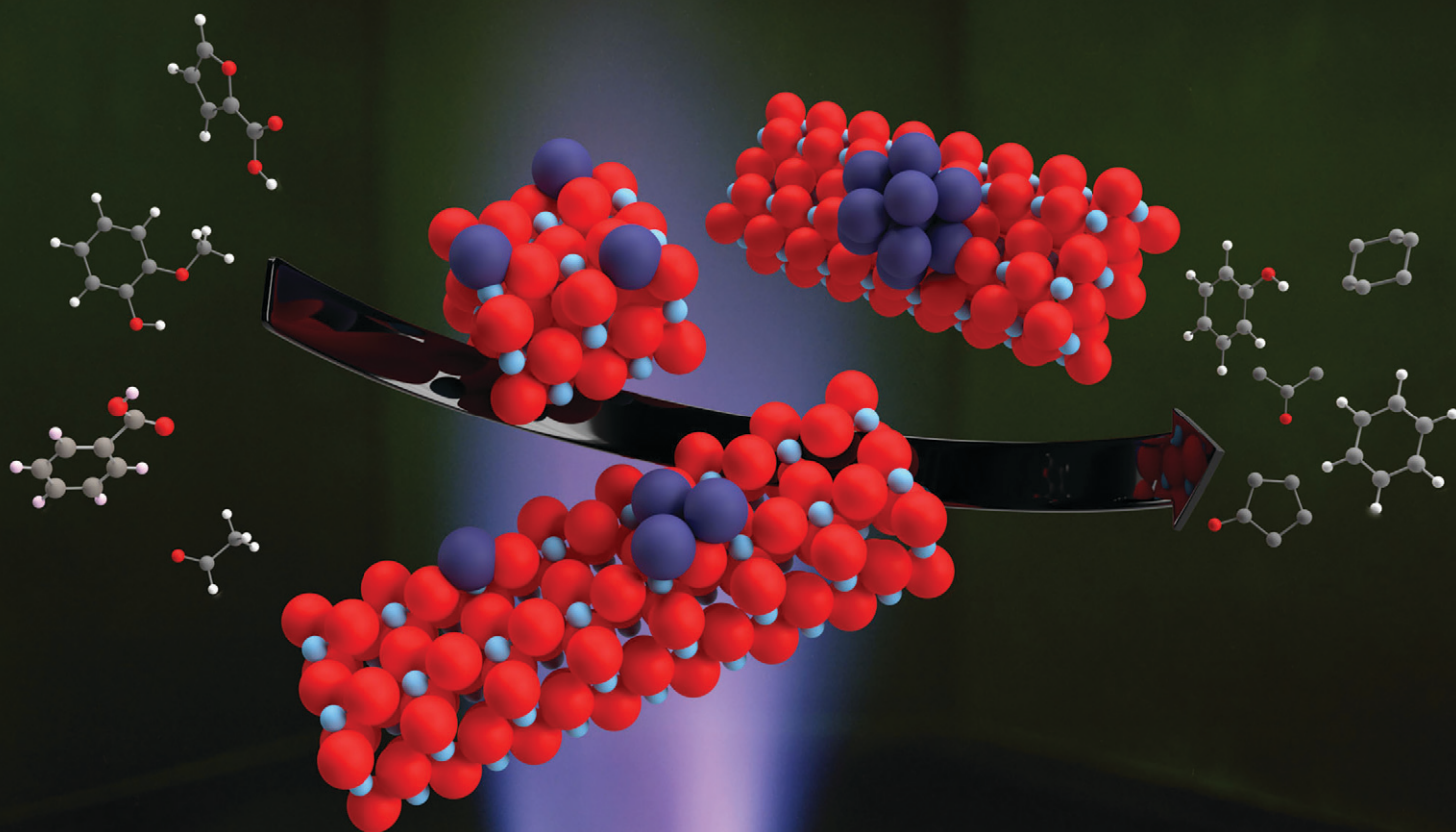


Catalysis Science & Technology

Volume 13
Number 17
7 September 2023
Pages 4885–5142

rsc.li/catalysis



ISSN 2044-4761

PAPER

Frederick G. Baddour *et al.*
Synthesis, performance evaluation, and economic assessment
of tailored Pt/TiO₂ catalysts for selective biomass vapour
upgrading *via* a scalable flame spray pyrolysis route

Cite this: *Catal. Sci. Technol.*, 2023,
13, 4941

Synthesis, performance evaluation, and economic assessment of tailored Pt/TiO₂ catalysts for selective biomass vapour upgrading *via* a scalable flame spray pyrolysis route†

Anurag Kumar,^a Rongyue Wang,^b Kellene A. Orton,^a Kurt M. Van Allsburg,^{iD}^a
Calvin Mukarakate,^{iD}^a Evan C. Wegener,^c Qiyuan Wu,^a Susan E. Habas,^{iD}^a
Krzysztof Z. Pupek,^b Joseph A. Libera^b and Frederick G. Baddour^{iD}^{*a}

Commercialization of next-generation nanostructured catalysts for biomass conversion is largely limited by the unique challenges associated with their large scale manufacture. In this work, we present the scalable, continuous synthesis of nanostructured Pt/TiO₂ catalysts in a single step *via* flame-spray pyrolysis (FSP) and demonstrate their utility in the upgrading of biomass vapours *via* catalytic fast pyrolysis (CFP). Spectroscopic and microscopic characterization of the catalysts prepared *via* FSP and conventional incipient wetness impregnation (IW) demonstrated that the FSP method enabled scalable production of Pt/TiO₂ with distinct Pt speciation (single atoms, clusters, and nanoparticles) that could be tuned by varying the Pt loading. This compositional control by FSP resulted in catalytic activity for upgrading of pine pyrolysis vapours to aromatics, oxygenates, and alkenes that was comparable to the IW analogues' performance, with the addition of more control over the product slate by modifying the FSP synthesis conditions. Specifically, the FSP-synthesized catalysts exhibited decreased hydrogenation performance and greater selectivity towards partially deoxygenated products, aromatics, (cyclo)alkanes, and (cyclo)alkenes with total hydrocarbon yields similar to IW-synthesized catalysts at equivalent Pt loadings. A preliminary cost analysis comparing the FSP and IW processes identified the major cost drivers, including solvent usage, process gas requirements, and Pt utilization, that can be addressed to improve the economic viability of the FSP synthesis developed herein.

Received 20th April 2023,
Accepted 9th July 2023

DOI: 10.1039/d3cy00550j

rsc.li/catalysis

Introduction

Catalytic upgrading of biomass to drop-in liquid fuels represents a significant research challenge. In particular, it requires the removal of oxygen from a complex mixture of oxygenated molecules (*e.g.*, phenols, alcohols, syringols, ketones, and aldehydes) due to the detrimental effects of oxygenated species on fuel properties such as a low heating value, high acidity, and chemical instability.^{1–3} *Ex situ* catalytic fast pyrolysis (CFP) is a promising conversion

technology wherein the vapours of pyrolyzed biomass are passed over a catalyst at high temperature and near-ambient pressure (450 °C, 1 atm) to generate a stabilized bio-oil with lower oxygen content that is more suitable for down-stream upgrading to hydrocarbon fuels or high value chemicals. However, techno-economic analyses of this process have revealed the need for further development of next-generation catalysts if the goal of sustainable, low-cost CFP-derived biofuels is to be realized.³ Notably, several key catalytic properties have been identified, including the importance of bifunctionality (*i.e.*, combination of acidic and metallic sites), such that H₂ can be activated at the low-pressure-high-temperature conditions of CFP with selectivity that favours cleavage of C–O bonds over C–C bonds.¹

Bifunctional catalysts possessing metallic and acidic sites such as noble metals (*e.g.*, Pt, Pd, Ru)^{3–7} supported on acidic reducible oxide supports (*e.g.*, TiO₂, ZrO₂) have been identified as effective catalysts for biomass pyrolysis vapour upgrading at lab-scale because of their ability to activate hydrogen and perform deoxygenation reactions.^{3–7}

^a Catalytic Carbon Transformation and Scale-up Center, National Renewable Energy Laboratory, 15013 Denver West Parkway, Golden, Colorado 80401-3305, USA. E-mail: Frederick.Baddour@nrel.gov

^b Energy Systems Division, Argonne National Laboratory, Argonne, Illinois 60439, USA

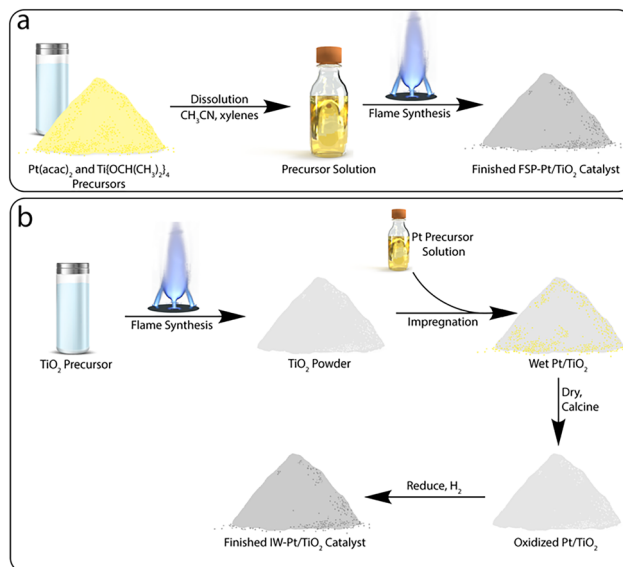
^c Chemical Sciences and Engineering Division, Argonne National Laboratory, Lemont, IL, 60439 USA

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3cy00550j>



Traditionally, these catalysts are synthesized by incipient wetness impregnation (IW), co-precipitation, sol-gel, or hydrothermal methods for their ease of preparation.^{8–10} However, recent work has highlighted the performance enhancements that are attainable in the catalytic upgrading of bio-derived feedstocks by employing advanced synthetic methods such as solution phase synthesis^{11,12} and strong electrostatic adsorption^{3,13,14} that afford more control over catalyst physical properties (e.g., particle size, dispersion, acid-site strength, and density).^{15,16} However, the realization and deployment of any biomass conversion technology requires not only tailored high-performance catalysts, but methods to sustainably produce them at industrial scales. A commercial biorefinery for the conversion of biomass *via* CFP sized to compete with incumbent fuel production technologies would require catalyst loads of 50 tons or more, which poses a significant scaling challenge for the production of tailored catalysts by advanced synthetic methods.¹⁷

There are few chemical manufacturing techniques deployed at commercial scales that are amenable to the production of catalysts with targeted physical properties at quantities suitable for the needs of a conceptual biorefinery. For example, high surface area catalyst supports exhibiting the aforementioned physical and chemical properties are commonly produced using flame synthesis methods such as vapour-fed aerosol flame synthesis, in which precursor vapours are continuously combusted in a controlled stream. The versatility^{18–20} and scalability^{21–23} of this process is highlighted by its industrial maturity, with flame-synthesized materials manufactured on the scale of millions of tons per year to produce a wide variety of engineered materials including activated carbons and fumed metal oxides (e.g., SiO₂, TiO₂, Al₂O₃). A particularly promising method for advanced catalyst synthesis is flame spray pyrolysis (FSP), in which a continuously fed solution of metal or metal oxide precursor(s) in organic solvent is combusted to form nanoparticles (NPs) from the aerosolized precursors. This methodology has been adapted to prepare multi-component heterogeneous catalysts in a single step (*i.e.*, metal active phase and a metal oxide support) by co-feeding a judicious choice of metal and metal oxide precursor solutions,^{24–26} including single-atom and nanostructured Pt on TiO₂.^{27,28} This approach offers substantial benefits for green synthesis of catalytic materials, as it combines two conventional steps—flame synthesis of the catalyst support and loading of an active phase—into a single FSP step. FSP is particularly promising in the application of catalyst synthesis because it is a continuous, readily scalable process that can significantly decrease the number of processing steps involved in catalyst manufacturing compared to traditional solution-based routes (Scheme 1). Furthermore, it offers precise control over catalytically relevant physical properties (e.g., spatial distribution of active components) that may not be as readily controlled through traditional synthetic methods.²⁴ However, despite the maturity of flame synthesis technologies and the development of “green manufacturing” technologies for NPs afforded by FSP, the



Scheme 1 Simplified comparison of the synthetic steps involved in the preparation of a Pt/TiO₂ catalyst by (a) flame spray pyrolysis and (b) a traditional incipient wetness impregnation approach.

viability and potential for widespread market adoption of this technology for catalytic applications is uncertain without a detailed economic assessment.

Technoeconomic evaluation of new synthesis pathways is an effective approach to assess catalyst viability at industrial scales and identify barriers to commercialization.^{3,29} To proliferate a new method offering reduced environmental impact compared to traditional wet chemistries,^{30,31} it is crucial to demonstrate that the method can achieve cost parity with incumbent methods *via* a preliminary technoeconomic analysis. Moreover, cost assessment allows the quantification of benefits and cost reduction opportunities for novel synthesis approaches, such as the FSP route described herein. We employed the CatCost catalyst cost estimation tool^{32,33} to better understand the holistic impact of the FSP synthesis method.

Herein, we report the continuous synthesis of a family of bifunctional Pt/TiO₂ catalysts in a single step *via* FSP and evaluation of their performance for the upgrading of biomass pyrolysis vapours in *ex situ* CFP. Catalysts synthesized by this method were found to exhibit higher selectivity towards aromatics, partially deoxygenated compounds, and alkenes compared to traditionally prepared IW-synthesized materials, which primarily yielded alkanes. Moreover, FSP-synthesized catalysts resulted in similar or higher total yields of carbon products when compared to IW-synthesized materials at comparable Pt loadings, indicating better utilization of the expensive noble metal. Detailed microscopy and spectroscopic characterization of the FSP-synthesized catalysts revealed notable differences in Pt speciation and dispersion compared to IW-based catalysts which we hypothesize to be responsible for the differences in observed selectivity and deactivation behaviour. Facile control over the



relative speciation of single Pt atoms, clusters, and NPs as a function of FSP reaction conditions, and the impact of these features on the catalytic performance, highlight the ability of this synthetic method to tune for a desirable product slate. Finally, we performed a cost assessment to compare the FSP synthesis process with a conventional IW method, providing an economic outlook on the FSP synthesis method and areas in which further research is needed to lower cost and improve the commercial viability of this process.

Experimental section

Materials

Titanium tetraisopropoxide (TTiP, 98%, ACROS Organics), and platinum(II) acetylacetonate (Pt(acac)₂, 97%, Sigma-Aldrich) were used as received without further purification. Titania support (Aeroxide P25) used in the IW impregnation synthesis was supplied by Evonik and used as received. All gases used in the synthesis and catalytic evaluation were ultra-high purity (UHP) unless otherwise noted. Loblolly pine was supplied by Idaho National Laboratory and elemental analyses showed that the feed contained 49.3 wt% carbon, 6.1 wt% hydrogen, 0.1 wt% nitrogen, and 44.5 wt% oxygen (by difference) on a dry biomass basis.

Catalyst synthesis

Flame spray pyrolysis synthesis of Pt/TiO₂. The pyrolytic synthesis of Pt/TiO₂ was conducted in a custom-built FSP apparatus that is presented in Fig. S1.† A combustible mixture of TTiP, acetonitrile, and xylene was prepared in the volumetric ratio 20:28:52. Platinum acetylacetonate (Pt(acac)₂) was dissolved in 300 mL batches of this mixture at concentrations of 0.30, 0.75, and 1.51 mM targeting nominal Pt loadings of 0.10, 0.25, and 0.50 wt%. Actual Pt loadings determined by inductively coupled-optical emission spectroscopy (ICP-OES) were found to be 0.07, 0.14, 0.33 wt%, respectively, as shown in Table 1 and denoted as FSP-Pt(x)/TiO₂ (x = 0.07, 0.14, and 0.33). A neat mixture of the TTiP, acetonitrile, and xylene was prepared to give the control FSP titania (FSP-TiO₂). The prepared solutions were combusted in an enclosed FSP system using an array of six premixed methane–oxygen pilot flames arranged circumferentially around the spray nozzle and oriented to converge on the flame axis 35 mm from the tip of the spray nozzle, as shown in Fig. S1.† The pilot flames were

operated continuously to ignite and support the combustion. Oxygen, used as a carrier gas flowing at 10 LPM, was used to atomize the liquid feed and provide a portion of the combustion oxidizer. An additional 25 LPM of oxygen was provided by a sheath flow located circumferential to the atomizing nozzle (Fig. S1.†). The pilot methane and oxygen flows were 2 and 4 LPM respectively and the liquid feed was 8 mL min⁻¹. Combustion was quenched with N₂ gas that was fed circumferentially to the overall burner at 50 LPM. Finally, an additional 50 LPM of N₂ quench gas was added to the combustion exhaust flow prior to collecting the produced Pt/TiO₂ powder on a glass paper filter pad.

Incipient wetness impregnation synthesis of Pt/TiO₂. A series of 3 Pt/TiO₂ catalysts were prepared using traditional IW synthesis methods, denoted as IW-Pt(x)/TiO₂, where x represents actual loadings of 0.07, 0.14, and 0.27, respectively. The aqueous IW point of the titania (Evonik, Aeroxide P25) was determined to be 0.21 mL g⁻¹. Briefly, an aqueous solution containing Pt(NH₃)₄(NO₃)₂ in concentrations corresponding to theoretical Pt loadings of 0.08, 0.16, and 0.32 wt% was added dropwise to the TiO₂ powder. The impregnated material was dried in air at 50 °C for ca. 12 h. The dried material was then calcined in air at 500 °C (heating at 5 °C min⁻¹) for ca. 3 h.

Catalyst characterization

Platinum metal content of the prepared materials was determined by inductively coupled plasma optical emission spectroscopy (ICP-OES) performed by Galbraith Laboratories (Knoxville, TN).

Powder X-ray diffraction (XRD). XRD measurements were performed using a Rigaku Ultima IV diffractometer with a Cu Kα source (40 kV, 44 mA). Diffraction patterns were collected in the 2θ range of 10–100° at a scan rate of 4° min⁻¹. Diffraction patterns were compared to powder diffraction files (PDFs) of the reference materials from the International Centre for Diffraction Data (ICDD).

Transmission Electron Microscopy (TEM). Samples for TEM were prepared from powder samples of TiO₂ and Pt/TiO₂ that were dispersed in ethanol by brief sonication at room temperature. One drop of the suspension was drop-cast onto a carbon-coated copper grid (Ted Pella, 01885-F) and allowed to evaporate. TEM imaging was performed on a JEOL JEM-2100F microscope at an accelerating voltage of 200 kV.

Table 1 Physico-chemical properties of FSP- and IW-Pt/TiO₂ catalysts

Catalyst	Pt loading (wt%)	Surface area (m ² g ⁻¹)	Acid-site density (μmol g ⁻¹)	Acid-site density (μmol m ⁻²)
FSP-TiO ₂	—	49	377	7.7
P25-TiO ₂	—	56	457	8.2
FSP-Pt(0.07)/TiO ₂	0.07	42	297	7.1
FSP-Pt(0.14)/TiO ₂	0.14	40	278	7.0
FSP-Pt(0.33)/TiO ₂	0.33	41	266	6.5
IW-Pt(0.07)/TiO ₂	0.07	53	377	7.1
IW-Pt(0.14)/TiO ₂	0.14	52	369	7.1
IW-Pt(0.27)/TiO ₂	0.27	56	387	6.9



Scanning transmission electron microscopy (STEM). Samples for high angle annular dark field (HAADF)-STEM were prepared analogously to those analysed by TEM, with the addition of a vacuum drying step, which was performed in a vacuum oven at 82 °C overnight. STEM images were obtained with a probe aberration corrected 200 kV STEM/TEM equipped with a cold field emission source with 0.35 eV energy resolution (JEOL JEM-ARM200CF). At least 3 locations were surveyed for each sample analysed by STEM. The Pt NP and sub-nm cluster sizes, as well as presence of single Pt atoms, were determined by manually measuring and counting with the Gatan Microscopy Suite Software by analysing at least 10 STEM images for each sample. The diameter was measured horizontally regardless of the shape/orientation of the Pt NP. The frequency of Pt NPs and single atoms was binned with a 0.25 nm interval to produce the histograms presented in Fig. 2 and S6.†

Surface area and acid-site quantification. N₂ physisorption data were collected at 77 K using a Quantachrome Autosorb 1-C instrument. The Pt/TiO₂ samples were pretreated under vacuum for 12 h at 150 °C. Surface area was determined using the Brunauer–Emmett–Teller (BET) method, and pore volume was determined from the adsorption isotherm data using the Barrett–Joyner–Halenda (BJH) method. The acid site densities of the Pt/TiO₂ catalysts were determined *via* temperature-programmed desorption of ammonia (NH₃-TPD) using a dedicated microflow reactor system (Altamira AMI-390) equipped with a thermal conductivity detector (TCD). The catalysts were reduced in 10% H₂/Ar by heating at 10 °C min⁻¹ from RT to 450 °C for 2 h prior to NH₃ adsorption at 100 °C and subsequent TPD in He flow to 450 °C. A sample loop of known volume (5 mL) was used to calibrate the TCD response for 10% NH₃/He and, thereby, quantify the amount of NH₃ desorbed from the catalyst sample. Surface acidic sites were quantified by assuming adsorption stoichiometry of one NH₃ molecule adsorbed per acid site.

Diffuse Reflectance Infrared Spectroscopy (DRIFTS). DRIFTS measurements were performed on a Thermo-Nicolet iS50 FTIR spectrometer fitted with a Harrick “Praying Mantis” high-pressure/high-temperature cell equipped with CaF₂ windows. Spectra were collected from 64 scans of both the background and sample at 4 cm⁻¹ resolution and as-synthesized catalysts were activated *in situ* by heating to 450 °C at 5 °C min⁻¹ in flowing 5% H₂/N₂ (100 mL min⁻¹). Catalysts were held at 450 °C then cooled to collect a background spectrum. Samples were dosed with 1% CO/N₂ flowing at 110 mL min⁻¹ for *ca.* 20 min, then the cell was purged with N₂ (100 mL min⁻¹) for *ca.* 20 min prior to data collection. All spectra were collected at 50 °C.

X-ray absorption spectroscopy (XAS). XAS experiments were performed on the insertion device beamline (10-ID) of the Materials Research Collaborative Access Team at the Advanced Photon Source, Argonne National Laboratory. Measurements were performed in fluorescence mode using a Dectris Pilatus 100k detector downstream of a bent Laue energy analyzer to reject scattering and fluorescence from the

TiO₂ support. The spectrum of a Pt foil (edge energy 11562.76 eV) was collected simultaneously with each measurement for energy calibration. Spectra of the fresh catalysts were collected at room temperature in air. Reductive pretreatments (450 °C in 50 sccm 100% H₂ for 0.5 h) were performed using a custom built *in situ* reactor.³⁴ Following reduction, the samples were cooled to room temperature and spectra were collected under flowing H₂.

XAS data processing and analysis were performed using the Demeter software suite.³⁵ Edge energies were determined from the position of maximum intensity of the derivative of X-ray absorption near edge structure (XANES) spectra. X-ray absorption fine structure (EXAFS) coordination parameters were determined from simultaneous fits in *R*-space of the *k*¹, *k*², and *k*³ weighted Fourier transformed EXAFS using theoretical phase shift and backscattering amplitudes calculated from FEFF.³⁶ Pt–O and Pt–Pt scattering were modeled from the structures of PtO₂ and FCC Pt, respectively. The amplitude reduction factor for Pt was determined to be 0.86 from a fit of the metal foil and this value was held fixed in fits of the samples to determine coordination numbers.

Catalytic conversion of biomass pyrolysis vapours

Catalytic evaluation of the prepared Pt/TiO₂ catalysts was performed in a tandem micro-furnace pyrolyzer (PY-2020iS, Frontier Laboratories, Japan) coupled with a gas chromatograph-mass spectrometer/flame ionization detector (GC-MS/FID). Pyrolysis of pine particles (at 500 °C and 1 atm under inert flow) and subsequent catalytic upgrading of pyrolysis vapours (at 400 °C and 1 atm in 50% H₂/50% He) over the Pt/TiO₂ catalysts was performed using a tandem micropyrolyzer-GC-MS system.^{37,38} A detailed description of its operation has been previously reported.^{39–41} The micropyrolyzer apparatus consists of a pyrolysis zone for generating biomass vapours coupled to a downstream zone for catalytic upgrading. Condensable products were collected in a liquid nitrogen trap downstream of the reactor where they were subsequently vapourised into the inlet of the GC-MS/FID during GC oven heat up. Samples of biomass (*ca.* 500 µg each) in deactivated stainless-steel cups were dropped automatically in intervals of 30 min into the preheated pyrolysis zone operating at 500 °C. The products from the pyrolysis zone were entrained in 54 mL min⁻¹ of He carrier gas and passed over a fixed bed of 10 mg catalyst. For experiments conducted in reducing gas atmospheres, 54 mL min⁻¹ of H₂ was added prior to the fixed-bed reactor for a total flow of 108 mL min⁻¹. The total pressure was 115 kPa. For each experiment, three successive cups of biomass were passed over the same catalyst bed for a total mass ratio of biomass fed to catalyst of 0.15 g g⁻¹. For deactivation experiments, a biomass:catalyst weight ratio (B:C) of 1.0 was reached by successive injection of biomass cups. The catalysts were pretreated for 60 min in a mixture of 50% He/50% H₂. The temperature of the catalysis zone was maintained at 400 °C. The condensable vapours produced



during pyrolysis and upgrading were adsorbed on the liquid nitrogen trap, but light gases such as CH₄ and CO passed through the trap and were separated in a GS-GasPro column and measured by a TCD. The condensable vapours were desorbed from the liquid nitrogen trap and separated in an Ultra-Alloy-5 capillary column, which utilized a stationary phase consisting of 5% phenyl and 95% dimethyl polysiloxane. The GC oven was programmed to hold at 40 °C for 4.5 min, then ramp to 300 °C at a rate of 20 °C min⁻¹. The separated upgraded species were identified using the NIST GC-MS library and quantified based on calibration of twenty compounds representative of upgraded pyrolysis vapours. The CFP product carbon yields were calculated using eqn (1).

$$\text{Carbon yield} = \frac{\text{total carbon in the product formed}}{\text{total carbon in the biomass feed}} \quad (1)$$

Economic assessment

Cost estimates were prepared in Microsoft Excel v16 using the spreadsheet version of CatCost v1.0.4. Further discussion of the CatCost methods and assumptions has been previously published.³² All prices were adjusted to 2016 USD using the U.S. Bureau of Labor Statistics Chemical Producer Price Index or, for equipment costs, the *Chemical Engineering Plant Cost Index*.

Raw materials prices at ≥1000 kg scale were estimated through a combination of vendor quotations, freely available and proprietary price databases, and estimates from industry experts. Generally, several sources were consulted to develop an average and/or verify each assumed price. A factor of 3% was added to the raw materials costs to account for waste and spoilage.

Processing costs were estimated using the CapEx & OpEx Factors method (catalyst manufacture at a dedicated new-build plant).³² A production scale of 100 kg h⁻¹ (788 000 kg per year accounting for downtime) was used for all scenarios. Equipment was scaled from a base design automatically to the specified production scale by increasing the number of equipment items when the upper bound of the validity of that cost correlation was exceeded; similarly, when the equipment was required at less than the correlation's minimum size, the cost of the minimum size was used. Installation costs were estimated using the equipment-specific installation factors of Garrett,⁴² and labor factors (number of operators required per shift for that equipment item) were taken from Ulrich and Vasudevan.⁴³ In both cases, the average value was used, or the midpoint if only a range was provided. The remainder of the capital costs—including direct capital costs like installation, piping, instrumentation, and buildings; indirect costs like engineering, legal, and contingencies; and working capital—were estimated as fixed factors (multipliers) of the total purchased equipment cost using the modified Lang factors^{44,45} of Peters and

Timmerhaus.⁴⁶ A similar calculation approach and factors were taken from the same source⁴⁶ to determine operating costs such as supervisory labor and maintenance supplies, fixed/indirect costs such as insurance and overhead, and general expenses such as distribution and marketing. Direct labor was calculated by summing the labor factors of all the equipment items after scaling to the specified production rate, and then rounding up to the nearest whole number to determine the number of operators. Year-round operation (8760 hours) with full staffing during maintenance downtime (10%) was assumed. A labor rate, including benefits, of \$48 per hour for US Gulf Coast production was used.

Results and discussion

Catalyst characterization

The FSP approach investigated here involves combustion of an aerosolized continuous feed of a solvent mixture comprised of liquid precursors (of the active metal phase and oxide support). The metals (Ti and Pt) are atomized into the gas phase followed by condensation (nucleation and agglomeration) into nano-sized catalyst particles. The relative volatility, or conversely, the temperature at which the individual metals condense into solid particles, establishes an order of condensation which results in Pt tending to be located on the TiO₂ surface due to the higher volatility of Pt metal vapour compared to Ti. In this work, we synthesized a series of Pt/TiO₂ catalysts *via* the FSP method modified from previous reports.²⁸ Briefly, a solution of Pt(acac)₂ was prepared in a mixture of TTiP, acetonitrile, and xylene and combusted in a custom enclosed FSP system schematically presented in Fig. S1.†

Catalysts prepared by FSP with weight loadings of 0.07, 0.14, and 0.33 wt% Pt were attained by controlling the concentration of the Pt precursor solution. Actual loadings are presented in Table 1, with catalysts denoted as FSP-Pt(x)/TiO₂ (x = 0.07, 0.14, and 0.33) based on these loadings. A lower feed-rate was employed to prepare the 0.33 wt% Pt catalyst to prevent further Pt particle aggregation at the higher required concentration. To assess the impact of catalyst synthesis method on the observed catalyst properties and performance, catalysts with similar nominal loadings were prepared *via* traditional IW impregnation and are presented in Table 1, denoted as IW-Pt(x)/TiO₂ (x = 0.07, 0.14, and 0.27). The physico-chemical properties of all the catalysts are summarized in Table 1. The BET surface area measurements *via* N₂ physisorption studies reveal that all catalysts synthesized by FSP, including the Pt-free TiO₂ (FSP-TiO₂), exhibit a consistent surface area between 41–49 m² g⁻¹, which is comparable to that of the commercially available Aerioxide P25 TiO₂ support (56 m² g⁻¹). Notably, the TiO₂ morphologies are similar between P25 and FSP-TiO₂ (as observed by TEM and described below) despite their syntheses utilizing different titania precursors (TiCl₄ and TTiP, respectively). Therefore, this commercial material was chosen as the support to prepare the IW-Pt/TiO₂ catalysts.



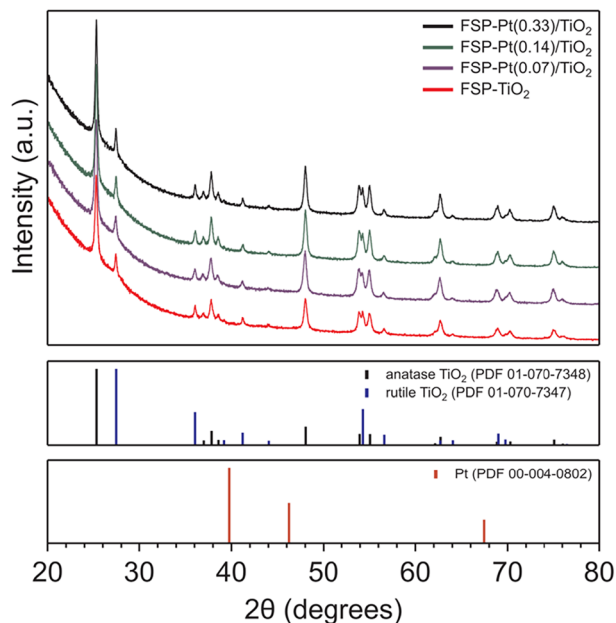


Fig. 1 XRD patterns of FSP-derived Pt/TiO₂ and TiO₂ with reference patterns provided below.

Inclusion of the Pt precursor in the FSP synthesis resulted in a minor reduction in the measured surface area from 49 m² g⁻¹ (FSP-TiO₂) to 40–42 m² g⁻¹ (FSP-Pt/TiO₂), although no relationship was observed between Pt weight loading and surface area. In addition, all FSP-Pt/TiO₂ catalysts were found to have pore sizes and pore volumes consistent with the parent FSP-TiO₂ (Table S1 and Fig. S2†) suggesting that the addition of Pt to the FSP synthesis does not disrupt the formation of the FSP-TiO₂ pore structure.

The acid site density, as measured by ammonia TPD (NH₃-TPD) (profiles shown in Fig. S3†), of the FSP-TiO₂ (377 μmol g⁻¹) was found to be lower than that of the commercial IW-TiO₂ (457 μmol g⁻¹) on a per-gram-catalyst basis, which is consistent with the lower measured surface area of the FSP-TiO₂. The acid site density normalized by the surface area for FSP-TiO₂ (7.7 μmol m⁻²) was within the experimental measurement error when compared to P25-TiO₂ (8.2 μmol m⁻²). Pt incorporation at different metal loadings reduced the acid site density for both FSP-TiO₂ and IW-TiO₂ by 20–30% as compared to the parent TiO₂.

XRD analysis of the as-synthesized FSP-Pt/TiO₂ catalysts revealed the presence of titania with a predominantly anatase structure that corresponds well with the ICDD PDF 01-070-7348, with a minor fraction adopting the rutile phase (ICDD PDF 01-070-7347) in all cases, as shown in Fig. 1. The average composition of anatase and rutile phases, as determined by single-peak reference intensity ratio analysis, of all materials prepared by FSP (83.8 ± 1.5% anatase) were slightly below that of the P25-derived IW materials (91.9 ± 0.5% anatase), as shown in Table S2.† Unsurprisingly, the addition of Pt to the TiO₂ support *via* IW had no impact on the resulting distribution of phases as determined by XRD (Fig. S4†). However, at all weight loadings studied, the introduction of

Pt to the FSP synthesis increased the amount of rutile present (83.7–85.2%) compared to the bare FSP-TiO₂ (81.7%). This observation is consistent with previous literature reports,^{28,47} and has been attributed to structural similarity between PtO₂ and rutile TiO₂, whereby PtO₂ seeds promote the formation of the rutile phase.^{48–51} In addition, the presence of surface Pt has been found to have a dehydroxylating effect on TiO₂ at the high temperatures of the FSP process, which has also been shown to promote rutile formation.²⁸ Scherrer analysis of the (101) diffraction peak at 25.3° 2θ for the anatase titania revealed crystallites with diameters (d_{XRD}) in the range of 26–34 nm, with a trend toward increasing size observed with higher Pt loadings. This trend is in good agreement with previous reports that reveal an increasing d_{XRD} over a range of low Pt loadings (0–1 atom%) and is consistent with the materials synthesized herein (0.03–0.14 atom% Pt).²⁸ This effect of increased crystallite size as a function of Pt loading is mirrored by the rutile phase as determined by Scherrer analysis of the (110) diffraction peak at 27.4° 2θ where the crystallite size was observed to increase from 33 to 43 nm over the interrogated weight loadings. All crystallite sizes determined by XRD are presented in Table S2.† Unsurprisingly, no diffraction peaks corresponding to Pt metal were observed in any of the Pt/TiO₂ catalysts because of the low Pt loadings and small particle size, as well as the high crystallinity of the support.

The as-synthesized FSP-Pt/TiO₂ and IW-Pt/TiO₂ catalysts were imaged with high resolution TEM and HAADF-STEM to assess any morphological changes induced by either Pt weight loading or synthetic method. All FSP-Pt/TiO₂ materials were comprised of aggregates of titania particles approximately 35 ± 15 nm in diameter which is consistent with the titania particle size determined by XRD. TEM images revealed that the titania aggregates of FSP-Pt/TiO₂ (Fig. S5a–c†) have similar particle size and size distributions for both FSP-TiO₂ (Fig. S5d†) and P25-TiO₂ (Fig. S5e†). HAADF-STEM was employed to elucidate the size distribution of the Pt species present on the FSP-TiO₂ catalysts. As expected, the Pt speciation and size distribution was found to be dependent on concentration of the Pt(acac)₂ solution used during synthesis (*i.e.*, Pt weight loading).⁵² Analysis of the HAADF-STEM images of the as-synthesized FSP-Pt(0.07)/TiO₂ revealed only the presence of highly dispersed single atoms of Pt (Fig. 2a and g), with no sub-nm clusters or NPs of Pt observed. Exemplary single atoms are highlighted with yellow circles in Fig. 2a. Conversely, both FSP-Pt(0.14)/TiO₂ and FSP-Pt(0.33)/TiO₂ (Fig. 2b and c) were found to possess a mixture of single-atoms, sub-nm clusters, and NPs with size distributions presented in the histograms in Fig. 2h and i, respectively. To characterize the morphology of the Pt-species present under conditions more relevant to CFP, the as-synthesized FSP-Pt/TiO₂ catalysts were subjected to a reductive thermal treatment (450 °C, 100% H₂, 6 h) and imaged with HAADF-STEM. For all FSP-Pt/TiO₂ catalysts there was an increase in the average particle sizes and shift in the particle size distributions (Fig. 2g–i). After reduction of the



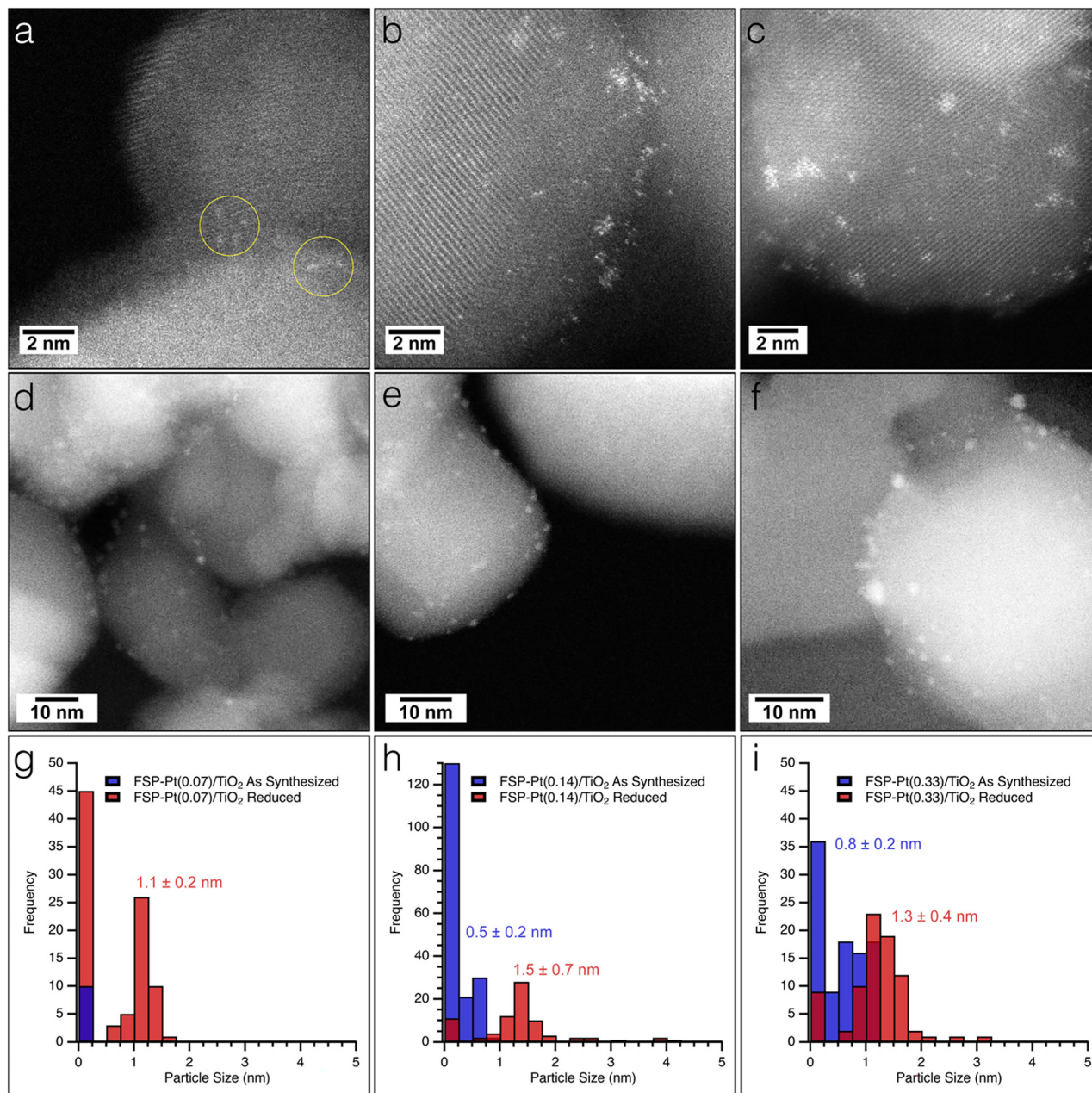


Fig. 2 HAADF-STEM images of as-synthesized (a) FSP-Pt(0.07)/TiO₂, (b) FSP-Pt(0.14)/TiO₂, and (c) FSP-Pt(0.33)/TiO₂ and after reduction in hydrogen (d) FSP-Pt(0.07)/TiO₂, (e) FSP-Pt(0.14)/TiO₂, and (f) FSP-Pt(0.33)/TiO₂. Particle size histograms for as-synthesized and reduced (g) FSP-Pt(0.07)/TiO₂, (h) FSP-Pt(0.14)/TiO₂, and (i) FSP-Pt(0.33)/TiO₂. Exemplary regions of single atoms are highlighted with yellow circles.

FSP-Pt(0.07)/TiO₂ material, both small NPs and Pt clusters were observed with an average size of 1.1 ± 0.2 nm, as shown in Fig. 2d, whereas only atomically dispersed Pt could be identified in the as-synthesized material. Despite the appearance of larger particles, the predominant species present in the FSP-Pt(0.07)/TiO₂ catalyst remained atomically dispersed Pt. The observed particle growth was more severe in the cases of FSP-Pt(0.14)/TiO₂ and FSP-Pt(0.33)/TiO₂ (Fig. 2e and f), where, after reduction the ratio of sub-nm clusters-to-NPs of Pt decreased from 2.5 to 0.2 and from 0.6 to 0.1, respectively. HAADF-STEM of the analogous samples

prepared by IW revealed Pt particles with sizes of 0.9 ± 0.2 nm and 1.1 ± 0.2 nm for IW-Pt(0.07)/TiO₂ and IW-Pt(0.14)/TiO₂, respectively (Fig. S6[†]). The average Pt particle size of IW-Pt(0.27)/TiO₂ was found to be slightly larger at 1.3 ± 0.5 nm due to a population of particles approximately 2 nm in size. All Pt and titania particle sizes are presented in Table S3.[†]

XAS experiments were performed to complement the microscopy data to better understand changes in Pt oxidation state in the as-synthesized FSP-Pt/TiO₂ catalyst and reduced FSP-Pt/TiO₂ under reaction relevant pretreatment conditions. Notably, the as-synthesized FSP-Pt/TiO₂ catalysts have an



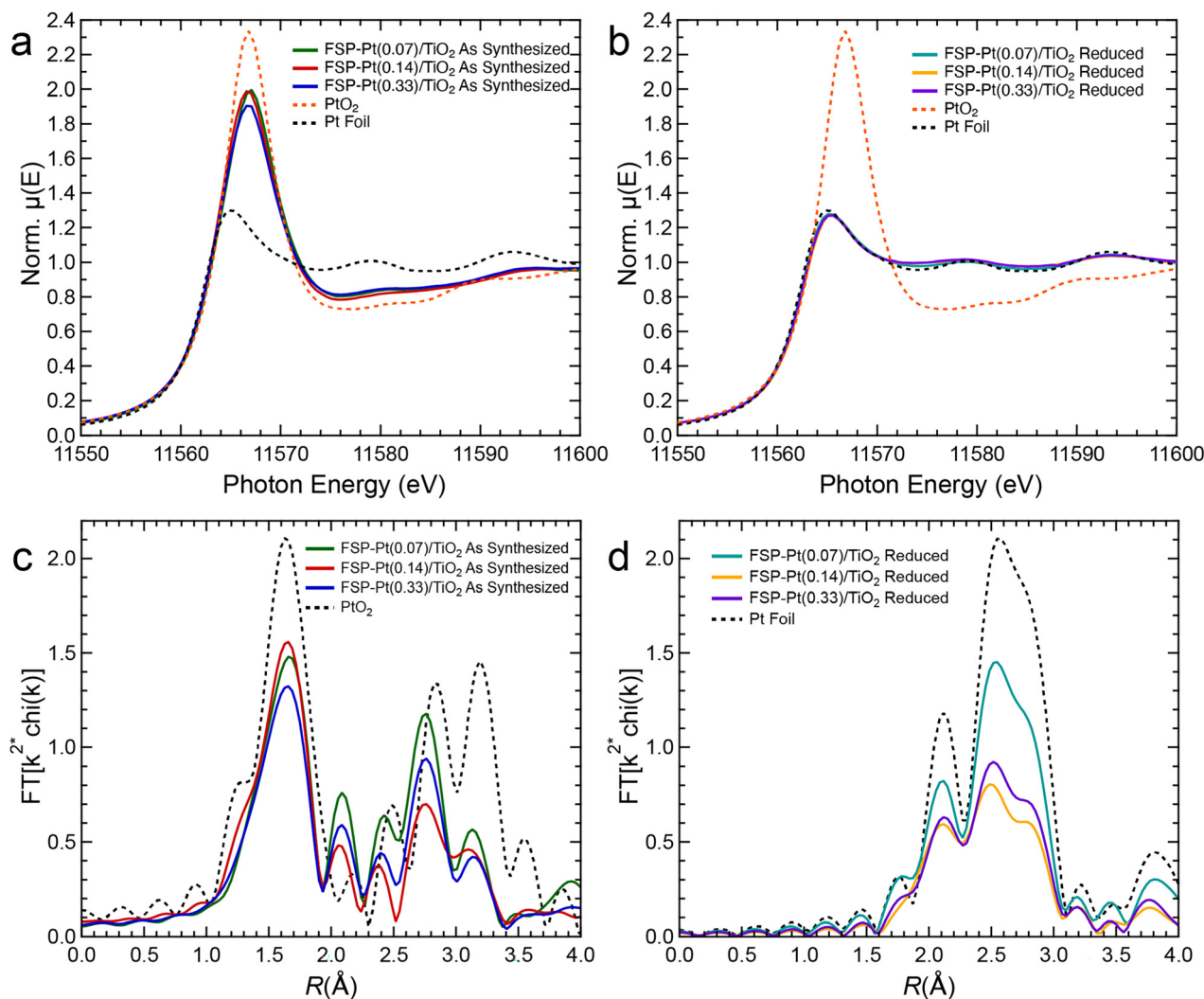


Fig. 3 XANES spectra of as (a)-synthesized and (b) reduced FSP-Pt/TiO₂ catalysts compared to Pt foil and PtO₂ standards; and magnitude of the Fourier transformed k^2 -weighted Pt L_{III} edge EXAFS of (c) as prepared and (d) FSP-Pt/TiO₂, Pt L_{III}-edge EXAFS.

intermediate white line intensity compared to PtO₂ and Pt foil (Fig. 3a). Because the white line intensity in the Pt L_{III}-edge XANES spectrum depends on the density of unoccupied 5d states, it has been frequently used as a metric to estimate the oxidation state of Pt.^{53,54} The white line intensity of the normalized XANES of all the as-synthesized samples (1.9–2.0) was found to be closer to that of the bulk PtO₂ (2.3) than measured for the Pt foil (1.3). It is important to note that due to the nature of XAS as a bulk technique, the observed intermediate white line intensity cannot prove the presence of a mixed population of Pt⁴⁺ (*i.e.*, PtO₂) and Pt⁰, as such a mixed population could be indistinguishable from a homogenous population of Pt²⁺. However, this result does indicate the presence of a significant population of oxidized Pt present in the as-synthesized samples. This result is in contrast with the reduced samples, as upon treatment with H₂ at 450 °C the white line intensities (1.3 norm. μeV) for all weight loadings are consistent with that of Pt⁰ suggesting complete reduction of the oxidized Pt species (Fig. 3b). The EXAFS of the as-

synthesized FSP-Pt/TiO₂ catalysts shown in Fig. 3c, reveals a Pt–O scattering peak at *ca.* 1.7 Å. However, despite a similar position to that of the first-shell peak in PtO₂, the magnitude is lower, suggesting a lower coordination number than the bulk oxide. Likewise, the second-shell peaks (*ca.* 2.2–3.4 Å) of the three samples are similar in shape, but the intensities are also significantly different than PtO₂, indicating the average local structure of Pt in the as-synthesized catalysts is not the same as the bulk oxide, which is consistent with the XANES analysis. Aside from intensity differences, the shapes of the real components of the Fourier transformed EXAFS of the second shell are essentially identical (Fig. S7[†]), suggesting that the as-synthesized materials contain the same species. Furthermore, similarities in the real components of the as-synthesized catalysts and Pt foil from 4.4–6.0 Å suggest a fraction of the Pt is present in metallic particles. After reduction, however, the EXAFS spectra (Fig. 3d) are consistent with metallic NPs (*i.e.*, similar shape as foil, but lower intensity). In agreement with the XANES, there is no evidence



of Pt–O scattering indicating all the Pt is reduced. Quantitative EXAFS fitting parameters for the FSP-Pt/TiO₂ catalysts are summarized in Table S4.†

DRIFTS was performed to investigate the adsorption of CO on the FSP-Pt/TiO₂ catalysts compared to their IW analogues (Fig. S8a and b†). Spectra were collected at 50 °C after a 450 °C pretreatment in 5% H₂ to better emulate the speciation of the catalyst prior to reaction. All catalysts analyzed exhibited two distinct spectral features, a minor band at 2109–2115 cm⁻¹ and a larger peak at 2071 cm⁻¹. The band at 2071 cm⁻¹ can be assigned to CO linearly adsorbed to Pt⁰ on NP surfaces.^{54,55} The relative intensity of this peak increased as a function of Pt loading in all cases, which is consistent with the increased population of metallic NPs present in the higher weight-loading materials as observed by STEM and TEM. The minor peak/shoulder observed at 2109–2115 cm⁻¹ may be attributed to more oxidized Pt²⁺, Pt^{δ+}, single Pt atoms, or very small clusters.^{54,56,57} Notably, the relative intensity of this feature is significantly higher for both FSP and IW catalysts at a Pt loading of 0.07 wt%. In the case of the FSP catalysts, this feature at 2109 cm⁻¹ is diminished almost entirely at Pt loadings of 0.14 and 0.3 wt%, consistent with the sintering and loss of dispersed Pt atoms observed after pretreatment. In contrast, the IW catalysts retain this feature at all loadings.

Taken as a whole, the results of the spectroscopic characterization and imaging data reveal several distinct physical properties that differentiate the FSP-synthesized materials from their IW analogues. This ability to tune the distribution of Pt species (*i.e.*, the population of Pt clusters, single atoms, and metallic NPs) as a function of FSP conditions highlights the potential of this synthetic method to afford control over the physico-chemical properties of the synthesized catalysts. In particular, differences in Pt speciation that were introduced by the FSP synthesis process may significantly alter their catalytic behaviour. To assess the impact of the physico-chemical changes afforded by the FSP method, we evaluated the performance of the FSP- and IW-synthesized materials for the upgrading of whole biomass pyrolysis vapours.

Whole biomass vapour upgrading

The catalytic performance of the FSP-Pt/TiO₂ materials was evaluated with whole biomass vapours of pine particles using a micropyrolyzer-GC-MS system that has previously been described in detail.^{39–41} The biomass was pyrolyzed at 500 °C in He flow, while the catalytic upgrading reaction was operated at 400 °C in H₂ flow at 115 kPa total pressure. Experiments were continued until a B:C of *ca.* 0.9–1.0 was reached based on literature precedent for evaluating the utility of catalysts for biomass conversion to fuels and chemicals, as well as to study the stability of these catalysts for continuous catalytic operation.^{40,58–60}

The hydrocarbon products observed were grouped into 6 categories: alkanes, alkenes, aromatics, partially upgraded

(intermediate) oxygenates, primary (unreacted) vapours, carbon monoxide, carbon dioxide, and a small fraction of unidentified species (unknowns, ≤1.5%). Representative species, that account for >90% of each category, are shown in Table S5.† Partially upgraded oxygenates are incompletely deoxygenated products formed from C–O cleavage *via* deoxygenation or hydrodeoxygenation of complex oxygenates present in the pyrolysis vapours (*i.e.*, primary vapours) such as furanones, acids, methoxyphenols, acetaldehyde, hydroxybutanones. Aromatics are formed by aromatic C–O bond cleavage (*via* direct deoxygenation or hydrodeoxygenation) of the oxygenates without subsequent hydrogenation to form cycloalkenes (partial hydrogenation) and cycloalkanes (complete hydrogenation). Alkenes are a result of deoxygenation of linear oxygenates present in pyrolysis vapours or C–C bond breaking (cracking) of alkyl chains in branched oxygenates. Complete hydrogenation of the alkenes, due to high hydrogen activation and utilization, leads to the formation of alkanes. A generalized reaction scheme is presented in Fig. 4.

As shown in Fig. 5, the total carbon yield for FSP-TiO₂ and P25-TiO₂ was low (<25%), with the product consisting mainly of primary vapours (not included in the yield) and CO₂, and alkanes and aromatics as minor products. Unsurprisingly, the absence of hydrogen-activating Pt metal sites resulted in limited hydrodeoxygenation leading to the formation of intermediate oxygenates through direct C–O and C–C bond cleavage on the acid sites of TiO₂ and/or thermolytic reaction. Incorporation of Pt resulted in an increased total carbon yield for all catalysts (Fig. 5a) with a concomitant enhancement of the yield of aromatics, alkanes, and intermediate oxygenates (Fig. 5b and c) due to hydrogen activation on the metal sites and subsequent hydrodeoxygenation, hydrogenation, and hydrogenolysis reactions. Considering the hydrocarbon products observed, excluding the primary oxygenates, FSP-Pt(0.07)/TiO₂ resulted in a carbon yield significantly higher than IW-Pt(0.07)/TiO₂ (34.3 ± 0.9% *vs.* 22.3 ± 0.7%) at the same Pt loading (0.07

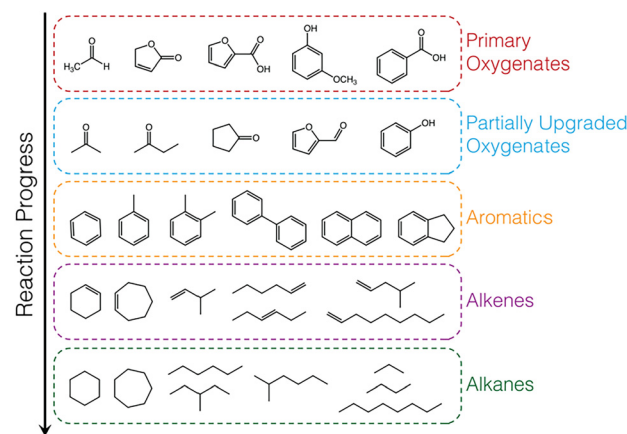


Fig. 4 Generalized reaction scheme for biomass vapour upgrading with exemplary species shown for each category of product observed.



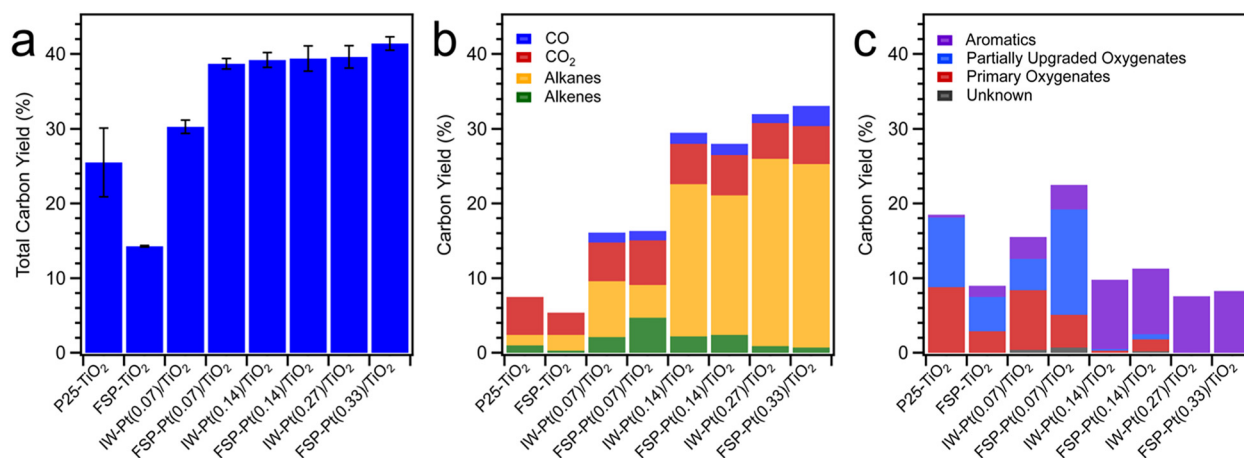


Fig. 5 Category plots of (a) total carbon yield, (b) carbon yield of CO, CO₂, alkanes, and alkenes, and (c) aromatics, partially upgraded oxygenates, and primary oxygenates for all catalysts.

wt%). This is suggestive of enhanced metal utilization resulting from the FSP-based synthesis method as evidenced by the increased population of Pt clusters observed for this material. The carbon yield for the intermediate Pt loading catalysts, FSP-Pt(0.14)/TiO₂ (39.4 ± 1.7%) and IW-Pt(0.14)/TiO₂ (39.2 ± 1.0%), are within the error of the micro-pyrolyzer experiments, which is consistent with the predominance of Pt NPs as the form of Pt for both of these catalysts as observed by TEM after reduction. At the highest Pt loading of 0.3 wt%, both FSP-Pt(0.33)/TiO₂ and IW-Pt(0.27)/TiO₂ produced similar carbon yields of 41.4 ± 0.9% and 39.6 ± 1.5%, respectively, suggesting the enhanced dispersion afforded by FSP synthesis is not preserved at higher loadings after reduction. This conclusion is in agreement with the imaging and spectroscopic results detailed above that revealed a loss of atomic and clustered Pt species upon reduction for the FSP-Pt(0.33)/TiO₂ catalyst. These results indicate that the FSP-synthesized Pt/TiO₂ catalysts exhibit comparable activity to conventional IW-synthesized Pt/TiO₂ catalysts at weight loadings ≥ 0.14%, with significantly greater activity at lower loadings. We hypothesize that the enhanced metal utilization at lower loadings observed for the FSP-synthesized materials results from the preservation of sub-nm Pt-clusters, as less sintering is observed in the case of the FSP-Pt(0.07)/TiO₂ catalyst as revealed by the microscopy study above.

Analysis of the CFP product distributions reveals that the FSP-Pt/TiO₂ catalysts generate a distinct product slate compared to the IW-Pt/TiO₂ catalysts. At a Pt loading of 0.07 wt% and cumulative B:C of ca. 0.3, CFP over FSP-Pt(0.07)/TiO₂ lead to greater formation of alkenes (4.7 ± 1.2 C%) and intermediate oxygenates (14.1 ± 2.1 C%) as compared to IW-Pt(0.07)/TiO₂ (alkenes (2.1 ± 0.1 C%); intermediate oxygenates (4.2 ± 0.3 C%)) as shown in Fig. 5b and c. Furthermore, FSP-Pt(0.07)/TiO₂ resulted in lower alkane production (4.4 ± 1.8 C%) and lower breakthrough of primary vapours (4.4 ± 1.9 C%) as compared to IW-Pt(0.07)/TiO₂ (alkane (7.5 ± 0.6 C%) and primary vapours (8.0 ± 1.6 C%)). At higher Pt loadings

(0.14 and 0.3 wt%) and similar B:Cs of ca. 0.3, the product distribution for FSP-synthesized and IW-synthesized Pt/TiO₂ was similar as shown in Fig. 5b and c. The differences in product distribution at lower Pt loadings suggest that the single-atom site and Pt clusters present on the FSP catalysts (at low Pt loadings) reduce the hydrogenation activity of the FSP-Pt/TiO₂ catalyst formulations and minimize the production of alkanes, thereby, leading to higher formation of partially upgraded oxygenates. This enables a facile synthetic handle for tuning the product slate in *ex situ* CFP towards ketones and alkenes, without adversely affecting C yield compared to IW synthesized catalysts. At higher Pt loadings of 0.14 wt% and 0.3 wt% Pt, similar product distributions for FSP and IW Pt/TiO₂ catalysts are in line with the similar catalyst characterization results (acid site density, Pt speciation, and particle size), as noted above.

We investigated the evolution of catalyst performance with increasing B:C (representative of reaction time-on-stream) for FSP- and IW-synthesized Pt/TiO₂. The results of these experiments are presented in Fig. 6 and S9–S11.† As shown in Fig. 6, FSP-Pt(0.07)/TiO₂ resulted in increasing yields of intermediate oxygenates (11.5–18.9 C%) and primary vapours (2.5–10.0% C%) over the course of reaction (Fig. 6b and d), whereas at all B:Cs, IW-Pt(0.07)/TiO₂ catalysts allowed breakthrough of more primary vapours (6.3–17.1 C%) and lower production of intermediate oxygenates (4.3–5.7 C%; Fig. 6a). These results indicate that FSP-Pt(0.07)/TiO₂ enables enhanced catalytic deoxygenation of primary oxygenates to partially deoxygenated intermediates *via* C–O bond cleavage as compared to IW-Pt(0.07)/TiO₂. Further, as shown in Fig. S9,† FSP-Pt(0.07)/TiO₂ showed a consistently higher yield (0–3.5 C%) of phenolic molecules (alkyl phenols and phenol) and a lower yield (0–0.4 C%) of methoxyphenols for B:C ca. 0.1–1.0 whereas IW-Pt(0.07)/TiO₂ produced higher yields (4.8–5.4 C%) of methoxyphenols and a lower yield (0–1.1 C%) of phenols from B:C ca. 0.1–0.9, further evidence of the enhanced deoxygenation functionality of FSP catalyst. Both the FSP- and IW-synthesized Pt/TiO₂ catalysts exhibited low



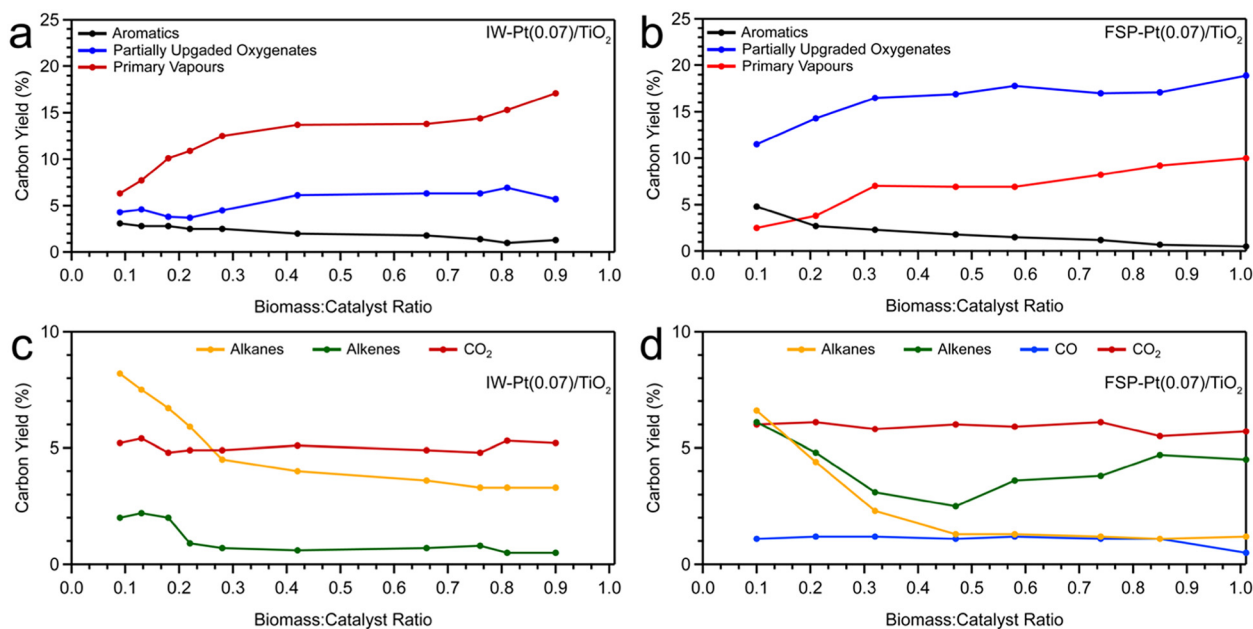


Fig. 6 Carbon yield to aromatics, partially upgraded oxygenates, and primary vapours as a function of biomass-to-catalyst ratio for (a) IW-Pt(0.07)/TiO₂ and (b) FSP-Pt(0.07)/TiO₂ catalysts and carbon yield to alkenes, alkanes, CO, and CO₂, as a function of biomass-to-catalyst ratio for (c) IW-Pt(0.07)/TiO₂ and (d) FSP-Pt(0.07)/TiO₂ catalysts.

yields of 1-ring aromatics (Fig. S9a and b†). As shown in Fig. 6c and d, FSP-Pt(0.07)/TiO₂ and IW-Pt(0.07)/Pt/TiO₂ demonstrated low yields of alkanes and alkenes, however, FSP-Pt(0.07)/TiO₂ maintained significantly higher yields of alkenes over the course of the reaction compared to the IW analogue (2.5–6.1 C% vs. 0.2–2.2 C%, respectively). This comparison further indicates the lower hydrogenation functionality of FSP Pt/TiO₂ catalyst in comparison to IW catalyst at low Pt loadings.

Fig. S10a–d† shows the results of deactivation studies for FSP-Pt(0.14)/TiO₂ and IW-Pt(0.14)/TiO₂. At B:C < 0.39, both catalysts demonstrate similar activity and product distribution. However, at B:C > 0.39, FSP-Pt(0.14)/TiO₂ exhibited increasing yields of intermediate oxygenates (increases from 2.1% to 16.5% over the course of reaction) with a concurrent decrease in aromatics yield (from 9.7 to 4.0 C%). The yield of intermediate oxygenates produced over FSP-Pt(0.14)/TiO₂ was observed to decrease starting at a B:C ca. 0.77 with a concurrent increase in the yield of primary vapours, indicative of catalyst deactivation. In contrast, CFP over the IW-Pt(0.14)/TiO₂ resulted in a consistent yield of aromatics (8.6 ± 0.8 C%) until a B:C ca. 0.96 with a low intermediate oxygenates yield (2.0 ± 1.8%) and minimal breakthrough of primary vapours (<0.7%). Moreover, as shown in Fig. S10d,† the alkane yields decreased rapidly for FSP-Pt(0.14)/TiO₂ after B:C ca. 0.17, while IW-Pt(0.14)/TiO₂ showed consistent alkane yields until a B:C ca. 0.78. These results suggest a more rapid loss of hydrogenation activity for FSP(0.14)/TiO₂ compared to the IW-Pt(0.14)/TiO₂ analogue. FSP-Pt(0.33)/TiO₂ and IW-Pt(0.33)/Pt/TiO₂ catalysts result in similar product distribution and carbon yields with increasing B:C from ca. 0.08–0.98 (Fig. S11a–d†). Both

catalysts demonstrated increased hydrogenation activity compared to their lower Pt loading analogues with high C yields to aromatics and alkanes and low yields of intermediate oxygenates, primary vapours, and alkenes. Both catalyst systems exhibited minimal deactivation over the B:Cs evaluated, with sustained high hydrocarbon yields (carbon yields > 28.5%) until a B:C ca. 0.83.

These performance data coupled with the detailed structural characterization suggest that (1) the effect of highly dispersed single-atom/sub-nm Pt clusters is more pronounced at low Pt loadings on FSP-synthesized catalysts leading to higher hydrocarbon yields and catalyst stability, (2) at higher Pt loadings, FSP catalysts are as active and stable as the conventional IW catalysts, and (3) FSP-synthesized catalysts show lower hydrogenation activity than IW-synthesized catalysts. We hypothesize that the changes in the product slate at increasing B:C for the FSP-Pt/TiO₂ catalysts result from continued structural evolution under reaction conditions that is distinct from the deactivation observed for the traditionally prepared IW catalysts. Further study will seek to elucidate the basis of this structural evolution and its catalytic consequences.

Cost analysis of FSP synthesis

A cost estimate for the FSP-synthesized Pt/TiO₂ material at a 0.07% weight loading was developed in CatCost³² on the basis of the lab-scale procedures. The lab-scale FSP equipment has a throughput of 200 g h⁻¹ and approximate cost of \$200 000 and requires a dedicated operator. Unsurprisingly, this small-scale, labour-intensive equipment results in a very high estimated catalyst purchase cost of



Table 2 Purchase cost analysis results for FSP-Pt(0.07)/TiO₂ and IW-Pt(0.07)/TiO₂ materials

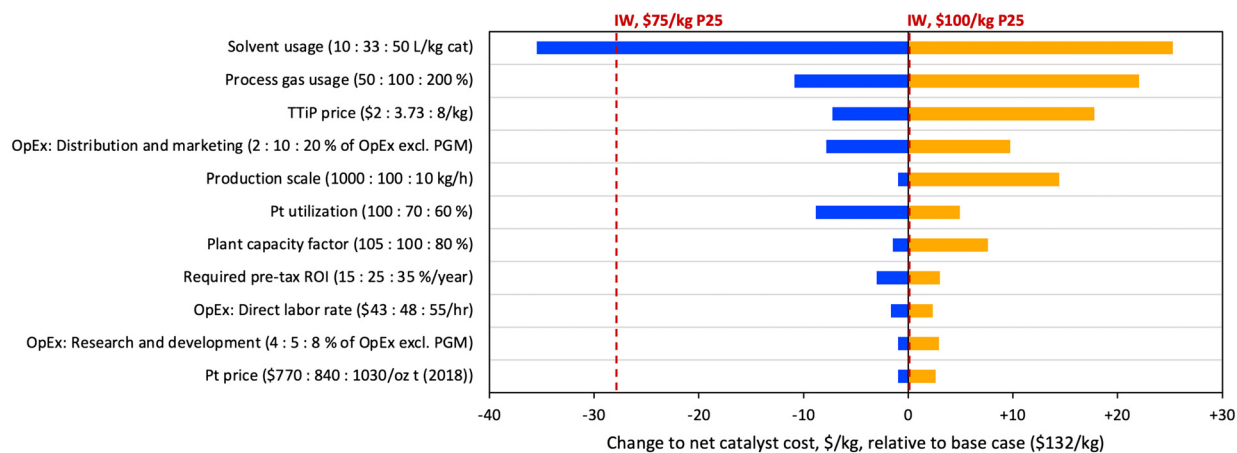
Inputs		Costs, 2016 \$ per kg catalyst							
Scenario	Details	Pt	TiO ₂ /precursor	Solvent	Process gases	CapEx	OpEx	Margin	Total
A	FSP, 200 g h ⁻¹	26	14	44	16	65	822	164	1150
B	FSP, 200 g h ⁻¹ , baghouse	26	14	44	16	56	810	140	1105
C	FSP, 10 kg h ⁻¹ , baghouse	26	14	44	16	3	33	8	142
D	IW, \$3 per kg TiO ₂	18	3	0.01	3	0.16	3	0.40	27
E	IW, \$50 per kg P25	18	51	0.01	3	0.16	10	0.40	83
F	IW, \$75 per kg P25	18	77	0.01	3	0.16	14	0.40	113
G	IW, \$100 per kg P25	18	103	0.01	3	0.16	18	0.40	142

\$1150 per kg, driven principally by operating expenditures (OpEx, Table 2, scenario A). However, we estimate that system throughput could be increased to 10 kg h⁻¹ for only twice the purchase cost while maintaining the single operator. This increase in scale results in a dramatic decrease in both capital (CapEx) and OpEx, for a total cost of \$142 per kg (scenario C). As part of the scale-up, the product collection equipment used in the lab-scale system is replaced with a more industrially relevant baghouse (bag filters), which produces a cost savings on its own (scenario B vs. A). Because Pt metal is routinely recovered from spent catalysts, we used the spent catalyst value (SCV) module of CatCost to estimate the recoverable value of Pt at the end of the catalyst's useful life. The catalysts were estimated to have a spent catalyst value of \$10.23 per kg across all scenarios, which is not included in Table 2.

The cost to produce Pt/TiO₂ *via* FSP was compared to that of a conventional IW synthesis. The IW-Pt(0.07)/TiO₂ cost estimates ranged from \$27–142 per kg, depending on the type and cost of the titania support used for IW. The most relevant comparison is to a flame-synthesized titania powder such as the P25 titania support we have used throughout this work. Through a survey of quotes and publicly available prices, we determined a price range for P25 of *ca.* \$50–100 per kg, depending on scale, purity, and other specifications. At the upper end of this range, the resulting catalyst cost of

\$142 per kg is identical to that of the FSP-derived material (Table 2, scenarios C and G). We also included a scenario using a low-cost (\$3 per kg), non-flame-synthesized titania. The lower Pt cost for IW-Pt(0.07)/TiO₂ relative to FSP-Pt(0.07)/TiO₂ results from differences in Pt utilization (yield), which was assumed to be quantitative for IW but was observed to be 70% for FSP (targeted loading: 0.10%, actual loading: 0.07%).

In order to assess major cost drivers and identify R&D opportunities for the FSP material, a sensitivity analysis was performed based on scenario C, and the sensitivity results for the top contributors to cost uncertainty are shown in Fig. 7. The figure uses net catalyst cost, after spent catalyst value is subtracted. The costs of IW materials based on P25 prices of \$75 and \$100 per kg are included on the plot for reference. Consistent with the large contribution of solvent costs in Table 2, the largest opportunity for cost reduction is in solvent usage. A proportional reduction of solvent usage by approximately 3-fold (33 to 10 L kg⁻¹) would yield cost savings of over \$35 per kg, resulting in a purchase cost lower than the \$75 per kg P25 IW scenario. Further opportunities for cost reduction through R&D include process gas usage and Pt utilization; on the latter point, it should be noted that industrial syntheses are expected to be substantially (*i.e.*, 90+%) more materials efficient. Combining the modelled savings from these three areas would reduce purchase cost by

**Fig. 7** Tornado plot depicting the results of a sensitivity analysis on the FSP-Pt(0.07)/TiO₂ material.

\$55 per kg total, to a cost of \$87 per kg, approaching the level of the \$50 per kg P25 IW scenario (\$83 per kg). Other factors driving significant cost variation include TTiP price, distribution and marketing costs, production scale, and plant capacity factor (the ratio of actual to expected plant outputs). It is noteworthy that the TTiP precursor price was only the third most significant factor contributing to variation in catalyst cost, revealing that commercial applications of a TTiP/FSP route to titania may be limited more by solvent cost than TTiP cost.

Conclusions

The continuous FSP synthesis presented here highlights the utility and tunability of this approach for the scalable production of high-performance Pt/TiO₂ catalysts for biomass upgrading applications. The *ex situ* CFP reaction studies demonstrate the (1) high deoxygenation activity of FSP-Pt/TiO₂ catalysts revealed by formation of intermediate oxygenates, aromatics, (cyclo)alkanes, and (cyclo)alkenes as the dominant hydrocarbon products, and (2) decreased hydrogenation as revealed by the high alkene and low alkane product yield when compared to IW-Pt/TiO₂. We hypothesize that the observed shift towards unsaturated hydrocarbons (alkenes, intermediate oxygenates, and aromatics) and improved deoxygenation performance at the lowest Pt loading compared to the IW materials results from the observed differences in Pt speciation and particle size. A preliminary cost analysis revealed several opportunities to improve the scalable FSP synthesis to make it more cost-competitive with traditionally synthesized IW catalysts including (1) reductions to solvent usage, (2) improving process gas usage, and (3) increasing Pt utilization. In conclusion, the FSP synthesis of Pt catalysts demonstrated process intensification of the synthesis procedure while offering changes to catalyst morphology that may be beneficial for tuning activity.

Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The usage of Electron Microscopy Service at University of Illinois at Chicago with help of Dr. Fengyuan Shi is gratefully acknowledged. This work was authored in part by the National Renewable Energy Laboratory, operated by Alliance for Sustainable Energy, LLC, and in part by Argonne National Laboratory, operated by The University of Chicago for the U.S. Department of Energy (DOE) under Contract No. DE-AC36-

08GO28308 and DE-AC02-06CH11357, respectively. This research used resources of the Advanced Photon Source; a U.S. DOE Office of Science User Facility operated for the DOE Office of Science by Argonne National Laboratory under Contract DE-AC02-06CH11357. MRCAT operations were supported by the DOE and the MRCAT member institutions. Funding provided by the U.S. Department of Energy Office of Energy Efficiency and Renewable Energy Bioenergy Technologies Office. This research was conducted in collaboration with the Chemical Catalysis for Bioenergy (ChemCatBio) Consortium, a member of the Energy Materials Network (EMN). The views expressed in this article do not necessarily represent the views of the DOE or the U.S. Government. The U.S. Government retains and the publisher, by accepting the article for publication, acknowledges that the U.S. Government retains a nonexclusive, paid-up, irrevocable, worldwide license to publish or reproduce the published form of this work, or allow others to do so, for U.S. Government purposes.

References

- 1 D. A. Ruddy, J. A. Schaidle, J. R. Ferrell III, J. Wang, L. Moens and J. E. Hensley, *Green Chem.*, 2014, **16**, 454–490.
- 2 J.-M. Ha, K.-R. Hwang, Y.-M. Kim, J. Jae, K. H. Kim, H. W. Lee, J.-Y. Kim and Y.-K. Park, *Renewable Sustainable Energy Rev.*, 2019, **111**, 422–441.
- 3 M. B. Griffin, K. Iisa, H. Wang, A. Dutta, K. A. Orton, R. J. French, D. M. Santosa, N. Wilson, E. Christensen, C. Nash, K. M. Van Allsburg, F. G. Baddour, D. A. Ruddy, E. C. D. Tan, H. Cai, C. Mukarakate and J. A. Schaidle, *Energy Environ. Sci.*, 2018, **11**, 2904–2918.
- 4 S. Boonyasuwat, T. Omotoso, D. E. Resasco and S. P. Crossley, *Catal. Lett.*, 2013, **143**, 783–791.
- 5 M. B. Griffin, G. A. Ferguson, D. A. Ruddy, M. J. Bidy, G. T. Beckham and J. A. Schaidle, *ACS Catal.*, 2016, **6**, 2715–2727.
- 6 M. Lu, H. Du, B. Wei, J. Zhu, M. Li, Y. Shan and C. Song, *Energy Fuels*, 2017, **31**, 10858–10865.
- 7 C. A. Teles, R. C. Rabelo-Neto, G. Jacobs, B. H. Davis, D. E. Resasco and F. B. Noronha, *ChemCatChem*, 2017, **9**, 2850–2863.
- 8 J. A. Schwarz, C. Contescu and A. Contescu, *Chem. Rev.*, 1995, **95**, 477–510.
- 9 B. A. T. Mehrabadi, S. Eskandari, U. Khan, R. D. White and J. R. Regalbuto, in *Advances in Catalysis*, ed. C. Song, Academic Press, 2017, vol. 61, pp. 1–35.
- 10 P. Munnik, P. E. de Jongh and K. P. de Jong, *Chem. Rev.*, 2015, **115**, 6687–6718.
- 11 F. G. Baddour, E. J. Roberts, A. T. To, L. Wang, S. E. Habas, D. A. Ruddy, N. M. Bedford, J. Wright, C. P. Nash, J. A. Schaidle, R. L. Brutchey and N. Malmstadt, *J. Am. Chem. Soc.*, 2020, **142**, 1010–1019.
- 12 S. E. Habas, F. G. Baddour, D. A. Ruddy, C. P. Nash, J. Wang, M. Pan, J. E. Hensley and J. A. Schaidle, *Chem. Mater.*, 2015, **27**, 7580–7592.
- 13 M. Schreier and J. R. Regalbuto, *J. Catal.*, 2004, **225**, 190–202.
- 14 J. T. Miller, M. Schreier, A. J. Kropf and J. R. Regalbuto, *J. Catal.*, 2004, **225**, 203–212.



- 15 Z. Li, S. Ji, Y. Liu, X. Cao, S. Tian, Y. Chen, Z. Niu and Y. Li, *Chem. Rev.*, 2020, **120**, 623–682.
- 16 F. G. Baddour, C. P. Nash, J. A. Schaidle and D. A. Ruddy, *Angew. Chem., Int. Ed.*, 2016, **55**, 9026–9029.
- 17 A. Dutta, J. A. Schaidle, D. Humbird, F. G. Baddour and A. Sahir, *Top. Catal.*, 2016, **59**, 2–18.
- 18 W. Y. Teoh, R. Amal and L. Mädler, *Nanoscale*, 2010, **2**, 1324–1347.
- 19 T. Johannessen, J. R. Jensen, M. Mosleh, J. Johansen, U. Quaade and H. Livbjerg, *Chem. Eng. Res. Des.*, 2004, **82**, 1444–1452.
- 20 S. Li, Y. Ren, P. Biswas and S. D. Tse, *Prog. Energy Combust. Sci.*, 2016, **55**, 1–59.
- 21 K. Wegner and S. E. Pratsinis, *Chem. Eng. Sci.*, 2003, **58**, 4581–4589.
- 22 R. Mueller, L. Mädler and S. E. Pratsinis, *Chem. Eng. Sci.*, 2003, **58**, 1969–1976.
- 23 G. D. Ulrich, *Chem. Eng. News*, 1984, **62**, 22–29.
- 24 R. Koirala, S. E. Pratsinis and A. Baiker, *Chem. Soc. Rev.*, 2016, **45**, 3053–3068.
- 25 R. Strobel, A. Baiker and S. E. Pratsinis, *Adv. Powder Technol.*, 2006, **17**, 457–480.
- 26 A. Varma, A. S. Mukasyan, A. S. Rogachev and K. V. Manukyan, *Chem. Rev.*, 2016, **116**, 14493–14586.
- 27 S. Ding, H.-A. Chen, O. Mekasuwandumrong, M. J. Hülsey, X. Fu, Q. He, J. Panpranot, C.-M. Yang and N. Yan, *Appl. Catal., B*, 2021, **281**, 119471.
- 28 W. Y. Teoh, L. Mädler, D. Beydoun, S. E. Pratsinis and R. Amal, *Chem. Eng. Sci.*, 2005, **60**, 5852–5861.
- 29 D. A. Ruddy, J. E. Hensley, C. P. Nash, E. C. D. Tan, E. Christensen, C. A. Farberow, F. G. Baddour, K. M. Van Allsburg and J. A. Schaidle, *Nat. Catal.*, 2019, **2**, 632–640.
- 30 S. E. Pratsinis, *AIChE J.*, 2010, **56**, 3028–3035.
- 31 P. Weyell, H. D. Kurland, T. Hülser, J. Grabow, F. A. Müller and D. Kralisch, *Green Chem.*, 2020, **22**, 814–827.
- 32 K. M. Van Allsburg, E. C. D. Tan, J. D. Super, J. A. Schaidle and F. G. Baddour, *Nat. Catal.*, 2022, **5**, 342–353.
- 33 F. G. Baddour, L. Snowden-Swan, J. D. Super and K. M. Van Allsburg, *Org. Process Res. Dev.*, 2018, **22**, 1599–1605.
- 34 T. B. Bolin, T. Wu, N. Schweitzer, R. Lobo-Lapidus, A. J. Kropf, H. Wang, Y. Hu, J. T. Miller and S. M. Heald, *Operando IV 4th Int. Congr. Operando Spectrosc.*, 2013, vol. 205, pp. 141–147.
- 35 B. Ravel and M. Newville, *J. Synchrotron Radiat.*, 2005, **12**, 537–541.
- 36 J. J. Rehr, J. Mustre de Leon, S. I. Zabinsky and R. C. Albers, *J. Am. Chem. Soc.*, 1991, **113**, 5135–5140.
- 37 C. Mukarakate, M. J. Watson, J. ten Dam, X. Baucherel, S. Budhi, M. M. Yung, H. Ben, K. Iisa, R. M. Baldwin and M. R. Nimlos, *Green Chem.*, 2014, **16**, 4891–4905.
- 38 C. Mukarakate, X. Zhang, A. R. Stanton, D. J. Robichaud, P. N. Ciesielski, K. Malhotra, B. S. Donohoe, E. Gjersing, R. J. Evans, D. S. Heroux, R. Richards, K. Iisa and M. R. Nimlos, *Green Chem.*, 2014, **16**, 1444–1461.
- 39 C. Mukarakate, K. Orton, Y. Kim, S. Dell'Orco, C. A. Farberow, S. Kim, M. J. Watson, R. M. Baldwin and K. A. Magrini, *ACS Sustainable Chem. Eng.*, 2020, **8**, 2652–2664.
- 40 A. R. Stanton, K. Iisa, C. Mukarakate and M. R. Nimlos, *ACS Sustainable Chem. Eng.*, 2018, **6**, 10030–10038.
- 41 K. Iisa, Y. Kim, K. A. Orton, D. J. Robichaud, R. Katahira, M. J. Watson, E. C. Wegener, M. R. Nimlos, J. A. Schaidle, C. Mukarakate and S. Kim, *Green Chem.*, 2020, **22**, 2403–2418.
- 42 D. E. Garrett, *Chemical Engineering Economics*, Van Nostrand-Reinhold, New York, 1989.
- 43 G. D. Ulrich and P. T. Vasudevan, *Chemical Engineering Process Design and Economics*, Durham, New Hampshire, 2nd edn, 2004.
- 44 H. J. Lang, *Chem. Eng.*, 1947, 117–121.
- 45 H. J. Lang, *Chem. Eng.*, 1948, 112–113.
- 46 M. S. Peters and K. D. Timmerhaus, *Plant Design and Economics for Chemical Engineers*, McGraw-Hill, New York, 5th edn, 2003.
- 47 E. Sánchez, T. López, R. Gómez, Bokhimi, A. Morales and O. Novaro, *J. Solid State Chem.*, 1996, **122**, 309–314.
- 48 K. J. Range, F. Rau, U. Klement and A. M. Heyns, *Mater. Res. Bull.*, 1987, **22**, 1541–1547.
- 49 K. Sugiyama and Y. Takéuchi, *Z. Kristallogr. Cryst. Mater.*, 1991, **194**, 305–313.
- 50 T. Sasaki, N. Koshizaki, S. Terauchi, H. Umehara, Y. Matsumoto and M. Koinuma, *Nanostruct. Mater.*, 1997, **8**, 1077–1083.
- 51 T. Sasaki, N. Koshizaki, M. Koinuma and Y. Matsumoto, *Nanostruct. Mater.*, 1999, **12**, 511–514.
- 52 X. Sun, J. Lin, Y. Chen, Y. Wang, L. Li, S. Miao, X. Pan and X. Wang, *Commun. Chem.*, 2019, **2**, 27.
- 53 T. I. Hyde, P. W. Ash, D. A. Boyd, G. Randschofer, K. Rothenbacher and G. Sankar, *Platinum Met. Rev.*, 2011, **55**, 233–245.
- 54 Y. Zhou, D. E. Doronkin, M. Chen, S. Wei and J.-D. Grunwaldt, *ACS Catal.*, 2016, **6**, 7799–7809.
- 55 A. D. Allian, K. Takanabe, K. L. Fajdala, X. Hao, T. J. Truex, J. Cai, C. Buda, M. Neurock and E. Iglesia, *J. Am. Chem. Soc.*, 2011, **133**, 4498–4517.
- 56 K. Ding, A. Gulec, A. M. Johnson, N. M. Schweitzer, G. D. Stucky, L. D. Marks and P. C. Stair, *Science*, 2015, **350**, 189–192.
- 57 P. Bazin, O. Saur, J. C. Lavalley, M. Daturi and G. Blanchard, *Phys. Chem. Chem. Phys.*, 2005, **7**, 187–194.
- 58 C. A. Teles, P. M. de Souza, R. C. Rabelo-Neto, M. B. Griffin, C. Mukarakate, K. A. Orton, D. E. Resasco and F. B. Noronha, *Appl. Catal., B*, 2018, **238**, 38–50.
- 59 M. Xu, C. Mukarakate, K. Iisa, S. Budhi, M. Menart, M. Davidson, D. J. Robichaud, M. R. Nimlos, B. G. Trewyn and R. M. Richards, *ACS Sustainable Chem. Eng.*, 2017, **5**, 5477–5484.
- 60 K. Murugappan, C. Mukarakate, S. Budhi, M. Shetty, M. R. Nimlos and Y. Román-Leshkov, *Green Chem.*, 2016, **18**, 5548–5557.

