversion first and that, after the
203 oxidation has started, its reaction rate can still be affected by competition with other species. It can be
204 noticed that the more the start of oxidation is delayed, the greater the reaction rate is.

205 Finally, once CO oxidation starts (Figure 2), some inflections can be noticed in RME and diesel
206 combustion with EGR light-off curves at 120°C and 130°C respectively, as it has been noticed in
207 other studies [10]. This change in oxidation rate can be related to an increase in medium-heavy
208 hydrocarbon conversion happening at the same temperature (Figure 6). Once hydrocarbons are
209 released from the zeolites and become active for oxidation, they compete for the same sites as CO,
210 limiting its oxidation. This is particularly affecting RME as its hydrocarbon oxidation starts at lower
211 temperatures and the hydrocarbons produced are on average heavier compared to other exhaust gases,
212 possibly affecting CO accessibility to the active sites. Therefore, the competition for active sites takes
213 place when the catalyst is not yet able to fully oxidise CO (only 55% conversion efficiency), affecting
214 more strongly CO oxidation. This inflection in CO reaction rate can also be noticed for diesel with
215 EGR. However, this effect is to a lower extent as hydrocarbons oxidise at higher temperatures
216 (130°C), when more catalytic sites are available and active for both CO and HC oxidation, reducing
217 the inhibition effect on CO. A minor slowdown in CO conversion can also be noticed at 135°C for
218 diesel without EGR and 150°C for 0.2% propane, when hydrocarbon oxidation starts to increase. This
219 effect cannot be noticed for the other exhaust gases due to either lower hydrocarbon concentration or
220 hydrocarbon conversion starting at higher temperatures when a greater proportion of catalytic sites is
221 already active for CO oxidation.

222 A schematic summarising exhaust species interactions affecting CO oxidation as presented above can
223 be found as Figure S2 in the Supporting Information section of this paper.

HC Oxidation

Medium-Heavy Hydrocarbons

226 The oxidation catalyst used in this study contains zeolites which can trap medium-heavy
227 hydrocarbons at low temperature and release them later on. Therefore, any low temperature
228 conversion can be considered as 'virtual' conversion as the hydrocarbons are not actually oxidised but
229 only removed from the exhaust gas and stored momentarily within the zeolites [24].

230 In Figure 6, it can be noticed that zeolites behave similarly for most of the exhaust gases used in the
231 study, showing 60%-75% conversion efficiencies at 70°C. The light-off curve from the 0.5% propane
232 dual fuel combustion behaves differently even though it contains a similar concentration of medium-
233 heavy hydrocarbons. This highlights some strong inhibition effects as zeolite trapping efficiency is

234 reduced. A comparison between the exhaust composition from the combustion using 0.2% and 0.5%
235 propane in the engine shows higher CO and light HC species concentration for 0.5% propane (Figure
236 1 b)). Nevertheless, these exhaust species would not have a direct deactivation effect on the zeolites
237 activity as they are short hydrocarbon chains and therefore not trapped by zeolites. Further
238 investigations are required in that case, in order to understand what would cause such a great
239 inhibition in zeolites trapping.

240 Once the trapped hydrocarbons are released from the zeolites, different oxidation behaviours are
241 recorded as the reaction activation energy and rate depend on the hydrocarbon nature, molecular
242 structure (saturated, unsaturated, cyclic, aromatics, etc.) and chain length which influence their
243 adsorption strength [12]. HC light-off curves from the combustion of RME and GTL show a plateau
244 in conversion efficiency while hydrocarbon conversions from the rest of the studied fuels show a drop
245 in conversion efficiency. This drop can be explained by the higher concentration of aromatic
246 hydrocarbons in comparison to RME and GTL. Aromatic hydrocarbons require on average higher
247 temperatures to get oxidised as they are more weakly adsorbed due to their ring shape and can suffer
248 from inhibition from other species [13,21,25]. The depth of the loss in conversion efficiency depends
249 on how efficiently these hydrocarbons are adsorbed on the zeolites and oxidised as they are released.
250 The greater drop in conversion efficiency recorded is from dual fuel combustion exhaust gas which
251 could be due to the presence of more numerous short chain hydrocarbons limiting the oxidation of the
252 released hydrocarbons [9].

253 It has been previously reported that NO could compete with hydrocarbons for adsorption on the
254 catalytic sites and limit their oxidation [8,26,27]. Nevertheless, several studies have also investigated
255 the capacity of hydrocarbons to partially or completely reduce NO_2 to NO, depending on the type of
256 hydrocarbon used [10,16,26,28]. Thus, the availability of NO_2 within the catalyst should be
257 considered. It can be noticed on Figure 4 that no NO_2 is recorded at the catalyst outlet, even though
258 NO_2 is present in the feed gas while the NO level increases. At low temperature, hydrocarbons are
259 preferentially oxidised by NO_2. As the temperature increases, they start reacting preferably with
260 molecular oxygen from O_2 rather than NO_2, which can be noticed by NO_2 concentration increasing
261 again at the catalyst outlet and NO level reducing to its inlet value. However, it has to be pointed out
262 that, based on the stoichiometric ratio required for hydrocarbon oxidation with NO_2, this reaction
263 would account for only a limited oxidation of some hydrocarbons. This mechanism could explain the
264 opposite behaviour of CO and HC light-off for diesel combustion with and without EGR, as diesel
265 with EGR shows a greater drop in hydrocarbon conversion while having an early CO light-off
266 compared to diesel without EGR. Therefore, the higher NO concentration which is detrimental for CO
267 light-off could be promoting low temperature hydrocarbon oxidation through NO_2 production. NO
268 and NO_2 can also be stored on the alumina washcoat and form nitrites or nitrates which hydrocarbons
269 can react with [29,30]. This could also explain the improvement in the hydrocarbon conversion at low

270 temperature from diesel combustion without EGR and the lower NO concentration at the catalyst
271 outlet.

272 CO has also been considered as an inhibitor for other species oxidation, as previously mentioned.
273 From CO and HC light-off graphs (Figures 2 and 6), it seems that the start of hydrocarbon oxidation
274 happens when a portion of CO had already been oxidised, around 50 % of the inlet CO, for most of
275 the exhaust gases (Figure 7). Catalytic sites become available for hydrocarbons only after CO
276 oxidation is already well developed [13]. As CO conversion efficiency increases, its concentration
277 along the catalyst length decreases. This reduces CO inhibition effect on hydrocarbon oxidation by
278 allowing them to adsorb on available catalytic sites at the rear of the catalyst brick. Therefore, before
279 this threshold, as hydrocarbons are released from zeolites, most of the active sites can already be
280 occupied by strongly adsorbed CO, covering the catalyst and limiting the ability of released
281 hydrocarbons to adsorb on the catalytic sites.

282 *Light Hydrocarbons*

283 From propane, propylene and ethylene light-off curves, it can be observed that zeolites only
284 selectively trap medium-long chain hydrocarbons at low temperature (Figure 6).

285 Propylene and ethylene oxidation starts at lower temperature than medium-heavy hydrocarbons and
286 follows a steady increase until the maximum conversion efficiency, in a similar trend to CO light-off
287 curve. Light alkene hydrocarbons are short chains of unsaturated hydrocarbons and therefore are more
288 easily oxidised at lower temperatures than other hydrocarbons. Moreover, their oxidation does not
289 suffer from competitive adsorption with other hydrocarbons but it can still be inhibited by a high light
290 alkene concentration at low temperatures due to their strong adsorption onto the active sites [9,21],
291 similar to CO. Light alkenes can also suffer from adsorption competition with CO at low
292 temperatures, as previously mentioned, and it can be noticed that their oxidation commences only
293 after CO oxidation has started, as more active sites become available for them to adsorb. The fact that
294 they are less affected by competitive adsorption with other exhaust species justifies why the general
295 trend of their light-off curve is not much altered when changing from 0.2% to 0.5% propane exhaust
296 composition. The slightly delayed start of oxidation for 0.5% propane (around 20°C) can be justified
297 by its higher alkene and CO content. Nevertheless, similarly to CO, once the oxidation starts, a greater
298 reaction rate is recorded for 0.5% propane, leading to both dual fuel combustion exhaust gases
299 reaching their maximum conversion efficiency at similar temperatures.

300 Propane, an alkane hydrocarbon, is a stable component due to its carbon atom being saturated by
301 hydrogen, requiring more energy to be broken down and therefore greater temperatures to be
302 oxidised. As previously mentioned, light alkanes are poorly adsorb on active sites and can be affected
303 by the presence of other species. Therefore, their oxidation can only start when most of the other

304 exhaust components are already oxidised, especially strongly adsorbed CO and light alkenes. This is
305 the reason why propane light-off and reaction rate are greatly affected by the change of exhaust
306 composition and CO and hydrocarbon concentration increase from 0.2% to 0.5% propane.

307 **Strategies to Enhance DOC Low Temperature Pollutant Removal**

308 The above studied exhaust gas components and catalyst surface interactions allow the design of
309 strategies to limit the quantity of pollutants emitted to the atmosphere at low exhaust temperatures. To
310 promote low temperature CO oxidation, NO should be temporarily removed from the exhaust gas to
311 reduce CO-NO_x adsorption competition. The use of EGR strategy also allows an improvement in CO
312 light-off, by reducing NO engine out concentration, but care should be taken to limit possible fouling
313 of the EGR cooler at low temperature (use of high pressure loop or temporarily by-pass the cooler at
314 low temperature). Higher CO concentration can inhibit CO start of oxidation but can also increase its
315 reaction rate once the reaction has started, due to the heat produced by exothermic CO oxidation.
316 Moreover, trapping medium-heavy hydrocarbons through the use of zeolites can reduce the
317 competition for active sites between hydrocarbons and CO, affecting CO oxidation rates (inflection in
318 CO light-off curve) and delaying hydrocarbon start of oxidation. It is proposed in this study that
319 promoting CO oxidation at lower temperature would allow more active sites to be available for
320 adsorption, when hydrocarbons are released from the zeolites. In the meantime, hydrocarbons can
321 react with stored NO_x and partially reduce them at low-medium temperatures. Light alkene
322 hydrocarbons show early light-off and are not much affected by competition with other species apart
323 from themselves and CO. Light alkanes on the contrary require higher temperatures to be oxidised.
324 They do not affect other species oxidation but suffer from stronger adsorption of other species. Due to
325 their short carbon chains, they are not efficiently trapped in zeolites and therefore are released into the
326 atmosphere until the temperature is high enough to allow their oxidation. This study did not find any
327 strategies directly promoting their oxidation, but lower temperature oxidation of other species (e.g.
328 CO, light alkenes, etc.) can reduce competitive adsorption, increase active site availability and local
329 temperature which can indirectly promote light alkanes oxidation.

330 This study gives tools to qualitatively predict how efficiently CO and different hydrocarbon species
331 will be oxidised, based on the exhaust gas composition. It also gives an insight for the diesel exhaust
332 aftertreatment design requirements, for selective momentary control of some species in order to
333 promote the oxidation activity of the catalyst at low temperature. Thus, this study can generate ideas
334 for identifying synergies between engines calibration (i.e. promotion of light alkenes during
335 combustion instead of alkanes), fuels specifications and catalyst efficiency.

336

<u>Abbreviations</u>

CO   Carbon Monoxide

CO_2 Carbon Dioxide

DOC   Diesel Oxidation Catalyst

DPF   Diesel Particulate Filter

EGR   Exhaust Gas Recirculation

GTL   Gas to Liquid

HC   Hydrocarbons

NO   Nitric Oxide

NO_2 Nitrogen Dioxide

NO_x Nitrogen Oxides

RME   Rapeseed Methyl Ester

SCR   Selective Catalytic Reduction

THC   Total Hydrocarbons

<u>Supporting Information Available</u>

Figure S1 is a schematic of the experimental set up used in this study. Figure S2 represents a schematic summary of the different interactions affecting CO oxidation. Before the oxidation starts, the catalyst is covered by carbonaceous species from CO, strongly adsorbed onto the active sites, restricting oxygen access and limiting CO oxidation. Later, once CO starts desorbing, other species adsorb onto the free active sites (first light alkenes and then nitrogen oxides) limiting CO adsorption and oxidation. Finally, when the oxidation is developing, medium-heavy hydrocarbons can also start

adsorbing on the active sites and compete with CO, affecting the reaction rate of CO oxidation (inflection in the light-off curve at higher temperatures). This information is available free of charge via the Internet at http://pubs.acs.org.

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Figure Caption

454   **Figure 1:** Engine exhaust gas compositions (a) and exhaust hydrocarbon species concentration from

455   the engine operation on dual fuelling (0.2% and 0.5% propane with diesel) (b).

456   **Figure 2:** CO light-off curves from the different exhaust gas produced.

457   **Figure 3:** Light-off curves for light alkene (ethylene and propylene) and alkane (propane)

458   hydrocarbons for the dual fuel combustion mode.

459   **Figure 4:** NO and NO_2 catalyst outlet concentration over the temperature ramp (0°C representing

460   engine-out concentrations).

461   **Figure 5:** Catalyst inlet temperature required to reach 10%, 50% and 100% CO conversion for the

462   different tested fuels.

463   **Figure 6:** Medium-heavy hydrocarbon light-off curves.

464   **Figure 7:** Catalyst inlet temperature required for 50% CO oxidation and for medium-heavy
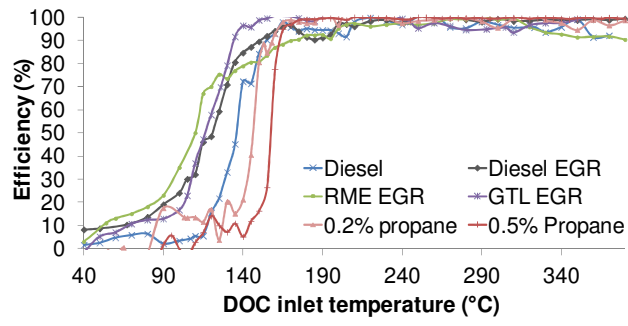
465   hydrocarbon start of oxidation.
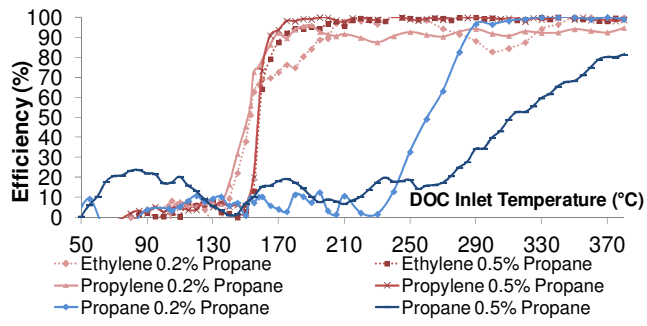
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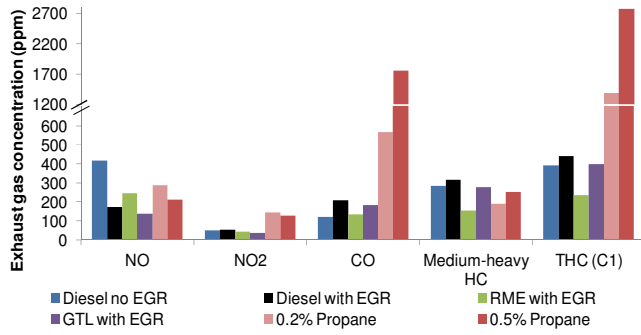
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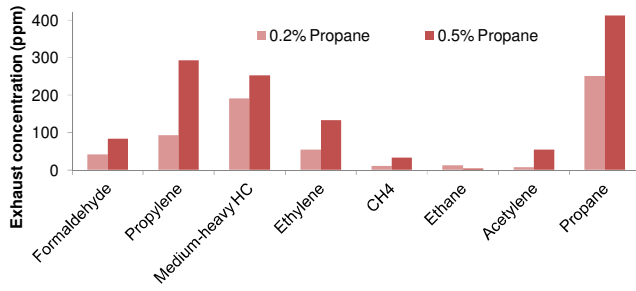
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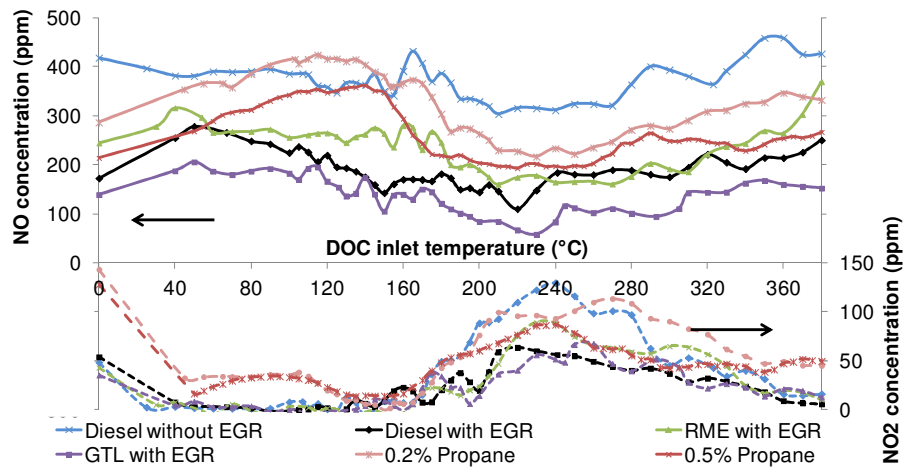
469 **Figure 1.**

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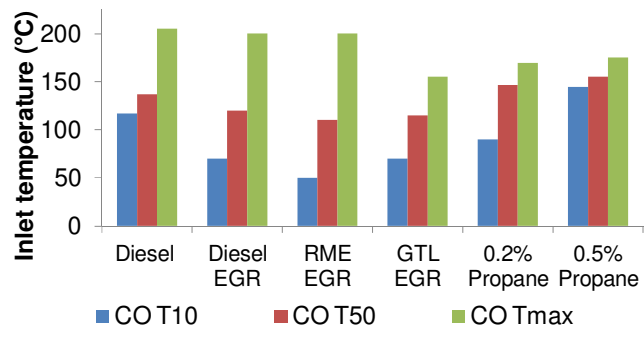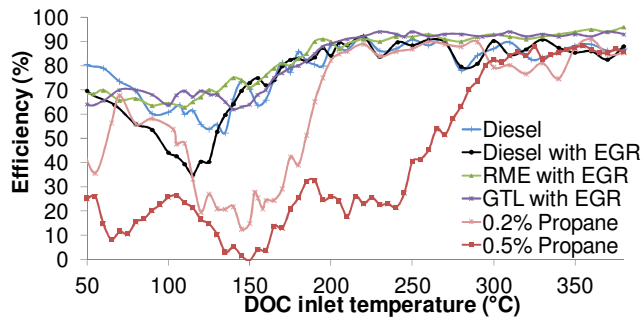471 **Figure 2.**

472

473   **Figure 3.**
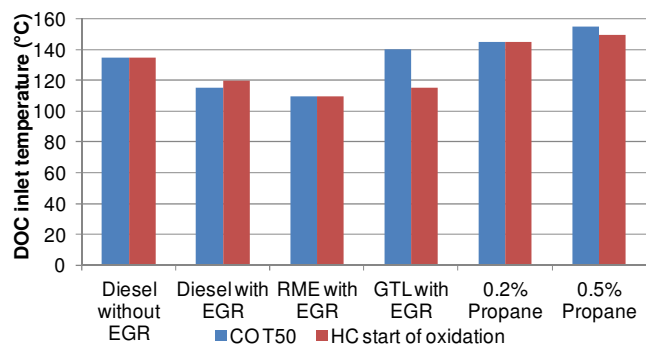
474

475 **Figure 4.**

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477   **Figure 5.**

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479   **Figure 6.**

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481   **Figure 7.**