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Supported nanocatalysts: recent developments in microwave synthesis for application in heterogeneous catalysis†

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The synthesis of nanocatalysts on high surface area solid supports is gaining attention from synthetic chemists for application in heterogeneous catalysis due to their high catalytic performance and easy recoverability. However, the synthesis of nanocatalysts on solid supports mostly composed of metals/metal oxides/hybrids is challenging for researchers considering their durability and discrete nano-dispersibility. Microwave (MW)-assisted synthetic methods, attractive as fast and green approaches, are being widely studied for the synthesis of solid-supported nanocatalysts. This review article describes recent advances devoted towards MW-assisted synthetic strategies for creating nanostructured catalysts on solid supports, as well as their characterization and applications in catalyzing various organic transformations, including electrocatalysis. The sustainable use of solid-supported nanocatalysts is then discussed in comparison with similar supported nanocatalysts prepared by conventional methods. The pros and cons of MW-assisted strategies for the synthesis of solid-supported nanocatalysts are then summarized, and their limitations as well as future directions for further investigation are emphasized.

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1. Introduction

To meet the rising demand of fine chemicals and to reduce the dependency on the use of expensive and non-recyclable homogeneous catalysts, synthetic chemists are searching for more durable nanostructured heterogeneous catalysts. Compared to bulk materials, nanostructured materials have

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proved beneficial in catalysis because they offer improved physicochemical properties such as high surface-to-volume ratio, surface energy, active sites and better composition-based selectivity towards particular reaction pathways.^{1,2} Additionally, the vast improvement in characterization tools has also enabled researchers to follow catalytic reactions from nanoparticles (NPs) down to even single atoms.³

Transition metals (Ti, Cr, Mn, Fe, Ni, Cu, Ru, Pd, Pt, Au and Ag), which can easily donate and accept electrons from molecules, have been considered ideal for heterogeneous catalysis in many organic and bioorthogonal syntheses.^{4,5} However, nanocatalysts derived from transition metals/metal oxides/blends are prone to aggregation due to stronger inter-particle dipolar forces compared to their bulk counterparts, thereby reducing their potential for long-time application and preservation. Some transition metals (Cu, Ni) are also unstable and readily oxidized in air, preventing their sustainable use.⁶ The modification of transition metal nanocatalysts with functionalizing agents such as surfactants^{7,8} or polymers⁹ has helped to improve their stability against aggregation, but such modification often deteriorates their catalytic properties *via* changes in the electronic structure of the metal nanocatalysts.¹⁰ An alternative way proposed to overcome aggregation and the deterioration of catalytic properties is the implanting or anchoring of nanocatalysts on inert low-density solid supports. Solid supports help to maintain the size, shape, and morphology of the nanocatalysts and also limit the leaching of metal ions, which is favorable for achieving durable and enhanced catalytic performance. The solid supports used prominently include graphene,^{11,12} zeolites,¹³ cellulose,^{14–16} activated coke/carbon,^{17,18} sepiolite,¹⁹ bentonite,²⁰ SiO₂,^{21,22} and Al₂O₃.^{23,24} In general, the preparation of supported nanocatalysts is usually conducted in two steps. In the first step, the deposition or impregnation of the metal precursor on a solid support is achieved *via* impregnation/precipitation/coprecipitation/electrostatic/co-ordination, depending on the support functional groups.^{25,26} The second step involves calcination/reduction/both by different techniques such as chemical reduction,^{24,27} sonochemical reduction,^{28,29} laser ablation,^{30,31} and thermal decomposition.³² These methods generally require high temperatures, long reaction times and inert atmosphere. All these factors adversely affect the large-scale synthesis of supported nanocatalysts, and therefore the search for simple and scalable synthetic methods has become imperative. Recently, microwave (MW) irradiation/heating has gained popularity for the synthesis of metal/metal oxide nanocatalysts due to the fast and uniform heating of the reaction medium, which results in fast reaction kinetics,^{33,34} uniform growth of the NPs,^{35–37} easy operation and green synthesis.³⁸

In this review, we discuss the recent articles on supported nanocatalysts prepared using MW strategies. The nanocatalysts discussed herein include metals, metal oxides, multi-metals, mixed metal–metal oxides and metal carbides. However, the few studies available on metal sulfides, nitrides and phosphides are not included. Some review articles on the optimization of catalytic performance by controlling the size, shape and morphology of different metal nanocomposites, both supported

and non-supported, are available.^{39–43} However, these reviews hardly considered all types of metals and metal oxides, rather focusing on a group of metals or metal oxides. Jamkhande *et al.* reviewed the different methods used for the preparation of metal NPs and focused on their advantages, disadvantages and applications.⁴⁴ In a recent review, MW chemistry was discussed briefly to understand the formation mechanism and growth of particles, focusing on improving the performance of MW methods for future research.⁴⁵ In this review, we discuss the latest articles on the MW-assisted synthesis of transition metals, metal oxides, hybrids and perovskite-type nanocatalysts supported on solid supports. Initially, the chemistry of MW synthesis is shortly outlined, and then the preparation of various types of nanocatalysts is discussed. Subsequently, an explicit review is presented on the applications of supported nanocatalysts in various organic transformations. Particular emphasis is given to the recyclability/durability of supported nanocatalysts and the comparison of their performance with that of similar nanocatalysts prepared *via* other methods. Finally, the future challenges and ways of further improving the catalytic performance of nanocatalysts prepared *via* the MW method are described in the Scope and outlook section.

2. MW chemistry and its applications

Since the 1950s, MW energy was found to be useful for heating foodstuffs, food processing, chemical drying, *etc.* Upon irradiation with MW, the dielectric heating creates a heating effect on matter and the heating efficiency depends on the ability of the substance to interact with MW radiation.⁴⁶ In 1986, two research groups first reported the utilization of MW warming to speed up organic chemical changes.^{47,48} In the early days, a household oven was used to perform experiments in sealed Teflon or glass vessels with no mechanism for controlling the temperature or pressure. This fast and uncontrolled heating of organic solvents later encouraged researchers to use solvent-free MW chemistry.⁴⁹ The solvent free methodology is based on the pre-adsorption of reagents and catalysts on MW transparent inorganic supports, enabling controlled reactions in MW-assisted organic synthesis. With continuous improvement in MW technology and the availability of online temperature and pressure control, in the mid-90s Strauss developed a method of organic synthesis in a sealed vessel using a conventional solvent.⁵⁰ Consequently, MW heating/irradiation has now become a recognized tool in all areas of organic synthesis including both solvent-free and water-mediated reactions.^{51–54} Additionally, the further development of MW reactor technology now facilitates the synthesis of nanomaterials.^{55–58} Recently, the continuous flow microfluidic approach has been attempted in MW-assisted synthesis to improve the control of the temperature gradient, rate of heating and mixing, and also to enable fast quenching to stop prolonged nucleation events.⁵⁹

The frequency and wavelength of MW radiation (*i.e.*, electromagnetic radiation) are in the range of 0.3–300 GHz and 1 mm–1 m, which are between the infrared and radio region in the electromagnetic range. Commercial and domestic



MW ovens operate at a frequency of up to 2.45 GHz to eliminate interference from the communication frequency bands of cellular phones. MW radiation travels with the speed of light and provides energy that is insufficient to break a chemical bond, which in most cases does not affect the framework of chemical molecules.⁶⁰ MW heating depends on the capacity of materials to absorb and transform MW radiation into heat. Three categories of materials including MW reflectors, MW transmitters and MW absorbers describe the interaction with MW. Metals classified as MW reflectors are not effectively heated by MW. Materials that are transparent to MW (e.g., Teflon and polystyrene) are classified as MW transmitters and ideal for use as MW reactor and vessel materials. Materials known as MW absorbers can absorb energy from the MW field (e.g., solvents/reactants) and are heated very quickly in the process.⁶¹ The addition of a small amount of ionic liquid, a strong MW absorbing agent, to a non-polar solvent generally enhances the heating rate, and hence increases the reaction rate and product yield.⁶²

MW-assisted reactions proceed at lightning speed, reducing the reaction time from several hours to minutes due to the reduction in the activation energy barrier.⁶³ For example, the MW-irradiated green synthesis of Ag nanoclusters (2 nm) was possible within just 70 s,⁶³ whereas the sonochemical method required 1 h to produce AgNPs,²⁸ and chemical reduction of the precursor AgNO₃ required an even longer reaction time (6 h).⁶⁴ The conventional heating process using hot plates or mantles is slow and inefficient for transferring energy to the reactants, given that it depends on convective currents and the conductive nature of various compounds and the materials to be penetrated. This often produces a temperature gradient between the reaction medium and vessel, which is responsible for non-uniform growth, hindering scaled-up production. In contrast, MW irradiation causes volumetric heating, i.e., uniform temperature rise throughout the liquid volume *via* direct coupling of MW energy to the molecules present in the reaction vessel.^{65–67} This volumetric heating in the synthesis of nanomaterials causes rapid and homogeneous heating, which reduces the temperature gradient and provides instantaneous nucleation, controlled and uniform growth with good reproducibility in terms of size, shape and morphology.⁴⁵ The shorter reaction time reduces the probability of the formation of byproducts through side reactions and increases the product yield, homogeneity and crystallinity.³⁸

Thus far, MW-assisted irradiation/heating has been attempted for the synthesis of a large variety of unsupported mono- and bi-metallic NPs as well as different types of simple to quaternary metal oxides.^{68–76} The morphology and average size can be controlled by the precursor type, concentration, solvent type, pH and MW heating conditions such as power, pressure, temperature and time.

3. Synthesis and characterization

3.1. Simple metal catalysts

3.1.1. Carbon supported. Carbon materials are considered attractive for applications in the fields of adsorption,

separation, sensors, energy storage, hydrogen storage, catalysts and catalyst supports given that they exhibit some interesting properties such as large specific surface area, large pore volume, tunable mesoporous structure, high electron conductivity and high thermal and chemical durability.^{77–80} Different types of carbon materials such as graphite (Grp), carbon black, activated carbon (AC), nanotubes, and graphene (G) have been used as support materials for catalysts in MW-assisted synthesis.

Chuang *et al.* observed changes in the morphology of a Cu/AC catalyst due to a variation in MW irradiation power (400, 700 and 900), irradiation time (45, 60 or 110 s), and pH (4, 7, 10 and 12) of the polyol precursor during its preparation from a homogeneous solution of Cu precursor salt in ethylene glycol (EG).⁸¹ The obtained solids from the MW reactor were washed and dried, and then calcined under an H₂/He atmosphere at 475 °C for 4 h before characterization. The maximum surface area and pore size were observed when the MW power was set at 700 W for 60 s and the pH of the polyol precursor was 7 (Cu/AC-700 W-60 sec-7). Under these conditions, Cu⁰ was the dominant phase. In contrast, the minimum 16 nm-sized Cu cubes (some aggregated) were formed when the pH of the polyol precursor was 12. Ren *et al.* employed pre-impregnation of AC with an aqueous Cu(NO₃)₂·3H₂O solution and carbothermic MW irradiation at three different temperatures of 150 °C, 410 °C and 540 °C.⁸² The characteristic diffraction signal due to Cu⁰ was the strongest in Cat-48-540 among the catalysts, and optimistically a decrease in the average size of the cubic Cu₂O and Cu⁰ particles from 43 ± 12 to 37 ± 12 nm was observed with an increase in irradiation temperature from 410 °C to 540 °C. The higher irradiation temperature was favorable to produce a higher Cu⁰ content. The same group also prepared hollow Cu/AC *via* template-free carbothermic MW reduction using a homogeneous solution of Cu(NO₃)₂·3H₂O.⁸³ The formation mechanism of hollow CuNPs was explained based on the Ostwald ripening effect and self-assembly under MW-induced reduction (Fig. 1). Under the optimum conditions, hollow CuNPs with large cavities of ~35 nm diameter and 50 nm outer diameter were obtained within 8 min when the precursor was heated from room temperature to 360 °C.

An Fe/AC catalyst was prepared *via* the impregnation of Fe(NO₃)₃·9H₂O in AC, drying and MW treatment for 2 h at 400 °C under an He flow.⁸⁴ The theoretical ratio of Fe to AC was 10 wt%. The Brunauer–Emmett–Teller (BET) surface area of the catalyst obviously decreased in the Fe loading process; however, microscopic evidence was not provided.

Veerakumar *et al.* generated 3D ordered mesoporous carbon materials (CPMs) for supporting RuNPs by subjecting a cured mixture of polymer and Ru(acac)₃ to MW thermal reduction and carbonization.⁸⁵ The catalysts were labeled as Ru/CPM-1 and Ru/CPM-2 based on 0.5 and 1.0 wt% Ru loading, respectively. Hexagonal close-packed (hcp) Ru⁰NPs on a 3D CPM matrix with a crystallite size of 3–5 nm were detected and the average size of the RuNPs from transmission electron microscopy (TEM) was 2.6 ± 0.5 nm. The same group employed porous plastic-derived carbon (PDC) to prepare a PDC@RuNP catalyst by applying MW irradiation to a mixture of Ru(acac)₃ and PDC in tetrahydrofuran



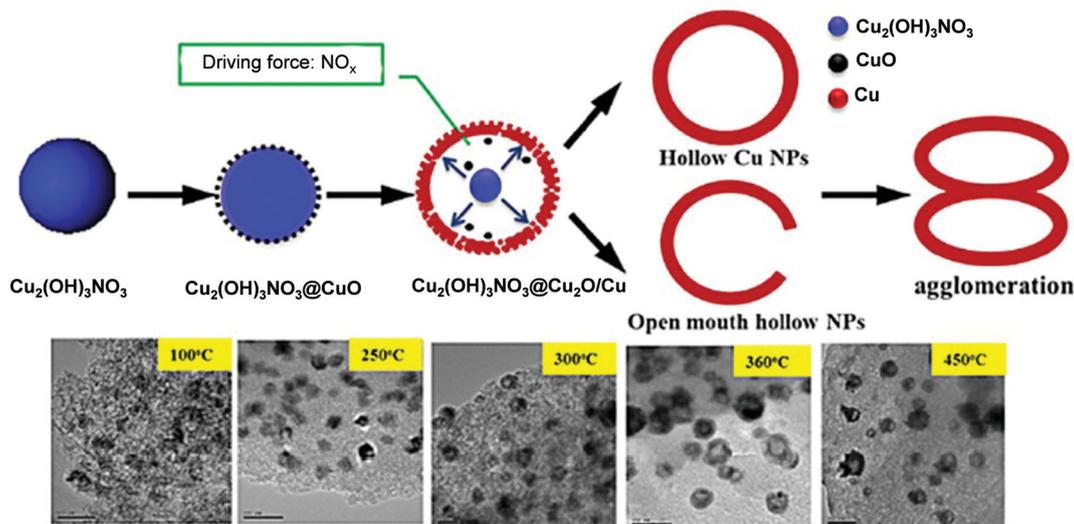


Fig. 1 Schematic representation of the Ostwald ripening-coupled self-assembly process of the hollow Cu structures.⁸³ (Copyright 2016, Wiley-VCH.)

at 300 W for 1 h.⁸⁶ Very fine Ru^0 crystallites with an average size of 5–6 nm were formed under these conditions.

The chemistry of supported Pt catalysts on carbon-based materials *via* MW synthesis started at the beginning of the 21st century.^{87,88} Here, they discussed the preparation of homogeneously distributed <4 nm sized PtNPs on either carbon or carbon nanotubes (CNT). In most cases Pt had a face centered cubic structure (fcc) and contained Pt^0 as the major component. Later, Bo *et al.* used coal-derived granular AC (GAC) for the preparation of a Pt/GAC catalyst *via* MW irradiation.⁸⁹ GAC was impregnated with an $\text{H}_2\text{PtCl}_6 \cdot 6\text{H}_2\text{O}$ aqueous solution, evaporated to near-dryness in an MW oven at 500 W, and then calcined by MW at 500 W for 5 min. The Pt particles on the porous GAC support had an asymmetric dispersion and irregular shape with a size of 100–800 nm. Comparatively, Gan *et al.* obtained homogeneously distributed PtNPs (3–5 nm) on CNT *via* a one-pot MW polyol method using a mixture of H_2PtCl_6 , EG and sodium dodecyl sulfate (SDS).⁹⁰ The anchored Pt had an fcc crystalline structure and average crystallite size of 4.6 nm. A similar polyol method also produced well-dispersed spherical 2.7 nm-sized PtNPs in a Pt/carbon aerogel (CA) catalyst.⁹¹ Pt had an identical fcc structure and the average crystallite size (2.88 nm) matched the microscopic image diameter. Sharma *et al.* observed an improvement in the Pt loading up to 23.7 wt% in Pt/C electrocatalysts when the $(\text{NH}_4)_2\text{PtCl}_6$ precursor was used in the MW-assisted polyol method.⁹² In contrast, the H_2PtCl_6 -sourced and commercial 20% Pt/C catalysts contained 19.5 and 18.8 wt% Pt, respectively. Independent of the nature of the precursor, the average size of the PtNPs was around 2 nm. Thus, the precursor type is important during the MW synthesis of supported catalysts.

Pentsak *et al.* studied the deactivation process of Pd/C catalytic systems during catalysis under MW and conventional heat treatment (CHT).⁹³ The catalyst preparation method was not detailed, but most likely wet impregnation and thermal decomposition were employed. Substantial transformation of

the carbon support *via* the formation of pits, trenches, nanofibers and nanowalls during catalysis under both MW and CHT was observed. The narrow size distribution of PdNPs (6 nm) increased to 62 nm after 30 min of MW catalysis. The movement of hot metal particles on the carbon surface turned the carbon surface into a porous spongy framework. The initial Pd^0 also turned into PdO (88%) after 30 min of MW treatment in air. These transformations of the metal support were more dominant upon MW treatment due to the easy formation of metal hot spots. The same research group further showed that the modification of the carbon support could be minimized if MW irradiation is carried out under vacuum and that the selection of an appropriate metal salt is also crucial.⁹⁴

Antonetti *et al.* used a low-boiling point alcohol or a mixture of alcohol (isopropanol/ethanol)/water as the solvent and a reducing agent to prepare RuNPs/CNT under MW irradiation.⁹⁵ The MW-assisted synthesis of unsupported RuNPs in isopropanol was first optimized using poly(*N*-vinyl-2-pyrrolidone), PVP, as a stabilizing agent, and finally the idea of MW synthesis was extended to the *in situ* preparation of RuNPs/CNT. Two types of CNTs were used, one was purified CNT (*via* treatment with HCl) and the other was HNO_3 -treated purified CNT (FCNT). Independent of the CNT type, the average diameter of the RuNPs was between 2.7 to 3.6 nm. However, the carboxylic and oxygenated groups on FCNT produced a more homogeneous distribution of RuNPs and sufficient to stabilize the NPs without the need for PVP. The same *in situ* approach using FCNT was also evaluated in aqueous ethanol (EtOH/ H_2O 20/80) at 150 °C under excess pressure. Again, small-sized particles with a homogeneous distribution were confirmed. In contrast, the solvent-free one-pot MW thermolytic route was found to be unsuitable for the production of well-dispersed Ru on FCNT.^{96,97} Hemraj-Benny *et al.* used non-functional single-walled CNT (SWCNT) to prepare an SWCNT-Ru catalyst *via* the reduction of RuCl_3 in an ethanolic solution of NaBH_4 .⁹⁸ The average size of the RuNPs was dependent on the MW temperature, where 100 °C yielded smaller (2.0 ± 0.5 nm)



Ru⁰NPs. In contrast, a higher temperature caused partial agglomeration. In another investigation, the ethanol reduction of RuCl₃ at 150 °C was examined to determine the effect of the weight percentage (0.05–1.00 wt%) of RuCl₃ on the size distribution of the RuNPs in an SWCNT-Ru catalyst.⁹⁹ An increase in RuCl₃ content from 0.05 to 1.00 wt% resulted in an increase in particle size from 2.0 ± 0.5 nm to around 45 nm Ru clusters.

Graphene (G), an allotrope of carbon with a hexagonal atomic layer structure, is another important support material with unique properties such as high surface area and good thermal, chemical and mechanical stability.^{100–104} Fei *et al.* prepared a series of embedded monodispersed atomic transition metals in nitrogen-doped G (NG) *via* MW heating.¹⁰⁵ The prepared nanocomposites were named M-NG-MW, where M represents Co/Ni/Cu. Initially, an aqueous solution of GO was reacted with ammonia solution at 70 °C to produce amine-functionalized GO (AGO). Typically, a certain amount of CoCl₂ solution was well mixed with the AGO solution and the freeze-dried product was MW irradiated in an argon atmosphere. The amino N in Co-AGO was the dominant species, but in Co-NG-MW the amino N signal vanished due to its interaction with the uniformly distributed metal atoms.

In 2011, Siamaki *et al.* prepared Pd/G *via* batch-wise MW irradiation for 60 s.¹⁰⁶ For this preparation, a mixture of dried graphite oxide (GrpO), requisite amount of Pd(NO₃)₂ and HNO₃ was microwaved with N₂H₄ reducing agent following sonication. The complete reduction of GrpO to G by the reducing agent visibly improved the thermal stability of Pd/G compared to Pd/GrpO. Subsequently, the success of the continuous scalable flow approach was tested with respect to the reproducibility and yield of the Pd/G catalyst.¹⁰⁷ A homogeneous suspension of GrpO in water containing Pd(NO₃)₂ and N₂H₄ solution was fed from two separate vessels at an equal rate into the MW reactor operating at 80 °C with an overall flow rate of 4 mL min⁻¹. The produced Pd/G had an average particle diameter of 9.37 nm and contained 80% Pd⁰ with the rest being Pd(II). Alternatively, Yang *et al.* employed a batch MW-assisted reduction approach under 1.2 MPa H₂ pressure to prepare Pd/G, Pd/AC, Pd/Grp and Pd/SiO₂ catalysts using a mixture of support and [Pd(OAc)₂]_n in 0.1 M SDS.¹⁰⁸ They found that the nature of the support influenced the Pd content in the catalyst.

Recently, Elazab *et al.* used a one-step MW irradiation approach to prepare magnetic Pd-Fe₃O₄/G from reduced GO sheets (rGO).¹⁰⁹ In a typical procedure, Pd(NO₃)₂ was added to an aqueous solution of FeCl₃ and reduced under MW condition using N₂H₄. Well-dispersed Fe₃O₄ particles were observed. A similar method was also tested to prepare Pd/G, Fe₃O₄/G nanocomposites using Pd(NO₃)₂ and FeCl₃ precursors.

Sulfonated G and GrpO (100 g) were evaluated to prepare Pt/G and Pt/GrpO by heating a mixture of crushed K₂PtCl₆ (variable: 14.5, 355, and 15 mg) and ionic liquid, 2-hydroxyethanaminium formate, (5.0 g) in a MW oven.¹¹⁰ PtNPs with a size of 5 to 30 nm were formed on sulfonated G or GrpO. The shape of the PtNPs on G was spherical, whereas that on GrpO was cubic. A higher amount of functional oxygen groups on GrpO increased the anchoring efficiency of Pt and also

influenced the shape and size of the produced NPs. The one-pot MW-assisted polyol synthesis of Pt/G nanocomposite and subsequent heating at 500 °C produced flaky crystalline G sheets.¹¹¹ The Pt/G3 (0.50%) nanocomposite was comprised a comparatively uniform distribution of 7–11 nm-sized PtNPs.

3.1.2. Oxide-supported catalysts. In MW synthesis, different types of oxide minerals such as clay minerals (*e.g.* SiO₂, vermiculite, and bentonite) and metal oxides (*e.g.* Al₂O₃, MgO, TiO₂, and CeO₂) have been used as solid supports for various metal NPs. As support materials, many of them possess interesting and desirable properties including high thermal and chemical stability, high volume to surface area, easy functionalization capacity, low cost and easy availability.

3.1.2.1. Silica. Zuliani *et al.* used ALSBA-15, a mesoporous Al-coordinated SiO₂ material, as a support for the preparation of MW-2%Ag@ALSBA-15 and MW-2%Au@ALSBA-15 *via* the simple MW *in situ* reduction of an ethanolic mixture of AgNO₃ (0.04 mmol)/AuBr₃ (0.02 mmol) at 150 °C for 3 min.¹¹² The abundant hydroxyl groups on the support surface stabilized the metal NPs.^{113,114} Alternatively, the mechanochemical ball milling (BM) technique was used to prepare the BM-2%Ag@ALSBA-15 and BM-2%Au@ALSBA-15 nanocatalysts developed by other researchers.¹¹⁵ However, a detailed characterization in terms of size distribution, valence state and crystallinity was not presented. Manno *et al.* had two different MW synthetic approaches, *i.e.*, batch synthesis (three different times: 70, 35 and 17 s) and continuous flow microfluidic synthesis (Fig. 2), to anchor Ag nanoclusters (AgNCs) on mesoporous amine-functionalized SBA-15 using an acrylic acid Na salt as a reducing and stabilizing agent.^{116,117} In the batch synthesis, an increase in irradiation time resulted in an increased in the mean particle size and the most uniform AgNCs (1.6 ± 0.7 nm) were obtained at 17 s. Slow cooling after MW heating in the batch system caused aggregation and size evolution of the clusters. In contrast, in a continuous flow reactor, rapid quenching to 4 °C within 21 s favored a reduction in the average diameter to 1.2 ± 0.3 nm. The continuous microfluidic approach produced better reproducibility and comparable inter-batch size distribution.

Jiang *et al.* used 40% H₂SO₄-treated bentonite (AB) to prepare a Ni/AB catalyst *via* a MW-assisted drying process (250 W, 10 min) and subsequent calcination (550 °C, 3 h).¹¹⁸ The impregnation slurry was made from an ethanol–water solution of Ni(NO₃)₂·6H₂O, cetyltrimethylammonium bromide (CTAB) and AB suspension and the obtained catalyst was named Ni/AB-MW. Another batch of catalysts, named Ni/AB-CH, was prepared *via* conventional drying and calcination. MW-assisted drying improved the loading amount of Ni (16.1%) in Ni/AB-MW and also favored the formation of smaller Ni and NiO crystallites due to the better interaction of Ni with the AB support.

Akay *et al.* used the combination of MW radiation, heat treatment and plasma reduction under a stream of H₂ for 24 h at 400 °C and 550 °C to prepare porous Co/SiO₂ catalysts.¹¹⁹ The reduction under an H₂ stream at 550 °C produced scattering signals due to the Co crystal lattice and CoO but reduction at



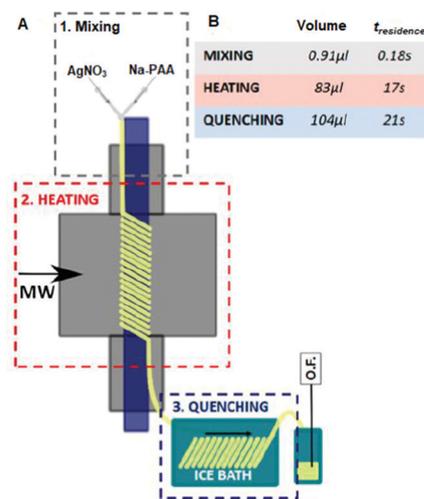


Fig. 2 (A) Schematic representation of the flow diagram for the continuous synthesis of AgNCs. Three different zones: mixing stage (grey color), MW heating zone (red color) and quenching stage (blue color). (B) Volume and calculated residence time for each stage (total flow rate 0.292 mL min⁻¹).¹¹⁶ (Copyright 2020, the American Chemical Society.)

400 °C did not result in any detectable metallic Co. The crystallite size of metallic Co at 550 °C was 15.6 nm, larger than that of CoO (5.01 nm). Simultaneously, another group attempted to prepare Co/SiO₂ *via* a similar four-step process, including the impregnation of porous SiO₂ in Co(NO₃)₂·6H₂O, MW drying, furnace drying at 550 °C, and finally reduction under an H₂ flow at 550 °C for 24 h.¹²⁰ The scanning electron microscopy (SEM) and TEM images revealed that the MW catalyst had a better distribution and uniformity (crystallite size, 3–8 nm) and loading of CoNPs compared to that prepared *via* the conventional drying method. Similar to the previous literature,¹¹⁹ this investigation also suggested that the final reduction under an H₂ flow is necessary to produce the Co metallic phase.

The MW *in situ* reduction of a dried SiO₂-impregnated H₂PtCl₆·6H₂O precursor in EG containing N₂H₄ produced fine well-dispersed ≤ 2 nm-sized PtNPs on an SiO₂ support at the loading of 0.3 wt%.¹²¹ An increase in the Pt loading to 0.9 wt% slightly increased the particle size to <10 nm. Comparatively, the rotary chemical evaporation (RCE) method produced larger (≥ 20 nm)-sized PtNPs. The comparative study revealed that the MW-assisted reduction method is superior with respect to the organized distribution of smaller PtNPs on the surface of SiO₂.

Wang *et al.* prepared a series of Ti-MCM-41 with different Si/Ti ratios (100 to 5) under MW irradiation using CTAB template, TEOS and tributyl titanate (TBOT).¹²² The obtained materials were calcined at 550 °C for 6 h in air to decompose the template. The Ti contents in the catalyst samples closely resembled the theoretical values. The arrangement of channels was slightly disturbed with an increase in the Ti content. Earlier, another research group prepared ZrMCM-41 mesoporous molecular sieves using Zr(SO₄)₂ and CTAB under MW irradiation.¹²³ The BET specific surface area and pore volume decreased with an increase in

the incorporated Zr. The mesoporous ordering deteriorated with an increase in the thermal treatment temperature.

3.1.2.2. Fe₃O₄ and analogous oxides. Fe₃O₄ NPs possess interesting application properties as a catalyst support because the presence of Fe²⁺ ions provides some reducible oxide characters.¹²⁴ Additionally Co₃O₄ and Ni(OH)₂ also show good activity in the oxidation of CO.^{125,126} Elazab *et al.* proposed a method for the preparation of PdNPs supported on Fe₃O₄, Co₃O₄ and Ni(OH)₂ NPs and nanoplates *via* MW irradiation.¹²⁷ In a typical process for the preparation of Pd/Fe₃O₄ containing 5, 20, 40 and 50 wt% Pd, mixtures of Fe(NO₃)₃·9H₂O and Pd(NO₃)₂ solutions (10 wt% in 10 wt% HNO₃) were sonicated for 1 h, and then irradiated in the presence of N₂H₄ in 60 s cycles for a total 7 min. Similarly, with slight modification, Pd/Co₃O₄ and Pd/Ni(OH)₂ catalysts containing variable wt% of Pd were prepared. The majority of the PdNPs were deposited on the surface of nanoplates. In another study, Elazab *et al.* reported the synthesis of a Pd/Fe₃O₄ catalyst *via* continuous flow synthesis under MW irradiation.¹²⁸ In this approach, two separate solutions, one containing the required amount of Fe(NO₃)₃·6H₂O and another containing N₂H₄, were injected and mixed before being introduced in the reactor. Different operating temperatures (80 °C, 120 °C and 150 °C) and flow rates were studied. The XPS binding energy showed that a fraction of Pd was in PdO. An increase in MW temperature resulted in an increase in size and the saturation magnetization was reduced when the particle size was smaller. Continuous flow reaction was found to be advantageous for producing smaller (7–9 nm) PdNPs than that (12–14 nm) prepared under batch conditions.

Morad *et al.* prepared AuNPs on an Mn₃O₄ support by reducing an HAuCl₄ hydrate solution using N₂H₄ under MW irradiation.¹²⁹ Mn(OH)₂ gel was prepared *via* direct precipitation from Mn(NO₃)₂·4H₂O at pH 12. The obtained gel was divided into two parts. One part was subjected to hydrothermal treatment (HT) at 150 °C, filtered, washed, and then dried to obtain Mn₃O₄ HT. The other part of the gel, represented as Mn₃O₄, was directly filtered and dried at 100 °C. The TEM images indicated the formation of cubic-, spheroid- and plate-shaped nanocatalysts (~33 nm) without any Au aggregates. Almost simultaneously, another research group reported identical cubic-, spheroid- and plate-shaped PdNPs/Mn₃O₄ nanostructures (38 nm) using an identical preparation method.¹³⁰ The presence of PdNPs could not be confirmed from the TEM images, but the XPS results indicated the presence of Pd⁰ and Pd²⁺ species on the Mn₃O₄ support.

3.1.2.3. Other metal oxides. In an early attempt, Glaspell *et al.* observed that the use of a capping agent, poly(ethylene glycol), PEG, or PVP in the preparation of Au and Pd NPs on CeO₂/CuO/ZnO support materials accelerated the nucleation process of metal NPs and produced a narrow size distribution.¹³¹ The respective support was derived from precursor salts (Ce(NO₃)₄, Cu(NO₃)₂ and Zn(NO₃)₂, respectively) and dopant Au or PdNPs were prepared *via* the addition of an



appropriate amount of metal salt (HAuCl_4 or $\text{Pd}(\text{NO}_3)_2$) to obtain the desired dopant concentration (2%, 5% and 10%). Comparatively, the 10% Pd/CeO_2 catalyst had the narrowest size distribution with average size 3–4 nm. Chuang *et al.* used $\text{CeO}_2\text{-TD}$, which was obtained *via* the thermal decomposition of Ce salt, to prepare $\text{Cu}/\text{CeO}_2\text{-TD}$ (Cu 5 wt%) by MW heating of an EG solution of $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ and PVP.¹³² Finally, the dried product was calcined at 475 °C for 4 h. For comparison, commercial ceria ($\text{CeO}_2\text{-C}$) and AC were also used as support materials. The size of the Cu particles was dependent on the nature and size of the support materials, where the minimum size (33 nm) was found for the $\text{Cu}/\text{CeO}_2\text{-TD}$ catalyst. Elazab *et al.* used different amounts (347, 329, 292, and 256 mg) of $\text{Cu}_2\text{H}_{10}\text{N}_4\text{O}_{17}$ precursor salt to prepare a Pd/CuO catalyst with variable Pd loadings (5, 10, 20, and 30 wt%).^{133,134} A simple one-step MW process was applied in the presence of N_2H_4 . The TEM images revealed the formation of a 20–40 nm-sized catalyst and X-ray photoelectron spectroscopy (XPS) confirmed the valence states as Cu^{2+} and Pd^0 . The estimated Pd content in the catalyst was much lower than the theoretical content calculated from its formula.

The different forms of Al_2O_3 (α , β , and γ) are perhaps the most important as catalyst supports due to their favorable combination of textural properties such as specific surface area, pore volume, pore size distribution, local microstructure, phase composition and acid/base characteristics.¹³⁵ Song *et al.* prepared an $\text{Ni}/\text{Al}_2\text{O}_3$ catalyst using PVP as a stabilizer and MW absorbent.¹³⁶ The catalysts were denoted as $\text{Ni}20\text{-Al}_2\text{O}_3\text{-M}$ and $\text{Ni}20\text{-Al}_2\text{O}_3\text{-I}$, where 20 represents the loading amount of Ni, and M and I stand for the MW and impregnation techniques, respectively. The MW method produced smaller (~ 10 nm) NiNPs in $\text{Ni}20\text{-Al}_2\text{O}_3\text{-M}$ compared to that (15 nm) in $\text{Ni}20\text{-Al}_2\text{O}_3\text{-I}$. The BET surface area and pore volume of $\text{Ni}20\text{-Al}_2\text{O}_3\text{-M}$ were also higher. This indirectly confirmed that the impregnation method (IM) produced slight agglomeration and blocking of the pore structure. In another investigation, $\text{Pd}/\text{Al}_2\text{O}_3$ catalysts were prepared *via* fixed-bed MW-induced plasma irradiation under three different conditions.¹³⁷ Fig. 3 illustrates the fixed-bed reactor with MW-induced plasma jet. The particles prepared under different conditions were named PF (fixed bed

reactor). The distance between the waveguide and particle bed influenced the distribution and size of Pd (Fig. 3). Spherical-shaped tiny Pd particles and the highest dispersion (25.3%) were observed in sample PF-1 when a shorter distance was applied (3 cm). It is obvious that a short distance produced a higher temperature. The plasma treatment reduced PdO to Pd in a short time, which had an fcc structure. The crystalline phase of $\gamma\text{-Al}_2\text{O}_3$ was detected in all the catalysts. The same authors reported the use of an MW-induced plasma-spouted bed reactor for the preparation of $\text{Pd}/\text{Al}_2\text{O}_3$ under various operating conditions.¹³⁸ The same catalyst was also prepared *via* CHT in an electric arc furnace. The MW-combined plasma treatment produced smaller PdNPs than that prepared by conventional heating at 500 °C. The rapid heat change induced by an increase in MW power reportedly produced surface cracks in $\text{Pd}/\text{Al}_2\text{O}_3$, whereas CHT at 500 °C did not produce surface cracks in the $\text{Pd}/\text{Al}_2\text{O}_3$ samples.

Wang *et al.* observed that MW hydrothermal (MWH) treatment is superior to conventional IM in terms of the better dispersion of WO_3 on an Al_2O_3 support and oxalic acid stabilizer had little effect on the dispersion.¹³⁹ The catalyst prepared by IM also had a higher agglomeration of WO_3 . After MWH treatment, the dried $\text{WO}_3/\text{Al}_2\text{O}_3$ was calcined at 550 °C for 4 h to produce $\text{W}/\text{Al}_2\text{O}_3$.

TiO_2 as a catalyst support is known to possess high thermal and chemical stability and offer outstanding resistance to corrosion in electrolytic media.¹⁴⁰ Howe *et al.* prepared a 1 wt% Ru/TiO_2 catalyst in an MW reactor from stock solutions of $\text{Ru}(\text{acac})_3$ or $\text{RuCl}_3 \cdot x\text{H}_2\text{O}$ in EG and TiO_2 at 150 °C, 170 °C and 200 °C for 5, 10 and 15 min.¹⁴¹ The obtained catalysts were air dried at 110 °C for 16 h. The catalysts were represented by the reductant-preparation temperature (°C)-preparation time, *e.g.* Acac-200-10 . Independent of the precursor salt, MW treatment for a longer time at 200 °C caused sintering of the RuNPs, given that their mean diameter increased slightly. However, despite the sintering effect, the average size of the RuNPs was still less than 3 nm. In either case, the catalysts exhibited a high surface carbon concentration.

To understand the detailed formation mechanism of various hybrids, Pt and AuNPs supported on CeO_2 , TiO_2 , ZnO , MgO , Al_2O_3 , and SiO_2 were prepared *via* the one-pot MW-assisted

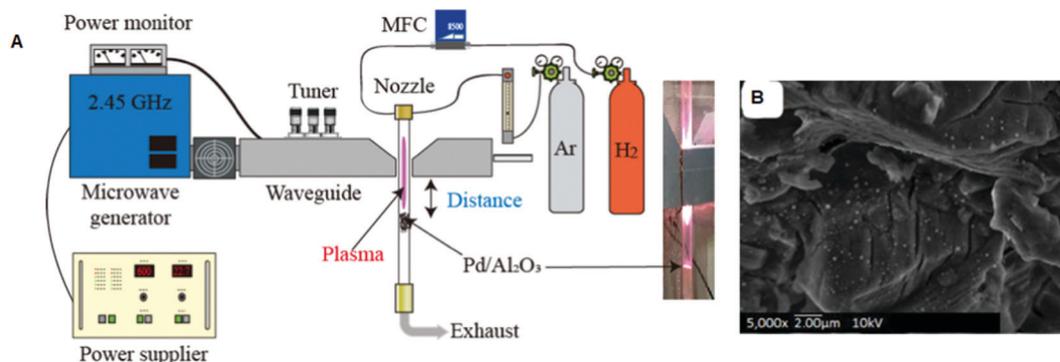


Fig. 3 (A) Schematic representation of the fixed bed reactor with plasma irradiation. (B) A typical SEM image of the $\text{Pd}/\text{Al}_2\text{O}_3$ catalyst (PF-1).¹³⁷ (Copyright 2019, MDPI.)



reduction of metal salts in EG.¹⁴² The size of the metal NPs was influenced by the efficiency of MW absorption by the support materials. Smaller and fine 2–3 nm-sized PtNPs were formed on CeO₂, TiO₂ and ZnO but not on the SiO₂ particles. Comparatively, the low dielectric loss substrate SiO₂ did not heat up rapidly. Likewise, on MgO and TiO₂, the uniform decoration of AuNPs was observed, whereas on Al₂O₃ and Si substrates, very few particles were present. This is because the Al₂O₃ and Si substrates did not absorb MW radiation.

3.1.3. Metal carbide-supported catalysts. Transition metal carbides possess excellent catalytic properties, and thus have great potential as high surface area support materials for metal catalysts.^{143,144} However, only a few works are available in this area. For example, Shi *et al.* reportedly used two types of mesoporous WC/C for loading 10 wt% Pt.¹⁴⁵ One was WC/C-F using the triblock copolymer Pluronic F127 as a soft template and the other was WC/C-S using mesoporous SBA-15 as a hard template. In both cases, the support material was dispersed in H₂PtCl₆ aqueous solution, mixed with EG, and then microwaved at 180 °C for 10 min. In both the Pt-WC/C-F and Pt-WC/C-S catalysts, Pt had an fcc crystalline structure, but the latter was more crystalline. The average pore size of WC/C-S was 4.7 nm, while that of WC/C-F was comparatively larger (8.0 nm). In both systems, the PtNPs were evenly distributed with little aggregation on the WC/C-S support. In a similar type of investigation, the stability of WC during the preparation of Pt@WC/organic mesoporous carbon abbreviated as Pt@WC/OMC-MP electrocatalyst *via* a pulse MW-assisted polyol method was investigated.¹⁴⁶ The loading of Pt in the Pt@WC/OMC-MP electrocatalyst was 20 wt%. MW irradiation in absence of the Pt precursor had little effect on the stability of WC/OMC; however, in the presence of the Pt precursor, the stability of WC was significantly affected. XPS confirmed the partial oxidation of WC to WO₃ during the preparation of the Pt@WC/OMC-MP catalyst. Asahara *et al.* studied the utility of Pd/SiC.¹⁴⁷ Here, [Pd(OAc)₂]_n dissolved in acetonitrile and small pieces of SiC taken in a screw-capped test tube were MW-heated at 100 °C for 3 h. The catalyst was represented as MW3. Catalyst MW1 was also prepared *via* the same method. For comparison, two catalysts, OB3 and OB10, were prepared *via* CHT for 3 and 10 h, respectively. However, information on the size distribution, surface porosity, and valence state of Pd was not available, only the catalyst activity was detailed.

3.2. Metal carbide catalysts

In terms of application performance, metal carbides such as W and Mo carbides are considered alternatives to noble metal catalysts, particularly Pt.¹⁴⁸ In 1973, Levy *et al.* showed that WC and Pt had similarities in electronic structure and catalysis behavior in water formation from H₂ and O₂ at room temperature.¹⁴⁹ Following that groundbreaking work, these carbides were proven to be useful in a wide variety of catalytic reactions such as hydrogenation, hydrodeoxygenation, hydrodesulfurization, hydrodenitrogenation and isomerization.^{150–153} In this perspective, a few works carried out on the preparation and applications of supported metal carbides are discussed below.

3.2.1. Carbon-supported catalysts. Despite the fact that the catalytic properties of Mo₂C resemble that of noble metals, the quick deactivation of bulk Mo₂C is a problem.¹⁵⁴ Accordingly, an MW-assisted single-source route was investigated for the preparation of highly dispersed Mo₂C/AC using β-Mo₂C as the precursor.¹⁵⁵ XPS showed the existence of Mo(vi) species in Mo₂C/AC, possibly resulting from surface oxidation in air. The TEM images confirmed the formation of highly dispersed 3 nm-sized Mo₂C NPs. Mnatsakanyan *et al.* used two different MW-assisted routes to prepare Mo₂C/C and WC/C catalysts.¹⁵⁶ The complete formation of Mo₂C occurred after 3 min irradiation, whereas for the WC system, an irradiation time of 8 min was required. The initial size of the carbon from the SEM image was ~50 nm, and after irradiation the sizes of the Mo₂C/C and WC/C catalysts increased to 50–70 nm and 300–500 nm, respectively.

In 2009, Liang *et al.* described the preparation of WC/CNT *via* MW-assisted metal-organic chemical vapor decomposition (MOCVD) under atmospheric pressure in a fluidized reactor.¹⁵⁷ A precursor mixture made from W(CO)₆ and CNT was placed in a quartz-tube reactor and fluidized with an argon flow for 2 h before irradiation in an MW oven. CNTs without pre-oxidized and pre-oxidized were evaluated as a support. A uniform distribution of 2–5 nm-sized WC was found on the surface of the CNTs. Overall, the results indicated that preoxidation of CNTs is not essential in the MW-assisted MOCVD method. In the following year, Pang *et al.* described the preparation of Mo₂C/CNTs utilizing the above-mentioned method.¹⁵⁸ However, the CNTs were not pre-oxidized before use. In 16.7 wt% Mo₂C/CNT, well-dispersed 5 nm-sized Mo₂C was detected. The average crystallite size of Mo₂C increased with an increase in the Mo loading. Pt-Mo₂C/CNT was also prepared and 3–6 nm monodispersed PtNPs were formed.

3.3. Metal oxide catalysts

3.3.1. Carbon-supported catalysts. The use of carbon materials as support materials for metal oxide catalysts has scarcely been studied to date. Metal oxide catalysts gained prominence in the mid-1950s, given that they were found effective in catalyzing oxidation and acid-base reactions. Metal oxides have now found applications in the petrochemical industry, in synthesizing chemicals and improving environmental issues.¹⁵⁹

Zhong *et al.* demonstrated the suitability of MW-assisted 3D heating in air for the rapid *in situ* synthesis of metal oxides (CoO_x, NiO and Fe₃O₄) using carbonized wood (C-wood) as a 3 D heating substrate.⁷⁵ In a typical experiment, the Co(NO₃)₂ precursor was dissolved in a 1:1 ethanol-water mixture, and then a piece of C-wood was immersed in this mixture for 10 min and dried for 20 min to obtain the C-wood/Co(NO₃)₂ precursor. The C-wood/Co(NO₃)₂ precursor was then microwaved at 1000 W for 4 s to produce C-wood/CoO_x. Around 11 nm-sized uniformly distributed CoO_x NPs were formed on the C-wood substrate. The potential scalability of the 3D heating method was also verified with a larger-sized C-wood substrate.

Motshekga prepared MWCNT/SnO₂ NPs *via* the conventional and MW-assisted wet impregnation for application in optoelectronic devices.¹⁶⁰ The SEM and TEM images confirmed



the high loading of SnO₂ NPs (2–4 nm) on MWCNT *via* MW method within 10 min. There is also a report on the preparation of TiO₂/G from GrpO, titanium isopropoxide, TTIP (variable) in isopropyl alcohol and N₂H₄ *via* MW irradiation in a closed Pyrex glass bottle.¹⁶¹ The obtained dried precipitate was finally calcined at 400 °C for 4 h.

3.3.2. Silica-supported catalysts. Mohamed *et al.* studied the effect of MW irradiation time (min) during preparation of CeO₂-SiO₂ NPs from an acidic solution of CTAB, TEOS and Ce(NO₃)₃·6H₂O in ethanol.¹⁶² Finally, the washed and dried product was heat treated at 500 °C for 4 h. Irrespective of the irradiation time, all the MW-irradiated samples contained a pure CeO₂ phase. An increase in the irradiation time resulted in an increase in the crystallinity and average crystallite size, but a decrease in the BET surface area. The TEM image analysis for sample M-30 (irradiation time 30 min) indicated the formation of cubic-shaped particles with an average size of 8 nm, whereas conventional HT for 24 h (H-24 h) produced comparatively larger spherical (15 nm) particles.

Li *et al.* prepared an NiO/vermiculite or NiO/Vmt catalyst with 10 wt% loading from a dried precursor comprising Vmt in Ni(NO₃)₂·6H₂O aqueous solution.¹⁶³ The precursor was divided into two parts, where one part was subjected to CHT and the other part was subjected to MW treatment to obtain CHT-NiO/Vmt and MW-NiO/Vmt, respectively. The NiO NPs were well dispersed, but MW irradiation produced smaller particles (3 nm) compared to that from CHT (6 nm). The estimated crystallite size in the MW-NiO/Vmt catalyst was 2.4 nm, lower than that in the CHT-NiO/Vmt catalyst (2.7 nm). In a similar investigation, 2-D layered Vmt was used to prepare MW-assisted MW-NiO/2D-Vmt and compared with muffle furnace (MF) calcination.¹⁶⁴ The overall size distribution and crystallinity again confirmed that the MW method is superior for producing smaller-grained highly crystalline NiO.

3.3.2. Metal-supported catalysts. A Co₃O₄@stainless steel-mesh (SUS) micro-rod array electrocatalyst was prepared with the aid of diethylenetriamine (DETA) as an anchoring agent for Co²⁺ (Fig. 4).¹⁶⁵ For comparison, Co₃O₄ layered nanosheets

(Co₃O₄-LNS) were prepared *via* the same method without the SUS substrate. MW heating allowed recrystallization of the rod-shaped Co(OH)₂ in the Co₃O₄@SUS electrocatalyst, while in the absence of the SUS substrate, a flower-like Co(OH)₂ nanomaterial was formed. In either case, the annealing at 400 °C resulted in the complete conversion of Co(OH)₂ to Co₃O₄. The average crystallite size of Co₃O₄ in Co₃O₄@SUS and Co₃O₄-LNS was 33.41 and 29.33 nm, respectively. Fig. 4 confirms the different appearances of Co₃O₄ in the presence and absence of the SUS substrate.

May-Masnou discussed the preparation of anatase TiO₂ NPs attached to larger anisotropic Au *via* an MW-assisted method.¹⁶⁶ Firstly, TiO₂ NPs were synthesized from a homogeneous solution of titanium butoxide (TBOT) and PVP in benzyl alcohol (BA) in an MW reactor at 50 °C, and then again heated at 190 °C for 10 min. In the second step, two different sized AuNPs (Fig. 5) were prepared in the presence of TiO₂ NPs dispersed in EG using (a) 100 and (b) 600 mg of PVP. Part of the sample was dried overnight and the other part was calcined in air at 400 °C, 450 °C, 550 °C, and 600 °C for 2 h to remove PVP and increase the contact between Au and TiO₂. The PVP-capped TiO₂ NPs possessed a pure crystalline anatase phase with an average size of 9 ± 2 nm. The TEM photographs (Fig. 5) indicated that 100 mg of PVP produced >50 nm-sized Au NPs, whereas 600 mg of PVP produced smaller (8 ± 2 nm) AuNPs. The larger AuNPs were randomly decorated with smaller TiO₂ NPs but the smaller AuNPs were only surrounded by TiO₂. Calcination had a larger effect on the shape of Au than that of the TiO₂ NPs.

3.4. Binary metal catalysts

Binary and ternary metal catalysts, which are considered more active compared to simple metal catalysts, are particularly useful as electrocatalysts in proton exchange membrane fuel cells (PEMFCs) and direct alcohol fuel cells.^{167,168} In early investigations, electrocatalysts based on precious metals such as Pt or Au were mostly investigated. However, the low durability and high cost Pt and Au metal catalysts hinder their

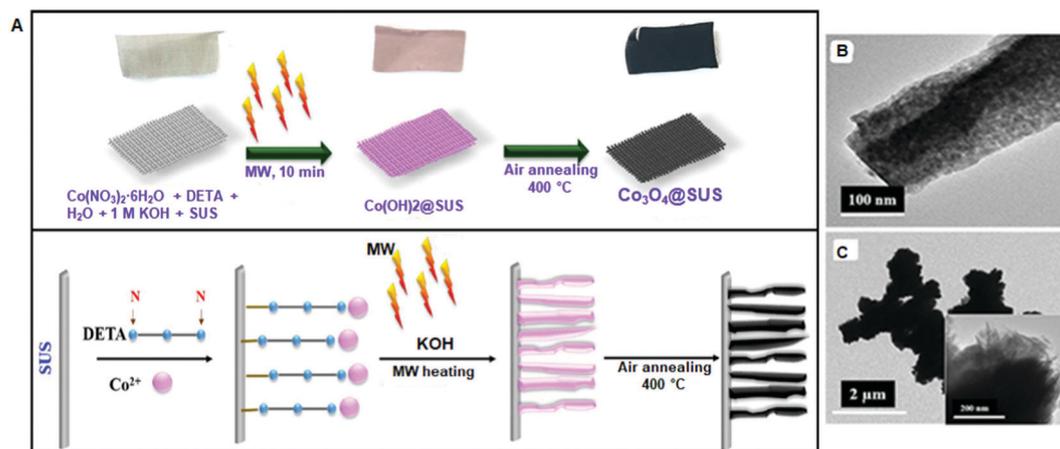


Fig. 4 (A) Proposed mechanism for the controlled synthesis of Co₃O₄@SUS. TEM images of (B) Co₃O₄@SUS and (C) Co₃O₄-LNS.¹⁶⁵ (Copyright 2017, the American Chemical Society.)



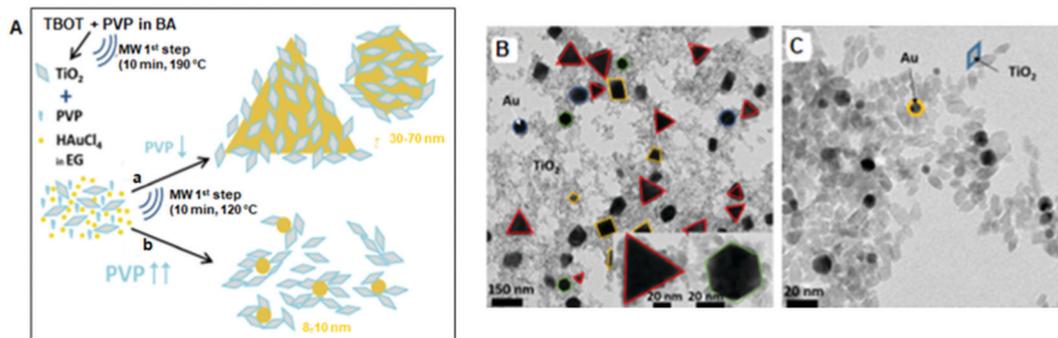


Fig. 5 (A) Schematic diagram of the two-step MW-assisted synthesis of Au/TiO₂ nanostructures. (a) TiO₂:PVP:H AuCl₄ = 1:0.097:0.155 and (b) TiO₂:PVP:H AuCl₄ = 1:0.583:0.155 (molar ratios). TEM images of the as-synthesized (B) sample 'a' and (C) sample 'b'.¹⁶⁶

commercial application. Additionally, supported Pt-based catalysts exhibit sluggish oxygen reduction reaction (ORR) activity at the cathode. Accordingly, doping with transition metals containing Pt or Au reportedly improved their durability.^{169,170} The search for alternative low-cost materials with high electrocatalytic activity and good durability has led to the development of ternary metal catalysts either by replacing or reducing the Pt or Au content.

3.4.1. Carbon-supported catalysts. Stevanović *et al.* used a carbon support for the synthesis of Pt, PtRh and PtSn catalysts with 20 wt% metal loading.¹⁷¹ Initially, the bimetallic NPs first prepared *via* polyol MW-treatment of H₂PtCl₆ and SnCl₂/RhCl₃ in EG. The supported catalysts were prepared by blending bimetallic NPs with a carbon support, acid treatment, and finally drying in an N₂ atmosphere. The alloy structure of the PtSn/C and PtRh/C catalysts was confirmed, and the TEM images showed the uniform distribution of ~3 nm-sized NPs on the carbon support for all the catalysts. The valence state of the metals was not identified. In 2017, Tsuji *et al.* tried to prepare PtY/C alloy NPs *via* MW co-reduction of a precursor mixture of *cis*-[Pt(NH₃)₂(NO₂)₂] or Pt(C₅H₇O₂)₂ and Y(CH₃COO)₃·4H₂O in the presence of a carbon support in EG.¹⁷² Irrespective of the Pt precursor, the composition revealed the formation of mono-dispersed Pt-YO_x/C catalysts. The obtained catalysts were denoted as Pt-YO_x/C(A):5:5 and Pt-YO_x/C(B):5:5, where A and B represent the *cis*-[Pt(NH₃)₂(NO₂)₂] and Pt(C₅H₇O₂)₂ precursor, respectively. To overcome the solubilization of the YO_x component in acid fuel solution, the samples were treated with HNO₃ to produce PtY/C alloy catalysts. After acid treatment, both samples contained a small amount of Y component and the catalyst size with a narrow distribution was retained. The addition of Y(CH₃COO)₃·4H₂O was found to be beneficial for producing monodispersed Pt/C(A, B) particles. To enhance the binding between the PtNPs and carbon supports, a Pt₃Cu/C catalyst was prepared with a theoretical Pt loading of 20 wt%.¹⁷³ A mixture of metallic precursors (H₂PtCl₆·6H₂O and CuCl₂·2H₂O) and carbon support in an EG-water mixture was microwaved at pH 10 for approximately 6 min at 125 °C. The Pt₃Cu NPs (2.31 nm) were uniformly distributed on the support. Pt and Cu had an fcc crystalline alloy structure. The metallic state of Pt and Cu was also confirmed. By just changing the amount of Cu precursor in the recipe, Pt₄Cu/C (2.14 ± 0.88 nm) and Pt₂Cu/C (3.49 ± 1.097 nm) were also prepared

and characterized. In the synthesis of an Pd-Au/C catalyst, relative to CHT, catalyst activation by MW irradiation produced a much higher degree of Pd-Au alloying and better homogeneity of the bimetallic phase.¹⁷⁴ The use of acetone as a solvent also improved the homogeneity of the Pd-Au phase compared to the aqueous solution. The catalyst samples produced by MW irradiation also contained an increase Pd content at the surface of the NPs. The poor ORR activity of 20% CoSe₂/C encouraged the preparation of the corresponding CoSe₂/C.^{175–177} Accordingly, a homogeneous precursor solution of Co(CH₃COO)₂ and SeO₂ was placed in an MW oven at a power of 800 W for 180 s, and the product was washed and dried before mixing with carbon black. Diffraction signals of the CoSe₂ phase were not observed up to an Se/Co ratio of 4.0 due to their uniform distribution. In another investigation, FeSe₂/C NPs were prepared *via* an MW method under identical conditions with the FeSe₂ loading adjusted to 36%.¹⁷⁸ The average particle diameter was in the range of 3–12 nm.

Sulfonated MWCNT (SF-MWCNT) was used to prepare SF-MWCNT-PdSn and SF-MWCNT-PdNi *via* MW-assisted solvothermal co-reduction (1000 W, 170 °C, 60 s) of the precursor salts in EG.¹⁷⁹ Pd had a well-defined fcc crystalline structure, but Ni and Sn possessed a partially amorphous structure. The alloy structure was confirmed and the XPS spectra suggested the presence of Pd in the metallic and oxide forms with Ni in the oxo-nickel species (*i.e.*, Ni_xO_y and NiOOH). However, the uniform distribution of the bimetallic catalysts was not analyzed. Nassr *et al.* used oxygen-functionalized CNT (FCNT) to prepare a PtNi/FCNT catalyst *via* the polyol MW reduction of metal salts, H₂PtCl₆ and Ni(NO₃)₃·6H₂O, either continuously for a defined time or in pulse mode with a duration of 20 s.¹⁸⁰ The transition of the NiO peaks to PtNi alloy occurred at 90 s. However, in pulse mode, the formation of the PtNi alloy was observed even at a lower pulse number of 6. An increase in the irradiation time enhanced the particle size of PtNi/CNT from 2.7 to 3.3 nm. Comparatively, the size of PtNi/CNT prepared in the pulse mode was almost independent of the pulse number given that the average size was around 2.5 nm. Rahsepar *et al.* prepared a PtRu/MWCNT catalyst using MW-assisted reduction at pH ~12.5 from a water solution of metal precursors, pretreated MWCNT and HCHO reducing agent.¹⁸¹ After re-acidification and drying, additional heating at 300 °C under a flow of H₂ helped to form well-dispersed



PtRu alloy NPs (1.5–3.5 nm) on the surface of MWCNT. The XPS signals indicated the presence of a large amount of Pt⁰ (72.15%) and Ru⁰ (68.07%) with a small amount of Pt(II), Pt(IV), RuO₂ and RuO₂·xH₂O species. Al-Tememy had MWCNT-doped G nanoplatelets (MWCNT-GNP) to support a PtPd bimetallic catalyst by microwaving a mixture of metal precursors and the support dispersed in an EG-isopropanol mixture at pH 12.¹⁸² For comparison, PtPd/GNP and PtPd/MWCNT catalysts were also investigated. The doping of G nanoplatelets by MWCNT improved the metal loading (35.1%), dispersity and crystallinity index of the PtPd NPs. Independent of the support material, all the catalysts contained graphitic carbon and fcc-structured metals. The average sizes were in the range of 2.47–2.70 nm.

3.4.2. Oxide-supported catalysts

3.4.2.1. Silica. Verma *et al.* discussed the preparation of Pd-Ag/SiO₂ and Pd-Au/SiO₂ bimetallic catalysts in two steps.¹⁸³ Firstly, 1 wt% of either Au or Ag was incorporated on SBA-15 *via* the MW-assisted polyol method, and then decorated with PdNPs *via* a localized surface plasmon resonance-assisted deposition method under visible light irradiation. Spherical NPs of Pd-Ag and Pd-Au with diameters of 4.2 and 4.9 nm were formed. The formation of the alloy *via* the electron transfer mechanism was proved by XPS. Tripathi *et al.* demonstrated the preparation of Cu-Ag/MgO-SiO₂ *via* four different techniques including conventional impregnation, MW-assisted method, PVA/NaBH₄-assisted method and benzoxazine-assisted method.¹⁸⁴ In the MW method, drying was carried out at 80 °C in an MW oven before calcination at 400 °C for 4 h. MW drying and benzoxazine-assisted reduction efficiently produced a higher surface area and pore volume. Partial oxidation of Cu to CuO was observed in Cu-Ag/MgO-SiO₂. However, other characterizations were not presented. In another investigation, Ti-inserted ordered mesoporous SiO₂ (Ti-SBA-15) was used to support a CoMo catalyst.¹⁸⁵ 20Ti-SBA-15 (Si/Ti molar ratio 20) was prepared *via* hydrolysis and aging in an MW reactor from TEOS and TTIP in the presence of

Pluronic P123 as a structure-directing agent. The crystallized product was further calcined at 450 °C for 6 h. As expected, the insertion of Ti reduced the surface area and pore size of SBA-15. The Ti loading in 20Ti-SBA-15 and CoMo/20Ti-SBA15 was 2.9 and 2.4 wt%, respectively, which was close to the theoretical value.

3.4.2.2. Metal oxides. In an early investigation, Ru-Sn/ZnO catalysts were synthesized by subjecting a mixture of RuCl₃·3H₂O dissolved in EG containing SnCl₂·2H₂O and ZnO (targeted metal loading 2%) to variable MW power.¹⁸⁶ XPS analysis showed the formation of Sn⁰ in all the Ru-Sn/ZnO catalysts prepared at 450 W for 4 and 8 min and Ru⁰ was detected only when the MW power was set at 150 W for 20 min. The rest of the samples mainly contained Ru^{III}. The morphological data for the Ru-Sn/ZnO catalysts was not available.

3.4.3. Metal-organic framework-supported catalysts. Tan *et al.* used zeolite-imidazole frameworks (ZIFs) to construct dispersed Co-N-C catalysts, and then prepared an Ni₃Fe/Co-N-C catalyst *via* a rapid MW method (Fig. 6).¹⁸⁷ The fcc structure of Ni₃Fe together with a broad diffraction plane of carbon was detected. The Ni, Fe and Co contents (ICP analyses) in Ni₃Fe/Co-N-C were 20.04%, 6.53% and 1.26%, respectively. After decoration with Ni₃Fe, the size of the respective catalysts did not change (Fig. 6). The bimetal particle size was around 11.2 nm.

3.5. Mixed metal oxide catalysts

3.5.1. Carbon-supported catalysts. Elazab *et al.* prepared three different catalysts (Pd-Fe₃O₄/G, Pd-Co₃O₄/G and Pd-Ni(OH)₂/G) by irradiating a mixture of GO nanosheets, precursor salts and N₂H₄ for 60 s.¹⁸⁸ In the Pd-Fe₃O₄/G and Pd-Co₃O₄/G catalysts, well dispersed 35 ± 2 nm and 25 ± 2 nm PdNPs were formed, while the sizes of Fe₃O₄ and Co₃O₄ were (52 ± 2 nm) and (45 ± 2 nm), respectively. The binding energies confirmed the valence states as Pd⁰ and Pd²⁺. The magnetic property of Pd-Fe₃O₄/G was also confirmed. The same group

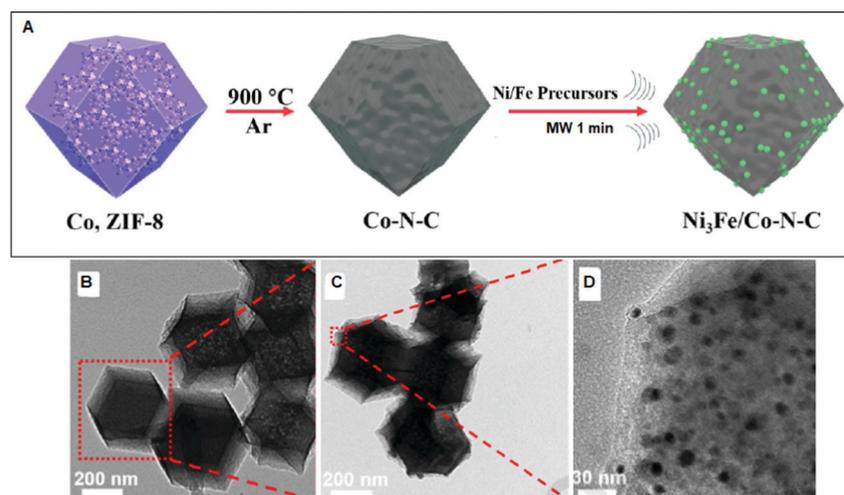


Fig. 6 (A) Preparation scheme of the Ni₃Fe/Co-N-C bifunctional electrocatalyst. TEM (B and C) and HRTEM (D) images of (B) Co-N-C and (C and D) Ni₃Fe/Co-N-C.¹⁸⁷ (Copyright 2020, Elsevier B.V.)



also prepared four magnetically recyclable Pd-Fe₃O₄/G catalysts labeled as 1–4 with the Pd loading varying from 50 wt% in Catalyst 1 to 6 wt% in Catalyst 4.¹⁸⁹ Catalyst 3 contained the highest Pd(0)/Pd(II) ratio. Khalid *et al.* prepared Ag-TiO₂/G composites *via* an MW-assisted hydrothermal method from a GO dispersion in an ethanol-water (1:1) mixture containing anatase TiO₂ and varying amounts of 0.01 mol L⁻¹ AgNO₃.¹⁹⁰ The weight ratio of Ag to the TiO₂-G composite was controlled at 0, 0.03, 0.06, 0.09, 0.12, and 0.15 and composites were named TG, ATG3, ATG6, ATG9, ATG12, and ATG15, respectively. A crystalline anatase TiO₂ phase was found, but G and Ag could not be detected in Ag-TiO₂/G owing to their low content. The valence states were confirmed to be Ti⁴⁺ and Ag⁰ in the composites. Pan *et al.* used N-doped coal-derived carbon to support Co/CoO_x.¹⁹¹ Depending on the heat treatment temperature, the coal-derived carbons were labeled as MC800, MC950, MC1050 and MC1100. The high-temperature treatment of the carbon (MC1050) was found to be favorable for producing structural disorder and subsequent introduction of Co/CoO_x. The Co nucleation was followed by strong electrostatic interaction between the oxygen functional groups on the surface and Co cations. The Co/CoO_x particles were homogeneously distributed at a lower Co content. Galal *et al.* discussed the preparation of Ru-based perovskites supported on RGO.¹⁹² The preparation was carried out *via* the redox reaction of metal salts (Sr(NO₃)₂/Ba(NO₃)₂/Ca(NO₃)₂), RuCl₃ and GO *via* either conventional or MW heating. Both conventional and MW heating allowed the formation of SrRuO₃ in combination with RuO₂ or RuNPs. Highly crystalline nanosized CaRuO₃ was detected as the main phase using MW heating. In comparison, around 1 nm-sized BaRuO₃ particles with RuO₂ were formed *via* conventional heating, while MW heating produced amorphous BaRuO₃.

Li *et al.* used carboxymethyl cellulose (CMC), an inexpensive biobased polymer, to support Ag-Fe₃O₄.¹⁹³ Firstly, Ag@CMC was prepared *via* the MW heating of Ag(NH₃)₂⁺ salt in the presence of aqueous CMC sodium salt.^{194,195} Secondly, Fe₃O₄ was co-precipitated under MW irradiation and alkaline conditions to afford Ag-Fe₃O₄@CMC (Fig. 7). Monodispersed AgNPs (2.3 ± 0.3 nm) were formed in Ag@CMC, but the magnetic Fe₃O₄ NPs (15.3 ± 3.5 nm) were slightly polydispersed (Fig. 7).

3.5.2. Oxide-supported catalysts

3.5.2.1. Metal oxide. Jing *et al.* prepared a CuNiZnO/γ-Al₂O₃ catalyst *via* the MW-assisted thermal hydrolysis of urea.¹⁹⁶ The required amounts of Ni(NO₃)₂·6H₂O and γ-Al₂O₃ powder were taken in Cu(NO₃)₂·3H₂O and Zn(NO₃)₂·6H₂O aqueous solution, sonicated, and finally heated in an MW oven at 80 °C, 85 °C, 90 °C or 95 °C for 2 h. The recovered dried product was calcined at 350 °C for 5 h. The composition analysis of the four catalysts indicated that the Cu content gradually decreased with an increase in the MW temperature, whereas both the Ni and Zn contents increased. The surface area and pore volume of the γ-Al₂O₃ support decreased as usual following modification. A higher MW temperature produced uniform smaller particles with better dispersion.

A Ce_{0.8}Zr_{0.2}O₂ solid solution was synthesized *via* a rather long process of citrate complexation (17 h) and supported on different loadings of Al₂O₃ *via* the impregnation (CI) or MW combustion technique (CC), and finally calcined at 800 °C for 5 h.¹⁹⁷ The CZ loading was found to be optimum at 40 wt%, and a greater CZ loading produced agglomeration. Accordingly, samples CZ40A-CC and CZ40A-CI contained relatively finer particles.

3.6. Ternary metal catalysts

3.6.1. Carbon-supported catalysts. Zheng *et al.* prepared a series of ternary (Co_xNi_{1-x})Se₂/C (0 < x < 1.0) catalysts *via* a simple MW polyol method to replace the poor-performing

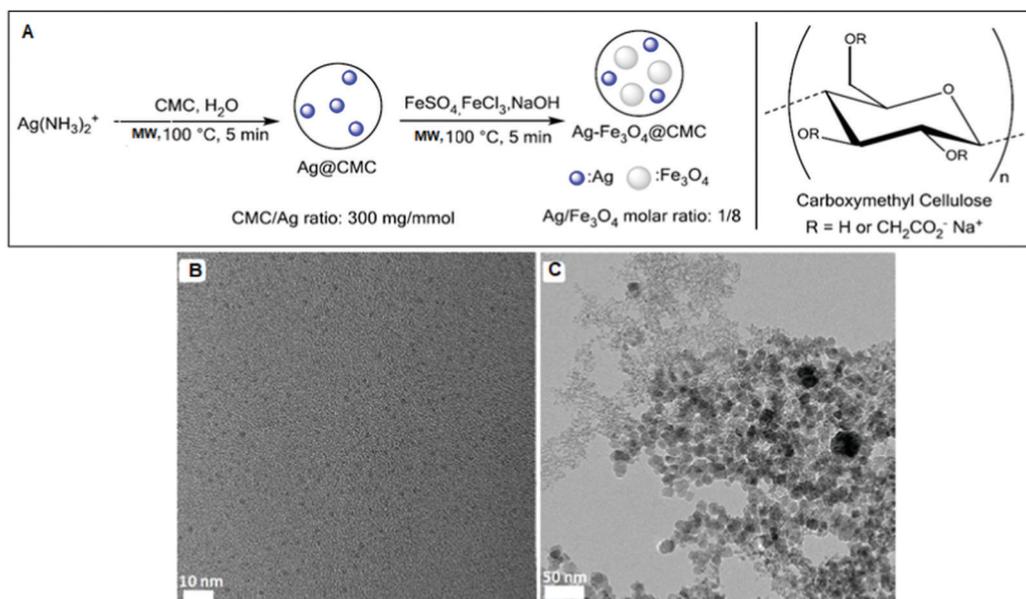


Fig. 7 (A) Preparation scheme of Ag@CMC and Ag-Fe₃O₄@CMC catalysts. TEM images of (B) Ag@CMC and (C) Ag-Fe₃O₄@CMC catalysts.¹⁹³ (Copyright 2016, the American Chemical Society.)



binary NiSe₂/C electrocatalyst.¹⁹⁸ In the synthesis, Co(CH₃COO)₂, Ni(CH₃COO)₂ and SeO₂ were used as the starting materials. The ternary catalyst solid solution was comprised mainly of cubic NiSe₂ and CoSe₂. An increase in the Co content reduced the average crystallite size of (Co_xNi_{1-x})Se₂/C from 32.1 nm to 20.6 nm. The TEM image confirmed the uniform distribution of NPs (<2 nm) on the carbon support. In another study, ternary ultra-small composites named Pt_{4.5}Sn_{1.5}Rh₁ USNP/C, Pt₃Sn₁Rh₁ USNP/C and Pt_{4.5}Sn₁Rh_{1.5} USNP/C containing 20% Pt loading were prepared *via* a rapid and simple one-step MW polyol method (Fig. 8A).¹⁹⁹ The TEM images of Pt_{4.5}Sn_{1.5}Rh₁ USNP/C, Pt₃Sn₁Rh₁ USNP/C and Pt_{4.5}Sn₁Rh_{1.5} USNP/C showed the uniform distribution of NPs on the carbon support with average diameters of 1.57 ± 0.97, 1.48 ± 1.18 and 1.75 ± 1.14 nm (Fig. 8B–D), respectively. XPS analysis of Pt_{4.5}Sn_{1.5}Rh₁ USNP/C revealed that Pt and Rh were in the metallic state, while Sn was in the oxidized state.

4. Catalysis by supported simple metal catalysts

The previous sections revealed that a variety of supported nanocatalysts has been prepared *via* MW-assisted methods. The supported nanocatalysts include simple metal catalysts such as Cu,^{81–83,105,132} Ni,^{105,118,136} Co,^{105,119,120} Fe,⁸⁴ Ru,^{85,86,95–99,141} Ag,^{112,116} Au,^{112,129,131,142} Pt,^{87–92,106,111,121,142,145,146} Ti,¹²² Zr,¹²³ Pd,^{93,106–108,127,128,130,133,134,137,138,147} and W.¹³⁹ Several attempts have also been made to prepare supported simple metal oxide (CoO_x, Co₃O₄, NiO, Fe₃O₄, SnO₂, CeO₂, and TiO₂) nanocatalysts for application in the petrochemical industry.^{75,160–166} To improve the catalytic activity and versatility of the application potential of simple metal or metal oxide nanocatalysts, investigations on supported bimetal,^{171–175,177–187} mixed metal oxide^{188–193,196,197} and ternary metal^{198,199} nanocatalysts have also been performed. MW-assisted synthesis has been claimed

to be an economically viable, short, green method to produce supported nanocatalysts. However, to achieve high catalytic efficiency, strong bonding of the nanocatalyst to the support materials without damaging the support materials due to MW irradiation is essential. Moreover, the loading amount, nanocatalyst size and distribution, and reduction efficiency of the precursor salt during the preparation of catalysts are also known to contribute to their catalytic activity. Consequently, the following section covers the usefulness of various MW-synthesized supported nanomaterials in catalysis with emphasis on their catalytic performance, stability and recycling efficiency. The performance is compared with that prepared *via* conventional methods, if available. For better comparison, this section is divided into subsections under different reaction types such as oxidation, reduction/reductive degradation, hydrodesulfurization/hydrodearomatization, photocatalytic degradation, hydrogenation, cross couplings, synthetic reaction (cycloisomerization, dehydrogenation, transesterification, and photocatalytic hydrogen generation) and electrocatalysis. Some of these reactions are illustrated in Scheme S1 (ESI[†]). It is worth mentioning that only the articles which published after 2012 are considered in the following section.

4.1. Oxidation reactions

Chuang *et al.* observed that smaller (16 nm) CuNPs and a cubic morphology in Cu/AC-700 W-60 s-12 prepared under the optimized MW conditions were favorable for obtaining the highest oxidation of CO.⁸¹ The same observation was made with the Cu/CeO₂-TD catalyst during CO oxidation (95.8% conversion) due to the better dispersion and smaller CuNPs (33 nm).¹³² The catalytic efficiency of Cu/CeO₂-TD was comparable to that of Cu/CeO₂ nanorods prepared *via* the wet IM and hydrothermal method.^{200,201} A comparative study on Pd/Co₃O₄, Pd/Fe₃O₄ and Pd/Ni(OH)₂ catalysts for the CO oxidation in a programmable flow tube furnace reactor indicated that both 30 wt% Pd/Co₃O₄ and 50 wt% Pd/Fe₃O₄ catalysts can produce 100% CO oxidation at 127 °C given that the Co₃O₄ support

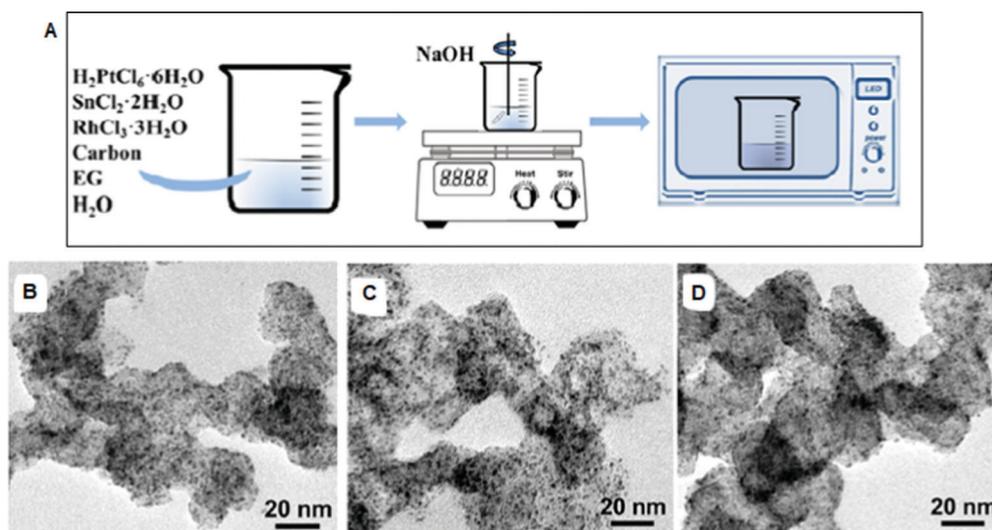


Fig. 8 (A) Scheme of the preparation of ternary Pt–Sn–Rh USNP/C composites. TEM images of (B) Pt_{4.5}Sn_{1.5}Rh₁ USNP/C, (C) Pt₃Sn₁Rh₁ USNP/C and (D) Pt_{4.5}Sn₁Rh_{1.5} USNP/C composites.¹⁹⁹ (Copyright 2020, Elsevier B.V.)



itself has significant conversion efficiency.¹²⁷ In a related report, the Pd/Fe₂O₃-573 catalyst (Pd 1.87 wt%) prepared *via* coprecipitation followed by calcination at 573 K produced 100% CO oxidation efficiency at only 60 °C.²⁰² This was due to the smaller (~ 10 nm) crystalline size of the Fe₂O₃ support compared to Fe₃O₄ (25–30 nm). The deposition of AuNPs on the Mn₃O₄ support improved the catalytic oxidation of CO and the 4-Au/Mn₃O₄ catalyst (4 wt% Au) resulted in 100% conversion at 98 °C.¹²⁹ The stability test of 4-Au/Mn₃O₄ at 50 °C and 125 °C exhibited its excellent stability up to 20 h, and then it started to deactivate slowly, losing 4% of its initial activity after 30 h. Likewise, the best catalytic activity was obtained over the 4-Pd/Mn₃O₄ (containing 4 wt% Pd) catalyst given that 100% CO oxidation occurred at only 20 °C, a much lower temperature than that of the previous 4-Au/Mn₃O₄ catalyst.¹³⁰ This activity was almost on par with the meso-structured 2.7 wt% Pd-loaded Pd/Mn₃O₄ catalyst prepared *via* impregnation and reduction.²⁰³ The stability of 4-Pd/Mn₃O₄ was also better and found to be stable up to 30 h without any remarkable decrease in its activity.

Both the loading method (*e.g.*, complexation-MW assisted combustion, CC, and complexation-impregnation, CI) and weight percentage of Ce–Zr solid solution in the Ce_{0.8}Zr_{0.2}O₂/Al₂O₃ catalyst influenced the oxidation of CO.¹⁹⁷ Obviously, the conversion degree increased with the loading of Ce_{0.8}Zr_{0.2}O₂ and the MW-assisted (CC) method resulted in better oxidation activity. CZ40A-CC converted 98% and 100% CO at 250 °C and 275 °C, respectively, but CZ40A-CI converted 100% CO at only the higher temperature (275 °C). The maximum conversion achieved by the CZ40A catalysts was due to the proper amount of Al₂O₃, reasonable surface area and higher dispersion of ceria and zirconia. However, the stability of the catalysts was not studied.

Ren *et al.* evaluated the performance of a hollow Cu/AC nanocatalyst in the oxidative carbonylation of methanol to dimethyl carbonate (DMC).⁸³ Compared to solid Cu/AC-4, the hollow structure of CuNPs with exposed Cu(111) crystals improved the selectivity.^{204,205} An irradiation temperature of 360 °C for a longer duration during the preparation of catalyst was found to be undesirable given that the catalytic efficiency was reduced due to the change in initial morphology.²⁰⁶

An MW-NiO/2D-Vmt catalyst exhibited better activity than an MF-NiO/2D-Vmt catalyst for acetylene carbonylation in a constant volume sealed container for the synthesis of acrylic acid (AA).¹⁶⁴ Again, smaller NiO grains in the MW-NiO/2D-Vmt catalyst produced a higher yield of AA (86%), whereas the MF-NiO/2D-Vmt catalyst produced 68% yield. However, the MW-NiO/2D-Vmt catalyst was unstable, demonstrating a drastic change in morphology and disappearance of the NiO signal after the first cycle.

An Fe/AC catalyst removed 93% of added phenol (concentration unknown) after 180 min at 30 °C during oxidative degradation but its stability was not that good given that the removal efficiency dropped to 20% after the fourth recycle due to severe leaching of its Fe ions.⁸⁴ However, the Fe/AC catalyst (4 wt% Fe), prepared by heat treatment in air, was worse in terms of performance and stability.²⁰⁷

4.2. Reduction/reductive degradation reactions

Independent of the Ru content, both Ru/CPM-1 (0.5 wt%) and Ru/CPM-2 (1.0 wt%) catalysts completely reduced *p*-nitroaniline (*p*-NA) and crystal violet (CV) using NaBH₄ reductant within 15 min and 30 s, respectively.⁸⁵ However, a smaller size of embedded RuNPs in the PDC@RuNP catalyst and porous structure helped to attain 98% elimination of the yellow color of K₃Fe(CN)₆ within 30 s.⁸⁶ Comparatively, commercial Ru/C took a longer time (>30 s) due to its insufficient porous structure. The PDC@RuNPs catalyst was fairly stable, retaining ~85% catalytic activity after six consecutive cycles. The specific surface area decreased slightly after the 6th run but no leaching of Ru was detected. The complete degradation of new fuchsin (NF) dye was also possible over the PDC@RuNP catalyst.

The smaller (2.0 ± 0.5 nm) size and dense distribution of RuNPs in the SWCNT-Ru catalyst produced through MW reduction at 100 °C for 15 min degraded Congo red (CR) dye at both low concentration (0.01 mM, 63.8%, 2.1 min) and high concentration (0.06 mM, 91.0%, 3.2 min).⁹⁹ The reusability of the SWCNT-Ru catalyst did not show any significant drop in degradation after the third cycle. The same research group observed that the SWCNT-Ru catalyst prepared using ethanol as a reducing agent degraded up to 98.2% of 0.04 mM CR dye solution within 4 min.¹⁰⁰ The formation of larger Ru clusters at a higher Ru loading reduced its activity.

Independent of the NP incorporation method (IM or DR), Pd–Au/C catalysts activated by MW irradiation exhibited better hydrodechlorination, H₂Cl, activity than that activated by thermal treatment in hydrogen.¹⁷⁴ The improved catalytic activity was attributed to the better homogeneity in the Pd–Al alloy. The deactivation of the PdAu-DR-MW catalyst was also lower owing to the better homogeneity of the Pd–Au alloy phase.

4.3. Photocatalytic/thermal degradation

A smaller size and homogeneous distribution of PtNPs and Mo₂C in Pt/G and Mo₂C/C catalysts, respectively, again showed good to excellent methylene blue (MB) photodegradation and hydrous hydrazine decomposition activities.^{111,156} The MW-assisted catalytic decomposition with a power of 180 W produced complete decomposition over the Mo₂C/C catalyst within 4 min. The WC/AC-H catalyst prepared by the rather long procedure of impregnation and carbothermal hydrogen reduction, produced complete hydrazine conversion at 50 °C.²⁰⁸ CeO₂–SiO₂ NPs abbreviated as M-30 (irradiation time 30 min), exhibited better MB decomposition activity under UV light, also due to their lower particle size and higher surface area and pore volume.¹⁶² In this case, the degradation efficiency was 99.9% at pH 7. On the contrary, the same CeO₂–SiO₂ catalyst (H-24 h) prepared *via* the hydrothermal process showed only 85% degradation efficiency.

4.4. Hydrogenation reaction

The selective hydrogenation reaction of *p*-chloronitrobenzene (*p*-CNB) to *p*-chloroaniline (*p*-CAN) in the presence of either Ru/CNT or Ru/FCNT catalysts indicated that the Ru/FCNT catalysts prepared without/with a PVP stabilizer had better activity with



100% selectivity at 100% conversion after 30 min of reaction.⁹⁵ These findings were consistent with the more homogeneous distribution of RuNPs and stronger anchoring of the RuNPs on the FCNT surface. Further, the selectivity (96%) of commercial Ru/CNT was comparatively lower. In a study on the effect of the precursor type (Ru(acac)₃/RuCl₃) on the catalytic activity of an Ru/TiO₂ catalyst, it was found that an increase in the irradiation time resulted in an increase in the particle size.¹⁴¹ This reduced the yield of γ -valerolactone (GVL) from the hydrogenation of levulinic acid (LA).

An MW-synthesized Ni/AB-MW catalyst was found to be superior to an Ni/AB-CH catalyst in the hydrogenation of nitrobenzene to aniline with respect to conversion (98.7%) and selectivity (93.3%).¹¹⁸ MW synthesis produced a smaller and uniform size as well as a higher loading of metallic Ni and NiO particles. The stability of the Ni/AB-MW catalyst was also better given that the Ni crystallite size indicated an increase in size by 27.4% after 60 h of reaction, while in the Ni/AB-CH catalyst, the size increase was more dominant (26.5% after 36 h).

Pd/Al₂O₃ catalysts prepared *via* plasma-induced MW irradiation, designated as PF (plasma fixed-bed reactor) and PS (plasma spouted bed reactor), were used for the hydrogenation of acetylene and compared with CM (conventional thermal treatments).¹³⁷ In the plasma fixed-bed reactor, three catalysts named PF-1, PF-2 and PF-3 were prepared by controlling the distance between the waveguide and particle bed in the range of 3 to 7 cm. Overall, PF-1 possessed the maximum activity due to formation of the Pd (100) plane and a better dispersion. The highest selectivity for consecutive acetylene hydrogenation towards ethane was also possible in the presence of PF-1. In another investigation, the authors proved that although plasma spouted bed treatment produced a higher gas temperature than conventional heat treatment at 500 °C, the former method prevented Pd metal from sintering due to its shorter reaction time.¹³⁸

4.5. Hydrodesulfurization/hydrodearomatization

Hydrodesulfurization (HDS) is an industrial process required for reducing the sulfide content in fuel. A W/Al₂O₃ catalyst (identified as MWH-B) prepared *via* the part-wise addition of oxalic acid showed enhanced HDS activity due to the good dispersion of WO₃.¹³⁹ The overall analysis suggested that the catalyst prepared *via* the MWH method had superior HDS activity relative to that prepared by IM. Nguyen *et al.* found that the HDS conversion of 4,6-dimethyldibenzothiophene (4,6-DMDBT) over the CoMo/Ti-SBA-15 catalyst was 1.35-times higher than that on the CoMo/SBA-15 catalyst.¹⁸⁵ Alternatively, the insertion of Ti in the CoMo/SBA-15 catalyst reduced the hydrodearomatization (HDA) activity of 1-methylnaphthalene (1-MN) and phenanthrene by half. Hence, the inclusion of Ti promised control of the HDA activity over the CoMo/SBA-15 catalyst.

4.6. Coupling reactions

To understand the deactivation process, Pd/C and Pd/Grp catalysts were subjected to MW irradiation for variable times (30 s, 5 min and 30 min) before the Suzuki–Miyaura

(phenylboronic acid and *p*-nitrobromobenzene) and Mizoroki–Heck (iodobenzene, butyl acrylate, and triethylamine) coupling transformations.⁹³ A longer MW irradiation time between 5 to 30 min produced a considerable loss in catalytic activity for the Pd/C catalyst, which was more severe for the Pd/Grp catalyst after irradiation for 30 s. The deactivation of the catalyst resulted from an increase in particle size, dissolution of carbon inside the metal NPs and formation of metal oxide. In another investigation, 1 mol% of Pd/G catalyst prepared by continuous flow synthesis produced 100% conversion in the MW-assisted batch Suzuki–Miyaura coupling reaction between bromobenzene and phenylboronic acid.¹⁰⁷ However, the recyclability test was not promising given that conversion decreased from 100% to 77% in the fourth cycle of the coupling reaction. Comparatively, the MW-assisted Suzuki–Miyaura coupling reaction between bromobenzene and phenylboronic acid over a magnetic Pd–Fe₃O₄/G nanocomposite showed better stability with 100% conversion up to the 3rd cycle.¹⁰⁹ Further recycling reduced the conversion to around 81% in the 9th cycle, which is still good. Controlling the MW temperature between 25–80 °C is important given that an increase in temperature to 100 °C reduced the conversion after 10 min of reaction, indicating destabilization of the catalyst.

Elazab *et al.* observed that a Pd content of 20 wt% in Pd/CuO catalysts is optimum for obtaining 100% conversion in the MW-assisted Suzuki–Miyaura coupling reaction between aryl halides and phenylboronic acid.¹³³ The application of the 20 wt% Pd/CuO catalyst for other substrates also produced good conversion (82–90%). The conversion drastically dropped to 60% in the sixth run due to the agglomeration of the PdNPs. Elazab *et al.* also investigated the applicability of Pd/CuO catalysts for the Mizoroki–Heck (aryl bromide and alkene) and Sonogashira–Hagihara (aryl bromide and alkyne) coupling reactions under MW irradiation.¹³⁴ The Mizoroki–Heck coupling reaction between bromobenzene and alkene on 2 mol% of the 20 wt% Pd/CuO catalyst produced the maximum conversion of 93% at the optimized temperature of 150 °C. The catalyst was found to be fairly stable up to 5 cycles.

Interestingly, the Pd/SiC catalyst, labeled as MW3, exhibited high catalytic activity up to 20 reaction cycles with more than >90% yield in the Mizoroki–Heck (iodobenzene and methyl acrylate) coupling reaction.¹⁴⁷ Comparatively, the OB3 catalyst, which was prepared *via* conventional heating, produced 100% yield in the first cycle, but this was drastically reduced to 75% yield. In contrast, the commercial Pd/C catalyst was usable for only one Mizoroki–Heck reaction cycle. The better catalytic activity, *i.e.*, longer catalytic life of MW3 was due to larger amounts of PdNPs on the support. MW3 was also found applicable in the Suzuki–Miyaura coupling reaction.

In comparison to the Pd/Ag/SBA-15 catalyst, the Pd/Au/SBA-15 catalyst showed superior activity in the Suzuki–Miyaura coupling reaction between iodobenzene and aryl boronic acid and produced ~70% conversion.¹⁸³ The enhanced activity of aryl halides followed the order of I > Br > Cl. A comparative study among the Pd–Fe₃O₄/G, Pd–Co₃O₄/G and Pd–Ni(OH)₂/G catalysts for the Suzuki–Miyaura coupling reaction (bromobenzene with phenyl boronic acid) showed the lowest (<60%) conversion



over the Pd-Ni(OH)₂/G catalyst but both the Pd-Fe₃O₄/G and Pd-Co₃O₄/G catalysts produced 100% conversion within 10 min at 120 °C.¹⁸⁸ The good nano-size distribution of the catalyst on G nanosheets in both cases substantially improved the catalytic conversion. The Pd-Fe₃O₄/G catalyst was fairly stable, producing 90% conversion in the fifth cycle. In an early article, Elazab *et al.* demonstrated that 0.3 mol% of magnetically recyclable Pd-Fe₃O₄/G catalyst, labeled as catalyst 3 (Pd 7.6 wt%), produced 100% conversion within 45 min in the Suzuki-Miyaura coupling reaction.¹⁸⁹ Despite having a low Pd content, catalyst 3 had the highest activity because it contained the smallest Pd (3–4 nm) and Fe₃O₄ NPs (12–16 nm).

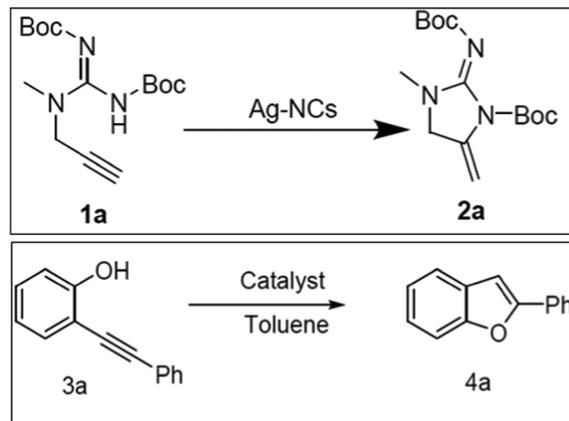
4.7. Synthesis reactions

The catalytic cycloisomerization of substituted propargylic urea to produce substituted imidazolone under optimized reaction conditions was carried out over MW-2%Ag@ALSBA-15, MW-2%Au@ALSBA-15, BM-2%Ag@ALSBA-15 and BM-2%Au@ALSBA-15 catalysts (Scheme 1).¹¹² Irrespective of their preparation methods, the supported Au nanocomposites were almost inactive with a yield of less than 5%. Alternatively, the ball-milled BM-2%Ag@ALSBA-15 catalyst was more active (yield, 56%) than the MW-synthesized MW-2%Ag@ALSBA-15 catalyst (yield 35%). However, despite the lower activity of the MW-synthesized catalyst, the authors claimed that the MW-assisted synthesis of the catalyst is favorable because it requires a few minutes for the one-step reduction compared to several hours required by ball milling-assisted synthesis. However, information on the size distribution, valence state and crystallinity of the nanocatalyst is not available, and thus it is difficult to predict the exact reason for the poor performance by the MW-2%Ag@ALSBA-15 catalyst. However, another author reported 99% yield of 2a (Scheme 2) with 100% conversion at 50 °C using a similar type of 0.96 wt% AgNCS/SBA-15 catalyst during the cyclization of propargylguanidine.¹¹⁶ This result is remarkable compared to that of Ag salts and AgNCS, although the metal loading is much lower.^{209,210} The conversion was retained at >90% conversion after 4 cycles, and no Ag leaching was detected and the morphology with the size distribution of the recovered catalyst remained unchanged. The cyclization of 2-(phenylethynyl)phenol in the presence of 0.96% AgNCS/SBA-15 catalyst also resulted in 100% conversion with 93% selectivity for 2-phenylbenzofuran (4a) at 70 °C (Scheme 2).

Song *et al.* observed superior CO₂ methanation activity over an Ni-Al₂O₃-M catalyst compared to that over an Ni-Al₂O₃-I



Scheme 1 The catalytic cycloisomerization of substituted propargylic urea.¹¹² (Copyright 2019, the American Chemical Society.)



Scheme 2 The catalytic cyclization of propargylguanidine (1a) and 2-(phenylethynyl)phenol (3a).¹¹⁶ (Copyright 2020, the American Chemical Society.)

catalyst.¹³⁶ The maximum conversion efficiency of 91.6% was possible at 325 °C over the Ni₂₀-Al₂O₃-M catalyst (20 wt% Ni) and the selectivity for CH₄ in the temperature range of 200–400 °C was around 99.3%. The catalyst was stable at 325 °C for up to 72 h given that the conversion remained high (91.6%). The anti-sintering stability of the Ni₂₀-Al₂O₃-M catalyst was confirmed.

Akay *et al.* tested a Co/SiO₂ catalyst in non-thermal plasma-promoted Fisher-Tropsch synthesis (FTS) involving CO and H₂ gases.¹¹⁹ Catalyst-ABC-550, where metal oxide was reduced in a plasma reactor under a stream of H₂ for 24 h at 550 °C, was found to be the best, producing 100% CO conversion. Catalyst-ABC-550 also possessed better stability given that the CO conversion of 100% remained constant over the complete test period of 175 h. It was possible to restore the complete catalytic activity of the deliberately deactivated Catalyst-ABC-550 by reduction with H₂ (25 mL min⁻¹) at a plasma power of 90 W for 24 h at 250 °C. In contrast, the Co/SiO₂ catalyst prepared by depositing Co monolayers on a 10 nm-thick SiO₂ film demonstrated poor stability in the FTS reaction.²¹¹ In another investigation, Co/SiO₂ catalysts prepared *via* conventional drying offered better CO conversion (68.3%) compared to that prepared *via* the MW method (42.1%) but the selectivity towards liquid hydrocarbons of C₅₊ was higher (63.2%) for the catalyst prepared *via* the MW method.¹²⁰ However, it is not clear whether the MW method caused any damage to the support or agglomeration.

A comparative study between MW-NiO/Vmt and CHT-NiO/Vmt catalysts for CO methanation in a stainless steel tubular microreactor in the temperature range of 250–500 °C showed the better CO conversion (99.6%) and CH₄ selectivity (93.8%) by the MW-synthesized MW-NiO/Vmt catalyst at 400 °C.¹⁶³ MW-NiO/Vmt was also found to be comparatively more stable.

The ethanol conversion in the range of 250–450 °C with a constant H₂ flow over Pt/SiO₂ catalysts, which were prepared with different Pt loadings *via* the MW-assisted solution and RCE methods, increased markedly with an increase in temperature, but decreased slightly with the Pt loading, which was



affected by the dispersion behavior of the PtNPs.¹²¹ The selectivity toward ethylene (dehydration) at 450 °C for the 0.6 and 0.9 wt% Pt-loaded catalysts prepared *via* the MW method was higher (50–53%) compared to that (47%) for the samples prepared *via* the RCE method.

The photocatalytic activity of TiO₂/Au NPs for H₂ production from a mixture of ethanol and water in the gas phase through a tubular reactor suggested that the removal of the capping agent, PVP, *via* calcination is necessary to improve the efficiency.¹⁶⁶ The optimal calcination temperature was 450 °C for the best H₂ evolution of ~5.3 H₂ g_{cat}⁻¹ h⁻¹ (sample A, 100 mg of PVP) and ~3.5 H₂ g_{cat}⁻¹ h⁻¹ (sample B, 600 mg of PVP). A higher calcination temperature (>550 °C) reduced the catalytic performance *via* aggregation. The higher catalytic activity of sample A was due to the larger interface area between Au and TiO₂, which favored the interaction between the Au and TiO₂ NPs.²¹² A comparative study of the catalytic property between Pd/Ag/SBA-15 and Pd/Au/SBA-15 catalysts for hydrogen production from ammonia borane (AB) under both dark and visible light irradiation conditions indicated the superior catalytic activity of the Pd/Ag catalyst.¹⁸³ Khalid *et al.* observed that the catalysis power of an Ag–TiO₂/G composite for hydrogen production from methanol aqueous solution was superior to that of the pure TiO₂, Ag–TiO₂ and TiO₂/G samples.¹⁹⁰ Hence, the inclusion of a G support and Ag played a vital role in enhancing the photocatalytic activity. The hydrogen production efficiency from methanol over four different copper-based catalysts (CuNiZnO/γ-Al₂O₃), which were prepared at different MW temperatures (80 °C, 85 °C, 90 °C and 95 °C) indicated that the content of Cu, Ni and Zn had an effect on the catalyst power.¹⁹⁶ The MW-Cu/Ni-80 catalyst contained a higher amount of Cu (47.92 wt%), resulting in higher activity at a lower reaction temperature (200 °C). In contrast, the other catalysts processed at higher MW temperatures contained larger amounts of Ni and Zn than the MW-Cu/Ni-80 catalyst, which played the main role for increasing methanol conversion at a higher temperature (≥225 °C). The catalytic selectivity with respect to H₂ production increased with an increase in the reaction temperature for MW-Cu/Ni-80, but for MW-Cu/Ni-90 and MW-Cu/Ni-95 this trend could not be distinctly confirmed and the overall yield was relatively high for both MW-Cu/Ni-90 and MW-Cu/Ni-95. The formation of an increasing amount of Ni possibly inhibited the formation of methyl formate and increased the selectivity for H₂ and CO in methanol decomposition.^{213,214} The MW-Cu/Ni-90 catalyst was fairly stable and methanol conversion was maintained at 100% for up to 26 h, only slightly decreasing to 98.8% after 30 h.

4.8. Electrocatalysis

In the last few decades, a considerable amount of research has been devoted towards the application of Pt NPs in PEMFCs,^{215,216} water splitting,²¹⁷ and direct alcohol fuel cells.²¹⁸ However, due to the high cost of Pt, its application is limited, and thus researchers are focusing more on the development of multicomponent (binary or ternary) catalysts to promote the catalytic performance. The combination of Pt with non-precious metals can also reduce

the dosage of Pt and is beneficial for improving its resistance to poisoning.^{219,220}

Sharma *et al.* studied the electrocatalytic activity of Pt/C electrocatalysts prepared using different precursors.⁹² A comparative study of the cyclic voltammograms (CV) suggested that relative to the commercial Pt/C catalyst, the H₂PtCl₆- and (NH₄)₂PtCl₆-sourced MW-synthesized Pt/C catalysts possessed a higher and almost identical electrochemical surface area (ECSA). The (NH₄)₂PtCl₆-sourced Pt/C catalyst exhibited a comparable ORR performance with that of the commercial and H₂PtCl₆-sourced catalysts. The electrocatalytic activity of CoSe₂/C catalysts prepared with different Se/Co ratios showed that an Se/Co ratio of 3.0 is the best for ORR activity and stability.¹⁷⁷ An excess amount of Se oxide reduced the ORR activity due to the occurrence of severe aggregation of the CoSe₂ NPs. Likewise, FeSe₂/C NPs prepared with different Fe/Se ratios exhibited the best ORR activity at Se/Fe ratios of 2.0 and 4.0, respectively.¹⁷⁸ These ratios were equivalent to 47.21% and 48.2% of the total amount of FeSe₂ and FeSe, respectively. An increase in Co content in a series of ternary (Co_xNi_{1-x})Se₂/C catalysts improved their electrocatalytic activities towards the ORR.¹⁹⁸ The beneficial effect of the incorporation of Co was further confirmed by the increase in the ORR potential (*E*_{ORR}). The incorporation of Co in binary NiSe₂/C was also advantageous for the enhanced stability of the (Co_{0.75}Ni_{0.25})Se₂/C electrocatalyst.

The addition of Sn to a Pt/C catalyst greatly enhanced its electrocatalytic activity for ethanol oxidation (EO), whereas the addition of Rh only modestly improved the electrochemical performance.¹⁷¹ Chronoamperometric tests suggested that PtSn/C is considerably less poisoned compared to the Pt/C or PtRh/C catalysts. Similarly, the inclusion of Cu in Pt/C (Pt3Cu/C, Pt4Cu/C, and Pt2Cu/C) catalysts improved their electrocatalytic performances compared to the commercial Pt/C catalyst for EO and MO.¹⁷³ The Pt3Cu/C catalyst had the highest mass activity, MA, (1.61 A mg⁻¹) and surface specific activity, SA, (3.23 mA cm⁻²) towards EO. The Pt3Cu/C catalyst also displayed the highest MA (1.92 A mg⁻¹) towards MO. The inclusion of Cu also improved the durability of PtCu/C towards the EO and MO reactions. SF-MWCNT-Pd and its mixed bimetallic SF-MWCNT-PdSn_{mix} and SF-MWCNT-PdNi_{mix} nanocatalysts were found to be more active towards EO compared to alloyed SF-MWCNT-PdNi and SF-MWCNT-PdSn catalysts produced *via* MW assisted co-reduction.¹⁷⁹ The addition of Sn to Pd produced the maximum influence on electrocatalytic property. However, chronopotentiometric experiments showed that SF-MWCNT-PdNi_{mix} was more tolerant to CO poisoning than SF-MWCNT-PdSn_{mix}. The electrocatalytic activity of PtNi/FCNT analyzed in MO was dependent on the metal loading and mode of irradiation.¹⁸⁰ The catalysts prepared at different irradiation times had almost the same stability, but higher than that of the reference Pt/C catalyst. Three different ternary catalysts labeled as Pt_{4.5}Sn_{1.5}Rh₁ USNP/C, Pt₃Sn₁Rh₁ USNP/C and Pt_{4.5}Sn₁Rh_{1.5} USNP/C also showed an improvement in electrocatalytic performance towards both MO and EO than 20 wt% Pt/C.¹⁹⁹ Notably, the highest MA and SA were observed for the Pt_{4.5}Sn_{1.5}Rh₁ USNP/C catalyst. The stability tests also



confirmed the superiority of the Pt_{4.5}Sn_{1.5}Rh₁ USNP/C catalyst. The structures and morphologies after 1000 sweeping cycles did not change for all the Pt–Sn–Rh/C catalysts, whereas for the 20 wt% Pt/C catalyst, severe aggregation was observed. A trimetallic AuPdPt@WC/C catalyst also showed an improvement in electrocatalytic MO reaction compared to the commercial Pt/C catalyst.²⁰⁰ The anti-poisoning ability of trimetallic AuPdPt@WC/C was 5-times better than that of the commercial Pt/C catalyst. The formation of highly dispersed AuPdPt metals on a WC support and the coupling effect between NPs enhanced the electrocatalytic performance.

The MW synthesis of PtRu/MWCNT NPs resulted in an improvement in poisoning resistance to CO following the formation of a homogeneously dispersed alloy phase.¹⁸¹ In a similar investigation, the PtPd/MWCNT-GNP catalyst was found to possess the highest electrochemical stability and durability compared to PtPd/MWCNT and PtPd/GNP catalysts.¹⁸² This behavior was ascribed to its better dispersion, smallest NP size and highest specific surface area (164 m² g⁻¹).

The electrochemical activity of the Co₃O₄@SUS electrode towards the OER in water splitting was compared with reference materials including commercial RuO₂@SUS and Co₃O₄-LNS@SUS catalysts.¹⁶⁵ Co₃O₄@SUS exhibited the most favorable OER kinetics due to its rod-like morphology and highest surface to volume ratio. The stability of the Co₃O₄@SUS catalyst was good towards the OER with 78% current retention after 24 h and the initial morphology of the catalyst was maintained. Meanwhile, a metal organic framework, Co–N–C, supported Ni₃Fe catalyst offered the best OER activity and best reaction kinetics compared to Ni₃Fe/N–C, IrO₂, and Co–N–C.¹⁸⁷ A negligible decay in current density was observed for the Ni₃Fe/Co–N–C catalyst after 3000 potential sweeps. Subsequently, the Ni₃Fe/Co–N–C showed a comparable ORR performance. Hence, the Ni₃Fe/Co–N–C catalyst can be an efficient bifunctional electrocatalyst for the OER and ORR. Pan *et al.* also demonstrated an improvement in OER activity for a supported Co/CoO_x catalyst following the inclusion of Co.¹⁹¹

5. Scope and outlook

This review article focused on evaluating the feasibility of MW-assisted methods over conventional methods for the preparation of durable and effective supported heterogeneous metal/metal oxide/their hybrid nanocatalysts. It was already mentioned that unsupported nanocatalysts are prone to aggregation, are unstable in air, and promote leaching, and thus deactivate easily during catalysis. In this regard, the anchoring or mobilization of nanocatalysts on an inert or catalytically active support could help to improve the catalyst lifetime by inhibiting leaching, enhancing its stability, reducing its poisoning and maintaining its nano-size distribution.

Numerous variations in MW-assisted routes have been developed and evaluated for the preparation of supported nanocomposite catalysts. The predominantly used methods studied include MW-assisted polyol/alcohol reduction of metal

salts,^{81,87,88,90–92,95,99,111,112,132,141,142,145,146,158,162,166,171–173,175–180,182,183,186,190,198,199} MW-assisted reduction in the presence of reducing agents or under H₂ pressure,^{98,106–109,116,121,127–130,133,134,161,181,188,189} MW heating with the aid of an MW absorber (ionic liquid),¹¹⁰ carbothermic MW reduction of impregnated metal salts,^{82,83} MW thermal reduction of impregnated dried precursors,^{75,84,85,96,97,105,154,163,164} drying and calcination under MW conditions followed by impregnation,⁸⁹ MW drying followed by impregnation,^{118,120,184} MW thermal decomposition of metal salts with or without subsequent drying and/or calcination,^{86,122,123,131,136,139,147,155,157,158,165,187,191,193,196} MW activation followed by wet impregnation of metal NPs,^{174,197} and MW-induced plasma irradiation.^{119,137,138} Most of these methods are carried out in a batch-wise fashion, except a few in continuous flow mode.^{107,116,128} The comparative study of MW-assisted methods and conventional methods, whenever investigated, demonstrated that MW methods promoted faster nucleation, produced more homogeneous smaller-size NPs within a shorter time, improved the catalyst loading and better alloying in the binary/ternary metallic phase because of the high loss tangent values, strong MW absorption, and thus faster heating and lower activation energy.^{112,118,120,121,136,138,139,160,162,163,174,184}

The usefulness of MW-assisted methods is dependent on the ability to produce a perfectly homogeneous distribution of uniform nanosized catalyst with a controlled morphology on a solid support. The intimate contact with strong fixation is also a desirable requirement from the view point of stability and good dispersity of the nanocatalyst on the support. In almost all the above-mentioned articles, it was found that MW-assisted batch methods under optimum conditions favorably produced a well-dispersed, uniform, nano-sized distribution of metal/metal oxide/their hybrid catalyst, reportedly down to even 1.6 ± 0.7 nm, which was observed for AgNCs.¹¹⁶ The addition of stabilizer was considered not essential to improve the size distribution, and thus the surface property of the nanocatalyst was easily maintained. The MW power, irradiation time, temperature, loading amount, functionality/porosity of the support materials, pH of the reaction medium, nature of the precursor salt and reducing agent all influenced the size distribution of the supported nanocatalysts. The use of a too high MW irradiation power and too long irradiation time sometimes caused sintering of the nanocatalysts^{116,141,157,162,180} and even damaged the support materials.⁹⁴ Thus, a balance is needed between MW power and irradiation time to achieve the optimum nanosize distribution. MW irradiation under vacuum and the selection of an appropriate precursor salt were also found to be crucial in minimizing damage to the support.⁹⁴ In some instances, functionality or porosity in the support materials was found beneficial for improving the metal loading, dispersibility and crystallinity index of the anchored nanocatalyst.^{86,95,110,116,120,121,162,180,191,193} Interestingly, a variation in the shape and morphology (*e.g.* spheroid, cubic, rod, plate, hollow, and porous) of nanocatalysts was also achieved by controlling the MW reaction conditions such as pH, irradiation temperature and time, precursor and support materials.^{81–83,110,129,130,162,165} However, despite the successful



preparation of various supported, well-dispersed nanocatalysts, the wide variations in MW-assisted methods needed to be optimized in terms of simplicity, scalability, economic feasibility, processability and reproducibility. In this regard, methods such as the one-pot MW polyol/alcohol reduction method, one-pot MW-assisted reduction in the presence of reducing agents and one-pot thermal decomposition of the precursor without additional calcination can be preferentially suitable for mass production. These particular methods have been proven beneficial for preparing a large variety of supported well-dispersed nanocatalysts (including Pd, Ag, Cu, Pt, Ru, Au, Pd-Fe₃O₄, Pd-Co₃O₄, Pt-Mo₂C, PtRh, PtSn, CoSe₂, and PdNi) without aggregation. In the above-mentioned *in situ* methods, the addition of a solvent is desirable because the solvent-free MW method was not found to be conducive for obtaining a homogeneous distribution due to the aggregation of the particles.^{96,97} Although scarcely studied, the continuous flow MW-assisted method, which is favorable for improving the reproducibility, batch consistency and yield, was found to be promising for preparing supported, well-dispersed finely tuned, uniform nanocatalysts on a mass scale.^{107,116,128} Importantly, the average size of the nanocatalysts produced in the continuous flow mode was smaller compared to that prepared *via* the batch method.^{116,128}

The second point is the analysis of the catalytic power of various MW-synthesized supported nanocatalysts, tested for different organic transformation reactions including electrocatalysis. In general, the MW irradiation conditions, particle size, dispersibility, morphology, loading amount, valence state of the active catalyst and nature of the support materials affect the catalyst efficiency. For example, irradiation for longer time reduced the catalyst efficiency due to a change in morphology, dispersity, damage to the support and formation of metal oxide.^{83,94,141} Morphology also plays a key role, given that cubic-shaped smaller nanocatalyst particles showed better catalytic activity, which is ascribed to their higher surface area.⁸¹ Table S1 (ESI[†]) summarizes the activity of various supported nanocatalysts prepared under MW irradiation and conventional reaction conditions. Interestingly, in most cases, the MW-assisted-synthesized supported nanocatalysts possessed better catalytic activity compared to the commercial^{95,101,147,162,186} and/or corresponding nanocatalysts prepared *via* conventional methods.^{118,120,121,136,139,147,162,163,174,221–223} Although the comparative data on the corresponding catalysts prepared by conventional methods is still inadequate, it is reasonable to conclude based on the available results that the MW-assisted-synthesized supported nanocatalysts possess good prospects for industrial use. Some authors even carried out the catalysis reactions under MW reaction conditions over supported nanocatalysts and found fairly good conversion and/or yield.^{112,119,120,128,156} The better catalytic activity of MW-synthesized supported nanocatalysts can be assumed to be due to the improved homogeneous distribution of smaller nanocatalyst particles, larger volume to surface ratio, surface heterogeneity, synergistic interaction between metals and stronger interaction between the nanocatalyst and support materials. The data on the stability of MW-synthesized nanocatalysts during recycle is scarce, and thus more investigation is needed in the area.

However, according to the limited available data, the stability and recyclability were found to be comparable to both the commercial and corresponding nanocatalysts prepared by conventional methods. In some instances, even better recycle stability was observed for MW-assisted synthesized supported nanocatalysts.^{118,164,174} However, although the data on the stability and recyclability of MW-synthesized nanocatalysts is encouraging, much more effort is necessary to further improve their performance, given that in some cases the activity decreased significantly after a few reaction cycles.^{85,86,95,98,99,107,116,133,174} The decrease in activity during recycle in either case occurred mostly due to the change in morphology, aggregation, graphitization and loss of the active catalyst *via* leaching of metal ions.^{84,86,118,133,134,155,164} In this regard, the one-pot synthesis of supported nanocatalysts under mild MW conditions without additional calcination and use of a functional support for strong fixation can be attempted to further improve the recycle stability, thus reducing both the aggregation and leaching of the metal ions. It is noteworthy to mention that in batch catalysis reactions in a constant volume reactor, the activity of the nanocatalyst decreases sharply compared to that under continuous flow reaction conditions in a flow tube reactor (Table S1, ESI[†]). The shorter residence time, more efficient heat transfer from the heating/cooling source due to faster mixing, faster reaction kinetics than their batch counterpart and reduction in frequent abrasion/collision due to the absence of additional stirring may have improved the catalyst life in the continuous flow reaction condition.

A considerable volume of research is also available on the testing of the electrocatalytic activity of MW-synthesized supported binary and ternary metal nanocatalysts, with the aim to reduce the cost/dosage of precious Pt-based supported nanocatalysts for applications in PEMFCs, water splitting and alcohol fuel cells. In many cases, improved electrochemical performances with improved durability were observed for MW-synthesized electrocatalysts compared to the reference/commercial Pt/C electrocatalyst.^{92,145,165,171,173,178,180–182,187,199} The synergistic coupling effect, smaller size, and electronic and ionic effect helped to improve the electrocatalytic performance of binary and ternary metal nanocatalysts. Continuous irradiation rather than pulsed mode of irradiation was found to result in better electrocatalytic activity and better CO anti-poisoning ability.¹⁸⁰ The electrocatalytic activity suggested that a porous support enhanced the property due to the improved mass-transport property.¹⁰⁵ Overall analyses suggested that bimetallic: FeSe₂/C, Pt-WC/C-F, PtSn/C, PtRh/C, Pt3Cu/C, PtNi/FCNT, PtRu/MWCNT, PtPd/MWCNT-GNP, Ni₃Fe/Co-N-C, and Co/CoO_x-40-MC1050 and trimetallic: Pt-Sn-Rh/C, AuPdPt@WC/C can be good replacement for the commercial Pt/C catalyst.

6. Conclusion

The MW-assisted methods available for the preparation of supported metal/metal oxide/their blend nanocatalysts were



reviewed with the aim to evaluate their suitability and present suggestions for improving their performance, if any. Numerous variations in MW-assisted methods have been developed for the preparation of a broad spectrum of supported nanocatalysts. In almost all the methods, well-dispersed, smaller, uniform supported nanocatalysts were successfully prepared with good catalytic properties. The limited data on durability and recyclability also indicated the better performance of MW-synthesized nanocatalysts compared to commercial and the corresponding supported nanocatalysts prepared *via* conventional methods. Thus, it can be concluded that MW-assisted methods can be used on an industrial scale for the preparation of supported nanocatalysts, and from now, efforts should be made to optimize the various processes into a few that can be used for mass scale production. Accordingly, the wide variations in MW-assisted methods need to be simplified and optimized into scalable, one-step *in situ* methods, where all the reactants (precursor salt, solvent, MW absorber, and polyol/alcohol/reducing agent) in the presence of the support materials can be MW treated. There is also scope to further improve the stability/recyclability of the used nanocatalyst, at least for up to 10 recycles. Presently only 5–6 recycles are possible, which are insufficient from an economic point. This can be done by eliminating high temperature calcination, which is employed on many occasions after MW synthesis, using a functional support to improve anchoring and by developing MW reactors for efficient heat transfer so that the MW power can be reduced to the minimum. Continuous flow MW reactors have also been tried on a limited scale, and this idea is interesting for mass scale production but more investigation is needed to check the efficiency of the reactor for synthesizing a wide variety of supported nanocatalysts with durable properties.

Author contributions

Hasan Ahmad: conceptualization, data curation, formal analysis, resources, writing – original draft, writing – review & editing, supervision. Mohammad Kawsar Hossain: visualization, software.

Conflicts of interest

The authors declare that there are no conflicts to declare.

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