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Enhancing the performance of Pd/zeolite-based H₂-SCR catalysts: the role of noble metal loading, promoter addition, and combination with a conventional Fe-BEA NH₃-SCR catalyst

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The direct utilization of hydrogen (H₂) as a reductant in the selective catalytic reduction (SCR) of nitrogen oxides (NO_x) presents a promising strategy for mitigating emissions from hydrogen-fueled internal combustion engines (H₂-ICEs). Despite recent advancements, H₂-SCR suffers from a narrow operational temperature window and limited product selectivity. In this study, Pd/TiO₂/HY catalysts with varying palladium loadings were synthesized, characterized, and evaluated for their catalytic performance. A strong correlation between NO conversion and H₂ activation was observed, particularly at low temperatures, with higher Pd content enhancing pollutant abatement. The incorporation of tungsten oxide (WO₃) as a promoter improved catalytic activity and suppressed ammonia (NH₃) formation under both dry and humid conditions, albeit at the cost of increased nitrous oxide (N₂O) emissions. Alternatively, vanadia (V₂O₅) addition reduced secondary emissions but compromised NO_x conversion efficiency. A bifunctional catalyst system combining Pd/TiO₂/HY with a conventional NH₃-SCR catalyst, Fe-BEA, achieved high NO conversion and with more than 90% N₂ selectivity between 180–300 °C also superior product selectivity. Spatially resolved concentration profiles in monolithic samples revealed that Fe-BEA mediates N₂O suppression and facilitates the rapid consumption of NH₃ species formed *in situ* over Pd/TiO₂/HY. These findings underscore the potential of advanced catalyst combinations to overcome current limitations of H₂-SCR and pave the way for cleaner hydrogen-based combustion technologies.

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Introduction

On the way towards a decarbonization of the transport sector, lean-burn hydrogen internal combustion engines (H₂-ICE) represent a promising technology, as hydrogen combustion yields only water as a reaction product.^{1,2} However, the elevated in-cylinder temperatures associated with hydrogen combustion promote the formation of thermal nitrogen oxides (NO_x), necessitating downstream catalytic abatement. In addition to established technologies like NO_x storage and reduction (NSR) catalysts (commonly Pt/BaO/Al₂O₃ or similar formulations) and selective catalytic reduction (SCR) of NO_x using ammonia (NH₃) provided through thermolysis and hydrolysis of urea-water solution (typically employing V₂O₅-WO₃-TiO₂ or Cu and Fe incorporated into zeolites as catalysts), the selective catalytic reduction of NO_x using hydrogen (H₂-SCR) gains increasing

attention.^{3–5} The inherent advantage of H₂-SCR lies in the direct utilization of hydrogen fuel as the reductant. Nevertheless, two major challenges impede its widespread adoption: the relatively low selectivity of the H₂-SCR reaction, which can lead to the formation of nitrous oxide (N₂O), a potent greenhouse gas, as well as toxic ammonia, and diminished catalytic activity at elevated temperatures due to the preferential oxidation of hydrogen by oxygen (O₂).

Catalyst formulation plays a critical role in modulating both the activity and selectivity of the H₂-SCR process. Among the elements that gained particular attention in recent years for their outstanding activity are platinum (Pt),⁶ palladium (Pd),⁷ and iridium (Ir).^{8,9} A comparative study by Shao *et al.*⁹ demonstrated that these metals exhibit optimal performance across distinct temperature regimes: Pt at low temperatures, Pd at intermediate temperatures, and Ir at high temperatures. Also, rhodium (Rh) has been tested in the past.¹⁰ Among all these elements, Pd is particularly attractive due to its high activity in the medium-temperature range and its comparatively superior selectivity, especially when contrasted with the highly active but less selective Pt.¹¹ These attributes position Pd as a particularly

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°C with 2 K min⁻¹. Furthermore, prior to the measurement of each profile, the catalyst was subjected to the oxidation treatment described above. In contrast to the powder measurement and in order to prevent the condensation of water inside the capillary, SpaciPro measurements in the presence of steam were carried out using only 5 vol% H₂O instead of 10 vol% H₂O.

Results and discussion

Catalyst characterization

Compared to a TiO₂-free 1%Pd/HY catalyst from an earlier study that exhibited a BET surface area of 510 m² g⁻¹,³⁶ the surface areas of the catalysts studied herein (Table 1) range from 171 m² g⁻¹ (1%Pd/V₂O₅/TiO₂/HY) to 445 m² g⁻¹ (1%Pt/WO₃/TiO₂/HY) and are thus generally reduced. While this can mainly be attributed to blockage of zeolitic pores by the precipitated TiO₂ layer, the surface area variations throughout the series of TiO₂-containing catalyst samples are likely linked to subtle differences in each individual synthesis. This is evident from the correlation between the measured acidity and the specific surface area (*cf.* Table 1). The observed linear relationship between acidity and BET surface area (Fig. S7) indicates progressive blockage of the zeolite pores during incipient wetness impregnation of the promoters and the noble metal. The Fe-BEA catalyst, on the other hand, exhibits a very high surface area of 666 m² g⁻¹ after the ion exchange.

For the sake of clarity, the nominal compositions are used throughout this study to refer to the investigated catalyst samples. However, the precise contents of noble metals and promoters are detailed in Table 1. Although elemental analysis indicates slightly elevated noble metal concentrations compared to those expected based on the synthesis protocol, the relative trends across different loading levels remain consistent. This discrepancy may be attributed to the hygroscopic nature of the zeolite support, which could have influenced the accuracy of mass measurements during sample preparation.

XRD analysis (*cf.* Fig. S1–S3) reveals diffraction peaks exclusively associated with TiO₂ and Zeolite Y, suggesting a high degree of dispersion of the active metals, Pd and Fe, within their respective zeolitic frameworks. Notably, no diffraction signals corresponding to the promoter oxides

WO₃ and V₂O₅ were observed. While this absence may also reflect a fine dispersion of these oxides, the relatively high loading levels (>5 wt%) suggest that these species may be present in an amorphous phase rather than in a crystalline form detectable by XRD.

Influence of noble metal loading

TiO₂/HY-supported catalysts with different palladium loading (0.1 wt%, 0.5 wt%, 0.75 wt%, and 1 wt%) were evaluated for their performance in a water-free (1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid (1000 ppm NO, 5000 ppm H₂, 10 vol% O₂, 10 vol% H₂O in N₂) gas mixture. As underscored by the kinetic data depicted in Fig. 1, increasing the Pd-loading from 0.1 wt% to 0.5 wt% and 0.75 wt% shifts the NO light-off toward lower temperature, both in dry and humid conditions. While the 0.75 wt% Pd sample exhibits the highest NO_x conversion, the sample with a Pd loading of 1 wt% shows lower activity, at least for temperatures above 160 °C, despite the higher noble metal content. Taking the characterization data from Table 1 into account, our findings suggest an optimum palladium loading that is influenced by two key properties: the dispersion of the active species and the surface area of the catalyst. In general, an increasing noble metal loading typically results in larger particles, but only if the surface area is constant. Both 0.75%Pd/TiO₂/HY and 1%Pd/TiO₂/HY were found to have a dispersion ≤5%, corresponding to a mean Pd particle size of >22 nm. However, among the samples tested herein, 1%Pd/TiO₂/HY exhibits the lowest specific surface area (264 m² g⁻¹), whereas the most active 0.75 wt% Pd sample has a specific surface area of 339 m² g⁻¹. Since the NO reduction efficiency of Pd strongly correlates with the Pd particle size and since the noble metal-support interactions, which are commonly characterized by the interface of Pd and the support, strongly govern catalytic behavior,^{44,45} we can assume a complex interplay of microscopic properties accounting for the performance maximum of the 0.75 wt% Pd catalyst. Moreover, the size of carbon-supported Pd particles was reported to determine the efficiency of nitrite reduction in aqueous environment both in terms of activity and product selectivity.⁴⁶ This is of particular importance in the context of

Table 1 BET surface area and pore volume obtained from N₂ physisorption measurements, Pd dispersion and Pd particle size obtained from CO-chemisorption measurements, and elemental compositions of the catalyst formulations determined by ICP-OES. Information on standard deviations can be found in the SI

Catalyst formulation	A _{BET} [m ² g ⁻¹]	V _{pore} [cm ³ g ⁻¹]	Acidity [μmol g ⁻¹]	Dispersion [%]	d _{particle} [nm]	Active metal [wt%]	Promoter [wt%]	TiO ₂ [wt%]	SiO ₂ [wt%]	Al ₂ O ₃ [wt%]
1%Pd/TiO ₂ /HY	264	0.16	320	5.0	22.4	1.25	—	23.6	55.2	19.9
0.75%Pd/TiO ₂ /HY	339	0.19	560	3.3	33.7	0.94	—	27.2	52.8	19.0
0.5%Pd/TiO ₂ /HY	417	0.19	560	12.6	8.8	0.62	—	27.1	53.0	19.3
0.1%Pd/TiO ₂ /HY	310	0.19	480	10.9	10.2	0.134	—	28.0	51.6	20.3
1%Pd/WO ₃ /TiO ₂ /HY	427	0.24	580	9.6	11.5	1.30	5.61	27.1	47.6	18.4
1%Pd/V ₂ O ₅ /TiO ₂ /HY	171	0.13	290	3.4	32.7	1.16	5.65	26.6	48.2	18.4
Fe-BEA	666	0.33	470	—	—	3.73	—	—	92.1	4.2

Active metal is either Pd or Fe; promoter is either WO₃ or V₂O₅.





Fig. 1 NO and H₂ conversion (solid lines) and product selectivities (dotted lines) during the light-off measurements of the $x\%Pd/TiO_2/HY$ catalysts with $x = 0.1$ (a and b), 0.5 (c and d), 0.75 (e and f), and 1 (g and h) in the dry gas mixture (a, c, e and g; 1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid gas mixture (b, d, f and h; +10 vol% H₂O); GHSV = 60 000 h⁻¹.

H₂-SCR, because species such as nitrates, nitrites, and nitro-compounds were reported to be formed and stored especially

at the interface of PdO and TiO₂ under H₂-SCR conditions.⁴⁷ Not only the NO conversion, but also the dry (Fig. 1a, c, e and g)





Fig. 2 Turnover frequency (TOF) for H₂ conversion (solid lines, left y-axis) and NO conversion (dotted lines, right y-axis) during the light-off measurements of the Pd/TiO₂/HY catalysts with different Pd loading in the dry gas mixture (1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂); GHSV = 60 000 h⁻¹.

0.1%Pd/TiO₂/HY, notably, at temperatures exceeding 200 °C. Despite a slightly higher Pd dispersion of 12.6% and an even greater specific surface area, the 0.5%Pd/TiO₂/HY sample exhibits much lower TOF values and is outperformed by any other sample tested herein in terms of atom efficiency. Consistent with the activity data presented in Fig. 1, 0.75%Pd/TiO₂/HY exhibits the highest TOF values overall. The maximum TOF for NO conversion of 0.7 s⁻¹ is found at approx. 195 °C, which coincides with the peak absolute NO conversion of about 90% (Fig. 1e). For H₂ activation, TOF values of approx. 4 s⁻¹ are found once complete H₂ conversion is reached, with a plateau beyond 195 °C. Conversely, despite its higher noble metal loading, generally lower TOF values are observed for 1%Pd/TiO₂/HY, and the maximum TOF for NO (~0.3 s⁻¹) is found at around 213 °C, matching the maximum NO conversion of 77% (Fig. 1g). Given that CO chemisorption data indicate similar Pd dispersion for 1%Pd/TiO₂/HY (5%) and 0.75%Pd/TiO₂/HY (3.3%), other physicochemical parameters such as specific surface area (264 m² g⁻¹ for 1%Pd/TiO₂/HY versus 339 m² g⁻¹ for 0.75%Pd/TiO₂/HY) likely account for the observed

differences in TOF data. As elaborated later, aspects like surface acidity play a critical role in H₂-SCR. Since the differences in surface area correlate with the number of acid surface sites available for adsorption of intermediate species, the surface area can impact the overall catalytic performance.

Addition of promoter materials

Since previous research pointed to a performance improvement if H₂-SCR catalyst formulations were modified with promoter materials such as WO₃ or V₂O₅, their addition to Pd/TiO₂/HY was evaluated in a next step. Compared to 1%Pd/TiO₂/HY (Fig. 1g), the addition of WO₃ results in a clear increase in activity in both the low- and high-temperature regime for the dry mixture (Fig. 3a). In particular, the T₅₀ for hydrogen conversion is shifted toward lower temperatures, namely from 146 °C to 129 °C, and also the low-temperature NO conversion peak is shifted from 149 °C to 136 °C. Between 175 °C and 300 °C – the high-temperature regime – the activity for NO reduction is consistently higher as well. Only in the narrow temperature window between the two peaks (around 160 °C) the NO conversion falls below the levels of the unpromoted Pd catalyst by a small margin. The overall activity boost is assumed to originate from the electronic interaction between WO₃ and the noble metal; in analogy to what has been reported for WO₃-promoted Pt/TiO₂ H₂-SCR catalysts,⁴⁹ we can speculate about a higher share of metallic palladium sites (Pd⁰) upon the introduction of WO₃. This is of high relevance because reduced palladium has been observed to be more active for the H₂-SCR reaction than palladium oxide for various Pd-based catalyst formulations.^{27,41,50} With regard to product selectivity, the WO₃-promoted catalyst shows negligible ammonia formation and a slightly higher NO oxidation than the unpromoted one during the transition between the two NO conversion maxima around 160 °C. Nevertheless, N₂ and N₂O selectivities are overall comparable.

The addition of water to the reaction gas mixture does not alter these general trends significantly, although NO reduction efficiency below 150 °C is reduced to relevant extent in humid conditions (Fig. 3b). In contrast to the unpromoted 1%Pd/TiO₂/



Fig. 3 NO and H₂ conversion (solid lines) and product selectivities (dotted lines) during the light-off measurements of the 1%Pd/WO₃/TiO₂/HY catalyst in the dry gas mixture (a, 1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid gas mixture (b, +10 vol% H₂O); GHSV = 60 000 h⁻¹.



HY (Fig. 1h), NH_3 formation was found irrelevant over 1%Pd/ $\text{WO}_3/\text{TiO}_2/\text{HY}$ (Fig. 3b) and the presence of H_2O did not reduce the selectivity to the undesired product N_2O , but increased it even slightly for temperatures of 190 °C and above. Thus, despite promising performance of 1%Pd/ $\text{WO}_3/\text{TiO}_2/\text{HY}$ in dry environment and higher NO_x conversion than its unpromoted counterpart once temperatures exceed 180 °C, the fact that in real-world applications humidity will be omnipresent calls for further optimization of the catalyst formulation in order to suppress the generation of the strong greenhouse gas N_2O .

To address the need for enhanced selectivity in NO_x reduction, vanadium oxide (V_2O_5 , VO_x) was tested as an alternative promoter, since it has been reported to facilitate the formation of NH_4^+ intermediates that presumably promote selective catalytic reduction pathways.^{32–34} Although the performance data shown in Fig. 4 show that the selectivity of 1%Pd/ $\text{V}_2\text{O}_5/\text{TiO}_2/\text{HY}$ toward the target product N_2 can compete with that of the unpromoted 1%Pd/ TiO_2/HY (Fig. 1g and h) sample both in dry and humid conditions, the addition of V_2O_5 results in a dramatic drop in catalytic activity especially in the presence of water (Fig. 4b). Considering the very low specific surface area ($171 \text{ m}^2 \text{ g}^{-1}$) of the vanadia-containing catalyst sample, we can speculate about a blockage of zeolitic pores and thus a reduction of acidic surface sites due to the addition of non-zeolitic material.³⁶ The NH_3 -TPD-derived acidity data (cf. Table 1) support this hypothesis. Considering that NO reduction over noble metals such as Pt and Pd has been claimed to follow an NH_4^+ -mediated reaction mechanism,^{20,33,35} a lower number of Brønsted acid sites suppresses the formation of NH_4^+ -species that could then react with NO_x to form N_2 , hereby impeding efficient NO_x reduction. Taking the low dispersion of 3.4% (cf. Table 1) into account, we can also speculate about a partial blockage of highly active Pd^{2+} sites. Such Pd^{2+} ions can form as highly dispersed species in the sodalite cages of HY materials⁵¹ and are believed to contribute to the overall NO_x conversion during H_2 -SCR as well as to preserve high activity under humid conditions.³⁶ In conclusion, while we acknowledge the beneficial contribution of V_2O_5 to the N_2 selectivity, a simple deposition of vanadia onto high surface area supports as conducted herein has proven insufficient for H_2 -SCR catalysts.

However, advanced preparation techniques that allow for a more precise control of vanadia distribution on structured support materials may result in better performance.³⁴

Combining H_2 -SCR and NH_3 -SCR catalyst formulations

Based on the data discussed thus far, the formation of the strong greenhouse gas N_2O constitutes a significant limitation in the implementation of H_2 -SCR for NO_x abatement. To mitigate N_2O emissions during NO_x reduction *via* hydrogen, the integration of a secondary catalytic system capable of N_2O decomposition presents a viable strategy. Among the catalyst formulations reported in the literature, ion-exchanged iron zeolites – in particular Fe/ZSM-5 and Fe-BEA – have demonstrated relatively low activation temperatures for the N_2O decomposition reaction.^{52,53} In addition, Fe-BEA was reported to catalyze NO reduction and N_2O decomposition simultaneously,^{54,55} making it particularly promising for combination with H_2 -SCR catalyst formulations prone to N_2O formation. To reconcile these findings with the need for hydrothermal stability under SCR conditions typical for H_2 -ICE exhausts with high levels of humidity, Fe-BEA was selected as the more robust catalyst formulation.⁵⁶ This choice aims to minimize the N_2O levels while maintaining reasonable structural integrity under highly humid operational conditions. However, given that effective N_2O conversion requires temperatures exceeding 200 °C, the Pd-based H_2 -SCR catalyst and the Fe-BEA catalyst were combined in a mechanical mixture rather than arranged sequentially. This configuration enables direct utilization of local temperature increases resulting from the exothermic total oxidation of hydrogen in the presence of oxygen. Throughout the tests discussed in the following, the catalyst bed was composed of 300 mg of Pd-based H_2 -SCR catalyst, 300 mg of Fe-BEA catalyst, and 400 mg of SiO_2 as a diluent (reduced from the previously used 700 mg) to maintain a constant total mass of 1000 mg. Note that 1%Pd/ TiO_2/HY was chosen over 0.75%Pd/ TiO_2/HY for its superior water tolerance, which is particularly critical when evaluating the performance of the material combination under realistic humid conditions.



Fig. 4 NO and H_2 conversion (solid lines) and product selectivities (dotted lines) during the light-off measurements of the 1%Pd/ $\text{V}_2\text{O}_5/\text{TiO}_2/\text{HY}$ catalyst in the dry gas mixture (a, 1000 ppm NO , 5000 ppm H_2 , 10 vol% O_2 in N_2) and humid gas mixture (b, +10 vol% H_2O); GHSV = 60 000 h^{-1} .



The data depicted in Fig. 5a indicate that the combination of 1%Pd/TiO₂/HY and Fe-BEA in a 1:1 mass ratio exhibits substantial NO conversion in the dry gas mixture at temperatures exceeding 180 °C, reaching a peak conversion of 92% at 213 °C. The hydrogen light-off for this catalyst combination is slightly shifted to higher temperatures, with a T_{50} of 153 °C, compared to 146 °C observed for the H₂-SCR catalyst alone. Overall, a markedly lower N₂O selectivity, and consequently enhanced N₂ selectivity, was observed for the combined catalyst system. However, at temperatures exceeding 250 °C, an increase in NO oxidation activity appears to occur, leading to a decline in N₂ selectivity due to the formation of NO₂. While this effect is particularly pronounced in dry reaction environment (Fig. 5a), humidity mitigates NO₂ evolution (Fig. 5b). In the presence of water vapor, N₂ selectivity remains above 90% across the high-activity temperature range of 180–300 °C. Despite a slight overall shift of NO conversion toward higher temperatures in the presence of water vapor, which we attribute to competitive adsorption of H₂O and surface hydroxyl groups on the active sites, the high selectivity toward N₂ remains particularly encouraging. This observation is of practical relevance, as real-world exhaust gas streams typically contain substantial levels of water vapor. Therefore, the robustness of N₂ selectivity under humid conditions underscores the potential applicability of the catalyst system in realistic operating environments.

It is also noteworthy that no ammonia was detected in the effluent gas stream when employing the combined Pd-based H₂-SCR and Fe-BEA catalyst system. This observation may be attributed to two factors. First, iron-exchanged zeolites are well-established NH₃-SCR catalysts and may facilitate *in situ* consumption of NH₃ generated within the catalyst bed for NO reduction.^{57–59} Second, NH₃ can react with N₂O in the presence of iron zeolites,⁶⁰ hereby potentially representing another relevant reaction pathway, particularly at elevated temperatures.

Due to the significantly lower gravimetric density of Fe-BEA compared to the SiO₂ diluent, the catalyst bed length increased in the mixed catalyst configuration used in the aforementioned

experiments. To eliminate potential effects arising from increased residence time, a second sample was prepared consisting of 300 mg of Pd-based catalyst and 215 mg of Fe-BEA, without any diluent. This configuration maintained the same bed length as that used in measurements with the H₂-SCR catalyst alone. The corresponding results, presented in the SI (Fig. S4), confirm the previously described trends. Notably, in contrast to the diluted mixture, hydrogen conversion in the undiluted system is shifted toward lower temperatures (Fig. S4), which may be attributed to elevated intra-catalyst temperatures resulting from reduced thermal dissipation in the absence of the diluent.

Mechanistic considerations derived from experiments with spatial resolution

Although the Pd-based catalyst likely continues to generate relevant quantities of ammonia or ammonium intermediates during the NO reduction process, no NH₃ was detected at the outlet of the catalytic bed comprising both Pd- and Fe-based components. This absence is presumably due to the rapid consumption of NH₃ *via* reactions with NO, NO₂, or N₂O over the Fe-BEA catalyst. To elucidate the complex interplay of reactions occurring within this bifunctional catalytic system and to identify the specific reactions involved, spatially resolved concentration measurements were performed. These measurements were conducted on monolithic samples coated either with 1%Pd/TiO₂/HY alone or with a combination of 1%Pd/TiO₂/HY and Fe-BEA in a 2:1 mass ratio. Note that this mass ratio differs from that of the powder catalyst tests; due to the above-mentioned low gravimetric density of Fe-BEA, a reduction of its content was necessary in order to achieve a reasonable washcoat thickness (approx. 35 μm) that keeps the relevance of internal diffusion low. To ensure catalyst stability during spatially resolved steady-state measurements, both the Pd-based and Fe-based catalysts were subjected to a 12-hour long-term test at 200 °C and a gas hourly space velocity (GHSV) of 75 000 h⁻¹ under both dry and humid conditions.

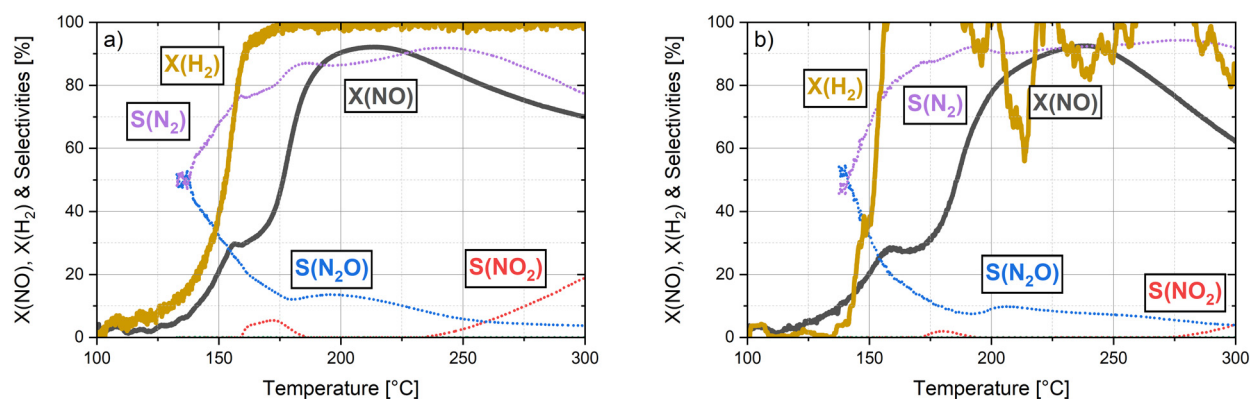


Fig. 5 NO and H₂ conversion (solid) and product selectivities (dotted lines) during the light-off measurements of the combination of 1%Pd/TiO₂/HY and Fe-BEA (1:1, m:m) catalysts in the dry gas mixture (a, 1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid gas mixture (b, +10 vol% H₂O); GHSV = 60 000 h⁻¹.



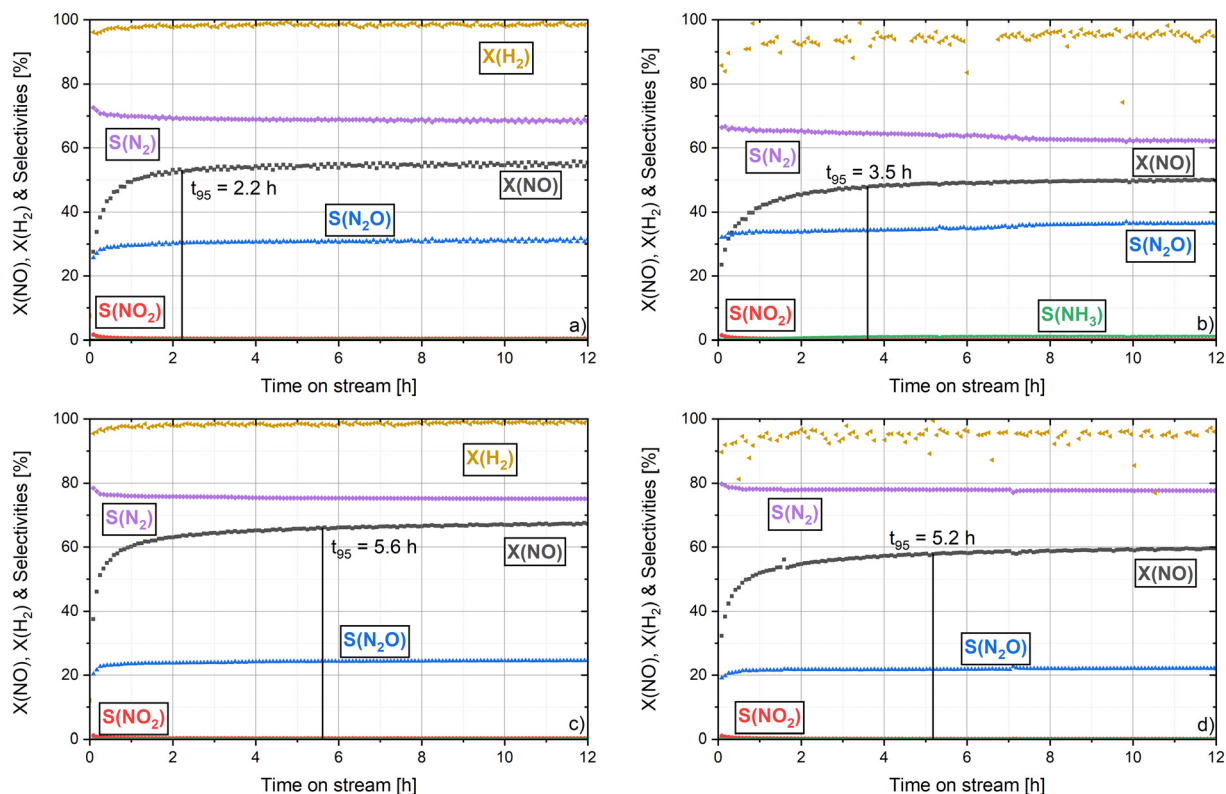


Fig. 6 Long-term trends of NO, H₂ conversion, and product selectivities of 1%Pd/TiO₂/HY (a and b) and 1%Pd/TiO₂/HY + Fe-BEA (2:1, m:m) (c and d) in the dry gas mixture (a and c: 1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid gas mixture (b and d: +5 vol% H₂O) at a GHSV = 75 000 h⁻¹ and T = 200 °C.

As shown in Fig. 6, all monolithic samples exhibited lower NO conversion under steady-state conditions compared to the light-off measurements conducted with powder samples. This observation is consistent with the higher GHSV employed in the long-term and spatially resolved experiments with monolithic samples, which results in reduced contact time between the gas and the catalyst surface. Also, the different mass ratio of 1%Pd/TiO₂/HY and Fe-BEA is expected to contribute to such variations. Furthermore, it is plausible that despite the low temperature ramp rate of 2 K min⁻¹ as chosen herein, light-off measurements may not accurately reflect true steady-state performance at a given temperature. This discrepancy may arise from the catalytic system's relaxation time exceeding the characteristic time of the temperature ramp, thereby preventing full equilibration before data acquisition. More importantly, the catalysts demonstrated stable performance during the 12-hour exposure to the reaction gas mixture, with no observable loss in activity. In fact, NO conversion increased over time. For the Pd-based H₂-SCR catalyst alone, 95% of the steady-state NO conversion (*t*₉₅) – the steady-state NO conversion is defined as the absolute conversion achieved after 12 h – was reached after 2.2 h under dry conditions (Fig. 6a) and 3.5 h under humid conditions (Fig. 6b). In contrast, the physical mixture of the Pd catalyst and Fe-BEA required 5.6 h and 5.2 h to reach *t*₉₅ under dry (Fig. 6c) and humid conditions (Fig. 6d), respectively.

Despite the differing space velocities employed in the powder and monolith experiments, which account for the lower NO conversion during spatial profiling, the observed trends remained consistent. Specifically, water vapor exposure led to a reduction in NO conversion and an increase in N₂O selectivity for the Pd-only catalyst (*cf.* Fig. 1g and h and 6a and b), whereas the Pd/Fe mixed catalyst exhibited a decrease in N₂O selectivity under the same conditions (*cf.* Fig. 5 and 6a and b). Additionally, NH₃ formation was detected in the effluent of the pure Pd catalyst under humid conditions (*cf.* Fig. 1h and 6b), while no ammonia was observed in the end-of-pipe measurements for both the powder and the monolithic 1%Pd/TiO₂/HY + Fe-BEA (2:1, m:m) catalyst under either dry or humid conditions (*cf.* Fig. 5 and 6c and d).

Fig. 7 presents the spatially resolved concentration profiles of NO, H₂, and reaction products along the monolith channel at temperatures of 150 °C, 200 °C, and 250 °C, at a GHSV of 75 000 h⁻¹. In analogy to the light-off tests discussed above, measurements were conducted under both dry (Fig. 7a–c) and humid (Fig. 7d–f) conditions using a monolithic 1%Pd/TiO₂/HY catalyst. At 150 °C, the NO reduction activity is minimal under both dry and humid conditions, with NO conversion not exceeding 10%. The predominant reaction products are N₂O and N₂, with some NO₂ detected as well, consistent with data at this temperature from light-off experiments. It is important to note that the quantification of H₂ was indirectly inferred by balancing with H₂O measurements obtained *via* FTIR





Fig. 7 NO, H₂, and product concentration profiles of the 1%Pd/TiO₂/HY monolithic catalyst in the dry gas mixture (a–c: 1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid gas mixture (d–f: +5 vol% H₂O) at $T = 150\text{ °C}$ (a and d), $T = 200\text{ °C}$ (b and e), and $T = 250\text{ °C}$ (c and f); GHSV = 75 000 h⁻¹.

spectroscopy. This approach introduced significant uncertainty due to the low signal-to-noise ratio of the H₂O signal. Furthermore, the additional dilution with N₂ during spatially resolved measurements exacerbated the signal degradation across all species. At 200 °C, the NO conversion reaches its maximum, in agreement with light-off data. The small amount of NO₂ observed at the catalyst inlet (~16 ppm) is fully consumed. In addition to the formation of N₂O and N₂, NH₃ is detected in the early segment of the catalytic channel. For example, under dry conditions, a peak NH₃ concentration of 22 ppm is observed approximately 3 mm downstream from the channel inlet (Fig. 7b), followed by its subsequent consumption along the channel length. At 250 °C, NO conversion declines due to competitive reduction reactions involving O₂ and NO

with H₂. Nevertheless, NH₃ formation increases, with the peak concentration rising to 32 ppm and shifting closer to the channel inlet (Fig. 7c), indicating enhanced NH₃ formation at elevated temperatures. Due to limitations in spatial resolution, the true peak NH₃ concentration cannot be precisely determined. However, while the presence of water vapor does not appear to significantly alter the position or magnitude of the NH₃ peak, H₂O seems to inhibit NH₃ consumption, resulting in a minor NH₃ slip at both 200 °C and 250 °C under humid conditions.

Fig. 8 illustrates the spatially resolved concentration profiles for the bifunctional catalyst system 1%Pd/TiO₂/HY + Fe-BEA (2:1, m:m) under identical experimental conditions. At 150 °C, both dry and humid environments yield approximately 10% NO



Fig. 8 NO, H₂, and product concentration profiles of the 1%Pd/TiO₂/HY + Fe-BEA (2:1, m:m) monolithic catalyst in the dry gas mixture (a–c: 1000 ppm NO, 5000 ppm H₂, 10 vol% O₂ in N₂) and humid gas mixture (d–f: +5 vol% H₂O) at $T = 150\text{ °C}$ (a and d), $T = 200\text{ °C}$ (b and e), and $T = 250\text{ °C}$ (c and f); GHSV = 75 000 h⁻¹.



conversion, similar to the monofunctional 1%Pd/TiO₂/HY catalyst. The primary reaction products are N₂ and N₂O, while NO₂ concentrations exhibit a slight decline; however, this change remains within the margin of experimental error. At 200 °C, the NO conversion reaches its peak both in dry and humid conditions, with clearly enhanced N₂ formation compared to the monofunctional catalyst (*cf.* Fig. 7b and e and 8b and e). This improvement in both activity and selectivity is attributed to the incorporation of Fe-BEA. Zeolite-incorporated iron is known to be active for selective NO reduction in the context of conventional NH₃-SCR, where not H₂ but NH₃ serves as reducing agent,⁶¹ and Fe-BEA in particular was found feasible for catalyzing simultaneous NO and N₂O reduction by NH₃.^{54,55} Notably, compared to the monofunctional catalyst, a significantly higher NH₃ peak concentration of 73 ppm is observed over the bifunctional catalyst near the channel inlet (Fig. 8b). This increase is likely due to the presence of the zeolitic BEA support, which introduces a greater density of Brønsted acid sites and thereby facilitates the formation of NH₄⁺ intermediates.⁶² Their decomposition into NH₃ and subsequent desorption may account for the increased NH₃ levels. Moreover, the direct interaction of Fe and Pd-based catalysts within the washcoat may enhance desired reaction pathways. Dong *et al.*⁶³ investigated Pt/SSZ-13 formulations for their NO reduction efficiency when using H₂ as reductant and concluded that two reaction pathways occur in H₂-SCR conditions: (i) the formation of NH_x⁺ species, in particular NH₄⁺, and their subsequent oxidation with high selectivity towards the desired reaction product N₂; (ii) the catalytic reduction of NO by NH₃ and thus conventional NH₃-SCR-like reactions taking place on noble metal-based materials, yet with comparably poor selectivity and thus formation of high N₂O levels. In this regard, the promotion of NH₄⁺ intermediate formation as well as the NH₃-SCR reaction taking place over Fe with higher selectivity than over noble metal particles could explain the overall better performance of the sample washcoated with the material mixture compared to the sample coated with Pd/TiO₂/HY only. This hypothesis is substantiated by the spatially resolved measurements, which uncover that the

bifunctional catalyst formulation effectively converts the elevated NH₃ concentrations formed at the inlet, which we attribute to NH₃-SCR reaction pathways taking place over Fe sites and hereby promote the overall NO_x reduction efficiency. Similar to the trends observed for the monofunctional catalyst, the NH₃ concentration peak shifts upstream toward the channel entrance at 250 °C, indicating temperature-dependent NH₃ formation dynamics. In contrast to 1%Pd/TiO₂/HY, the addition of water does not appear to impair the NH₃ conversion efficiency of the bifunctional catalyst (Fig. 8e and f). This suggests that the addition of the Fe-BEA component improves robustness in the presence of water, likely due to enabling NH₃-SCR reaction pathways. Water vapor can promote these through a hydroxyl-induced transformation of Lewis acid sites into Brønsted acid sites that enable higher surface coverage of important intermediates, in particular NO₃⁻/NO₂⁻ and NH₄⁺.⁶⁴ Consistently, light-off experiments of the monolithic bifunctional catalyst system performed at water vapor concentrations representative of real hydrogen internal combustion engine exhaust (0–20 vol% H₂O)⁶⁵ reveal only a minor effect of increasing humidity levels on catalyst performance (*cf.* Fig. S8) and thus underscore the potential of the bifunctional catalyst proposed herein for efficient deNO_x in H₂-ICE applications (Fig. 9).

Conclusions

This study systematically evaluated Pd-based catalysts for H₂-SCR of NO_x under conditions representative of H₂-ICE exhaust. The key aspects identified in this work to govern catalyst performance are summarized in schematic Fig. 9. Catalysts supported on TiO₂-modified HY zeolite showed that decreasing Pd loading raised NO and H₂ light-off temperatures, indicating diminished low-temperature activity. Kinetic analysis indicates that NO conversion at low temperatures is limited by H₂ activation, whereas competitive H₂ oxidation dominates at higher temperatures, hereby hampering NO reduction. Water vapor suppressed NO conversion across all formulations. Pd dispersion strongly influenced product selectivity: lower Pd



Fig. 9 Illustrative summary of key factors driving catalyst performance and their mechanistic implications.



loadings promoted NH_3 formation above 200 °C, while N_2O remained a persistent byproduct regardless of the noble metal loading. Variations in surface area may also affect performance, likely through changes in surface site density of acidic sites. While all HY-supported catalysts were synthesized *via* simple TiO_2 precipitation and Pd impregnation for the present study, the results suggest that more advanced synthesis methods are needed in the future to better control support morphology and metal particle size for clearer structure–activity correlations.

Composite catalysts were also explored to broaden the effective temperature window. Adding V_2O_5 improved N_2 selectivity but reduced NO conversion, particularly under humid conditions. In contrast, WO_3 incorporation enhanced low-temperature (100–175 °C) NO conversion and minimized NH_3 formation, though N_2O production above 200 °C remained problematic.

To mitigate byproduct formation more efficiently, a bifunctional system combining Pd/ TiO_2 /HY with Fe-BEA was developed. This composite catalyst improved NO conversion under both dry and humid conditions and significantly reduced NH_3 and N_2O formation, achieving >90% N_2 selectivity between 180 and 300 °C in realistic humid conditions (powder catalyst testing). Monolith testing confirmed stable performance over 12 hours. Spatially resolved measurements showed NH_3 formation near the inlet, attributable to BEA Brønsted acid sites, and downstream consumption *via* Fe-mediated NH_3 -SCR pathways, collectively contributing to lower N_2O formation. The relative contributions of these pathways to the overall performance, however, require further clarification.

Future work should employ *operando* spectroscopic methods to – preferentially simultaneously – elucidate electronic and mechanistic features governing bifunctional catalyst behavior. The strong synergies observed in the Pd/ TiO_2 -HY + Fe-BEA system highlight its promise for advancing H_2 -SCR toward practical H_2 -ICE emission control applications.

Author contributions

Michael Borchers: conceptualization, data curation, formal analysis, investigation, validation, visualization, writing – original draft. Daniel Hodonj: data curation, formal analysis, investigation, methodology, validation, visualization, writing – original draft. Kathrin Schäfer: formal analysis, investigation. Frank Manuel Bauer: formal analysis, investigation. Patrick Lott: conceptualization, data curation, funding acquisition, methodology, project administration, resources, supervision, validation, visualization, writing – original draft.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the supplementary information (SI).

Supplementary information is available. See DOI: <https://doi.org/10.1039/d5cy01387a>.

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