



(51) International Patent Classification:

B01J 23/34 (2006.01) *B01J 35/04* (2006.01)
B01J 23/38 (2006.01) *B01D 53/56* (2006.01)
B01J 23/58 (2006.01) *B01D 53/62* (2006.01)
B01J 23/63 (2006.01) *B01D 53/72* (2006.01)
B01J 35/00 (2006.01) *B01D 53/92* (2006.01)
B01J 37/02 (2006.01) *B01D 53/94* (2006.01)
B01J 23/42 (2006.01) *F01N 3/022* (2006.01)
B01J 23/44 (2006.01)

(21) International Application Number:

PCT/GB2015/050145

(22) International Filing Date:

23 January 2015 (23.01.2015)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

1401115.9 23 January 2014 (23.01.2014) GB
1405871.3 1 April 2014 (01.04.2014) GB

(71) Applicant: **JOHNSON MATTHEY PUBLIC LIMITED COMPANY** [GB/GB]; 5th floor, 25 Farringdon Street, London EC4A 4AB (GB).

(72) Inventors: **CHIFFEY, Andrew Francis**; 8 Spring View Road, Ware Hertfordshire SG12 9LB (GB). **GOODWIN, John**; 33 Braeburn Walk, Royston SG8 5LY (GB). **LEELAND, James**; c/o Johnson Matthey PLC, Gate 20, Orchard Road, Royston Hertfordshire SG8 5HE (GB). **MOR-EAU, Francois**; 29 Madras Road, Cambridge CB1 3PX (GB). **REID, Stuart, David**; 19 Mayfield Way, Great Cambourne, Cambourne Cambridgeshire CB23 5JA (GB).

SWALLOW, Daniel; 37 Primrose Hill, Little Gransden, Sandy Bedfordshire SG19 3DP (GB).

(74) Agent: **TURBERVILLE, Simon**; Johnson Matthey PLC, Intellectual Property Department, Gate 20, Orchard Road, Gate 20, Orchard Road, Royston Hertfordshire SG8 5HE (GB).

(81) Designated States (*unless otherwise indicated, for every kind of national protection available*): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

— with international search report (Art. 21(3))

(54) Title: DIESEL OXIDATION CATALYST AND EXHAUST SYSTEM

(57) Abstract: An oxidation catalyst for treating an exhaust gas from a diesel engine, which oxidation catalyst comprises: a first washcoat region comprising a first platinum group metal (PGM), a first support material and a NO_x storage component; a second washcoat region comprising platinum (Pt), manganese (Mn) and a second support material; and a substrate having an inlet end and an outlet end.



WO 2015/110817 A1

DIESEL OXIDATION CATALYST AND EXHAUST SYSTEM

FIELD OF THE INVENTION

The invention relates to an oxidation catalyst for a diesel engine and to an exhaust
5 system for a diesel engine comprising the oxidation catalyst. The invention also relates
to methods and uses of the oxidation catalyst for treating an exhaust gas from a diesel
engine.

BACKGROUND TO THE INVENTION

10 Diesel engines produce an exhaust emission that generally contains at least four classes
of pollutant that are legislated against by inter-governmental organisations throughout
the world: carbon monoxide (CO), unburned hydrocarbons (HCs), oxides of nitrogen
(NO_x) and particulate matter (PM). Emissions standards for diesel engines, whether
stationary or mobile (e.g. vehicular diesel engines), are being progressively tightened.
15 There is a need to provide improved catalysts and exhaust systems that are able to meet
these standards, which are cost-effective.

Exhaust systems for diesel engines may include several emissions control devices.
Each emissions control device has a specialised function and is responsible for treating
20 one or more classes of pollutant in the exhaust gas. The performance of an upstream
emissions control device can affect the performance of a downstream emissions control
device. This is because the exhaust gas from the outlet of the upstream emissions
control device is passed into the inlet of the downstream emissions control device. The
interaction between each emissions control device in the exhaust system is important to
25 the overall efficiency of the system.

Oxidation catalysts, such as diesel oxidation catalysts (DOCs), typically oxidise carbon
monoxide (CO) and hydrocarbons (HCs) in an exhaust gas produced by a diesel engine.
Diesel oxidation catalysts can also oxidise some of the nitric oxide (NO) that is present in
30 the exhaust gas to nitrogen dioxide (NO₂). Even though nitrogen dioxide (NO₂) is itself a
pollutant, the conversion of NO into NO₂ can be beneficial. The NO₂ that is produced
can be used to regenerate particulate matter (PM) that has been trapped by, for
example, a downstream diesel particulate filter (DPF) or a downstream catalysed soot
filter (CSF). Generally, the NO₂ generated by the oxidation catalyst increases the ratio of
35 NO₂:NO in the exhaust gas from the outlet of the oxidation catalyst compared to the
exhaust gas at the inlet. This increased ratio can be advantageous for exhaust systems
comprising a downstream selective catalytic reduction (SCR) catalyst or a selective

catalytic reduction filter (SCRTM) catalyst. The ratio of NO₂:NO in the exhaust gas produced directly by a diesel engine may be too low for optimum SCR or SCRTM catalyst performance.

- 5 Whilst it is generally advantageous to include an oxidation catalyst that has good NO₂ generation activity in an exhaust system, the use of an oxidation catalyst in this way can be problematic when seeking to obtain optimum performance from a downstream emissions control device (e.g. an SCR or SCRTM catalyst). The average amount of NO₂ that is generated by an oxidation catalyst at a given exhaust gas temperature can vary
- 10 considerably over its lifetime. This can lead to difficulties in calibrating the dosing of the nitrogenous reductant for performing active SCR.

SUMMARY OF THE INVENTION

The inventors have found that excellent activity toward the oxidation of nitric oxide (NO)

15 in an exhaust gas from a diesel engine can be obtained when manganese (Mn) is combined with platinum (Pt) in an oxidation catalyst. Platinum is expensive and is often included in oxidation catalysts in relatively large amounts for, amongst other things, its oxidation activity toward NO. The inclusion of manganese (Mn) in combination with platinum (Pt) may result in an improvement in NO oxidation activity or allow the use of a

20 reduced amount of Pt to achieve a given level of NO oxidation. The oxidation catalyst, particularly the Mn and Pt containing region, may have good activity toward oxidising carbon monoxide (CO) and hydrocarbons (HCs) in an exhaust gas produced by a diesel engine.

25 The invention provides an oxidation catalyst for treating an exhaust gas from a diesel engine, which oxidation catalyst comprises: a first washcoat region comprising a first platinum group metal (PGM) and a first support material; a second washcoat region comprising platinum (Pt), manganese (Mn) and a second support material; and a substrate having an inlet end and an outlet end.

30 The first washcoat region is suitable for NO_x storage (i.e. NO_x adsorption). It may also be suitable for oxidising carbon monoxide (CO) and hydrocarbons (HCs). The second washcoat region is suitable for oxidising nitric oxide (NO), but it may also be suitable for oxidising carbon monoxide (CO) and/or hydrocarbons (HCs), particularly when the

35 second washcoat region is substantially free of palladium.

The oxidative activity of the combination of Pt and Mn toward nitric oxide (NO) and also CO and/or HCs can be particularly advantageous when certain alumina based materials are used as a support material.

- 5 The combination of a washcoat region comprising Pt and Mn together with a washcoat region formulated for NO_x storage (i.e. NO_x adsorption) may be advantageous. The Pt and Mn containing washcoat region may convert NO_x to NO₂ to facilitate NO_x storage and/or may assist in the conversion of any NO to NO₂ that passes through the NO_x storage region.

10

The invention further provides an exhaust system for a diesel engine. The exhaust system comprises an oxidation catalyst of the invention and an emissions control device.

15

The amount of NO₂ in an exhaust gas can affect the performance of a downstream emissions control device for the selective catalytic reduction of oxides of nitrogen (NO_x). Selective catalytic reduction (SCR) catalysts and selective catalytic reduction filter (SCRFTM) catalysts for treating NO_x (e.g. NO₂ + NO) often require the ratio of NO₂ to NO in the inlet gas to be within a specific range for optimum catalytic performance. The optimal NO₂ proportion of NO_x typically depends on the type of composition used in the SCR or SCRFTM catalyst, but the ratio of NO₂ to NO in an exhaust gas directly from a diesel engine is usually too low for optimal performance of the catalyst.

20

25

The oxidation catalyst of the invention can be used to convert NO into NO₂ and thereby increase the amount of NO₂ in the exhaust gas that exits the catalyst in comparison to the amount of NO₂ in the exhaust gas at the inlet of the catalyst (i.e. the ratios of NO₂:NO_x and NO₂:NO in the exhaust gas that exits the oxidation catalyst are greater than the corresponding ratios of the exhaust gas at the inlet of the oxidation catalyst). The oxidation catalyst can alter the NO_x composition of the exhaust gas for optimum SCR performance.

30

35

A problem with using an oxidation catalyst to “boost” the NO₂ content of an exhaust gas is that the NO oxidation activity of the oxidation catalyst usually varies over its lifetime. Generally, as the catalyst “ages” (i.e. the catalyst has been used over a prolonged period), the NO oxidation activity of the catalyst decreases. Whilst the amount of NO₂ in the exhaust gas that exits an “aged” oxidation catalyst may be sufficient for optimal performance of a downstream emissions control device (e.g. an SCR catalyst), this

variation in the amount of NO₂ that is generated is problematic for calibrating the dosing of the nitrogenous reductant for performing active SCR.

5 It has been found that the oxidation catalyst of the invention can show relatively stable NO oxidation activity over its lifetime. Thus, the difference in NO oxidation activity of the oxidation catalyst in a fresh state (i.e. when it is “new” and has not been subjected to repeated, prolonged use) and an aged state is typically small.

10 A further aspect of the invention relates to a vehicle or an apparatus (e.g. a stationary or mobile apparatus). The vehicle or apparatus comprises a diesel engine and either the oxidation catalyst or the exhaust system of the invention.

The invention also relates to several uses and methods.

15 A first method aspect of the invention provides a method of treating an exhaust gas from a diesel engine. The method comprises either contacting the exhaust gas with an oxidation catalyst of the invention or passing the exhaust gas through an exhaust system of the invention. The expression “treating an exhaust gas” in this context refers to oxidising carbon monoxide (CO), hydrocarbons (HCs) and nitric oxide (NO) in an
20 exhaust gas from a diesel engine.

A second method aspect provides a method of modulating the content of NO_x in an exhaust gas from a diesel engine for an emissions control device. The method comprises: (a) controlling the NO_x content of an exhaust gas by contacting the exhaust
25 gas with an oxidation catalyst of the invention to produce a treated exhaust gas; and (b) passing the treated exhaust gas to an emissions control device.

A first use aspect of the invention relates to the use of an oxidation catalyst to treat an exhaust gas from a diesel engine, optionally in combination with an emissions control
30 device. Generally, the oxidation catalyst is used to treat (e.g. oxidise) carbon monoxide (CO) and hydrocarbons (HCs) in an exhaust gas from a diesel engine.

In a second use aspect, the invention relates to the use of an oxidation catalyst to modulate the content of NO_x in an exhaust gas from a diesel engine for an emissions
35 control device (e.g. a downstream emissions control device).

A third use aspect relates to the use of an oxidation catalyst in the regeneration of an emissions control device having a filtering substrate (e.g. a downstream emissions control device having a filtering substrate).

- 5 A fourth use aspect relates to the use of manganese (Mn), preferably in combination with platinum (Pt), in an oxidation catalyst for a diesel engine to enhance oxidation of nitric oxide (NO) in an exhaust gas from the diesel engine.

- A fifth use aspect relates to the use of manganese (Mn), preferably in combination with platinum (Pt), in an oxidation catalyst for a diesel engine to stabilise the NO oxidation activity of the oxidation catalyst over its lifetime.
- 10

In the first to fifth use aspects, the oxidation catalyst is an oxidation catalyst in accordance with the invention.

15

BRIEF DESCRIPTION OF THE DRAWINGS

Figures 1 to 10 are schematic representations of oxidation catalysts of the invention.

The exhaust gas may enter from either the left or the right side of the oxidation catalyst.

- Figure 1 shows an oxidation catalyst comprising a first washcoat region (1) and a second washcoat region/zone (2) disposed on a substrate (3).
- 20

Figure 2 shows an oxidation catalyst comprising a first washcoat region (1) and a second washcoat region/zone (2). The first washcoat region (1) is disposed directly on to the substrate (3). The second washcoat region/zone (2) is disposed on the first washcoat region (1).

- Figure 3 shows an oxidation catalyst comprising a first washcoat region (1) and a second washcoat region/zone (2). There is an overlap between the first washcoat region (1) and the second washcoat region/zone (2). A part of the first washcoat region (1) is disposed on the second washcoat region/zone (2). Both the first washcoat region (1) and the second washcoat region/zone (2) are disposed on the substrate (3).
- 25

- Figure 4 shows an oxidation catalyst comprising a first washcoat region (1) and a second washcoat region/zone (2). There is an overlap between the first washcoat region (1) and the second washcoat region/zone (2). A part of the second washcoat region/zone (2) is disposed on the first washcoat region (1). Both the first washcoat region (1) and the second washcoat region/zone (2) are disposed on the substrate (3).
- 30

Figure 5 shows an oxidation catalyst comprising a first washcoat layer (1) and a second washcoat layer (2) disposed on a substrate (3). The second washcoat layer (2) is disposed on the first washcoat layer (1).

5 Figure 6 shows an oxidation catalyst comprising a first washcoat zone (1) and a second washcoat region/zone (2). Both the first washcoat region/zone (1) and the second washcoat region/zone (2) are disposed on a third washcoat region/layer (4). The third washcoat region/layer (4) is disposed on a substrate (3).

10 Figure 7 shows an oxidation catalyst comprising a first washcoat zone (1), a second washcoat zone (2) and a third washcoat zone (4). Both the first washcoat zone (1) and the second washcoat zone (2) are disposed on a substrate (3). The third washcoat zone (4) is disposed on the second washcoat zone (2).

15 Figure 8 shows an oxidation catalyst comprising a first washcoat zone (1), a second washcoat zone (2) and a third washcoat zone (4). Both the first washcoat zone (1) and the third washcoat zone (4) are disposed on a substrate (3). The second washcoat zone (2) is disposed on the third washcoat zone (2).

Figure 9 shows an oxidation catalyst comprising a first washcoat zone (1), a second washcoat zone (2) and a third washcoat zone (4). Both the first washcoat zone (1) and the second washcoat zone (2) are disposed on a substrate (3). The third washcoat zone (4) is disposed on the first washcoat zone (1).

20 Figure 10 shows an oxidation catalyst comprising a first washcoat zone (1), a second washcoat zone (2), a third washcoat zone (4) and a fourth washcoat zone (5). Both the first washcoat zone (1) and the third washcoat zone (4) are disposed on a substrate (3). The second washcoat zone (4) is disposed on the third washcoat zone (2). The fourth washcoat zone (5) is disposed on the first washcoat zone (1).

25

DETAILED DESCRIPTION OF THE INVENTION

The oxidation catalyst of the invention comprises, or may consist of, a first washcoat region for NO_x storage, a second washcoat region for oxidising nitric oxide (NO) and a substrate.

30

The first washcoat region comprises a first platinum group metal (PGM), a first support material and a NO_x storage component.

35

The first PGM may be selected from the group consisting of platinum, palladium, rhodium and a combination of any two or more thereof. Preferably, the first PGM is selected from

platinum, palladium, a combination of platinum and palladium, a combination of platinum and rhodium, a combination of palladium and rhodium, and a combination of platinum, palladium and rhodium.

- 5 It may be preferable that the first PGM is selected from the group consisting of platinum, palladium and a combination of platinum and palladium. The first PGM may be platinum. The first PGM may consist essentially of platinum (e.g. the first PGM is platinum only). The first PGM may be palladium. The first PGM may consist essentially of palladium (e.g. the first PGM is palladium only). The first PGM may be a combination of platinum and palladium. The first PGM may consist essentially of platinum and palladium (e.g. the first PGM is platinum and palladium only). It is preferred that the first PGM is selected from the group consisting of platinum and a combination of platinum and palladium.
- 10

The first washcoat region may comprise the first PGM as the only platinum group metal(s). Thus, the only PGM(s) present in the first washcoat region is/are defined by the first PGM.

15

When the first PGM is a combination of platinum and palladium, then the first PGM may be in the form of an alloy, preferably a bimetallic alloy. Thus, the first PGM may comprise, or consist essentially of, an alloy of platinum and palladium.

20

The first washcoat region typically has a total loading of the first PGM of 5 to 300 g ft⁻³. It is preferred that the first washcoat region has a total loading of the first PGM of 10 to 250 g ft⁻³ (e.g. 75 to 175 g ft⁻³), more preferably 15 to 200 g ft⁻³ (e.g. 50 to 150 g ft⁻³), still more preferably 20 to 150 g ft⁻³.

25

When the first PGM is a combination of platinum and palladium, then typically the first washcoat region comprises a ratio by weight of platinum to palladium of 20:1 to 1:20 (e.g. 15:1 to 1:15), preferably 10:1 to 1:10 (e.g. 7.5:1 to 1:7.5), more preferably 5:1 to 1:5 (e.g. 3:1 to 1:3), and even more preferably 2.5:1 to 1:1.

30

It is preferred that when the first PGM is a combination of platinum and palladium then the first washcoat region comprises a total weight of platinum that is greater than or equal to the total weight of palladium (e.g. the ratio of Pt:Pd by weight is $\geq 1:1$). More preferably, the first washcoat region comprises a total weight of platinum that is greater than the total weight of palladium (e.g. the ratio of Pt:Pd by weight is $> 1:1$).

35

Advantageous CO and/or HC light off activity can be obtained when the total weight of platinum is greater than or equal to the total weight of palladium in the first washcoat region.

- 5 It is generally preferred that the first washcoat region comprises a ratio by weight of platinum to palladium of 20:1 to 1:1 (e.g. 15.1:1 to 1.1:1), more preferably 10:1 to 1.25:1 (e.g. 7.5:1 to 1.5:1), and still more preferably 5:1 to 2:1.

It is intended that the primary function of the first washcoat region is NO_x storage.

- 10 However, it is appreciated that in some embodiments of the oxidation catalyst, the first washcoat region may also oxidise carbon monoxide (CO) and hydrocarbons (HCs).

- Typically, the first PGM is disposed or supported on the first support material. The first PGM may be disposed directly onto or is directly supported by the first support material
15 (e.g. there is no intervening support material between the first PGM and the first support material). For example, platinum and/or palladium can be dispersed on the first support material.

- When the first PGM is a combination of platinum and palladium, then the platinum may
20 be disposed or supported on the first support material and/or the palladium may be disposed or supported on the first support material. It may be preferable that both the platinum and the palladium are disposed or supported on the first support material (i.e. the same support material is used for both platinum and palladium).

- 25 When the first PGM is a combination of platinum and palladium, a combination of palladium and rhodium or a combination of platinum, palladium and rhodium, then the first washcoat region may further comprise a palladium support material. The first washcoat region may therefore comprise, or consist essentially of, a first PGM, a first support material, a palladium support material and a NO_x storage component. The
30 platinum and/or rhodium may be disposed or supported on the first support material and the palladium may be disposed or supported on the palladium support material. The first support material and the palladium support material are preferably different (e.g. different composition).

- 35 When the first PGM is a combination of platinum and palladium, a combination of platinum and rhodium or a combination of platinum, palladium and rhodium, then the first washcoat region may further comprise a platinum support material. The first washcoat

region may therefore comprise, or consist essentially of, a first PGM, a first support material, a platinum support material and a NO_x storage component. The platinum may be disposed or supported on the platinum support material and the palladium and/or rhodium may be disposed or supported on the first support material. The first support material and the platinum support material are preferably different (e.g. different composition).

Typically, the first support material comprises, or consists essentially of, a refractory metal oxide. Refractory metal oxides suitable for use as a catalytic component of an oxidation catalyst for a diesel engine are well known in the art.

The refractory metal oxide is typically selected from the group consisting of alumina, silica, titania, zirconia, ceria and a mixed or composite oxide thereof, such as a mixed or composite oxide of two or more thereof. For example, the refractory metal oxide may be selected from the group consisting of alumina, silica, titania, zirconia, ceria, silica-alumina, titania-alumina, zirconia-alumina, ceria-alumina, titania-silica, zirconia-silica, zirconia-titania, ceria-zirconia and alumina-magnesium oxide.

The first support material, or the refractory metal oxide thereof, may optionally be doped (e.g. with a dopant). The dopant may be selected from the group consisting of zirconium (Zr), titanium (Ti), silicon (Si), yttrium (Y), lanthanum (La), praseodymium (Pr), samarium (Sm), neodymium (Nd) and an oxide thereof.

The inclusion of a dopant can thermally stabilise the refractory metal oxide or support material. It is to be understood that any reference to "doped" in this context refers to a material where the bulk or host lattice of the refractory metal oxide is substitution doped or interstitially doped with a dopant. In some instances, small amounts of the dopant may be present at a surface of the refractory metal oxide. However, most of the dopant will generally be present in the body of the refractory metal oxide. The chemical and/or physical properties of a refractory metal oxide are often affected by the presence of a dopant.

When the first support material, or the refractory metal oxide thereof, is doped, the total amount of dopant is 0.25 to 5 % by weight, preferably 0.5 to 3 % by weight (e.g. about 1 % by weight).

The first support material, or the refractory metal oxide thereof, may comprise, or consist essentially of, alumina doped with a dopant. It is particularly preferred that the first support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina doped with a dopant when the first washcoat region comprises an alkaline earth metal.

The alumina may be doped with a dopant comprising silicon (Si), magnesium (Mg), barium (Ba), lanthanum (La), cerium (Ce), titanium (Ti), or zirconium (Zr) or a combination of two or more thereof. The dopant may comprise, or consist essentially of, an oxide of silicon, an oxide of magnesium, an oxide of barium, an oxide of lanthanum, an oxide of cerium, an oxide of titanium or an oxide of zirconium. Preferably, the dopant comprises, or consists essentially of, silicon, magnesium, barium, cerium, or an oxide thereof, particularly silicon, or cerium, or an oxide thereof. More preferably, the dopant comprises, or consists essentially of, silicon, magnesium, barium, or an oxide thereof; particularly silicon, magnesium, or an oxide thereof; especially magnesium or an oxide thereof.

Examples of alumina doped with a dopant include alumina doped with silica, alumina doped with magnesium oxide, alumina doped with barium or barium oxide, alumina doped with lanthanum oxide, or alumina doped with ceria, particularly alumina doped with silica, alumina doped with lanthanum oxide, or alumina doped with ceria. It is preferred that the alumina doped with a dopant is alumina doped with silica, alumina doped with barium or barium oxide, or alumina doped with magnesium oxide. More preferably, the alumina doped with a dopant is alumina doped with silica or alumina doped with magnesium oxide. Even more preferably, the alumina doped with a dopant is alumina doped with magnesium oxide. Alumina doped with a dopant can be prepared using methods known in the art.

When the alumina is alumina doped with silica, then the alumina is doped with silica in a total amount of 0.5 to 45 % by weight (i.e. % by weight of the alumina), preferably 1 to 40 % by weight, more preferably 1.5 to 30 % by weight (e.g. 1.5 to 10 % by weight), particularly 2.5 to 25 % by weight, more particularly 3.5 to 20 % by weight (e.g. 5 to 20 % by weight), even more preferably 4.5 to 15 % by weight.

When the alumina is alumina doped with magnesium oxide, then the alumina is doped with magnesium oxide in an amount as defined above or an amount of 1 to 40 % by weight (i.e. % by weight of the alumina), such as 1 to 30 % by weight, preferably 5 to

28 % by weight, such as 5 to 25 % by weight. More preferably, the alumina is doped with magnesium oxide in amount of 10 to 25 % by weight.

5 It is preferred that the first support material, or the refractory metal oxide thereof, is not doped with a dopant comprising, or consisting essentially of manganese. Thus, the first support material, or the refractory metal oxide thereof, is not promoted with a promoter, such as a promoter selected from the group consisting of tin, manganese, indium, group VIII metal (e.g. Fe, Co, Ni, Ru, Rh, Pd, Os, Ir and Pt, particularly Ir) and combinations thereof.

10

Alternatively or additionally, the first support material, or refractory metal oxide thereof, may comprise, or consist essentially of, an alkaline earth metal aluminate. The term "alkaline earth metal aluminate" generally refers to a compound of the formula MAI_2O_4 where "M" represents the alkaline earth metal, such as Mg, Ca, Sr or Ba. Such
15 compounds generally comprise a spinel structure. These compounds can be prepared using conventional methods known in the art or by using a method described in EP 0945165, US 6,217,837 or US 6,517,795.

20

Typically, the alkaline earth metal aluminate is magnesium aluminate ($MgAl_2O_4$), calcium aluminate ($CaAl_2O_4$), strontium aluminate ($SrAl_2O_4$), barium aluminate ($BaAl_2O_4$), or a mixture of two or more thereof. Preferably, the alkaline earth metal aluminate is magnesium aluminate ($MgAl_2O_4$).

25

In general, when the first support material, or the refractory metal oxide thereof, comprises or consists essentially of a mixed or composite oxide of alumina (e.g. silica-alumina, alumina-magnesium oxide or a mixture of alumina and ceria), then preferably the mixed or composite oxide of alumina comprises at least 50 to 99 % by weight of alumina, more preferably 70 to 95 % by weight of alumina, even more preferably 75 to 90 % by weight of alumina.

30

When the first support material, or refractory metal oxide thereof, comprises or consists essentially of ceria-zirconia, then the ceria-zirconia may consist essentially of 20 to 95 % by weight of ceria and 5 to 80 % by weight of zirconia (e.g. 50 to 95 % by weight ceria and 5 to 50 % by weight zirconia), preferably 35 to 80 % by weight of ceria and 20 to
35 65 % by weight zirconia (e.g. 55 to 80 % by weight ceria and 20 to 45 % by weight zirconia), even more preferably 45 to 75 % by weight of ceria and 25 to 55 % by weight zirconia.

In general, the palladium support material comprises, or consists essentially of, a refractory metal oxide. The palladium support material, or the refractory metal oxide thereof, may be a support material as herein defined above with reference to the first support material. When the first washcoat region comprises a palladium support material, it is preferred that the palladium support material, or the refractory metal oxide thereof, comprises, or consists essentially of, ceria and/or a mixed or composite oxide of ceria, such as ceria-zirconia.

Typically, the platinum support material comprises, or consists essentially of, a refractory metal oxide. The platinum support material, or the refractory metal oxide thereof, may be a support material as herein defined above with reference to the first support material. When the first washcoat region comprises a platinum support material, it is preferred that the platinum support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina, wherein the alumina is optionally doped with a dopant, such as described above. When the platinum support material comprises alumina doped with a dopant, it is preferred that the dopant comprises, or consists essentially of, silicon, magnesium, cerium, lanthanum or an oxide thereof, more preferably magnesium or an oxide thereof.

The first support material (or the refractory metal oxide thereof), the platinum support material (or the refractory metal oxide thereof) and the palladium support material (or the refractory metal oxide thereof) are not doped with a dopant comprising, or consisting essentially of manganese. Thus, the platinum support material (or the refractory metal oxide thereof) and/or the palladium support material (or the refractory metal oxide thereof) is not promoted with a promoter, such as a promoter selected from the group consisting of tin, manganese, indium, group VIII metal (e.g. Fe, Co, Ni, Ru, Rh, Pd, Os, Ir and Pt, particularly Ir) and combinations thereof.

The first washcoat region may comprise an amount of support material (e.g. the total amount of the first support material and, when present, platinum support material and palladium support material) of 1.0 to 8.0 g in⁻³ (e.g. 4.0. to 7.5 g in⁻³), preferably 1.5 to 7.5 g in⁻³ (e.g. 4.5 to 7.0 g in⁻³), still more preferably 2.0 to 7.0 g in⁻³ (5.0 to 7.0 g in⁻³ or 4.5 to 6.5 g in⁻³), and even more preferably 2.5 to 6.5 g in⁻³ (e.g. 4.0 to 6.5 g in⁻³).

The first washcoat region may further comprise a hydrocarbon adsorbent material. The hydrocarbon adsorbent material may be a zeolite.

It is preferred that the zeolite is a medium pore zeolite (e.g. a zeolite having a maximum ring size of ten tetrahedral atoms) or a large pore zeolite (e.g. a zeolite having a maximum ring size of twelve tetrahedral atoms). It may be preferable that the zeolite is not a small pore zeolite (e.g. a zeolite having a maximum ring size of eight tetrahedral atoms).

Examples of suitable zeolites or types of zeolite include faujasite, clinoptilolite, mordenite, silicalite, ferrierite, zeolite X, zeolite Y, ultrastable zeolite Y, AEI zeolite, ZSM-5 zeolite, ZSM-12 zeolite, ZSM-20 zeolite, ZSM-34 zeolite, CHA zeolite, SSZ-3 zeolite, SAPO-5 zeolite, offretite, a beta zeolite or a copper CHA zeolite. The zeolite is preferably ZSM-5, a beta zeolite or a Y zeolite.

When the first washcoat region comprises a hydrocarbon adsorbent, the total amount of hydrocarbon adsorbent is 0.05 to 3.00 g in⁻³, particularly 0.10 to 2.00 g in⁻³, more particularly 0.2 to 1.0 g in⁻³. For example, the total amount of hydrocarbon adsorbent may be 0.8 to 1.75 g in⁻³, such as 1.0 to 1.5 g in⁻³.

However, it may be preferable that the first washcoat region does not comprise a hydrocarbon adsorbent material.

Typically, the NO_x storage component comprises an alkali metal, alkaline earth metal and/or a rare earth metal, wherein the rare earth metal is selected from the group consisting of lanthanum, yttrium and a combination thereof. The NO_x storage component generally comprises, or consists essentially of, (i) an oxide, a carbonate or a hydroxide of an alkali metal; (ii) an oxide, a carbonate or a hydroxide of an alkaline earth metal; and/or (iii) an oxide, a carbonate or a hydroxide of a rare earth metal, wherein the rare earth metal is selected from the group consisting of lanthanum, yttrium and a combination thereof.

When the NO_x storage component comprises an alkali metal (or an oxide, a carbonate or a hydroxide thereof), then preferably the alkali metal is selected from the group consisting of potassium (K), sodium (Na), lithium (Li), caesium (Cs) and a combination of two or more thereof. It is preferred that the alkali metal is potassium (K), sodium (Na) or lithium (Li), more preferably the alkali metal is potassium (K) or sodium (Na), and most preferably the alkali metal is potassium (K).

When the NO_x storage component comprises an alkaline earth metal (or an oxide, a carbonate or a hydroxide thereof), then preferably the alkaline earth metal is selected from the group consisting of magnesium (Mg), calcium (Ca), strontium (Sr), barium (Ba) and a combination of two or more thereof. It is preferred that the alkaline earth metal is calcium (Ca), strontium (Sr), or barium (Ba), more preferably strontium (Sr) or barium (Ba), and most preferably the alkaline earth metal is barium (Ba).

It is preferred that the NO_x storage component comprises potassium or barium (e.g. an oxide, a carbonate or a hydroxide of potassium or an oxide, a carbonate or a hydroxide of barium). More preferably, the NO_x storage component comprises barium (e.g. an oxide, a carbonate or a hydroxide of barium).

The NO_x storage component may be disposed or supported (e.g. directly disposed or directly supported) on the first support material. It is preferred that the NO_x storage component is disposed or supported on the first support material when the first support material or the refractory metal oxide thereof comprises or consists essentially of alumina, alumina doped with a dopant, magnesium aluminate, ceria or a mixed or composite oxide of ceria, which is preferably ceria-zirconia. The alumina doped with a dopant is preferably alumina doped with magnesium or an oxide thereof.

When the first washcoat region comprises a platinum support material, the NO_x storage component may be disposed or supported (e.g. directly disposed or directly supported) on the platinum support material (the NO_x storage component may be disposed or supported on the platinum support material in addition or as alternative to the NO_x storage component being disposed or supported on the first support material).

Preferably the platinum support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina, alumina doped with a dopant or magnesium aluminate. The alumina doped with a dopant is preferably alumina doped with lanthanum (or an oxide thereof) or alumina doped with magnesium (or an oxide thereof). It is preferred that the platinum support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina doped with magnesium oxide.

When the NO_x storage component is disposed or supported on the platinum support material, then preferably the first support material or the refractory metal oxide thereof comprises, or consists essentially of, ceria or a mixed or composite oxide of ceria (e.g. ceria-zirconia). More preferably, the palladium support material or the refractory metal oxide thereof comprises, or consists essentially of, ceria.

The first washcoat region may comprise a palladium support material. When the first washcoat region comprises a palladium support material, the NO_x storage component may be disposed or supported (e.g. directly disposed or directly supported) on the palladium support material (the NO_x storage component may be disposed or supported on the platinum support material in addition or as alternative to the NO_x storage component being disposed or supported on the first support material). Preferably the palladium support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina, alumina doped with a dopant, magnesium aluminate, ceria or a mixed or composite oxide of ceria, which is preferably ceria-zirconia. The alumina doped with a dopant is preferably alumina doped with lanthanum (or an oxide thereof) or alumina doped with magnesium (or an oxide thereof). It is preferred that the palladium support material or the refractory metal oxide thereof comprises, or consists essentially of, ceria or a mixed or composite oxide of ceria (e.g. ceria-zirconia). More preferably, the palladium support material or the refractory metal oxide thereof comprises, or consists essentially of, ceria.

When the NO_x storage component is disposed or supported on the palladium support material, then preferably the first support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina, alumina doped with a dopant or magnesium aluminate. The alumina doped with a dopant is preferably alumina doped with lanthanum (or an oxide thereof) or alumina doped with magnesium (or an oxide thereof). It is preferred that the first support material or the refractory metal oxide thereof comprises, or consists essentially of, alumina doped with magnesium oxide.

Additionally or alternatively, the first washcoat region may further comprise a support material for the NO_x storage component. The NO_x storage component is disposed or supported (e.g. directly disposed or directly supported) on the support material for the NO_x storage component. The first PGM is not disposed or supported on the support material for the NO_x storage component.

Typically, the support material for the NO_x storage component comprises, or consists essentially of, ceria or a mixed or composite oxide of ceria, such as a mixed or composite oxide of ceria as defined above. The mixed or composite oxide of ceria is preferably ceria-zirconia. It is preferred that the support material for the NO_x storage component comprises, or consists essentially of, ceria.

When the first PGM comprises rhodium (e.g. is a combination of Pt and Rh, a combination of Pd or Rh, or a combination of Pt, Pd and Rh), the rhodium may be disposed or supported on the first support material and/or the platinum support material (if present) and/or the palladium support material (if present) and/or the support material
5 for the NO_x storage component (if present).

In general, the oxidation catalyst or the first washcoat region comprises a total amount of NO_x storage component of the alkaline earth metal of 250 to 900 g ft⁻³ (e.g. 250 to 500 g ft⁻³), particularly 300 to 850 g ft⁻³ (e.g. 300 to 450 g ft⁻³), more particularly 400 to
10 800 g ft⁻³, such as 450 to 600 g ft⁻³.

When the first PGM comprises, or consists essentially of, platinum (Pt), then preferably the first washcoat region comprises a total weight of the alkaline earth metal that is greater than the total weight of the platinum (Pt).
15

It may, however, be preferable that the first washcoat region does not comprise rhodium and/or an alkali metal. Thus, the first washcoat region may not comprise rhodium and/or an alkali metal, particularly an alkali metal disposed or supported on a support material.

20 For the avoidance of doubt, a general feature of the first washcoat region (i.e. in any or all of the embodiments above) is that the first washcoat region is substantially free of manganese or an oxide thereof. The first washcoat region preferably does not comprise manganese or an oxide thereof.

25 The oxidation catalyst of the invention comprises a second washcoat region for oxidising nitric oxide (NO) to nitrogen dioxide (NO₂). The second washcoat region comprises or may consist essentially of platinum (Pt), manganese or an oxide thereof, and a second support material.

30 The second washcoat region may further comprise palladium, such as palladium disposed or supported on the second support material. When the second washcoat region comprises palladium, then the ratio of platinum to palladium by total weight is generally $\geq 2:1$ (e.g. Pt:Pd 1:0 to 2:1), more preferably $\geq 4:1$ (e.g. Pt:Pd 1:0 to 4:1). Advantageous CO and/or HC light off activity can be obtained when the total weight of
35 platinum is greater than the total weight of palladium in the first washcoat region.

It is generally preferred that the second washcoat region is substantially free of palladium, particularly substantially free of palladium (Pd) disposed or supported on the second support material. More preferably, the second washcoat region does not comprise palladium, particularly palladium disposed or supported on the second support material. The presence of palladium, particularly in a large amount, in the second washcoat region can be detrimental to NO oxidation activity. The NO oxidising activity of palladium is generally poor under the typical usage conditions for a diesel oxidation catalyst. Also, any palladium that is present may react with some of the platinum that is present to form an alloy. This can also be detrimental to the NO oxidation activity of the second washcoat region because platinum-palladium alloys are not as active toward NO oxidation as platinum is by itself.

Generally, the second washcoat region comprises platinum (Pt) as the only platinum group metal. The second washcoat region preferably does not comprise one or more other platinum group metals, such as ruthenium (Ru), rhodium (Rh), palladium (Pd), osmium (Os) and/or iridium (Ir). More preferably, the second washcoat region does not comprise one or more other platinum group metals, such as ruthenium (Ru), rhodium (Rh), palladium (Pd), osmium (Os) and/or iridium (Ir), supported on the second support material.

The platinum (Pt) is typically disposed or supported on the second support material. The platinum may be disposed directly onto or is directly supported by the second support material (e.g. there is no intervening support material between the platinum and the second support material). For example, platinum can be dispersed on the second support material.

The second washcoat region typically has a total loading of platinum of 5 to 300 g ft⁻³. It is preferred that the second washcoat region has a total loading of platinum of 10 to 250 g ft⁻³ (e.g. 75 to 175 g ft⁻³), more preferably 15 to 200 g ft⁻³ (e.g. 50 to 150 g ft⁻³), still more preferably 20 to 150 g ft⁻³.

It is intended that the primary function of the second washcoat region is oxidising nitric oxide (NO) to nitrogen dioxide (NO₂). However, it is appreciated that in some embodiments of the oxidation catalyst, the second washcoat region may also oxidise some carbon monoxide (CO) and/or some hydrocarbons (HCs) during use.

The second washcoat region comprises manganese (Mn). The manganese may be present in an elemental form or as an oxide. The second washcoat region typically comprises manganese or an oxide thereof.

- 5 The manganese (Mn) is typically disposed or supported on the second support material. The manganese (Mn) may be disposed directly onto or is directly supported by the second support material (e.g. there is no intervening support material between the Mn and the second support material).
- 10 The second washcoat region typically has a total loading of manganese (Mn) of 5 to 500 g ft⁻³. It is preferred that the second washcoat region has a total loading of manganese (Mn) of 10 to 250 g ft⁻³ (e.g. 75 to 175 g ft⁻³), more preferably 15 to 200 g ft⁻³ (e.g. 50 to 150 g ft⁻³), still more preferably 20 to 150 g ft⁻³.
- 15 Typically, the second washcoat region comprises a ratio of Mn:Pt by weight of $\leq 5:1$, more preferably $< 5:1$.

In general, the second washcoat region comprises a ratio of Mn:Pt by weight of $\geq 0.2:1$ (e.g. $\geq 0.5:1$), more preferably $> 0.2:1$ (e.g. $> 0.5:1$).

20

The second washcoat region may comprise a ratio by total weight of manganese (Mn) to platinum of 5:1 to 0.2:1, such as 5:1 to 0.5:1 (e.g. 5:1 to 2:3 or 5:1 to 1:2), preferably 4.5:1 to 1:1 (e.g. 4:1 to 1.1:1), more preferably 4:1 to 1.5:1. The ratio of Mn:Pt by weight can be important in obtaining the activity toward NO described herein.

25

Typically, the second support material comprises, or consists essentially of, a refractory metal oxide. The refractory metal oxide is typically selected from the group consisting of alumina, silica, titania, zirconia, ceria and a mixed or composite oxide thereof, such as a mixed or composite oxide of two or more thereof. For example, the refractory metal

30 oxide may be selected from the group consisting of alumina, silica, titania, zirconia, ceria, silica-alumina, titania-alumina, zirconia-alumina, ceria-alumina, titania-silica, zirconia-silica, zirconia-titania, ceria-zirconia and alumina-magnesium oxide.

35

The second support material, or the refractory metal oxide thereof, may optionally be doped (e.g. with a dopant). The dopant may be selected from the group consisting of zirconium (Zr), titanium (Ti), silicon (Si), yttrium (Y), lanthanum (La), praseodymium (Pr), samarium (Sm), neodymium (Nd) and an oxide thereof.

When the second support material, or the refractory metal oxide thereof, is doped, the total amount of dopant is 0.25 to 5 % by weight, preferably 0.5 to 3 % by weight (e.g. about 1 % by weight).

- 5 The second support material, or the refractory metal oxide thereof, may comprise, or consist essentially of, alumina doped with a dopant. It is particularly preferred that the second support material, or the refractory metal oxide thereof, comprises, or consists essentially of, alumina doped with a dopant. It has been found that the combination of manganese (Mn), platinum (Pt) and a doped alumina support material, particularly an alumina support material doped with silica, provides excellent NO oxidation activity and
10 can stabilise NO oxidation activity of the oxidation catalyst over its lifetime.

- The alumina may be doped with a dopant comprising silicon (Si), magnesium (Mg), barium (Ba), lanthanum (La), cerium (Ce), titanium (Ti), or zirconium (Zr) or a
15 combination of two or more thereof. The dopant may comprise, or consist essentially of, an oxide of silicon, an oxide of magnesium, an oxide of barium, an oxide of lanthanum, an oxide of cerium, an oxide of titanium or an oxide of zirconium. Preferably, the dopant comprises, or consists essentially of, silicon, magnesium, barium, cerium, or an oxide thereof; particularly silicon, cerium, or an oxide thereof. More preferably, the dopant
20 comprises, or consists essentially of, silicon, magnesium, barium, or an oxide thereof; particularly silicon, magnesium, or an oxide thereof; especially silicon or an oxide thereof.

- Examples of alumina doped with a dopant include alumina doped with silica, alumina doped with magnesium oxide, alumina doped with barium or barium oxide, alumina
25 doped with lanthanum oxide, or alumina doped with ceria, particularly alumina doped with silica, alumina doped with lanthanum oxide, or alumina doped with ceria. It is preferred that the alumina doped with a dopant is alumina doped with silica, alumina doped with barium or barium oxide, or alumina doped with magnesium oxide. More preferably, the alumina doped with a dopant is alumina doped with silica or alumina
30 doped with magnesium oxide. Even more preferably, the alumina doped with a dopant is alumina doped with silica.

- When the alumina is alumina doped with silica, then the alumina is doped with silica in a total amount of 0.5 to 45 % by weight (i.e. % by weight of the alumina), preferably 1 to
35 40 % by weight, more preferably 1.5 to 30 % by weight (e.g. 1.5 to 10 % by weight), particularly 2.5 to 25 % by weight, more particularly 3.5 to 20 % by weight (e.g. 5 to 20 % by weight), even more preferably 4.5 to 15 % by weight.

When the alumina is alumina doped with magnesium oxide, then the alumina is doped with magnesium in an amount as defined above or an amount of 1 to 30 % by weight (i.e. % by weight of the alumina), preferably 5 to 25 % by weight.

- 5 It is preferred that the second support material, or the refractory metal oxide thereof, is not doped with a dopant comprising, or consisting essentially of manganese. Thus, the second support material, or the refractory metal oxide thereof, is not promoted with a promoter, such as a promoter selected from the group consisting of tin, manganese, indium, group VIII metal (e.g. Fe, Co, Ni, Ru, Rh, Pd, Os, Ir and Pt, particularly Ir) and
10 combinations thereof.

Alternatively or additionally, the second support material, or refractory metal oxide thereof, may comprise, or consist essentially of, an alkaline earth metal aluminate, such as an alkaline earth metal aluminate having a spinel structure.

15

Typically, the alkaline earth metal aluminate is magnesium aluminate (MgAl_2O_4), calcium aluminate (CaAl_2O_4), strontium aluminate (SrAl_2O_4), or barium aluminate (BaAl_2O_4), or a mixture of two or more thereof. Preferably, the alkaline earth metal aluminate is magnesium aluminate (MgAl_2O_4).

20

- In general, when the second support material, or the refractory metal oxide thereof, comprises or consists essentially of a mixed or composite oxide of alumina (e.g. silica-alumina, alumina-magnesium oxide or a mixture of alumina and ceria), then preferably the mixed or composite oxide of alumina comprises at least 50 to 99 % by weight of
25 alumina, more preferably 70 to 95 % by weight of alumina, even more preferably 75 to 90 % by weight of alumina.

- When the second support material, or refractory metal oxide thereof, comprises or consists essentially of ceria-zirconia, then the ceria-zirconia may consist essentially of 20
30 to 95 % by weight of ceria and 5 to 80 % by weight of zirconia (e.g. 50 to 95 % by weight ceria and 5 to 50 % by weight zirconia), preferably 35 to 80 % by weight of ceria and 20 to 65 % by weight zirconia (e.g. 55 to 80 % by weight ceria and 20 to 45 % by weight zirconia), even more preferably 45 to 75 % by weight of ceria and 25 to 55 % by weight zirconia.

35

Typically, the second washcoat region comprises an amount of the second support material of 0.1 to 4.5 g in⁻³ (e.g. 0.25 to 4.0 g in⁻³), preferably 0.5 to 3.0 g in⁻³, more preferably 0.6 to 2.5 g in⁻³ (e.g. 0.75 to 1.5 g in⁻³).

- 5 In some applications, it may generally be preferable that the second washcoat region is substantially free of a hydrocarbon adsorbent material, particularly a zeolite. Thus, the second washcoat region may not comprise a hydrocarbon adsorbent material.

10 It may be further preferable that the second washcoat region is substantially free of a zeolite catalyst. Thus, the second washcoat region may not comprise the zeolite catalyst.

15 The second washcoat region typically does not comprise indium and/or iridium. More preferably, the second washcoat region does not comprise indium, iridium and/or magnesium.

20 It may be preferable that the second washcoat region does not comprise cerium oxide or a mixed or composite oxide thereof, such as (i) a mixed or composite oxide of cerium oxide and alumina and/or (ii) a mixed or composite oxide of cerium oxide and zirconia.

25 Additionally or alternatively, the second washcoat region may be substantially free of rhodium, an alkali metal and/or an alkaline earth metal, particularly an alkali metal and/or an alkaline earth metal disposed or supported on the second support material. Thus, the second washcoat may not comprise rhodium, an alkali metal and/or an alkaline earth metal, particularly an alkali metal and/or an alkaline earth metal disposed or supported on the second support material.

30 Typically, the first washcoat region comprises > 25 % of the total concentration of platinum group metal (i.e. of the oxidation catalyst). It is preferred that the first washcoat region comprises > 30 %, more preferably ≥ 40 %, of the total concentration of platinum group metal.

35 In general, the total concentration of the first PGM is greater than the total concentration of platinum in the second washcoat region.

The first washcoat region and/or the second washcoat region may be disposed or supported on the substrate.

The first washcoat region may be disposed directly on to the substrate (i.e. the first washcoat region is in contact with a surface of the substrate; see Figures 1 to 5). The second washcoat region may be:

- 5
- (a) disposed or supported on the first washcoat region (e.g. see Figures 2, 4 and 5); and/or
 - (b) disposed directly on to the substrate [i.e. the second washcoat region is in contact with a surface of the substrate] (e.g. see Figures 1, 3, 4); and/or
 - (c) in contact with the first washcoat region [i.e. the second washcoat region is adjacent to, or abuts, the first washcoat region].

10

When the second washcoat region is disposed directly on to the substrate, then a part or portion of the second washcoat region may be in contact with the first washcoat region or the first washcoat region and the second washcoat region may be separated (e.g. by a gap).

15

When the second washcoat region is disposed or supported on the first washcoat region, all or part of the second washcoat region is preferably disposed directly on to the first washcoat region (i.e. the second washcoat region is in contact with a surface of the first washcoat region). The second washcoat region may be a second washcoat layer and the first washcoat region may be a first washcoat layer.

20

The second washcoat region may be disposed directly on to the substrate (i.e. the second washcoat region is in contact with a surface of the substrate; see Figures 1, 3 and 4). The first washcoat region may be:

- 25
- (i) disposed or supported on the second washcoat region (e.g. see Figures 3 and 4); and/or
 - (ii) disposed directly on to the substrate [i.e. the first washcoat region is in contact with a surface of the substrate] (e.g. see Figures 3 and 4); and/or
 - (iii) in contact with the second washcoat region [i.e. the first washcoat region is adjacent to, or abuts, the second washcoat region].
- 30

The first washcoat region may be disposed directly on to the second washcoat region (i.e. the first washcoat region is in contact with a surface of the second washcoat region).

35

It is preferred that only a portion or part of the first washcoat region is disposed or supported on the second washcoat region. Thus, the first washcoat region does not completely overlap or cover the second washcoat region.

In general, it is possible that both the first washcoat region and the second washcoat region are not directly disposed on the substrate (i.e. neither the first washcoat region nor the second washcoat region is in contact with a surface of the substrate).

- 5 The first washcoat region may be a first washcoat layer or a first washcoat zone. When the first washcoat region is a first washcoat zone, then typically the first washcoat zone has a length of 10 to 90 % of the length of the substrate (e.g. 10 to 45 %), preferably 15 to 75 % of the length of the substrate (e.g. 15 to 40 %), more preferably 20 to 70 % (e.g. 30 to 65 %, such as 25 to 45 %) of the length of the substrate, still more preferably 25 to 10 65 % (e.g. 35 to 50 %). It is preferred that the length of the first washcoat zone is greater than the length of the second washcoat zone.

- The second washcoat region may be a second washcoat layer or a second washcoat zone. When the second washcoat region is a second washcoat zone, then typically the 15 second washcoat zone has a length of 10 to 90 % of the length of the substrate (e.g. 10 to 45 %), preferably 15 to 75 % of the length of the substrate (e.g. 15 to 40 %), more preferably 20 to 70 % (e.g. 30 to 65 %, such as 25 to 45 %) of the length of the substrate, still more preferably 25 to 65 % (e.g. 35 to 50 %).

- 20 The first washcoat zone may adjoin the second washcoat zone. Preferably, the first washcoat zone is contact with the second washcoat zone. When the first washcoat zone adjoins the second washcoat zone or the first washcoat zone is in contact with the second washcoat zone, then the first washcoat zone and the second washcoat zone may be disposed or supported on the substrate as a layer (e.g. a single layer). Thus, a layer 25 (e.g. a single) may be formed on the substrate when the first and second washcoat zones adjoin or are in contact with one another. Such an arrangement may avoid problems with back pressure.

- The first washcoat zone may be separate from the second washcoat zone. There may 30 be a gap (e.g. a space) between the first washcoat zone and the second washcoat zone.

- The first washcoat zone may overlap the second washcoat zone. Thus, an end portion or part of the first washcoat zone may be disposed or supported on the second washcoat zone. The first washcoat zone may completely or partly overlap the second washcoat 35 zone. When the first washcoat zone overlaps the second washcoat zone, it is preferred that first washcoat zone only partly overlaps the second washcoat zone (i.e. the top,

outermost surface of the second washcoat zone is not completely covered by the first washcoat zone).

5 Alternatively, the second washcoat zone may overlap the first washcoat zone. Thus, an end portion or part of the second washcoat zone may be disposed or supported on the first washcoat zone. The second washcoat zone generally only partly overlaps the first washcoat zone.

10 It is preferred that the first washcoat zone and the second washcoat zone do not substantially overlap.

15 In the oxidation catalyst of the invention, the second washcoat region may be arranged to contact the exhaust gas at the outlet end of the substrate, typically after contact of the exhaust gas with the first washcoat region. This may, for example, be achieved by disposing the second washcoat region/layer/zone at an outlet end of the substrate. The first washcoat region may be arranged or oriented to contact exhaust gas before the second washcoat region. Thus, the first washcoat region may be arranged to contact exhaust gas as it enters the oxidation catalyst and the second washcoat region may be arranged to contact the exhaust gas as it leaves the oxidation catalyst.

20

The oxidation catalyst may show advantageous oxidative activity (e.g. toward CO, HCs and NO), and particularly toward NO, when it has an arrangement that facilitates the contact of the exhaust gas with the washcoat region containing the Pt and Mn shortly before the exhaust gas exits the catalyst and after it has been in contact with the washcoat region for oxidising CO and HCs. In such an arrangement of the oxidation catalyst, as the exhaust gas enters the catalyst it first comes into contact with the first washcoat region. After the exhaust gas has passed through or over the first washcoat region, it comes into contact with the second washcoat region for oxidising NO before it finally passes through the outlet of the oxidation catalyst. It has further been found that when the oxidation catalyst is arranged in such a way, the Mn-containing washcoat region is surprisingly tolerant to poisoning by sulfur from diesel fuel and can retain its NO oxidation activity.

30

35 The second washcoat region is arranged or oriented to contact exhaust after it has contacted the first washcoat region when:

- (a) the second washcoat region is a second washcoat zone disposed at an outlet end of the substrate and optionally the first washcoat region is a first washcoat zone disposed at an inlet end of the substrate; or
- (b) the first washcoat region is a first washcoat layer and the second washcoat region is a second washcoat zone, wherein the second washcoat zone is disposed on the first washcoat layer at an outlet end of the substrate; or
- (c) the first washcoat region is a first washcoat layer and the second washcoat region is a second washcoat layer, and wherein the second washcoat layer is disposed on the first washcoat layer.

10

Typically, the first washcoat region is arranged or oriented to contact exhaust gas before the second washcoat region. Thus, the first washcoat region may be arranged to contact exhaust gas as it enters the oxidation catalyst and the second washcoat region may be arranged to contact the exhaust gas as it leaves the oxidation catalyst. The zoned arrangement of the first and second washcoat regions shown in Figures 1 and 6 to 10 are particularly advantageous in this respect (the exhaust gas enters at the left hand side).

15

The second washcoat region is arranged to contact the exhaust gas at the outlet end of the substrate and after contact of the exhaust gas with the first washcoat region in any one of the first to third oxidation catalyst arrangements described herein below.

20

It is preferred that the second washcoat region is a second washcoat zone. More preferably, the second washcoat zone is disposed or supported at or near an outlet end of the substrate.

25

In a first oxidation catalyst arrangement, the first washcoat region is disposed or supported upstream of the second washcoat zone. Preferably, the first washcoat region is a first washcoat zone. More preferably, the first washcoat zone is disposed or supported at or near an inlet end of the substrate. The Mn-containing second washcoat zone may show good tolerance to sulfur when the oxidation catalyst has such a "zoned" arrangement.

30

In a second oxidation catalyst arrangement, the first washcoat region is a first washcoat layer. It is preferred that the first washcoat layer extends for an entire length (i.e. substantially an entire length) of the substrate, particularly the entire length of the channels of a substrate monolith.

35

The second washcoat zone is typically disposed or supported on the first washcoat layer. Preferably the second washcoat zone is disposed directly on to the first washcoat layer (i.e. the second washcoat zone is in contact with a surface of the first washcoat layer).

- 5 When the second washcoat zone is disposed or supported on the first washcoat layer, it is preferred that the entire length of the second washcoat zone is disposed or supported on the first washcoat layer. The length of the second washcoat zone is less than the length of the first washcoat layer.
- 10 In a third oxidation catalyst arrangement, the first washcoat region is a first washcoat layer. It is preferred that the first washcoat layer extends for an entire length (i.e. substantially an entire length) of the substrate, particularly the entire length of the channels of a substrate monolith.
- 15 The second washcoat region is a second washcoat layer. It is preferred that the second washcoat layer extends for an entire length (i.e. substantially an entire length) of the substrate, particularly the entire length of the channels of a substrate monolith.

The second washcoat layer is disposed or supported on the first washcoat layer.

- 20 Preferably the second washcoat layer is disposed directly on to the first washcoat layer (i.e. the second washcoat layer is in contact with a surface of the first washcoat layer).

- Alternatively, in the oxidation catalyst of the invention, the second washcoat region may be arranged to contact the exhaust gas at the inlet end of the substrate, typically before
25 contact of the exhaust gas with the first washcoat region. This may, for example, be achieved by disposing the second washcoat region/layer/zone at an inlet end of the substrate. The first washcoat region may be arranged or oriented to contact exhaust gas after the second washcoat region. Thus, the second washcoat region may be arranged to contact exhaust gas as it enters the oxidation catalyst and the first washcoat region
30 may be arranged to contact the exhaust gas as it leaves the oxidation catalyst.

- The oxidation catalyst can have an arrangement that facilitates the contact of the exhaust gas with the washcoat region containing the Pt and Mn before the exhaust gas is brought into contact with the washcoat region for NO_x storage. In such an
35 arrangement, as the exhaust gas enters the catalyst it first comes into contact with the second washcoat region, which can oxidise NO to NO₂. After the exhaust gas has passed through or over the second washcoat region, it comes into contact with the first

washcoat region for NO_x storage before it finally passes through the outlet of the oxidation catalyst. For the NO_x storage component in the first washcoat region to adsorb or store NO_x, the NO component of NO_x must be converted into NO₂. The second washcoat region facilitates this conversion.

5

The second washcoat region is arranged or oriented to contact exhaust before it has contacted the first washcoat region when:

- (a) the second washcoat region is a second washcoat zone disposed at an inlet end of the substrate and optionally the first washcoat region is a first washcoat zone disposed at an outlet end of the substrate; or
- (b) the first washcoat region is a first washcoat layer and the second washcoat region is a second washcoat zone, wherein the second washcoat zone is disposed on the first washcoat layer at an inlet end of the substrate; or
- (c) the first washcoat region is a first washcoat layer and the second washcoat region is a second washcoat layer, and wherein the first washcoat layer is disposed on the second washcoat layer.

10

15

Typically, the first washcoat region is arranged or oriented to contact exhaust gas after the second washcoat region. Thus, the first washcoat region may be arranged to contact exhaust gas as it leaves the oxidation catalyst and the second washcoat region may be arranged to contact the exhaust gas as it enters the oxidation catalyst. The zoned arrangement of the first and second washcoat regions shown in Figures 1 and 6 to 10 are particularly advantageous in this respect (the exhaust gas enters at the right hand side).

25

The second washcoat region is arranged to contact the exhaust gas at the inlet end of the substrate and after contact of the exhaust gas with the first washcoat region in any one of the fourth to sixth oxidation catalyst arrangements described herein below.

30

It is preferred that the second washcoat region is a second washcoat zone. More preferably, the second washcoat zone is disposed or supported at or near an inlet end of the substrate.

35

In a fourth oxidation catalyst arrangement, the first washcoat region is disposed or supported downstream of the second washcoat zone. Preferably, the first washcoat region is a first washcoat zone. More preferably, the first washcoat zone is disposed or supported at or near an outlet end of the substrate.

In a fifth oxidation catalyst arrangement, the second washcoat region is a second washcoat layer. It is preferred that the second washcoat layer extends for an entire length (i.e. substantially an entire length) of the substrate, particularly the entire length of the channels of a substrate monolith.

5

The first washcoat region is a first washcoat zone, which is typically disposed or supported on the second washcoat layer. Preferably the first washcoat zone is disposed directly on to the second washcoat layer (i.e. the first washcoat zone is in contact with a surface of the second washcoat layer).

10

When the first washcoat zone is disposed or supported on the second washcoat layer, it is preferred that the entire length of the first washcoat zone is disposed or supported on the second washcoat layer. The length of the first washcoat zone is less than the length of the second washcoat layer.

15

In a sixth oxidation catalyst arrangement, the first washcoat region is a first washcoat layer. It is preferred that the first washcoat layer extends for an entire length (i.e. substantially an entire length) of the substrate, particularly the entire length of the channels of a substrate monolith.

20

The second washcoat region is a second washcoat layer. It is preferred that the second washcoat layer extends for an entire length (i.e. substantially an entire length) of the substrate, particularly the entire length of the channels of a substrate monolith.

25

The first washcoat layer is disposed or supported on the second washcoat layer. Preferably the first washcoat layer is disposed directly on to the second washcoat layer (i.e. the first washcoat layer is in contact with a surface of the second washcoat layer). The Mn-containing second washcoat layer may show good tolerance to sulfur when the oxidation catalyst has such a "layered" arrangement.

30

In general (including the first to sixth oxidation catalyst arrangements above), the oxidation catalyst may further comprise a third washcoat region. Thus, at least one of the first washcoat region and the second washcoat region may be disposed or supported on a third washcoat region. The third washcoat region may or may not comprise a

35

platinum group metal.

The third washcoat region may be disposed directly on to the substrate (i.e. the third washcoat region is in contact with a surface of the substrate; see Figures 6, 8 and 10).

The second washcoat region may be:

- 5 (a) disposed or supported on the third washcoat region (e.g. see Figures 6, 8 and 10); and/or
- (b) disposed directly on to the substrate [i.e. the second washcoat region is in contact with a surface of the substrate]; and/or
- (c) in contact with the third washcoat region [i.e. the second washcoat region is adjacent to, or abuts, the third washcoat region].

10

Additionally or alternatively, the third washcoat region may be disposed or supported on the first washcoat region and/or the second washcoat region (e.g. see Figures 7 and 9).

15

When the third washcoat region is disposed directly on to the substrate, then the first washcoat region may be disposed or supported on the third washcoat region (e.g. see Figure 6). The second washcoat region may also be disposed or supported on the third washcoat region. The third washcoat region may be a third washcoat layer, the first washcoat region may be a first washcoat zone and the second washcoat region may be a second washcoat zone. The second washcoat region/zone may be in contact with the first washcoat region/zone [i.e. the second washcoat region/zone is adjacent to, or abuts, the first washcoat region/zone]. Alternatively, the first washcoat region/zone and the second washcoat region/zone may be separated (e.g. by a gap).

20

25

When the third washcoat region is disposed directly on to the substrate, then the first washcoat region may be disposed directly on to the substrate (e.g. see Figures 8 and 10). The second washcoat region may be disposed or supported on the third washcoat region and/or the first washcoat region, preferably the second washcoat region is disposed or supported on the third washcoat region. The third washcoat region may be a third washcoat zone and the first washcoat region may be a first washcoat zone. The third washcoat region/zone may be in contact with the first washcoat region/zone [i.e. the third washcoat region/zone is adjacent to, or abuts, the first washcoat region/zone]. Alternatively, the first washcoat region/zone and the third washcoat region/zone may be separated (e.g. by a gap).

30

35

The second washcoat region may be a second washcoat layer or a second washcoat zone, preferably a second washcoat zone. When the second washcoat region is a second washcoat zone, the oxidation catalyst may further comprise a fourth washcoat

region (e.g. see Figure 10). The fourth washcoat region may be disposed or supported on the first washcoat zone. The fourth washcoat region may be in contact with the second washcoat zone [i.e. the fourth washcoat region is adjacent to, or abuts, the second washcoat zone]. Alternatively, the fourth washcoat region and the second washcoat zone may be separated (e.g. by a gap).

The fourth washcoat region may be a fourth washcoat zone.

The third washcoat region may be disposed or supported on the second washcoat region (e.g. see Figure 9). The second washcoat region may be a second washcoat zone and the first washcoat region may be a first washcoat zone. The second washcoat region/zone may be in contact with the first washcoat region/zone [i.e. the second washcoat region/zone is adjacent to, or abuts, the first washcoat region/zone]. Alternatively, the first washcoat region/zone and the second washcoat region/zone may be separated (e.g. by a gap).

The third washcoat region may be a third washcoat layer or a third washcoat zone.

When the third washcoat region is a third washcoat zone, then the third washcoat zone typically has a length of 10 to 90 % of the length of the substrate (e.g. 10 to 45 %), preferably 15 to 75 % of the length of the substrate (e.g. 15 to 40 %), more preferably 20 to 70 % (e.g. 30 to 65 %, such as 25 to 45 %) of the length of the substrate, still more preferably 25 to 65 % (e.g. 35 to 50 %).

When the third washcoat region is a third washcoat zone, then the third washcoat zone may be disposed or supported on the second washcoat zone (e.g. see Figure 7). The first washcoat region may be a first washcoat zone. The third washcoat zone may be in contact with the first washcoat zone [i.e. the third washcoat zone is adjacent to, or abuts, the first washcoat zone]. Alternatively, the first washcoat zone and the third washcoat zone may be separated (e.g. by a gap).

When the third washcoat region is a third washcoat zone, then the third washcoat zone may be disposed or supported on the first washcoat zone (e.g. see Figure 9). The second washcoat region may be a second washcoat zone. The third washcoat zone may be in contact with the second washcoat zone [i.e. the third washcoat zone is adjacent to, or abuts, the second washcoat zone]. Alternatively, the second washcoat zone and the third washcoat zone may be separated (e.g. by a gap) and/or are not in

contact (e.g. the second washcoat zone is not in physical contact with the third washcoat zone).

5 When the third washcoat region is a third washcoat layer, then the third washcoat layer may be disposed or supported on both the first washcoat region/zone and the second washcoat region/zone.

10 Substrates for supporting oxidation catalysts for treating an exhaust gas from a diesel engine are well known in the art. Methods of making washcoats and applying washcoats onto a substrate are also known in the art (see, for example, our WO 99/47260, WO 2007/077462 and WO 2011/080525).

15 The substrate typically has a plurality of channels (e.g. for the exhaust gas to flow through). Generally, the substrate is a ceramic material or a metallic material.

It is preferred that the substrate is made or composed of cordierite ($\text{SiO}_2\text{-Al}_2\text{O}_3\text{-MgO}$), silicon carbide (SiC), Fe-Cr-Al alloy, Ni-Cr-Al alloy, or a stainless steel alloy.

20 Typically, the substrate is a monolith (also referred to herein as a substrate monolith). Such monoliths are well-known in the art. The substrate monolith may be a flow-through monolith or a filtering monolith.

25 A flow-through monolith typically comprises a honeycomb monolith (e.g. a metal or ceramic honeycomb monolith) having a plurality of channels extending therethrough, which channels are open at both ends. When the substrate is a flow-through monolith, then the oxidation catalyst of the invention is typically a lean NO_x trap (LNT) [also referred to as a NO_x storage catalyst (NSC)] or is for use as a lean NO_x trap (LNT).

30 A filtering monolith generally comprises a plurality of inlet channels and a plurality of outlet channels, wherein the inlet channels are open at an upstream end (i.e. exhaust gas inlet side) and are plugged or sealed at a downstream end (i.e. exhaust gas outlet side), the outlet channels are plugged or sealed at an upstream end and are open at a downstream end, and wherein each inlet channel is separated from an outlet channel by a porous structure.

35 When the monolith is a filtering monolith, it is preferred that the filtering monolith is a wall-flow filter. In a wall-flow filter, each inlet channel is alternately separated from an

outlet channel by a wall of the porous structure and vice versa. It is preferred that the inlet channels and the outlet channels are arranged in a honeycomb arrangement. When there is a honeycomb arrangement, it is preferred that the channels vertically and laterally adjacent to an inlet channel are plugged at an upstream end and vice versa (i.e. the channels vertically and laterally adjacent to an outlet channel are plugged at a downstream end). When viewed from either end, the alternately plugged and open ends of the channels take on the appearance of a chessboard.

It is preferred that the substrate is a flow-through monolith.

In principle, the substrate may be of any shape or size. However, the shape and size of the substrate is usually selected to optimise exposure of the catalytically active materials in the catalyst to the exhaust gas. The substrate may, for example, have a tubular, fibrous or particulate form. Examples of suitable supporting substrates include a substrate of the monolithic honeycomb cordierite type, a substrate of the monolithic honeycomb SiC type, a substrate of the layered fibre or knitted fabric type, a substrate of the foam type, a substrate of the crossflow type, a substrate of the metal wire mesh type, a substrate of the metal porous body type and a substrate of the ceramic particle type.

In general, the oxidation catalyst of the invention is for use as a lean NO_x trap (LNT).

It is to be understood that any reference to an oxidation catalyst of the invention for use as a lean NO_x trap (LNT) may include passive NO_x adsorber activity (DOC-PNA) or cold start catalyst activity (DOC-CSC).

The invention also provides an exhaust system comprising the oxidation catalyst and an emissions control device. The emissions control device is preferably downstream of the oxidation catalyst.

Examples of an emissions control device include a diesel particulate filter (DPF), a lean NO_x trap (LNT), a lean NO_x catalyst (LNC), a selective catalytic reduction (SCR) catalyst, a diesel oxidation catalyst (DOC), a catalysed soot filter (CSF), a selective catalytic reduction filter (SCRTM) catalyst, an ammonia slip catalyst (ASC) and combinations of two or more thereof. Such emissions control devices are all well known in the art. It is preferred that the emissions control device is a selective catalytic reduction (SCR) catalyst or a selective catalytic reduction filter (SCRTM) catalyst

Some of the aforementioned emissions control devices have filtering substrates. An emissions control device having a filtering substrate may be selected from the group consisting of a diesel particulate filter (DPF), a catalysed soot filter (CSF), and a selective catalytic reduction filter (SCRTM) catalyst.

5

It is preferred that the exhaust system comprises an emissions control device selected from the group consisting of a lean NO_x trap (LNT), an ammonia slip catalyst (ASC), diesel particulate filter (DPF), a selective catalytic reduction (SCR) catalyst, a catalysed soot filter (CSF), a selective catalytic reduction filter (SCRTM) catalyst, and combinations
10 of two or more thereof. More preferably, the emissions control device is selected from the group consisting of a diesel particulate filter (DPF), a selective catalytic reduction (SCR) catalyst, a catalysed soot filter (CSF), a selective catalytic reduction filter (SCRTM) catalyst, and combinations of two or more thereof. Even more preferably, the emissions control device is a selective catalytic reduction (SCR) catalyst or a selective
15 catalytic reduction filter (SCRTM) catalyst.

When the exhaust system of the invention comprises an SCR catalyst or an SCRTM catalyst, then the exhaust system may further comprise an injector for injecting a nitrogenous reductant, such as ammonia, or an ammonia precursor, such as urea or
20 ammonium formate, preferably urea, into exhaust gas downstream of the oxidation catalyst and upstream of the SCR catalyst or the SCRTM catalyst. Such an injector may be fluidly linked to a source (e.g. a tank) of a nitrogenous reductant precursor. Valve-controlled dosing of the precursor into the exhaust gas may be regulated by suitably programmed engine management means and closed loop or open loop feedback
25 provided by sensors monitoring the composition of the exhaust gas. Ammonia can also be generated by heating ammonium carbamate (a solid) and the ammonia generated can be injected into the exhaust gas.

Alternatively or in addition to the injector, ammonia can be generated in situ (e.g. during
30 rich regeneration of the LNT disposed upstream of the SCR catalyst or the SCRTM catalyst). Thus, the exhaust system may further comprise an engine management means for enriching the exhaust gas with hydrocarbons.

The SCR catalyst or the SCRTM catalyst may comprise a metal selected from the group
35 consisting of at least one of Cu, Hf, La, Au, In, V, lanthanides and Group VIII transition metals (e.g. Fe), wherein the metal is supported on a refractory oxide or molecular sieve.

The metal is preferably selected from Ce, Fe, Cu and combinations of any two or more thereof, more preferably the metal is Fe or Cu.

5 The refractory oxide for the SCR catalyst or the SCRFTM catalyst may be selected from the group consisting of Al₂O₃, TiO₂, CeO₂, SiO₂, ZrO₂ and mixed oxides containing two or more thereof. The non-zeolite catalyst can also include tungsten oxide (e.g. V₂O₅/WO₃/TiO₂, WO_x/CeZrO₂, WO_x/ZrO₂ or Fe/WO_x/ZrO₂).

10 It is particularly preferred when an SCR catalyst, an SCRFTM catalyst or a washcoat thereof comprises at least one molecular sieve, such as an aluminosilicate zeolite or a SAPO. The at least one molecular sieve can be a small, a medium or a large pore molecular sieve. By "small pore molecular sieve" herein we mean molecular sieves containing a maximum ring size of 8, such as CHA; by "medium pore molecular sieve" herein we mean a molecular sieve containing a maximum ring size of 10, such as ZSM-
15 5; and by "large pore molecular sieve" herein we mean a molecular sieve having a maximum ring size of 12, such as beta. Small pore molecular sieves are potentially advantageous for use in SCR catalysts.

20 In the exhaust system of the invention, preferred molecular sieves for an SCR catalyst or an SCRFTM catalyst are synthetic aluminosilicate zeolite molecular sieves selected from the group consisting of AEI, ZSM-5, ZSM-20, ERI including ZSM-34, mordenite, ferrierite, BEA including Beta, Y, CHA, LEV including Nu-3, MCM-22 and EU-1, preferably AEI or CHA, and having a silica-to-alumina ratio of about 10 to about 50, such as about 15 to about 40.

25

In a first exhaust system embodiment, the exhaust system comprises the oxidation catalyst of the invention and a catalysed soot filter (CSF). The oxidation catalyst is typically followed by (e.g. is upstream of) the catalysed soot filter (CSF). Thus, for example, an outlet of the oxidation catalyst is connected to an inlet of the catalysed soot
30 filter.

In a second exhaust system embodiment, the exhaust system comprises the oxidation catalyst of the invention, a catalysed soot filter (CSF) and a selective catalytic reduction (SCR) catalyst. The oxidation catalyst is typically followed by (e.g. is upstream of) the catalysed soot filter (CSF). The catalysed soot filter is typically followed by (e.g. is upstream of) the selective catalytic reduction (SCR) catalyst. A nitrogenous reductant injector may be arranged between the catalysed soot filter (CSF) and the selective

catalytic reduction (SCR) catalyst. Thus, the catalysed soot filter (CSF) may be followed by (e.g. is upstream of) a nitrogenous reductant injector, and the nitrogenous reductant injector may be followed by (e.g. is upstream of) the selective catalytic reduction (SCR) catalyst.

5

In a third exhaust system embodiment, the exhaust system comprises the oxidation catalyst of the invention, a selective catalytic reduction (SCR) catalyst and either a catalysed soot filter (CSF) or a diesel particulate filter (DPF).

10 In the third exhaust system embodiment, the oxidation catalyst of the invention is typically followed by (e.g. is upstream of) the selective catalytic reduction (SCR) catalyst. A nitrogenous reductant injector may be arranged between the oxidation catalyst and the selective catalytic reduction (SCR) catalyst. Thus, the oxidation catalyst may be followed by (e.g. is upstream of) a nitrogenous reductant injector, and the nitrogenous reductant
15 injector may be followed by (e.g. is upstream of) the selective catalytic reduction (SCR) catalyst. The selective catalytic reduction (SCR) catalyst are followed by (e.g. are upstream of) the catalysed soot filter (CSF) or the diesel particulate filter (DPF).

A fourth exhaust system embodiment comprises the oxidation catalyst of the invention
20 and a selective catalytic reduction filter (SCRFTM) catalyst. The oxidation catalyst of the invention is typically followed by (e.g. is upstream of) the selective catalytic reduction filter (SCRFTM) catalyst. A nitrogenous reductant injector may be arranged between the oxidation catalyst and the selective catalytic reduction filter (SCRFTM) catalyst. Thus, the oxidation catalyst may be followed by (e.g. is upstream of) a nitrogenous reductant
25 injector, and the nitrogenous reductant injector may be followed by (e.g. is upstream of) the selective catalytic reduction filter (SCRFTM) catalyst.

When the exhaust system comprises a selective catalytic reduction (SCR) catalyst or a selective catalytic reduction filter (SCRFTM) catalyst, such as in the second to fourth
30 exhaust system embodiments described hereinabove, an ASC can be disposed downstream from the SCR catalyst or the SCRFTM catalyst (i.e. as a separate substrate monolith), or more preferably a zone on a downstream or trailing end of the substrate monolith comprising the SCR catalyst can be used as a support for the ASC.

35 Another aspect of the invention relates to a vehicle or an apparatus. The vehicle or apparatus comprises a diesel engine. The diesel engine may be a homogeneous charge compression ignition (HCCI) engine, a pre-mixed charge compression ignition (PCCI)

engine or a low temperature combustion (LTC) engine. It is preferred that the diesel engine is a conventional (i.e. traditional) diesel engine.

5 The vehicle may be a light-duty diesel vehicle (LDV), such as defined in US or European legislation. A light-duty diesel vehicle typically has a weight of < 2840 kg, more preferably a weight of < 2610 kg.

10 In the US, a light-duty diesel vehicle (LDV) refers to a diesel vehicle having a gross weight of \leq 8,500 pounds (US lbs). In Europe, the term light-duty diesel vehicle (LDV) refers to (i) passenger vehicles comprising no more than eight seats in addition to the driver's seat and having a maximum mass not exceeding 5 tonnes, and (ii) vehicles for the carriage of goods having a maximum mass not exceeding 12 tonnes.

15 Alternatively, the vehicle may be a heavy-duty diesel vehicle (HDV), such as a diesel vehicle having a gross weight of > 8,500 pounds (US lbs), as defined in US legislation.

20 The invention also provides a method of modulating the content of NO_x in an exhaust gas from a diesel engine for an emissions control device. The step of (b) passing the treated exhaust gas to an emissions control device typically involves directly passing the treated exhaust gas to the emissions control device. Thus, an outlet of the oxidation catalyst is directly connected (e.g. without intermediary) to an inlet of the emissions control device.

25 Typically, the emissions control device is a selective catalytic reduction (SCR) catalyst, a selective catalytic reduction filter (SCRTM) catalyst, a diesel particulate filter (DPF), or a catalysed soot filter (CSF). It is preferred that the emissions control device is a selective catalytic reduction (SCR) catalyst or a selective catalytic reduction filter (SCRTM) catalyst.

30 Any reference to "modulating the content of NO_x" as used herein, particularly in relation to method or use aspects of the invention, refers to changing (i.e. adjusting) or maintaining, preferably changing, the ratio (in ppm or % volume, typically at the temperature and pressure of the exhaust gas) of NO:NO₂ to be within a predefined ratio at a specific exhaust gas temperature or temperature range. The predefined ratio is
35 typically less than 17:3, preferably to be from 5:1 to 1:5, more preferably 2.5:1 to 1:2.5, and even more preferably 2:1 to 1:2 (e.g. 1.5:1 to 1:1.5 or about 1:1).

The invention also relates to the use of the oxidation catalyst in the regeneration of an emissions control device having a filtering substrate (e.g. a downstream emissions control device having a filtering substrate).

- 5 The emissions control device having a filtering substrate may be selected from the group consisting of a diesel particulate filter (DPF), a catalysed soot filter (CSF), a selective catalytic reduction filter (SCRTM) catalyst and a combination of two or more thereof.

10 When the oxidation catalyst of the invention is used in the regeneration of an emissions control device having a filtering substrate, it may be used in the active or passive regeneration of the emissions control device, preferably the active regeneration.

The oxidation catalyst may be used to regenerate the emissions control device having a filtering substrate at a temperature of at least 220°C, preferably at least 240°C, more
15 preferably at least 260 °C, still more preferably at least 280 °C, by oxidising nitric oxide (NO) to nitrogen dioxide (NO₂).

DEFINITIONS

20 The term “washcoat” is well known in the art and refers to an adherent coating that is applied to a substrate usually during production of a catalyst.

The term “washcoat region” as used herein refers to an area of washcoat on a substrate. A “washcoat region” can, for example, be disposed or supported on a substrate as a “layer” or a “zone”. The area or arrangement of a washcoat on a substrate is generally controlled during the process of applying the washcoat to the substrate. The “washcoat
25 region” typically has distinct boundaries or edges (i.e. it is possible to distinguish one washcoat region from another washcoat region using conventional analytical techniques).

Typically, the “washcoat region” has a substantially uniform length. The reference to a “substantially uniform length” in this context refers to a length that does not deviate (e.g.
30 the difference between the maximum and minimum length) by more than 10 %, preferably does not deviate by more than 5 %, more preferably does not deviate by more than 1 %, from its mean value.

It is preferable that each “washcoat region” has a substantially uniform composition (i.e. there is no substantial difference in the composition of the washcoat when comparing
35 one part of the washcoat region with another part of that washcoat region). Substantially uniform composition in this context refers to a material (e.g. washcoat region) where the

difference in composition when comparing one part of the washcoat region with another part of the washcoat region is 5% or less, usually 2.5% or less, and most commonly 1% or less.

5 The term “washcoat zone” as used herein refers to a washcoat region having a length that is less than the total length of the substrate, such as $\leq 75\%$ of the total length of the substrate. A “washcoat zone” typically has a length (i.e. a substantially uniform length) of at least 5% (e.g. $\geq 5\%$) of the total length of the substrate.

The total length of a substrate is the distance between its inlet end and its outlet end (e.g. the opposing ends of the substrate).

10 Any reference to a “washcoat zone disposed at an inlet end of the substrate” used herein refers to a washcoat zone disposed or supported on a substrate where the washcoat zone is nearer to an inlet end of the substrate than the washcoat zone is to an outlet end of the substrate. Thus, the midpoint of the washcoat zone (i.e. at half its length) is nearer to the inlet end of the substrate than the midpoint is to the outlet end of the substrate.

15 Similarly, any reference to a “washcoat zone disposed at an outlet end of the substrate” used herein refers to a washcoat zone disposed or supported on a substrate where the washcoat zone is nearer to an outlet end of the substrate than the washcoat zone is to an inlet end of the substrate. Thus, the midpoint of the washcoat zone (i.e. at half its length) is nearer to the outlet end of the substrate than the midpoint is to the inlet end of the substrate.

20 When the substrate is a wall-flow filter, then generally any reference to a “washcoat zone disposed at an inlet end of the substrate” refers to a washcoat zone disposed or supported on the substrate that is:

- 25 (a) nearer to an inlet end (e.g. open end) of an inlet channel of the substrate than the washcoat zone is to a closed end (e.g. blocked or plugged end) of the inlet channel, and/or
- (b) nearer to a closed end (e.g. blocked or plugged end) of an outlet channel of the substrate than the washcoat zone is to an outlet end (e.g. open end) of the outlet channel.

30 Thus, the midpoint of the washcoat zone (i.e. at half its length) is (a) nearer to an inlet end of an inlet channel of the substrate than the midpoint is to the closed end of the inlet channel, and/or (b) nearer to a closed end of an outlet channel of the substrate than the midpoint is to an outlet end of the outlet channel.

35 Similarly, any reference to a “washcoat zone disposed at an outlet end of the substrate” when the substrate is a wall-flow filter refers to a washcoat zone disposed or supported on the substrate that is:

(a) nearer to an outlet end (e.g. an open end) of an outlet channel of the substrate than the washcoat zone is to a closed end (e.g. blocked or plugged) of the outlet channel, and/or

5 (b) nearer to a closed end (e.g. blocked or plugged end) of an inlet channel of the substrate than it is to an inlet end (e.g. an open end) of the inlet channel.

Thus, the midpoint of the washcoat zone (i.e. at half its length) is (a) nearer to an outlet end of an outlet channel of the substrate than the midpoint is to the closed end of the outlet channel, and/or (b) nearer to a closed end of an inlet channel of the substrate than the midpoint is to an inlet end of the inlet channel.

10 A washcoat zone may satisfy both (a) and (b) when the washcoat is present in the wall of the wall-flow filter (i.e. the washcoat zone is in-wall).

The acronym "PGM" as used herein refers to "platinum group metal". The term "platinum group metal" generally refers to a metal selected from the group consisting of Ru, Rh, Pd, Os, Ir and Pt, preferably a metal selected from the group consisting of Ru, Rh, Pd, Ir and Pt. In general, the term "PGM" preferably refers to a metal selected from the group consisting of Rh, Pt and Pd.

15 The term "mixed oxide" as used herein generally refers to a mixture of oxides in a single phase, as is conventionally known in the art. The term "composite oxide" as used herein generally refers to a composition of oxides having more than one phase, as is conventionally known in the art.

20 Any reference to washcoat zones that do not "substantially overlap" as used herein refers an overlap (i.e. between the ends of neighbouring zones on a substrate) of less than 10 % of the length of the substrate, preferably less 7.5 % of the length of the substrate, more preferably less than 5 % of the length of the substrate, particularly less than 2.5 % of the length of the substrate, even more preferably less than 1 % of the length of the substrate, and most preferably there is no overlap.

30 The expression "consist essentially" as used herein limits the scope of a feature to include the specified materials, and any other materials or steps that do not materially affect the basic characteristics of that feature, such as for example minor impurities. The expression "consist essentially of" embraces the expression "consisting of".

The expression "substantially free of" as used herein with reference to a material, typically in the context of the content of a washcoat region, a washcoat layer or a washcoat zone, means that the material in a minor amount, such as $\leq 5\%$ by weight, preferably $\leq 2\%$ by weight, more preferably $\leq 1\%$ by weight. The expression "substantially free of" embraces the expression "does not comprise".

Any reference to an amount of dopant, particularly a total amount, expressed as a % by weight as used herein refers to the weight of the support material or the refractory metal oxide thereof.

EXAMPLES

5 The invention will now be illustrated by the following non-limiting examples.

Example 1

Silica doped alumina powder was slurried in water and milled to a $d_{90} < 20$ micron.

Barium acetate was added to the slurry followed by appropriate amounts of soluble platinum and palladium salts. Beta zeolite was added such that the slurry comprised 78%
10 silica doped alumina and 22% zeolite by mass. The slurry was then stirred to homogenise. The resulting washcoat was applied to the inlet channels of a cordierite flow through monolith having 400 cells per square inch using established coating techniques, and then dried.

15 A second slurry was prepared by taking silica-doped alumina powder and milling to a $d_{90} < 20$ micron. Soluble platinum salt was added to the slurry followed by a manganese nitrate solution. The slurry was stirred to homogenise. This second slurry was applied to the outlet channels of the flow through monolith using conventional coating techniques. The part was dried and then calcined at 500°C . The finished catalyst had a total PGM
20 loading of 75 g ft^{-3} and a manganese loading on the outlet coating of 100 g ft^{-3} . The weight ratio of platinum to manganese was 3:5.

Example 2

Silica doped alumina powder was slurried in water and milled to a $d_{90} < 20$ micron.

Barium acetate was added to the slurry followed by appropriate amounts of soluble
25 platinum and palladium salts. Beta zeolite was added such that the slurry comprised 78% silica doped alumina and 22% zeolite by mass. The slurry was then stirred to homogenise. The resulting washcoat was applied to the inlet channels of a cordierite flow through monolith having 400 cells per square inch using established coating techniques, and then dried.

30 A second slurry was prepared by taking silica-doped alumina powder and milling to a $d_{90} < 20$ micron. An appropriate amount of soluble platinum salt was added to the slurry and it was stirred to homogenise. This second slurry was applied to the outlet channels of the flow through monolith using conventional coating techniques. The part was dried and
35 then calcined at 500°C . The finished catalyst had a total PGM loading of 75 g ft^{-3} .

Experimental Results*Measurement of NO oxidation*

Core samples were taken from the catalysts of Examples 1 and 2. All cores were hydrothermally “aged” in an oven at 750°C for 15 hours. Additional cores were taken and kept in the “fresh” condition (i.e. they were kept without any oven thermal treatment).

The catalytic activity was determined using a synthetic gas bench activity test (SCAT). The fresh and aged cores were tested in a simulated catalyst activity testing (SCAT) gas apparatus using the inlet gas mixtures in Table 1. In each case the balance is nitrogen.

Table 1

CO	1500 ppm
HC (as C ₁)	430 ppm
NO	100 ppm
CO ₂	4%
H ₂ O	4%
O ₂	14%
Space velocity	55000/hour

Results

The results from the SCATs are shown in Tables 2 and 3 below.

Table 2

Example No.	NO oxidation at 200°C (%) (Fresh condition)	NO oxidation at 200°C (%) (Aged condition)
1	31	39
2	44	25

The results in Table 2 show that the catalyst of Example 1 (comprises manganese in the Pt zone) in the aged condition at 200°C has greater NO oxidation performance than the catalyst of Example 2.

Table 3

Example No.	NO oxidation at 220°C (%) (Fresh condition)	NO oxidation at 220°C (%) (Aged condition)
1	55	55
2	59	47

The results in Table 3 show that the catalyst of Example 1 in the aged condition at 220°C shows better NO oxidation activity than the catalyst of Example 2.

5

Tables 2 and 3 also show that the difference in NO oxidation activity between the fresh and aged conditions of the catalyst is smaller for Example 1 than for Example 2. Thus, the catalyst of Example 1 shows more stable NO oxidation performance than Example 2. A stable NO oxidation performance between fresh and aged conditions is advantageous

10 for downstream SCR dosing calibrations.

Example 3

A first slurry was prepared as follows: silica-alumina powder was slurried in water and milled to a $d_{90} < 20$ micron. Soluble platinum salt was added followed by manganese nitrate. The mixture was stirred to homogenise.

15

The slurry was applied to the outlet end of a cordierite flow through monolith having 400 cells per square inch using established coating techniques. It was then dried and calcined at 500°C. The Pt loading on the part was 15 g ft⁻³. The manganese loading on the part was 50 g ft⁻³.

20

A second slurry was prepared by taking silica-alumina powder and milling to a $d_{90} < 20$ micron. Barium acetate was added to the slurry followed by appropriate amounts of soluble platinum and palladium salts. Beta zeolite was added such that the slurry comprised 77% silica-alumina and 23% zeolite by mass. The slurry was then stirred to homogenise. The resulting washcoat was applied to the inlet channels of the flow through monolith using established coating techniques. The part was dried and calcined at 500°C. The finished catalyst had a total Pt loading of 45 g ft⁻³ and a Pd loading of 15 g ft⁻³.

25

Example 4

A first slurry was prepared and applied to a cordierite flow through monolith having 400 cells per square using the same methods of Example 3. The Pt loading on the part was 15 g ft⁻³. The manganese loading on the part was 50 g ft⁻³.

5

A second slurry was prepared by taking silica-alumina powder and milling to a d₉₀ < 20 micron. Soluble platinum salt was added followed by manganese nitrate. Beta zeolite was added such that the slurry comprised 77% silica-alumina and 23% zeolite by mass. The slurry was then stirred to homogenise. The resulting washcoat was applied to the inlet channels of the flow through monolith using established coating techniques. The part was dried and calcined at 500°C. The finished catalyst had a total Pt loading of 54 g ft⁻³ and manganese loading of 100 g ft⁻³.

10

Experimental Results*Measurement of oxidation activity*

15

Core samples were taken from each of the catalysts of Examples 3 and 4. The cores were hydrothermally “aged” in an oven at 750°C for 15 hours using 10% water. Catalytic activity was determined using a synthetic gas bench test. The cores were tested in a simulated catalyst activity testing (SCAT) gas apparatus using the inlet gas mixture in Table 1. In each case the balance is nitrogen. The oxidation activity for CO and HC is determined by the light off temperature whereby 50% conversion is achieved (T50). The oxidation activity for NO is determined as the percentage conversion at 300°C.

20

Results

The results from the SCATs are shown in Table 4 below.

25

Table 4

Example No.	T50 CO light off (°C)	T50 HC light off (°C)	NO oxidation at 300°C (%)
3	173	190	19
4	176	190	30

30

Results in Table 4 show that Examples 3 and 4 have very similar T50 light off temperatures for CO and HC. This means Examples 3 and 4 have very similar CO and HC oxidation activity. Example 4 comprises both platinum and manganese on the inlet coating as well as the outlet coating. Example 3 comprises both platinum and manganese on the outlet coating only. Example 4 has higher NO oxidation performance at 300°C than Example 3.

Example 5

Silica-alumina powder was slurried in water and milled to a $d_{90} < 20$ micron. Appropriate amounts of soluble platinum and palladium salts were added. The mixture was stirred to homogenise. The washcoat was applied to a cordierite flow through monolith having 400
5 cells per square inch using established coating techniques. It was then dried and calcined at 500°C. The overall PGM loading for the part was 60 g ft⁻³. The ratio of Pt:Pd was 4:1 by mass.

Example 6

Silica-alumina powder was slurried in water and milled to a $d_{90} < 20$ micron. Appropriate
10 amounts of soluble platinum and palladium salts were added followed by manganese nitrate. The mixture was stirred to homogenise. The washcoat was applied to a cordierite flow through monolith having 400 cells per square inch using established coating techniques. It was then dried and calcined at 500°C. The overall PGM loading for the part was 60 g ft⁻³. The ratio of Pt:Pd was 4:1 by mass. The manganese loading
15 was 100 g ft⁻³.

Example 7

Silica-alumina powder was slurried in water and milled to a $d_{90} < 20$ micron. Appropriate amounts of soluble platinum and palladium salts were added. The mixture was stirred to homogenise. The washcoat was applied to a cordierite flow through monolith having 400
20 cells per square inch using established coating techniques. It was then dried and calcined at 500°C. The overall PGM loading for the part was 60 g ft⁻³. The ratio of Pt:Pd was 10:1 by mass.

Example 8

Silica-alumina powder was slurried in water and milled to a $d_{90} < 20$ micron. Appropriate
25 amounts of soluble platinum and palladium salts were added followed by manganese nitrate. The mixture was stirred to homogenise. The washcoat was applied to a cordierite flow through monolith having 400 cells per square inch using established coating techniques. It was then dried and calcined at 500°C. The overall PGM loading for the part was 60 g ft⁻³. The ratio of Pt:Pd was 10:1 by mass. The manganese loading
30 was 100 g ft⁻³.

Experimental Results

Measurement of oxidation activity

Core samples were taken from each of the catalysts of Examples 5, 6, 7 and 8. The cores were hydrothermally “aged” in an oven at 800 °C for 16 hours using 10% water.
35 The catalytic activity was determined using a synthetic gas bench test. The cores were tested in a simulated catalyst activity testing (SCAT) gas apparatus using the inlet gas

mixture in Table 1. In each case the balance is nitrogen. The oxidation activity for CO and HC is determined by the light off temperature whereby 50% conversion is achieved (T50).

5 Results

The results from the SCATs are shown in Table 5 below.

Table 5

Example No.	T50 CO light off (°C)	T50 HC light off (°C)
5	182	191
6	172	191
7	196	202
8	172	197

- 10 Results in Table 5 show the CO and HC T50 light off temperatures for Examples 5, 6, 7 and 8. Examples 5 and 6 have the same precious metal loading and a Pt:Pd weight ratio of 4:1. Example 6 which comprises manganese has a lower CO T50 light off temperature than Example 5 which does not comprise manganese. The HC T50 light off temperature for Examples 5 and 6 are identical. Examples 7 and 8 have the same
- 15 precious metal loading and Pt:Pd weight ratio of 10:1. Example 8 which comprises manganese has lower CO and HC T50 light off temperatures than Example 7 which does not comprise manganese.

- A greater improvement in light off temperature is seen with the catalyst containing
- 20 manganese at a Pt:Pd weight ratio of 10:1 than at 4:1.

For the avoidance of any doubt, the entire content of any and all documents cited herein is incorporated by reference into the present application.

CLAIMS

1. An oxidation catalyst for treating an exhaust gas from a diesel engine, which oxidation catalyst comprises:
 - a first washcoat region comprising a first platinum group metal (PGM), a first
5 support material and a NO_x storage component;
 - a second washcoat region comprising platinum (Pt), manganese (Mn) and a second support material; and
 - a substrate having an inlet end and an outlet end.
- 10 2. An oxidation catalyst according to claim 1, wherein the manganese (Mn) is disposed or supported on the second support material.
3. An oxidation catalyst according to claim 1 or claim 2, wherein the second
15 washcoat region has a ratio by total weight of manganese (Mn) to platinum of 5:1 to 0.2:1.
4. An oxidation catalyst according to any one of the preceding claims, wherein the second washcoat region comprises palladium.
- 20 5. An oxidation catalyst according to any one of the preceding claims, wherein the second washcoat region has a ratio of platinum to palladium by total weight of 1:0 to 2:1.
6. An oxidation catalyst according to any one of the preceding claims, wherein the
25 second support material comprises a refractory metal oxide selected from the group consisting of alumina, silica, titania, zirconia, ceria and a mixed or composite oxide thereof, wherein the refractory metal oxide is optionally doped with a dopant.
7. An oxidation catalyst according to claim 6, wherein the refractory metal oxide is
30 selected from the group consisting of alumina, silica, titania, zirconia, ceria, silica-alumina, titania-alumina, zirconia-alumina, ceria-alumina, titania-silica, zirconia-silica, zirconia-titania, ceria-zirconia and alumina-magnesium oxide.
8. An oxidation catalyst according to claim 6, wherein the refractory metal oxide is
35 alumina doped with silica.
9. An oxidation catalyst according to any one of the preceding claims, wherein the NO_x storage component comprises an alkali metal, alkaline earth metal and/or a rare

earth metal, wherein the rare earth metal is selected from the group consisting of lanthanum, yttrium and a combination thereof.

10. An oxidation catalyst according to claim 9, wherein the NO_x storage component
5 comprises an alkaline earth metal selected from the group consisting of magnesium (Mg), calcium (Ca), strontium (Sr), barium (Ba) and a combination of two or more thereof.

11. An oxidation catalyst according to any one of the preceding claims, wherein the
10 first PGM is disposed or supported on the first support material.

12. An oxidation catalyst according to any one of the preceding claims, wherein the
first PGM is selected from platinum, palladium, a combination of platinum and palladium,
a combination of platinum and rhodium, a combination of palladium and rhodium, and a
combination of platinum, palladium and rhodium.

13. An oxidation catalyst according to any one of the preceding claims, wherein the
15 first support material comprises a refractory metal oxide comprising an alkaline earth metal aluminate.

14. An oxidation catalyst according to any one of claims 1 to 12, wherein the first
20 support material comprises a refractory metal oxide selected from the group consisting of alumina, silica, titania, zirconia, ceria and a mixed or composite oxide thereof, optionally doped with a dopant.

15. An oxidation catalyst according to claim 14, wherein the refractory metal oxide is
25 selected from the group consisting of alumina, silica, titania, zirconia, ceria, silica-alumina, titania-alumina, zirconia-alumina, ceria-alumina, titania-silica, zirconia-silica, zirconia-titania, ceria-zirconia and alumina-magnesium oxide.

16. An oxidation catalyst according to claim 14, wherein the refractory metal oxide is
30 alumina doped with magnesium oxide.

17. An oxidation catalyst according to any one of the preceding claims, wherein the
35 first PGM is a combination of platinum and palladium, and the platinum is disposed or supported on the first support material and/or the palladium is disposed or supported on the first support material.

18. An oxidation catalyst according to any one of the preceding claims, wherein the first PGM is a combination of platinum and palladium, a combination of palladium and rhodium or a combination of platinum, palladium and rhodium, and wherein the first washcoat region further comprises a palladium support material.

5

19. An oxidation catalyst according to claim 18, wherein the platinum and/or the rhodium is disposed or supported on the first support material and the palladium is disposed or supported on the palladium support material.

10 20. An oxidation catalyst according to claim 18 or 19, wherein the NO_x storage component is disposed or supported on the palladium support material.

21. An oxidation catalyst according to any one of claims 18 to 20, wherein the palladium support material comprises a refractory metal oxide comprising ceria or a
15 mixed or composite oxide of ceria.

22. An oxidation catalyst according to any one of the preceding claims, wherein the second washcoat region is a second washcoat zone disposed at an outlet end of the
20 substrate.

20

23. An oxidation catalyst according to claim 22, wherein the first washcoat region is a first washcoat zone disposed at an inlet end of the substrate.

24. An oxidation catalyst according to any one of claims 1 to 21, wherein the second
25 washcoat region is a second washcoat zone disposed at an inlet end of the substrate.

25. An oxidation catalyst according to claim 24, wherein the first washcoat region is a first washcoat zone disposed at an outlet end of the substrate.

30 26. An oxidation catalyst according to claim 22 or 24, wherein the first washcoat region is a first washcoat layer, and wherein the second washcoat zone is disposed on the first washcoat layer.

35 27. An oxidation catalyst according to any one of claims 1 to 21, wherein the first washcoat region is a first washcoat layer and the second washcoat region is a second washcoat layer, and wherein the second washcoat layer is disposed on the first washcoat layer.

28. An oxidation catalyst according to any one of the preceding claims, wherein the substrate is a through-flow substrate.
- 5 29. An exhaust system comprising an oxidation catalyst as defined in any one of claims 1 to 28 and an emissions control device.
30. An exhaust system according to claim 26, wherein the emissions control device is selected from the group consisting of emissions control device selected from the group
10 consisting of a diesel particulate filter (DPF), a lean NO_x trap (LNT), a lean NO_x catalyst (LNC), a selective catalytic reduction (SCR) catalyst, a diesel oxidation catalyst (DOC), a catalysed soot filter (CSF), a selective catalytic reduction filter (SCRTM) catalyst, an ammonia slip catalyst (ASC) and combinations of two or more thereof.
- 15 31. A vehicle or apparatus comprising a diesel engine and an oxidation catalyst as defined in any one of claims 1 to 28 or an exhaust system as defined in claim 29 or claim 30.

1/2

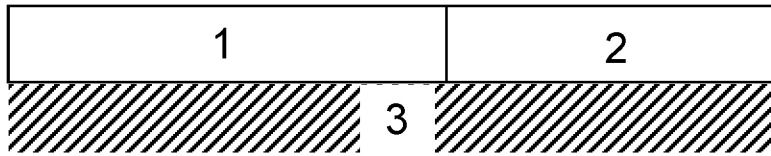


Figure 1

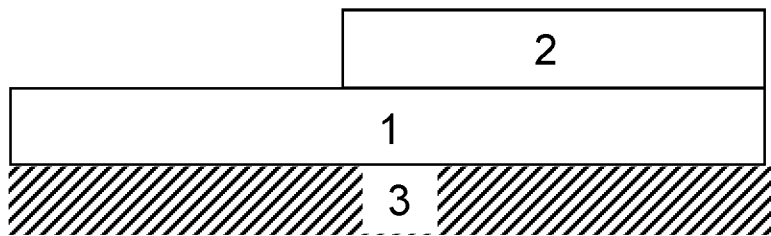


Figure 2

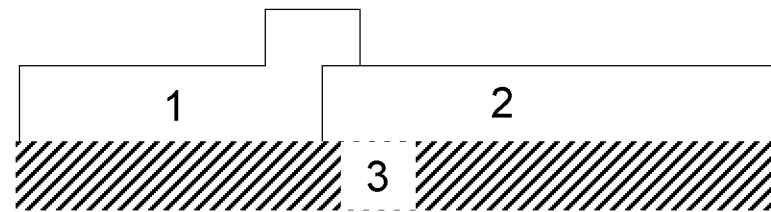


Figure 3

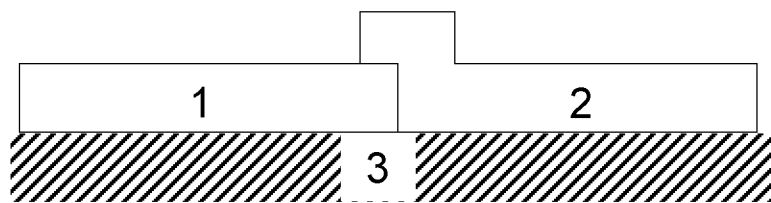


Figure 4



Figure 5

2/2

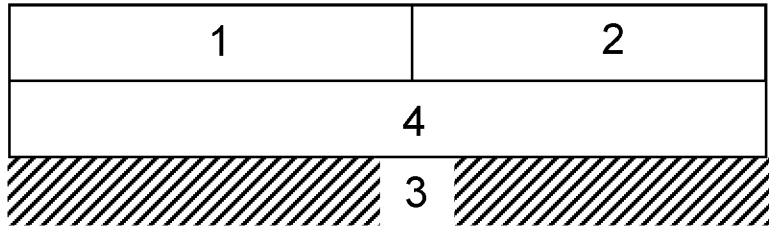


Figure 6

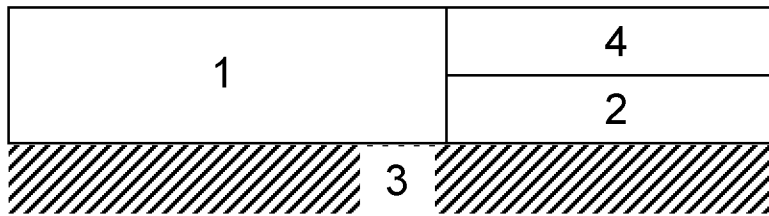


Figure 7

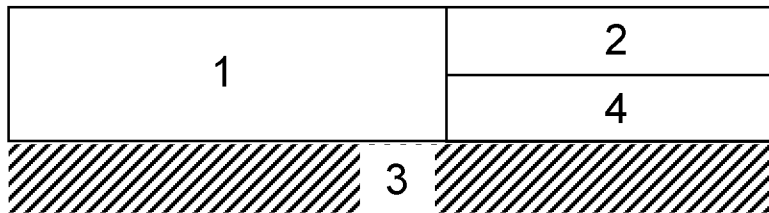


Figure 8

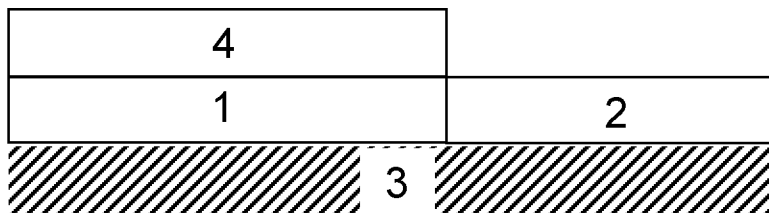


Figure 9

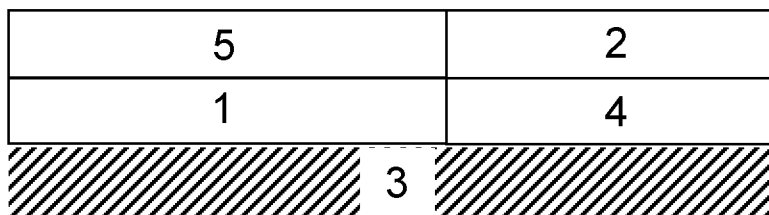


Figure 10

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2015/050145

A. CLASSIFICATION OF SUBJECT MATTER					
INV.	B01J23/34	B01J23/38	B01J23/58	B01J23/63	B01J35/00
	B01J37/02	B01J23/42	B01J23/44	B01J35/04	B01D53/56
	B01D53/62	B01D53/72	B01D53/92	B01D53/94	F01N3/022

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED
Minimum documentation searched (classification system followed by classification symbols) B01J B01D F01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, WPI Data
--

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2006/044974 A1 (ENGELHARD CORP [US]) 27 April 2006 (2006-04-27)	1-31
Y	the whole document page 15, lines 4-30 page 35, lines 1-6 page 38, lines 25-32 page 53, lines 22-31 page 56, line 4 - page 61, line 4 examples 1,2 claims ----- -/--	1-31

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents :	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 2 April 2015	Date of mailing of the international search report 20/04/2015
---	--

Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Gosselin, Daniel
--	--

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2015/050145

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 1 046 423 A2 (DEGUSSA [DE] OMG AG & CO KG [DE] UMICORE AG & CO KG [DE]) 25 October 2000 (2000-10-25) the whole document paragraphs [0001], [0012], [0024], [0031], [0033], [0041] examples 1,4 claims	1-3, 6-15, 22-31
X	US 6 221 804 B1 (YAMADA HIROSHI [JP] ET AL) 24 April 2001 (2001-04-24) the whole document column 1, lines 45-57 column 6, lines 1-45 column 11, lines 4-16 column 13, lines 5-44 column 15, lines 21-54 examples I,III,IX claims	1-12,14, 15,17-31
X	EP 1 264 629 A1 (MITSUBISHI MOTORS CORP [JP]) 11 December 2002 (2002-12-11) the whole document paragraphs [0009], [0020] - [0021], [0023] figures 4,5 claims	1,2,6,7, 9-12,14, 15, 26-29,31
Y	US 8 449 852 B1 (SUNG SHIANG [US] ET AL) 28 May 2013 (2013-05-28) the whole document column 2, lines 5-13 column 3, lines 4-10 column 4, lines 45-67 column 6, lines 43-52 column 7, line 10 - column 8, line 43 column 9, line 46 - column 11, line 12 figures 1,2,6 claims	1-31
A	US 2012/023909 A1 (LAMBERT CHRISTINE KAY [US] ET AL) 2 February 2012 (2012-02-02) the whole document figures 2,3 paragraphs [0021] - [0026]	1-31
A	US 2012/240554 A1 (QI GONGSHIN [US] ET AL) 27 September 2012 (2012-09-27) the whole document paragraph [0067]	1-31
	----- -/--	

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2015/050145

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	GB 2 481 057 A (JOHNSON MATTHEY PLC [GB]) 14 December 2011 (2011-12-14) the whole document figures claims	1-31
A	----- WO 01/14046 A1 (ENGELHARD CORP [US]) 1 March 2001 (2001-03-01) the whole document	1-31
A	----- US 2013/156669 A1 (HAN HYUN-SIK [KR] ET AL) 20 June 2013 (2013-06-20) the whole document -----	1-31

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/GB2015/050145

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2006044974	A1	27-04-2006	EP 1809407 A1 25-07-2007
			JP 2008516768 A 22-05-2008
			KR 20070073598 A 10-07-2007
			US 2005164879 A1 28-07-2005
			WO 2006044974 A1 27-04-2006
EP 1046423	A2	25-10-2000	AR 023446 A1 04-09-2002
			AU 778156 B2 18-11-2004
			BR 0001659 A 31-10-2000
			CA 2306499 A1 23-10-2000
			CN 1271618 A 01-11-2000
			EP 1046423 A2 25-10-2000
			JP 4873582 B2 08-02-2012
			JP 2000312825 A 14-11-2000
			KR 20010049282 A 15-06-2001
			MY 138320 A 29-05-2009
			PL 339830 A1 06-11-2000
			TR 200001071 A2 21-11-2000
			TW 570838 B 11-01-2004
US 6221804	B1	24-04-2001	DE 69919917 D1 14-10-2004
			DE 69919917 T2 02-03-2006
			EP 0931590 A1 28-07-1999
			US 6221804 B1 24-04-2001
EP 1264629	A1	11-12-2002	DE 60201707 D1 02-12-2004
			DE 60201707 T2 10-03-2005
			EP 1264629 A1 11-12-2002
			JP 4645786 B2 09-03-2011
			JP 2002361094 A 17-12-2002
			KR 20020095079 A 20-12-2002
			US 2003008773 A1 09-01-2003
US 8449852	B1	28-05-2013	CA 2857496 A1 06-06-2013
			CN 104053870 A 17-09-2014
			EP 2785988 A1 08-10-2014
			JP 2015500736 A 08-01-2015
			KR 20140105495 A 01-09-2014
			US 8449852 B1 28-05-2013
			WO 2013082367 A1 06-06-2013
US 2012023909	A1	02-02-2012	CN 102953786 A 06-03-2013
			DE 102012214197 A1 21-02-2013
			RU 125628 U1 10-03-2013
			US 2012023909 A1 02-02-2012
			US 2013302232 A1 14-11-2013
US 2012240554	A1	27-09-2012	CN 102688691 A 26-09-2012
			DE 102012204524 A1 11-10-2012
			US 2012240554 A1 27-09-2012
GB 2481057	A	14-12-2011	NONE
WO 0114046	A1	01-03-2001	AT 411097 T 15-10-2008
			AU 6795300 A 19-03-2001
			BR 0013516 A 15-07-2003
			EP 1206312 A1 22-05-2002
			JP 5490342 B2 14-05-2014

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/GB2015/050145

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		JP 2003507177 A	25-02-2003
		JP 2013176774 A	09-09-2013
		KR 20020026597 A	10-04-2002
		WO 0114046 A1	01-03-2001

US 2013156669 A1	20-06-2013	EP 2313196 A2	27-04-2011
		KR 20100013534 A	10-02-2010
		US 2011150742 A1	23-06-2011
		US 2013156669 A1	20-06-2013
		WO 2010013856 A2	04-02-2010
