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Additional Information

Pollutants emission and particle behavior in a pre-turbo aftertreatment light-duty diesel engine

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Abstract

Diesel particulate filters are a standard technology used in diesel engines in order to comply with actual and forthcoming regulations, regarding soot emissions and particulate matter in exhaust gases. In recent years, pre-turbo aftertreatment response has been investigated as opposed to the traditional aftertreatment location downstream from the turbine but just regarding engine performance. Previous studies do not deal in detail with gaseous and particle emission analysis in a pre-turbo aftertreatment configuration. This paper focuses on these topics. The gaseous pollutant and particle emissions have been assessed in a 4-cylinder, light-duty diesel EURO 4 engine typically used in European passenger car vehicles. Different steady-state operating points have been considered in order to extend the study over a wide range of operating conditions. Additionally, the New European Driving Cycle has been performed with the aim of reaching a comprehensive understanding of the aftertreatment dynamic response in terms of pollutant emissions. An increase in the amount of NO_2 converted from NO and a reduction in emitted CO have been found at low load steady-state operating conditions with pre-turbo aftertreatment placement. In driving cycle conditions, a shift from nucleation to accumulation mode particles have been found, being the filtration efficiency scarcely affected.

Keywords: Diesel engines, pre-turbo aftertreatment, pollutants emission, particle emission, DPF efficiency

1. Introduction

In the automotive sector, diesel engines are a major source of particulate matter (PM) and nitrogen oxides (NO_x). It is well known that these pollutants are precursors for other chemical compounds, producing significant damage to the environment [1] and human health [2]. In recent years, different strategies have been developed to reduce pollutant emissions: exhaust gas recirculation (EGR) [3], common-rail injection [4], improvements in the fuel-air mixture process, optimized engine control [5], exhaust manifolds with improved heat transfer losses [6], etc. Owing to the application of the results obtained in these studies, and the progress carried out in the combustion process [7], a reduction has been achieved in the soot mass emitted by diesel engines. In addition, with the introduction of diesel particulate filters (DPF) in the aftertreatment systems of the exhaust gas, soot mass emission has reduced by about

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99% [8]. However, a decrease in mass emission is not always accompanied by a reduction in particle emission. In this sense, the reduction in mass due to soot mass trapped at the DPF may be accompanied by an increase in the number of particles emitted with a size below 100 nm. According to Setter et al. [9], a new conceptualization of particle size distribution shows an increase in particle numbers below 100 nm for the new generation of engines. These small particles can pass more easily through the DPF [10], so that an increase in total particle number at the DPF outlet has been observed. Furthermore, due to the detrimental effect on human airways that this type of particles can have [11], emission regulations are becoming increasingly stringent on this issue [12].

Nowadays, most DPFs are made of ceramic substrates with a honeycomb wall-flow structure. These systems are characterized by high filtration efficiency and low penalty in fuel consumption due to the active regeneration strategies that are carried out when the soot load inside the DPF reaches certain level [13]. In this regard, studies are mainly conducted to increase the maximum amount of ash that the DPF can tolerate [14], to better understanding on the pressure drop dynamics across this element [15], to analyse heat transfer phenomena [16], to the development of advanced catalytic materials [17], to the analysis of the soot filtration efficiency [18], or to study the relative position of the aftertreatment system with respect to the turbine [19].

Recently, some studies have attempted to find the optimal position of the aftertreatment system by varying the diesel oxidation catalyst (DOC) and DPF placement. Windsor et al. [20] propose that these be located at the cylinder outlet before the turbocharger. In turbocharged engines, the aftertreatment system is traditionally placed downstream from the turbine, with the DOC upstream of the DPF. This configuration uses the NO_2 produced in the DOC for further oxidation of particulate matter in the DPF. Placing the aftertreatment system upstream from the turbine offer certain advantages such as clean high pressure EGR. In this sense, the pre-turbo aftertreatment approach ensures that the exhaust gas first passes through the DPF, and it is then recirculated into the intake manifold before passing through the turbocharger. It provides a soot-free exhaust gas to the high pressure EGR system thus the durability of the elements of this system is increased [21]. Other advantages of the pre-turbo aftertreatment configuration are the high temperatures reached at the DPF inlet, which can serve to promote passive regenerations and reduce time up to DOC light-off [22]; and the pressure drop reduction along the aftertreatment system due to the higher gas density [23]. On one hand, this effect leads to fuel savings because it mitigates the increase of exhaust back-pressure especially when soot accumulation within the DPF increases [24]. On the other hand, an additional reduction in fuel consumption is obtained due to the reduction of active regenerations required to clean the DPF [25].

2. Experimental setup

The measurements were carried out in the exhaust aftertreatment system of a 2.0 L, 4-cylinders, high-speed direct injection (HSDI) turbocharged diesel engine for passenger car application which complies with EURO 4 standards. Further detailed specifications of the engine characteristics and the aftertreatment system used are given in Table 1 and Table 2. The engine was installed in a fully equipped test cell with all the auxiliary devices required for appropriate

engine operation and control. The test cell was designed using the criteria detailed by Martyr and Plint [26]. Regarding engine fuel, ultralow sulfur diesel (USLD) has been used in all the test being its properties shown in Table 3.

The test bench was equipped with a 250 kW asynchronous dynamometer SHENCK®DYNAS Li250, which allows instant engine speed and torque control. In order to perform modifications to the engine operating parameters, the ETAS®INCA software was used to modify the engine calibration in an engine control unit (ECU). K type thermocouples and KISTLER®4045A5 piezoresistive sensors with the original cooler adaptor, whose main characteristics are shown in Table 4, were suitably located in the intake and exhaust zones of the engine. Fuel consumption was determined by two systems. The first consists of a fuel gravimetric system with an AVL®733S Dynamic Fuel Meter. The measurement device consists of a measuring vessel filled with the fuel and suspended on a balance system. Fuel consumption values are obtained by calculating the vessel's weight loss over time. Since the response time of this system might possibly be too long for measurement under transient operation, the fuel consumption signal provided by the ECU was calibrated in steady-state operating conditions. After calibration, the ECU was used as a second fuel consumption measurement system [27]. The Sensyflow®P Sensycon hot-plate anemometer system was used to measure the flow rate of the intake air mass. The measurement range of the anemometer is 0 – 720 kg/h.

Different equipment has been used in this study for measuring particle and pollutant emissions. For particle measurement, a TSI®EEPS (Engine Exhaust Particle Sizer) Spectrometer has been employed in order to obtain fast response particle measurements during dynamic cycles [28]. EEPS is capable to measure particle size distribution at a sample-rate of up to 1 Hz and providing a measurement range between 5.6 to 560 nm. A valve system was installed in order to provide measurements upstream and downstream of the DPF in steady-state conditions and allow filtration efficiency calculation.

A similar valve system concept was installed to assess the DOC conversion during the steady-state operating points. In this case, a HORIBA®MEXA 6000 F-TIR analyzer was used to provide a figure of unregulated pollutant emissions. Finally, an HORIBA®MEXA 7100 DEGR was installed at the end of the exhaust line downstream of the aftertreatment to measure tailpipe emission of regulated pollutants. Figure 1 shows the experimental setup for this study.

2.1. Operating mode

Extensive experimental work has been carried out in order to assess the effect of aftertreatment system placement in the exhaust line in terms of gaseous pollutants and particle emissions. The study was conducted comparing the pre-turbo aftertreatment configuration versus the post-turbo aftertreatment configuration (traditional). The original post-turbo aftertreatment configuration is composed of a close-coupled DOC followed by underfloor DOC and DPF. In the case of the pre-turbo aftertreatment configuration, the underfloor DOC and DPF are placed upstream of the turbine being the close-coupled DOC removed in this configuration.

In order to assess a wide range of different engine behavior, different steady-state operating points and dynamic tests were performed. As shown in Figure 2, five engine operating modes have been selected from the New European

Driven Cycle (NECD) for passenger cars. The selection criteria of these points were the flatlands occurring during the NEDC cycle. Thus, these points remain stable for 3 seconds, except K point, which is located on the first acceleration ramp in the EUDC (Extra Urban Driving Cycle). These points have different engine speed, load, EGR rate and SOI (Start of Injection), in order to obtain a representative range of the engine emission. Table 5 shows the main characteristics of the selected points.

2.2. Particle measurement method

The methodology used to measure the particle size distribution in transient conditions, which is also applicable to steady-state conditions, is performed in the laboratory according to Desantes et al. methodology [29], which is sketched in Figure 3.

The dilution system used for this study was a Dekati® Fine Particle Sampler FPS-4000 [30]. This system dilutes the sample in two stages. A porous tube (PTD) is used as the primary diluter, and subsequent an ejector diluter (ED) carries out as the secondary dilution (hot dilution + cold dilution), as shown Figure 3 from stage A→B and B→C ways. The particles within the size range of 10-50 nm are very unstable and are significantly affected by the dilution temperature due to the fact that they consist of a solid core that contains some volatile fractions. In this sense, the dilution ratio affects gas-to-particle conversion phenomena through the nucleation and adsorption of the soluble organic fractions (SOF) on the soot [31] or the supersaturate vapor condensation [32].

3. Results and discussion

Firstly, the assessment of the different partial steady-state operating points selected from the NEDC cycle was carried out. For this task, samples of the exhaust gas were taken after a given period of engine stabilization in order to guarantee a steady thermal and fluid dynamic state in each test. To ensure that the analyzed steady-state operating conditions were the same in both post-and pre-turbo aftertreatment configurations, tests were carried out at iso-torque conditions and with constant intake manifold pressure. Figure 4 shows some of the control variables measured in different configurations. The gaseous emission measurement was performed upstream and downstream from the DOC, while the particle measurement was carried out upstream and downstream from the DPF.

3.1. Gaseous emission. Steady-state operating points.

- Nitrogen Oxides: NO , NO_2 and NO_x

Firstly, Figure 5.A shows that total NO_x at both DOC inlet and outlet are practically invariant. However, when the analysis is focused on each compound separately (NO and NO_2), which are shown in Figure 5.B and Figure 5.C, components of total NO_x do not remain invariant but there are significant variations.

On one hand, Figure 4 shows that the EGR rate is slightly lowered in a pre-turbo configuration in the case of K and Q operating points with respect to post-turbo aftertreatment configuration, but the air to fuel ratio (AF) is

similar. It causes an increase of NO and a decrease of NO_2 at the DOC inlet due to the higher density of the air in the intake manifold induced by the lower EGR rate. On the other hand, the DOC outlet shows a decrease in NO and an increase in NO_2 , stronger in the pre-turbo than post-turbo configuration. As the conversion ratio of NO to NO_2 is strongly dependent on temperature, one can say that it was caused by the DOC inlet temperature, which at the pre-turbo configuration is higher because it is located upstream of the turbine and the expansion process has not occurred yet.

- Ammonia (NH_3), sulfur dioxide (SO_2) and dinitrogen oxide (N_2O).

The concentrations of these pollutants are negligible. Since the sulfur content in the fuel is less than 10 ppm, as mandated by the Spanish law, sulfur dioxide formation is negligible. Ammonia is usually present when SCR (Selective Catalytic Reduction) aftertreatment systems are used [33], using urea as an additive for NO_x reduction reaction but this is not the case in the present study. N_2O emission is affected by the presence of NH_3 and HCN . Some results show that the presence of NH_3 helps in the conversion of NO to N_2O , whereas HCN acts as the activating source of N_2O formation [34]. This compound has only relevance when the SCR system is used, so that its emission is not present as in the case of the ammonia.

- Carbon monoxide (CO) and total hydrocarbons (THC)

Carbon monoxide is a very harmful compound to human health, thus any reduction is advantageous for the environment. CO emissions occur due to non-complete combustion, and CO emissions are strongly linked with THC emissions, since both are results of low quality combustion.

Figure 6.A shows that CO emissions at the DOC inlet are similar in both configurations of the exhaust line. When it is measured downstream from the DOC, there is a very significant reduction at the F and J points in the case of the pre-turbo aftertreatment architecture. In this oxidation catalyst, the activation temperature is approximately $150^\circ C$. These two steady-state operating points do not reach this temperature level in the traditional post-turbo aftertreatment placement, so an increase of the DOC inlet temperature at these conditions brings a very pronounced decrease in the amount of CO emitted to the atmosphere.

Concerning THC emission, only DOC outlet data are available due the fact that HORIBA®MEXA 7100 was installed at the end of the exhaust line, and HORIBA®6000 F-TIR cannot provide THC results. Figure 6.B shows the results for THC DOC outlet emission with a similar trend than that obtained with CO .

- Light hydrocarbons (<7 Carbon atoms)

During the combustion process in diesel engines, the fuel does not mix completely with the required oxygen to burn the total amount of fuel. Unburned hydrocarbons are emitted as a result of the incomplete combustion process, in which part of the fuel is not oxidized. Figure 7 only shows the concentration of methane (CH_4) and iso-butene (*iso* - C_4H_8), as measured concentration of other hydrocarbons analyzed (ethylene (C_2H_4),

ethane (C_2H_6), propylene (C_3H_6), 1,3 Butadiene (1,3 - C_4H_6), n-Butane ($n - C_4H_{10}$), benzene (C_6H_6) and toluene (C_7H_8) are negligible and below the analyzer error (2 ppm). In this case, methane shows a flat trend emission despite the steady-state operating point performed. The iso-butene variations in different steady-state operating conditions demonstrate that the highest levels of hydrocarbons occur at low load and low speed. In such operating conditions, the combustion efficiency is lower because the too lean mixture damages flames stability in last stages of the injection process.

3.2. Steady-state operating points. Particle emission.

This section evaluates the change produced by the aftertreatment placement in terms of particle emissions. As shown in Figure 4, the engine performance is similar in pre-turbo and post-turbo aftertreatment configuration. Proof of this are the particle size distributions upstream of the DPF are practically the same independently on the exhaust line architecture, as shown in Figure 8.

When the DPF inlet particle emission is evaluated in detail by separating the nucleation and the accumulation mode [35], a reduction is observed in the pre-turbo aftertreatment configuration both in accumulation and nucleation modes.

Usually, the particle size grows along the exhaust line, mainly due to agglomeration and aggregation phenomena. These processes cause the growth of the particles due to the temperature drop along the exhaust system. It is related to the heat transfer losses taking place in the different elements placed in the exhaust line [36]. Figure 9 shows the temperature drop in each element of the aftertreatment architecture as a function of the steady-state operating point. Figure 9 clearly shows that in pre-turbo configuration the aftertreatment temperature is higher than in the post-turbo aftertreatment. Due to the fact that the turbine uses large amount of the exhaust gas energy [24], the largest temperature drop occurs in the expansion process. Therefore, the temperature at the aftertreatment system inlet with pre-turbo architecture is much higher than in post-turbo aftertreatment configuration. It contributes to avoid adsorption and condensation phenomena in the pre-turbo configuration under steady-state operation.

On one hand, the raw emission of particles comprising the accumulation mode, which are those with $D_p > 30$ nm, are reduced in the pre-turbo aftertreatment configuration (Figure 10.A). This reduction in the accumulation mode and the significant decrease occurring in the nucleation mode suggests that some particles finish their formation along the exhaust line in the post-turbo aftertreatment configuration but this does not happen in the pre-turbo aftertreatment configuration because of the shorter path up to the DPF [37]. On the other hand, homogenous nucleation occurs when enough free energy exists in the vapor to form a new spherical cluster. The nucleation can be modeled by the RKC (Reiss-Katz-Cohen) theory [38]. This theory states that the nucleation rate of supersaturated vapor is given by:

$$J = \frac{a}{\rho} \sqrt{\left(\frac{2N_A^3 \sigma M}{\pi}\right) \left(\frac{S P_e}{RT}\right)^2} \exp\left[\frac{-16\pi N_A}{3(\ln S)^2} \left(\frac{M}{\rho}\right)^2 \left(\frac{\sigma}{RT}\right)^3\right] \quad (1)$$

In equation (1), J is the rate of nucleation, N_A is the Avogadro's number, P_e is the equilibrium vapor pressure, T is the gas temperature, M represents the molecular weight of the vapor, ρ is the liquid density, σ is the surface tension, R is the gas constant, a is the condensation coefficient and S is the supersaturation, which is defined as $S = P/P_e$, being P the local pressure.

Pressure and temperature of the exhaust gas at DPF inlet are higher in the pre-turbo than in the post-turbo aftertreatment configuration, as shown in Figure 9. An increase in these two variables establishes a reduction in the nucleation rate as it can be concluded from Figure 10.A. In the case of the post-turbo aftertreatment placement, the adsorption and condensation phenomena of organic compounds of high molecular weight, such as unburned hydrocarbons, oxygenated hydrocarbons or polycyclic aromatic hydrocarbons, cause an overall increase of the particles diameter and also the generation of new particles, according to the sketch shown in Figure 11. However, these phenomena are not taking place in the pre-turbo aftertreatment configuration under steady-state operation, since the particles perform a shorter route in the exhaust line in comparison to the post-turbo aftertreatment configuration. Consequently the nucleation mode particles may be not still formed passing across the DPF shaped as hydrocarbons. Hence the overall balance shows a reduction in cores percentage in the pre-turbo aftertreatment configuration (Figure 10.B) and an increase of the geometric mean diameter (GMD) in particle size distribution (Figure 10.C) with respect to the post-turbo aftertreatment configuration.

The particle size distribution at DPF outlet vary much more than at DPF inlet between both aftertreatment configurations. Nevertheless, the filtration efficiency, which is shown in Figure 12, is very high in both configurations and inside the well-known characteristics of wall-flow DPFs, being the minimum measured value at discrete diameters in 97%. At first glance, as filtration efficiency indicates, the DPF acts similarly in both configurations in a range between 20 and 150 nm. In contrast, the DPF filtration efficiency suffers a decrease for particles smaller than 20 nm in the pre-turbo aftertreatment configuration in most operating points. The reason can be found in the passive regeneration process according to Beatrice et al. [39]. To verify the occurrence of passive regeneration the DPF is weighed before and after the sequential testing of the steady-state operating points. In this regard, a reduction of the weight of the DPF in pre-turbo placement with respect to the traditional post-turbo configuration has been found.

Lastly, in the low load operating points F, J, and L (engine load below 10%), there is also a significant decrease in filtration efficiency in the range between 150 nm and 180 nm [40] in the case of the pre-turbo aftertreatment configuration. Since the most penetrating particle size seems to be located at a smaller size, this result may be related to measurement uncertainty in this operating region, since the filtration efficiency in operating points K and Q (engine load from 21% to 25%) is identical.

3.3. New European Driving Cycle. Gaseous emission.

Two types of tests have been performed for this cycle. On one hand, the first comprises the evaluation of the aftertreatment position in terms of pollutant limits dictated by the European standards [12]. In order to do that, a cold NEDC was tested for both exhaust line configurations and was compared concerning the approval of regulated gaseous pollutants. This kind of test allows assessing the position of the aftertreatment system, showing variables like the influence of warm-up engine strategy or DOC light-off. On the other hand, NEDC cycles were tested in order to provide a comprehensive baseline for unregulated pollutants and particles emission analysis when the aftertreatment placement is varied.

- Nitrogen oxides (NO , NO_2 and NO_x).

The overview of Figure 13 points out that the total accumulated NO_x is slightly higher in the pre-turbo aftertreatment configuration than in the post-turbo aftertreatment configuration. As Table 6 shows, both aftertreatment configurations are fulfilling EURO 4 regulation concerning NO_x emissions.

On one hand, DOC inlet NO_x emission is slightly decreased during UDC (Urban Driving Cycle) cycles when the pre-turbo aftertreatment configuration is evaluated (Figure 13.A). This behavior is due to the fact that the EGR valve is slightly open in order to keep the intake manifold pressure at the same level as in the case of the post-turbo aftertreatment configuration. This response reduces the fresh air mass and consequently decreases NO_x , specially in these stretches. On the other hand, when EUDC (Extra Urban Driving Cycle) emission was analyzed, an increase in total NO_x mass at the end of NEDC was observed.

From the point of view of NO concentration, the higher EGR rate during UDC has driven to NO reduction at the inlet of the DPF in pre-turbo aftertreatment configuration. (Figure 13.C). A drop by almost half in the NO concentration at the DOC outlet is observed in Figure 13.D in also in the case of the pre-turbo aftertreatment configuration. It is also interesting worth highlighting the fact that the NO to NO_2 conversion rate are greater in the case of pre-turbo placement of the aftertreatment (40% conversion in the pre-turbo configuration vs. 8% conversion in the post-turbo configuration), as is shown in Figure 13.F. As explained in Section 3.1, the DOC conversion rate is determined by the temperature of the exhaust gas. The higher the temperature is, the higher the conversion rate. Observing the DOC inlet temperature behavior, there was an increase in the pre-turbo aftertreatment configuration since the exhaust gas does not pass through the turbocharger before the DOC. Therefore, it is avoided the temperature drop that the flow undergoes in the turbine expansion.

- Ammonia (NH_3), sulfur dioxide (SO_2) and dinitrogen oxide (N_2O).

As stated in Section 3.1, the emission of these unregulated contaminants during steady-state operating points are negligible. Nevertheless, the emission of these compounds may occur during a transient operation so that its evaluation during NEDC is advised. Figure 14 shows results at the DOC inlet in the case of pre-turbo aftertreatment configuration. It confirms also for transient operation the low emission for these pollutants,

showing values under the analyzer error (2 ppm). In this sense, it is not possible to extract any conclusion about them, since they are emitted in a very low concentration.

- Carbon monoxide (CO) and Total hydrocarbons (THC).

During NEDC it can be observed that with a pre-turbo aftertreatment configuration there is a decrease in carbon monoxide and hydrocarbons emission at the DOC outlet. Regarding CO (Figure 15.A) and THC (Figure 15.B), both compounds show a similar trend, being different the oxidation rate between CO and THC . In the pre-turbo aftertreatment configuration, the DOC placement leads to reduce the required time to reach the DOC light-off (at 150°C). This allow a reduction of 11% in CO and a reduction of 2% in THC emitted when pre-turbo aftertreatment configuration is tested, as shown Table 6.

- Light hydrocarbons (<7 Carbon atoms)

Unregulated light hydrocarbons emission have some relevance in terms of quantity. The pollutants analyzed in Figure 16 are methane (CH_4), ethylene (C_2H_4), propene (C_3H_6), iso-butene ($iso - C_3H_8$) and benzene (C_6H_6). Other compounds are not taken into account in the study since the measurement is within the analyzer error (2 ppm).

Methane (CH_4), whose emission is shown in Figure 16.A, is a greenhouse gas which remains in the atmosphere for 9-15 years [41]. Methane may be appearing due to thermal cracking of paraffins and olefins. Thermal cracking occurs when a long-chain paraffinic bond is broken and consequently separated into two pairs of paraffins and olefins [42]. In the pre-turbo aftertreatment configuration, the DOC inlet temperature is higher than in the post-turbo aftertreatment configuration assisting thermal cracking and promoting CH_4 emission. Olefins like C_3H_6 , shown in Figure 16.B, have a strong tendency to split into like C_2H_4 , which explains the increase in ethylene emission, as it can be observed in Figure 16.C. The change in iso-butene ($iso - C_3H_8$) emission, which is represented in Figure 16.D, shows the same trend like that under steady-state operation, showing a constant emission level during whole NEDC.

Focusing on benzene (C_6H_6), which is a polycyclic aromatic hydrocarbon (PAH), is one of the most harmful compounds to human health due to its carcinogenic effect. Recent studies have shown the difficulty in oxidizing benzene in a DOC [43]. Taking into account such a difficulty, it is expected that most of the $TLHC$ emission, shown in Figure 17, are due to benzene contribution. It is confirmed by the benzene tailpipe emission, also shown in Figure 16.E. In fact, as raw benzene emission is the same whatever the aftertreatment placement, the lower catalytic surface in pre-turbo placement is damaging the benzene conversion efficiency. Considering the raw emission of lighter hydrocarbons and the tailpipe $TLHC$ trend, these must have higher conversion efficiency in pre-turbo placement; i.e. due to the fact that they are lighter the temperature increase in pre-turbo placement is dominating on dwell time.

3.4. New European Driving Cycle. Particle emission.

From a general point of view, both aftertreatment configurations fulfill the future EURO 6 particle number limit despite of the EURO 4 engine calibration. The particle emission does not exceed $2 \cdot 10^{11}$ #/km in any event, clearly lower than the EURO 6 limit set at $6 \cdot 10^{11}$ #/km.

As explained in Section 2.2, a very important factor when the particle measurement is carried out is the temperature at which the sampled exhaust gas is diluted with the primary dilution air [29]. Another factor of influence in particle size distribution measurement is the dilution ratio required to condition the sample at the spectrometer inlet. In this case, as shown in Figure 18, dilution ratios (DR) vary depending on both sampling point, upstream or downstream of the DPF, and on the aftertreatment configuration. This occurs because the pressure at the DPF inlet varies with the analyzed configuration. The dilution system employed works by pressure difference between the sample inlet and the pressure that reaches the diluted sample due to the suction taking place in the secondary dilution. As a consequence, greater sample pressure is reflected in a smaller dilution ratio. In the pre-turbo configuration, the gases passing through the DPF have a higher pressure than in the post-turbo configuration, since they have not undergone the expansion process occurring across the turbine. In any case, the dilution ratio was selected at the beginning of the cycle (at idle), being 170: 1 for the post-turbo aftertreatment configuration, and 100: 1 for the pre-turbo aftertreatment configuration at the DPF inlet.

When the measurement is carried out downstream of the DPF, it is not possible to establish high dilution ratios, since the intrinsic low particle concentration of the sample makes it hardly detectable by the spectrometer if the sample is diluted. Therefore, the dilution ratio applied was the maximum possible ensuring that the sample could be detected by the EEPS. The dilution ratio was set to 8: 1 for the post-turbo aftertreatment configuration and to 2: 1 in the case of pre-turbo DPF placement.

As seen previously, there is a slightly increase in hydrocarbons at the DPF inlet in the case of the pre-turbo aftertreatment configuration in comparison with the post-turbo aftertreatment configuration. This behavior is also seen in the nucleation particle mode ($D_p < 30$ nm), shown in Figure 19.A for DPF inlet, which are the particles whose formation is closely linked to hydrocarbons. However the accumulation particle mode ($D_p > 30$ nm) and the total particles remain at the same level in both configurations, as Figure 19.C and Figure 19.D show.

During the particle transport along the particle measurement line, it may occur that certain nucleation particles were generated due to nucleation of hydrocarbons. Figure 20 (A→B→D way), which shows the phase diagram in the dilution system, points out that the particles remain within the two steps of the dilution system in the pre-turbo aftertreatment configuration. A larger quantity of hydrocarbons present in the exhaust gas (Y axis in Figure 20), and a higher exhaust temperature (X axis in Figure 20) at DPF inlet favour thermochemical processes comprising the adsorption-nucleation phenomena of hydrocarbons in the secondary stage of dilution (ED at Figure 3). Due to this fact, some particles detected by the spectrometer are actually hydrocarbon formed by condensation. The combination of the increase in hydrocarbons and particle growing phenomena are the reason for the increase in the nucleation mode at the DPF inlet pre-turbo configuration.

However, when comparing pre-turbo versus post-turbo aftertreatment configurations at the DPF outlet, a decrease can be observed in nucleation particle mode and an increase in accumulation particles mode. The increase in accumulation mode downstream of the DPF is due to the thermal gradient between exhaust gas and ceramic wall temperature. In this sense, during the NEDC assessed with the pre-turbo aftertreatment configuration, the temperature inside the DPF outlet channels was cold so coagulation of certain nucleation particles mode present at DPF inlet was expected. As a consequence of coagulation of nucleation particles mode at DPF outlet, a decrease in nucleation particles mode measurement was observed.

Although an increase in passive regeneration process and an increase in nucleation mode at DPF outlet was found under steady-state conditions, there is no similar trend during NEDC. This is due to the fact that the nucleation rate during transient operation at the DPF outlet in pre-turbo configuration is much lower than in the post-turbo aftertreatment configuration because of the shift to accumulation mode. In this sense, although passive regeneration is favored and consequently an increase in nucleation mode would be expected, the overall balance between passive regeneration event and particles size growth due to high-hydrocarbons condensation leads to a reduction in the nucleation mode particles number emitted. One can say that the reduction in this particle mode is a clear advantage due to the fact that these particles are the most harmful to human health.

4. Summary and conclusions

The experiments performed in this work have been carried out in a EURO 4 passenger car engine considering post-turbo and pre-turbo aftertreatment placements. The original post-turbo aftertreatment configuration is composed of a close-coupled DOC and underfloor DOC and DPF. In the case of the pre-turbo aftertreatment configuration the underfloor DOC and DPF are placed upstream of the turbine removing the close-coupled DOC. Firstly, the pollutant emissions and the DPF filtration efficiency have been assessed in five steady-state operating points. After presenting the most relevant conclusions under steady-state conditions, the main results on engine performance during NEDC has been highlighted.

4.1. Steady-state operating points

- In terms of nitrogen oxides, it should be noted that placing the aftertreatment system in pre-turbo location has not shown effects in total NO_x emission. In contrast, when NO and NO_2 compounds are analyzed separately, an increase in NO_2 at the DOC outlet at low load operating points is obtained. The reason is the increased ratio of NO to NO_2 conversion, favored by the higher temperature at the inlet of the DOC. This is advantageous because it contributes, together with higher temperature upstream the DPF, to NO_2 passive regeneration leading to extend the time between active regenerations events. This is translated in fuel savings because of the decrease in both DPF average soot load and regeneration events frequency.
- An increased CO and THC oxidation ratio in the low load steady-state operating points has been found in the pre-turbo aftertreatment configuration.

- The light hydrocarbon emissions remained virtually unchanged irrespective of the aftertreatment system position, being the analyzed experimental values at the same level in both configurations.
- Concerning particles measurement; nucleation particle emission, smaller in size, is reduced at the DPF inlet in the case of the pre-turbo aftertreatment configuration due to the fact that these particles were never being formed upstream the DPF+VGT configuration. Contrarily, a slight increase of this mode emission appears at the VGT outlet due to the effect of the passive regeneration process.

4.2. Dynamic cycles. NEDC

- In the same way under steady-state operating points, there is much more NO_2 available for DPF passive regeneration in pre-turbo aftertreatment placement due to the higher DOC inlet temperature during the NEDC, leading to higher NO to NO_2 conversion rate.
- In terms of regulated pollutants, there is a decrease in both CO and THC emission during NEDC because the positive effect of the higher DOC inlet temperature leading to a faster DOC light-off.
- Concerning particle emission, both aftertreatment architectures fulfill next future EURO 6 legislation with an engine calibration for EURO 4, as the emission limit is 6.10^{11} #/km and in any event an emission of 2.10^{11} #/km is exceeded.
- If the particles are separated into nucleation particles ($D_p < 30$ nm) and accumulation particles ($D_p > 30$ nm), there is a decrease of the former. The reason lies in the coagulation of nucleation particles across the DPF due to fact that thermal inertia has at the DPF outlet channels, cooling down the exhaust gases. Hence, the primary particles undergoes a diameter growth becoming part of the accumulation particles mode. This is a clear advantage because recent studies show that smaller particles are the most harmful to human health [44, 45] even they are not regulated.

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Nomenclature

Subscripts

<i>inlet</i>	inlet channel
<i>outlet</i>	outlet channel
<i>ac</i>	accumulated mass

Abbreviations

AF	Air to fuel ratio
CRT	Continuous regeneration trap
DOC	Diesel oxidation catalyst
DR	Dilution ratio
DPF	Diesel particulate filter
ECU	Electronic control unit
EEPS	Engine exhaust particle sizer
EGR	Exhaust gas recirculated
EUDC	Extra urban driving cycle
FPS	Fine particle sampler
FTIR	Fourier transform infrared spectroscopy
GMD	Geometric mean diameter
HSDI	High speed direct injection
NEDC	New european driving cycle
SCR	Selective catalytic reduction
SOI	Start of injection
UDC	Urban driving cycle
USLD	Ultra low sulfur Diesel

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Table 1: Engine's main characteristics.

Type	4-cycle
Displacement	1997 [cm^3]
Diameter	85 [mm]
Stroke	88 [mm]
Number of cylinders	4 [-]
Valves by cilinder	4 [-]
Compression ratio	18:1 [-]
Maximun power	100 [kW] at 4000 rpm
Maximun torque	320 [Nm] at 1750 rpm
Spray cone angle	140 [$^\circ$]
Nozzle holes	6 x 146 [μm]

Table 2: Aftertreatment characteristics.

DOC		DPF	
Length	0.105 [m]	Length	0.165 [m]
Diameter	0.145 [m]	Diameter	0.135 [m]
Volume	1.73 [L]	Volume	2.36 [L]
Cell density	400 [cpsi]	Cell density	200 [cpsi]
Cell size	0.94 [mm]	Cell size	1.47 [mm]
Wall thickness	0.33 [mm]	Wall thickness	0.32 [mm]
Number of channels	10240 [-]	Number of channels	4470 [-]
Catalytic surface	4.04 [m^2]	Filtration surface	2.17 [m^2]
		Permeability	3.175×10^{-13} [mm]

Table 3: Fuel properties.

Cetane number	51.6 [-]
Viscosity at 40°C	2.46 [mm^2/s]
Density at 15°C	0.843 [kg/l]
Lower heating value	42.055 [MJ/kg]
Sulfur content	6.6 [ppm]
Water content	96 [ppm]

Table 4: Technical data of piezoresistive pressure sensors.

Range	0 - 5 [$bar(abs)$]
Overload	12.5 [$bar(abs)$]
Sensitivity	25 [$mV/bar/mA$]
Linearity	≤ 0.1 [% FSO]
Temperature range	0 - 140 [$^{\circ}C$]

Table 5: Steady-state operating points. Main parameters.

Point	Speed	Torque	Load	SOI	EGR	Particle mode
	[rpm]	[N.m]	[%]	[$^{\circ}$]	[%]	[-]
F	1370	15	7	-1.62	48.5	Bimodal
J	1960	16	8	-0.40	36	Bimodal
K	1660	104	21	-3.54	12.5	Accumulation
L	1580	42	10	-2.98	36.5	Accumulation
Q	2240	89	25	-1.2	12.85	Accumulation

Table 6: Regulated pollutants emission.

Pollutant	EURO 4 limit	EURO 5 limit	Post-turbo	Pre-turbo
	[mg/km]	[mg/km]	[mg/km]	[mg/km]
NOx	250	180	188.1	197.2
THC+NOx	300	230	234.5	239.1
CO	500	500	460.1	409.1

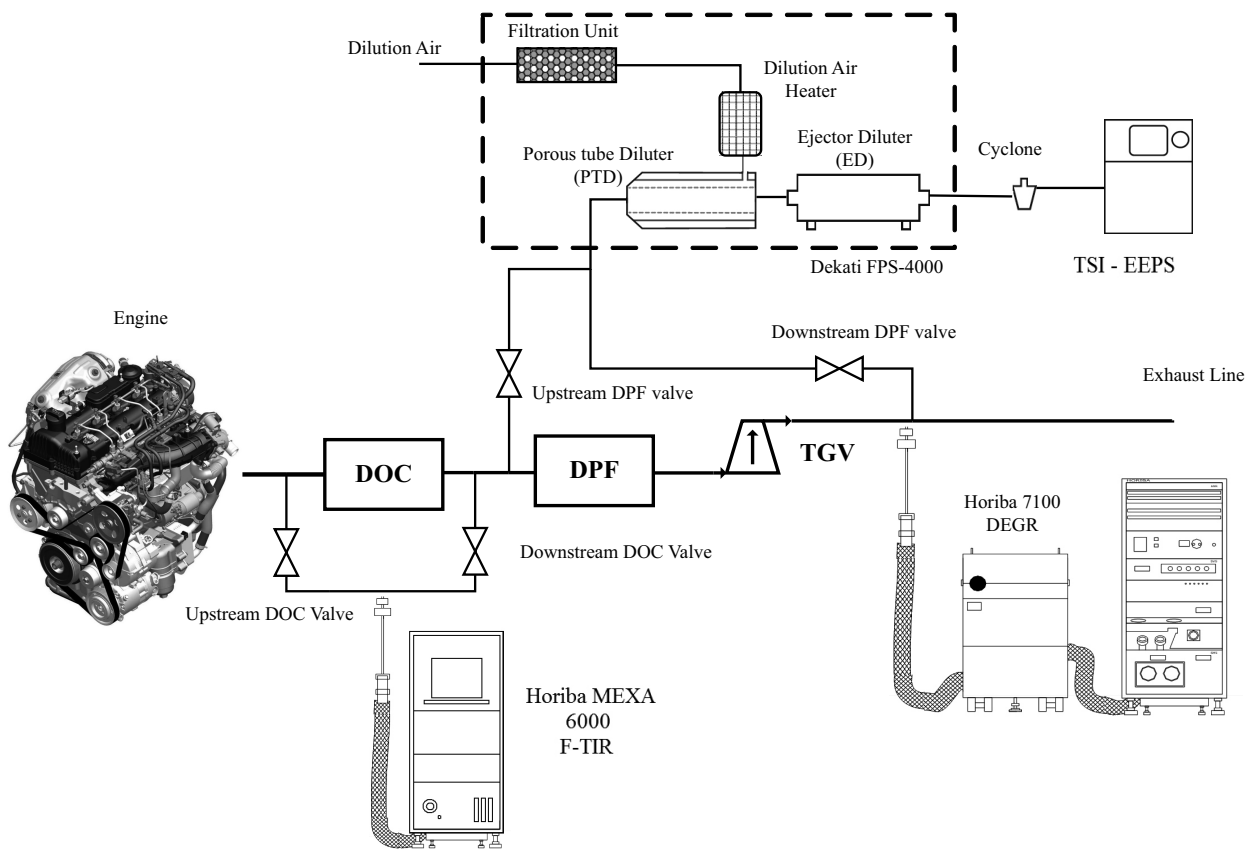


Figure 1: Experimental setup for the evaluation of the aftertreatment response.

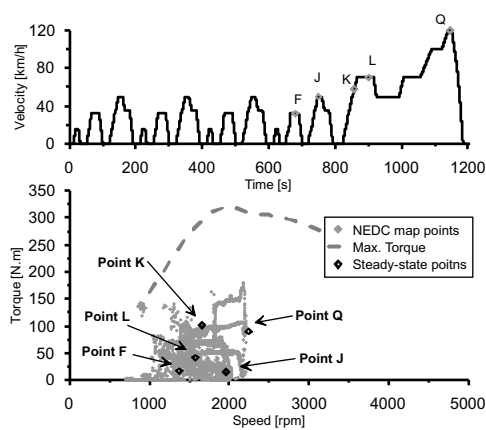


Figure 2: Representation in the NEDC map of the operating points selected for the evaluation of the aftertreatment response.

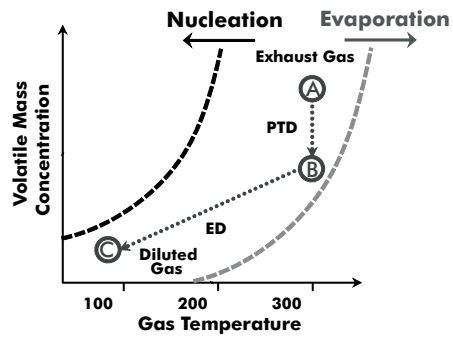


Figure 3: Particle evolution at dilution system. Theoretical phase-diagram used in the methodology for measuring particle distribution size [29].

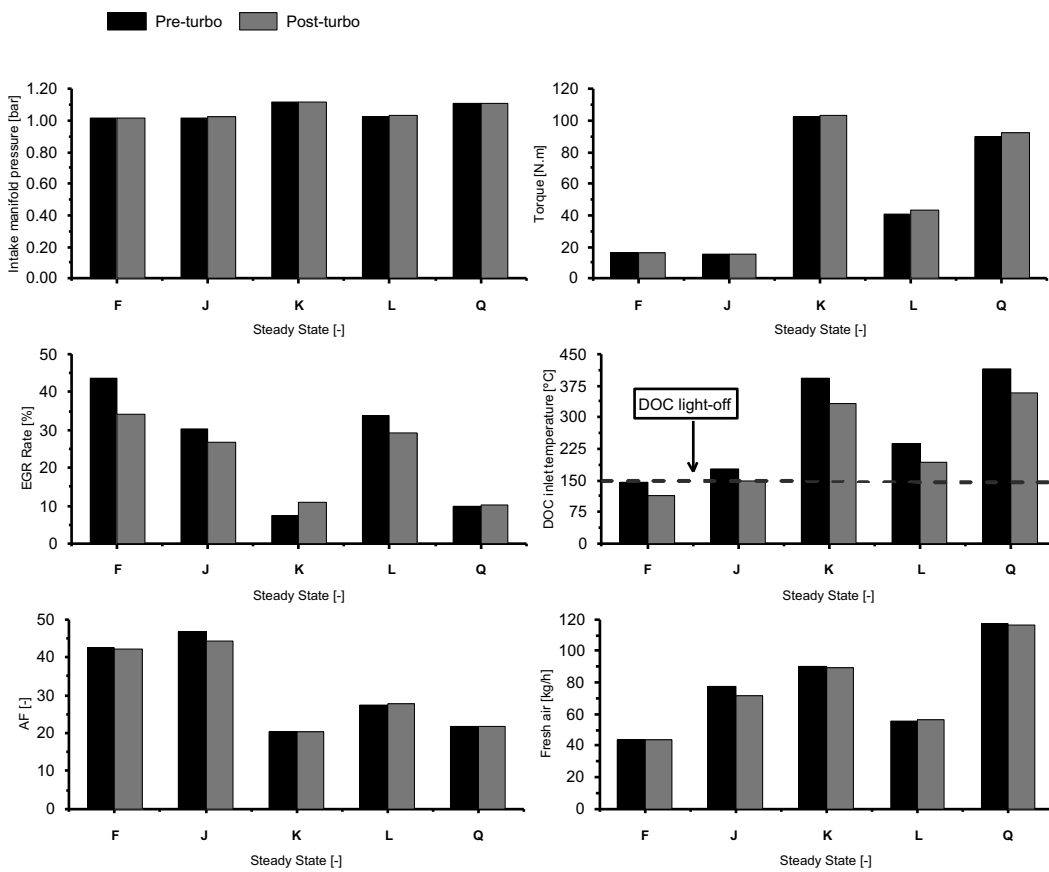


Figure 4: Steady-state operating points. Control variables analyzed.

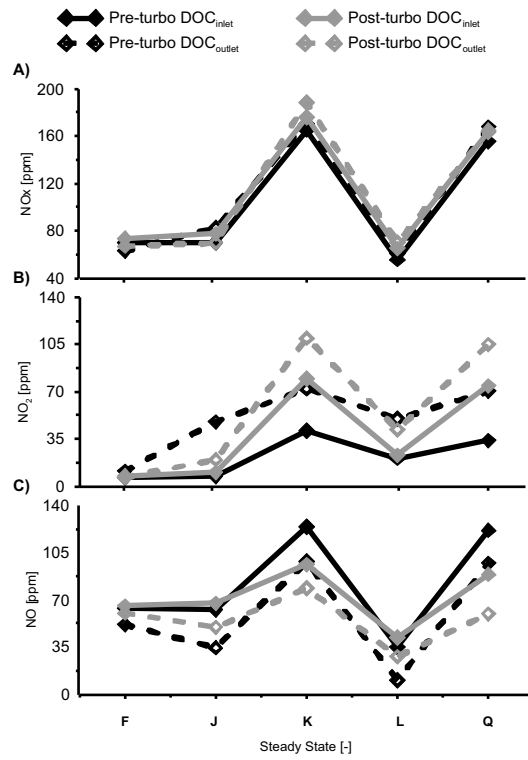


Figure 5: Nitrogen oxides emission at steady-state operating conditions. A) Total nitrogen oxides (NO_x). B) Nitrogen dioxide (NO_2). C) Nitrogen monoxide (NO). Pre-turbo vs. Post-turbo aftertreatment configuration.

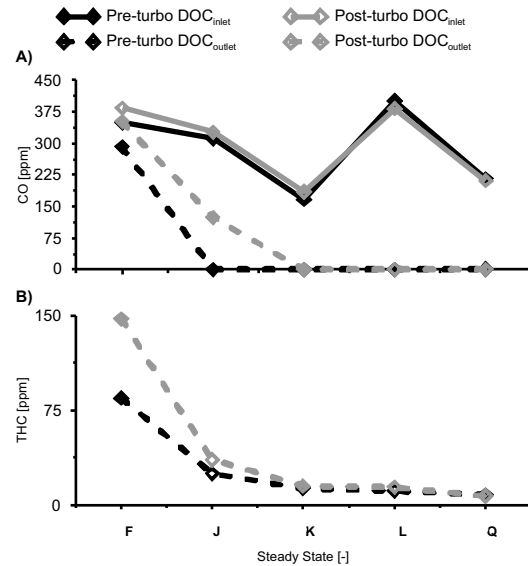


Figure 6: Carbon monoxide and total hydrocarbons emission at steady-state operating conditions. Pre-turbo vs. Post-turbo aftertreatment configuration.

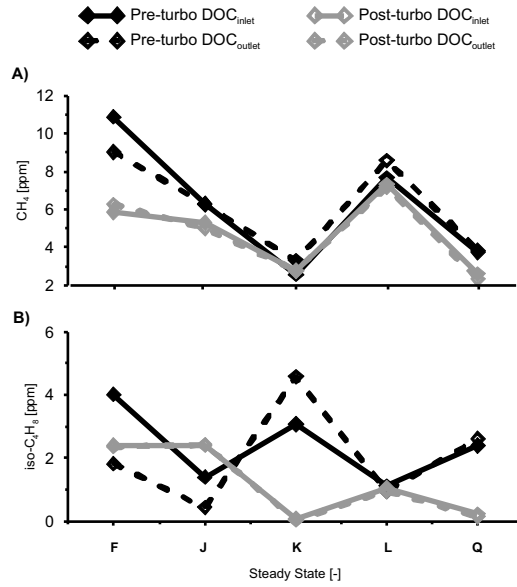


Figure 7: Light hydrocarbons (<7 Carbon atoms) at steady-state operating conditions. Methene (CH_4) and iso-butene ($iso - C_4H_8$). Pre-turbo vs. Post-turbo aftertreatment configuration.

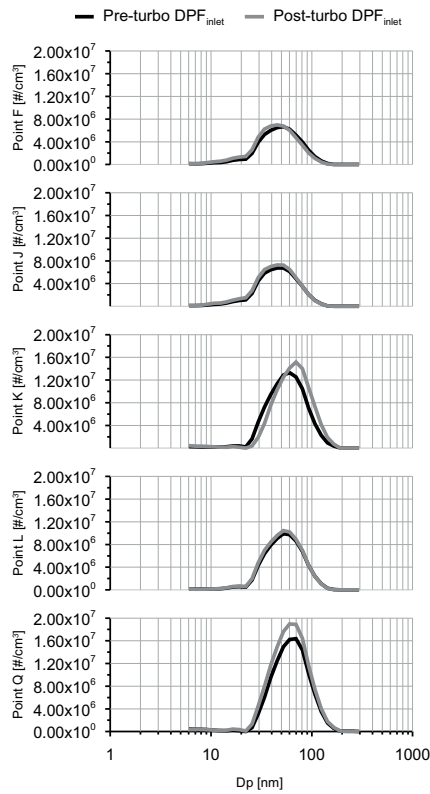


Figure 8: Particle distribution size upstream the DPF at steady-state operating points. Pre-turbo vs. Post-turbo aftertreatment configuration.

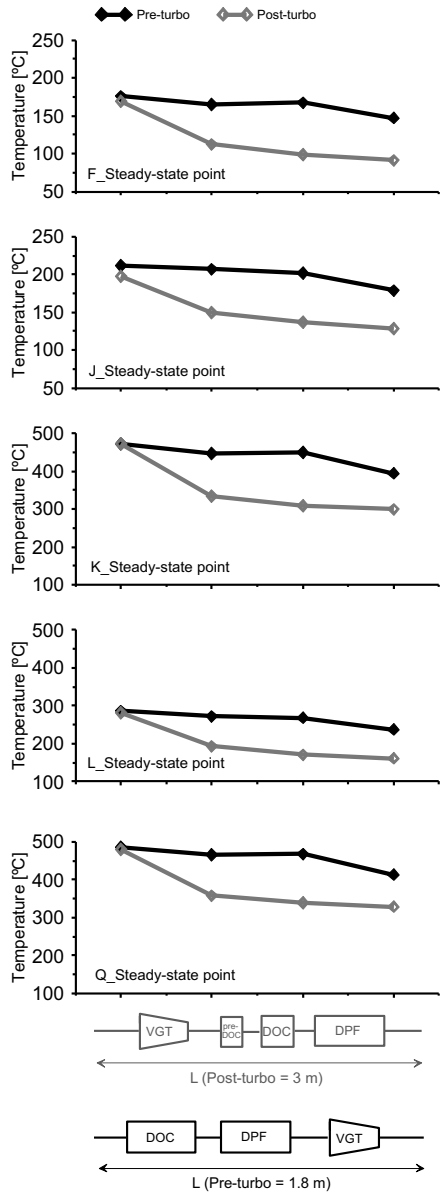


Figure 9: Temperature evolution along the exhaust architecture at steady-state operating points. Pre-turbo vs. Post-turbo aftertreatment configuration

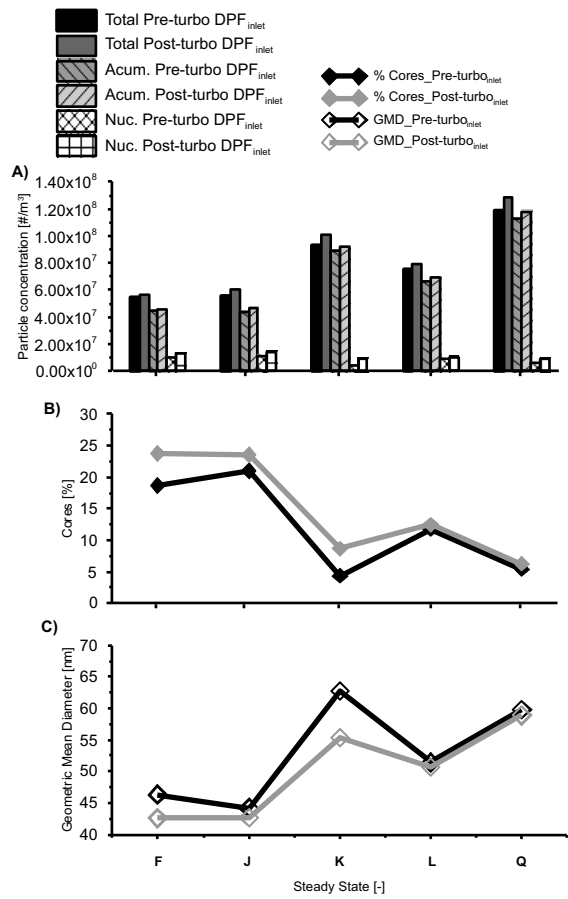


Figure 10: DPF inlet particle emission analysis at steady-state operation conditions. A) Particle concentration. B) Percentage of cores. C) Geometric mean diameter (GMD). Pre-turbo vs. Post-turbo aftertreatment configuration.

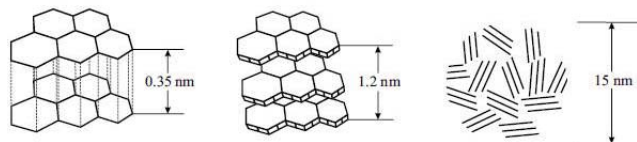


Figure 11: Nuclei-mode formation.

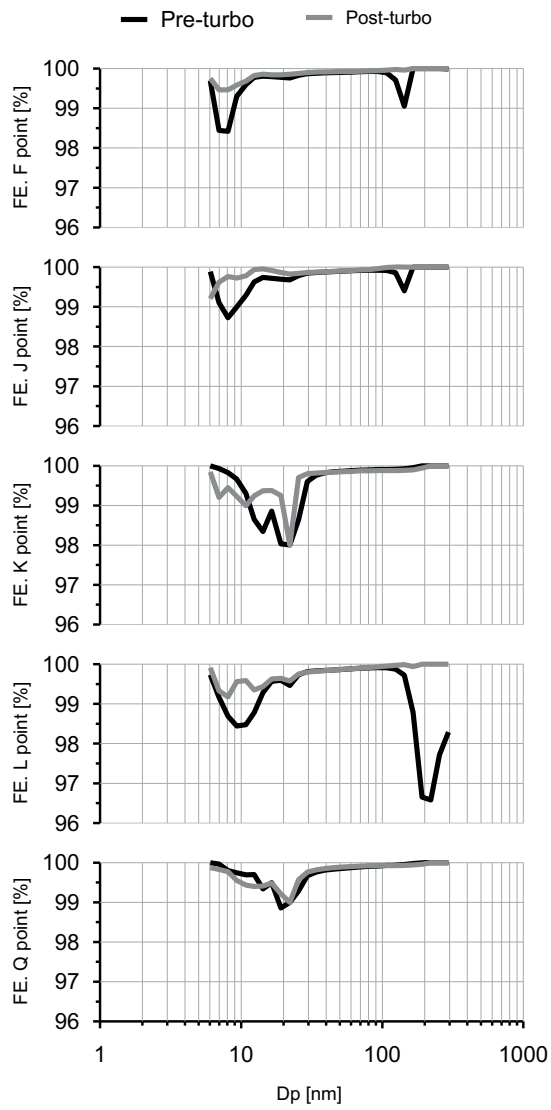


Figure 12: DPF filtration efficiency (FE) at steady-state operating conditions. Pre-turbo vs. Post-turbo aftertreatment configuration.

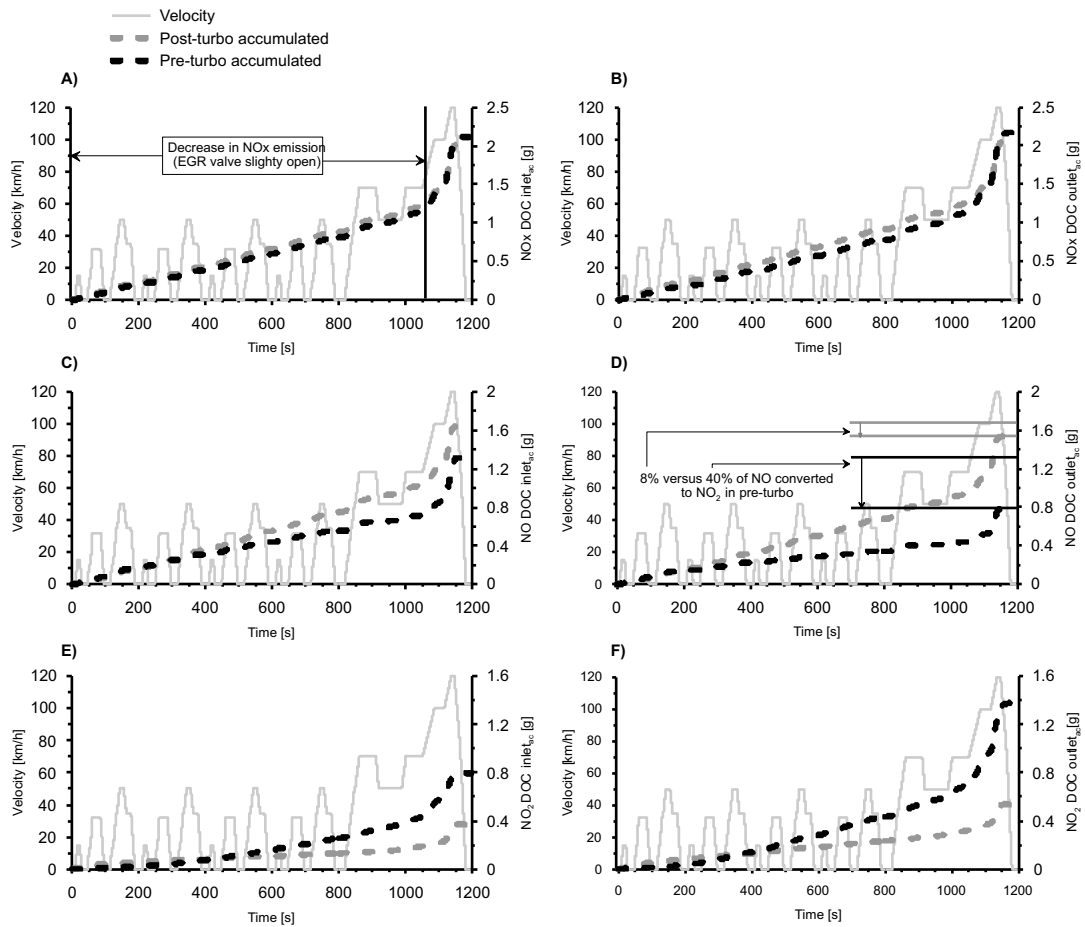


Figure 13: Nitrogen oxides evaluation during NEDC. A) NO_x upstream DOC. B) NO_x downstream DOC. C) NO upstream DOC. D) NO downstream DOC. E) NO₂ upstream DOC. F) NO₂ downstream DOC. Pre-turbo vs. Post-turbo aftertreatment configuration.

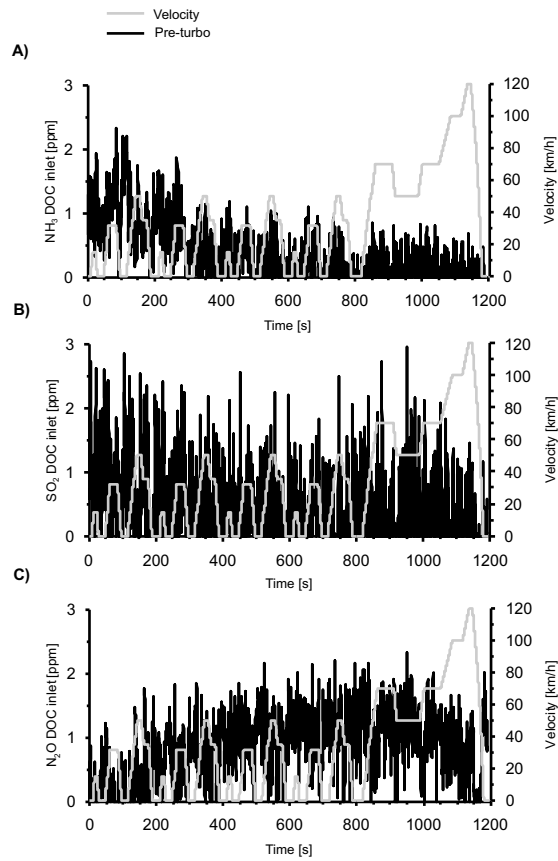


Figure 14: Ammonia (NH_3), sulfur dioxide (SO_2) and dinitrogen oxide (N_2O) emission during NEDC assessed in a pre-turbo aftertreatment configuration. A) NH_3 emission. B) SO_2 emission. C) N_2O emission

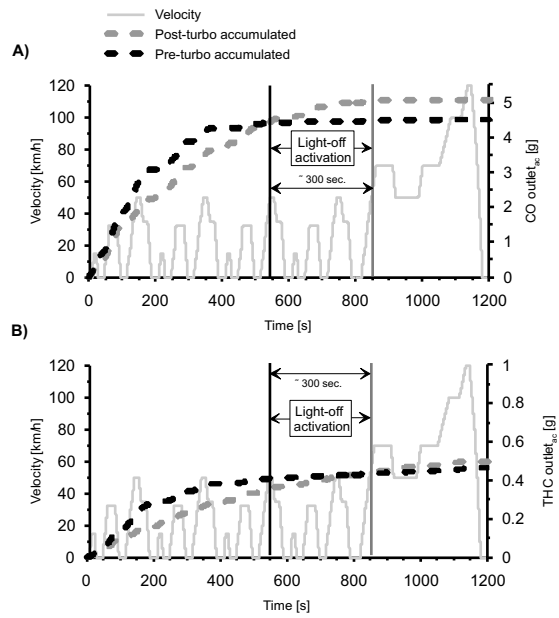


Figure 15: Carbon monoxide and total hydrocarbons evaluation during a NEDC. A) CO emission. B) THC emission. Pre-turbo vs. Post-turbo aftertreatment configuration.

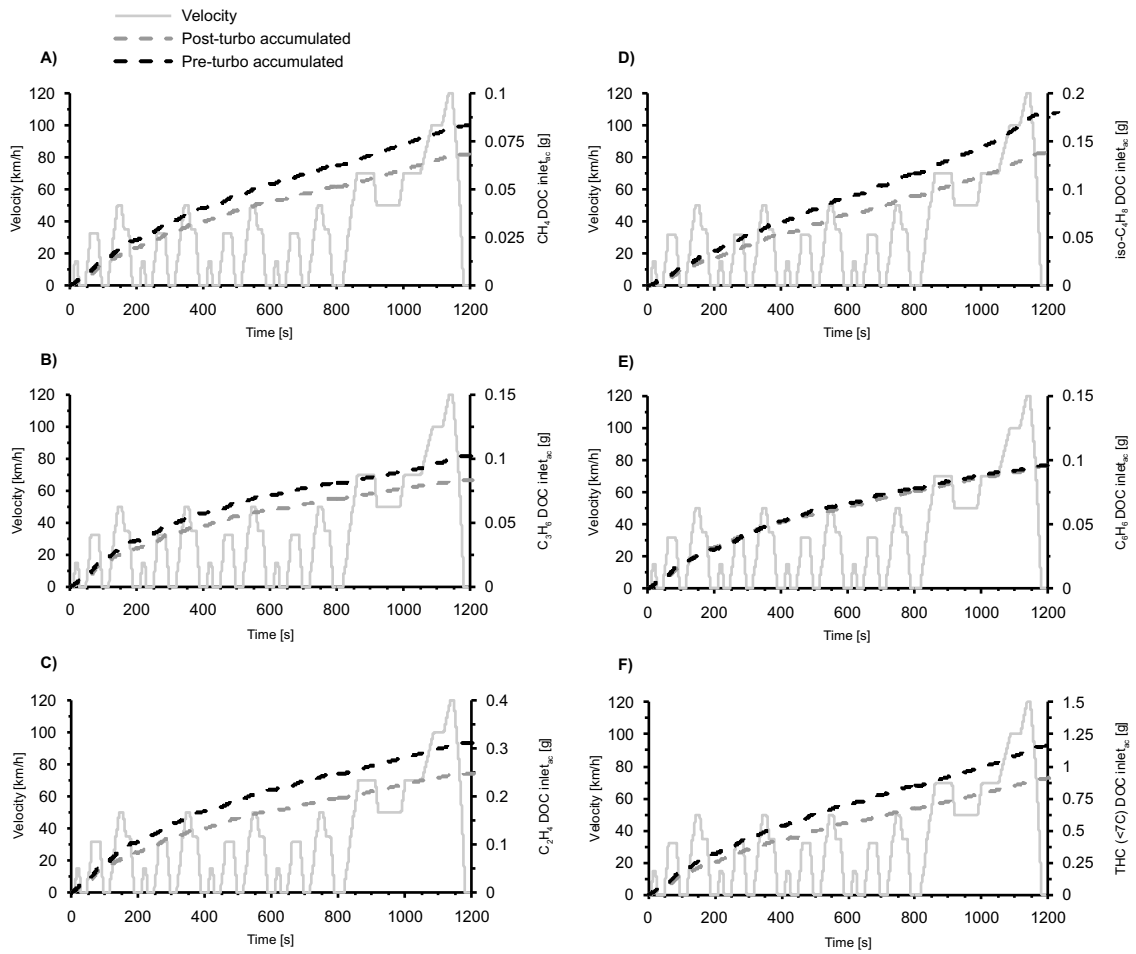


Figure 16: Light HC (<7 Carbon atoms) evaluation at DOC inlet during NEDC. A) Methane (CH_4). B) Ethylene (C_2H_4). C) Propene (C_3H_6). D) Iso-butene (*iso* – C_3H_8). E) Benzene (C_6H_6). F) Total light HC. Pre-turbo vs. Post-turbo aftertreatment configuration.

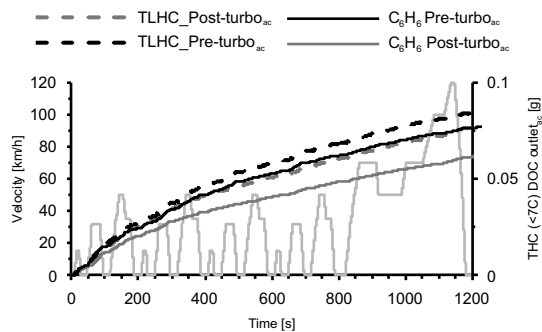


Figure 17: Light HC (<7 Carbon atoms) evaluation at DOC outlet during NEDC. Pre-turbo vs. Post-turbo aftertreatment configuration.

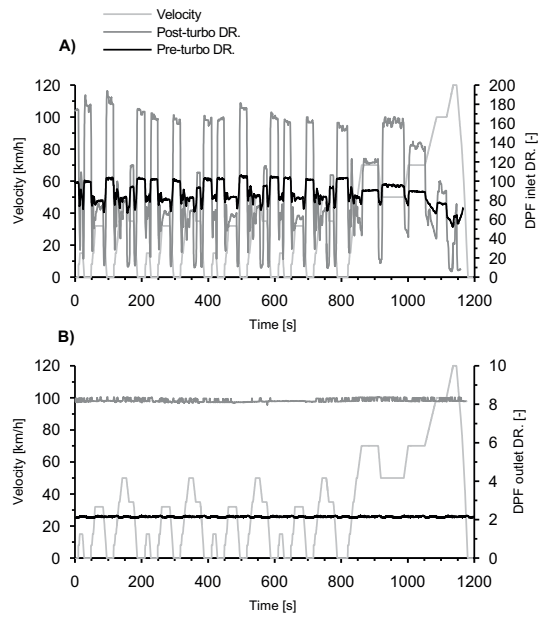


Figure 18: Dilution ratio during NEDC. A) Dilution ratio upstream the DPF. B) Dilution ratio downstream the DPF. Pre-turbo vs. Post-turbo aftertreatment configuration.

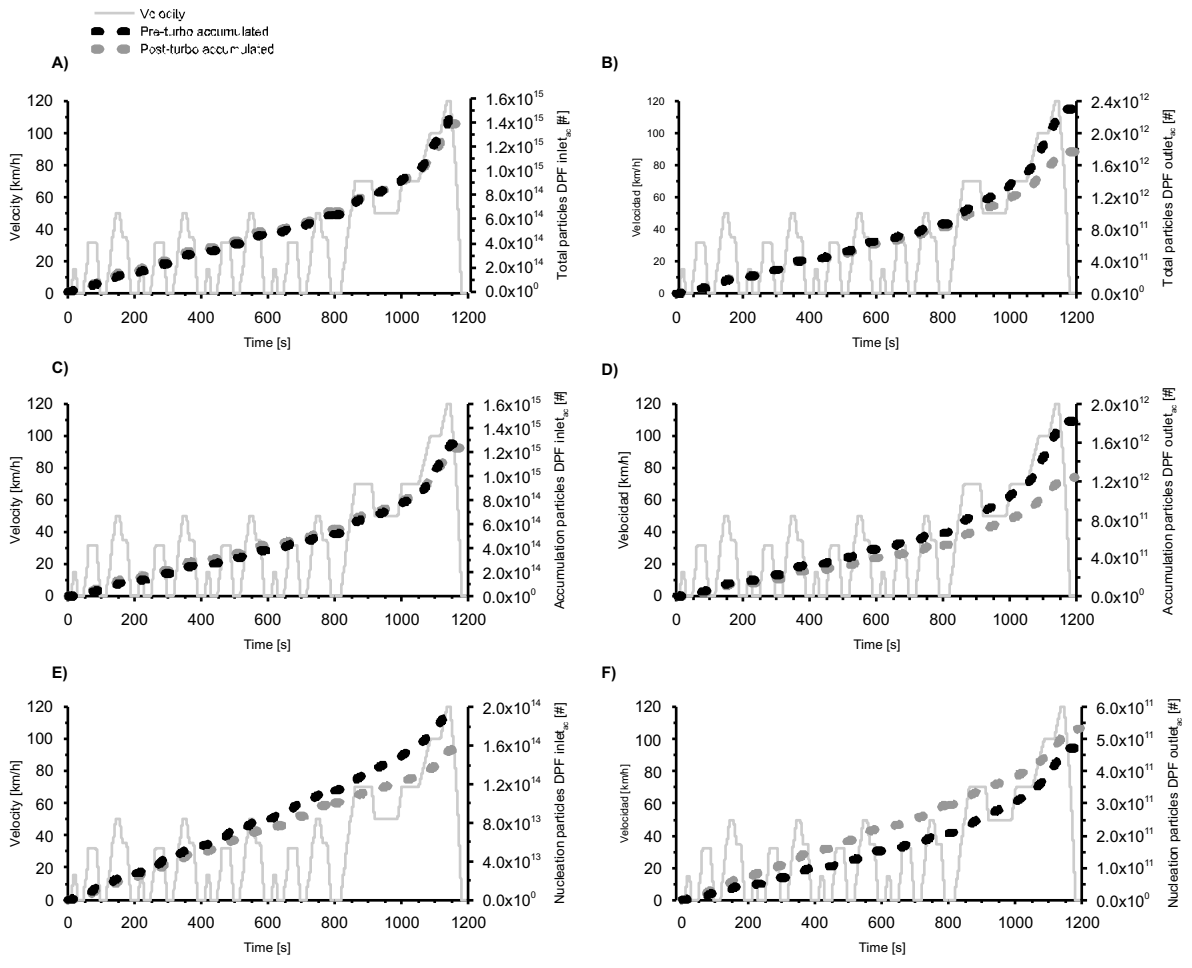


Figure 19: Particle emission evaluation during NEDC. A) Total particles at DPF inlet. B) Total particles at DPF outlet. C) Accumulation particles at DPF inlet. D) Accumulation particles at DPF outlet. E) Nucleation particles at DPF inlet. F) Nucleation particles at DPF outlet. Pre-turbo vs. Post-turbo aftertreatment configuration.

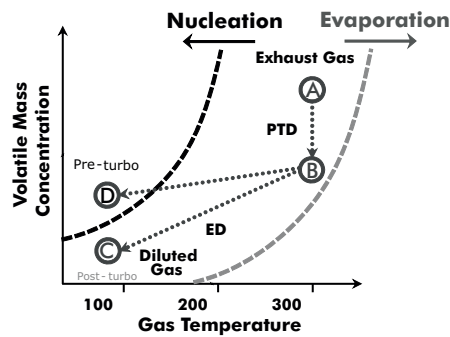


Figure 20: Particle evolution at the dilution system. Pre-turbo vs. Post-turbo aftertreatment configuration.