Dual catalyst system for selective vinyl chloride production via ethene oxychlorination†

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A dual system comprising two catalytic reactors connected in series was developed for the direct conversion of ethene to vinyl chloride monomer (VCM). The first reactor uses ZrO2-supported ceria (CeO2/ZrO2) to perform ethene oxychlorination to 1,2-dichloroethene (EDC) that is dehydrochlorinated to VCM in the second reactor over calcium-promoted γ-Al2O3. The choice of carrier for ceria is of critical importance to maximize the EDC production by reducing combustion products. While MgO, SiO2, SiC, TiO2, ZSM-5, and γ-Al2O3 carriers induced higher overoxidation compared to bulk ceria, ZrO2 was the only carrier that suppressed CO2 formation. Moreover, the latter carrier led to the highest oxychlorination activity. The unique performance of the CeO2/ZrO2 catalyst was rationalized by its ability to promote chlorine evolution and to suppress the combustion of chlorinated products, as inferred from the activity evaluation in HCl and VCM oxidation, respectively. The outstanding redox properties, enabling operation at low temperature and thus high selectivity, are associated with the formation of defective CeO2 nanoparticles, contrasting the low activity over Ce–Zr mixed oxide. In order to subsequently form VCM, an efficient EDC dehydrochlorination catalyst was designed by moderating the acidity of γ-Al2O3 via calcium doping and used in a reactor after CeO2/ZrO2. This dual catalyst system displayed 100% selectivity to VCM at 25% ethene conversion, surpassing the space time yield of the best ethene-to-VCM catalyst EuOCl by a factor of four, where the first step is operated at an elevated temperature of about 100–150 K with respect to cupric chloride benchmarks. In addition, the catalytic dehydrochlorination, operated at a lower temperature of 100 K than the current non-catalytic process, showed minimized coke formation. This developed system rendered stable after slight initial deactivation, offering promising potential to intensify VCM production.

Introduction

Ethene oxychlorination, i.e., the reaction of a former alkene with HCl and oxygen, constitutes an important step of the so-called balanced technology for the production of vinyl chloride monomer (VCM) that is the key precursor in the manufacture of polyvinyl chloride (PVC).1,2 Since its commercialization in the late 1950s,1–3 the oxychlorination process has been industrially practiced over catalysts comprising an active cupric chloride (CuCl2) phase supported on γ-alumina carriers and promoted by (oxy)chlorides of alkali and rare-earth metals as these materials provide high activity and selectivity to 1,2-dichloroethene (EDC, C2H4Cl2),1,2,4–11 which is subsequently dehydrochlorinated to VCM in a separate non-catalytic step.1,2 Nonetheless, cupric chloride species display an inherently low melting point.1,2,6 In combination with the high exothermicity of the oxychlorination reaction, this factor can cause particle stickiness in fluidized bed reactors, or gradual loss of the active phase due to volatilization, linked to hot-spot formation in fixed-bed reactors. Efforts to remediate both these effects by the addition of promoters were only partially effective. This limited stability of CuCl2-based catalysts triggered a search for alternative active phases that could overcome this challenge while still exhibiting exceptional activity and selectivity for ethene oxychlorination. The quest was additionally motivated by the recent progress in the development of stable non-copper catalytic systems involving RuO2- and CeO2-based catalysts that are able to preserve the structural integrity under the similarly harsh reaction environment of HCl oxidation (Deacon reaction).12,13 Notably, the evaluation of bulk RuO2 and CeO2 in ethene...
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dehydrochlorination catalysts exhibit limited VCM selectivity at reactions in a single catalytic system. Nonetheless, existing integration of oxychlorination and dehydrochlorination success.

magnitude higher costs of europium compared to those of benchmark CuCl₂- oxychlorination. In the first step, CeO₂ was nanostructured for highly selective direct VCM production systems for highly selective direct VCM production via ethene oxychlorination. As demonstrated in previous studies, carriers can modify the redox properties of CeO₂ via geometric and electronic effects, thus offering the possibility to curb combustion. In addition, similar to the beneficial effects that ceria provides when used as a carrier, appropriate carriers for CeO₂ could promote the EDC-to-VCM conversion step by introducing acid functionalities that are required for this reaction. In fact, as reported in the literature, several acid materials, such as Al₂O₃ or Pt-Cu/SiO₂, are active in EDC dehydrochlorination. If applied as active carriers or co-catalysts for CeO₂, these materials can allow integration of oxychlorination and dehydrochlorination reactions in a single catalytic system. Nonetheless, existing dehydrochlorination catalysts exhibit limited VCM selectivity at high EDC conversion levels and a prominent propensity for deactivation due to coking.

This study aims at developing CeO₂-based catalytic systems for highly selective direct VCM production via ethene oxychlorination. In the first step, CeO₂ was nanostructured over a broad range of carriers in order to suppress CO₂ formation and to determine the potential of different carriers to promote EDC dehydrohalogenation. Hereby, it is shown that overoxidation can be completely avoided only by supporting CeO₂ over a ZrO₂ carrier that, however, leads to a marginal EDC-to-VCM conversion. To foster the latter step, a calcium doped-γ-Al₂O₃ co-catalyst that exhibits a combination of high activity and unprecedented selectivity to VCM in EDC dehydrochlorination with a low propensity for coking was developed. Finally, a dual-bed reactor system that comprises CeO₂/ZrO₂ and calcium doped-γ-Al₂O₃ catalysts was designed enabling direct ethene to VCM conversion via oxychlorination with absolute selectivity to this polymer precursor, outperforming the best reported catalyst to date, EuOCl, by a factor of four in terms of space time yield.

Experimental

Catalyst preparation

Bulk CeO₂ (Sigma-Aldrich, nanopowder, 99.5%) and carriers, MgO (~20 nm, Strem Chemicals, ≥99%), SiO₂ (20–60 μm, Evonik, AEROPEARL 300/30, ≥99.0%), TiO₂ (≤60 μm, Alfa-Aesar, 99%), Al₂O₃ (≤60 μm, Alfa-Aesar, 99%), γ-Al₂O₃ (5–100 μm, Sasol, PURALOX SCFa 140, ≥98%), ZSM-5-15 (Zeolyst, CBV3024E), ZSM-5-40 (Zeolyst, CBV8014), ZSM-5-1040 (Tosoh, HSZ-690HOA), ZSM-5-1040 (Tosoh, HSZ-890HOA), FER-29 (Zeolchem, CP905), Y-40 (Zeolyst, CBV780), and MOR-110 (Tosoh, HSZ-690HOA), were calcined prior to their use in catalytic tests or in catalyst preparation. Dry impregnation was used as a standard method for catalyst synthesis. Herein, an appropriate amount of Ce(NO₃)₃·6H₂O (Aldrich, 99.9%) was dissolved in a volume of deionized water equal to the carrier pore volume to achieve the desired wt% loading of Ce. The precursor solutions were then added dropwise to the carrier, which was stored under vacuum (50 mbar) at 373 K. The wetted materials were periodically mixed at room temperature for 1 h, then dried and finally calcined. The resulting catalysts are denoted as CeO₂-x/C-SA-T, where x denotes the Ce loading, C the carrier, SA the Si:Al ratio if C is a zeolite, and T the calcination temperature. The standard loading of 10 wt% and calcination temperature (T = 873 K) are omitted for clarity only specified in the case of deviations, where 5 wt% and 20 wt% of Ce are indicated by letters l (low) and h (high), respectively. Promoted alumina catalysts were prepared following the same dry impregnation protocol as described for the preparation of supported CeO₂ catalysts, using Ce(NO₃)₃·6H₂O, KNO₃·6H₂O (Aldrich, 99.9%), La(NO₃)₃·6H₂O (Aldrich, 99.9%), and Ca(NO₃)₂·6H₂O (Aldrich, 99.9%) as precursors. Drying was performed under vacuum (50 mbar) at 373 K for 2 h, while calcination was performed in static air at 873 K for 5 h using a heating rate of 5 K min⁻¹. The resulting catalysts are denoted as M-x/Al₂O₃, where M denotes the metal promoter and x its loading. The standard promoter loading was 2 wt% and is omitted in the catalyst naming, while the reduced promoter loading of 0.5 wt% is indicated by letter l (low).

Catalyst characterisation

Powder X-ray diffraction (XRD) was measured using a PANalytical X’Pert PRO-MPD diffractometer with the Bragg–Brentano geometry by applying Cu Kα radiation (λ = 1.54060 Å). The data were recorded in the 10–70° 2θ range with an
The detailed spectra of the Ce 3d core levels was 46.95 eV to 200 CeO₂ nanoparticles were examined to estimate their average size, using the ImageJ software. Raman spectroscopy was performed on a WITec CRM200 confocal Raman system using a 532 nm laser with a 20 mW power, a 100× objective lens with a numerical aperture NA = 0.9 (Nikon Plan), and a fiber-coupled grating spectrometer (2400 lines per mm), giving a spectral sampling resolution of 0.7 cm⁻¹. N₂ sorption at 77 K was measured using a Micromeritics TriStar analyser. Prior to the measurements, the samples were degassed to 50 mbar at 573 K for 12 h. The Brunauer–Emmett–Teller (BET) method was applied to calculate the total surface area, S_BET in m² g⁻¹. Temperature-programmed desorption of ammonia (NH₃-TPD) and temperature-programmed reduction with hydrogen (H₂-TPR) were performed using a Micromeritics Autochem II 2920 unit equipped with a thermal conductivity detector coupled to a Pfeiffer Vacuum OmniStar mass spectrometer. The powder sample (0.1 g) was loaded into a U-shaped quartz micro-reactor, pre-treated in He (20 cm³ STP min⁻¹) at 573 K for 3 h, and cooled to 373 K in He. For NH₃-TPD experiments, ammonia was chemisorbed at 473 K in three consecutive cycles of saturation with 5 vol.% NH₃/He (20 cm³ STP min⁻¹) for 30 min followed by purging with He (20 cm³ STP min⁻¹) at the same temperature for 30 min. Desorption of NH₃ was monitored in the range of 473–1273 K using a heating rate of 20 K min⁻¹ and a He flow of 20 cm³ STP min⁻¹. For H₂-TPR experiments, the sample was pre-treated in He (20 cm³ STP min⁻¹) at 423 K for 1 h, and cooled to room temperature followed by ramping the temperature at 10 K min⁻¹ up to 1273 K in 5 vol% H₂ (20 cm³ STP min⁻¹). The resulting profiles of NH₃-TPD and H₂-TPR were normalised to the sample weight. Thermogravimetric analysis (TGA) was used to determine the amount of coking was performed on a Linseis STA-PT-1600 apparatus. The sample (30 mg) was loaded on an alumina pan and pretreated in Ar (100 cm³ STP min⁻¹) at 373 K for 1 h, then the temperature was ramped up to 1173 K with a heating rate of 10 K min⁻¹ under a flow of synthetic air (100 cm³ STP min⁻¹). X-ray photoelectron spectroscopy (XPS) measurements were performed on a Physical Electronics Quantum 2000 XPS using monochromatic Al-Kα radiation generated from an electron beam operated at 15 kV, and equipped with a hemispherical capacitor electron-energy analyser. The powdered sample was firmly pressed onto indium foil. The area analysed was 150 μm in diameter and the electron take-off angle was 45°. The pass energy used for the detailed spectra of the Ce 3d core levels was 46.95 eV to yield a total analyser energy resolution of 0.95 eV. The spectrometer energy scale was calibrated for the Au 4f electrons to be at 84.0 ± 0.1 eV. Partial compensation of surface charging during spectra acquisition was obtained by the simultaneous operation of electron and argon ion neutralizers. Extended X-ray absorption fine structure spectroscopy (EXAFS) and X-ray absorption near-edge structure spectroscopy (XANES) measurements were performed at the Ce L3-edge on the XAFCA beamline at the Singapore Synchrotron Light Source (SSLS). The storage ring of the SSLS was operated at 0.7 GeV with a maximum current of 200 mA. The data were collected in transmission mode (to k_max = 10.5) using a Si(111) double crystal monochromator and ion chamber detectors. To ascertain the reproducibility of the experimental data, at least three scan sets were collected. EXAFS and XANES data processing was performed using the Demeter software package (Athena and Artemis). The amplitude reduction factor, S₀₂ was derived from the EXAFS data analysis of the commercial CeO₂ standard.

Catalyst testing

Steady-state oxychlorination of ethene was investigated at ambient pressure in a continuous-flow fixed-bed reactor (Scheme 1). The set-up consists of (i) digital mass flow controllers (Brookhors®) to dose C₂H₄ (PanGas, 3.5%), HCl (Air Liquide, 2.8%, anhydrous), O₂ (Messer, 19.96% in He), He (PanGas, 5.0%) as a carrier gas, and Ar (PanGas, 5.0%) as an internal standard at a total volumetric flow, F_total of 100 cm³ STP min⁻¹, (ii) a syringe pump (Nexus 6000, Chemxy) to feed EDC (Fluka, 99.5%), (iii) a vapouriser operated at 403 K, accommodating a quartz T-connector filled with glass beads to vapourize EDC, (iv) an electrically heated oven hosting a quartz micro-reactor equipped with a K-type thermocouple whose tip reaches the centre of the catalyst bed, (v) downstream heat-tracing to avoid any condensation of the reactants and products, and (vi) a gas chromatograph coupled to a mass spectrometer (GC-MS) for on-line analysis. The effluent steam was neutralised by passing it through an impinging bottle containing an aqueous NaOH solution (1 M). The catalyst (W_cat = 0.5 g, particle size d_p = 0.4–0.6 mm) was loaded in the micro-reactor (10 mm inner diameter) and pre-treated in He at 473 K for 30 min. Prior to the analysis of the reaction mixtures, the catalysts were equilibrated for at least 1 h under each condition. Carbon-containing compounds in the effluent gas stream at the reactor outlet (C₂H₄, C₂H₃Cl, C₂H₄Cl₂, CO, and CO₂) and Ar were quantified using an on-line gas-chromatograph, equipped with a GS-Carbon PLOT column, which was coupled to a mass spectrometer (Agilent GC 7890B, Agilent MSD 5977A) with a triple-axis detector and an electron multiplier. The conversion of reactant i, Xᵢ, (i: C₂H₄, C₂H₃Cl, and C₂H₄Cl₂) was calculated using eqn (1),

\[
X_i = \frac{n_{\text{inlet}}^i - n_{\text{outlet}}^i}{n_{\text{inlet}}^i} \times 100\%,
\]

where \(n_{\text{inlet}}^i\) and \(n_{\text{outlet}}^i\) are the molar flows of reactant i at the inlet and outlet of the reactor, respectively. The selectivity to gaseous...
products, $S_j$ and yield, $Y_j$ of product $j$ (j: $\text{C}_2\text{H}_4\text{Cl}_2$, $\text{C}_2\text{H}_4\text{Cl}_3$, $\text{C}_2\text{H}_4\text{Cl}$, CO, and $\text{CO}_2$) were determined according to eqn (2) and (3),

\[
S_j = \frac{n_{\text{outlet}}^j \times N_{C_{ij}}}{\sum n_{\text{outlet}}^j \times N_{C_{ij}}} \times 100\%,
\]

\[
Y_j = \frac{X_i \times S_j}{100}.
\]

where $n_{\text{outlet}}^j$ is the molar flow of product $j$ at the reactor outlet. $N_{C_{ij}}$ is the number of carbon atoms in compound $j$. The error of the carbon balance, $\varepsilon_{\text{C}}$, was determined using eqn (4),

\[
\varepsilon_{\text{C}} = \frac{n_{\text{inlet}}^{\text{C}} \times N_{C_{ij}} - \left(n_{\text{inlet}}^{\text{HCl}} \times N_{C_{ij}} + \sum n_{\text{outlet}}^j \times N_{C_{ij}} \right)}{n_{\text{inlet}}^{\text{HCl}} \times N_{C_{ij}}} \times 100\%.
\]

where $N_{C_{ij}}$ is the number of carbon atoms in compound $i$ or $j$, which was less than 5% in all experiments, i.e., the carbon mass balance reached $\geq 95\%$. Coke was not considered a product, and therefore not included in the standard selectivity calculation. However, the amount of coking that a catalyst experience is expressed as selectivity as the ratio of the rate of coke formation in the catalyst to the rate of ethene feed. In the HCl oxidation tests, Cl$_2$ production was quantified using a Mettler Toledo G20 compact titrator by off-line iodometric titration of triiodide, which is formed by purging a Cl$_2$-containing reactor outlet through a 0.10 M aqueous KI solution (Sigma-Aldrich, 99.5%), with a 0.01 M sodium thiosulfate solution (Sigma-Aldrich, 99.99%). The conversion of HCl in HCl oxidation experiments, $X(\text{HCl})$, was calculated using eqn (5):

\[
X_{\text{HCl}} = \frac{2n_{\text{outlet}}^{\text{Cl}_2}}{n_{\text{inlet}}^{\text{HCl}}} \times 100\%.
\]

where $n_{\text{inlet}}^{\text{HCl}}$ and $n_{\text{outlet}}^{\text{Cl}_2}$ denote the molar flow of HCl and Cl$_2$ at the reactor inlet and outlet, respectively. Calculation of the Weisz–Prater and Mears criteria indicated that all catalytic tests were performed in the absence of heat and mass transport limitations. After the tests, the reactor was quenched to room temperature in a He flow.

Results and discussion

Supported ceria catalysts for selective oxychlorination

To provide a comprehensive assessment of the carrier impact on the oxychlorination performance of ceria, a range of supported catalysts with a nominal cerium loading of 10 wt% was prepared including CeO$_2$/MgO, CeO$_2$/SiO$_2$, CeO$_2$/SiC, CeO$_2$/TiO$_2$, CeO$_2$/ZrO$_2$, CeO$_2$/Al$_2$O$_3$, and CeO$_2$/ZSM-5-15. X-ray diffraction (XRD) analysis of the catalysts evidenced the ceria dispersion in the form of nanoparticles (Fig. 1a), exhibiting a size range of 5–16 nm (Table 1). The only exception was CeO$_2$/ZrO$_2$ which displayed no characteristic diffraction lines of the CeO$_2$ phase. Consistent with the XRD patterns, the Raman spectra of fresh CeO$_2$, CeO$_2$/MgO, CeO$_2$/SiO$_2$, CeO$_2$/SiC, CeO$_2$/Al$_2$O$_3$, and CeO$_2$/ZSM-5-15 (Fig. 1b) exhibited a pronounced peak at 461–464 cm$^{-1}$, which corresponds to the F$_{2g}$ mode of the fluorite CeO$_2$, while the spectra of CeO$_2$/ZrO$_2$ exhibited only contributions that are ascribed to monoclinic ZrO$_2$, further corroborating high Ce dispersion. Notably, the spectra of fresh CeO$_2$/SiC and CeO$_2$/TiO$_2$ show only the peaks that can be ascribed to the Si–C bonds (ca. 796 cm$^{-1}$) and TiO$_2$-anatase phase, with no observable CeO$_2$ contribution. Considering the prominent diffraction lines of CeO$_2$ that were observed in these two samples, the absence of the characteristic active phase vibrations in the spectra of these samples can be rationalised by their significantly weaker Raman intensity as compared to their respective carriers.

To confirm the structural integrity of the investigated materials under the oxychlorination environment, all the catalyst samples were equilibrated in ethene oxychlorination at 673 K and $F_T = 100$ cm$^3$ STP min$^{-1}$ g$_{\text{cat}}^{-1}$ for 4 h using an inlet feed composition as detailed in Table 2. Most of the equilibrated catalysts displayed similar textural and crystallographic properties as the corresponding fresh materials, indicating the absence of significant structural changes such as sintering or extensive chlorination of the ceria phase (Fig. 1, Table 1). Nonetheless, more prominent changes with respect to the fresh catalysts were observed in the Raman spectra of the used CeO$_2$/SiC and CeO$_2$/MgO catalysts, which displayed new peaks at around 512 and 240 cm$^{-1}$, respectively (Fig. 1b). The former band can be ascribed to Mg–Cl bond vibrations, and the latter can be associated with the Si–Si bond vibration, suggesting the partial restructuring of these carriers. Moreover, the spectra of used CeO$_2$/Al$_2$O$_3$, and CeO$_2$/ZSM-5-15 exhibited strong fluorescence which is consistent with the pronounced propensity of these catalysts towards coking (vide infra). High-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) was used to examine the differences in ceria dispersion between the equilibrated catalysts more closely (Fig. 2). The surface-averaged size of CeO$_2$ nanoparticles showed a good agreement with those estimated by the Scherrer equation from the respective diffraction lines. However, in contrast to the low-surface area catalysts, CeO$_2$/MgO (5 nm) and CeO$_2$/TiO$_2$ (5 nm), that comprise spherical-like ceria nanoparticles with uniform size and spatial distribution, higher-surface area materials, such as CeO$_2$/SiO$_2$ (7 nm), CeO$_2$/Al$_2$O$_3$ (12 nm), and CeO$_2$/ZSM-5-15 (16 nm), exhibited a broader range of particle sizes, which are distributed in the form of small clusters and larger agglomerates leading to a significant fraction of uncovered carrier areas. In addition, CeO$_2$/ZSM-5-15 displays elongated nanoparticles, which is likely the reason for the more significant deviation between the particle size estimates between XRD and microscopy observed only for this sample. This broader particle size distribution can be attributed to the initial high dispersion of the cerium precursor (i.e. small particles) after impregnation. Depending on the affinity of CeO$_2$ for the support and its mobility on the surface upon calcination, sintering of contacting particles leads to larger clusters or agglomerates, where smaller particles sinter more easily than larger particles, which results in a broad particle size distribution on the high surface area materials.
After assessing the structural stability of supported ceria catalysts, their oxychlorination performance was investigated in broader ranges of space velocities (50–200 cm$^3$ STP min$^{-1}$ g$_{\text{cat}}^{-1}$) and temperatures (623–723 K) that closely match the typical operating window of the state-of-the-art copper chloride-based materials. The absence of deactivation-induced measurement bias was ascertained by measuring the catalytic performance under fixed standard conditions ($T = 673$ K and $F_T = 200$ cm$^3$ STP min$^{-1}$ g$_{\text{cat}}^{-1}$, feed composition as shown in Table 2) before and after the variations of temperature and space velocity, which showed identical ethene conversion levels and product selectivities.

Examination of the recorded activity and selectivity patterns in the form of a circular plot allows unambiguous benchmarking with respect to the bulk ceria (highlighted regions) and simple comparison between the different classes of catalysts. Herein, the central white part of the plot presents the specific rates of ethene oxychlorination measured at constant oxychlorination feed composition (Table 2), temperature (673 K), and space velocity (200 cm$^3$ STP min$^{-1}$ g$_{\text{cat}}^{-1}$), while the grey region indicates the corresponding selectivity to coke, determined after 4 h on stream under standard conditions (vide supra). The four outer rings provide the information on the selectivity (radial direction) to different products, i.e., VCM, EDC, DCE, and CO$_2$ as a function of ethene conversion that increases in a

Fig. 1  a) XRD patterns and b) Raman spectra of bulk and supported CeO$_2$ catalysts prior to (black) and after use (red). Reference patterns in a) are shown below the measured diffractograms in orange (CeO$_2$, ICDD-PDF 81-0792) and black (supports, ICDD-PDF numbers are shown correspondingly).
Table 1  Particle size, surface area, and pore volume of fresh and used supported catalysts with 10 wt% Ce loading and bulk ceria as the reference material

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>Particle size/µm</th>
<th>(S_{\text{BET}})/m² g⁻¹</th>
<th>(V_{\text{pore}})/cm³ g⁻¹</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Fresh</td>
<td>Used</td>
<td>Fresh</td>
</tr>
<tr>
<td>CeO₂</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CeO₂/MgO</td>
<td>5</td>
<td>7</td>
<td>(5)</td>
</tr>
<tr>
<td>CeO₂/SiO₂</td>
<td>8</td>
<td>8</td>
<td>(7)</td>
</tr>
<tr>
<td>CeO₂/SiC</td>
<td>10</td>
<td>10</td>
<td>26</td>
</tr>
<tr>
<td>CeO₂/TiO₂</td>
<td>5</td>
<td>6</td>
<td>(5)</td>
</tr>
<tr>
<td>CeO₂/ZrO₂</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>CeO₂/Al₂O₃</td>
<td>8</td>
<td>10</td>
<td>(12)</td>
</tr>
<tr>
<td>CeO₂/ZSM-5-15</td>
<td>6</td>
<td>8</td>
<td>(16)</td>
</tr>
</tbody>
</table>

a As determined by the Scherrer equation and for the selected materials, by microscopy on a particle sample size of 200 (in brackets). b As determined by N₂ sorption.

clockwise direction. Within the top half of the plot, the CeO₂/MgO, CeO₂/SiO₂, CeO₂/SiC, CeO₂/TiO₂, CeO₂/ZrO₂, CeO₂/Al₂O₃, and CeO₂/ZSM-5-15 catalysts are divided into four sections (a–d) based on their performance. The first catalyst section (a) is represented by CeO₂/MgO, showing a lower activity than bulk ceria and high CO₂ selectivity. The former could be attributed to the high degree of chlorination, or formation of chloride species, as evidenced by Raman spectroscopy. Furthermore, MgO can cover parts of small CeO₂ nanoparticles, further reducing the active phase area, while coking is pronounced, likely linked to the basicity of MgO. In section (b), exhibiting slightly higher ethene oxychlorination rates, CeO₂/SiC and in particular CeO₂/SiO₂ supported catalysts selectively yield EDC at conversion levels of up to 8%, while promoting combustion and VCM formation. The higher activity of CeO₂/SiO₂ could be linked to a difference in the surface area and CeO₂ particle structure, even though the CeO₂ particle size differs only marginally. However, in contrast to the MgO supported catalyst, CeO₂/SiO₂ exhibited no coking, as also evidenced by fully preserved pore volume after the reaction. In section (c), catalysts that mirror the performance of bulk CeO₂, are shown along with this benchmark. The performance patterns of CeO₂/TiO₂ are shifted towards lower conversion levels, yet indicate higher VCM selectivity, at the cost of stronger combustion compared to bulk ceria. CeO₂/ZrO₂, on the other hand, exhibits no coking, decreases CO₂ and DCE formation, and exceeds any other catalyst formulation in terms of activity, although the selectivity to VCM does not exceed 25%. The increased activity of CeO₂/ZrO₂ (Fig. 3 and 4a) however, allows operation at lower temperature, resulting in fully selective EDC formation at 25% ethene conversion. This might be attributed to a substantial CeO₂–ZrO₂ interaction, as shown in Fig. 1, where no CeO₂ reflections could be distinguished. Considering the well-known propensity of CeO₂ to form solid solutions with ZrO₂, high resolution, XRD was performed on CeO₂/ZrO₂ and co-precipitated Ce–Zr mixed oxide (Ce–ZrO₂) with 10 wt% nominal cerium loading to shed light on ceria dispersion in this sample (Fig. 4b). While the most prominent reflection of Ce–ZrO₂ displays a clear shift with respect to the ZrO₂ reference, the most prominent reflection of CeO₂/ZrO₂ coincides with the reference. This indicates the absence of substantial cerium inclusion in the ZrO₂ lattice in the CeO₂/ZrO₂ catalyst. Consistently, in contrast to the model solid solution material Ce–ZrO₂, EDX and HRTEM analyses revealed the presence of CeO₂ particles in CeO₂/ZrO₂ (Fig. 4c–h). To determine the descriptors for the outstanding activity of CeO₂/ZrO₂ in comparison to the mixed oxide and bulk ceria, XPS analysis revealed that bulk CeO₂ exhibits the highest Ce⁴⁺ fraction, followed by Ce–ZrO₂, and CeO₂/ZrO₂ (Fig. 4i), which allows two considerations: first, under ultra-high vacuum conditions, the material that exhibits a lower Ce⁴⁺ fraction is more easily reducible, which explains why CeO₂/ZrO₂ is much more reactive than Ce–ZrO₂ and bulk CeO₂. Second, this good reducibility of CeO₂/ZrO₂ in comparison with Ce–ZrO₂ could arise from the fact that nanoparticle systems exhibit a considerable number of defects, in comparison to the Ce–Zr mixed oxide. The latter hypothesis was further supported by XANES and EXAFS analyses (Fig. 4j), where XANES evidenced a red-shift of the Ce L₃ edge of CeO₂/ZrO₂ confirming more oxygen vacancies or defects in the surface, while EXAFS showed distinct characteristics of the Ce–O paths. Even though the errors are not negligible, the Ce–ZrO₂ mixed oxide exhibits the highest number of neighbouring atoms, followed by bulk CeO₂ and CeO₂/ZrO₂ (Table 3), which indicates that in particular, CeO₂/ZrO₂ exhibits strongly defective CeO₂ nanoparticles, in contrast to the mixed oxide, which shows an even higher number of neighbouring atoms than the reference value (8). In addition, CeO₂/ZrO₂ exhibits a slightly decreased Ce–O distance (2.31 Å) in comparison to the used bulk CeO₂ (2.34 Å) due to the strongly defective surface, while the mixed

Table 2  Reactions and conditions used in this work

<table>
<thead>
<tr>
<th>Reaction</th>
<th>Concentration/a/vol%</th>
<th>T/K</th>
<th>(C_2H_4)</th>
<th>EDC</th>
<th>VCM</th>
<th>HCl</th>
<th>O₂</th>
<th>Ar (^b)</th>
<th>He (^c)</th>
</tr>
</thead>
<tbody>
<tr>
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<td></td>
<td>623–723</td>
<td>3</td>
<td>0</td>
<td>0</td>
<td>4.8</td>
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<td>3</td>
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\(^a\) \(F_y/\text{cat}^{-1} = 200 \text{ cm}^3 \text{ STP min}^{-1} \text{ g}_{\text{cat}}^{-1}\) unless indicated otherwise. \(^b\) Internal standard. \(^c\) Carrier gas.
oxide shows even lower Ce–O distances (2.27 Å), most likely related to the accommodation of additional atoms in the lattice. This incorporation of Ce into the ZrO$_2$ matrix is indicated by a peak in the EXAFS spectra, ascribed to the Ce–Zr scattering path. In addition, the average Zr–O bond length was found shorter than that of Ce–O, while in general, the metal–oxygen bond renders much shorter in the mixed oxide than in CeO$_2$ or CeO$_2$/ZrO$_2$, suggesting again that CeO$_2$/ZrO$_2$ predominantly features CeO$_2$ nanoparticles.

The fourth section (d) of the supported ceria catalysts in Fig. 3 is represented by CeO$_2$/Al$_2$O$_3$, and CeO$_2$/ZSM-5-15, which both exhibit an encouraging performance with respect
to VCM selectivity (>75%, CO\textsubscript{x} and EDC) at 10–20% conversion, while differences between the two materials arise outside this region. In contrast to the Al\textsubscript{2}O\textsubscript{3} supported sample, the higher activity of CeO\textsubscript{2}/ZSM-5-15 is impaired by...
strongly reduced VCM selectivity at higher conversions. In addition, both materials form combustion products under all circumstances, in increased amounts at high temperature or low flow rates, and coking of the catalysts is considerable, especially for CeO$_2$/ZSM5–15. However, the activity and selectivity patterns show no clear correlation with particle size, total surface area, pore volume (Table 1), or particle surface area (Fig. S1†). By aiming to better understand the
observed activity and selectivity differences, we evaluated three distinct catalyst key descriptors: (i) activity in ethene oxychlorination, likely linked to the redox properties of the material, (ii) VCM selectivity, linked to the dehydrochlorination of EDC and material acidity, and (iii) propensity towards combustion, again linked to the redox properties (Fig. 5).

In order to address the first point, HCl oxidation was adopted as a suitable probe reaction to characterise the redox properties of a material for ethene oxychlorination, due to its similarities in reaction conditions and the requirement for Cl evolution in both reactions. In contrast, the typical application of H2-TPR to assess the redox properties of the used catalysts led to no convincing correlation (Fig. S2†), which is ascribed to the chlorination of the materials under reaction conditions.40,41 Fig. 5a depicts the rate of ethene consumption in oxychlorination as a function of the rate of HCl oxidation over pre-equilibrated catalysts, evidencing a clear positive correlation between the two rates. This indicates that the activity of a catalyst in ethene oxychlorination for VCM formation is primarily determined by its ability to activate ethene and HCl, which requires two sites to form EDC, in line with previous theoretical studies.16 Thus, the activity of a catalyst in ethene oxychlorination is mainly determined by its ability to activate HCl, in line with the light-off curves in HCl oxidation (Fig. 5a, S3a†), the availability of chlorinated vacancies in ethene activation,16 and the ease of chlorine or EDC evolution from such vacancies.16 Yet, considering that all carriers were inactive in ethene oxychlorination, promotional effects of the support that go beyond the determination of the previously described CeO2 particle shape and speciation with respect to the Ce4+ fraction are unlikely. Secondly, rationalising the VCM selectivity, primarily influenced by the material acidity,24 was performed by using the EDC dehydrochlorination activity as an acidity descriptor that can be evaluated in an environment similar to ethene oxychlorination. This resulted in a clear positive trend between high VCM selectivity in ethene oxychlorination and low temperature where 20% conversion is reached in EDC dehydrochlorination (Fig. 5b). In addition to rationalising the VCM selectivity by the EDC dehydrochlorination performance, acidity characterisation by temperature programmed desorption of ammonia (NH3-TPD, Fig. 6) corroborated the found relationship, where only CeO2/ZrO2 renders as an outlier, likely due to the tight Ce–Zr integration, which is likely influenced by the presence of chlorine, and thus not correctly probed by ammonia. On the other hand, when considering the amount of acid sites (Fig.

<table>
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<th>Catalyst</th>
<th>N/−</th>
<th>σ²[A²]</th>
<th>R/Å</th>
<th>R-factor</th>
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</table>

Table 3 Number of neighbouring atoms (N), disorder in the neighbour distance (σ²), distance to neighbouring atoms (R), and R-factor, determined by EXAFS for the Ce–O path. The corresponding errors are reported in brackets.
S4 instead of the average strength (Fig. 5b), CeO2/MgO, which exhibits strong coking, contains the lowest number. Nevertheless, both correlations lead to the same conclusion that acid features of the material enhance VCM selectivity in accordance with EDC dehydrochlorination tests (Fig. 4b). However, the increase in acidity also promotes coking. Thus, optimal performance necessitates moderated acid properties. Furthermore, pure supports were tested and compared with the catalysts in EDC dehydrochlorination, evidencing only slight shifts of T20 to a higher temperature for materials with high VCM selectivity and shifts of 50 K for materials with low VCM selectivity (Fig. 5b). This indicates that indeed EDC dehydrochlorination is promoted by the presence of CeO2 in the catalyst formulation, but the major part of VCM formation seems to happen on the carrier, implying the existence of true bifunctional, two-site catalysts. Considering the ceria nanoparticle distribution as shown in Fig. 2, big particles or clusters of nanoparticles of ceria with large uncovered carrier surfaces in between therefore seem to facilitate the two-step reaction oxychlorination–dehydrochlorination on a hybrid surface. Nevertheless, the pronounced coking of CeO2/Al2O3 and CeO2/ZSM-5-15 was also observed for the corresponding pure carriers, indicating that CeO2 in the catalyst formulation is not the cause, but rather the result of coke formation (Fig. 6b), that is caused by the support acidity, as also observed for dehydrochlorination catalysts reported in the literature.20,26 Thirdly, the propensity towards combustion in ethene oxychlorination is likely not only determined by the presence of free or chlorinated vacancies, which are known to reduce combustion to some extent,16 but primarily influenced by the ease of oxygen removal from the surface.16,42 As this property is heavily influenced by the degree of chlorination and thus difficult to quantify under relevant reaction conditions,16 another probe reaction is used to rationalise the selectivity towards combustion products. While ethene oxidation primarily yields CO2 on CeO2, chlorinated hydrocarbons yield CO2 and CO.16 Therefore, the oxidation of VCM (Fig. S3b†) was studied on pre-equilibrated catalysts and correlated with the CO2 selectivity in ethene oxychlorination, as depicted in Fig. 5c. Therein, the temperature where 20% conversion in VCM oxidation was reached serving as a measure for the ease of combustion, resulting in a linear correlation between bulk CeO2 and CeO2/ZSM-5-15, CeO2/Al2O3, CeO2/ZrO2, and CeO2/
SiO₂. In contrast to HCl oxidation, CeO₂/MgO emerged as the only outlier, surprisingly displaying the least activity in VCM oxidation while exhibiting the highest COₓ selectivity in ethene oxychlorination, which might be linked to oxidation of ethene or EDC. Nevertheless, the correlation of the other materials exemplifies how strongly dehydrochlorination and oxidation functions are intertwined or influence each other, as catalysts with high VCM selectivity also produce a high fraction of combustion products. Thus, in accordance with studies on bulk ceria, a very simplified reaction pathway for nanostructured CeO₂ catalysts can be formulated as C₂H₄ → EDC → VCM → COₓ, clarifying the need for a catalyst which enables steps one and two, but suppresses step three, while two and three seem to be linked.

Inspired by the high activity and high selectivity towards the VCM of CeO₂/ZSM5-15, as well as the goal to satisfy the main priority to suppress combustion, i.e. designing an active catalyst that would lie in the lower right corner of Fig. 5c, zeolite supports were further investigated with the aim of understanding the impact of the framework, Si:Al ratio, Ce loading, and calcination temperature on the catalytic performance, as shown in Fig. 3e–h. First, considering different framework types (FER, Y, MOR, and ZSM-5) with as close as possible Si:Al ratios and 10 wt% Ce loading, it becomes evident that their activity at the same reaction temperature is very different, ranging from 4.5 (CeO₂/FER-29) to 10 (CeO₂/MOR-110) molC₂H₄ h⁻¹ molCe⁻¹. However, none of the catalysts reached more than 70% VCM selectivity, or fully suppressed combustion or DCE formation. In addition, the observed differences in the conversion–selectivity relationships are marginal considering the hugely different frameworks and the number of changing parameters in the system. Therefore, in the Pareto set of conversion–selectivity curves, the ZSM-5 framework was selected as a carrier material for further studies, due to (i) its good selectivity towards VCM at reasonable conversions and most importantly, (ii) its broad available range of Si:Al ratios. Therein, four carriers with Si:Al ratios of 15, 40, 140, and 1040 were selected, of which the corresponding catalysts with 10 wt% Ce loading showed no linear trend in terms of activity. Instead, a maximum could be observed for CeO₂/ZSM5-140 (the order with respect to ethene oxychlorination rate: CeO₂/ZSM5-140 > CeO₂/ZSM5-40 > CeO₂/ZSM5-15 ≈ CeO₂/ZSM5-1040), which might be attributed to better dispersion of ceria nanoparticles due to optimal acid properties for Ce anchoring. In terms of selectivity, a higher Si:Al ratio of the carrier led to lower COₓ and higher EDC selectivity, while the selectivity of VCM decreased (as evidenced also by the probe reaction in Fig. 5b) and DCE formation, already at very low levels, was not affected. The selectivity to coke decreased with a rising Si:Al ratio as expected, since acidic sites in the zeolite and aromatics formation on its external surface are known to generate coke. The fact that strong acidity is necessary for EDC dehydrochlorination, while it seems to promote COₓ and coke formation, excludes the as-prepared zeolite based catalysts from successful application in ethene oxychlorination to VCM. One remaining strategy to tune the

![Fig. 7](https://example.com/fig7.png)

**Fig. 7** a) VCM yield as a function of temperature in EDC dehydrochlorination over the Al₂O₃-based catalysts. b) Selectivity to VCM (top) and coke (bottom) in a dual-catalyst configuration as a function of the temperature of 20% (solid symbols) and 90% (open symbols) conversion in EDC dehydrochlorination. In the dual-catalyst configuration for ethene oxychlorination, CeO₂/ZrO₂ was used in the first bed, and the corresponding Al₂O₃-based catalyst in the second bed, operated at 623 K. Conditions are detailed in Table 2.
The performance of zeolite supported ceria systems might depend on adjusting the ceria particle size and coverage of the acidic carrier through optimisation of Ce loading and calcination temperature (Fig. 3g and h). However, while an increase of the calcination temperature slightly decreased the activity, not affecting the selectivity significantly, lower (5%) or higher (20%) Ce loading negatively affected the performance by strongly decreasing activity, or increasing combustion, respectively.

Overall, considering the observed catalytic performance and the descriptors in Fig. 5, it indicates that all the catalysts promoting VCM formation also promote combustion and coking, which leaves only CeO$_2$/ZrO$_2$ as a candidate that remains highly active and selective towards EDC at low temperatures. In addition to forming no other chlorinated hydrocarbons than EDC, no Cl$_2$ evolved on any catalyst below 673 K, preserving a high chlorine efficiency.

**Dual catalytic system for selective VCM production**

In order to overcome the above described obstacles, while refraining from using thermal but instead catalytic EDC dehydrochlorination, a promising catalyst for the latter was developed. Pure Al$_2$O$_3$ was selected as the starting material (Fig. 7a), since CeO$_2$/Al$_2$O$_3$ was found to coke less in ethene oxychlorination compared to CeO$_2$/ZSM-5-15 (Fig. 3). Moreover, Al$_2$O$_3$ enables the same dehydrochlorination performance as ZSM-5-15, reaching 100% yield at 623 K (Fig. S3c†). In order to reduce this coking, Al$_2$O$_3$ was promoted with additives known to improve selectivity for copper based oxychlorination catalysts,$^{1,2,3,4,5}$ in particular Ca, K, La, and Ce. Doping with 2 wt% metal led to reduced coke formation, with an effectiveness order K > Ca > Ce > La, while the VCM yield decreased in the almost opposite order: Ce/Al$_2$O$_3$ > La/Al$_2$O$_3$ > Ca/Al$_2$O$_3$ > K/Al$_2$O$_3$ (Fig. 7a). Using a dual catalyst configuration with 0.5 g CeO$_2$/ZrO$_2$ and a 0.5 g Al$_2$O$_3$-based bed in series in one reactor (similar to illustration in Fig. 9), the VCM selectivity could again be correlated with the temperature of 20% (and 90%) conversion in EDC dehydrochlorination, as shown in Fig. 7b. This suggests that the used dopants mask the acid sites, which are responsible for coking and EDC dehydrochlorination. In addition, the NH$_3$-TPD of fresh dehydrochlorination catalysts (Fig. 8a) evidences a decrease in total acidity upon doping with different cations, which is in accordance with the dehydrochlorination performance (Fig. 7). However, catalysts where coking was reduced to values close to zero did not show high VCM yields in the desired temperature region below 673 K. Therefore, further optimisation was performed by reducing the dopant loading to 0.5 wt% on the intermediate Ca promoted Al$_2$O$_3$, with the aim of masking primarily the strongest sites, while keeping weaker sites available for EDC dehydrochlorination, resulting in Ca-I/Al$_2$O$_3$, which showed a 40% VCM yield at 623 K (Fig. 7a). In addition, reaching 90% conversion in EDC dehydrochlorination was enabled in contrast to...
to Ca/Al$_2$O$_3$. The coke selectivity was slightly increased, yet very limited, following a correlation between acidity availability for EDC dehydrochlorination and coke selectivity in ethene oxychlorination (Fig. 7b). However, as the yield of this catalyst was insufficient for application in VCM selective dual catalytic systems, different strategies to increase the EDC conversion were explored. In the first step, the amount of Ca-l/Al$_2$O$_3$ was increased by a factor of 6, still not yielding full EDC conversion and producing new products, in particular ethyl chloride (EtCl), in parallel with increased coke selectivity (Fig. 9). Tighter integration of the same amounts of the two catalysts in a single, physically mixed bed reduced the overall coke selectivity, likely due to the fact that the VCM is a major contributor to coke formation, yet also reduced VCM selectivity and increased the formation of combustion products. Increasing the temperature to achieve higher EDC conversion on the dehydrochlorination catalyst, while reducing the amount of the oxychlorination catalyst for keeping similar conversion levels (Fig. 9, panel three), led as expected to higher EDC conversion, but also increased combustion and DCE selectivity. Therefore, the solution to preserve the selective performance of the oxychlorination step, while ensuring selective and full EDC dehydrochlorination to VCM in the second step with minimal coking, was to operate CeO$_2$/ZrO$_2$ at 623 K, coupled to a separate reactor, containing Ca-l/Al$_2$O$_3$ and operated at 673 K. This indeed resulted in full and 100% selective EDC to VCM conversion (at 25% ethene conversion) and a coke selectivity of <0.3% (Fig. 9).

This system was then subjected to a stability test, as depicted in Fig. 10, where the conversion, determined by CeO$_2$/ZrO$_2$, rendered stable for 160 h on stream. However, while no combustion or formation of undesired chlorinated products occurred, the selectivity to VCM, and thus EDC, changed over the course of the test. The observed decrease of VCM selectivity can be linked to a moderate loss of acidity upon use (Fig. 8b), and a decrease in surface area and pore volume (Table 3), consistent with the formation of coke species. Yet, the decrease levelled off, reaching stable values of about 80% VCM and 20% EDC selectivity, which
outperforms the current dehydrochlorination yield by 60% at a 100 K lowered temperature.\textsuperscript{1,2,6}

Conclusions

In this work, we developed copper-free dual catalytic systems for direct VCM production from ethene. In order to suppress combustion, chlorinated side-product formation, and coking, while improving the VCM selectivity to make thermal cracking obsolete, nanostructured ceria catalysts were designed in the first step. In-depth rationalisation and exploration of carrier effects led to the result that achieving all the goals is not feasible in a single material. CeO\textsubscript{2}/ZrO\textsubscript{2} proved to be a stable oxychlorination catalyst, exhibiting full selectivity to EDC and extraordinary activity even at low temperature. In order to catalytically dehydrochlorinate EDC to VCM, a dedicated catalyst for this second function was developed, consisting of the promoted solid acid catalyst Ca–Al\textsubscript{2}O\textsubscript{3} which led to a minimal coking, and 100% EDC selectivity to EDC and extraordinary activity even at low temperature. In order to catalytically dehydrochlorinate EDC to VCM, a dedicated catalyst for this second function was developed, consisting of the promoted solid acid catalyst Ca–Al\textsubscript{2}O\textsubscript{3} which led to a minimal coking, and 100% EDC conversion to VCM. Some degree of deactivation was observed during the first 100 h on stream, decreasing the overall selectivity to VCM from 100% to stable 80% at 25% ethene conversion. Nevertheless, the developed dual system outperforms EuOCl as the best ethene oxychlorination catalyst to directly form VCM by a factor of four in terms of space time yield. Thus, this study provides the basis for potential intensification of the current industrial-scale VCM.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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References