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# Improving the conversion of petroleum coke to solid acid catalysts – effectiveness of hydrogen peroxide and impact of silica contamination

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Petroleum coke (petcoke) samples containing silica sand (SiO<sub>2</sub>) in the concentrations of ~50 and 7 wt% were functionalized with nitric acid and hydrogen peroxide to produce carbon-based solid acid catalysts by converting the inherent sulfur species in petcoke to sulfonic acid sites (–SO<sub>3</sub>H). The effect of SiO<sub>2</sub> content and functionalization agent on the physical, surface, and acidic properties of the prepared materials were evaluated, and their catalytic activities were assessed in the esterification reaction of octanoic acid and methanol at 60 °C, yielding 17–54% of ester after 6 h. The functionalization with H<sub>2</sub>O<sub>2</sub> is more environmentally friendly than with HNO<sub>3</sub> and produces more active catalysts (ester yields of 54% and 45%, respectively). The activities of the prepared materials were more dependent on having fewer oxygen-containing carbon groups (related to total acidity) on the surface than on a higher concentration of sulfonic acid sites. Thus, the improved activity of the catalyst prepared with H<sub>2</sub>O<sub>2</sub> was due to a lower total acidity (~1 mmol g<sup>–1</sup>) than the catalyst prepared with HNO<sub>3</sub> (~5 mmol g<sup>–1</sup>). The stabilities of petcoke-derived catalysts were poor, and regeneration was not possible, which is a common problem for carbon-based solid acid catalysts. The petcoke with a higher content of SiO<sub>2</sub> resulted in better esterification performance. The silica sand promoted the reduction of particle size during the ball-milling process, leading to reduced diffusion limitations and the generation of silanol groups that are hydrogen-bond donors, helping in the esterification reaction.

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## Introduction

Petroleum coke (petcoke) is a solid by-product from the refining of crude oil and upgrading of bitumen. This carbon-rich material (>80 wt%) also contains hydrogen, nitrogen, oxygen, sulfur (1–8 wt%), and several trace elements,<sup>1</sup> and is stockpiled at a rate of ~6 million tonnes per year globally.<sup>2</sup> The sulfur content limits the direct use of petcoke, and as the piles of petcoke are open to the atmosphere, the release of pollutants – volatile organic and aromatic compounds – to the air and groundwater is a problem.<sup>3</sup> Thus, researchers have been exploring ways to valorize this feedstock, including as an adsorbent for heavy metals sequestration,<sup>4</sup> a precursor for the preparation of carbon-based nanomaterials,<sup>5</sup> supercapacitors,<sup>6</sup> a support for photocatalysts,<sup>7</sup> a co-gasification feed with biomass,<sup>8</sup> adducts in construction materials,<sup>9</sup> and a solid acid catalyst.<sup>10–13</sup> While various reactants can be used to convert the petcoke to these products, many of these reactants are not environmentally friendly.

The sulfur species in petcoke were reported to be mainly organic compounds, accounting for more than 90 wt% of the

total sulfur, and the remaining were inorganic species (sulfates). The organic fraction contains dominantly thiophene, benzothiophene, and dibenzothiophene molecules (60–80 wt%), 10–30 wt% of sulfoxide, and a minor fraction (<10 wt%) of sulfides/disulfides.<sup>12,14–16</sup> These thiophenes have high thermal stability, making their removal hard to accomplish. However, they can be converted through an oxidation process. Strong acids are often used as oxidizers; nitric acid, for example, is effective at converting the inherent organic sulfur into sulfonic acid sites (–SO<sub>3</sub>H)<sup>12,13</sup> but releases NO<sub>x</sub> gases and promotes SO<sub>2</sub> formation during its decomposition in the petcoke functionalization process.<sup>13,17</sup> After oxidation with nitric acid, the sulfur content in petcoke was reported to decrease around 20–40 wt% due to SO<sub>2</sub> formation, and the sulfur species were reported to be exclusively organic, in which 10–30 wt% were oxidized sulfur (–SO<sub>3</sub>H), and the remaining was thiophenic compounds.<sup>12,13</sup>

Solid-acid catalysts from petcoke were also prepared in a two-step process involving chemical activation with KOH and sulfonation with H<sub>2</sub>SO<sub>4</sub>.<sup>18,19</sup> Another study applied chemical activation with KOH and impregnation with ammonium heptamolybdate to enhance the mesoporosity, then used functionalization with HNO<sub>3</sub> to convert petcoke into a mesoporous solid acid catalyst.<sup>11</sup> These activation processes resulted in the removal of sulfur from petcoke.<sup>20</sup> For the conversion of other

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carbon materials, such as agricultural waste, researchers have used mainly concentrated acids (*e.g.*, H<sub>2</sub>SO<sub>4</sub>, HNO<sub>3</sub>, and H<sub>3</sub>PO<sub>4</sub>) *via* a sulfonation process to introduce acidity to the surface of the carbon material. Before the sulfonation, however, most waste carbon feedstocks were modified with carbonization, pyrolysis or activation.<sup>21–26</sup> Based on sulfur conversion pathways predicted *via* density functional theory (DFT) calculations, 'OH is formed and could contribute to the oxidation of sulfur in petcoke.<sup>13</sup> Therefore, hydrogen peroxide, which decomposes to water, may be effective because it produces these radicals but is less active for carbon oxidation compared to HNO<sub>3</sub><sup>27,28</sup> and so will result in fewer undesirable surface groups (*i.e.*, carbonyl, carboxyl, and hydroxyl).

Aside from the components within the petcoke, other contaminants may be introduced from the extraction, processing, and storage steps.<sup>29–35</sup> Petcoke is one of many waste materials that contains silica. Agricultural waste, such as rice husk, sugar cane bagasse, wheat husk, and corn straw, may contain up to 50–86 wt% silica/silicon in the ash.<sup>36–39</sup> Similar to petcoke, the harvesting, handling, and storage processes of these agricultural wastes can incorporate soil, resulting in a SiO<sub>2</sub> concentration of 2–25 wt% in these feedstocks.<sup>38,39</sup> Silica is generally an inert material for many reactions but may damage the process equipment because of its hardness. This material is also detrimental to the valorization of other waste feedstocks, affecting the feasibility of the pyrolysis of waste tires to value-added products,<sup>40</sup> and inhibiting the biogas production from sewage sludge.<sup>41</sup> Separation of the silica before processing the material, however, adds complexity and cost to the overall process, possibly negating the benefits of using a waste feedstock.

In this study, petcoke-derived solid acid catalysts were prepared using hydrogen peroxide to access the sulfur species and convert them to sulfonic acid sites and were characterized with various techniques. Two batches of petcoke with different silica contents were used to evaluate the effect of SiO<sub>2</sub> on the functionalization process and subsequent activity of the produced catalysts. The reaction of octanoic acid and methanol to produce an ester was used to evaluate the catalysts. Esterification reactions such as this one are part of the route to convert biomass, a renewable resource, into biodiesel.

## Experimental

### Materials and methods

**Materials.** The delayed petcoke (PC) from the oil sands industry used in this work was supplied by Suncor Energy Inc., Alberta, Canada. The two batches contained different levels of silica. The batch with the higher ash content (~50 wt%) is referred to as PCSi, while the batch with the lower ash content (~7 wt%) is referred to as PC. Before any experiments, samples from both batches were ground and sieved into particles in the range of 20–100 μm. Nitric acid (HNO<sub>3</sub>, 70 wt%, Sigma-Aldrich), hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>, 30% w/w, VWR Chemicals BDH), and hydrochloric acid (HCl, 37 wt%, Sigma-Aldrich) were used to prepare petcoke-derived acid catalysts. Methanol (MeOH, HPLC ≥ 99.9%, Sigma-Aldrich) and octanoic acid (OA, 99%, Thermo

Scientific) were the reactants for the esterification reaction. A commercial solid-acid catalyst, Amberlyst-15 ion exchange resin (Thermo Scientific), was also tested for comparison to the petcoke-derived catalysts. A 0.1 N standard solution of sodium hydroxide (NaOH, J. T. Baker-Avantor), a 0.01 N standard solution of hydrochloric acid (J. T. Baker-Avantor), and phenolphthalein (1% w/v in 95% v/v Alcohol, RICCA Chemical Company) were utilized in the titration experiments to quantify total acidity.

**Preparation of petcoke-derived solid acid catalysts.** Before any treatment, PC and PCSi were wet ball milled using a Planetary Ball Mill (Pulverisette 6, Fritsch), with 5 mm zirconia balls at 300 rpm for 4 h at a sample-to-ball mass ratio of 1 : 25 (4 g of sample: 100 g of ZrO<sub>2</sub> balls) and 25 mL of isopropanol. The samples were dried in a vacuum drying oven at 65 °C to remove the isopropanol and named BPC and BPCSi, respectively. The ball-milled samples were functionalized with nitric acid as previously reported.<sup>12,13</sup> In a round-bottom flask, 5 g of BPCSi were mixed with 50 mL of HNO<sub>3</sub> and stirred at 600 rpm during refluxing at 90 °C for 8 h. Then, this mixture was filtered using a 0.22 μm membrane filter, and the solid was washed with reverse osmosis water and a 0.6 wt% HCl solution. After that, the solid was dried in a vacuum oven at 65 °C overnight to remove moisture and HCl, producing the BPCSi-HNO<sub>3</sub> catalyst. Functionalization with hydrogen peroxide was done for both BPCSi and BPC, and involved mixing 5 g of ball-milled petcoke with 50 mL of 30% w/w H<sub>2</sub>O<sub>2</sub> followed by stirring at 600 rpm at room temperature for 20 h. The filtration, washing, and drying processes were conducted as previously described, producing the BPCSi-H<sub>2</sub>O<sub>2</sub> and BPC-H<sub>2</sub>O<sub>2</sub> materials.

**Characterization.** Thermogravimetric analysis (TGA, SDT Q600, TA Instruments, New Castle, DE, USA) was performed to determine the ash content and check the thermostability of the samples. Approximately 15 mg of material was heated under an air flow of 50 mL min<sup>-1</sup> from ambient temperature to 750 °C at 20 °C min<sup>-1</sup> and held at this temperature for 30 min. The structural properties of petcoke-derived catalysts were investigated by X-ray diffraction of powder samples (XRD, D8 Advanced ECO, Bruker, Billerica, MA, USA), collected using a CuKα X-ray source, λ = 0.15418 nm, 2θ angles between 5 to 75°, 40 kV, and 25 mA. The samples were analyzed as received and loaded into a 25 mm polymethyl methacrylate (PMMA) sample holder. Chemical composition (CHNS) was determined by ultimate elemental analysis using a Flash 2000 CHNS analyzer (Thermo Scientific, Waltham, MA, USA). The morphology of the surface was investigated by scanning electron microscopy (SEM, Phenom Pro X, Thermo Fisher Scientific Phenom-World BV) at an acceleration voltage of 15 kV. Nitrogen adsorption and desorption isotherms were collected by N<sub>2</sub> physisorption at -196 °C using a Tristar II Plus analyzer (Micromeritics Instrument Corporation, Norcross, GA, USA), and the non-local density functional theory (NLDFT) model was used to determine surface area and pore volume. All samples were degassed at 150 °C under vacuum overnight before the analysis.

Surface functional groups on the samples were identified using a Fourier Transform Infrared (FTIR, Nicolet iS50, Thermo



Fisher Scientific, Waltham, MA, USA) spectrometer with attenuated transmission reflectance (ATR). The spectra were collected in the wavenumber range of 4000–400  $\text{cm}^{-1}$ , at a resolution of 4  $\text{cm}^{-1}$  and 128 scans. Additional surface analysis was performed by X-ray photoelectron spectroscopy (XPS, Kratos Axis Spectrometer, Kratos Analytical Limited, Manchester, UK) with monochromatized Al  $K\alpha$  ( $h\nu = 1486.71$  eV). Surface elemental compositions of the samples were obtained from the survey spectra, considering major elemental peaks. The peak fitting of components in the C 1s and S 2p narrow-scan spectra was done using CasaXPS (Version 2.3.22PR1.0, Casa Software Ltd, Teignmouth, UK), in which each component represents different chemical bonds. The peaks in the narrow scan spectra were fitted according to chemical and physical constraints.<sup>16,42</sup> For the C 1s fitting, the constraints applied were the same FWHM for all peaks (0.8–1.8 eV), with peak position as reported in literature,<sup>13,43–45</sup> with a shift of  $\pm 0.3$  eV for each component. While the fitting of peaks for the S 2p narrow scan followed the approximate 2 : 1 relative area of the spin-orbit doublets ( $2p_{3/2}$  and  $2p_{1/2}$ ) for each species, separated by 1.18–1.2 eV with equal FWHM (1.2–1.6 eV), and peak position as reported in literature.<sup>44–51</sup> The total acidity (number of total surface acidic groups) on the petcoke-derived catalysts was determined using a modified Boehm titration method. A mixture of 0.05 g of sample and 5 mL of a 0.1 N NaOH standard solution was oscillated in a shaker at 25 °C, 250 rpm for 24 h. Then, 1 mL of this solution was filtered and back-titrated with a 0.01 N standard solution using phenolphthalein as indicator.

**Esterification reaction.** The esterification reactions between octanoic acid and methanol were performed in a 100 mL two-neck round-bottom flask connected to a reflux condenser. The system was heated in a water bath and constantly stirred at 850 rpm. For a typical run, the methanol to octanoic acid molar ratio used was 10 : 1 (~23.0 g of MeOH and ~10.0 g of OA), mixed with ~0.5 g of catalyst (5 wt% of OA) for 6 h at 60 °C. Periodically (every 30 min to 90 min then every 60 min), 0.2 mL samples were withdrawn from the mixture and filtered with a membrane filter (0.22  $\mu\text{m}$ , nylon) to remove the solid catalyst. Once withdrawn, the solution cooled to room temperature at which the rates of the heterogeneous and homogeneous reactions were negligible. A fraction of the withdrawn sample (44  $\mu\text{L}$  of solution) was diluted in 10 mL of methanol and quantified by gas chromatography with Flame Ionization Detection (GC-FID, 8860 GC-FID, Agilent Technologies, Santa Clara, CA, USA) using a DB-FATWAX UI (polyethylene glycol-type, 30 m  $\times$  0.25 mm  $\times$  0.25  $\mu\text{m}$ ) capillary column. The temperature program of the GC consisted of 0.5 min at 50 °C, then a ramp of 40 °C  $\text{min}^{-1}$  to 200 °C, holding for 2 min. Helium (He) was used as the carrier gas, in which an aliquot of 1  $\mu\text{L}$  of each sample was injected in the split mode (ratio of 100 : 1), with an inlet temperature of 250 °C. The detector mode was at a constant makeup flow of 25 mL  $\text{min}^{-1}$  of He, 450 mL  $\text{min}^{-1}$  of air, and 40 mL  $\text{min}^{-1}$  of hydrogen ( $\text{H}_2$ ). The samples were analysed in triplicate, and the error of the analysis was estimated to be <1%. The only chemicals identified by GC-FID were octanoic acid (OA) and methyl octanoate (MOA).

The ester yield was calculated based on the following equation:

$$\text{Ester Yield}(\%) = \frac{[\text{MOA}]_{\text{actual}}}{[\text{MOA}]_{\text{theoretical}}} \times 100\% \quad (1)$$

where  $[\text{MOA}]_{\text{actual}}$  is the concentration of methyl octanoate formed and  $[\text{MOA}]_{\text{theoretical}}$  is the theoretical maximum concentration of ester, which is equal to the initial concentration of octanoic acid. The initial turnover frequencies (TOF), which represent the catalyst activity per site, were calculated as follows:<sup>13</sup>

$$\text{TOF}(\text{h}^{-1}) = \frac{k \times [\text{OA}]_0 \times V}{m_{\text{catalyst}} \times [-\text{SO}_3\text{H}]} \quad (2)$$

where  $k$  is the pseudo first-order rate constant ( $\text{h}^{-1}$ ),  $[\text{OA}]_0$  is the initial concentration of octanoic acid ( $\text{mmol mL}^{-1}$ ),  $V$  is the total volume of the reaction solution (mL),  $m_{\text{catalyst}}$  is the mass of catalyst utilized in the reaction (g), and  $[-\text{SO}_3\text{H}]$  is the concentration of sulfonic acid sites on the catalyst surface ( $\text{mmol g}^{-1}$ ). Eqn (3) shows the pseudo first-order model, which assumes a constant methanol concentration and negligible reverse reaction rate:<sup>10,52</sup>

$$[\text{OA}] = [\text{OA}]_0 e^{-kt} \quad (3)$$

where  $[\text{OA}]$  is the concentration of OA ( $\text{mmol mL}^{-1}$ ) at time  $t$  (h) and  $k$  is the rate constant ( $\text{h}^{-1}$ ) for the pseudo first-order model, in which all assumptions were satisfied with the conditions applied at the initial stages of the reaction.

**Catalyst reuse and regeneration.** To evaluate the stability of the catalysts, a reuse experiment was performed with the spent catalyst recovered after the first reaction cycle. After the reaction, the catalyst was filtered, washed with methanol to eliminate any chemical adsorbed on the surface of the material, and dried overnight in the fume hood. Then, the catalyst was used again for the esterification reaction under the same conditions previously described.

For the regeneration of catalysts, the procedure was modified from the literature.<sup>53</sup> The deactivated material (after the first cycle) was washed with methanol and acetone. Then, the catalyst was immersed in 50 mL of 1 M  $\text{H}_2\text{SO}_4$  solution for 24 h and stirred at room temperature. After that, the catalyst was filtered and washed with reverse osmosis water until the residual solution reached pH 7. The samples were dried in a vacuum oven at 60 °C overnight before being used in the esterification reaction once more.

## Results and discussion

### Esterification reaction

Fig. 1 shows the catalytic activity of the petcoke-derived solid acid catalysts prepared by chemical treatment with nitric acid or hydrogen peroxide in the esterification reaction of octanoic acid and methanol. Each ester yield data point is an average of 3 runs, and the error bars represent one standard deviation. Without functionalization, PCSi, and the ball-milled sample (BPCSi) were inactive for the esterification reaction, and the



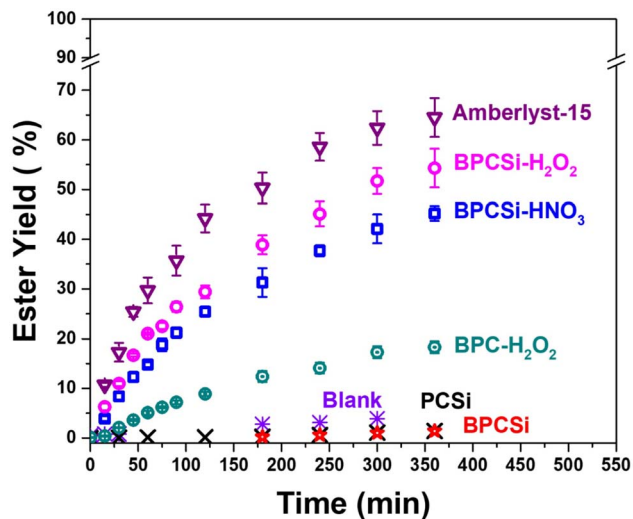


Fig. 1 Esterification of octanoic acid using petcoke-derived catalysts. Reaction conditions: MeOH : OA molar ratio = 10 : 1, 5 wt% of catalyst (based on the weight of octanoic acid), 60 °C, 800 rpm.

ester yield obtained in both cases (1.5%) were lower than the blank test (4.4%). The blank test results indicate that there was a relatively slow homogeneous reaction occurring. The lower yield obtained in the presence of PCSi and BPCSi can be explained by the adsorption of the product, from the homogeneous reaction, on the oxygen-containing carbon groups (*i.e.*, hydroxyl and ether groups) on the surface of petcoke.<sup>13</sup>

After chemical treatment, the activity improved in the order of samples BPC-H<sub>2</sub>O<sub>2</sub> < BPCSi-HNO<sub>3</sub> < BPCSi-H<sub>2</sub>O<sub>2</sub>, with ester yields after 6 h of 17%, 45%, and 55%, respectively. All prepared materials had a lower yield than the 65% yield obtained with the commercial Amberlyst-15 catalyst. Ester yields negatively correlate with the number of oxygen-containing carbon

functional groups on the petcoke-derived catalyst surface,<sup>13</sup> which might explain the poorer performance of BPCSi-HNO<sub>3</sub>. The performance of the high silica-containing petcoke was over three times better than that of the petcoke with a lower silica content. Table S1 summarizes other recent studies on waste-derived carbon-based solid acid catalysts, with a series of materials prepared *via* the extensively studied sulfonation with sulfuric acid (H<sub>2</sub>SO<sub>4</sub>),<sup>10,22–24,26</sup> applied in the esterification reaction of octanoic or oleic acid with methanol. The difference between these carboxylic acids is their carbon chain length – oleic acid has 18 carbons and octanoic acid has only 8 carbons. The reactivity of these molecules, under the same conditions, decreases with the growth of the carbon chain due to steric hindrance.<sup>54</sup> So, a material that is active for the esterification of oleic acid should be more active for the reaction with octanoic acid. The estimated TOF for BPCSi-H<sub>2</sub>O<sub>2</sub> (308 h<sup>-1</sup>) and BPCSi-HNO<sub>3</sub> (192 h<sup>-1</sup>) were higher than the other waste-derived materials (between 1.3–68 h<sup>-1</sup>) and Amberlyst-15 (6 h<sup>-1</sup>). The fact that higher yields were reported for these catalysts<sup>21–24,26</sup> suggests that an excess of catalyst was used in the experiments. In addition, the reaction conditions were different with higher temperatures,<sup>21–23</sup> and a higher excess of alcohol.<sup>10,22,23</sup> Comparing the TOFs, the activity per site of the –SO<sub>3</sub>H groups originated from the inherent sulfur species in petcoke is better than that on the surface of Amberlyst-15, and that generated from sulfonated waste-derived carbon materials.

Even though the chemical functionalization of petcoke with hydrogen peroxide produced a more active material than the nitric acid treatment, the catalyst stability was not improved (Fig. 2 and S1). All samples, including Amberlyst-15, were deactivated after the first cycle. The deactivation was more pronounced for the petcoke-derived solid acid catalysts, for which the ester yields decreased to 10–15%. This loss in activity could be due to the partial deactivation of acid sites or the leaching of sulfonic acid sites into the solution.<sup>53,55</sup> The

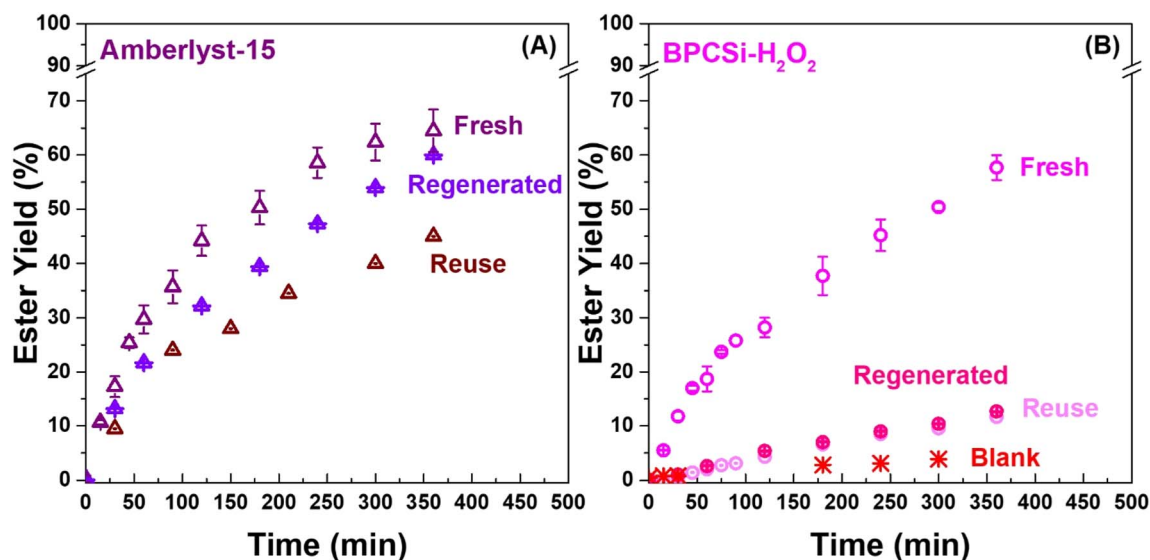


Fig. 2 Esterification of octanoic acid using fresh catalyst, spent catalyst (reuse curve), and catalyst after regeneration with H<sub>2</sub>SO<sub>4</sub>: Amberlyst-15 (A) and BPCSi-H<sub>2</sub>O<sub>2</sub> (B). Reaction conditions: MeOH : OA molar ratio = 10 : 1, 5 wt% of catalyst, 60 °C, 800 rpm.



performance of the regenerated catalysts is shown in Fig. 2. The deactivation of the Amberlyst-15 was reversible, and the majority of the activity could be recovered after acid treatment (Fig. 2A). The BPCSi-H<sub>2</sub>O<sub>2</sub> catalyst, however, was irreversibly deactivated, consistent with leaching of sulfonic acid sites (-SO<sub>3</sub>H) to the reaction solution. These results are consistent with those of other waste-derived carbon-based materials (Table S1). Sulfonated activated carbon from coffee residue<sup>25</sup> and from spent coffee grounds<sup>26</sup> lost ~70% of their initial activity after the 5th and 4th cycle, respectively, which is comparable with the loss for BPCSi-HNO<sub>3</sub> and BPCSi-H<sub>2</sub>O<sub>2</sub> after the 2nd use. Again, the reaction conditions and amounts of catalyst used were different in each study. Catalysts that are more porous may have active sites that take longer to diffuse through the material and into solution, as well as more active sites per mass of catalyst. For example, the loss of activity for the higher porosity materials (*i.e.*, activated carbon) made from potato peel,<sup>22</sup> ginger straw,<sup>23</sup> and garlic peel<sup>24</sup> was lower (~30% of the initial ones) after 4 to 5 cycles, but still significant. The sulfonated activated carbon produced from corn cob,<sup>21</sup> with the highest porosity and initial -SO<sub>3</sub>H concentration, had the lowest loss in activity of only 9%

**Table 1** Surface area, pore volume, and ash content of petcoke before and after functionalization with different agents

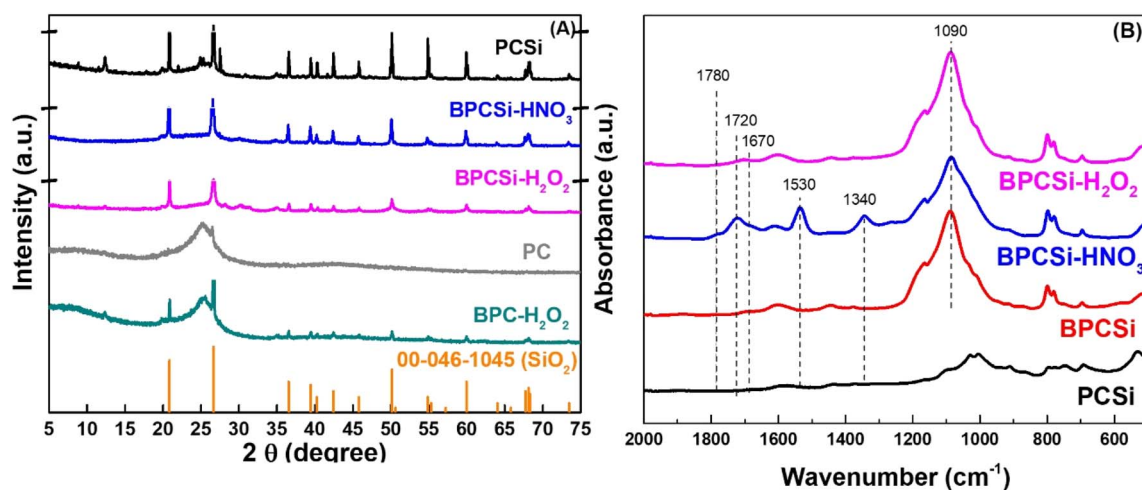
Sample	N <sub>2</sub> physisorption		Ash (wt%)
	Surface area (m <sup>2</sup> g <sup>-1</sup> )	Pore volume (cm <sup>3</sup> g <sup>-1</sup> )	
PCSi	1	0.008	47.5
BPCSi	23	0.068	48.2
BPCSi-HNO <sub>3</sub>	20	0.084	37.7
BPCSi-H <sub>2</sub> O <sub>2</sub>	26	0.082	44.7
PC	1	0.004	6.9
BPC	2	0.008	10.5
BPC-H <sub>2</sub> O <sub>2</sub>	2	0.007	14.2

after 8 cycles. The higher TOF observed for BPCSi-HNO<sub>3</sub> and BPCSi-H<sub>2</sub>O<sub>2</sub> may originate from the faster detachment of -SO<sub>3</sub>H groups from the surface for these lower porosity materials (Table 1). The rate of the homogeneous reaction contributes to the overall activity of the system but is not considered in most studies. In this study, it is not known if the sulfonic acid sites catalyze the esterification of octanoic acid before detaching from the surface. While the sites are not stable in methanol (leaching tests), in the presence of the reactant (octanoic acid), the material may act as a solid acid catalyst.

### Catalysts properties: differences between functionalization with HNO<sub>3</sub> and H<sub>2</sub>O<sub>2</sub>

The XRD patterns and FTIR spectra of petcoke before (PCSi) and after functionalization with HNO<sub>3</sub> and H<sub>2</sub>O<sub>2</sub> are shown in Fig. 3. In the XRD analysis, a broad peak at  $2\theta \approx 25.0^\circ$ , characteristic of petcoke, and corresponding to the graphite C (002) plane, is evident. In addition, several sharp peaks at  $2\theta = 21^\circ, 26.6^\circ, 36.6^\circ, 39.6^\circ, 40.4^\circ, 42.6^\circ, 45.6^\circ, 50.2^\circ, 55.1^\circ, 60.0^\circ$ , and  $67.9^\circ$  are present, which are attributed to the crystalline structure of SiO<sub>2</sub>. Moreover, PCSi and BPCSi-H<sub>2</sub>O<sub>2</sub> samples have a peak at  $2\theta = \sim 12^\circ$ , which is characteristic of the kaolinite mineral structure (Al<sub>2</sub>Si<sub>2</sub>O<sub>5</sub>(OH)<sub>4</sub>).<sup>56–58</sup> This mineral is usually present in silica sand composition<sup>56</sup> and has been reported as one of the minerals in Alberta oil sands.<sup>58</sup> The relative intensities of these peaks between samples are consistent with the ash values shown in Table 1, which contains the physical properties of the materials. Due to the high intensity of the SiO<sub>2</sub> diffraction peaks, it was not possible to observe any differences in the structure of the petcoke-derived catalysts after chemical functionalization. These results are consistent with previous findings<sup>59</sup> where a blend of petcoke and SiO<sub>2</sub> with a carbon-to-silicon elemental ratio of 1 : 1 had only SiO<sub>2</sub> peaks in the XRD pattern.

The surface area and pore volume were calculated from N<sub>2</sub> physisorption experiments. After the functionalization of PCSi,



**Fig. 3** Characterization of bulk and surface properties of petcoke-derived samples. (A) XRD patterns of catalysts and reference pattern for SiO<sub>2</sub> (PDF-5+ card of quartz). The peaks at  $21^\circ$  and  $26.6^\circ$  of PCSi, BPCSi-HNO<sub>3</sub>, and BPCSi-H<sub>2</sub>O<sub>2</sub> samples were truncated to enhance visibility of minor peaks; (B) ATR-FTIR spectra of high silica petcoke before and after functionalization with different agents.



there was an overall increase in the surface areas and pore volumes of the prepared samples (BPCSi-H<sub>2</sub>O<sub>2</sub> and BPCSi-HNO<sub>3</sub>), but these materials are much less porous than activated carbon. The decrease in ash content (Table 1 and Fig. S2) for BPCSi-HNO<sub>3</sub> (38%) when compared to PCSi (47.5%) was likely due to the removal of some mineral impurities during the treatment with nitric acid,<sup>13</sup> which is reinforced by the absence of a diffraction peak at  $2\theta = \sim 12^\circ$  for this sample. In contrast, no significant change in ash content was detected for BPCSi-H<sub>2</sub>O<sub>2</sub> (45%). Overall, there were not many differences between the physical properties of BPCSi-HNO<sub>3</sub> and BPCSi-H<sub>2</sub>O<sub>2</sub>, so the better activity for the latter is related to its chemical properties.

The S/C and O/C ratios were calculated using the ultimate elemental analysis of the catalysts (Table S2) – the PCSi sample had an S/C of 0.073 and O/C of 0.03. After functionalization with both hydrogen peroxide and nitric acid, there was a slight decrease in S/C ratio to 0.062 and 0.071 for BPCSi-HNO<sub>3</sub> and BPCSi-H<sub>2</sub>O<sub>2</sub>, respectively. The major change was in the O/C ratio of BPCSi-HNO<sub>3</sub> that increased to 0.47, while that of BPCSi-H<sub>2</sub>O<sub>2</sub> increased only marginally from 0.03 to 0.06. After the functionalization with HNO<sub>3</sub> and H<sub>2</sub>O<sub>2</sub>, the concentration of oxygen was expected to increase due to the formation of sulfonic acid sites and other oxygen-containing carbon functional groups, consistent with the increased O/C ratio for both samples.

The chemical states of functional groups after HNO<sub>3</sub> and H<sub>2</sub>O<sub>2</sub> functionalization were investigated by FTIR (Fig. 3B) and

XPS (Fig. 4). The FTIR spectrum of PCSi had a band centered at 1600 cm<sup>-1</sup> associated with poly-aromatic rings of hydrocarbon molecules and bands between 900–600 cm<sup>-1</sup>, which correspond to aromatic hydrogen.<sup>10</sup> After functionalization, the infrared spectra of BPCSi-HNO<sub>3</sub> and BPCSi-H<sub>2</sub>O<sub>2</sub> had absorption bands at 1670, 1720, and 1780 cm<sup>-1</sup> assigned to C=O stretching of –COOH, C–O, and C=O groups,<sup>60,61</sup> and vibration in the region of 1280–940 cm<sup>-1</sup>. Although these spectra have intense bands at  $\sim 1030$  cm<sup>-1</sup> and  $\sim 1200$  cm<sup>-1</sup>, which are related to the S=O stretching vibration of sulfonic groups,<sup>12,13,18</sup> the –SO<sub>3</sub>H formation could not be confirmed using only the FTIR technique. Silica has Si–O–Si vibration absorption between 1200–1000 cm<sup>-1</sup><sup>62,63</sup> that overlaps with the sulfonic acid functional group. The FTIR spectrum of silica is shown in Fig. S3, confirming that the peaks between 1200–1000 cm<sup>-1</sup> in the FTIR spectra of BPCSi and the functionalized materials are mainly due to the presence of SiO<sub>2</sub>. Also, after ball-milling treatment, the peak intensity of Si–O–Si stretching vibration increased when compared to PCSi (Fig. S3). This behaviour has also been reported for biochar made from sugarcane bagasse and wheat straw after ball-milling and is associated with the high silica content of these materials.<sup>64</sup> The acid-treated catalyst (BPCSi-HNO<sub>3</sub>) had the most intense bands of C=O stretching as well as bands at 1340 cm<sup>-1</sup> and 1530 cm<sup>-1</sup> associated with NO<sub>2</sub> stretching modes, indicating the presence of nitro groups.<sup>13</sup> Moreover, the FTIR results suggested that this material has

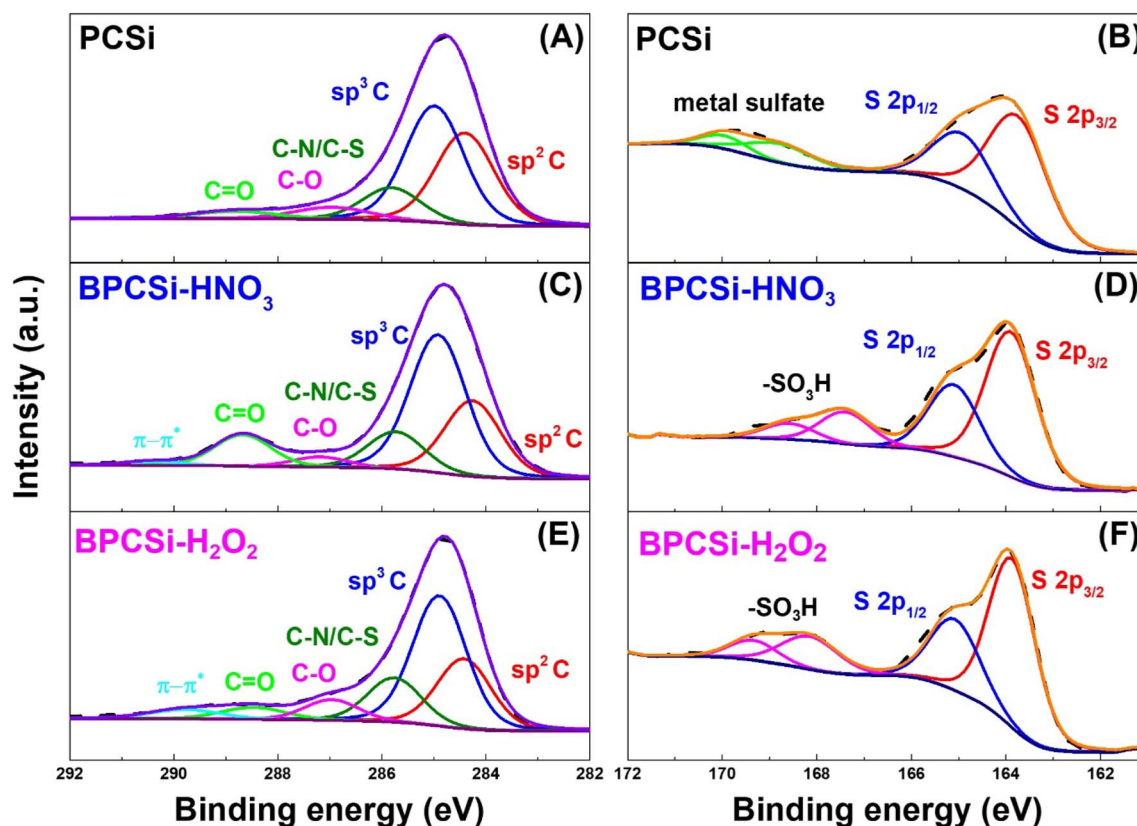


Fig. 4 Narrow scan XPS spectra for pristine high-silica petcoke (PCSi) and after ball-milling and functionalization with HNO<sub>3</sub> and H<sub>2</sub>O<sub>2</sub> C 1s (left-A, C, E) and S 2p (right-B, D, F).



more oxygen-containing groups on its surface than BPCSi-H<sub>2</sub>O<sub>2</sub>, which may explain the poorer activity of BPCSi-HNO<sub>3</sub>.

The surface properties of the petcoke-derived catalysts were also analyzed by X-ray photoelectron spectroscopy (XPS). The surface elemental composition was calculated in weight percentage from the survey spectra (Fig. S4), and the results are shown in Table S3. In addition to carbon, oxygen, nitrogen, and sulfur, two other elements were detected on the surface of petcoke-derived samples: silicon and aluminium. PCSi contained 16.6 wt% Si. After treatment, BPCSi-HNO<sub>3</sub> had an increase in the nitrogen content, as expected, and an increase in the O/C ratio (from 0.50 to 0.76) while the S/C ratio did not significantly change. The significant decrease in the Si/C and Al/C ratio after acid functionalization is consistent with mineral impurity removal (*i.e.*, lower ash content, Table 1). The petcoke functionalization with hydrogen peroxide, on the other hand, resulted in a material (BPCSi-H<sub>2</sub>O<sub>2</sub>) with a lower sulfur content on the surface, demonstrated by the decrease in S/C ratio. This decrease is likely due to partial oxidative desulfurization promoted by H<sub>2</sub>O<sub>2</sub> during preparation,<sup>65,66</sup> and an increase in the O/C ratio (from 0.50 to 0.89). The greater O/C ratio for BPCSi-H<sub>2</sub>O<sub>2</sub> was associated with the higher SiO<sub>2</sub> content in this sample, as demonstrated by its ash percentage and more intense peaks in FTIR related to Si–O–Si vibration (Fig. 3B). In addition, the higher Si/C and Al/C ratios for this sample were related to a reduction in the carbon content after functionalization, and not to mineral removal.

The fitting of peaks for the XPS narrow scan spectra are shown in Fig. 4 and were used to determine the oxidation states of carbon and sulfur in the samples. The C 1s spectra (Fig. 4A, C, and E) contained five peaks for the PCSi sample, centred at 284.9 eV, 284.4 eV, 285.8 eV, 287.0 eV, and 288.7 eV corresponding to sp<sup>3</sup> carbon, sp<sup>2</sup> carbon, carbon hybridized with N and S, C–O, and C=O,<sup>13,43–45</sup> respectively. After functionalization with HNO<sub>3</sub> and H<sub>2</sub>O<sub>2</sub>, one additional peak was detected centered at ~290 eV corresponding to π–π\* transition (O=C–O), indicating the addition of oxygen-containing carbon groups on the surface. The percentage of carbon in the C–O bond changed from 5.6% for PCSi to 3.4% for BPCSi-HNO<sub>3</sub>, and 7.1% for BPCSi-H<sub>2</sub>O<sub>2</sub>. The C=O bond content increased from 3.8% in petcoke (PCSi) to 10.7% for BPCSi-HNO<sub>3</sub> and 5.4% for BPCSi-H<sub>2</sub>O<sub>2</sub> (Table S4). The sulfur species are shown in Fig. 4B, D, and F, and the PCSi S 2p spectrum was fitted into four peaks, corresponding to two different species. The peaks centred at 163.9 eV and 165.0 eV are attributed to C–S–C (2p<sub>3/2</sub> and 2p<sub>1/2</sub>, respectively), characteristic of sulfur in the thiophene structure,<sup>46,47</sup> while the peaks at ~169 eV and 170 eV are the binding energy of SO<sub>4</sub><sup>2-</sup> species related to metal sulfates in petcoke.<sup>48–50</sup> After functionalization, the sulfate peak disappeared, and two additional peaks at 167.7 eV and 168.8 eV related to –SO<sub>3</sub>H groups<sup>44,45,51</sup> appeared in the S 2p high-resolution spectra of BPCSi-HNO<sub>3</sub> and BPCSi-H<sub>2</sub>O<sub>2</sub>. This result confirmed that hydrogen peroxide can access and convert the inherent sulfur species in petcoke into sulfonic acid sites, in the same way as has been demonstrated for nitric acid.<sup>12</sup> Even though BPCSi-HNO<sub>3</sub> had a higher concentration of oxygen groups, as demonstrated by elemental analysis, the XPS spectrum was not

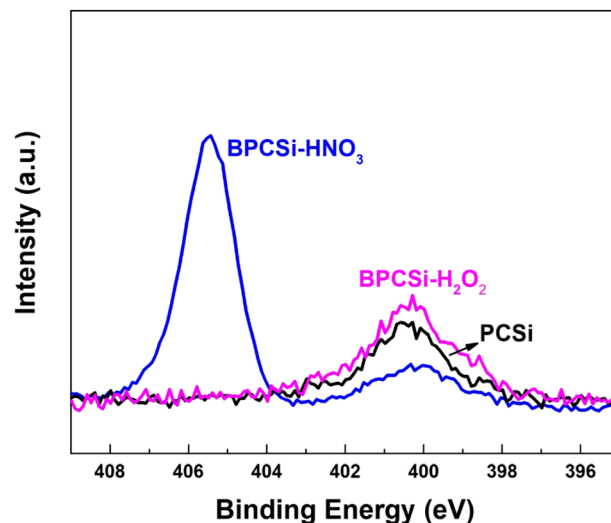


Fig. 5 Narrow scan XPS spectra of N 1s for PCSi and petcoke-derived solid acid catalysts.

significantly different from BPCSi-H<sub>2</sub>O<sub>2</sub>. The main difference was in the C=O intensity, which is higher for the former in accordance with the FTIR results.

The N 1s narrow scan spectra of PCSi and chemically treated samples are shown in Fig. 5. Before (PCSi) and after functionalization with hydrogen peroxide (BPCSi-H<sub>2</sub>O<sub>2</sub>), the amount of nitrogen on the surface was ~1 wt% (Table S3), and one broad peak centered at ~400.0 eV can be seen and is reported as N–O, pyrrolic or pyridinic structures.<sup>67,68</sup> The nitric acid treatment resulted in the increase of nitrogen content on the surface (3.6 wt%) due to the formation of nitro groups (–NO<sub>2</sub>) demonstrated by the additional peak at the binding energy of 405.5 eV<sup>13,68</sup> and confirmed by the FTIR results of the BPCSi-HNO<sub>3</sub> samples (Fig. 3B).

The number of total surface acidic groups on the samples was calculated using the Boehm titration method, and the concentration of sulfonic acid sites was estimated based on the XPS results, as shown in Table 2. Initially, PCSi had no sulfonic acid sites on the surface, based on XPS, and a low total acidity of 0.30 mmol g<sup>-1</sup>. After the ball milling treatment, the total acidity of the resulting sample (BPCSi) increased to 0.67 mmol g<sup>-1</sup>. This increase in the total acidity was due to the exposure of silanol groups (Si–OH) on the surface of silica sand, which is a common effect when in contact with water<sup>69</sup> – isopropanol applied in the ball-milling process is hygroscopic. The ATR-FTIR spectra of BPCSi (Fig. S5B) had a band at ~3450 cm<sup>-1</sup> attributed to the vibration of Si–OH groups.<sup>70</sup> Functionalization with nitric acid generated a catalyst (BPCSi-HNO<sub>3</sub>) with an increased total acidity of ~5 mmol g<sup>-1</sup> and a 0.14 mmol g<sup>-1</sup> concentration of –SO<sub>3</sub>H sites. The petcoke functionalization with hydrogen peroxide resulted in a material (BPCSi-H<sub>2</sub>O<sub>2</sub>) with a much lower total acidity (1.29 mmol g<sup>-1</sup>), and a lower density of sulfonic acid sites (0.09 mmol g<sup>-1</sup>).

The XPS C 1s and S 2p narrow scan spectra of the spent catalysts produced from a high-silica petcoke are shown in Fig. 6. After use, the sulfonic acid sites on the surface of BPCSi-



Table 2 Acidity and turnover frequency (TOF) of Amberlyst-15, petcoke, and functionalized petcoke

Sample	Acidity (mmol g <sup>-1</sup> )		Fresh catalyst		Reused catalyst
	[-SO <sub>3</sub> H] <sup>a</sup>	Total <sup>b</sup>	Ester yield <sup>c</sup> (%)	TOF (h <sup>-1</sup> )	Ester yield <sup>c</sup> (%)
Amberlyst-15	4.70 <sup>f</sup>	4.70 <sup>f</sup>	64.5	6	44.8
PCSi	0	0.30 ± 0.14	1.5	— <sup>d</sup>	— <sup>e</sup>
BPCSi	0	0.67 ± 0.06	— <sup>e</sup>	— <sup>d</sup>	— <sup>e</sup>
BPCSi-HNO <sub>3</sub>	0.14	5.01 ± 0.21	45.1	192	16.1
BPCSi-H <sub>2</sub> O <sub>2</sub>	0.09	1.29 ± 0.26	54.2	308	17.2
PC	0	0.24 ± 0.04	— <sup>e</sup>	— <sup>d</sup>	— <sup>e</sup>
BPC	0	0.33 ± 0.06	— <sup>e</sup>	— <sup>d</sup>	— <sup>e</sup>
BPC-H <sub>2</sub> O <sub>2</sub>	0.20	0.97 ± 0.04	17.3	42	— <sup>e</sup>

<sup>a</sup> Estimated from XPS spectra. <sup>b</sup> Estimated by Boehm titration. <sup>c</sup> After 6 h. <sup>d</sup> Not calculated for materials without -SO<sub>3</sub>H acidity. <sup>e</sup> Not determined. <sup>f</sup> From the product data sheet of the manufacturer. In Amberlyst-15 all the acidity is attributed to acid sulfonic acid groups.

HNO<sub>3</sub>-spent were completely gone, as demonstrated by the absence of -SO<sub>3</sub>H peaks in the narrow scan S 2p spectra of this sample (Fig. 6B) and a decrease in the sulfur content on the surface of this catalyst from 2.5 wt% in the fresh sample to 0.6 wt% in the spent material (Table S5), demonstrating the complete leaching of -SO<sub>3</sub>H sites during the esterification reaction. The BPCSi-H<sub>2</sub>O<sub>2</sub>-spent, on the other hand, still presented peaks at ~168.0 eV and ~169.0 eV corresponding to sulfonic acid sites. However, the total sulfur concentration decreased considerably from 1.4 wt% to 0.7 wt% (Table S5), also indicating leaching of the active sites, but not completely as for BPCSi-HNO<sub>3</sub>-spent. For comparison, the density of sulfonic acid

sites in the fresh sample was 0.09 mmol g<sup>-1</sup>, which decreased to 0.03 mmol g<sup>-1</sup> for the spent material. The C 1s narrow scan spectra of the spent samples were similar to the fresh ones, and the distribution of functional groups did not significantly change (Table S6).

Although BPCSi-H<sub>2</sub>O<sub>2</sub> had a lower concentration of -SO<sub>3</sub>H sites than BPCSi-HNO<sub>3</sub>, the former had a higher activity per site (TOF) of 308 h<sup>-1</sup> versus 192 h<sup>-1</sup> for the nitric acid catalyst. The lower activity of BPCSi-HNO<sub>3</sub> can be attributed to its higher total acidity, which includes contributions from acidic oxygen-containing carbon groups such as lactonic and carboxylic functionalities,<sup>71</sup> as demonstrated by the FTIR bands at 1670,

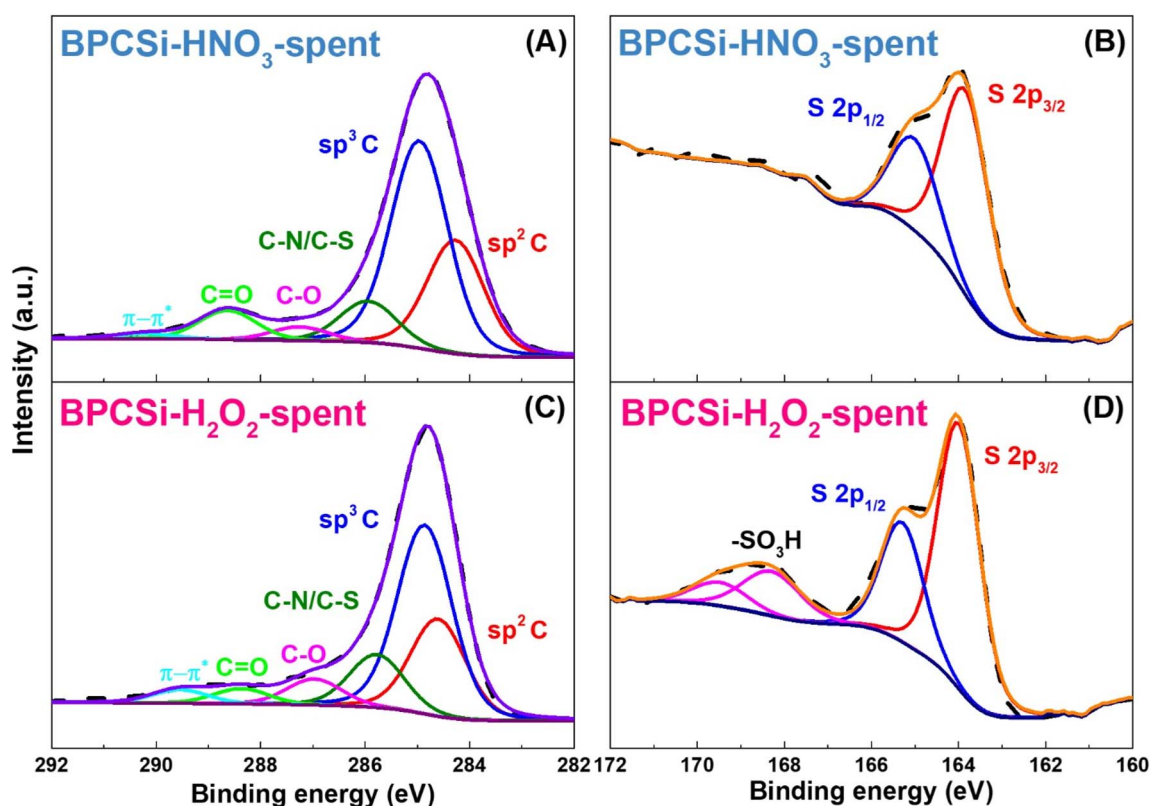


Fig. 6 Narrow scan XPS spectra for spent catalysts prepared from a high-silica petcoke C 1s (left-A and C) and S 2p (right-B and D).



1720, and 1789  $\text{cm}^{-1}$  (Fig. 3), as well as a small contribution of the silanol groups. The oxygen-containing carbon groups hinder the reaction by promoting the adsorption of reactants and products through hydrogen bonding interactions, particularly with methanol, octanoic acid, and the ester formed, causing steric hindrance.<sup>13,72,73</sup> In addition, the TOF calculation was done in terms of the sulfonic acid site concentration only, because these oxygen-containing carbon groups present weak acidity that cannot protonate the carboxyl group of the octanoic acid, which is the first step of the reaction. This protonation requires strong Brønsted acid sites; therefore, only  $-\text{SO}_3\text{H}$  groups are considered truly active sites for the reaction.<sup>55,74–78</sup>

Therefore, the functionalization with hydrogen peroxide resulted in a more active catalyst than the functionalization with nitric acid because of the formation of fewer oxygen-containing carbon groups on the surface of  $\text{BPCSi-H}_2\text{O}_2$  when compared to  $\text{BPCSi-HNO}_3$ . This observation supports the hypothesis that enhancing the activity of petcoke-derived catalysts is highly dependent on a reduction in the number of oxygen-containing carbon groups formed during material preparation. Specifically, these groups seem to negatively influence the catalyst activity to a much greater extent than the positive effect of a higher concentration of sulfonic acid sites. Ideally, to achieve even better catalytic performance with these petcoke-derived materials, it is desirable to keep the total acidity as low as the one obtained for  $\text{BPCSi-H}_2\text{O}_2$ , while promoting the formation of additional  $-\text{SO}_3\text{H}$  groups. The maximum possible concentration of sulfonic acid sites was calculated to be  $0.83 \text{ mmol g}^{-1}$ , assuming that all sulfur in  $\text{PCSi}$  was converted to  $-\text{SO}_3\text{H}$ . Thus, there is potential for improvement, as only about 10% of this maximum value has been achieved with  $\text{BPCSi-H}_2\text{O}_2$ .

### Catalyst properties: effect of initial $\text{SiO}_2$ content on the petcoke feedstock

The presence of silica in the petcoke feedstock arises from different stages of production, processing, and handling. During the coking step, the equipment used (siliceous refractory linings) and silica-based additives applied in the process might contribute to the silica content in the final product.<sup>34,35</sup> More than likely, however, the major contribution comes from handling and storage. Petcoke is largely stockpiled in an open field,<sup>29–31</sup> in direct contact with the ground. If the storage area is rich in silica sand, and precautions are not taken,  $\text{SiO}_2$  can get mixed with petcoke. The area from which the petcoke used in this study was obtained (Fort McMurray – Alberta, Canada) is abundant in silica sand.<sup>79</sup> So, a batch of petcoke from the bottom of the pile can have a much higher fraction of sand mixed with it.

The esterification results (Fig. 1) showed that  $\text{BPCSi-H}_2\text{O}_2$ , generated from a feedstock containing a higher silica content ( $\text{PCSi}$ ), was much more active than  $\text{BPC-H}_2\text{O}_2$ . Both samples were prepared in the same way, and silica ( $\text{SiO}_2$ ) is generally chemically inert at the reaction conditions used in this study.<sup>80</sup> Thus, the results suggest that while the  $\text{SiO}_2$  component does not directly catalyze the esterification reaction, it contributes to

enhancing the activity of the resulting catalyst. To investigate how silica sand is affecting the overall performance of petcoke-derived samples in the esterification reaction, the structural properties of hydrogen peroxide-treated materials were investigated by XRD (Fig. 3).  $\text{PC}$  has the characteristic broad peak at  $2\theta = 25^\circ$  related to graphite C (002), and a sharp peak at  $26.6^\circ$   $2\theta$  corresponding to microcrystals of  $\text{SiO}_2$ . After ball milling and  $\text{H}_2\text{O}_2$  functionalization ( $\text{BPC-H}_2\text{O}_2$ ), the  $\text{SiO}_2$  crystalline structure, as well as the kaolinite characteristic peak, became more evident and reflect the increase in the ash content from 6.9 wt% for  $\text{PC}$  to 14.2 wt% for  $\text{BPC-H}_2\text{O}_2$  (Table 1), suggesting that part of the carbon is decomposed during the chemical functionalization process. The difference between the XRD patterns of  $\text{BPC-H}_2\text{O}_2$  and  $\text{BPCSi-H}_2\text{O}_2$  is mainly the intensity of  $\text{SiO}_2$  peaks, which does not explain the better performance of the latter.

The ultimate elemental analysis of  $\text{PCSi}$  and  $\text{PC}$  (Table S2) revealed similar S/C (0.073 and 0.076, respectively) and O/C (0.03 and 0.01, respectively) ratios for both samples, corroborating the existence of  $\text{SiO}_2$  as a physical mixture in  $\text{PCSi}$ . The surface chemical groups of  $\text{PC}$  and  $\text{BPC-H}_2\text{O}_2$  were investigated by FTIR-ATR (Fig. S5) and XPS C 1s and S 2p high-resolution spectra (Fig. S6). The functional groups existing on the surface of the materials before and after functionalization with hydrogen peroxide were the same as those of  $\text{PCSi}$  and  $\text{BPCSi-H}_2\text{O}_2$ , respectively. Even the distribution of carbon species ( $\text{sp}^2$  C,  $\text{sp}^3$  C, C–N/C–S, C–O, C=O, and  $\pi$ – $\pi^*$ ) and sulfur species (C–S–C, C=S,  $-\text{SO}_3\text{H}$ , and  $\text{SO}_4^{2-}$ ) in Table S4 were quite similar for  $\text{BPC-H}_2\text{O}_2$  and  $\text{BPCSi-H}_2\text{O}_2$ , suggesting that  $\text{SiO}_2$  has no influence on the oxidation pathway of petcoke. After functionalization with hydrogen peroxide, the S/C ratio decreased and the O/C, Si/C, and Al/C ratios increased for both  $\text{BPCSi-H}_2\text{O}_2$  and  $\text{BPC-H}_2\text{O}_2$  (Table S3).

Comparing the number of total surface acidic groups and the density of sulfonic acid sites (Table 2) of the hydrogen peroxide-treated samples,  $\text{BPC-H}_2\text{O}_2$  showed a total acidity of  $0.97 \text{ mmol g}^{-1}$  and sulfonic acid sites concentration of  $0.20 \text{ mmol g}^{-1}$ . The  $\text{BPCSi-H}_2\text{O}_2$  sample had values of  $1.29 \text{ mmol g}^{-1}$  and  $0.09 \text{ mmol g}^{-1}$  (total and  $-\text{SO}_3\text{H}$ , respectively). The total acidity, which is mainly related to the oxygen-containing carbon groups, of both samples is similar, and the density of sulfonic acid sites is twice as high for  $\text{BPC-H}_2\text{O}_2$ . From these results, it was expected that the activity of this catalyst would be much higher than  $\text{BPCSi-H}_2\text{O}_2$ . On the contrary, the TOF of  $\text{BPCSi-H}_2\text{O}_2$  is sevenfold better than  $\text{BPC-H}_2\text{O}_2$  ( $308 \text{ h}^{-1}$  compared to  $42 \text{ h}^{-1}$ , respectively). Although the concentration of sulfonic acid sites in  $\text{BPC-H}_2\text{O}_2$  is higher than in  $\text{BPCSi-H}_2\text{O}_2$ , normalization by the  $\text{SiO}_2$  content in each sample (ash% in Table 1) yields comparable values, indicating that the presence of silica did not significantly affect the resulting concentration of sulfonic groups. Also, the reuse test of  $\text{BPC-H}_2\text{O}_2$  (Fig. S1), XPS S 2p spectrum of the spent catalyst (Fig. S7), and sulfur content on its surface (Table S5) showed a leaching of  $-\text{SO}_3\text{H}$  groups after the first cycle. The fresh catalyst had a sulfur content of 3.0 wt% that decreased to 1.9 wt% in the spent material, representing a significant loss in sulfonic acid site concentration that resulted in a density of  $0.06 \text{ mmol g}^{-1}$  in  $\text{BPC-H}_2\text{O}_2$ -spent – much lower than the



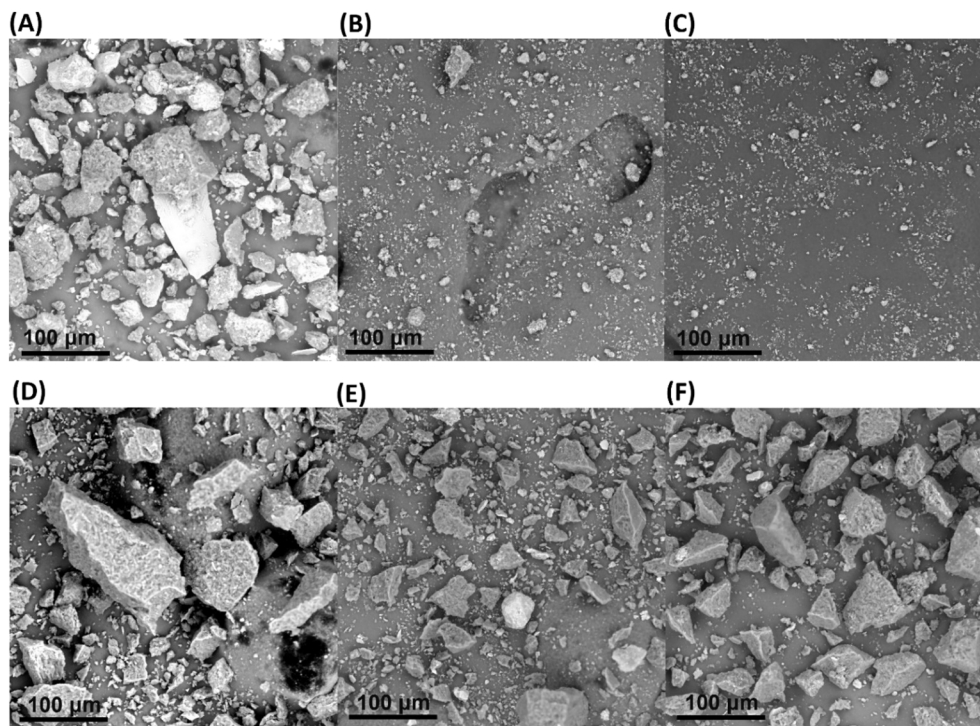


Fig. 7 SEM images PCSi (A), BPCSi (B), BPCSi-H<sub>2</sub>O<sub>2</sub> (C), PC (D), BPC (E), and BPC-H<sub>2</sub>O<sub>2</sub> (F). Brighter particles are silica sand (SiO<sub>2</sub>).

0.20 mmol g<sup>-1</sup> in the fresh material (BPC-H<sub>2</sub>O<sub>2</sub>). If all the sulfonic acid sites were on the surface of BPC-H<sub>2</sub>O<sub>2</sub> and were leached to the reaction solution, this material would have a higher activity than BPCSi-H<sub>2</sub>O<sub>2</sub> – considering the higher sulfonic acid sites density of the former. Therefore, a fraction of -SO<sub>3</sub>H groups in the BPC-H<sub>2</sub>O<sub>2</sub> samples was likely to be inaccessible for the reaction of octanoic acid molecules. Comparing the total acidity of the BPCSi (0.67 mmol g<sup>-1</sup>) and BPC (0.33 mmol g<sup>-1</sup>), however, the presence of a higher content of SiO<sub>2</sub> generates more silanol groups that have a weak acidity, behaving as hydrogen-bond donors<sup>81,82</sup> that facilitate the protonation of the octanoic acid and methanol, but the silanol groups alone were not active in the esterification reaction, as the ester yield of BPCSi after 6 h was only 1.5% (Fig. 1). Therefore, the hydrogen-bond donation effect of silanol groups alone could not be responsible for such a difference in the performance between BPCSi-H<sub>2</sub>O<sub>2</sub> and BPC-H<sub>2</sub>O<sub>2</sub>.

The SEM images of PCSi, BPCSi, BPCSi-H<sub>2</sub>O<sub>2</sub>, PC, BPC, and BPC-H<sub>2</sub>O<sub>2</sub> are shown in Fig. 7. Initially, PCSi and PC were sieved into particles of 20–100 μm (Fig. 7A and D, respectively). After the ball-milling process, a significant reduction of particle size was noticeable for the feedstock containing a higher silica content (Fig. 7B and C). This effect was not observed at the same level for BPC and BPC-H<sub>2</sub>O<sub>2</sub> (Fig. 7E and F). The production of smaller particles can also be confirmed by the increase in surface area (SA) after ball-milling and functionalization (Table 1) of PCSi (SA = 1 m<sup>2</sup> g<sup>-1</sup>) to BPCSi-H<sub>2</sub>O<sub>2</sub> (SA = 26 m<sup>2</sup> g<sup>-1</sup>), while the transformation of PC to BPC-H<sub>2</sub>O<sub>2</sub> did not impact the surface area (~1 m<sup>2</sup> g<sup>-1</sup>). Silica is a hard material that enhances the efficiency of ball-milling<sup>10,64,83</sup> by promoting particle size reduction, which may also explain the improved activity of

BPCSi-H<sub>2</sub>O<sub>2</sub>, as smaller particles are associated with reduced diffusion limitations.<sup>84</sup> Therefore, the promotional effect of SiO<sub>2</sub> in petcoke-derived solid acid catalysts was attributed to a combined effect of enhancing particle size reduction during mechanical treatment (ball milling) and formation of silanol groups that act as hydrogen-bond donors for the reactants, helping the catalysis – effects that are less pronounced in samples with a small content of SiO<sub>2</sub>.

Although the stability of petcoke-derived solid acid catalysts was limited, valuable insights were gained regarding the functionalization of petcoke with hydrogen peroxide and nitric acid. These findings can be applied broadly to the production of carbon materials, enabling tailored control of surface oxygen-containing groups to meet specific application requirements. If a higher degree of oxidation is required on the surface, HNO<sub>3</sub> can be applied, while H<sub>2</sub>O<sub>2</sub> will generate fewer oxygen-containing carbon groups on the surface. Moreover, the leaching of sulfonic acid sites observed for these materials is promoted by water in solution. Hence, applying these catalysts to gas-phase reactions, such as dehydration, may offer improved site stability. Additionally, the SiO<sub>2</sub> promoting effect emphasizes the importance of considering the impact of feedstock contamination on the valorization of waste materials. In the context of petcoke catalysts, the observed effects were advantageous for a better activity, however, this outcome may not necessarily apply to other catalytic systems.

## Conclusions

Hydrogen peroxide was used to oxidize inherent sulfur species in petcoke, producing sulfonic acid sites (-SO<sub>3</sub>H) that act as



active groups for esterification. Hydrogen peroxide is a more environmentally friendly functionalization agent than  $\text{HNO}_3$ . The  $\text{H}_2\text{O}_2$ -prepared catalyst (BPCSi- $\text{H}_2\text{O}_2$ ) achieved a 54% ester yield, higher than BPCSi- $\text{HNO}_3$  (45%), though both were less active than the commercial Amberlyst-15 catalyst (65%). Petcoke-derived catalysts were unstable and not regenerable, a common issue for carbon-based solid acid catalysts. BPCSi- $\text{H}_2\text{O}_2$  had fewer oxygen-containing carbon groups ( $\sim 1 \text{ mmol g}^{-1}$ ) than BPCSi- $\text{HNO}_3$  ( $\sim 5 \text{ mmol g}^{-1}$ ), which improved its activity. Despite a lower  $-\text{SO}_3\text{H}$  concentration ( $0.09 \text{ mmol g}^{-1}$  vs.  $0.14 \text{ mmol g}^{-1}$ ), BPCSi- $\text{H}_2\text{O}_2$  had a higher TOF than BPCSi- $\text{HNO}_3$  ( $308$  vs.  $192 \text{ h}^{-1}$ ). Increasing sulfonic acid site concentration while minimizing oxygen-containing carbon groups could enhance catalyst performance.

A higher initial  $\text{SiO}_2$  content in petcoke helped improving the catalytic performance, as silica reduced particle size during ball milling and generated more silanol groups that helped the catalytic process. The promoting effect of  $\text{SiO}_2$  illustrates the importance of not overlooking impurities in the feedstock when working with waste-derived materials, because they can play an important role in the overall process.

## Author contributions

N. M. C.: data curation; formal analysis; investigation; methodology; validation; visualization; writing – original draft. J. M. H.: conceptualization; 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 has been included as part of the supplementary information (SI). Supplementary information: Fig. S1: ester yield of petcoke-derived solid acid catalysts treated with  $\text{H}_2\text{O}_2$  for two consecutive runs. Reaction conditions: methanol : octanoic acid molar ratio of 10 : 1, 5 wt% of catalyst,  $60 \text{ }^\circ\text{C}$ , 800 rpm; Fig. S2: thermogravimetric analysis results: (A) TGA profile of PCSi, BPCSi- $\text{HNO}_3$ , and BPCSi- $\text{H}_2\text{O}_2$ , and (B) TGA profile of PC and BPC- $\text{H}_2\text{O}_2$ . Collected at  $20 \text{ }^\circ\text{C min}^{-1}$  to  $750 \text{ }^\circ\text{C}$  in a  $50 \text{ mL min}^{-1}$  flow of air; Fig. S3: ATR-FITR spectra of PCSi, ball-milled PCSi (BPCSi), and silica sand sample ( $\text{SiO}_2$ ) with particle size range 300–600  $\mu\text{m}$ ; Fig. S4: XPS survey spectra of PC, PCSi, and chemically treated samples; Fig. S5: ATR-FTIR (A) comparing PCSi versus PC, and BPCSi- $\text{H}_2\text{O}_2$  versus BPC- $\text{H}_2\text{O}_2$  in the 500–2000  $\text{cm}^{-1}$  region; (B) full range (4000–500  $\text{cm}^{-1}$ ) for all samples; Fig. S6: high-resolution XPS spectra of petcoke-derived samples before (PC and PCSi) and after treatment with hydrogen peroxide (BPCSi- $\text{H}_2\text{O}_2$  and BPC- $\text{H}_2\text{O}_2$ ), C 1s (left) and S 2p (right); Table S1: esterification reactions over different waste-derived carbon-based solid acid catalysts and Amberlyst-15; Table S2: elemental analysis of petcoke and petcoke-derived catalysts; Table S3: surface elemental analysis calculated from

XPS survey spectrum (wt%); Table S4: Area% of deconvoluted signals from XPS high-resolution data. See DOI: <https://doi.org/10.1039/d5ra05900c>.

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