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Nanomaterials in plant tissue culture: the disclosed and undisclosed

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Plant tissue cultures are the core of plant biology, which is important for conservation, mass propagation, genetic manipulation, bioactive compound production and plant improvement. In recent years, the application of nanoparticles (NPs) has successfully led to the elimination of microbial contaminants from explants and demonstrated the positive role of NPs in callus induction, organogenesis, somatic embryogenesis, somaclonal variation, genetic transformation and secondary metabolite production. This review aims to consolidate all of the current achievements made through the integration of nanotechnology into plant tissue culture and highlight the positive attributes of using NPs in plant tissue culture. Both the positive and adverse effects of using NPs in the culture medium are discussed and presented. The toxicity aspects and the safety concerns of exposing plants and the associated environment to NPs are recorded. Finally, future prospects through the involvement of not merely Ag, TiO₂, and ZnO NPs, but more recent innovations such as graphene, carbon nanotubes, SiO₂, quantum dots, and dendrimers are proposed. The undisclosed shadows hanging in the background, including the repercussions of using nanomaterials without proper awareness, as well as dosage-based adverse effects and nanotoxicity aspects, are highlighted. The need for more research in the pursuit of discrete answers to unresolved questions regarding mechanisms is emphasized as the key to real progress in plant nanobiotechnology.

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1. Introduction

Plant tissue culture is directed towards the growth of plant cells or parts of plants on a nutrient medium under a controlled, sterile, simulated environment. It is an important technique for both basic and applied areas of plant biology, such as cytology, embryogenesis, morphogenesis, nutrition, pathology and germplasm conservation, genetic manipulation, large-scale clonal propagation, and the production of pathogen-free plants and useful metabolites.¹ An efficient plant regeneration protocol is mandatory for genetic transformation and mass propagation. The success of *in vitro* plant culture depends on several factors, such as genotype, the physiological status of the donor plants, the type of explants, surface disinfection methods, the culture medium, plant growth regulators, the size of the culture vessels, spectral quality, light intensity, photo-period and temperature.² The composition of the culture medium strongly influences the morphogenetic potential of the explants. The medium generally consists of macro- and micro-nutrients, amino acids, organic supplements, vitamins, carbon sources, plant growth regulators, and solidifying agents. Optimization of the mineral elements in the culture medium

enhances the growth and morphogenesis of explants. It also improves cell proliferation, organogenesis, somatic embryogenesis, shoot quality and the bioactive compound content in cell and organ cultures.

We are living in the 'Nano Age', the era when every aspect of life has a touch of nano, be it the cosmetics we use, the textiles we wear, the appliances we use, the gadgets we employ, the food we eat, or the environment we live in; whether we like it or not, nanomaterials are already in us, on us and around us. Nanotechnology involves the study and manipulation of materials at length scales below 100 nm. The visionary who first predicted the nano era was Richard Feynman. In December of 1959, he gave a talk, "There's Plenty of Room at the Bottom" at an annual meeting of the American Physical Society at Caltech. It was in this historical classic lecture that Feynman laid the conceptual foundations for the field now called nanotechnology; this was when he imagined a day when things could be miniaturized. He urged scientists to start looking down to smaller invisible materials for bigger solutions and answers. Today, as it now stands, there is indeed plenty of room at the bottom; nanotechnology has bombarded all walks of life, and every discipline in science, be it biology, chemistry, physics, engineering or medicine. Nanomaterials, including nanoparticles, nano-coatings, nanofilms, nanosheets, and nanoclusters, have delivered what their bulk components had failed to. Today's nanotechnology harnesses progress in chemistry, physics,

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materials science and biotechnology to create novel materials that have unique properties because their structures are determined on the nanometer scale. Some of these materials have already found their way into consumer products, like sunscreens and stain-resistant paints and textiles. Others are being intensively researched for solutions to humanity's greatest problems – diseases, clean energy, clean water and a cleaner environment. Some of the noteworthy applications of nanomaterials include: cosmetics (*e.g.* sunscreen lotions with radiation absorbing properties); nanocomposite materials and nanotubes as reinforced fillers to improve the mechanical properties of nanocomposites and impart new properties (optical, electronic *etc.*); nanocoatings, where a nanometer thickness of a nanomaterial can be used to improve properties like wear and scratch-resistance, optoelectronic properties, and hydrophobic properties; hard cutting tools using metal nanocomposites such as tungsten carbide, tantalum carbide and titanium carbide, which have improved wear and erosion-resistance, and last longer than their conventional bulk materials; displays, using carbon nanotubes as emission devices for monitors and televisions (FEDs: field-emission displays); lightweight, high-energy density batteries; fuel additives and catalytically efficient materials; nanospheres as lubricants; nanoscale magnetic materials in data storage devices; nanostructured membranes for water purification; and last but not least, inputs into nanoelectronics, nanobiotechnology and nanomedicine.^{3–9}

In plant tissue culture, there are numerous reports that indicate positive inputs from nanotechnology. Nanoparticles (NPs) have been widely used to improve seed germination, enhance plant growth and yield, enable plant genetic modification, improve bioactive compound production and achieve plant protection.^{10,11} On treatment of tomato seeds with silicon dioxide (SiO_2), the NPs enhanced the percentage of seed germination and seedling growth.¹² The application of iron and magnesium nano-fertilizers significantly improved the number of seeds per pod and the seed protein content in black-eyed peas.¹³ Au-capped mesoporous silica NPs delivered DNA into the protoplasts, cells and leaves of tobacco.¹⁴ Treatment of licorice seedlings with copper oxide (CuO) and zinc oxide (ZnO) increased the content of anthocyanins, flavonoids, glycyrrhizin, phenolic compounds and tannins.¹⁵ Silica–silver NPs were reported to possess antimicrobial activity against several plant pathogens. The application of silica–silver NPs to infected plants of green squash was found to be useful for powdery mildew control.¹⁶ Recent studies have shown that surface disinfection of explants with NPs significantly reduces microbial contamination in various plants. The addition of NPs to tissue culture media can eliminate bacterial contamination and enhance the morphogenetic potential of explants. The inclusion of NPs in the medium also resulted in somaclonal variation. The treatment of plant cell, tissue or organ cultures with NPs increased their content of bioactive compounds. Engineered NPs have been used to deliver DNA or proteins into plants. Recently, Sarmast and Salehi¹⁷ reviewed the application of silver NPs in plant tissue culture. It was interesting to observe that although there were scattered reports across plant

biotechnology journals on the application of nanomaterials in plant tissue culture, there were no consolidated reviews on this topic. This review summarizes the current achievements regarding the use of NPs in plant tissue culture. The milestones achieved in the elimination of microbial contamination in plant cultures, callus induction, organogenesis, somatic embryogenesis, somaclonal variation, genetic transformation and metabolite production through the introduction of nanomaterials are presented. The need to incorporate more of the new-age nanomaterials, such as graphene and carbon buckyballs, and the possibility of creating nano-environments for effective plant tissue culture are speculated upon in the future prospects section.

2. Nanotechnological achievements in plant tissue culture (Fig. 1)

2.1 Input of nanomaterials for surface disinfection of explants

Microbial contamination is a serious problem when it comes to plant tissue culture (Fig. 2). Microbial contamination can extinguish the entire process and efficiency of the propagation system even before initialization. The explants themselves and the laboratory environment used are the sources of contaminants.¹⁸ Usually, the various organs collected from field- or greenhouse-grown plants are surface sterilized before the establishment of *in vitro* cultures. However, in many instances, this has proved inadequate and hence many experimental set

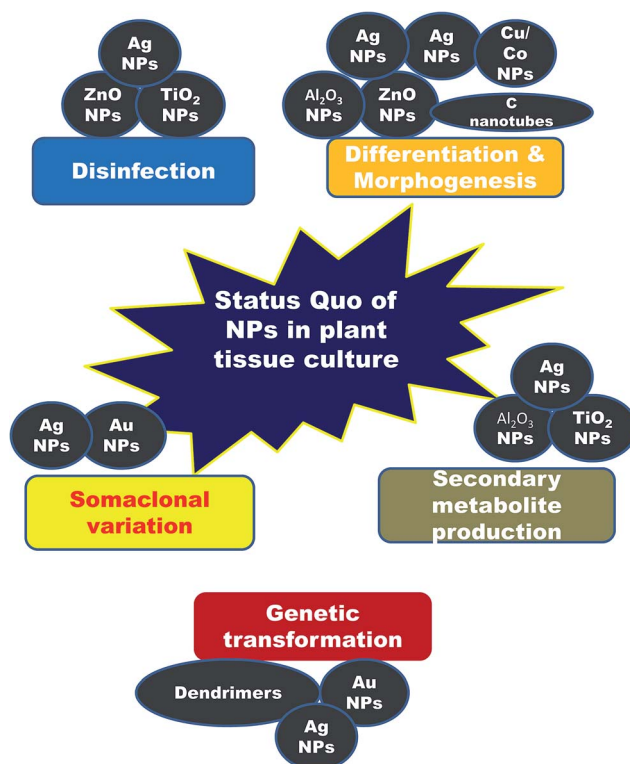


Fig. 1 Schematic showing an overview of the various NPs currently involved in plant tissue culture.



ups have been ruined, which has led to a loss of labor and time. Surface disinfection of the explants is a major step prior to *in vitro* culture initiation because microorganisms grow faster in tissue culture medium than the explants do, and they can seriously affect culture initiation. Several sterilizing agents such as bromine water (BW), calcium hypochlorite (CaOCl), ethanol (EtOH), hydrogen peroxide (H₂O₂), sodium hypochlorite (NaOCl), mercuric chloride (HgCl₂), silver nitrate (AgNO₃), antibiotics, and fungicides are in use to obtain sterile explants.¹⁸ The concentration of disinfectants and exposure time often affect the quality of the explants. Moreover, eliminating endogenous bacteria is often very challenging and therefore, several antibiotics and AgNO₃ are included in the culture medium to kill the bacteria or prevent bacterial growth.¹⁹ However, the phytotoxicity of antibiotics to cell, tissue and organ cultures of various plant species has also been reported.^{19–23} Some disinfecting agents have even failed to eliminate contaminants in explants, and instead they profoundly affected organogenesis due to their phytotoxicity. Metal and metal oxide NPs have been proven to be useful for the elimination of various microorganisms.²⁴ A wide range of NPs such as silver (Ag), aluminum oxide (Al₂O₃), CuO, iron oxide (Fe₃O₄), gold (Au), magnesium oxide (MgO), nickel (Ni), silicon (Si), SiO₂, titanium dioxide (TiO₂), and ZnO have been reported to possess antimicrobial activities against various microorganisms.²⁵ Abdi *et al.*²⁶ employed Ag NPs for the first time to control bacterial

contamination in *Valeriana officinalis*. Single node explants obtained from greenhouse-grown plants were surface disinfected with 70% EtOH for 1 min, 10% Clorox for 1 min and then 100 mg L⁻¹ of Ag NPs for 180 min; this process resulted in 89% contamination-free cultures. Moreover, the Ag NP treatment did not affect shoot multiplication and subsequent rooting. Similarly, treatment of the immature capitulum of *Gerbera jamesonii* with 1.5% NaOCl for 10 min and 200 mg L⁻¹ Ag NPs for 15 min successfully eliminated all contaminants and showed no adverse effects on organogenesis.²⁷ Leaf explants of *Vitis vinifera* 'Farkhi', 'Khoshnave' and 'Rashe' were successfully decontaminated with 95% EtOH for 30 s, and 1000 mg L⁻¹ Ag NPs for 20 min.²⁸

Mahna *et al.*²⁹ investigated the effect of Ag NP treatment on surface disinfection of *Arabidopsis* seeds, potato leaves and tomato cotyledons. The treatment of the explants with 100 mg L⁻¹ Ag NPs for 1 or 5 min was found to be ideal for decontaminating (100%) the seeds, leaves or cotyledons and this treatment had no side effects on explant viability. However, seed germination and leaf and cotyledon survival were reduced when they were treated with higher concentrations of Ag NPs. Elimination of microbes from *in vitro* cultures is a huge challenge when it comes to woody plants. Node and shoot tips obtained from 9 year-old olive plants were treated with 70% EtOH for 1 min, followed by 10% Clorox for 10 min, and this yielded 51.4% contamination-free culture. Treating the explants in 100 mg L⁻¹ Ag NPs for 60 min after EtOH and Clorox exposure completely prevented contamination, but only a few explants survived. On the other hand, supplementation of Ag NPs (4 mg L⁻¹) to the culture medium adequately controlled internal contaminants in the olive explants and no negative effects were observed on the explants and their growth.³⁰ Sarmast *et al.*³¹ reported that surface sterilization of explants of *Araucaria excelsa* followed by immersion in 200 mg L⁻¹ of Ag NPs for 180 min reduced the rate of contamination from 61.5 to 11.3%. The inclusion of 400 mg L⁻¹ of Ag NPs in the culture medium reduced the contamination from 81.25 to 18.75% (Table 1). Similarly, the application of 100 and 150 mg L⁻¹ Ag NPs, through both immersion and addition to the MS medium, significantly reduced internal and external contamination in G × N15 (hybrid of almond × peach) rootstocks.³²

The addition of 50 mg L⁻¹ Ag NPs to Murashige and Skoog (MS³³) medium significantly inhibited microbial growth.³⁴ Similarly, contamination-free cultures were observed when surface-disinfected shoot buds obtained from potato and tobacco were cultured on MS medium supplemented with Ag NPs or TiO₂ NPs.^{35,36} Ideal results were attained on MS medium containing 1% (w/w) TiO₂ NPs.³⁷ Bacterial contamination was notably reduced when *Rosa hybrida* explants were treated with 200 mg L⁻¹ Ag NP solution for 20 min. On the other hand, the supplementation of 100 mg L⁻¹ AgNPs to the medium reduced the rate of bacterial contamination and also phenolic exudation.³⁸ It has been reported that internal bacterial contamination in callus cultures of *Bacopa monnieri* was significantly reduced by the addition of 160 mg L⁻¹ Ag NPs to the MS medium.³⁹ The addition of 60 μg mL⁻¹ TiO₂ NPs to the MS medium eliminated bacterial contamination in callus cultures

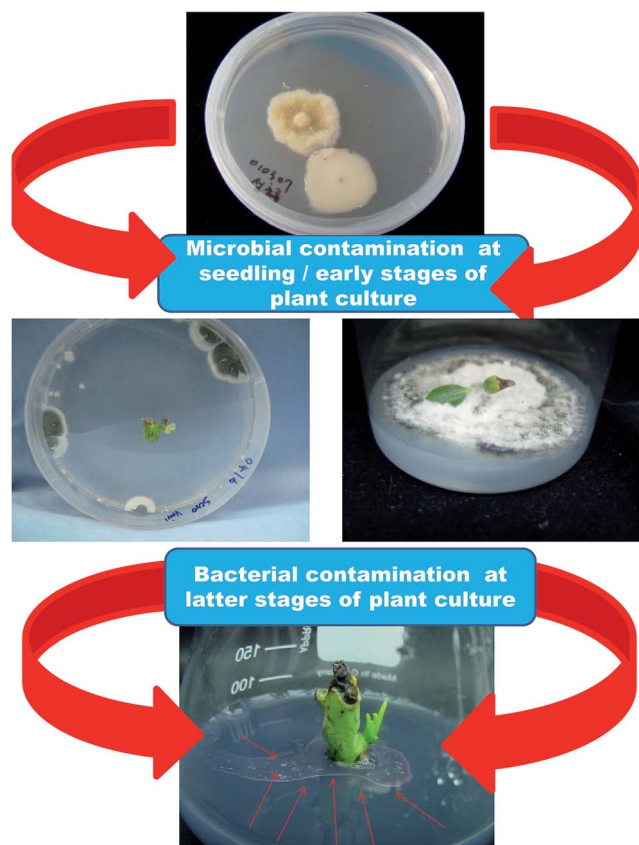


Fig. 2 Slides showing microbial contamination during different phases of the plant culturing process.





Table 1 Application of NPs in plant tissue culture

Plant species	NPs	Size (nm)	Conc.	Phytotoxicity	Role	References
<i>Nicotiana tabacum</i>	Au-capped mesoporous silica	3, 10–15	3.3 $\mu\text{g mL}^{-1}$	–	Delivered DNA into protoplasts, cells and leaves	14
<i>Valeriana officinalis</i> L.	Ag	35	25–100 mg L^{-1}	–	Reduced bacterial contamination	26
<i>Agrostis stolonifera</i> L.	Poly(amidoamine) dendrimer	4.5	–	–	Delivered the GFP gene to cells of creeping bent grass	76
<i>Arabidopsis thaliana</i>	C Nanotubes	9.5	10–600 mg L^{-1}	+	Decreased cell viability	63
<i>Oryza sativa</i> L.	C Nanotubes	10–30	20 & 40 mg L^{-1}	+	Decreased cell viability and dry weight	64
Olive 'Mission'	Ag	–	100–400 mg L^{-1}	+	Reduced the rate of contamination	30
			4–16 mg L^{-1}	–		
<i>Nicotiana tabacum</i> , <i>Oryza sativa</i>	Sharp carbon-supported Au	33 \pm 4	200–2000 ng	–	Delivered the GUS gene	75
<i>leucocephala</i>						
<i>Daucus carota</i> L.	Fe_3O_4	6	2.01–33.5 mg L^{-1}	+	Decreased cell proliferation and number of somatic embryos	71
Some plants	Ag	–	8–80 mg L^{-1}	–	Reduced microbial contamination	34
Tobacco	Ag	35	–	–	Reduced bacterial contamination	35
	TiO_2	10	–	–		
<i>Araucaria excelsa</i> R. Br. var. <i>glauca</i>	Ag	–	200 & 400 mg L^{-1}	–	Reduced bacterial contamination	31
<i>Brassica juncea</i> L.	CaP	20–50	0.5 mg mL^{-1}	–	Increased the transformation efficiency	77
<i>Brassica juncea</i>	Ag	29	25–400 mg L^{-1}	+	Enhanced seedling growth and decreased the content of hydrogen peroxide, malondialdehyde and proline	53
<i>Gerbera jamesonii</i>	Ag	–	25–200 mg L^{-1}	–	Reduced microbial contamination	27
<i>Hordeum vulgare</i> L.	TiO_2	24.5	10–60 $\mu\text{g mL}^{-1}$	–	Increased callus formation and callus size, and reduced bacterial contamination	40
<i>Linum usitatissimum</i> L.	C	5–20	1–3 g L^{-1}	–	Decreased the percentage of callus induction and somatic embryogenesis.	65
					Increased the number of tetraploid cells and the methylation level	
<i>Solanum tuberosum</i> L.	Ag	35	5–100 mg L^{-1}	–	Reduced bacterial contamination	36
Tobacco	TiO_2	10	–	–		
	C nanotubes	22	5–500 $\mu\text{g mL}^{-1}$	+	Stimulated cell growth at low concentration	62
<i>Tecomella undulata</i>	Ag	18.2	5–80 mg L^{-1}	–	Increased frequency of shoot induction, number of shoots per explant and plantlet survival	51
<i>Arabidopsis Nicotiana tabacum</i>	Mesoporous silica	40–50	–	–	Delivered the red fluorescence protein gene to protoplasts of tobacco and roots of <i>Arabidopsis</i>	79
<i>Arabidopsis thaliana</i>	Au	24	10 & 80 $\mu\text{g mL}^{-1}$	–	Improved the percentage of seed germination, vegetative growth and antioxidant activity	55
<i>Arabidopsis thaliana</i>	CeO_2	10–30	0–3000 mg L^{-1}	+	Increased plant biomass and anthocyanin content	68
	In_2O_3	20–70	–	–		



Table 1 (Contd.)

Plant species	NPs	Size (nm)	Conc.	Phytotoxicity	Role	References
<i>Arabidopsis thaliana</i>	Ag	—	100–2000 mg L ⁻¹	+	Reduced microbial contamination at lower concentrations and consequently, all decontaminated seeds germinated, and leaf and cotyledon explants survived	29
<i>Lycopersicon Esculentum</i>	Ag	—	25–100 mg L ⁻¹	+		
<i>Solanum tuberosum</i>	Ag	—	10–500 mg L ⁻¹	+	Enhanced artemisinin production in hairy root cultures	95
<i>Artemisia annua</i>	Ag-SiO ₂	101.8 ± 8.9	900 mg L ⁻¹	—	Delivered the GUS gene to protoplasts	80
<i>Brassica napus</i>	Magnetic Au	20–30	7.5–30 µg L ⁻¹	—		
<i>Daucus carota</i> L.	Au	—	1 mg L ⁻¹	—	Gold NPs increased callus growth, regeneration zones and somatic embryogenesis more than silver NPs	72
<i>Linum usitatissimum</i> L.	Ag	—	1 mg L ⁻¹	—	Increased phenolic content and cell density, and decreased cell viability	67
<i>Nicotiana tabacum</i>	Al ₂ O ₃	<50	10–100 µg mL ⁻¹	+	Enhanced aloin content in cell suspension cultures	98
<i>Aloe vera</i> L.	Ag	—	0.63 mg L ⁻¹	—	Enhanced root growth and accumulation of anthocyanin	66
<i>Arabidopsis</i>	TiO ₂	—	120 mg L ⁻¹	—		
<i>Bacopa monnieri</i>	Ag	29–33	16 × 10 ⁻³ to 16 × 10 ⁻¹⁰	+	Reduced the rate of contamination	39
Banana	Zn	35	50–200 mg L ⁻¹	—	Eliminated microbial contamination, and enhanced somatic embryogenesis, and chlorophyll and proline content	41
<i>Calendula officinalis</i> L.	ZnO	—	1–100 µM	—	Increased the fresh and dry weights of calli and secondary metabolite production	91
<i>Calendula officinalis</i> L.	AgNO ₃	—	0.3–1.2 mg L ⁻¹	—		
<i>Cynodon dactylon</i>	Ag	—	100 & 200 mg L ⁻¹	+	A combination of Ag NPs and thymol inhibited microbial growth.	42
G × N15 (hybrid of almond × peach) rootstock	Ag	—	50–200 mg L ⁻¹	+	Reduced internal and external contamination	32
<i>Nicotiana tabacum</i>	CaP	20–25	—	—	Delivered plasmid DNA into tobacco cells	78
<i>Rosa hybrida</i>	Ag	—	100–400 mg L ⁻¹	—	Reduced the rate of bacterial contamination and phenolic exudation	38
<i>Solanum tuberosum</i> L.	TiO ₂	10	1 & 2% (w/w)	—	Reduced microbial contamination	37
<i>Vitis vinifera</i> L.	Ag	—	100–1000 mg L ⁻¹	+	Reduced microbial contamination	28
<i>Artemisia annua</i>	Co	10	0.25–5 mg L ⁻¹	—	Increased artemisinin content in cell suspension cultures	96
<i>Cicer arietinum</i> L.	TiO ₂	—	0.5–6 mg L ⁻¹	—	Increased phenolic and flavonoid compounds in calli	92
<i>Datura metel</i>	Ag	50–60	20 mg L ⁻¹	—	Enhanced the atropine content in hairy roots	97
<i>Lilium ledebourii</i>	ZnO	—	10–100 mg L ⁻¹	+	Increased the content of chlorophyll, anthocyanin, flavonoid and phenol, bulb number, number of leaves, and length of leaves and roots	93
<i>Nicotiana tabacum</i>	Ag	96	1–15 mg L ⁻¹	+	Reduced microbial contamination and increased the yielded of viable protoplasts	74



Table 1 (Contd.)

Plant species	NPs	Size (nm)	Conc.	Phytotoxicity	Role	References
<i>Satureja khuzestanica</i>	C Nanotubes	—	25–500 $\mu\text{g mL}^{-1}$	—	Enhanced callus growth and secondary metabolite production	61
<i>Solanum nigrum</i> L.	Ag	30–90	2–8 mg L^{-1}	—	Increased callus formation frequency and fresh weight of the callus	60
<i>Stevia rebaudiana</i>	Zn	<100	50–1000 mg L^{-1}	+	Reduced stevioside production	94
<i>Tecomella undulata</i>	Ag	—	30–120 $\mu\text{g mL}^{-1}$	—	Increased frequency of shoot induction, number of shoots and shoot length	52
<i>Arabidopsis thaliana</i>	CeO ₂	10–30	250 & 1000 mg L^{-1}	+	Increased ROS levels in roots	69
<i>Brassica nigra</i>	In ₂ O ₃	20–70	500–1500 mg L^{-1}	+	Inhibited seed germination, seedling growth and callus biomass, while inducing roots from stem explants. Increased secondary metabolite content in callus and shoot and root tissues of seedlings	54
	ZnO	100	1–20 mg L^{-1}	—		
<i>Capsicum</i> sp	Ag	—	50 mg L^{-1}	—	Enhanced capsaicin production in cell suspension cultures	99
<i>Corylus avellana</i> L.	Ag	—	2.5–10 mg L^{-1}	+	Increased the production of taxol and decreased viability of cells	100
<i>Mentha longifolia</i>	Cu	—	0.5 mg L^{-1}	—	Increased number of shoots, quantity of internodes, shoot height, growth index and percent of essential oil output	57
	Co	—	0.8 mg L^{-1}	—		
<i>Oryza sativa</i> L.	CuO	40 ± 5	1–20 mg L^{-1}	—	Enhanced the frequency of callus induction and shoot regeneration	70
<i>Prunella vulgaris</i> L.	Au	25–35	30 $\mu\text{g L}^{-1}$	—	Enhanced callus proliferation, protein content, antioxidant activity and accumulation of flavonoids and phenolics	73
	AgAu	—	—	—		
<i>Solanum lycopersicum</i>	ZnO	30	15 and 30 mg L^{-1}	—	Improved callus growth and regeneration under salinity stress	59
<i>Hyoscyamus reticulatus</i>	Fe ₃ O ₄	—	450–3600 mg L^{-1}	+	Stimulated hyoscyamine and scopolamine production in hairy root cultures	101
<i>Linum usitatissimum</i> L.	Au	—	500 mg L^{-1}	—	Increased somaclonal variation percentage, callus growth and regeneration	87
	Ag	—	500 mg L^{-1}	—		
<i>Stevia rebaudiana</i>	ZnO	34	0.1–1000 mg L^{-1}	+	Enhanced percentage of shoot formation, quantity of steviol glycosides, total flavonoid and phenolic contents	58
<i>Vanilla planifolia</i>	Ag	35 ± 15	25–200 mg L^{-1}	+	Reduced bacterial contamination, and increased shoot growth, antioxidant capacity and total phenolic content	43
<i>Verbena bipinnatifida</i> Nutt.	CuSO ₄	—	5–15 $\mu\text{g L}^{-1}$	+	Increased shoot and root lengths, fresh weight and total phenolic content	56

of barley after the 4th subculture.⁴⁰ Helaly *et al.*⁴¹ reported that shoot tips obtained from 3 year-old banana suckers when surface-sterilized with 80% NaOCl, 95% EtOH and finally 0.1% HgCl₂ for 15 min failed to eliminate microbial contamination. The authors identified nine strains of bacterial contaminants in the *in vitro* cultures (Table 1). The incorporation of 100 mg L⁻¹ Zn NPs or ZnO NPs into the MS medium resulted in contamination-free cultures. Taghizadeh and Solgi⁴² reported that a combination of Ag NPs and thymol inhibited microbial growth in *Cynodon dactylon*. Contamination-free cultures were established by treating nodal explants with 200 mg L⁻¹ Ag NP solution containing 100 mg L⁻¹ thymol for 60 min. Spinoso-Castillo *et al.*⁴³ reported that zero contamination was observed in shoot cultures of *Vanilla planifolia* when MS medium was fortified with 50–200 mg L⁻¹ Ag NPs (Fig. 3).

The antibacterial property of NPs is no new theme; it is one of the most established and well-studied achievements of NPs, with the most renowned being Ag NPs, TiO₂ NPs and ZnO NPs.^{44–50} The effectiveness of NPs in the elimination of microbial contaminants in plant tissue cultures depends on their dimensions, size, distribution, and type. Several researchers have reported the survival and subsequent regeneration of explants that were affected negatively by NPs during application periods. Therefore, the effects of different types and levels of NPs on explants obtained from various plant species should be investigated in order to determine the best dose with no or minimal phytotoxicity. The effectiveness of NPs can also be improved by synergistic effects when they are combined with sterilizing agents or antibiotics.

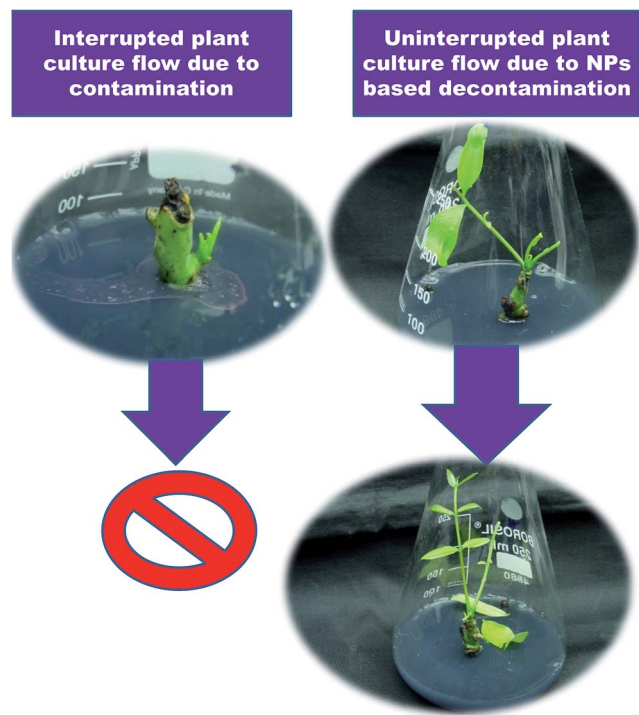


Fig. 3 A smooth and uninterrupted tissue culture process through integration of NPs into the medium for disinfection.

2.2 Input of nanomaterials towards callus induction, organogenesis, shoot growth and rooting

Several studies have shown positive effects of NPs on callus induction, shoot regeneration and growth. In *Tecomella undulata*, the shoot induction percentage, number of shoots and callus formation were increased when the stem explants were cultured on MS medium augmented with 10 mg L⁻¹ Ag NPs, 2.5 mg L⁻¹ BAP and 0.1 mg L⁻¹ IAA.⁵¹ The positive effects of Ag NPs on organogenesis may be due to inhibition of ethylene production. Shoot number, length of induced shoots and percentage of produced shoots were significantly enhanced when nodal explants of *T. undulata* were cultured on MS medium fortified with 60 mg L⁻¹ Ag NPs, 2.5 mg L⁻¹ BAP and 0.1 mg L⁻¹ IAA.⁵² Also, Ag NP treatment delayed explant senescence and increased survival by downregulation of the *TuACS* gene. However, at higher concentrations (above 60 mg L⁻¹) a detrimental effect on shoot regeneration was evident. Sharma *et al.*⁵³ reported that incorporation of Ag NPs at 50 mg L⁻¹ into MS basal medium improved the growth characteristics of *Brassica juncea* seedlings by decreasing the content of hydrogen peroxide, malondialdehyde (MDA) and proline through activating antioxidant enzymes. However, higher concentrations of Ag NPs (100–400 mg L⁻¹) had a detrimental effect on seedling growth. In *Brassica nigra*, adding ZnO NPs (500–1500 mg L⁻¹) to the MS medium significantly inhibited seed germination. In the presence of ZnO NPs, the shoot and root lengths were significantly affected. On the other hand, growth of stem explants of *B. nigra* on MS medium containing 1–20 mg L⁻¹ ZnO NPs resulted in root formation.⁵⁴ Kumar *et al.*⁵⁵ reported that the incorporation of Au NPs into MS basal medium improved the percentage of seed germination and seedling growth in *Arabidopsis thaliana*. Pod length and number of seeds were higher in plants treated with 10 µg mL⁻¹ Au NPs. The treatment with Au NPs improved the antioxidant enzyme activity and led to a decrease in the expression of microRNAs (miR398 and miR408) in *A. thaliana*. These physiological and molecular variations might be responsible for the beneficial effects of Au NPs. The inclusion of 5 µg L⁻¹ copper sulfate (CuSO₄) NPs into the MS medium significantly increased the shoot length (52% over control), root length (21% over control) and fresh weight (39% over control) of *Verbena bipinnatifida* seedlings.⁵⁶ The addition of 0.5 mg L⁻¹ Cu NPs and 0.8 mg L⁻¹ Co NPs to modified MS medium increased the number of shoots, shoot length and rooting in *Mentha longifolia*.⁵⁷ The highest frequency of shoot formation (89.6%) was obtained when nodal explants of *Stevia rebaudiana* were cultured on MS medium amended with 1 mg L⁻¹ ZnO NPs.⁵⁸ Spinoso-Castillo *et al.*⁴³ investigated the hormetic effects of Ag NPs (Argovit, a commercial product) on shoot regeneration of *Vanilla planifolia* using a temporary immersion bioreactor system. The highest number of shoots was obtained on MS medium supplemented with 25 or 50 mg L⁻¹ Ag NPs, with values of 14.33 and 14.89, respectively, whereas the lowest number of shoots (4.55) were observed on MS medium supplemented with 200 mg L⁻¹ of Ag NPs.

In tomato plants, callus growth and plant regeneration were maximized on medium containing 15 mg L⁻¹ ZnO NPs and 3.0 g



L^{-1} NaCl.⁵⁹ ZnO NPs mitigated the effects of NaCl by upregulating antioxidant enzymes like GPX and SOD. In *Solanum nigrum*, the frequency of callus formation (89%) and the fresh weight of the callus (4.67 g per leaf explant) were increased on MS medium augmented with 5 mg L^{-1} BA, 3 mg L^{-1} NAA and 8 mg L^{-1} Ag NPs.⁶⁰ Ghorbanpour and Hadian⁶¹ investigated the effect of multi-walled carbon nanotubes (25–500 $\mu\text{g mL}^{-1}$) on callus induction from leaf explants of *Satureja khuzestanica*. Callus growth improved significantly on B5 medium augmented with 25–50 $\mu\text{g mL}^{-1}$. However, the presence of multi-walled carbon nanotubes at 100–500 $\mu\text{g mL}^{-1}$ decreased the callus biomass. Khodakovskaya *et al.*⁶² reported that the incorporation of 100 $\mu\text{g mL}^{-1}$ multi-walled carbon nanotubes to a medium containing 1 mg L^{-1} 2,4-D increased the callus growth of tobacco explants (64% increase over control). The treatment with carbon nanotubes enhanced callus growth by upregulation of the genes involved in cell division (*CycB*), cell wall extension (*NtLRX1*) and water transport (*NtPIPI1*). However, carbon nanotube treatment (10–600 mg L^{-1}) decreased cell viability and dry weight in *Arabidopsis*.⁶³ Similarly, the addition of carbon nanotubes to rice cell suspension cultures decreased cell viability.⁶⁴ The presence of carbon NPs in the MS medium reduced the frequency of callus induction in *Linum usitatissimum*.⁶⁵ Syu *et al.*⁶⁶ investigated the effects of three different morphologies of Ag NPs, including decahedral (45 \pm 5 nm), spherical (8 \pm 2 nm) and triangular (47 \pm 7 nm), on seedling growth, gene expression and physiological changes in *Arabidopsis* (Table 1). Root growth was enhanced when the seedlings were treated with either triangular or decahedral Ag NPs, whereas it was unaffected following spherical Ag NP treatment. Ag NP treatment was reported to alter the content of antioxidant enzymes and expression of genes that are involved in the biosynthesis of auxin, abscisic acid and ethylene.

Poborilova *et al.*⁶⁷ reported that the addition of Al_2O_3 NPs (10–100 $\mu\text{g mL}^{-1}$) to tobacco cell suspension cultures significantly decreased cell viability through the generation of reactive oxygen (ROS) and nitrogen species (RNS). The presence of 250 mg L^{-1} cerium oxide (CeO_2) and 50–500 mg L^{-1} indium oxide (In_2O_3) in half-strength MS medium significantly increased the biomass of *Arabidopsis thaliana* seedlings.⁶⁸ However, seedling growth decreased at higher concentrations. ROS production significantly increased when seedlings were exposed to 250 mg L^{-1} CeO_2 and 1000 mg L^{-1} In_2O_3 , demonstrating clear oxidative stress in *Arabidopsis*.⁶⁹ In banana explants, the maximum frequency of somatic embryogenesis was observed in MS medium augmented with 100 mg L^{-1} Zn NPs followed by ZnO NPs. Shoot and root lengths were also increased when adding both NPs to the MS medium.⁴¹ However, higher concentrations decreased the callus growth and enhanced antioxidant enzyme activity. Anwaar *et al.*⁷⁰ reported that supplementation with CuO NPs (15–20 mg L^{-1}) increased organogenesis in rice cultivars. In *Daucus carota*, cell proliferation and the number of somatic embryos decreased on MS medium containing Fe_3O_4 .⁷¹ It has been reported that TiO_2 NPs may play a role similar to plant growth regulators (PGRs) like cytokinin and gibberellic acid.⁴⁰ The number and size of calli increased when barley mature embryos were grown in MS

medium augmented with 20 mg L^{-1} 2,4-D and 60 $\mu\text{g mL}^{-1}$ TiO_2 NPs. Kokina *et al.*⁷² prepared a BAP (cytokinin) powder coated with Au and Ag NPs and added it to the tissue culture medium. Stem segments of *Linum usitatissimum* were cultured on MS medium augmented with 1 mg L^{-1} 2,4-D plus 1 mg L^{-1} BAP or 1 mg L^{-1} Au BAP or Ag BAP. The inclusion of Au NPs and Ag NPs in the MS medium increased embryogenesis (70% and 50%, respectively). The authors confirmed deposition of NPs in the cells and hypothesized that the plants actively take up the plant growth regulators (PGRs) and the metal NPs are transported along with the PGRs. However, the actual mechanism *via* which Au NPs enhance embryogenesis is still unclear and needs further investigation. Fazal *et al.*⁷³ studied the effects of Au and Ag NPs individually or in combination on callus proliferation in *Prunella vulgaris*. The Ag (30 $\mu\text{g mL}^{-1}$), Ag–Au (1 : 2), and Ag–Au (2 : 1) NPs in combination with NAA (2.0 mg L^{-1}) enhanced callus proliferation (100%) compared to the control (95%). The highest biomass was obtained when the culture medium was augmented with Au NPs (Table 1).

Hence, from all of these reports it can be concluded that the addition of NPs to a plant tissue culture medium affects callus proliferation, shoot multiplication, somatic embryogenesis and rooting by altering antioxidant enzyme activities, gene expression, inhibition of ethylene production and production of ROS. However, the actual mechanisms of the promotive or inhibitory effects of NPs on each parameter need to be investigated in detail. The effects of various metal and metal oxide NPs on plants are well documented *in vivo*; such NPs can be used to promote or improve the morphogenetic potential of explants obtained from different plant species. The influence of different concentrations and combinations of NPs on various media (shoot induction, shoot multiplication and rooting media) also deserves evaluation, in order to gain a clear understanding of the underlying mechanisms behind the role of NPs in plant tissue culture.

2.3 Effect of nanomaterials on genetic transformation

Electroporation and particle bombardment (direct) and *Agrobacterium*-mediated (indirect) transformation methods are used for delivering a foreign gene into plant cells, tissues and organs. The electroporation technique has been widely used for transferring genes into protoplasts. However, the isolation and regeneration of protoplasts is not that easy. Bansod *et al.*⁷⁴ reported that the incorporation of 10 mg L^{-1} Ag NPs into the leaf incubation buffer enhanced the viable protoplast yield in tobacco. This is helpful in minimizing the damage caused by cellulolytic enzymes during protoplast isolation and improved costs (which is yet another constraint in protoplast isolation). Torney *et al.*¹⁴ demonstrated that mesoporous silica NPs could be used to deliver DNA into tobacco protoplasts by endocytosis; the gold-capped mesoporous silica NPs, DNA and chemicals were delivered into the callus and leaves using a biolistic gun. Vijayakumar *et al.*⁷⁵ showed that carbon-supported Au NPs delivered DNA more efficiently into *Nicotiana tabacum*, *Oryza sativa* and *Leucaena leucocephala* compared to regular gold particles using a gene gun. The amounts of gold and plasmid



present in the carbon-supported Au NPs were 4 and 3 times lower, respectively, compared to the commercial micrometer-sized gold particles. In addition, plant cell damage is minimal and therefore plant regeneration and the transformation frequency is increased. In *Agrobacterium*-mediated transformation, antibiotics such as carbenicillin, cefotaxime, rifampicin and thimentin have been used for the removal of bacteria after co-cultivation; however, their phytotoxic effects on explants affected the regeneration potential and genetic stability of the regenerated plantlets. Sarmast and Salehi¹⁷ reported that the growth of *A. rhizogenes* and *A. tumefaciens* was completely suppressed on LB medium containing $10 \mu\text{g mL}^{-1}$ Ag NPs. The authors also demonstrated that addition of Ag NPs to the culture medium successfully eliminated bacteria after co-cultivation during the *Agrobacterium*-mediated genetic transformation of *T. undulata* and tobacco. Other NPs require further testing in order to provide evidence for them eliminating *Agrobacterium* in the cells, tissues and organs of various plants following co-cultivation.

Pasupathy *et al.*⁷⁶ developed a novel gene delivery method in plants by using poly(amidoamine) dendrimer NPs. The authors successfully delivered green-fluorescent protein (GFP)-encoding plasmid DNA into turf-grass cells. The transfection efficiency was further enhanced by optimizing the pH of the culture medium and the molar ratio of the dendrimer to plasmid DNA. Naqvi *et al.*⁷⁷ used calcium phosphate (CaP) NPs for delivering pCambia 1301 harboring the GUS gene into *Brassica juncea*. The best transformation efficiency was observed with CaP NPs (80.7%) followed by *Agrobacterium tumefaciens* (54.4%) and naked DNA (8%). Recently, Ardekani *et al.*⁷⁸ reported that CaP NPs delivered pBI121 harboring the GFP gene into tobacco cells (Table 1). Mesoporous silica NPs have been used for delivering plasmid DNA into tobacco protoplasts and intact *Arabidopsis* roots.⁷⁹ Magnetic Au NPs can deliver plasmid DNA into the cells and protoplasts of canola and carrot plants.⁸⁰ NP-mediated gene delivery in plants has great importance in plant nanobiotechnology. Further research is warranted to study the effects of different NPs on genetic manipulation of various plant species.

2.4 Influence of nanomaterials on somaclonal variation

Changes observed in *in vitro* developed organs and plantlets are termed somaclonal variation. It is usually associated with changes in chromosome number, chromosome structure, DNA sequence, DNA methylation, mitotic crossing over and activation of transposable elements.^{81,82} Somaclonal variation has both advantages as well as disadvantages in plant tissue culture. Variants possess several useful characteristics like plant size, flower color, leaf variegation, fruit ripening, secondary metabolite production and resistance to biotic and abiotic stresses.⁸³ Several studies have demonstrated the phytotoxicity of NPs primarily applied at higher levels. NP treatments affect the mitotic index and DNA integrity, and alter the protein and DNA expression in plants.^{84–86} The ploidy level in *L. usitatissimum* calli was significantly affected on MS medium containing C NPs.⁶⁵ The supplementation of C NPs enhanced the number of

tetraploid cells. The level of DNA methylation was also higher in calli grown on medium with C NPs. Kokina *et al.*⁸⁷ investigated the effects of Au and Ag NPs on somaclonal variation in *L. usitatissimum*. The occurrence of somaclonal variation was higher in both calli and regenerated shoots grown on medium containing Au NPs than Ag NPs. Ewais *et al.*⁶⁰ reported that the addition of Ag NPs to tissue culture medium induces variation in callus morphology and anatomy by altering the protein and DNA profile in *Solanum nigrum* calli (Table 1). Further investigations are required in order to determine the effects of a wide range of NPs for enhancing somaclonal variation.

2.5 Nanomaterial enhancement of secondary metabolites

Plants are a rich source of various bioactive secondary metabolites, which play a significant role in the survival of plants in their respective environments. *In vitro* plant cell and organ culture has been proven to be advantageous for the production of secondary metabolites. The content of secondary compounds in cell and organ cultures was significantly enhanced by optimizing the composition of the culture medium, incorporation of precursors and elicitors and providing appropriate culture conditions.^{83,88–90} NPs added to the plant *in vitro* culture medium may act as a nutrient source and an elicitor. Poborilova *et al.*⁶⁷ reported that the addition of Al_2O_3 NPs ($10\text{--}100 \mu\text{g mL}^{-1}$) to tobacco cell suspension cultures significantly increased the phenolic content. The accumulation of phenolics in the cells was dose- and exposure time-dependent. Mohammed Al-oubaidi and Mohammed-Ameen⁹¹ reported that the essential oil content in *Calendula officinalis* calli was greatly enhanced in MS medium amended with 0.3 mg L^{-1} Ag NPs. The presence of TiO_2 NPs (4.5 or 6.0 mg L^{-1}) significantly increased the content of gallic acid, chlorogenic acid, *o*-coumaric acid, tannic acid and cinnamic acid in embryonic calli of *Cicer arietinum*.⁹² The application of Au–Ag NPs (1 : 3) induced maximum accumulation of total phenolic compounds and flavonoids in callus cultures of *Prunella vulgaris*.⁷³ Syu *et al.*⁶⁶ reported that the shape of Ag NPs plays a significant role in the production of anthocyanins in *Arabidopsis*. Treatment with spherical Ag NPs resulted in the highest levels of anthocyanin accumulation in seedlings. *Vanilla planifolia* shoots grown in MS medium supplemented with 25 and 50 mg L^{-1} Ag NPs showed a significant increase in total phenolic content.⁴³ Chamani *et al.*⁹³ reported the accumulation of specific bioactive compounds in *Lilium ledebourii* and its dependency on the concentration of ZnO NPs in the MS medium. The highest content of flavonoids, phenolics and anthocyanins was obtained on MS medium supplemented with 25, 75 and 100 mg L^{-1} ZnO NPs, respectively. The accumulation of steviol glycosides in shoot cultures of *S. rebaudiana* was significantly enhanced on MS medium fortified with 1 mg L^{-1} ZnO.⁵⁸ In addition, the total flavonoid and phenolic content also increased with ZnO treatment. However, higher concentrations of ZnO led to decreased secondary metabolite production due to the phytotoxic effects of ZnO. Desai *et al.*⁹⁴ reported that the addition of $50\text{--}1000 \text{ mg L}^{-1}$ Zn NPs decreased the production of stevioside in shoot cultures of *S. rebaudiana* (Table 1). The content of total phenolics in the



seedlings of *Verbena bipinnatifida* was enhanced two-fold when they were grown on MS medium containing 5 μM CuSO_4 NPs.⁵⁶ Application of Cu and Co NPs increased the essential oil content in *Mentha longifolia* by 2.226 and 2.19%, respectively.⁵⁷ In *S. khuzestanica* leaf cultures, the incorporation of multi-walled carbon nanotubes in B5 medium (25 or 50 $\mu\text{g mL}^{-1}$) gave maximum callus growth, while the highest content of phenolics, flavonoids, rosmarinic acid and caffeic acid was obtained on B5 medium with 100 or 250 $\mu\text{g mL}^{-1}$ multi-walled carbon nanotubes.⁶¹

Zhang *et al.*⁹⁵ investigated the elicitation potential of Ag NPs on increasing the content of artemisinin in hairy root cultures of *Artemisia annua*. The production of artemisinin showed a 3.9-fold increase when the cultures were treated with 900 mg L^{-1} Ag NPs for 3 days. Similarly, the highest artemisinin content (2.2-fold increase over the control) was obtained in cell suspension cultures of *A. annua* 24 h after treatment with 5 mg L^{-1} Co NPs.⁹⁶ The authors concluded that the Co NP treatment increased the artemisinin content as a result of downregulation of the *SQS* and *DBR2* genes. Shakeran *et al.*⁹⁷ investigated the effects of biotic (*Bacillus cereus* and *Staphylococcus aureus*) and abiotic (AgNO_3 and Ag NPs) elicitors on atropine production in hairy root cultures of *Datura metel*. Hairy root cultures were exposed to elicitors for 12, 24 and 48 h. Among the studied elicitors, Ag NPs were the most effective in enhancing the content of atropine in hairy roots. Raei *et al.*⁹⁸ reported that *Aloe vera* suspension cells interacting with 0.625 mg L^{-1} Ag NPs or 120 mg L^{-1} TiO_2 NPs for 48 h exhibited a significantly increased aloin content. The highest content of aloin was obtained subsequent to TiO_2 NP treatment. Bhat and Bhat⁹⁹ reported that addition of Ag NPs at 3 mg L^{-1} to a *Capsicum frutescens* cell suspension increased the content of capsaicin about 2-fold. Treatment of a *Corylus avellana* cell suspension culture with 5 mg L^{-1} Ag NPs significantly increased the production of taxol. However, treatment with 2.5 and 10 mg L^{-1} Ag NPs reduced the taxol production potential of the cells to 60 and 56% of the control, respectively.¹⁰⁰ Moharrami *et al.*¹⁰¹ investigated the elicitation potential of Fe NPs (450–3600 mg L^{-1}) for increasing the content of hyoscyamine and scopolamine in hairy root cultures of *Hyoscyamus reticulatus*. The exposure of hairy root cultures to 900 mg L^{-1} Fe NPs for 24 h and 450 mg L^{-1} Fe NPs for 48 h enhanced hyoscyamine and scopolamine production (Table 1). All of the studies mentioned above confirm the possibility of NPs being employed as successful and promising bioactive compound elicitors in plant cell and organ cultures. Further studies are needed to evaluate the elicitation potential of various other NPs on secondary metabolite production in plant tissue cultures and the corresponding mechanisms.

3. The toxicity concerns and safety issues surrounding exposure of plant tissue cultures to nanomaterials

There is no doubt that nanomaterials hold diverse promise and have been proven to have outstanding potential. However, working with invisible components does become more

challenging. Nanotoxicology is a flourishing branch which has evolved to record the negative aspects of nanomaterials. Because of quantum size effects and their large surface area to volume ratio, nanomaterials have unique properties compared to their larger counterparts; one such unique property is an added toxicological feature that nanomaterials may possess, unlike their corresponding bulk materials. Nanomaterials, even when made of inert elements like gold, become highly active at nanometer dimensions. Nanotoxicological studies determine whether and to what extent these properties may pose a threat to the environment and to human beings. There is an uncontrollable nano-boom that has brought about a wealth of positive changes, but simultaneously it is giving rise to a persistent form of pollution that is too small to detect or contain easily. Fighting a known enemy is much easier than fighting an invisible one. The health effects of nanopollution are yet to be fully understood, making nanopollution yet another man-made environmental disaster with uncertain long term effects in the making. Nanoparticles, being incredibly hard to isolate and quantify, require the invention of novel methods to fully assess the scale of the issue.

Thus far, the toxicity assessment of nanomaterials on plants has been mostly evaluated during *in vitro* seed germination and subsequent growth of seedlings. Researchers have elaborately reviewed the toxicity aspects of nanomaterials on plants.^{102–106} They suggest that the nanomaterials added to the culture medium can lead to significant and adverse effects on cell viability, organogenesis, shoot growth, seed germination, seedling development and explant survival. The phytotoxicity of nanomaterials is believed to depend on their chemical composition, dose, size, stability and type, the composition of the culture medium, the application method, and the explant type and plant species. *In vitro* seed germination and seedling growth of Alfalfa, barley, maize, rice, tomato and wheat was reported to be negatively impacted by high doses of carbon nanomaterials and metal NPs.^{104,107} Addition of NPs to the cell suspension reduced the viability of the cells by altering nucleic acid expression, inducing DNA damage, increasing the production of ROS, disturbing chlorophyll synthesis, inducing cell membrane damage and inducing electrolyte leakage.^{63,64,100,105} To date, the uptake of NPs into plants has not been assessed or documented. Research reports record their findings based on the addition of a broad range of concentrations to the growth medium for investigating the phytotoxicity of various nanomaterials. It has been reported that the uptake of NPs by plant cell, tissue and organ cultures is closely associated with the absorption of moisture and nutrients from the medium.^{108,109}

Most of these studies are restricted to *in vitro* conditions only; almost none or only a handful apply to *in vivo* studies. *In vivo*, laboratory or greenhouse conditions would completely differ from the field conditions; hence, more realistic and large-scale assessment of the nanomaterial life cycle in field environments will give a holistic picture to the toxicity aspects of nanomaterials on plants. More stringent dose-dependent studies are needed to identify the ideally safe NP doses that stay within the positive impacts on plant growth with lesser



negative impacts on the environment as well as plants. Thus, a series of safety evaluation and toxicological risk assessment standards should be formulated, including exposure route and safe exposure doses,¹⁰⁷ and meticulously applied for plant nanotechnology. The whole life cycle of these NPs should be evaluated, including their fabrication, storage and distribution, application and potential abuse, and disposal. Bio-accumulation, penetration, and translocation of NPs in plants should be evaluated in detail, specifically for individual NPs used, since each nanomaterial is unique and the results cannot be generalized. This being a cross-disciplinary area of research, appropriate awareness of the positive and negative aspects of nanoparticles is mandatory for the user.

4. Future prospects of nano-integrated plant tissue culture

From the disclosed reports citing the influence of NPs on plant tissue culture, there is no doubt that NPs have a lot to offer in terms of various aspects of plant biotechnology. Beginning from the first phase, through disinfection, to differentiation of the callus, genetic transformation, somaclonal variation and production of secondary metabolites, NPs have demonstrated their positive roles. Having set the stage, this is now just the tip of the iceberg; nanotechnology has more to offer, and it is time to explore and delve more into nano-resources and incorporate them into plant tissue culture. Nanotechnology is a multidisciplinary subject and offers limitless scope in multiple areas. For example, although a wide range of NPs have been reported to possess potent antimicrobial activity, only a few NPs, such as Ag, TiO₂, Zn and ZnO NPs, have been so far predominantly used for controlling microbial contaminants in plant tissue culture. With new-age materials like graphene, graphite, quantum dots, carbon nanotubes, polymer dendrimers and atomic clusters being developed, which all have established bactericidal and fungicidal activities, it is time to plunge deeper into these rich reservoirs of nanotechnology. Who knows? Maybe the best is yet undisclosed (Fig. 4).

Furthermore, as explained in Section 3, the choke in the pipeline is that whilst using these nanomaterials freely in living systems, the nanotoxicity aspects need to be well contemplated and thought upon. Several studies have demonstrated that NP treatment can severely damage explants and adversely affect the regeneration potential of the explants. While the incorporation of NPs into the culture medium eliminates microbial contaminants and hence could be useful for germplasm conservation, their presence in the medium and their possible uptake by the plants needs to be monitored. The effects of NPs on callus induction, organogenesis, shoot multiplication, shoot elongation, and rooting are also not well documented. Additionally, the subculture effects on callus proliferation and shoot multiplication in the presence of NPs need to be studied in detail. Moreover, a few studies have shown that the presence of NPs in the culture medium causes somaclonal variation; therefore, long-term exposure of plant cultures to NPs may negatively affect morphogenesis too. The addition of NPs can lead to enhanced bioactive compound content in cell, shoot and root cultures, possibly due

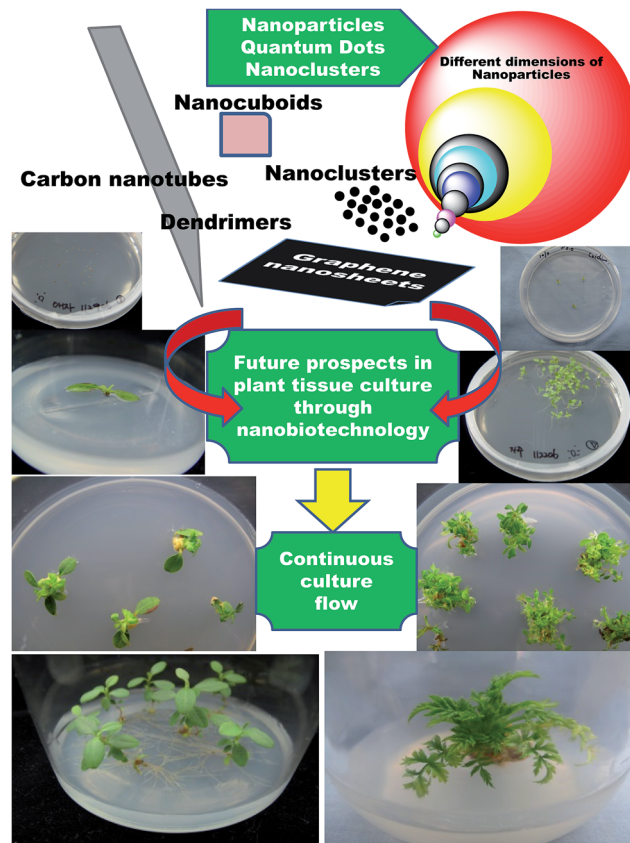


Fig. 4 Future prospects envisioned for nanotechnology in plant tissue culture to scale newer heights through suggested incorporation of new-age revolutionary nanomaterials.

to the production of ROS, activation of antioxidant enzymes and regulation of specific genes; however, the mechanism behind this is still unclear. The influence of various NPs on secondary metabolite production from several important plant species, and the chemical and physical properties and biological activities of compounds obtained from NP-treated plant cultures also need to be investigated in depth. The fate of NPs whilst interacting with plants needs to be fundamentally explored and transparent observations made and disclosed. Without this estimate, no real progress can be made without real time repercussions.

5. Conclusion

Even with all of these undisclosed facts looming in the backdrop as a shadow, there is still no question that plant nanobiotechnology is emerging as a prominent and promising field with excellent potential towards plant improvement. What is required is more targeted research to clarify and streamline the process to harness only the beneficial aspects without exposure to the adverse effects.

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References

- 1 T. Thorpe, *Mol. Biotechnol.*, 2007, **37**, 169–180.
- 2 I. Sivanesan and S. W. Park, *Ind. Crops Prod.*, 2015, **76**, 323–328.
- 3 K. Keren, R. S. Berman, E. Buchstab, U. Sivan and E. Braun, *Science*, 2003, **302**, 1380–1382.
- 4 Q. A. Pankhurst, J. Connolly, S. K. Jones and J. Dobson, *J. Phys. D: Appl. Phys.*, 2003, **36**, R167–R181.
- 5 W. J. Parak, D. Gerion, T. Pellegrino, D. Zanchet, C. Micheel, S. C. Williams, R. Boudreau, M. A. Le Gros, C. A. Larabell and A. P. Alivisatos, *Nanotechnology*, 2003, **14**, R15–R27.
- 6 H. Yan, S. H. Park, G. Finkelstein, J. Reif and T. LaBean, *Science*, 2003, **301**, 1882–1884.
- 7 C. McGovern, *Nanotechnol. Perceptions*, 2010, **6**, 158–178.
- 8 F. Dong, R. T. Koodali, H. Wang and W. K. Ho, *J. Nanomater.*, 2014, 276467, DOI: 10.1155/2014/276467.
- 9 P. Kerativitayanan, J. K. Carrow and A. K. Gaharwar, *Adv. Healthcare Mater.*, 2015, **4**, 1600–1627.
- 10 P. Wang, E. Lombi, F. J. Zjao and P. M. Kopittke, *Trends Plant Sci.*, 2016, **21**, 1–12.
- 11 B. Ruttkay-Nedecky, O. Krystofova, L. Nejdil and V. Adam, *J. Nanobiotechnol.*, 2017, **15**, 33, DOI: 10.1186/s12951-017-0268-3.
- 12 M. H. Siddiqui and M. H. Al-Wahaibi, *Saudi J. Biol. Sci.*, 2014, **21**, 13–17.
- 13 M. Delfani, M. B. Firouzabadi, N. Farrokhi and H. Makarian, *Commun. Soil Sci. Plant Anal.*, 2014, **45**, 530–540.
- 14 F. Torney, B. G. Trewyn, V. S. Lin and K. Wang, *Nat. Nanotechnol.*, 2007, **2**, 295–300.
- 15 H. Oloumi, R. Soltaninejad and A. Baghizadeh, *Indian J. Plant Physiol.*, 2015, **20**, 157–161.
- 16 H. J. Park, S. H. Kim, H. J. Kim and S. H. Choi, *Plant Pathol. J.*, 2006, **22**, 295.
- 17 M. K. Sarmast and H. Salehi, *Mol. Biotechnol.*, 2016, **58**, 441–449.
- 18 C. Leifert, C. E. Morris and W. M. Waites, *Crit. Rev. Plant Sci.*, 1994, **13**, 139–183.
- 19 C. Leifert and A. C. Cassells, *In Vitro Cell. Dev. Biol.: Plant*, 2001, **37**, 133–138.
- 20 C. Leifert, H. Cammota and W. M. Waites, *Plant Cell, Tissue Organ Cult.*, 1992, **29**, 153–160.
- 21 G. A. Teixeira da Silva, T. Duong, T. Michi and F. Seiichi, *Sci. Hortic.*, 2003, **97**, 397–410.
- 22 Y. H. Qin, J. A. T. da Silva, J. H. Bi, S. L. Zhang and G. B. Hu, *Plant Growth Regul.*, 2011, **65**, 183–193.
- 23 E. V. Tambarussi, M. Rogalski, F. T. S. Nogueira, G. E. Brondani, V. F. De Martin and H. Carrer, *Ann. Forest Res.*, 2015, **58**, 177–183.
- 24 L. Wang, C. Hu and L. Shao, *Int. J. Nanomed.*, 2017, **12**, 1227–1249.
- 25 N. Beyth, Y. Hourri-Haddad, A. Domb, W. Khan and R. Hazan, *Evid. base. Compl. Alternative Med.*, 2015, 246012, DOI: 10.1155/2015/246012.
- 26 G. H. Abdi, H. Salehiand and M. Khosh-Khui, *Acta Physiol. Plant.*, 2008, **30**, 709–714.
- 27 M. Fakhrfeshani, A. Bagheri and A. Sharifi, *Journal of Biological & Environmental Sciences*, 2012, **6**, 121–127.
- 28 A. Gouran, M. Jirani, A. A. Mozafari, M. K. Saba, N. Ghaderi and S. Zaheri, Effect of silver nanoparticles on grapevine leaf explants sterilization at *in vitro* conditions, *2nd National Conference on Nanotechnology from Theory to Application*, Isfahan, Iran, 20 February 2014, pp. 1–6.
- 29 N. Mahna, S. Z. Vahed and S. Khani, *J. Nanomed. Nanotechnol.*, 2013, **4**, 161, DOI: 10.4172/2157-7439.1000161.
- 30 A. A. Rostami and A. Shahsavar, *Asian J. Plant Sci.*, 2009, **8**, 505–509.
- 31 M. K. Sarmast, H. Salehi and M. Khosh-Khui, *Acta Biol. Hung.*, 2011, **62**, 477–484.
- 32 M. Arab, M. Yadollahi, A. Hosseini-Mazinani and S. Bagheri, *J. Genet. Eng. Biotechnol.*, 2014, **12**, 103–110.
- 33 T. Murashige and F. Skoog, *Physiol. Plant.*, 1962, **15**, 473–497.
- 34 K. Safavi, M. Esfahanizadeh, D. H. Mortzaeinezahad and H. Dastjerd, *The study of nano silver (NS) antimicrobial activity and evaluation of using NS in tissue culture media. Int Conf Life Sci Technol IPCBEE*, IACSIT Press, Singapore, 2011, vol. 3, pp. 159–161.
- 35 K. Safavi, F. Mortzaeinezahad, M. Esfahanizadeh and M. J. Asgari, *World Acad. Sci. Eng. Technol.*, 2011, **55**, 372–373.
- 36 K. Safavi, Evaluation of using nanomaterial in tissue culture media and biological activity, *2nd International Conference on Ecological, Environmental and Biological Sciences (EEBS'2012)*, Bali, Indonesia, 13–14 October 2012, pp. 5–8.
- 37 K. Safavi, *Bull. Environ., Pharmacol. Life Sci.*, 2014, **3**, 163–166.
- 38 S. Shokri, A. Babaei, M. Ahmadian, S. Hessami and M. M. Arab, *International Journal of Farming and Allied Sciences*, 2014, **3**, 50–54.
- 39 P. Kalsaitkar, J. Tanna, A. Kumbhare, S. Akre, C. Warade and N. Gandhare, *Asian J. Biol. Life Sci.*, 2014, **3**, 167–172.
- 40 M. Mandeh, M. Omid and M. Rahaie, *Biol. Trace Elem. Res.*, 2012, **150**, 376–380.
- 41 M. N. Helaly, M. A. El-Metwally, H. El-Hoseiny, S. A. Omar and N. I. El-Sheery, *Aust. J. Crop Sci.*, 2014, **8**, 612–624.
- 42 M. Taghizadeh and M. Solgi, *J. Hortic. Sci. Technol.*, 2014, **1**, 131–140.
- 43 J. L. Spinoso-Castillo, R. A. Chavez-Santoscoy, N. Bogdanchikova, J. A. Pérez-Sato, V. Morales-Ramos and J. J. Bello-Bello, *Plant Cell, Tissue Organ Cult.*, 2017, **129**, 195–207.
- 44 P. C. Maness, D. M. Smolinski, Z. Blake, E. J. W. Huang and W. A. Jacoby, *Appl. Environ. Microbiol.*, 1999, **65**, 4094–4098.
- 45 M. Heinlaan, A. Ivask, I. Blinova, H. C. Dubourguier and A. Kahru, *Chemosphere*, 2008, **71**, 1308–1316.
- 46 Y. W. Baek and Y. J. An, *Sci. Total Environ.*, 2011, **409**, 1603–1608.



- 47 S. J. Soenen, P. Rivera-Gil, J. M. Montenegro, W. J. Parak, S. C. De Smedt and K. Braeckmans, *Nano Today*, 2011, **6**, 446–465.
- 48 G. Applerot, J. Lellouche, N. Perkas, Y. Nitzan, A. Gedanken and E. Banin, *RSC Adv.*, 2012, **2**, 2314–2321.
- 49 M. Guzman, J. Dille and S. Godet, *Nanomedicine*, 2012, **8**, 37–45.
- 50 A. M. Allahverdiyev, E. S. Abamor, M. Bagirova and M. Rafailovich, *Future Microbiol.*, 2011, **6**, 933–940.
- 51 M. Aghdaei, H. Salehi and M. K. Sarmast, *Adv. Hortic. Sci.*, 2012, **26**, 21–24.
- 52 M. K. Sarmast, A. Niazi, H. Salehi and A. Abolmoghadam, *Plant Cell, Tissue Organ Cult.*, 2015, **121**, 227–236.
- 53 P. Sharma, D. Bhatt, M. G. H. Zaidi, P. P. Saradhi, P. K. Khanna and S. Arora, *Appl. Biochem. Biotechnol.*, 2012, **167**, 2225–2233.
- 54 H. Zafar, A. Ali, J. S. Ali, I. U. Haq and M. Zia, *Front. Plant Sci.*, 2016, **7**, 535, DOI: 10.3389/fpls.2016.00535.
- 55 V. Kumar, P. Guleria, V. Kumar and S. K. Yadav, *Sci. Total Environ.*, 2013, **46**, 462–468.
- 56 E. A. Genady, E. A. Qaid and A. H. Fahmy, *Int. J. Pharm. Res. Allied Sci.*, 2016, **5**, 196–202.
- 57 T. E. Talankova-Sereda, K. V. Liapina, E. A. Shkopinskij, A. I. Ustinov, A. V. Kovalyova, P. G. Dulnev and N. I. Kucenko, *Nanosci. Nanoeng.*, 2016, **4**, 31–39.
- 58 R. Javed, M. Usman, B. Yücesan, M. Zia and E. Gürel, *Plant Physiol. Biochem.*, 2017, **110**, 94–99.
- 59 H. F. Alharby, E. M. R. Metwali, M. P. Fuller and Y. A. Aldhebani, *Arch. Biol. Sci.*, 2016, **68**, 723–735.
- 60 E. A. Ewais, S. A. Desouky and E. H. Elshazly, *Nanosci. Nanotechnol.*, 2015, **5**, 45–56.
- 61 M. Ghorbanpour and J. Hadian, *Carbon*, 2015, **94**, 749–759.
- 62 M. V. Khodakovskaya, K. de Silva, A. S. Biris, E. Dervishi and H. Villagarcia, *ACS Nano*, 2012, **6**, 2128–2135.
- 63 C. Lin, B. Fugetsu, Y. Su and F. Watari, *J. Hazard. Mater.*, 2009, **170**, 578–583.
- 64 X. M. Tan, C. Lin and B. Fugetsu, *Carbon*, 2009, **47**, 3479–3487.
- 65 I. Kokina, E. Siedevskis, V. Gerbreder, D. Grauda, M. Jermaionoka, K. Valaine, I. Gavarâne, I. Pigiòka, M. Filipoviès and I. Rashal, *Proc. Latv. Acad. Sci., Sect. B*, 2012, **66**, 200–209.
- 66 Y. Syu, J. H. Hung, J. C. Chen and H. Chuang, *Plant Physiol. Biochem.*, 2014, **83**, 57–64.
- 67 Z. Poborilova, R. Opatrilova and P. Babula, *Environ. Exp. Bot.*, 2013, **91**, 1–11.
- 68 C. Ma, S. Chhikara, B. Xing, C. Musante, J. C. White and O. P. Dhankher, *ACS Sustainable Chem. Eng.*, 2013, **1**, 768–778.
- 69 C. Ma, H. Liu, H. Guo, C. Musante, S. H. Coskun, B. C. Nelson, J. C. White, B. Xing and O. M. Dhankher, *Environ. Sci.: Nano*, 2016, **3**, 1369–1379, DOI: 10.1039/c6en00189k.
- 70 S. Anwaar, Q. Maqbool, N. Jabeen, M. Nazar, F. Abbas, B. Nawaz, T. Hussain and S. Z. Hussain, *Front. Plant Sci.*, 2016, **7**, 1330, DOI: 10.3389/fpls.2016.01330.
- 71 L. Giorgetti, M. M. Castiglione, M. Bernabini and C. Geri, *Agrochimica*, 2011, **55**, 45–53.
- 72 I. Kokina, V. Gerbreder, E. Siedevskis and A. Bulanovs, *J. Biotechnol.*, 2013, **165**, 127–132.
- 73 H. Fazal, B. H. Abbasi, N. Ahmad and M. Ali, *Appl. Biochem. Biotechnol.*, 2016, **180**, 1076–1092.
- 74 S. Bansod, M. Bawskar and M. Rai, *IET Nanobiotechnol.*, 2015, **9**, 239–245.
- 75 P. S. Vijayakumar, O. U. Abhilash, B. M. Khan and B. L. V. Prasad, *Adv. Funct. Mater.*, 2010, **20**, 2416–2423.
- 76 K. Pasupathy, S. Lin, Q. Hu, H. Luo and P. C. Ke, *Biotechnol. J.*, 2008, **3**, 1078–1082.
- 77 S. Naqvi, A. N. Maitra, M. Z. Abdin, M. Akmal, I. Arors and M. Samim, *J. Mater. Chem.*, 2012, **22**, 3500–3507.
- 78 M. R. S. Ardekani, M. Z. Abdeni, N. Nasrullaah and M. Samim, *Int. J. Pharm. Sci.*, 2014, **6**, 605–609.
- 79 F. P. Chang, L. Y. Kuang, C. A. Huang, W. N. Jane, Y. Hung, Y. C. Hsing and C. Y. Mou, *J. Mater. Chem. B*, 2013, **1**, 5279–5287.
- 80 Y. Hao, X. Yang, Y. Shi, S. Song, J. Xing, J. Marowitch, J. Chen and J. Chen, *Botany*, 2013, **91**, 457–466.
- 81 M. W. Bairu, A. O. Aremu and J. Van Staden, *Plant Growth Regul.*, 2011, **63**, 147–173.
- 82 I. Sivanesan and B. R. Jeong, *Plant Cell, Tissue Organ Cult.*, 2012, **111**, 247–253.
- 83 B. R. Jeong and I. Sivanesan, *Plant Cell, Tissue Organ Cult.*, 2015, **123**, 607–618.
- 84 D. H. Atha, H. Wang, E. J. Petersen, D. Cleveland, R. D. Holbrook, P. Jaruga, M. Dizdaroglu, B. Xing and B. C. Nelson, *Environ. Sci. Technol.*, 2012, **46**, 1819–1827.
- 85 P. Landa, S. Prerostova, S. Petrova, V. Knirsch, R. Vankova and T. Vanek, *Environ. Sci. Technol.*, 2015, **49**, 14537–14545.
- 86 D. K. Tripathi, S. Shweta, S. Singh, R. Pandey, V. P. Singh, N. C. Sharma, S. M. Prasad, N. K. Dubey and D. K. Chauhan, *Plant Physiol. Biochem.*, 2017, **110**, 2–12.
- 87 I. Kokina, I. Mickevica, M. Jermalonoka, L. Bankovska, V. Gerbreder, A. Ogurcovs and I. Jahundovica, *Int. J. Genomics*, 2017, 1676874, DOI: 10.1155/2017/1676874.
- 88 M. Hussain, S. Fareed, S. Ansari, M. Rahman, I. Z. Ahmad and M. Saeed, *J. Pharm. BioAllied Sci.*, 2012, **4**, 10–20.
- 89 I. Sivanesan, R. K. Saini and D. H. Kim, *Ind. Crops Prod.*, 2016, **83**, 31–38.
- 90 I. Sivanesan, R. K. Saini, R. Noorzai, A. J. Zamany and D. H. Kim, *3 Biotech.*, 2016, **6**, 91.
- 91 H. K. Mohammed Al-oubaidi and A. S. Mohammed-Ameen, *Int. J. Pharmacol. Ther.*, 2014, **5**, 267–272.
- 92 H. K. Mohammed Al-oubaidi and N. M. Kasid, *World J. Pharm. Res.*, 2015, **4**, 1791–1799.
- 93 E. Chamani, S. K. Ghalehtaki, M. Mohebodini and A. Ghanbari, *Iranian Journal of Genetics and Plant Breeding*, 2015, **4**, 11–19.
- 94 C. V. Desai, H. B. Desai, K. P. Suthar, D. Singh, R. M. Patel and A. Taslim, *Appl. Biol. Res.*, 2015, **17**, 1–7.
- 95 B. Zhang, L. P. Zheng, W. Y. Li and J. W. Wang, *Curr. Nanosci.*, 2013, **9**, 363–370.
- 96 B. Ghasemi, R. Hosseini and F. D. Nayeri, *Turk. J. Bot.*, 2015, **39**, 769–777.



- 97 Z. Shakeran, M. Keyhanfar, G. Asghari and M. Ghanadian, *Turk. J. Biol.*, 2015, **39**, 111–118.
- 98 M. Raei, S. A. Angaji, M. Omid and M. Khodayari, *Int. J. Biosci.*, 2014, **5**, 74–81.
- 99 P. Bhat and A. Bhat, *J. Exp. Sci.*, 2016, **7**, 1–6.
- 100 M. Jamshidi, F. Ghanati, A. Rezaei and E. Bemani, *Cytotechnology*, 2016, **68**, 525–530.
- 101 F. Moharrami, B. Hosseini, A. Sharafi and M. Farjaminezhad, *In Vitro Cell. Dev. Biol.: Plant*, 2017, **53**, 104–111.
- 102 P. Miralles, L. T. Church and A. T. Harris, *Environ. Sci. Technol.*, 2012, **46**, 9224–9239.
- 103 G. Chichiricò and A. Poma, *Nanomaterials*, 2015, **5**, 851–873, DOI: 10.3390/nano5020851.
- 104 O. Zaytseva and G. Neumann, *Chem. Biol. Technol. Agric.*, 2016, **3**, 17, DOI: 10.1186/s40538-016-0070-8.
- 105 J. Yang, W. Cao and Y. Rui, *J. Plant Interact.*, 2017, **12**(1), 158–169.
- 106 S. Mishra, C. Keswani, P. C. Abhilash, L. F. Fraceto and H. B. Singh, *Front. Plant Sci.*, 2017, **8**, 471, DOI: 10.3389/fpls.2017.00471.
- 107 B. Ruttkay-Nedecky, O. Krystofova, L. Nejdil and V. Adam, *J. Nanobiotechnol.*, 2017, **15**, 33, DOI: 10.1186/s12951-017-0268-3.
- 108 C. W. Lee, S. Mahendra, K. Zodrow, D. Li, Y.-C. Tsai, J. Braam and P. J. J. Alvarez, *Environ. Toxicol. Chem.*, 2010, **29**, 669–675.
- 109 S. Lin, J. Reppert, Q. Hu, J. S. Hudson, M. L. Reid, T. A. Ratnikova, A. M. Rao, H. Luo and P. C. Ke, *Small*, 2009, **5**, 1128–1132.

