**Review**

**Phyto-Nanocatalysts: Green Synthesis, Characterization, and Applications**

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**Abstract:** Catalysis represents the cornerstone of chemistry since catalytic processes are ubiquitous in almost all chemical processes developed for obtaining consumer goods. Nanocatalysis represents nowadays an innovative approach to obtain better properties for the catalysts: stable activity, good selectivity, easy to recover, and the possibility to be reused. Over the last few years, for the obtaining of new catalysts, classical methods—based on potential hazardous reagents—have been replaced with new methods emerged by replacing those reagents with plant extracts obtained in different conditions. Due to being diversified in morphology and chemical composition, these materials have different properties and applications, representing a promising area of research. In this context, the present review focuses on the metallic nanocatalysts’ importance, different methods of synthesis with emphasis to the natural compounds used as support, characterization techniques, parameters involved in tailoring the composition, size and shape of nanoparticles and applications in catalysis. This review presents some examples of green nanocatalysts, grouped considering their nature (mono- and bi-metallic nanoparticles, metallic oxides, sulfides, chlorides, and other complex catalysts).

**Keywords:** phyto-nanocatalysts; plant extracts; environmental applications; mono- and bi-metallic catalysts; metallic oxides nanoparticles; complex catalysts

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1. **Phyto-Nanocatalysts (PNC)—Next Generation of Catalytic Materials**

Catalysis represents the cornerstone of chemistry since catalytic processes are ubiquitous in almost all chemical processes developed for obtaining consumer goods. Started about 200 years ago, as an empirical scientific field and no more than an unusual aspect of chemistry, catalysis gains more and more attention due to his importance in obtaining new compounds. More than 90% of industrial products (pharmaceutical, fibers, fuels, detergents, polymers) result through several steps which need an appropriate catalyst. The role of catalyst in the process development consists not only to accelerate the chemical reaction, but also to improve the whole process, by reducing energy consumption, the formation of undesired products, and production increase. Our society, as we know, does not exist in the absence of catalysis.

Over the last few decades, catalysis has been successfully applied in developing methods for environmental protection, with notable examples being the three ways catalyst for purification of exhaust gases [1] and water treatment using catalysts in the frame of advanced oxidation processes [2–4]. Catalysis evolution encompasses two main categories: homogeneous type, when reactants and catalysts are in the same phase, and heterogeneous type, when reactant and catalysts are in different phases.
Heterogeneous catalysis represents the real breakthrough in this field due to the numerous advantages; therefore, a large number of scientific papers and patents were focused on developing new catalytic processes. However, the unchallenged performances of heterogeneous catalysis come with some drawbacks. Catalyst comes in the form of porous solids with internal channels and pores, in different shapes and sizes, in order to ensure a large number of active centers [5]. This morphology leads to some difficulties in modeling and experimental set-up of catalytic reactors due to diffusion phenomena and fluid transport processes. Therefore, a new approach is required to surpass the disadvantages of both types of catalysis and to combine their impressive advantages. Nowadays, nanocatalysis represents a critical way to combine all features of previous approaches: accessibility to catalytic sites, stable activity, and selectivity, easy to recover, and with the possibility to be reused [6,7]. Nanotechnology represents a significant step forward in many domains, not only in catalysis. From bulk to nanoparticles smaller than 100 nm, the properties of materials are considerably changing; hence, there is the possibility of a wide variation of applications, especially in catalytic processes. For example, cobalt oxide is less active for the oxidation of CO to CO$_2$ in bulk state even at a high temperature, but very active at room temperature in nano form. Additionally, the gold nanoparticles are very active in an extensive range of reactions [8], however, while they are in bulk state, they are inactive. Utilization of nanoparticles brings many improvements in processes where they are involved: cost-efficient, safer catalyst, reduced chemical waste, lower energy consumption, reduced global warming [9]. Even in the range of nanometric dimensions, the nano stable properties are shifted to a large extent. The color of suspension of CdSe nanoparticles is shifted from blue to red when particle size change from 2 to 8 nm [10,11]. Nanomaterials with catalytic properties are classified after their dimensionality [7] such us zero-dimensional (0D) (nonporousnanoparticles) [12,13], one dimension (1D) (nanorods, nanofibers) [12], two dimensional (2D) (nanosheets) [14] and three-dimensional (3D) nanomaterials, which include all types of nanoparticles in different shapes (nanorods, nanoshells, nanocages) [6]. Another essential feature of nanoparticles is an outstanding possibility of functionalization, which increases the number of applications in all technical fields not only in catalysis. The versatility of functionalized nanomaterials makes them so valuable and fit perfectly the gap between homogeneous and heterogeneous catalysis. Preparation of nanoparticles tailored in different forms and composition would be difficult without thorough investigations using modern characterization tools [9,15,16]: ultraviolet–visible (UV-Vis) spectroscopy, Fourier transform infrared spectroscopy (FTIR), X-ray diffraction (XRD), X-ray fluorescence (XRF), X-Ray photoelectron spectroscopy (XPS), etc. These are complementary techniques that assure a complete assessment of particle properties and therefore an accurate modality to assign the appropriate applications. The main advantages of using nanoparticles (NPs) in catalysis are:

- The possibility of high scale production of nanoparticles.
- Long-term stability of the nanoparticles.
- Controlled functionalization of nanoparticles can be achieved.

In order to use nanoparticles in chemical reactions, adequate preparation methods are needed. Since these materials are diversified in morphology and chemical composition, different methods were developed. All these methods fall into two main categories: top down synthesis and bottom-up synthesis [17,18]. The top-down synthesis relies on destructive methods that bring bulk materials to nano dimensions state. For this category, mechanical milling, chemical etching, and sputtering are the most common methods used. The bottom-up synthesis uses an opposite approach, generally starting from atomic or molecular state and building up to nanoparticle dimensions (1–100 nm). Spinning, template-based synthesis, plasma or flame spraying, laser pyrolysis, atomic or molecular condensation, chemical vapor deposition, sol-gel, precipitation, solvothermal processing, microemulsion, and sonochemistry are the usual methods for bottom-up approach. These methods are state-of-the-art in nanoparticles manufacturing and allow the preparation of a broad range of solid materials. However, this technique either requires some unique and expensive equipment with high capital costs or expensive and environmentally hazardous reagents. Preparation of nanoparticles in the bottom-up mode with or
without the use of reagent is mainly based on the chemical reduction or electrochemical reduction processes. Generally, metal salts are reduced using sodium borohydride, cetyltrimethylammonium bromide, ascorbates, glucose, and citrates [19–21]. The nature of the reducing agent determines the size of the nanoparticles. Strong reducing agents produce bigger particles, while mild reducing agents induce formation of nanoparticles with smaller dimensions. Particle morphology control is achieved by utilizing appropriate surfactants (sodium dodecyl sulfate, Triton X, sodium formaldehyde sulfoxylate) and stabilizers (polyvinyl pyrrolidine (PVP), polymethylmethacrylate (PMMA), polyethylene glycol (PEG) and polymethyl acrylic acid (PMAA)). Sonochemical reduction of transitional metals occurs through acoustic cavitation and particle growth in a sonicated liquid medium. Improvement of nanotechnology field in general and nanocatalysis, in particular, require, besides the development of new catalytic systems, better preparation methods with less energy consumption and less harmful reagents. Hence, a clean, biocompatible, and eco-friendly process to synthesize nanoparticles is needed.

Over the last few years, new nanoparticle preparation methods have emerged by replacing the usual reagents with plant extracts obtained in different conditions. Plant extracts were used from centuries, as cures in folk medicine and since they have valuable properties, they are also employed in present days being a serious competitor for synthetic pharmaceuticals. The market of medicinal plants increases at a rapid pace each year and predicts for the year 2029 that it will surpass USD 129 billion in revenue. The herbal medicine field has experienced a revival in recent years due to the growing demand among consumers for ecological oriented chemical-free treatments. The main reason for the boost in utilization of plant extracts consists of the synergic effect of all components, which belong to a broad range of classes of organic compounds, and therefore, increase the therapeutic effects. The complexity of herbal extracts is precisely the reason why these liquids can be selected as substrates for nanoparticles synthesis, since they contain simultaneously reducing, stabilizers, and surfactants agents with natural origins, and hence, eco-friendly properties.

Transition to this approach was made based on substantial evolution over the last 20 years of research strategies in the area of natural product chemistry. Advanced extraction and separation methods [22,23] (hyphenated techniques: High-performance liquid chromatography-mass spectrometry (HPLC–MS), HPLC-nuclear magnetic resonance (HPLC-NMR), HPLC–MS–NMR, HPLC–diode array detector–MS–NMR, centrifugal partition chromatography) allow researchers to improve the isolation of secondary metabolites that help to build extensive databases of phytoconstituents present in different plants. Development of chemically engineered extract, dereplication, chemogenomics, metabolomics, fingerprinting, and chemical fingerprinting bring researchers a selection base for new compounds able to participate in nanoparticles synthesis. Diversification of extraction methods [24,25], with classic (Soxhlet extraction, percolation, maceration, hydro distillation, and steam distillation) or with modern instruments such as microwave-assisted extraction, accelerated solvent extraction, ultrasound-assisted extraction, enzyme assisted extraction, pulsed electric field extraction, supercritical fluid extraction, ohmic-assisted hydro distillation, and utilization of numerous solvents with high variation of polarity open the opportunity to obtain a more tailored extracts, with the appropriate composition for obtaining nanoparticles. The main reason for developing a plant-mediated synthesis of nanocatalysts is the staggering number of plat metabolites which can be used, and removing the use of potential hazardous reagents.

For example, more than 8000 polyphenolic compounds are already identified in different plant species [24]. The most majority occur in conjugated forms with sugar residues linking with compounds like lipids, organic acids, amines, or even other phenols is very usual. Other important vegetal constituents (alkaloids, tannins, iridoids, secoiridoids, coumarins, and terpenoids), also in large numbers, can sustain an ecological path for preparing nanocatalysts.

The field of nanomaterials phytosynthesis in general (and nanocatalysts phytosynthesis in particular) represented the focus of an increasing number of studies in the last decades, from all-over the world, as proven by the survey performed using Scopus database (Figure 1). The search was performed over the time period 2010–September 2019, using the keywords “nanoparticle extract”,...
“nanoparticle phytosynthesis”, and “nanoparticle natural extract”, with “catalyst” within the obtained results; multiple keywords were used, as the term “phytosynthesis” is not adopted by all authors.

Figure 1. Distribution of publications per year from Scopus Database according to the presented keywords, with duplicate data and “false-positive” results removal.

Considering these aspects, the current review focuses on the metallic nanocatalysts’ importance, different methods of synthesis with emphasis to the natural compounds used as support, characterization techniques, parameters involved in tailoring the composition, size and shape of nanoparticles, and possible applications (as reported in the literature). The review will briefly present some examples of green nanocatalysts, grouped considering their nature (mono- and bi-metallic nanoparticles, metallic oxides, sulfides, chlorides, and other complex catalysts).

2. Mono- and Bi-Metallic Catalysts: Phytosynthesis, Characterization, and Application

The most encountered materials, when speaking of phytosynthesized nanoparticles, are represented by the noble metal nanoparticles (especially silver and gold). Several review and research papers were published in the last years, covering the green synthesis (including phytosynthesis) of monometallic and bimetallic nanoparticles [26–32], with focus especially on their biological properties. The following subchapters will present some examples on the application of such nanoparticles in the area of catalysis.

2.1. Silver Nanoparticles (AgNP)

Phytosynthesis of silver nanoparticles (AgNP) was described by Gangula et al. [33] using the stem extract of Breynia rhamnoides, the reduction of the metal salts being attributed by the authors to the phenolic glycosides and reducing sugars present in the extract. The obtained nanoparticles were used for the reduction of 4-nitrophenol to 4-aminophenol (4-AP) in the presence of NaBH₄, reaction that was found to be depend on the nanoparticle size. In 2014, Vidhu and Philip [34] obtained almost spherical silver nanoparticles, with diameters of approximately 20 nm, having excellent catalyst properties for the reduction of Methylene blue.
Silver nanoparticles were obtained using aqueous extracts of dried *Alstonia macrophylla* leaves by Borase et al. [35], having an average diameter of 70 nm. The nanoparticles proved to be efficient in the catalytic reduction of 4-nitrophenol and *p*-nitroaniline to 4-aminophenol, and *p*-phenylenediamine, respectively. Similar silver nanoparticles were obtained by Vidu and Phillip [36] and Nasrollahzadeh et al. [37] using aqueous extracts of *Trigonella foenum-graecum* seeds and *Euphorbia condylacarpa* *M. bieb* root. The nanoparticles were evaluated by the authors for the catalytic degradation of methyl orange, methylene blue and eosin Y [36] for the catalytic synthesis of N-monosubstituted ureas in water [37], with promising results.

Reduction of 4-nitrophenol to 4-aminophenol was also studied by Naraginti and Sivakumar [38] and Seralathan et al. [39] using silver nanoparticles obtained via aqueous *Coleus forskohlii* root, and *Salicornia brachiata* extract, while the reduction of methylene blue by phytosynthesized silver nanoparticles was studied by Ashokkumar et al. [40,41]. The differences in terms of catalytic activity observed between the two studies can be assigned to the variation in particle size.

Using phytosynthesized silver nanoparticles, reduction of 4-nitrophenol to 4-aminophenol was achieved in 2015 by Bindhu and Umadevi [42] and Joseph and Mathew [43]. Gavade et al. [44] obtained silver nanoparticles (20–30 nm) using aqueous *Ziziphus jujuba* leaf extract for the reduction of the silver salt, which was successfully used for the reduction of 4-nitrophenol and methylene blue. Degradation of methylene blue was also achieved by Ajitha et al. [45,46] using silver nanoparticles phytosynthesized by *Momordica charantia* leaf broth and *Lantana camara* leaf extract. The authors also propose a mechanism for the phytosynthesis of AgNP using *M. charantia*, assigning the main role in the silver reduction process to the flavonoid content of the extract. Several environmentally hazardous organic dyes (methylene blue, methyl orange, and methyl red) were degraded by Ahmed et al. [47] using silver nanoparticles obtained using a Ferredoxin–NADP$^+$ reductase/ferredoxin (from spinach leaves). Two other studies published in the same year describes the catalytic application of phytosynthesized AgNP for the degradation of cellulose by cellulase [48] and for the synthesis of propargylamines [49].

Reduction of 4-nitrophenol using AgNP obtained by a greener route (involving the use of commercially available gallic acid, a phytoconstituent commonly found in natural extracts) was presented in 2016 by Park et al. [50]. In the same year, several other studies successfully applied the phytosynthesized AgNP for the degradation of dyes: direct yellow-12 [51], Acridine Orange [52], Congo red and methyl orange [53], cresyl blue [54], and methylene blue [55].

A reduction of 4-nitrophenol by AgNP was also presented in 2017 by Bello et al. [56] (reaching 95% degradation after 36 min), Arya et al. [57], Manjari et al. [58], Farhadi et al. [59], Francis et al. [60], Bonigala et al. [61], Muthu and Priya [62], Karthika et al. [63], Mohanty and Jena [64], Patil et al. [65], and Naraginti and Li [66]. Similar catalytic properties of silver nanoparticles were demonstrated for the degradation of Congo red [56,63], methylene blue and Congo red [57,58], methylene blue, methyl orange and methyl red [61], methyl orange [62], methylene blue [64,66], eosin Y (>97% degradation after 60 min) [67], Congo red [63,68], Coomassie Brilliant Blue G-250 [69], and a reduction of *p*-nitroaniline to *p*-phenylenediamine [70]. Phytosynthesized silver nanoparticles were also evaluated as catalysts for hydration of cyanamides in aqueous medium [71] and for the construction of pyrimido[1,2-b]indazole derivatives under solvent-free conditions [72].

In 2018 studies, Zaheer [73] and Arya et al. [74] presented the catalytic degradation of the carcinogenic dye 4-nitrophenol, using silver nanoparticles phytosynthesized in aqueous palm date fruit extract and aqueous *Prosopis juliflora* bark extract. The two studies noticed a 95% degradation in 8 min, correlated with the amount of AgNP used, respectively a 90% degradation in 80 min. Zaheer [73] proposed a mechanism for the phytosynthesis of the AgNP, assigning the main role in the reduction of the silver salt to the pyranoid fructose present in the fruit extract. Vijayan et al. [75,76] presented the reduction of *o/p*-nitroanilines, respectively, 4-nitrophenol and *o/p* nitroaniline catalyzed by AgNP obtained using leaf extract of *Indigofera tinctoria* and aqueous extract of *Orthosiphon aristatus* leaves, respectively. In the first study, the reaction was completed in 10 min for both *p*- and *o*-nitroaniline, while in the second case, the reactions were completed in 6 min for *o/p*- nitroaniline and in 10 min for
4-nitrophenol. Similar observations (but with longer reaction times) were made by Francis et al. [77] for the reduction of 4-nitrophenol, 2-/4-nitroaniline, and eosin Y, using AgNP obtained by aqueous leaves extract of *Elephantopus scaber*. Reduction of 4-nitrophenol, methylene blue, methyl orange and methyl red, and, respectively, 4- nitrophenol, methylene blue and Congo red, was described by Bonigala et al. [78] and Vijayan et al. [79], using AgNP obtained by the application of *Stemona tuberosa* Lour and *Myxopyrum serratum* A.W. Hill extracts.

Photocatalytic degradation of methylene blue was monitored by Khan et al. [80] and Sumitha et al. [81] using extracts of Longan fruit peel and *Durio zibethinus* seeds. Their results showed a 99% degradation of methylene blue in 7 min and 73.49% in 180 min, respectively. The differences between their results can most probably be attributed to the differences both in terms of morphology and average size between the obtained NPs (as presented in Table 1). Application of AgNPs for the degradation of several other dyes (Reactive Black 5, methyl orange, direct yellow-142; Coomassie Brilliant Blue; methylene blue, eosin yellowish, safranin, direct dye, reactive dyes; methyl orange, rhodamine B; Rose Bengal) was further investigated in a series of research papers, using AgNP phytosynthesized via *Convolvulus arvensis* [82], *Gardenia jasminoides* Ellis [83], *Allium cepa* [84], *Nervalia zeylanica* [85] and *Bauhinia tomentosa* Linn. [86] extracts.

An extensive study was also published by Nakkala et al. [87] on the phytosynthesis of silver nanoparticles using aqueous rhizome extract of *Acorus calamus*. The authors evaluated the obtained AgNP for *in vitro* anticancer activity, in vivo toxicity in rats, as well as for the catalytic degradation of several dyes: 4-nitrophenol, 3-nitrophenol, 2, 4, 6-trinitrophenol, picric acid, Coomassie brilliant blue, Congo red, eosin Y, rhodamine B, methylene blue, methyl red, methyl orange, cresol red, acridine orange, eriochrome black T, and phenol red. The degradation reactions were completed in 6–60 min, for all the tested dyes.

In a study published in 2019, Wang et al. [88] evaluated the catalytic properties of AgNP obtained using aqueous *Ginkgo biloba* leaves extract for the reduction of 4-nitrophenol, Congo red, methyl orange, and rhodamine B, achieving full conversion in 15–100 min. Vijayan et al. [89] applied an aqueous extract for the phytosynthesis of AgNP and evaluated the obtained materials for the catalytic degradation of methylene blue and rhodamine B. Yu et al. [90] evaluated the influence of reaction temperature on the phytosynthesis of AgNP via *Eriobotrya japonica* (Thunb.) leaves extract. Their observations were that the reaction temperature had a direct effect on the phytosynthesized nanoparticles (the higher the temperature, the larger the obtained NPs) and, as a consequence, on their catalytic properties.

Silver nanoparticles are, by far, the most encountered phytosynthesized metallic nanoparticles, mainly due to their known antimicrobial properties; the last years witnessed the increased application of the silver nanoparticles in other areas, including in catalytic applications. As is the case for the other nanoparticles, the efficiency of their final application and catalytic rates is related to their particle size and shape. The morphology of the obtained nanostructures depends on phytochemical composition of the extracts, which in turn varies depending on different extraction parameters. In perspective, for the improvement of AgNP phytosynthesis, it is necessary to optimize the synthesis process (in terms of used extract composition); another viable perspective is represented by the use of other groups of plant (less explored in comparison with common angiosperms and algae) like bryophytes and pteridophytes. These somewhat under-exploited plants contain highly oxidant agents and might be involved in producing stable nanoparticles.

2.2. Gold Nanoparticles (AuNP)

Although not as abundant as the papers regarding AgNP, several research articles offer numerous examples regarding the application of phytosynthesized gold nanoparticles (AuNP) in catalytic processes. In 2010, Gupta et al. [91] proposed the application of AuNP phytosynthesized using green tea for the catalytic reduction of methylene blue in the presence of stannous chloride. In 2012, Ghosh et al. applied phytosynthesized AuNP for the reduction of 4-nitrophenol [92], while in 2014, several studies present the application of phytosynthesized AuNP for the reduction of 4-nitrophenol [93,94] or
methylene blue [94]. Application of phytosynthesized AuNP for the degradation of 4-nitrophenol was also described in 2016 by Borhamdin et al. [95], Ghosh et al. [96] and Lim et al. [97]; the reduction of the anthropogenic pollutant o-nitroaniline using AuNP was presented by Dauthal and Mukhopadhyay [98], who also proved their catalytic recyclability, while Choudhary et al. [99] presented the phytosynthesis of AuNP using Lagerstroemia speciosa leaf extract and their application for the reduction of different pollutants in wastewater (methylene blue, methyl orange, bromophenol blue, bromocresol green, and 4-nitrophenol). The authors also suggested a mechanism for the phytosynthesis: according to the authors, the biomolecules from the extract (such as penta-O-galloyl-D-glucopyranose, lagerstroemin, corosolic acid) attract negatively charged gold anions, while (through the transformation of phenolic C-OH groups to C=O keto) releasing electrons involved in the reduction of gold anions. In two 2017 studies, phytosynthesized AuNP were applied for the reduction of 4-nitrophenol [100] and methylene blue [101]. The application of AuNP for the reduction of 4-nitrophenol was also presented by Wacławek et al. [102] in their 2018 study. The authors managed to tune the shapes and sizes of the nanoparticles as a function of the pH of the synthesis solution, extract and gold salt concentration. Additionally, the comparison of the catalytic properties of different shaped AuNP (spherical and triangular) showed that the reduction of 4-nitrophenol was achieved 1.5 times faster using triangular AuNP.

Several other studies describe the application of phytosynthesized gold nanoparticles, by comparison with silver nanoparticles. The gold nanoparticles phytosynthesized by Gangula et al. [33] using Breynia rhamnoides extract showed superior catalytic properties for the reduction of 4-nitrophenol compared with the silver nanoparticles. A similar observation was made by Joseph and Mathew [43], who observed superior catalytic activity of gold nanoparticles phytosynthesized using aqueous Aerva lanata leaves extract, compared with silver nanoparticles and by Francis et al. [60] for the degradation of 4-nitrophenol, rhodamine B and methyl orange. On the contrary, Park et al. [50] noted superior catalytic activity for AgNP, compared with AuNP, both types of NPs obtained using gallic acid as a reducing agent. This can also be explained by the lower dimensions of the AgNPs, compared with AuNPs. A similar observation was made by Manjari et al. [58], who observed minor differences between the degradation capacities of AuNP and AgNP towards methylene blue and Congo red (99% for AuNP, 99.9% for AgNP) and 4-nitrophenol (95% for AuNP and 97% for AgNP), also correlated with the variation of the particle dimensions. In a 2017 study, Karthika et al. [63] found gold nanoparticles to have inferior catalytic properties for the degradation of 4-nitrophenol and Congo red compared with AgNP.

The comparative study of Vijayan et al. [75] regarding the application of AgNP and AuNP for the reduction of o- and p-nitroaniline showed inferior results for AuNP (complete degradation time 18 min), that the authors assigned to the irregular shape of AuNP (although the NPs had similar dimensions). Similar, to the irregular shapes of the AuNP obtained was assigned by Bonigala et al. [78] and Vijayan et al. [79] the inferior catalytic properties, compared with AgNP, observed for the reduction of 4-nitrophenol, methylene blue, methyl orange and methyl red, and, respectively, 4-nitrophenol, methylene blue, and Congo red. The AuNP obtained by Vijayan et al. [88] using Bauhinia purpurea leaf extract presented longer times (when compared with AgNP obtained in the same conditions). Again, it can be assumed that the differences in terms of catalytic activity are mainly due to the differences observed in terms of morphology.

Recently, a very interesting review paper was published by Teimouri et al. [103], presenting the phytosynthesis of AuNP, their characterization, and application for the degradation of 4-nitrophenol from industrial wastewater, alongside their insecticidal activity.

Regarding the phytosynthesis of AuNP, it can be noticed the variation in terms of NP morphology. The study suggested that triangular-shaped AuNPs with small dimensions present the best catalytic properties.
2.3. Palladium Nanoparticles (PdNPs)

Palladium represents an important metal, widely used in environmental protection, for numerous catalytic applications \[104,105\]. Considering its catalytic potential, palladium became in the last years an important research topic, several works describing its phytosynthesis.

In 2014, Khan et al. \[106\] presented the phytosynthesis of palladium nanoparticles (PdNP) using Pulicaria glutinosa extract, as well as their catalytic application in the Suzuki coupling reaction, obtaining a complete conversion of bromobenzene to biphenyl in under 5 min. The authors assign the reduction of Pd to nanoparticles to the presence of flavonoids and polyphenols of the used extract. Veisi et al. \[107\] presented in 2015 the phytosynthesis of monodisperse PdNP using herbal tea extract, for the same catalytic application. The authors suggested that flavonoids and polyols play the main role in the NP production. More than that, by conducting the Suzuki reaction open to air, they were able to prove the PdNP stability, while the catalyst was able to support eight catalytic cycles, without any loss in activity. In the same year, Nasrollahzadeh et al. \[108\] applied phytosynthesized PdNP for Hiyama and Stille cross-coupling reactions, observing a minimum five cycles of reusability for the catalyst. Duan et al. \[109\] phytosynthesized PdNP using an aqueous extract of Eucommia ulmoides bark, obtaining spherical and quasi-spherical nanoparticles, with an average size of 12.2 nm. The obtained NPs showed very good activity for the excellent catalytic activity for the electro-catalytic oxidation of hydrazine and the catalytic reducing degradation of p-aminoazobenzene, alongside a good stability in time in aqueous solutions.

In 2016, Nasrollahzadeh and Mohammad Sajadi \[110\] presented the application of phytosynthesized PdNP as a heterogeneous catalyst for the Suzuki-Miyaura coupling reaction, while Tahir et al. \[111\] studied the influence of phytosynthesis temperature on the dimensions of PdNP and applied the optimized materials to the photodegradation of methylene blue, obtaining a 90% degradation in 70 min. The authors assign the rapid degradation of the dye to small sizes of the PdNP and to the phytoconstituents attached to their surface.

An interesting approach was adopted by Hazarika et al. \[112\], who used for the phytosynthesis Garcinia pedunculata Roxb as bio-reductant and starch as bio-stabilizer. They obtained small-size, dispersed NPs, with very good catalytic for the Suzuki-Miyaura cross-coupling reaction, alcohol oxidation and Cr(VI) reduction. In the same year, other groups presented the application of phytosynthesized PdNP for the degradation of different compounds (such as 4-nitrophenol or Congo red) \[113–115\] or for selective oxidation of alcohols \[116\].

In a recent work, Garole et al. \[117\] presented the phytosynthesis of PdNP assisted by Lagerstroemia speciosa leaves extract. The obtained nanocatalyst (1.36 nm) was used for the reduction of organic pollutants (methylene blue, methyl orange, and 4-nitrophenol) in the presence of sodium borohydride, obtaining complete reduction within 10 min.

2.4. Iron Nanoparticles (FeNP)

Njagi et al. \[118\] used aqueous sorghum bran extracts for the phytosynthesis of amorphous iron nanoparticles (FeNP), which were successfully used for the H₂O₂-catalyzed degradation of bromothymol blue. Machado et al. \[119\] obtained FeNP using black tea, grape marc, and vine leaves aqueous extracts that were applied for the degradation of ibuprofen both from aqueous solutions and sandy soils, obtaining degradation efficiency up to 66% for aqueous solutions (pH dependent) and over 95% for soils.

Lin et al. \[120\] studied the effect of reaction atmosphere on the dimensions of FeNP phytosynthesized using green tea extract. The authors noticed that larger particles were obtained in oxygen atmosphere, that, through XPS determinations were proved to be predominantly iron oxide in nature. The FeNP obtained in nitrogen atmosphere had smaller dimensions (approximately 84 nm) and had superior capacity for the removal of methylene blue (98.7%, 85.7% within 5 min).
Using phytosynthesized FeNP, different degradation of different dyes (such as methylene blue, methyl orange, allura red, brilliant blue, green S, or rhodamine B) was achieved by Garole et al. [121], Radini et al. [122], and Khan and Al-Thabaiti [123].

2.5. Copper Nanoparticles (CuNP)

Nasrollahzadeh and Sajadi [124] described the phytosynthesis of copper nanoparticles (CuNP) using Ginkgo biloba L. extract and their application for Huisgen [3 + 2] cycloaddition of azides and alkynes at room temperature. The authors obtained spherical nanoparticles, with a narrow size distribution (15–20 nm), and observed over 93% yields for the studied reaction, at 5 h and a concentration of the CuNP of 10 mol%. Prasad et al. [125] obtained, using broccoli extract, CuNP with spherical morphologies and average size of 4.8 nm. The CuNP were successfully applied for the reduction of 4-nitrophenol in the presence of NaBH₄ (reaction completed within a few minutes with high values of reaction rate constants) and degradation of methylene blue (68% degradation after 28 h) and methyl red (32% degradation after 28 h). At the same time, the authors performed catalyst re-usability studies, observing no significant change in the conversion percent even at the fifth cycle; however, a prolongation of the time necessary to complete the reaction (at the fifth cycle, the time necessary being of 12 min). Superior results were obtained by Nazar et al. [126] for the catalytic degradation of methylene blue (87.11%, after 3 h) using CuNP phytosynthesized via Punica granatum seeds extract.

Nasrollahzadeh et al. [127] used aqueous extract obtained from Plantago asiatica to phytosynthesize CuNP with spherical morphology and dimensions between 7-35 nm, in 5 min. The nanoparticles were applied as catalyst for the cyanation of aldehydes, obtaining yields ≥85% (pure isolated product) at reaction times 30–60 min, superior to the classical methods for the synthesis of aryl nitriles.

2.6. Bi-Metallic Nanoparticles

Besides the mono-metallic nanoparticles, another important area in nanomaterials phytosynthesis is represented by the bi-metallic nanoparticles. These types of nanoparticles can come in a wide variety of sizes and morphologies [28]. As can be expected, the majority of the studies regarding catalytic applications of bi-metallic nanoparticles are focused on Ag/Au NPs. Ravi et al. [128] described the phytosynthesis of crystalline Ag/AuNP using Silybum marianum seed extract as reducing and stabilizing agent. The spherical polycrystalline NPs had dimensions ≤40 nm and exhibited good catalytic performance for the reduction of 4-nitrophenol. The Ag/AuNP (10–20 nm) phytosynthesized by Karthika et al. [63] using bark extract of Guazuma ulmifolia L. also showed good reducing capability towards 4-nitrophenol and Congo red. More than that, the bimetallic nanoparticles were found to have the best catalytic properties for the decolorization of 4-nitrophenol, compared with individual nanoparticles.

Bi-metallic Fe/PdNP were obtained by Luo et al. [129] using grape leaf aqueous extract. The nanoparticles exhibited significant catalytic effect for the removal of Orange II dye (98% in 12 h). The kinetic study performed by the authors revealed that the removal fitted to the pseudo-first-order reduction and pseudo-second-order adsorption model, implying that the removal phenomenon involved both adsorption and catalytic reduction.

Al-Asfar et al. [130] phytosynthesized bi-metallic Ag/FeNP using Palm date fruit extract. The nanoparticles presented as disks and with irregular morphologies, with dimensions 5–40 nm. The authors observed a complete decolorization of the bromothymol blue dye in 60 min for the Ag/FeNP/H₂O₂ catalytic system. Similar bi-metallic nanoparticles were also obtained by Taghizadeh et al. [131] using Cupressus sempervirens extract and their catalytic activity was tested for methyl orange removal, reaching a 98.5% removal after 4 h.
3. Metallic Oxides and Chlorides Nanoparticles

Unlike mono-metallic nanoparticles, metal oxides nanoparticles cover a wider range of metals (such as Fe, Cu, Zn, Zr, Ce, Ni, and others). Their phytosynthesis was recently reviewed by Sharma et al. [132], Singh et al. [133], or Basnet et al. [134], covering the plausible mechanisms involved in their formation, as well as potential application in catalysis.

Phytosynthesis of Fe$_3$O$_4$ was presented by Basavegowda et al. [135]. The authors used aqueous extract of *Artemisia annua* leaves to obtain ferromagnetic nanoparticles, with average diameters of 6.4 nm. Their application as catalyst in organic synthesis led to the synthesis of benzoxazinone and benzthioxazinone derivatives in high yields (over 80%). Vasantharaj et al. [136] proposed the use of *Ruellia tuberosa* leaves extract for the phytosynthesis of hexagonal nanorods Fe$_3$O$_4$NP. The obtained NPs were used for the degradation of crystal violet dye under solar irradiation, obtaining a degradation of 80% after 150 min reaction.

Spherical copper oxide nanoparticles (CuONP) were obtained by Nasrollahzadeh et al. [137] in 2015 using an aqueous *Gundelia tournefortii* leaves extract. The nanoparticles were applied for the degradation of 4-nitrophenol (observing the disappearance of the specific 4-nitrophenolate ions UV-Vis peak in 70 seconds after the addition of CuONP) and for the hydration of cyanamides to N-monosubstituted ureas using acetaldoxime (with yields over 88%). Considering the catalyst reusability, the authors observed a decrease in the reaction yield of phenylcyanamide hydration (from 94% for fresh and 1st cycle catalyst to 90% for the 4th cycle). The same group [138] obtained CuONP using *Thymbus vulgaris* aqueous leaves extract and applied them for the N-arylation of indoles and amines. The obtained NPs (having diameters under 30 nm) exhibited good catalytic activity (58–98% yields for the arylation of indoles with aryl halides and 86–96% yields for the arylation of amines with aryl iodides). Regarding the catalyst reusability, the authors observed that the catalyst could be used for five times with almost consistent activity (97–92% yield) for the N-arylation of indoles.

Devi et al. [139] obtained CuO and ZnONP using the aqueous extract of *Centella asiatica*, considering as phytosynthesis responsible molecules the polyphenols and protein (or amino acids) constituents of the extract. When applying the obtained NPs as catalyst for the reduction of methylene blue using NaBH$_4$, the authors observed a higher catalytic activity of CuONP (rate constant 4.261, compared with 0.311 for ZnONP), that was attributed to their lesser work function value. The catalytic application of phytosynthesized CuONP was also studied by Sathiyavimal et al. [140], Prakash et al. [141], and Gu et al. [142], for the reduction of dyes, such as crystal violet and methyl red, bromothymol blue, and methylene blue or for the production of 3,4-dihydropyrimidinones by Biginelli reaction [141].

Zinc oxide nanoparticles (ZnONP) were phytosynthesized by Suresh et al. [143] using *Artocarpus gomezianus* fruit extract. The authors obtained porous nanoparticles, with dimensions dependent on the concentration of the extract used for the phytosynthesis. The nanoparticles exhibited good catalytic properties for the degradation of methylene blue, especially in alkaline pH, both under direct sunlight and UV light. The degradation of the same dye was studied by Anbuvannan et al. [144] using ZnONP phytosynthesized by aqueous *Phyllanthus niruri* leaves extract, obtaining an almost complete disappearance of the specific UV-Vis peak at 664 nm in under 30 min Patel et al. [145] obtained ZnO nanorods using *Aloe vera* gel and established its catalytic superiority for the thermal decomposition of potassium perchlorate, compared with nanorods of ZnO obtained using a commercial surfactant (poly(ethylene)glycol).

The same type of nanoparticles, obtained using different plant extracts were successfully applied by Raja et al. [146], Ishwarya et al. [147], and Ali et al. [148] for the photodegradation of dyes (methylene blue and methyl orange), all the authors observing an elevated degradation efficiency, especially towards methylene blue (over 90% degradation). Phukan et al. [149] obtained ZnO nanotapes, with widths within 9 nm using *Lantana camara* flowers extract and applied the nanostructures for ipso-hydroxylation of different aryl/heteroarylboronic acid to phenol. Khan et al. [150] obtained spherical ZnONP using *Trianthema portulacastrum* plant extract and applied them for the degradation of a textile dye (Synozol Navy Blue-KBF), obtaining a 91% degradation of the dye in 159 min. The author also proposed a
mechanistic pathway for the dye degradation, involving the physical adsorption of the dye on the nanoparticles surface, followed by the photocatalytic degradation under the action of OH radicals.

Tin oxide nanoparticles (SnO$_2$NP) were obtained by Haritha et al. [151] using *Catunaregam spinosa* bark extract. The nanoparticles were applied for the degradation of the Congo red dye, obtaining a 92% degradation after 45 min of reaction. The authors assign the main role in the phytosynthesis to the hydroxyl group containing phytomolecules. Begum and Ahmaruzzaman [152] used for the phytosynthesis of SnO$_2$NP using an aqueous extract of the pods of a tropical leguminous tree (*Parkia speciosa* Hassk). The obtained structures were applied for the degradation of the acid yellow 23 dye, reaching a 98% degradation under UV-C irradiation for 24 min and a reusability of five cycles, without any losses in terms of stability and morphology.

Phytosynthesized titania nanoparticles (TiO$_2$NP) were obtained by Thandapani et al. [153] and Udayabhanu et al. [154] and applied for the catalytic degradation of dyes (methylene blue, methyl orange, crystal violet, and alizarin red), obtaining degradation efficiencies over 77% after 6 h. In both studies, the best effect (over 92% degradation) was observed for methylene blue, while the lowest degradation efficiencies were observed for alizarin red (77.3%) and methyl orange (77.5%). Considering the comparable morphologies and sized of the NPs (Table 1), it can be concluded that the differences in terms of catalytic activity are due to the different phytochemicals from the used extracts.

Other types of metallic oxides nanoparticles (such as Mn$_3$O$_4$NP, CeO$_2$NP, ZrONP, or NiONP) were phytosynthesized by different groups [155–158] and used for different catalytic applications (catalytic thermal decomposition of ammonium perchlorate; degradation of crystal violet; degradation of methyl orange; photocatalytic degradation of 4-clorphenol).

Although not as common as the metallic oxides, phytosynthesized metallic chlorides are also encountered in the literature. For example, Huo et al. [159] obtained, using aqueous root extract of *Glycyrrhiza uralensis*, silver chloride nanoparticles (AgClNP). The obtained nanoparticles were successfully applied for the degradation of methylene blue, as a model test pollutant.

As many catalytic processes are related to the metal surface, NPs are usually more reactive than the bulk metal, a consequence of their smaller sizes and larger surface areas. Comparing metallic oxides nanoparticles, iron oxide magnetic nanoparticles could be considered a more desirable alternative for real-life applications, as they are easier to recover from the system, by applying an exterior magnetic field.

4. Complex Catalytic Structures Based on Phytosynthesized Nanoparticles

Besides the nanoparticles described in the previous chapters, the literature also presents complex structures, incorporating phytosynthesized nanomaterials, starting from the simple M$^1$/M$^2_x$N$_y$ catalysts (where M$^1$ and M$^2$ are metals, N = O, Cl) to composites such as membrane or zeolite-metallic nanoparticles catalysts. In the following paragraphs, we will present a short overview of the recent advances registered in this area.

Nasrollahzadeh et al. [160] used aqueous cocoa seeds extract for the phytosynthesis of Pd/CuO nanoparticles with average size of 40 nm, assigning the main role in the phytosynthesis process to different compounds (epicatechin, catechin, flavonoid and phenolic acids) present in the extract. The nanoparticles were applied for the reduction of 4-nitrophenol with NaBH$_4$, obtaining a complete disappearance of the specific absorption peak at 400 nm in 60 seconds, with a reusability of over six cycles. The nanoparticles were also proven to be a highly efficient and stable heterogeneous catalyst for phosphine-free Heck coupling reaction. Using aqueous *Silybum marianum* L. seeds extract, Sajadi et al. [161] obtained magnetic Cu/Fe$_3$O$_4$NP used as catalysts for the reduction of various nitroarenes (with yields of 88-97% and a recyclability of at least five cycles). Nasrollahzadeh and Sajadi [162] applied phytosynthesized Pd/TiO$_2$NP as a catalyst for the ligand-free Suzuki–Miyaura coupling reaction (registering yields up to 98% and a four cycles reusability without a significant yield decrease). Reduction of different compounds (4-nitrophenol, methyl orange, Congo red, and methylene
blue) was achieved by Atarod et al. [163] and Momeni et al. [164] in 2016, using phytosynthesized Ag/TiO$_2$NP and Cu/ZnONP, with very low reaction time.

Sajadi et al. [165] applied phytosynthesized Ag/Fe$_3$O$_4$NP for the [2+3] cycloaddition of arylcyanamides and sodium azide, while Hag et al. [166] used Ag/NiONP for the catalytic degradation of rhodamine B.

Ag/AgCl nanoparticles were obtained by Devi et al. [167,168] using Momordica charantia and Benincasa hispida extracts. The authors applied the nanoparticles for the reduction of 2-, 4-dinitrophenyl hydrazine for the degradation of malachite green oxalate under sunlight irradiation.

Other types of phytosynthesized catalytic material are represented by the metal oxide/metal oxide nanostructures. For example, Bharati and Suresh [169] proposed the application of phytosynthesized ZnO/SiO$_2$ nanostructures for the reduction of acenaphthylene from refinery waste water, while Maria Magdalane et al. [170] used CeO$_2$/CdO multilayered nanoplatelet for the degradation malachite green and catalytic hydrogenation of 4-nitrophenol.

Using aqueous Opuntia dilenii haw, Azadirachta indica, Ocimum sanctum, and Saraca indica extracts, Kombaiah et al. [171] and Garg et al. [172] obtained nano-ferrite (ZnFe$_2$O$_4$) and bismuth oxychloride (BiOCl), and demonstrated their potential for the oxidation of glycerol into formic acid for the degradation of methyl orange and bisphenol A.

Another widely encountered strategy is represented by the incorporation of phytosynthesized in porous materials (especially zeolites). For example, Zhan et al. [173] reported the obtaining of a AuNP/titanium silicalite-1 composite for vapor phase propylene epoxidation, Nasrollahzadeh et al. [174] reported the N-formylation of amines using CuNP/zeolite (Natrolite) as a “green” catalyst, and Hatamifard et al. [175] used AgNP/zeolite nanocomposite for the ligand-free hydroxylation of phenylboronic acid to phenol and the reduction of 4-nitrophenol (4-NP), methyl orange (MO), Congo red (CR), methylene blue (MB) and rhodamine B, while Das et al. [176] applied AgNP/zeolite (mesoporous silicate-1) catalyst for the reduction of 4-nitrophenol.

Catalytic composites constructed with reduced graphene oxides (RGO) and phytosynthesized nanoparticles were evaluated by Nasrollahzadeh et al. (PdNP/RGO) [177], Atarod et al. (Pd/RGO/Fe$_3$O$_4$) [178], Maham et al. (Ag/RGO/Fe$_3$O$_4$) [179], Nasrollahzadeh et al. (Cu/RGO/Fe$_3$O$_4$) [180], Khan et al. (Pd/RGO) [181], and Anasdass et al. [182] for the reduction of nitroarenes, reduction of 4-nitrophenol, reduction of 4-nitrophenol, Congo red and Rhodamine B, direct cyanation of aldehydes with K$_4$[Fe(CN)$_6$]$_4$, as catalyst for the Suzuki-Miyaura coupling, and as catalyst for Suzuki cross-coupling reactions.

Smuleac et al. [183] incorporated mono- and bi-metallic phytosynthesized nanoparticles (Fe and Fe/Pd) in a polycrylic acid-coated polyvinylidene fluoride membrane and applied them for the degradation of trichloroethylene, observing the preservation of the membrane reactivity after 4 months of use. Momeni et al. [184] applied Ag/bone nanocomposites for the hydration of cyanamides and Goswami et al. [185] used AgNP supported on cellulose for the degradation of methylene blue, methyl orange, bromphenol blue, Eosin Y and Orange G, while Naemi et al. [186] phytosynthesized MoO$_3$ nanoparticles and used them for the construction of MoO$_3$/Copper Schiff base complex with application in the oxidation of alcohols.

Another interesting approach, presented by several authors, is represented by the deposition of the phytosynthesized nanoparticles on different supports (by-products or wastes from different industries). Rostami-Vartooni [187] obtained CuONP deposited on seashell surface and applied the composites for the reduction of 4-nitrophenol and Congo red in the presence of NaBH$_4$, while Bordbar and Mortazavimanesh [188] applied PdNP/walnut shell composites for the reduction of 4-nitrophenol, Congo red, methylene blue, and rhodamine B, with reaction times under 1 min Khodadadi et al. [189] deposited PdNP on apricot kernel shells and used them as catalysts for the reduction of 4-nitrophenol, methyl orange, methylene blue, rhodamine B, and Congo red at room temperature (reaction completed in seconds); the same group [190] applied the developed strategy for obtaining AgNP/peach kernel
shells, for the reduction of 4-nitrophenol, methyl orange, and methylene blue (reaction times under 15 min).

The general procedure for developing the presented phytosynthesized nanocatalysts (as concluded from the reviewed articles) and the synthesis mechanisms are presented in Figure 2.

**Figure 2.** General procedure and phytosynthesis mechanisms for obtaining the “green” nanocatalysts (adapted from [28,32,127] and [167] and for the characterization steps (Transmission electron microscopy-TEM images of silver-AgNP, gold-AuNP and gold/silver nanoparticles-Ag/AuNP from the authors’ unpublished results).

Table 1 presents the main applications of phytosynthesized nanoparticles in the field of catalysis, including the extracts used for phytosynthesis, applied characterization techniques, and NPs characteristics, for quick reference. The schematic representation of some of the catalytic applications of nanoparticles identified in the present review is presented in Figure 3.
Table 1. Examples of the phytosynthesized nanoparticles, applied characterization techniques and their catalytic applications.

| NPs     | Plant Extract Used                  | NPs Characteristics            | Applied Characterization Techniques | Catalytic Activity                                                                 | Ref.   |
|---------|-------------------------------------|---------------------------------|-------------------------------------|-------------------------------------------------------------------------------------|--------|
| Ag, Au  | Breynia rhamnoides                  | Ag–spherical, 64 nm Au–spherical, 27 nm | UV-Vis, TEM, FTIR, particle size distribution | Catalytic conversion of 4-nitrophenol                                               | [33]   |
| Ag      | Aqueous Saraca indica flowers extract | Spherical, 18–22 nm              | UV-Vis, TEM, XRD, FTIR              | Reduction of Methylene blue                                                          | [34]   |
| Ag      | Aqueous Alstonia macrophylla leaf extract | Spherical, 70 nm                | UV-Vis, FTIR, XRD, SEM, particle size distribution | Reduction of 4-nitrophenol and p-nitroaniline                                         | [35]   |
| Ag      | Aqueous Trigonella foenum-graecum seeds extract | Spherical and quasi-spherical, 22–32 nm | UV-Vis, XRD, TEM, FTIR              | Catalytic degradation of methyl orange, methylene blue and eosin Y                          | [36]   |
| Ag      | Aqueous Euphorbia cordifolia M. bieb root extract | Spherical, diameters not determined | UV-Vis, FTIR, TEM                    | Catalytic synthesis of N-monosubstituted ureas in water                               | [37]   |
| Ag      | Coleus forskohlii root extract       | Triangular and spherical, 30–40 nm and 35–55 nm, depending on the extract quantity | UV-Vis, XRD, TEM, FTIR              | Catalytic reduction of 4-nitrophenol                                                | [38]   |
| Ag      | Aqueous Salicornia brachiata extract | Spherical, rod-like, prism, triangular, pentagonal and hexagonal, 30–40 nm | UV-Vis, SEM, TEM                    | Catalytic reduction of 4-nitrophenol                                                | [39]   |
| Ag      | Aqueous Leucas martinicensis leaves extract | Nearly spherical 20–30 nm         | UV-Vis, XRD, TEM, FTIR              | Reduction of methylene blue                                                          | [40]   |
| Ag      | Aqueous Tribulus terrestris extract  | Spherical, 15–40 nm              | UV-Vis, XRD, TEM, FTIR              | Reduction of methylene blue                                                          | [41]   |
| Ag      | Aqueous beetroot extract             | Spherical, 15 nm                 | UV-Vis, XRD, TEM                    | Reduction of 4-nitrophenol to 4-aminophenol                                          | [42]   |
| Ag, Au  | Aqueous Aerva lanata leaves extract  | Ag-Quasi-spherical, 18.62 nm Au–different morphologies, 17.97 nm | UV-Vis, FTIR, XRD, TEM              | Reduction of 4-nitrophenol to 4-aminophenol                                          | [43]   |
| Ag      | Aqueous Ziziphus jujuba leaves extract | Different shapes, 20–30 nm, hydrodynamic size 28 nm | UV-Vis, XRD, FT-IR, TEM, DLS, Zeta potential | Reduction of 4-nitrophenol and methylene blue                                        | [44]   |
| Ag      | Aqueous Monordica charantia leaves extract | Spherical, 16 nm                | XRD, SAED, SEM, XPS, UV-Vis, FTIR   | Reduction of methylene blue                                                          | [45]   |
| Ag      | Aqueous Lantana camara leaves extract | Almost spherical, 23–30 nm, varying with silver precursor concentration | XRD, SEM, TEM, UV-Vis, FTIR, Zeta potential | Reduction of methylene blue                                                          | [46]   |
| Ag      | Ferredoxin–NADP+ reductase/ferredoxin obtained from spinach extract | Spherical, 10–15 nm             | UV-Vis, TEM, FTIR                    | Degradation of methylene blue, methyl orange and methyl red                         | [47]   |
| Ag      | Aqueous extracts of Mentha arvensis var. piperascens, Buddleja officinalis Maximowicz, Epimedium koreanum Nakai, Artemisia messer-schmidtiana Besser, Magnolia kobus | Average particle sizes 24.7 to 40 nm | EDS, SEM, EDX, XPS, TEM              | Degradation of cellulose by cellulase                                                | [48]   |
| NPs     | Plant Extract Used                  | NPs Characteristics            | Applied Characterization Techniques | Catalytic Activity                                | Ref.     |
|---------|-------------------------------------|---------------------------------|-------------------------------------|---------------------------------------------------|----------|
| Ag      | Aqueous extract from dried Euphorbia helioscopia Linn leaves | Spherical, 2–14 nm              | TEM, XRD, FT-IR, UV-Vis              | Synthesis of propargylamines                      | [49]     |
| Ag      | Aqueous bark extract of Terminalia cuneata | Distorted spherical shape, 25–50 nm | FTIR, XRD, DLS, TEM, EDX            | Reduction of direct yellow-12 dye                | [51]     |
| Ag      | Aqueous extract of Erigeron bonariensis | Spherical, average size 13 nm   | SEM, TEM, XRD, SAED, AFM, FTIR, UV-Vis | Degradation of Acridine Orange                   | [52]     |
| Ag      | Aqueous extract of Anacardium occidentale testa | Distorted spherical shape, 25 nm | UV-Vis, FTIR, XRD, TEM              | Degradation of Congo red and methyl orange       | [53]     |
| Ag      | Aqueous extract of Crotonaria retus leaves | Spherical, 80 nm               | UV-Vis, TEM, DLS                    | Reduction of cresyl blue                         | [54]     |
| Ag      | Aqueous extract of Lychee (Litchi Chinensis) fruit peel | Spherical, 4–8 nm              | UV-Vis, XRD, EDX, SAED, TEM, FTIR   | Photocatalytic degradation of methylene blue      | [55]     |
| Ag      | Aqueous extract of Galera senegalensis | Spherical, 50 nm               | FTIR, TEM, SEM, XRD, EDX            | Degradation of 4-nitrophenol and Congo red       | [56]     |
| Ag, Au  | Aqueous extract of Cicer arietinum leaves | Spherical, 88.8 nm            | UV-Vis, SEM, TEM, DLS, EDX, FTIR,  | Degradation of 4-nitrophenol, methylene blue and Congo red | [57]     |
| Ag, Au  | Aqueous Aglaia elaeagnoides flower extract | Ag-spherical, 17 nm, Au-spherical, 25 nm | UV-Vis, FTIR, XRD, SEM, EDX, TEM | Degradation of 4-nitrophenol, methylene blue and Congo red | [58]     |
| Ag      | Aqueous Phoenix Dactylifera extract | Spherical, 25–60 nm           | UV-Vis, XRD, FTIR, SEM, TEM, AFM, EDX, Zeta potential | Degradation of 4-nitrophenol | [59]     |
| Ag, Au  | Aqueous leaf extract of Musaenda glabrata | Ag-spherical, 51.32 nm, Au-triangular and spherical, 10.59 nm | UV-Vis, FTIR, XRD, EDX, SAED, TEM, AFM | Degradation of 4-nitrophenol, rhodamine B and methyl orange | [60]     |
| Ag      | Aqueous leaf extract of Cascabela thevetia and Wrightia tomentosa | Not determined | UV-Vis | Degradation of 4-nitrophenol, methylene blue, methyl orange and methyl red | [61]     |
| Ag      | Fractionated Cassia auriculata L. flower aqueous extract | Spherical and triangular, 10–35 nm | UV-Vis, FTIR, XRD, TEM              | Degradation of 4-nitrophenol, methyl orange      | [62]     |
| Ag, Au, Ag/Au | Aqueous bark Guazuma ulmifolia L. extract | Ag-spherical, 10–15 nm, Au-spherical, 20–25 nm, Ag/Au-spherical, 10–20 nm | UV-Vis, FTIR, XRD, AFM, TEM | Degradation of 4-nitrophenol, Congo red | [63]     |
| Ag      | 90% ethanol extract of Dillenia indica bark | Spherical, 15–35 nm           | UV-Vis, XRD, TEM, FTIR              | Degradation of 4-nitrophenol, methylene blue     | [64]     |
| Ag      | Terminalia bellerica fruit aqueous extract | Spherical, ≤20.6 nm           | UV-Vis, FTIR, Zetasizer, SEM, EDX, XRD | Degradation of 4-nitrophenol                     | [65]     |

Table 1. Cont.
Table 1. Cont.

| NPs        | Plant Extract Used                  | NPs Characteristics                  | Applied Characterization Techniques | Catalytic Activity                                                                 | Ref. |
|------------|-------------------------------------|--------------------------------------|-------------------------------------|-------------------------------------------------------------------------------------|------|
| Ag, Au     | *Actinidia deliciosa* fruit extract | Ag–spherical, 25–40 nm Au–spherical, 7–20 nm | EDAX, XPS, XRD, FTIR               | Degradation of 4-nitrophenol, methylene blue                                        | [66] |
| Ag         | Aqueous extract of Camellia japonica leaves | Spherical, 12–25 nm                  | UV-Vis, XRD, FTIR, TEM, EDX        | Photocatalytic degradation of eosin-Y                                               | [67] |
| Ag         | Aqueous extract of Caulerpa serrulate algae | Spherical, 10 ± 2 nm                  | UV-Vis, FT-IR, XRD, TEM            | Degradation of Congo red                                                           | [68] |
| Ag         | *Artemisia tournefortiana* Rchb ethanol extract | Spherical, average diameter 22.89 ± 14.82 nm | UV-Vis, TEM, SEM, EDX, XRD, FTIR   | Degradation of Coomassie Brilliant Blue G-250 under UV light                      | [69] |
| Ag         | Aqueous bark extract of *Acanthopanax sessiliflorus* | Roughly-spherical, average diameter ~20 nm | UV-Vis, XRD, TEM                   | Reduction of p-nitroaniline to p-phenylenediamine                                  | [70] |
| Ag         | Aqueous extract of *Indigofera tinctoria* | Spherical, 15–45 nm                  | UV-Vis, FTIR, XRD, TEM, EDX, AMF   | Synthesis of N-monosubstituted ureas via the hydration of cyanamides in aqueous medium | [71] |
| Ag         | Aqueous extract of *Radix Puerariae* | Spherical, 10–35 nm                  | TEM, XRD, SEM, DLS, EDX, UV-Vis    | Construction of pyrimido[1,2-b]indazole derivatives under solvent-free conditions | [72] |
| Ag         | Aqueous extract of Aqueous palm date fruit pericarp extract | Spherical, 3–30 nm                  | SEM, TEM, SAED, EDX, DLS          | Catalytic degradation of 4-Nitrophenol                                             | [73] |
| Ag         | *Prosopis juliflora* bark extract     | Spherical, 10–50 nm                  | UV-Vis, DLS, SEM, FTIR            | Catalytic degradation of 4-Nitrophenol                                             | [74] |
| Ag, Au     | Aqueous leaf extract of *Indigofera tinctoria* | Ag–spherical, 9–26 nm Au–spherical, triangular, hexagonal, 6–29 nm | UV-Vis, FTIR, XRD, TEM, EDX, AMF   | Reduction of ortho-nitroanilines                                                   | [75] |
| Ag         | Leaf extract of Orthosiphon aristatus | Spherical, 15–45 nm                  | UV-Vis, FTIR, XRD, TEM, EDX        | Reduction of 4-nitrophenol and o-p-nitroaniline                                     | [76] |
| Ag         | Aqueous extract of *Elephantopus scaber* | Spherical, 20–60 nm                  | UV-Vis, FTIR, XRD, TEM, EDX, AMF   | Reduction of 4-nitrophenol, 2,4-nitroaniline and eosin Y                           | [77] |
| Ag, Au     | Aqueous extract of Stemona tuberosa Lour | Ag–spherical, 25 nm Au–irregular, 20–30 nm | UV-Vis, EDX, XRD, FTIR, SEM, TEM, SAED, Zeta potential | Degradation of 4-nitrophenol, methylene blue, methyl orange and methyl red | [78] |
| Ag, Au     | Aqueous leaves extract of *Myxopyrum serratum* A. W. Hill | Ag–spherical, 20–50 nm Au–irregular, 6–29 nm | UV-Vis, FTIR, XRD, TEM, EDX        | Reduction of 4-nitrophenol, methylene blue and Congo red                           | [79] |
| Ag         | Aqueous extract of Longan fruit peel | Spherical, 20 nm                     | UV-Vis, XRD, EDX, TEM, FTIR        | Degradation of methylene blue                                                      | [80] |
| Ag         | Aqueous extract of *Durio Zibethinus* seeds | Spherical, rod shaped, 20–75 nm      | UV-Vis, SEM, TEM, Zeta potential, XRD, EDX | Degradation of methylene blue                                                     | [81] |
| Ag         | Aqueous extract of *Consolida arvensis* leaves extract | Spherical, 10–30 nm                  | UV-Vis, SEM, TEM, EDX, XRD, DLS, Zeta potential | Reduction of Reactive Black 5, methyl orange, direct yellow-142                 | [82] |
Table 1. Cont.

| NPs | Plant Extract Used | NPs Characteristics | Applied Characterization Techniques | Catalytic Activity | Ref. |
|-----|--------------------|---------------------|-------------------------------------|--------------------|------|
| Ag  | Aqueous seed extract of *Gardenia jasminoides* Ellis | Spherical, average size 20 nm | UV-Vis, TEM, XRD, EDX, particle size analyses, FTIR | Reduction of Coomassie Brilliant Blue | [83] |
| Ag  | Aqueous extract of *Allium cepa* | Spherical, 50–100 nm | UV-Vis, SEM, TEM, Particle size, Zeta potential, EDX, XRD, FTIR | Degradation of methylene blue, eosin yellowish, safranin, direct dye, reactive dyes | [84] |
| Ag  | Aqueous *Nervilia zeylanica* leaves extract | Spherical, average particle size 34.2 nm | UV-Vis, FTIR, XRD, AFM, EDX, TEM | Degradation of methyl orange, rhodamine B | [85] |
| Ag  | Aqueous extract of *Bauhinia tomentosa* Linn. leaves | Spherical, 8–25 nm | UV-Vis, XRD, FTIR, SEM, EDX, TEM | Degradation of nitrophenol, 3-nitrophenol, 2, 4, 6-trinitrophenol, picric acid, Coomassie Brilliant Blue, Congo red, eosin Y, rhodamine B, methylene blue, methyl red, methyl orange, cresol red, acridine orange, eriochrome black T, and phenol red | [86] |
| Ag  | Aqueous rhizome extract of *Acorus calamus* | Spherical, hydrodynamic diameter 31.83 nm | UV-Vis, TEM, EDX, FTIR, zeta potential, DLS | Degradation of Reactive Red 120 and Reactive Black 5 | [87] |
| Ag  | Aqueous *Ginkgo biloba* leaves extract | Spherical, 20–40 nm | UV-Vis, XRD, XPS, SEM, TEM, DLS | Reduction of 4-nitrophenol, Congo red, methyl orange, and rhodamine B | [88] |
| Ag, Au | Aqueous leaf extract of *Bauhinia purpurea* | Ag– quasi-spherical Au– spherical, hexagonal, triangular, nanorods | UV-Vis, FTIR, XRD, TEM, EDX | Degradation of methylene blue and rhodamine B | [89] |
| Ag  | Aqueous *Eriobotrya japonica* (Thunb.) leaves extract | Spherical, 9.26 ± 2.72, 13.09 ± 3.66, and 17.28 ± 5.78 nm, temperature dependent | UV-Vis, SEM, XRD, TEM, FTIR, EDX, SAED | Degradation of Reactive Red 120 and Reactive Black 5 | [90] |
| Au  | Aqueous extract of green tea leaves | Mostly spherical, 20 nm | UV-Vis, TEM, XRD | Degradation of methylene blue | [91] |
| Au  | Aqueous extract of *Gnidia glauca* flower | Varying morphology, mostly spherical, 10 nm, with particles up to 150 nm | UV-Vis, TEM, DLS, XRD, FTIR | Degradation of 4-nitrophenol | [92] |
| Au  | 70% ethanol *Phoenix dactylifera* L. leaves extract | Spherical, 32–45 nm | UV-Vis, TEM, FTIR, AAS | Degradation of 4-nitrophenol | [93] |
| Au  | Aqueous *Salticornia brachiata* extract | Spherical, 22–35 nm | UV-Vis, SEM, EDX, TEM, XRD | Degradation of 4-nitrophenol and methylene blue | [94] |
| Au  | Aqueous leaf extract of *Polygonum minus* | Mostly icosahedral, 23 nm | UV-Vis, XRD, FTIR, TEM, EDX | Degradation of 4-nitrophenol | [95] |
| Au  | Aqueous *Gnidia glauca* leaves and stem extracts | Mostly spherical, 10–60 nm | UV-Vis, TEM, DLS, XRD, FTIR | Degradation of 4-nitrophenol | [96] |
| NPs   | Plant Extract Used                              | NPs Characteristics                      | Applied Characterization Techniques                      | Catalytic Activity                                                                 | Ref.   |
|-------|------------------------------------------------|------------------------------------------|---------------------------------------------------------|------------------------------------------------------------------------------------|--------|
| Au    | Aqueous *Artemisia capillaris* extracts, different conditions | Spherical, 16.88 ± 5.47–29.93 ± 9.80 nm | UV-Vis, TEM, particle dimension, zeta potential, XRD, FTIR | Degradation of 4-nitrophenol                                                        | [97]   |
| Au    | Aqueous *Delonix regia* leaves extract          | Spherical, 4–24 nm                       | UV-Vis, TEM, XRD, EDX, SEM, FTIR                        | Degradation of o-nitroaniline                                                       | [98]   |
| Au    | Aqueous *Lagerstroemia speciosa* leaves extract | Triangular and hexagonal, some spherical, 41–91 nm | TEM, UV-Vis, XRD, FTIR                               | Reduction of methylene blue, methyl orange, bromophenol blue, bromocresol green, and 4-nitrophenol | [99]   |
| Au    | Aqueous *Cotoneaster horizontalis* leaves       | Spherical, 18 ± 2 nm                     | UV-Vis, XRD, TEM, FTIR, SEM, EDX                      | Degradation of 4-nitrophenol                                                        | [100]  |
| Au    | Aqueous *Sueda fructicosa* extract              | Spherical, 2–12 nm                       | UV-Vis, FTIR, XRD, TEM                                | Degradation of methylene blue                                                       | [101]  |
| Au    | Ethanol *Artemisia dracunculus* extract         | Spherical, hexagonal, and triangular shapes, depending on phytosynthesis conditions, wide variety of dimensions | UV-Vis, SEM, DLS, zeta potential, FTIR                   | Reduction of 4-nitrophenol                                                          | [102]  |
| Pd    | Aqueous *Pulicaria glutinosa* extract           | Spherical, 20–25 nm                      | UV-Vis, XRD, TEM, EDX, FTIR                           | Catalytic conversion of bromobenzene to biphenyl                                    | [106]  |
| Pd    | Aqueous *Stachys lavandulifolia* extract        | Nearly spherical, 5–7 nm                 | UV-Vis, TEM, XRD, FTIR, SEM, EDX, ICP                 | Suzuki reaction, for a wide range of aryl halides                                  | [107]  |
| Pd    | Aqueous extract of leaves of *Euphorbia thymifolia* L. | Spherical, 20–30 nm                      | XRD, TEM, FTIR, UV-vis                                | Catalyst for Stille and Hiyama cross-coupling reactions                            | [108]  |
| Pd    | Aqueous extract *Eucommia ulmoides* bark        | Spherical and quasi-spherical, 10–20 nm   | TEM, EDX, XRD,                                         | Electro-catalytic oxidation of hydrazine and catalytic degradation of p-aminoazobenzene | [109]  |
| Pd    | Aqueous *Euphorbia granulate* leaves extract    | 25–35 nm                                | FTIR, TEM                                             | Catalyst for the Suzuki-Miyaura coupling reaction                                  | [110]  |
| Pd    | Aqueous *Sapium sebiferum* leaves extract       | Spherical, 2–14 nm, depending on the synthesis temperature | UV-Vis, XRD, TEM, SAED, TGA, DLS, FTIR               | Photodegradation of methylene blue                                                  | [111]  |
| Pd    | Aqueous *Garcinia pedunculata* Roxb and starch  | Spherical and non-spherical, 2–4 nm      | FTIR, XRD, TEM, XPS, SEM, EDX, UV-Vis, TGA, DLS       | Catalyst for Suzuki-Miyaura cross-coupling reaction, alcohol oxidation and Cr(VI) reduction | [112]  |
| Pd    | Aqueous *Santalum album* extract                | Spherical, 10–40 nm                      | UV-Vis, TEM, XRD, FTIR                               | Catalytic reduction of 4-nitrophenol                                                 | [113]  |
| Pd    | Aqueous *Camellia sinensis* leaves extract      | Near spherical, 5–8 nm                   | UV-Vis, XRD, FTIR, SEM, TEM, EDX                      | Catalytic reduction of 4-nitrophenol, heterogeneous catalyst for Suzuki coupling reactions | [114]  |
| Pd    | Aqueous *Pimpinella Tirupatiensis* leaves extract | Spherical, 12.25 nm                     | UV-Vis, XRD, FTIR, TEM                                | Catalytic degradation of Congo red                                                   | [115]  |
| NPs | Plant Extract Used                          | NPs Characteristics | Applied Characterization Techniques | Catalytic Activity                                                                 | Ref. |
|-----|--------------------------------------------|---------------------|-------------------------------------|------------------------------------------------------------------------------------|------|
| Pd  | Aqueous *Origanum vulgare* L. extract      | Spherical, 2–20 nm   | UV-Vis, FTIR, XRD, TEM, EDX, TGA     | Selective oxidation of alcohols                                                    | [116]|
| Pd  | Aqueous *Lagerstroemia speciosa* leaves extract | 136.5 nm            | UV-Vis, SEM, EDX, XRD, FTIR, TGA     | Reduction of methylene blue, methyl orange and 4-nitrophenol                      | [117]|
| Fe  | Aqueous sorghum bran                        | Spherical, 40–50 nm  | UV-Vis, SEM, TEM, XRD, zeta potential | Degradation of Bromothymol Blue                                                   | [118]|
| Fe  | Black tea, grape marc, and vine leaves aqueous extracts | 15–45 nm            | UV-Vis, TEM                          | Degradation of ibuprofen                                                          | [119]|
| Fe  | Aqueous green tea extract                   | Spherical, 84.7 ± 11.5 nm and 141.2 ± 26.3 nm, depending on the synthesis atmosphere | SEM, XPS, FTIR                      | Degradation of methylene blue                                                    | [120]|
| Fe  | Aqueous *Lagerstroemia speciosa* leaves extract | Spheroidal, 50–100 nm | UV-Vis, FTIR, EDX, SEM, XRD, TGA     | Degradation of methylene blue, methyl orange, allura red, brilliant blue, green S dyes | [121]|
| Fe  | Aqueous extract of *Trigonella foenum-graecum* seed | 7–14 nm              | UV-Vis, XRD, FTIR, TGA/DTG, TEM, magnetization, | Degradation of methyl orange                                                      | [122]|
| Fe  | *Hibiscus Sabdariffa* flower aqueous extract | Spherical, 18–44 nm, depending on the iron salt concentration | UV-Vis, TEM, FTIR                  | Degradation of rhodamine B                                                       | [123]|
| Cu  | Aqueous *Ginkgo biloba* L. extract          | Spherical, 15–20 nm  | TEM, EDX, FTIR, UV-Vis               | Huisgen [3 + 2] cycloaddition of azides and alkynes                              | [124]|
| Cu  | Aqueous broccoli extract                    | Spherical, average particle size 4.8 nm | UV-Vis, FTIR, TEM, DLS, XRD, cyclic voltammetry | Reduction of 4-nitrophenol, degradation of methylene blue and methyl red        | [125]|
| Cu  | Aqueous *Punica granatum* seeds extract     | Spherical, 40–80 nm, average size 43.9 | UV-Vis, XRD, SEM, EDX, FTIR, AFM     | Degradation of methylene blue                                                    | [126]|
| Cu  | Aqueous *Plantago asiatica* leaf extract    | Spherical, 7–35 nm   | FTIR, UV-Vis, TEM, XRD               | Cyanation of aldehydes usingK₄Fe(CN)₆                                          | [127]|
| Ag/Au| Aqueous *Silphium marianum* extract         | Spherical, ≤40 nm    | UV-Vis, DLS, XRD, TEM, SAED, EDX, XPS, FTIR | Reduction of 4-nitrophenol                                                      | [128]|
| Fe/Pd| Grape leaf aqueous extract                  | Quasi-spherical, 10–100 nm | SEM, FTIR                           | Removal of Orange II                                                             | [129]|
| Ag/Fe| Aqueous Palm dates fruit extract            | Disks, irregular, 5–40 nm | UV-Vis, TEM, EDX                     | Degradation of bromothymol blue                                                  | [130]|
| Ag/Fe| *Cupressus sempervirens* extract            | Core-shell, 22–72 nm, mean particle size 36.7 nm | UV-Vis, TEM, XRD, FTIR              | Degradation of methyl orange                                                    | [131]|
| Fe₂O₄| Aqueous *Artemisia annua* leaves extract     | Spherical, 3–10 nm   | UV-Vis, XRD, TEM, EDX, FTIR, TGA vibrating sample magnetometry | Catalytic synthesis of benzoxazinone and benzthiocyanazinone derivatives       | [135]|

Table 1. Cont.
| NPs  | Plant Extract Used                          | NPs Characteristics         | Applied Characterization Techniques                                      | Catalytic Activity                                                                 | Ref.  |
|------|--------------------------------------------|-----------------------------|---------------------------------------------------------------------------|----------------------------------------------------------------------------------|-------|
| Fe₃O₄ | Aqueous *Ruellia tuberosa* leaves extract   | Hexagonal nanorods, 20–80 nm | UV-Vis, FTIR, SEM, EDX, TEM, DSC, DLS                                      | Degradation of crystal violet                                                    | [136] |
| CuO  | Aqueous *Gundelia tournefortii* leaves extract | Spherical, not determined   | UV-Vis, SEM, TEM, EDX, XRD, FTIR                                         | Reduction of 4-nitrophenol and synthesis of N-monosubstituted ureas               | [137] |
| CuO  | Aqueous *Thymus vulgaris* leaves extract    | Quasi-spherical, ≤30 nm      | UV-Vis, TEM, EDX, XRD, FTIR, TGA, DTG                                     | N-arylation of indoles and amines                                                  | [138] |
| CuO, ZnO | Aqueous *Centella asiatica* extract       | Spherical, average diameter 7 nm | UV-Vis, XRD, SEM, TEM, DLS, FTIR                                         | Reduction of methylene blue                                                       | [139] |
| CuO  | Aqueous *Sida acuta* leaves extract        | Nanorods, 50 nm             | UV-Vis, FTIR, SEM, EDX, TEM                                                | Reduction of crystal violet and methyl red                                          | [140] |
| CuO  | Aqueous *Cordia sebestena* flowers extract | Spherical, 20–35 nm         | UV-Vis, FTIR, SEM, EDX, TEM, XRD, DLS, SAED, zeta potential               | Production of 3,4-dihydropyrimidinones by Biginelli reaction, degradation of bromothymol blue | [141] |
| CuO  | Aqueous *Cystoseira trinodis* algal extract | Spherical, 7–9 nm           | UV-Vis, FTIR, SEM, EDX, TEM, AFM, Raman, FTIR                             | Degradation of methylene blue                                                     | [142] |
| ZnO  | Aqueous *Phyllanthus niruri* leaves extract | Spherical, 25–90 nm         | UV-Vis, photoluminescence, XRD, SEM, TEM, FTIR,                             | Degradation of methylene blue                                                     | [144] |
| ZnO  | *Aloe vera* gel                            | Nanorods, 15–20 nm          | XRD, TEM, SAED, SEM, FTIR, DTG                                            | Thermal decomposition of potassium perchlorate                                    | [145] |
| ZnO  | Aqueous *Tabernaemontana divaricata* leaves extract | Spherical, 20–50 nm        | XRD, UV-Vis, TEM, FTIR                                                    | Degradation of methylene blue                                                     | [146] |
| ZnO  | *Ulva lactuca* seaweed extract             | Sponge-like asymmetrical shaped, 10–50 nm | XRD, UV-Vis, FTIR, SAED, TEM                                              | Degradation of methylene blue                                                     | [147] |
| ZnO  | Aqueous *Conyza canadensis* leaves extract | Spherical, ≤10 nm           | UV-Vis, FTIR, XRD, TEM, SEM, EDX                                          | Degradation of methylene blue and methyl orange                                   | [148] |
| ZnO  | Aqueous *Lantana camara* flowers extract   | Nanotapes, widths within 9 nm | UV-Vis, XRD, FTIR, TGA, TEM, specific surface area, photoluminescence     | *p* -hydroxylation of different aryl/heteroarylboronic acid to phenol            | [149] |
| ZnO  | Aqueous *Trianteha portulacastrum* extract | Spherical, 25–90 nm         | UV-Vis, XRD, FTIR, SEM, EDX, TEM, XPS                                      | Degradation of Synozol Navy Blue-KBF                                              | [150] |
| SnO₂ | Aqueous *Catunaregam spinosa* bark extract | Spherical, average size 47 ± 2 nm | UV-Vis, XRD, FTIR, TEM, EDX                                                | Degradation of Congo red                                                         | [151] |
| NPs         | Plant Extract Used                  | NPs Characteristics | Applied Characterization Techniques | Catalytic Activity                                                                 | Ref.  |
|-------------|------------------------------------|---------------------|-------------------------------------|------------------------------------------------------------------------------------|-------|
| SnO₂        | Aqueous Parkia speciosa Hassk pods extract | Spherical, average size 1.9 nm | UV-Vis, XRD, FTIR, TEM, EDX, SAED  | Degradation of acid yellow 23                                                     | [152] |
| TiO₂        | Aqueous Parthenium hysterophorus leaves extracts | Spherical, 20–50 nm | UV-Vis, FTIR, SEM, EDX, XRD        | Degradation of methylene blue, methyl orange, crystal violet, alizarin red        | [153] |
| TiO₂        | Aqueous Euphorbia hirta leaves extract | Spherical, 20–50 nm | UV-Vis, FTIR, XRD, SEM, EDX        | Degradation of methylene blue, methyl orange, crystal violet, alizarin red        | [154] |
| Mn₃O₄NP     | Aqueous Azadirachta indica leaves extract | Spherical, 20–30 nm | XRD, FTIR, XPS, SEM, TEM, specific surface area | Catalytic thermal decomposition of ammonium perchlorate   | [155] |
| CeO₂NP      | Methanolic Moringa oleifera peel extract | Spherical, average size 40 nm | UV-Vis, FTIR, XRD, TEM, SAED      | Degradation of crystal violet                                                     | [156] |
| ZrONP       | Aqueous Lagerstroemia speciosa Leaves extract | Tetragonal, few oval, average particle size 56.8 nm | UV-Vis, FTIR, XRD, SEM, TEM        | Degradation of methyl orange                                                      | [157] |
| NiONP       | Hydroalcoholic Aegle marmelos leaves extract | Spherical, 8–10 nm | UV-Vis, XRD, SEM, TEM              | Degradation of 4-chlorphenol                                                       | [158] |
| AgCINP      | Aqueous root extract of Glycyrrhiza uralensis | Spherical, 5–15 nm | UV-Vis, TEM, SAED, EDX, XRD, DLS, FTIR | Degradation of methylene blue                                                     | [159] |
| Pd/CuO      | Aqueous Theobroma cacao L. seeds extract | 40 nm               | FTIR, EDX, XRD, TEM, UV-Vis       | Reduction of 4-nitrophenol, catalyst for Heck coupling reaction under aerobic conditions | [160] |
| Cu/Fe₃O₄    | Aqueous Stizium marianum L. seeds extract | Spherical, 8.5–60 nm | XRD, TEM, EDX, UV-vis            | Catalytic reduction of nitroarenes                                                 | [161] |
| Pd/TiO₂     | Aqueous Myrtus communis L. extract | Spherical, 17–25    | SEM, TEM, FTIR, UV-Vis, EDX       | Catalyst for ligand-free Suzuki-Miyaura coupling reaction                         | [162] |
| Ag/TiO₂     | Aqueous Euphorbia heterophylla leaves extract | Core-shell, under 24 nm | FTIR, UV-Vis, XRD, SEM            | Reduction of 4-nitrophenol, methyl orange, Congo red, methylene blue              | [163] |
| Cu/ZnO      | Aqueous Euphorbia prolifica leaves extract | Core-shell, spherical CuNP 5–17 nm shell | FTIR, UV-Vis, XRD, SEM, EDX        | Reduction of Congo red, methylene blue                                           | [164] |
| Ag/Fe₃O₄    | Aqueous Euphorbia peplus L. leaves extract | Spherical, 5–10 nm | XRD, SEM, TEM, EDX, FTIR, UV-Vis  | Catalyst for the [2+3] cycloaddition of arylcyanamides and sodium azide          | [165] |
| Ag/NiONP    | Aqueous Azadirachta indica L. leaves extract | Irregular morphologies | XRD, SEM, TGA                   | Degradation of Rhodamine B                                                       | [166] |
| Ag/AgCl     | Aqueous Momordica charantia leaves extract | Spherical, average size 15 nm | UV-Vis, FTIR, SEM, EDX, XRD, TEM, SAED, TEM | Reduction of 2, 4-dinitrophenyl hydrazine                                         | [167] |
| Ag/AgCl     | Aqueous Benincasa hispida peel extract | Spherical, 25–30 nm | UV-Vis, XRD, FTIR, EDX, SEM, XPS  | Degradation of malachite green oxalate                                            | [168] |
Table 1. Cont.

| NPs | Plant Extract Used | NPs Characteristics | Applied Characterization Techniques | Catalytic Activity | Ref. |
|-----|-------------------|---------------------|-------------------------------------|--------------------|------|
| ZnO/SiO₂ | Aqueous *Butea monosperma* flowers extract | Spherical, 3–45 nm | Specific surface area, SEM, EDX, XRD, FTIR | Reduction of acenaphthylene | [169] |
| CeO₂/CdO | Concentrated *Citrus limonum* fruit extract | Nanoplatelets, diameter 10 nm, length 50–100 nm | XRD, FTIR, SEM, TEM, SAED, UV-Vis, TGA-DTG | Degradation of malachite green and catalytic hydrogenation of 4-nitrophenol | [170] |
| ZnFe₂O₄ | Aqueous *Opuntia dillenii haw* extract | Spherical, 23–50 nm for microwave assisted synthesis Spherical, 380–680 nm for classical temperature synthesis | FTIR, UV-Vis, EDX, XRD, Magnetization measurements, DLS | Oxidation of glycerol into formic acid | [171] |
| BiOCl | Aqueous extracts of *Azadirachta indica*, *Ocimum sanctum* and *Saraca indica* leaves | Nanoflower structures, 50–400 nm, depending on the extract | XRD, XPS, SEM, EDX, TEM, UV-Vis, specific surface area | Degradation of methyl orange and bisphenol A | [172] |
| AuNP/titanium silicalite-1 | *Cacumen Platycladi* extract | 1.7 ± 0.3–4.6 ± 0.5 nm | TEM, specific surface area, TGA, UV-Vis | Propylene epoxidation with H₂/O₂ mixture | [173] |
| CuNP/zeolite | Aqueous *Anthenis xylopoda* flowers extract | 28.5 ± 3 nm (NPs) | XRD, SEM, XRF, UV-Vis, FTIR, TEM, EDX | N-formylation of amines | [174] |
| AgNP/zeolite | Aqueous *Euphorbia prolifera* leaves extract | Semi-spherical, 15 nm (NPs) | FTIR, XRD, SEM, EDX, TEM, UV-Vis | Catalyst for ligand-free hydroxylation of phenylboronic acid to phenol, reduction of 4-nitrophenol, methyl orange, Congo red, methylene blue, rhodamine B | [175] |
| AgNP/zeolite | Aqueous extract of *Caromula* fruit | Spherical, 10–15 nm | SEM, TEM, EDX, XRD, FTIR, specific surface area, UV-Vis | Reduction of 4-nitrophenol, | [176] |
| PdNP/RGO | Barberry fruit extract | Spherical, average size 18 nm | UV-vis, XRD, FTIR, SEM, TEM, EDX | Reduction of nitroarenes | [177] |
| Pd/RGO/Fe₃O₄ | Aqueous *Withania coagulans* leaves extract | Spherical, ≤15 nm | FTIR, UV-Vis, XRD, SEM, TEM, EDX, magnetic properties, | Reduction of 4-nitrophenol | [178] |
| Ag/RGO/Fe₃O₄ | Aqueous *Lotus gurcinii* leaves extract | Spherical, 7–20 nm (AgNP) | FTIR, SEM, EDX, XRD, TEM, UV-Vis | Reduction of 4-nitrophenol, Congo red, Rhodamine B | [179] |
| Cu/RGO/Fe₃O₄ | Aqueous *Euphorbia bungei* Boiss leaves extract | 50–80 nm | UV-Vis, FTIR, XRD, SEM, EDX | Direct cyanation of aldehydes with K₄[Fe(CN)₆] | [180] |
| Pd/RGO | Aqueous *Pulicaria glutinosa* extract | Triangular, 15–18 nm, spherical, isotropic at higher Pd concentration, 7–8 nm | UV-Vis, TEM, EDX, XRD, FTIR, Raman | Catalyst for the Suzuki-Miyaura Coupling | [181] |
| Pd/RGO | Aqueous *Ficus carica* fruit extract | Spherical, 0.16 nm | UV-Vis, XRD, FTIR, TEM, Raman | Catalyst for Suzuki cross-coupling reactions | [182] |
| NPs            | Plant Extract Used                  | NPs Characteristics                  | Applied Characterization Techniques | Catalytic Activity                           | Ref.  |
|---------------|------------------------------------|--------------------------------------|-------------------------------------|---------------------------------------------|-------|
| Fe and Fe/Pd/PAA/PVDF membrane | Aqueous green tea extract           | Spherical, 20–30 nm, aggregation 80–100 nm | SEM, EDX, XRD                        | Degradation of trichloroethylene            | [183] |
| Ag/bone       | Aqueous *Myrica gale* L. extract    | Spherical, 5–10 nm                   | XRD, SEM, TEM, EDX, FTIR, UV-Vis    | Hydration of cyanamides                     | [184] |
| AgNP/cellulose| Aqueous seed extract of *Hibiscus* subdariffa | Spherical, 4 nm                      | UV-Vis, XRD, FTIR, TEM, SEM         | Degradation of methylene blue, methyl orange, bromophenol blue, Eosin Y, Orange G. | [185] |
| MoO₃/Copper complex | Aqueous *Sesbania sesban* extract | Cylindrical, 80 × 30 nm               | FTIR, EDX, SEM, TEM, AFM            | Oxidation of alcohols                       | [186] |
| CuONP/seashell| Aqueous *Rumex crispus* seeds extract | Spherical, 8–60 nm                   | FTIR, XRD, SEM, EDX, TEM, UV-Vis    | Reduction of 4-nitrophenol and Congo red    | [187] |
| PdNP/walnut shell | Aqueous *Equisetum arvense* L. leaves extract | Spherical, 5–12 nm                   | FTIR, UV-Vis, photoluminescence, XRD, SEM, EDX, TEM | Reduction of 4-nitrophenol, Congo red, methylene blue, and rhodamine B. | [188] |
| PdNP/apricot kernel shell | 70% ethanol *Salvia hydrangea* extract | Spherical, ≤10 nm                    | FTIR, SEM, UV-Vis, EDX, TEM, XRD    | Reduction of 4-nitrophenol, methyl orange, methylene blue, rhodamine B, and Congo red | [189] |
| AgNP/peach kernel shells | Aqueous *Achillea millefolium* L. extract | Spherical, ≤20 nm                    | FTIR, UV-Vis, XRD, SEM, EDX, TGA-DTG, TEM | Reduction of 4-nitrophenol, methyl orange, and methylene blue | [190] |

1 Where AAS—Atomic absorption spectroscopy; AFM—Atomic force microscopy; DLS—Dynamic light scattering; DSC—Differential scanning calorimetry; DTG—Differential thermogravimetry; EDX—Energy-dispersive X-ray spectroscopy; FTIR-Fourier-transform infrared spectroscopy; ICP-MS—Inductively coupled plasma mass spectrometry; Raman—Raman spectroscopy; SAED—Selected area electron diffraction; SEM—Scanning electron microscope; TEM—transmission electron microscopy; TGA—Thermogravimetric analysis; UV-Vis—Ultraviolet–visible spectroscopy; XPS—X-ray photoelectron spectroscopy; XRD—X-ray diffraction.
5. Conclusions and Perspectives

From all the presented examples, it can be concluded that the application of phytosynthesized nanoparticles represents a promising area of research in catalysis. When dealing with this type of nanoparticles, the authors should pay special attention to the complete characterization of the obtained materials. UV-Vis spectrometry can provide an important insights on the formation and dimensions of the nanoparticles (by the position of the specific absorbance peaks, as is the case, for example, for AuNP, CuNP, or AgNP, or by the disappearance of adsorption peaks specific for the metal salt, as is the case for Pd, for example). However, the formation of the nanoparticles should be confirmed by other techniques, such as XRD or selected area electron diffraction (SAED) (for the study of their crystalline structure, as well as for the calculation of crystallite size). As for catalytic studies, the size and shape of nanoparticles represents an important parameter, morphology studies should be employed in all studies (most appropriate being TEM, allowing both morphological studies and determination of size distribution of the NP). The main disadvantage of other techniques applied for the determination of particle size (such as Dynamic light scattering (DLS)) is that the technique offers the hydrodynamic diameter, not the “real” diameter of the nanoparticles.

The extracts used for the phytosynthesis also play an important role in the final application. Different types of extraction procedures lead to variations in the composition of the extracts, leading to different sizes and morphologies, thus influencing the catalytic application. Multiple extraction techniques (classical temperature extraction, microwave-assisted, ultrasounds assisted, and accelerated solvent extraction, just to name few examples) could and should be used, in order to optimize the phytosynthesis for the desired application. This observation leads to the major bottleneck in the phytosynthesis of the nanoparticles: although various groups proposes as responsible molecules the phenolic compounds, the flavonoids, or other biomolecules, very few articles applies the fractionation of the extracts and compounds separation, in order to fully elucidate the reaction mechanisms and the biomolecules acting as reducing and capping agents.
Considering the catalytic applications, most of the studies demonstrate the application of NP in synthetic aqueous matrices, not in real conditions. However, in order to reach this step, the proposed catalytic structures should be incorporated in water treatment installations, easily to separate from the water sources and to prove their re-usability. In spite the depicted drawbacks, we consider the area of NP application in catalysis as a promising research field, which could provide important technological advances and knowledge. It is important to get a thorough understanding of all of the aspects regarding specific interaction of biomolecules with inorganic materials in order to produce NPs with hierarchical structures. The quest of elucidating the specificity of a biomolecule and its adsorption on the surface of a particle still remain a challenge. Additionally, in the future, the plant extract-based catalyst preparation should focus on using vegetal wastes since they contain the same organic agent like raw materials and therefore the circle of sustainability can be complete.

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