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Plasma-digital nexus: plasma nanotechnology for digital manufacturing age

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Abstract: Digital transformation in manufacturing is one of the key megatrends in the development of global economy and society. Three-dimensional (3D) printing is a transformative digital technology poised to disrupt manufacturing and supply chains across major industries. Here we critically examine relevant insights into current and emerging applications of plasma nanotechnology in printing, including 3D printing. Plasma devices operated at atmospheric pressure coupled with printing processes may help strengthen 3D printing as an emerging fabrication technology that morphs diverse metal powders, polymers, plastics and other materials into digitally designed 3D shapes and patterns. We discuss how plasma applications may help overcome current limitations of 3D printing in various fields, e.g. limitations of sculpting composite materials, lack of mechanical strength and the need for post-processing. Our key focus is on the challenges, opportunities and physical mechanisms of the use of 3D printing in nano-manufacturing, defined as the fabrication of nanoscale building blocks, such as nanoparticles and nanomaterials; their assembly into higher-order (micro-scale) structures; and the integration of these structures into larger (macro-) scale devices and systems by controlling energy and matter at nanoscale. Moreover, we discuss the physico-chemical mechanisms that result in highly-conformal deposition of nanostructured materials onto 3D surfaces with microscopic (and possibly nanoscale) control of textures and inter-layer cross-linking, without the need for additional heating. We further highlight the arising opportunities for plasma nanotechnology to synergize with the emerging digital transformation platforms in surface micro- and nano-structuring using polymers, metals, metallic alloys, and other materials. These new findings in plasma-digital nanoscale fabrication may lead to a new digital manufacturing platform suitable for a number of cutting-edge applications in electronic, sensing and energy devices.

Keywords: Plasma printing, Plasma nanotechnology, Additive manufacturing, Digital technologies

1. Introduction

1.1. Digital manufacturing age

Our daily life significantly depends upon digital manufacturing, such as computer-aided design, robotics control, smart sensing and process simulation, which has tremendously accelerated growth and innovation in the traditionally linear and time-consuming manufacturing process (Fig. 1) (Chong, *et al.*, 2018).

Additive manufacturing (AM) is set to disrupt conventional manufacturing (Mohr and Khan, 2015) and supply chains in virtually all industrial sectors, completely changing the future landscape. Tunability of mechanical, electrical and other functional properties and expansion of the range of printable materials, coupled with programmable control of their composition and architecture through the various length scales (Truby and Lewis, 2016), drives innovation in a large number of applications. For example, biomimetic composites and shape-morphing systems (Gladman, *et al.*, 2016; Studart, 2016) for soft sensors and robotics applications can be produced by additive manufacturing (Truby and Lewis, 2016).

However, at the same time, there is a large gap in the existing technologies which needs to be overcome in order to reach the full potential of additive manufacturing. Lack of crosslinking properties between printed individual layers, high residual stresses and pores in 3D printed metal parts are common issues that usually require thermal, mechanical or electro-chemical post processing in order to improve the mechanical performance of the printed object (Bahnini, *et al.*, 2018). Even then, in micro- and nano-scale manufacturing, additive manufacturing is lacking in quality in comparison to conventional nanoscale fabrication (nanofabrication) techniques.

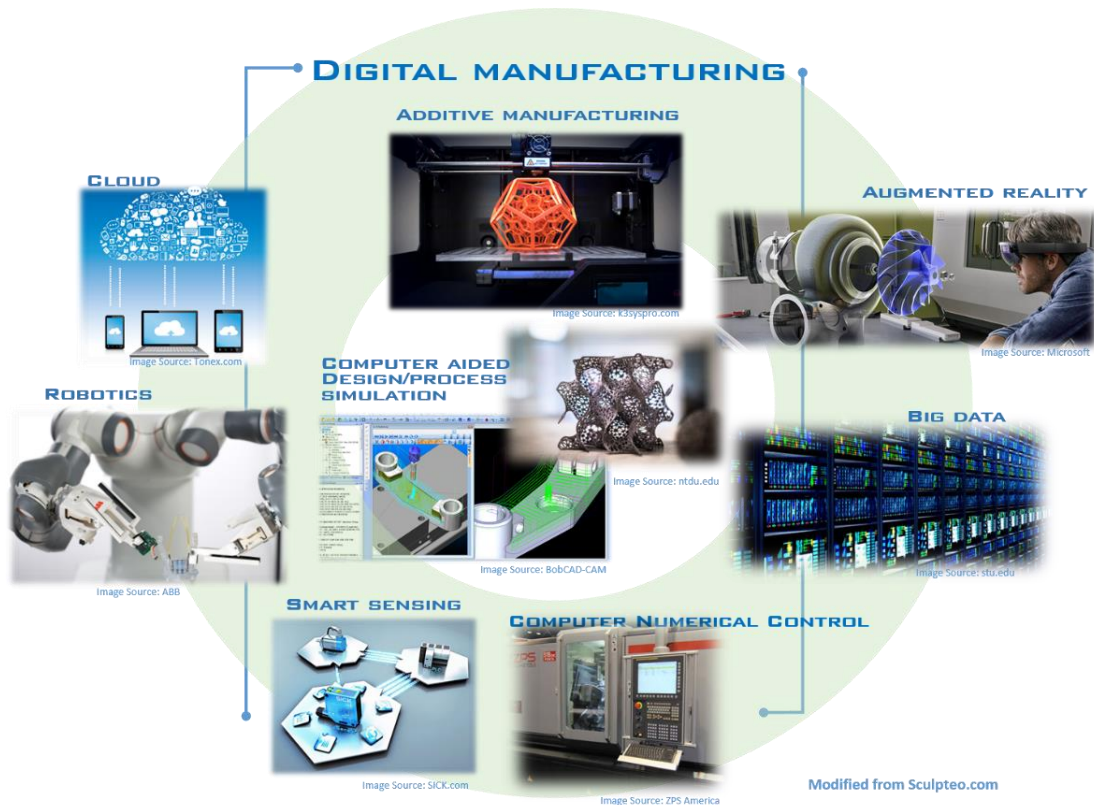


Fig. 1. Elements of digital manufacturing for future industry

In order to overcome the current limits of 3D printing technologies and meet the inspiring goal of creating architected matter that has qualitatively new properties with unprecedented control over the material's composition, structure, function and dynamics (Hirt, et al., 2017), more effective control factors are required. Potentially, low-temperature plasmas may provide a solution to these issues. Plasma has played a significant role in enabling the innovation of current high-end manufacturing processes, such as in semiconductor and display industries. Precisely controlled physical/chemical reactions involving high density electrons, energetic ionic species and various chemically reactive neutral radicals have made plasma an indispensable tool to the microelectronics industry, where one third of the manufacturing steps of microelectronic circuits and microprocessors are plasma-based (d'Agostino, *et al.*, 2005). In addition, due to the wide range of plasma characteristics – from the high-temperature, high-energy application of nuclear fusion processes to low temperature atmospheric-pressure surface modification – plasma technologies have shaped everyday lives through the numerous manufacturing processes they are employed in (John, 2005; Weltmann, *et al.*, 2019).

More recently, plasmas have contributed to the development of additive manufacturing. Examples of this include the commercialized thermal plasma system for nanopowder synthesis in metals, metal alloys and ceramics (Essentium; Tekna); and the low temperature plasma source combined with commercial polymer-based 3D printing to provide improved mechanical properties (Essentium; Innophysics; Plasma). Figure 2 summarises possible benefits of plasmas for additive manufacturing, particularly in nano- and microscale processes. This will be discussed in more detail in the following sections.

The non-equilibrium, low temperature plasmas are particularly suited to processes requiring temperature sensitive substrates or 3D objects to be processed at atmospheric pressure. Furthermore, being able to supply numerous chemically reactive species in a controlled manner meets the requirement for '*unprecedented control over the material's composition*' thanks to Plasma Enhanced Chemical Vapour Deposition (PECVD) technology.

The PECVD processes, which produce micro- and nano-scale films from gas, liquid vapour or powder precursors, can be controlled to create the required chemical/physical properties in the material. For example, a conformal multilayered thin-film structure can be deposited and printed on a 3D object. This technique could potentially transform many current additive manufacturing processes. The additional control using plasma electric fields is possible by changing the electrode configurations, both in the plasma PVD (Physical Vapour Deposition) and CVD. This enhancement can lead to the products with precisely customized physical properties as the exposure of the material to the energetic ionic species can be controlled. Furthermore, post-processing may not be required, as is often the case for laser processed film or 3D printed objects. Another advantage is the cost effectiveness in comparison to typical laser systems, making PECVD a promising technique as an assisting module for existing 3D printing systems or as a stand-alone plasma printing system.

In the following, we will discuss some of the recent developments in the digital manufacturing and critically examine the areas where the use of plasmas may contribute to the future development of this rapidly emerging technological field.

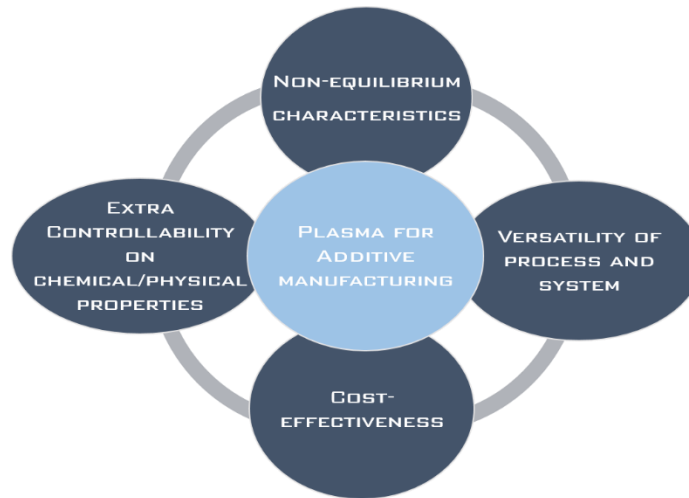


Fig. 2 Possible benefits of plasma processes for additive manufacturing

1.2. Printing and plasma: a synergy for digital manufacturing

The idea of adapting plasma technology for the printing process was proposed in early 2000s. Microplasmas using a dielectric barrier sandwiched between two electrodes enabled precise gravure printing with the 50-70 μm linewidth (Fig. 3) (Thomas, *et al.*, 2012). Using this technique, a desired pattern can be engraved in the conductive electrode surface or dielectric layer. It can further support Roll-to-Roll (R2R) continuous plasma processing on a flexible polymer film. The focus has mainly been towards surface modification to enhance hydrophilicity prior to the actual printing process. Using C_2H_2 mixed as a reactive gas, the ‘plasma stamping’ technique has been demonstrated to deposit a patterned polymer film with the minimum linewidth of 50 μm . However, the thickness of the printed layer was not uniform due to the inhomogeneous nature of the porous metal electrode through which the gas was transported.

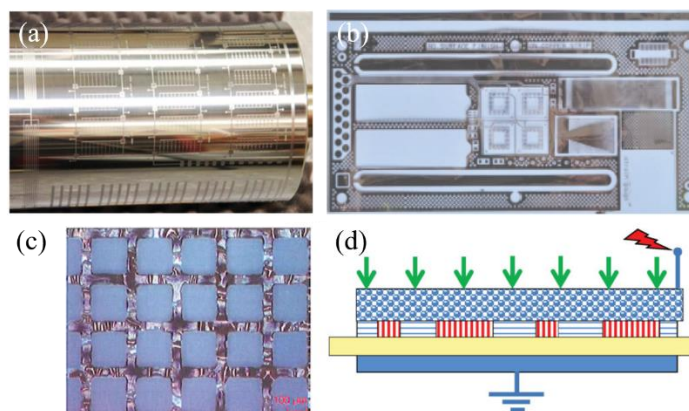


Fig. 3 (a) Printer roller showing the engraved pattern, (b) plasma printed and subsequently electroless copper metallized model structure with a line width as low as 70 μm , (c) plasma printed grid of an amorphous hydrogenated carbon film, obtained from 1% C_2H_2 in He, using a porous sinter metal electrode to provide gas supply and (d) schematic of plasma printing with gas-fed microplasmas using an electrode made from sintered metal powder. Vertical arrows indicate the gas flow (Thomas, *et al.*, 2012)

Significant efforts have been made to create flexible printing methods using atmospheric pressure plasmas without the limitations associated with fixed pattern or electrode configuration. Recently, as the growth of additive manufacturing has accelerated, the contribution from the plasma technology has also increased. Plasma printing of multiwalled carbon nanotubes was demonstrated on paper substrates using an atmospheric pressure plasma jet (Gandhiraman, *et al.*, 2016). The plasma-printed carbon nanotubes showed denser morphology and higher conductivity in comparison to non-plasma processed samples. The printed nanotubes showed good performance for a dopamine and ammonia sensing. The same group also reported the plasma jet printing and *in situ* reduction of graphene oxide (Dey, *et al.*, 2018) using He/H₂ gas, thus revealing the possibility to control chemical properties through the use of plasmas. This approach enabled removal of the multiple intermediate steps of chemical reduction and washing which are usually required to increase the conductivity of graphene oxide films.

Another significant advantage of using plasmas is the enhanced mechanical properties resulting from the improved chemical bond strength. It was shown that plasma can ‘weld’ different sheets of porous 3D graphene scaffolds to create strong intermolecular bonding (Chakravarty, *et al.*, 2016). The plasma welded scaffolds (Fig. 4) were produced to maintain a high porosity and ultra-low density with an interconnected, layered, open microstructure which had high yield strength and stiffness values (Chakravarty, *et al.*, 2016).

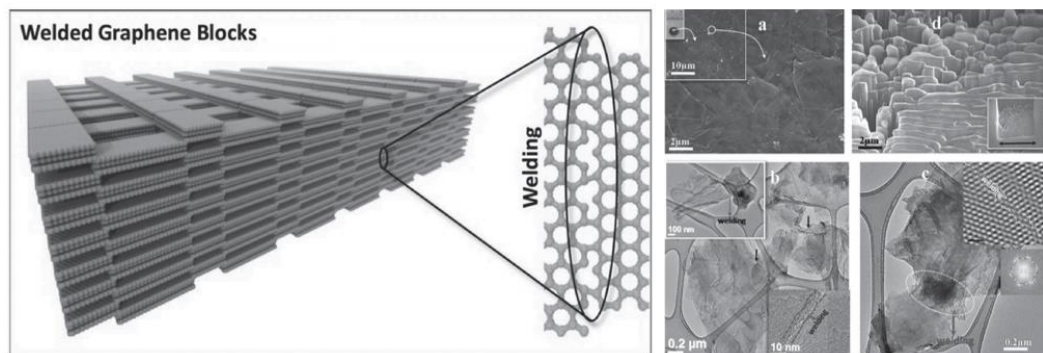


Fig. 4 (Left) Schematic representing the stacking of 2D graphene sheet (Right) (a) SEM image of the top surface of graphene sample at different magnifications (digital image and SEM image at low magnification as insets) (b) Bright field transmission electron microscopy image of graphene oxide showing randomly oriented sheets welded together (top left inset shows two sheets welded at the edge; inset at the bottom on right shows HRTEM image of the welded region) (c) Bright field TEM image of side-edge-welded graphene sheets, HRTEM image of the welded region showing defect, and FFT of the region showing misorientation of the two sheets; the line in inset corresponds to 0.5 nm distance and d) 50° tilted SEM image of the 3D architecture of the plasma welded sample by SPS (spark plasma sintering) technique observed after FIB at the center of the sample (shown as inset) (Chakravarty, *et al.*, 2016)

Researchers at the Fraunhofer Institute presented a 3D printing system combined with the plasma coating unit for bone implants (Fig. 5) (Fraunhofer, 2018). Using a plasma jet, it was possible to fabricate a cell growth-promoting coating functionalized with amino groups and apply these coatings to the interior and exterior surfaces of the implants.

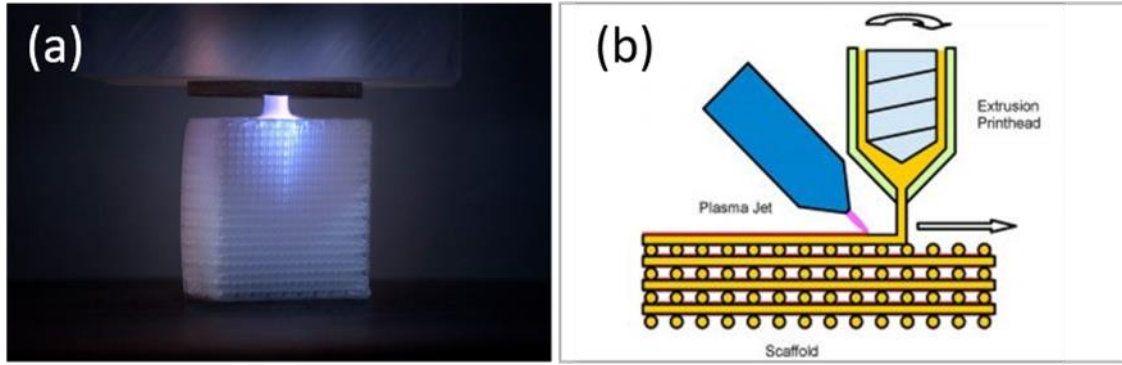


Fig. 5. (a) Plasma-jet coating of medical implant scaffolds (b) Schematic shows how to manufacture coated scaffolds: immediately after extrusion, the scaffold structures are treated with a cold plasma (Image source: Fraunhofer IST)

In recent years, there have appeared a few commercial 3D printer products which have been given the name ‘plasma printer’ as shown in Fig. 6. These printers are equipped with an additional plasma module for surface modification, welding and plasma assisted deposition of metal oxide film. These printers represent a promising platform to fabricate functionalized device elements by surface coating, scribing or etching of single/multiple layers and detailed discussion will be provided in section 5.1 and 5.3.

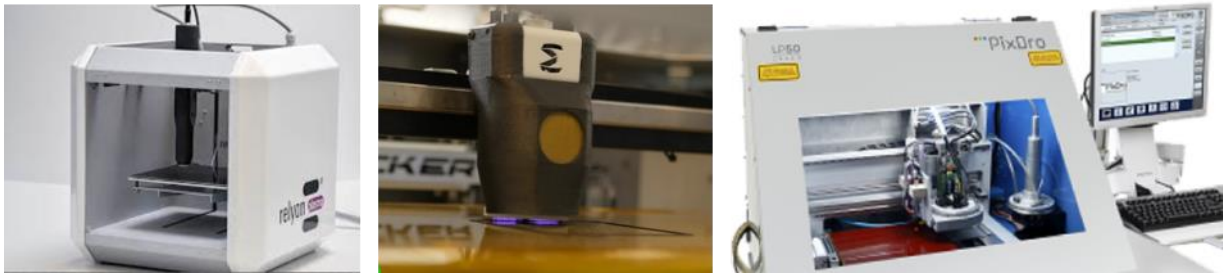


Fig. 6 Examples of commercial plasma printers. Image Sources (from left to right): Relyon Plasma, Essentium3d and Innophysics

To identify where and how plasma technology can potentially benefit the existing and future digital manufacturing platforms, we will first overview the selected current manufacturing processes.

2. Representative printing technologies

2.1 2D printing

To obtain a better insight on printing technology it is worthwhile to review some of the common 2D printing processes as a comparison to 3D manufacturing. Lithography is a common method for obtaining a pattern in layers. This technology helped establish modern microelectronics based industry and emerging internet of things (IoT) technologies. However, we will focus on conventional printing which can produce written patterns. Fig. 7 shows schematics of some examples of printing technology on a flat substrate: screen printing, gravure printing, inkjet printing and high-resolution electrohydrodynamic jet

printing (Park, *et al.*, 2007). As shown in Fig. 7 (a), in the screen printing process, the ink transfers through the patterned mesh onto a substrate by squeegee. In gravure printing, a pattern is etched onto a metal cylinder. The paper (or another substrate) is pressed against the inked cylinder on a rotary press, transferring the image directly to the substrate. Although the resolution of screen printing and gravure printing prevents these techniques from producing high-end products, they are still widely used techniques and the possibility of adapting plasma treatment to improve the printing quality through the increased wetting and adhesion properties has been explored.

Aside from the conventional office and household uses, inkjet printing has been widely used as an effective tool for the fabrication of flexible electronics, optical and photonic devices and more (Alaman, *et al.*, 2016). This is because of the attractive features of inkjet printing that support the additive operation. Indeed, this technique is able to pattern directly regardless of the type of substrates, even on biological materials that are incompatible with the established patterning methods such as photolithography. Further benefits of inkjet printing include the flexibility in structural design due to software-based control, compatibility with large-area substrates, and the potential for low-cost operation (de Gans, *et al.*, 2004; Forrest, 2004; Parashkov, *et al.*, 2005; Park, *et al.*, 2007). Inkjet printing is able to process not only organic materials, but also conductive inks such as metal nanoparticles (Magdassi, *et al.*, 2003; Lee, *et al.*, 2005), single-wall/multi-wall carbon nanotubes (SWCNTs/MWCNTs) (Wei, *et al.*, 2007; Mabrook, *et al.*, 2009) and graphene (Wang, *et al.*, 2016a).

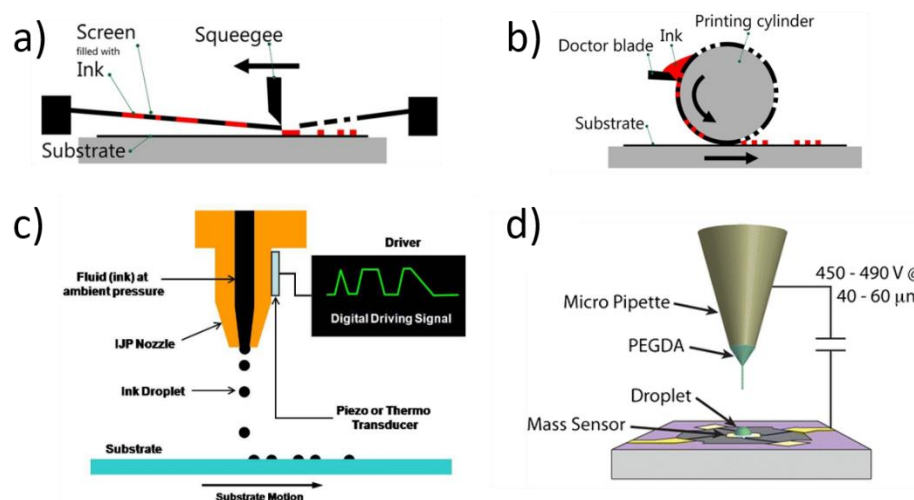


Fig. 7. Schematics showing working principle of (a) Screen printing (EMPA) (b) Gravure printing (EMPA) (c) inkjet printing (Xu, *et al.*, 2007) and (d) electrohydrodynamic printing where PEGDA indicates poly(ethylene glycol) diacrylate as an example of a printable material (Corbin, *et al.*, 2013)

However, the minimum achievable linewidth is typically in the range of 20–30 μ m (Szczech, *et al.*, 2002; Ling and Bao, 2004; Sanaur, *et al.*, 2006) due to the large droplet diameters of \sim 10–20 μ m with the placement errors typically \pm 10 μ m (Sirringhaus, *et al.*, 2000; Cheng, *et al.*, 2005). In order to

improve the printing resolution, various complementary methods have been suggested. Lithographically predefined assisting features (or simple masks) (Sirringhaus, *et al.*, 2000; Stutzmann, *et al.*, 2003; Wang, *et al.*, 2006) or surface functionalization significantly improved printing accuracy and resolution down to the sub μm scale (Sele, *et al.*, 2005) by enabling patterning of wettability of the surface (Stutzmann, *et al.*, 2003; Sele, *et al.*, 2005; Wang, *et al.*, 2006). However, these methods require additional patterning systems and processing steps which limits the benefits of the flexible inkjet printing technique.

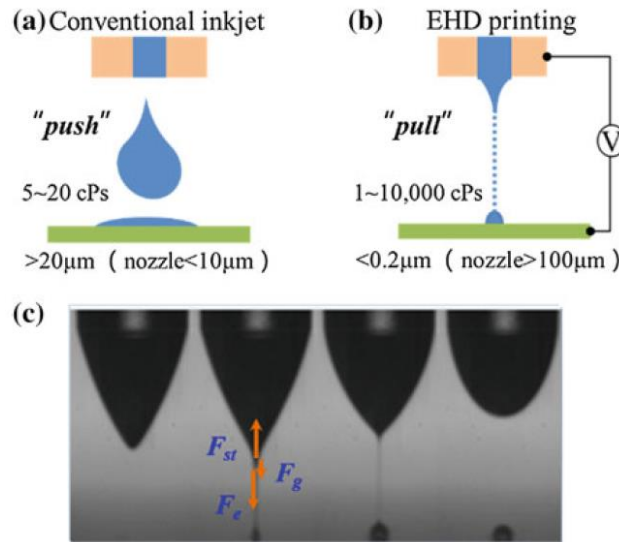


Fig. 8 Schematic of (a) conventional inkjet printing (b) EHD printing and (c) the liquid meniscus at the nozzle tip and the evolution of the shape of a Taylor cone during the EHD printing (Yin, *et al.*, 2018)

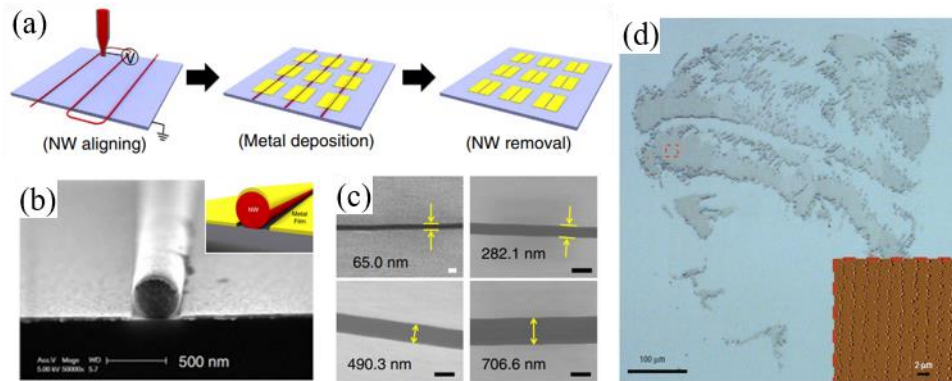


Fig. 9. (a) Schematic illustration of the process for micro/nano channel fabrication by using electrospun fiber as a template – NW (Nanowire) (b) The cross-section of the electrospun fiber after metal deposition (c) Nanochannels generated by removing the electrospun fiber (Min, *et al.*, 2013) (d) High-resolution EHD printing with printed feature sizes in the range from 240 nm to 5 μm . Optical micrograph of a portrait of the ancient scholar, Hypatia, printed using a polyurethane ink and a 500-nm-internal-diameter nozzle. The diameters of the dots are 490 nm. The inset shows an Atomic Force Microscopy (AFM) image of the printed dots (Park, *et al.*, 2007)

On the other hand, electrohydrodynamic (EHD) printing has the same advantage of the flexible writing in one step (as in inkjet printing) whilst also significantly improving the sub-micrometre scale resolution due to the effective control mechanism which relies on the electric field, as shown in Fig. 7

(d). In contrast to ‘push’ methodologies of typical inkjet processes, the electric potential in EHD printing induces a ‘pull’ into the process to confine the jetted ink flow, as shown in Fig. 8. When the gravity force F_g , surface tension force F_{st} and electric field force F_e are balanced, a liquid droplet forms in a stable shape and surface deformation is then triggered by further increase of the electric field. When the critical potential is reached, the repulsive electrical forces overcome the surface tension forces, and the liquid cone is distorted into a Taylor cone which forms the liquid jet. Emerging from the capillary, the jet is maintained at a high potential and can be disintegrated into droplets in various modes, depending on the applied voltage and the flow rate of the ink (Yin, *et al.*, 2018).

Figure 9 (a) - (c) shows an example of EHD printing combined with a lithography process to pattern nanowires for fabricating a shadow mask for large-scale flexible electronics applications. Using the EHD printing technique, organic nanowires are printed followed by metal deposition. The organic nanowires are then removed either by sonication in a solvent or by using an adhesive tape to detach them (Min, *et al.*, 2013). Figure 9(d) presents another example of one-step EHD printing for a graphic application using a nozzle of 500 nm in diameter to produce high resolution printing with individual dots of the similar size (Park, *et al.*, 2007). The use of an electric field for EHD printing significantly improves the printing resolution in comparison to other techniques. As the plasma also features electric fields, it may be possible to combine the EHD printing process with plasmas to further enhance it. With a relatively low AC voltage (0.5 – 3 kV) in a pulsed mode, it is possible to operate the process in a ‘drop-on-demand’ mode. Indeed, as the applied voltage increases, electrospinning (1 – 15 kV) and electrospraying (10 – 30 kV) can take place (Yin, *et al.*, 2018)

2.2 3D printing

2.2.1 Polymers

To enable 3D objects to be printed in an efficient and precise manner, many different fabrication techniques have been developed based on the existing 2D printing techniques. Figure 10 shows common polymer-based 3D printing methods. As shown in Fig. 10 (a) stereolithography (SLA) uses photocurable resin, which is polymerized layer by layer using a laser controlled by the raster image. Figure 10 (b) shows selective laser sintering (SLS) of polymeric powders in the powder bed. In order to facilitate spreading, granulated powders are used with typical diameters between 10 μm and 100 μm . As a result, the possible minimum feature size from SLS is usually a few times larger than what is possible in the SLA process (Truby and Lewis, 2016).

It is important to note that these processes support multiple material patterning in a single sequence because of the difficulties in manipulating the composition of a liquid photopolymer reservoir or powder bed during the printing (Choi, *et al.*, 2010). Ink-based printing methods, for example fused deposition

modelling (FDM), inkjet printing and direct ink writing (DIW), can however easily be used for multi-material 3D printing (Truby and Lewis, 2016).

In comparison to light-based printing, such as SLS or SLA, the resolution of ink-based 3D printing may appear to be limited. However, considering the flexibility of the process and the range of printable materials, especially in multiple material printing, ink-based processes can be the only practical solution in some cases. UV light can be used as an assisting tool for polymerization of photocurable resin whilst the resin is printed by the inkjet process, as shown in Fig. 10 (c). Figure 10 (d) shows a typical low-cost 3D printer using a thermoplastic filament in a technique known as fused deposition modelling (FDM). In this process, thermoplastic filaments are fed through a hot extrusion nozzle and begin to solidify, building a printed layer below their glass transition temperature (Zein, *et al.*, 2002). The polymer filaments used in this process can be filled with functional particles, such as carbon black, to enhance the functionality of the printed parts. (Farahani, *et al.*, 2016) This idea can be expanded to other types of nanomaterials for various applications. An important alternative to FDM printing is direct ink writing (DIW), as shown in Fig. 10 (e) (Lewis, 2006). Amenable for a range of structural (Clausen, *et al.*, 2015) and electrical materials (Frutiger, *et al.*, 2015) to biological materials (Kolesky, *et al.*, 2014), DIW is considered to have the broadest spectrum of printable materials. It is also possible to print multi-materials using DIW, either using multiple printheads equipped with different ink compositions, or by using microfluidic printheads with the embedded control of switching, mixing, core-shell printing, or printing multiple filament arrays etc. (Truby and Lewis, 2016). Mixing nozzles may allow adaptation of plasma-assisted printing as they enable tuneable gradients of mechanical, conductive or other material properties in the printed materials (Ober, *et al.*, 2015).

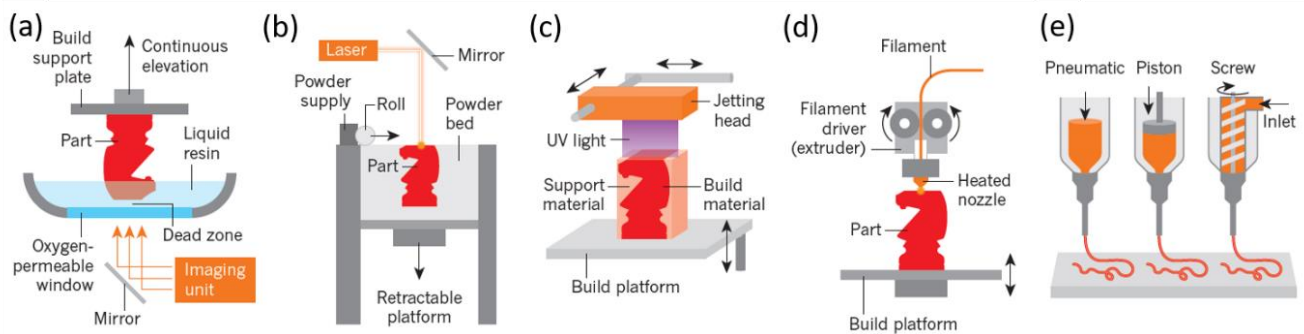


Fig. 10 Common light- and ink-based 3D printing methods. (a) The light-based 3D printing method known as continuous liquid interface production (b) Light-based selective laser sintering of powders (c) Light- and ink-based photocurable inkjet printing of photopolymerizable resins (d) Ink-based fused deposition modelling of thermoplastic filaments (e) Direct ink writing using viscoelastic inks. (Truby and Lewis, 2016)

Figure 11 shows some examples of shape-morphing systems, often called 4D printing, which reveal the benefit of combining programmed responsiveness with additive manufacturing. Responding to heat, light or moisture, the printed object can change its shape and properties over time; this can be useful in

smart textiles (Hu, *et al.*, 2012), biomedical devices (Randall, *et al.*, 2012) and robotic (Felton, *et al.*, 2014) systems.

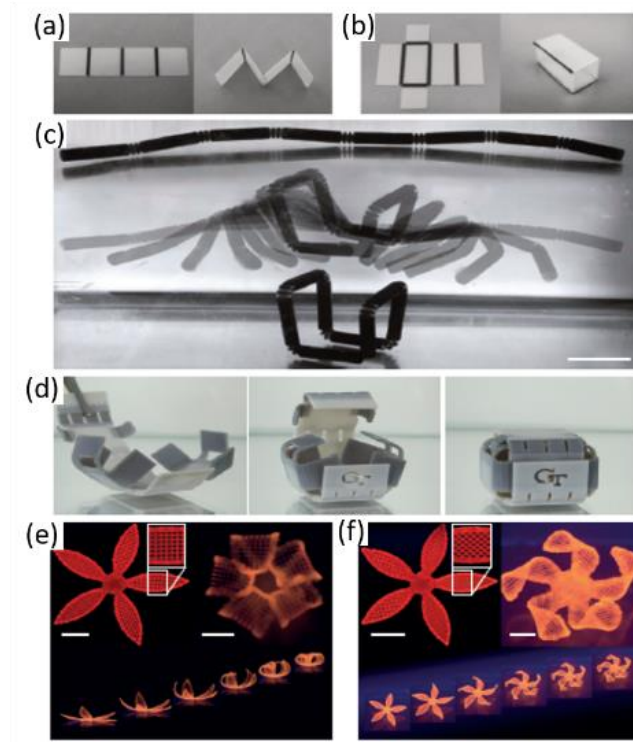


Fig. 11 Stimuli-responsive, morphing architectures (a) and (b) Prestrained polystyrene substrate with inkjet-printed hinges made of carbon black ink which autonomously folds into a 3D shape when illuminated with infrared light (scale bars, 10 mm; adapted from (Liu, *et al.*, 2012)) (c) 4D-printed composite with swellable hinges (top) that self-assembles from a linear into a box-like structure (bottom) when immersed in water (scale bar, 5 cm; adapted from (Tibbits, 2014)) (d) A 4D-printed unfolded box composed of shape-memory polymers that folds back into its original conformation when immersed in warm water (adapted from (Mao, *et al.*, 2015)) (e) and (f) Biomimetic 4D printing of hydrogel composites containing anisotropic cellulose fibrils that orient along the printing direction. They undergo anisotropic swelling to programmably change shape when immersed in water. The printed bilayer lattices transform into flowers, whose petals either bend or twist when the bilayer orientations are $90^\circ/0^\circ$ (e) or $-45^\circ/45^\circ$ (f) (scale bars, 5 mm; insets, 2.5 mm; adapted from (Gladman, *et al.*, 2016))

2.2.2 Metals

Among the most established methods for additive manufacturing of metals, are selective laser melting (SLM) and electron beam melting (EBM), based on the local fusion of metal particles to form a solid layer or object, as shown in Fig. 12 (a) (Murr, *et al.*, 2012; Herzog, *et al.*, 2016; Hirt, *et al.*, 2017). Although SLM and EBM technologies play a major role in commercialized AM at the macroscale, they are not considered to be the best solution for the nano- and micro-scale, as these techniques typically produce the minimum line width of several tens of micrometers (Exner, *et al.*, 2008; Vaezi, *et al.*, 2013). In comparison to SLM, which requires a powder bed, in the laser metal deposition (LMD) process the metallic materials, in the form of a wire or powder, are fed into the laser area through a separate nozzle as shown in Fig. 12(b). This is advantageous as LMD can produce composites or functionally graded materials efficiently. In this process, various materials are placed in different powder or wire feeders and delivered simultaneously or sequentially through the coaxial nozzles located beside the laser outlet. This

produces a composite with flexible composition relative to its location within the manufactured part (Mahamood, 2018).

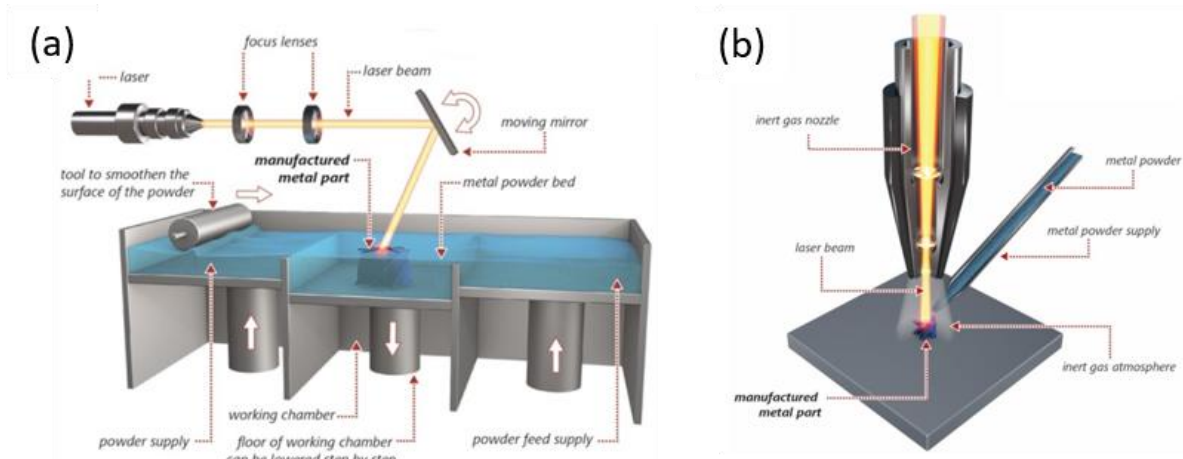


Fig. 12 Schematics for comparison of (a) SLM and (b) LMD technique. Image source <https://www.empa.ch/web/coating-competence-center>

As with 3D printing of polymers, direct ink writing (DIW) can be a suitable and simple method for microscale 3D printing of metallic materials. For example, several tens of nm sized nano-silver ink is used, dispersed in water and ethylene glycol. As shown in Fig. 13(a), to obtain a desirable surface morphology, high temperature post annealing processes are required which can limit the choice of substrate. Recently, a low power laser annealing technique was developed, demonstrating *in situ* annealing of the nanoparticles immediately after exiting the nozzle. This process resulted in the formation of a free-standing 3D structure with improved stability, further validating this technique. This process has some benefits over the electrohydrodynamic printing.

Fig. 13 (b) shows a schematic and a printed example of the EHD printing. As discussed in section 2, submicrometer-sized droplets of a nanoparticle ink are ejected due to electrohydrodynamic forces. In order to avoid clogging of the fine nozzle (~ 100 nm), much stricter particle loading is required for EHD (0.1 – 1 vol%) in comparison to DIW (typical concentration 75 wt%), and an annealing process is inevitable to improve the mechanical stability and electrical conductivity after printing (Galliker, *et al.*, 2012; Schneider, *et al.*, 2016). Another important factor to consider is that although EHD printing can be performed on nonplanar substrates or 3D objects, the possible electric field distortions must be taken into account (Hirt, *et al.*, 2017). The charged ink droplets involved in the EHD printing are significantly affected and controlled by the electric field (Richner, *et al.*, 2016). This effect, similar to what was frequently observed in dusty plasma research, can be an advantage for depositing high-aspect-ratio structures. However, if the extra charges are not removed swiftly, for example on a non-conductive substrate, the accumulated charges may repel the incoming droplets thus causing problems. Both DIW and EHD are relatively high temperature processes (typically 150 – 400 °C) and can therefore cause mechanical stresses, leading to warping or fracturing of printed layers or the fragile 3D printed nano- and micro-scale structures.

Another type of 3D metal printing is laser-assisted electrophoretic deposition (Fig. 13 (c)), which is based on an optical trap of locally concentrated nanoparticles in a solvent to which an electric field is applied to obtain dense morphology. Fig. 13 (d) shows a meniscus-confined electroplating technique which uses a metal precursor solution with metal particles which have not undergone reduction. Applying an electric potential to the substrate and inside the nozzle induces the electrochemical reduction of the metal ions in the meniscus. Controlling the retraction of the pipette therefore promotes the growth of the 3D metal structure (Hirt, *et al.*, 2017).

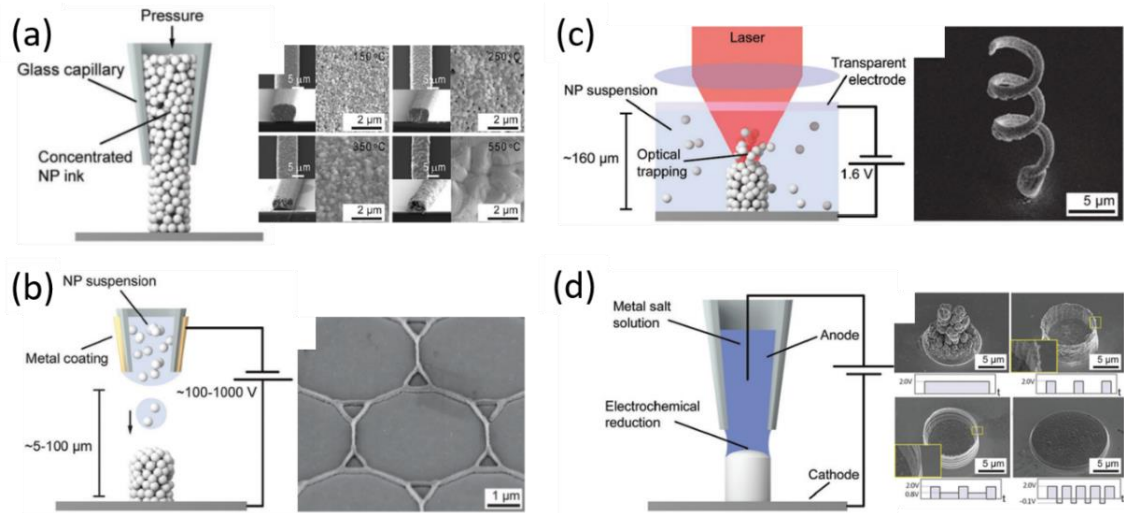


Fig. 13. Comparison of representative examples of microscale metal 3D printing techniques (Hirt, *et al.*, 2017) (a) Left: schematic of direct ink writing (DIW), right: SEM images of printed silver layer using DIW annealed at different temperatures (Skylar-Scott, *et al.*, 2016) (b) Left: schematic of electrohydrodynamic printing, right: gold grid transparent electrode printed using EHD technique with an aspect-ratio of 2.3 and a line diameter of 80 nm (Schneider, *et al.*, 2016) (c) Left: schematic of laser-assisted electrophoretic deposition, right: SEM image of a gold coil fabricated by laser-assisted electrophoretic deposition (Takai, *et al.*, 2014) (d) Left: schematic of meniscus-confined electroplating, right: the morphology of the deposits influenced by control of the meniscus evaporation rate as well as the deposition potential profile. Porous, dense, or hollow structures can be produced (Seol, *et al.*, 2015)

The formation of complex nano-architected metal scaffold recently demonstrated using metal (Ni)-rich photo-resist and two-photon lithography.(Vyatskikh, *et al.*, 2018) As shown in Fig. 14, the bath is first filled with hybrid organic–inorganic materials that contain Ni nanocrystals. Then, using two-photon lithography, a 3D structure is sculpted. After pyrolysis to remove the organic component, this process resulted in a >90 wt% Ni-containing architecture with the minimum 300 nm diameter beams made of 20-nm grained nanocrystalline Ni.

Aside from the limitations due to accessibility and high cost, focused electron/ion beam induced deposition (FEBID/FIBID) is a rare option that can offer nanometre scale metal printing (Hirt, *et al.*, 2017). Hence, it has been widely investigated for specific applications that other additive manufacturing technology may not be able to support, such as customized magnetic force microscopy tips, optical metamaterials, photonic crystals, scanning probe single electron transistors and nanoscale electron

sources (Koops, *et al.*, 2001; George, *et al.*, 2011; Gavagnin, *et al.*, 2014; Murakami and Takai, 2015; Tasco, *et al.*, 2016).

As shown in Fig. 15 (a), in the FEBID or FIBID process, a focused electron or ion beam interacts with adsorbed precursor molecules on the substrate and dissociates the precursor into metallic and organic products. This results in deposition of non-volatile (metal) products and, in principle, desorption of the volatile fragments. Figure 15(b)-(d) shows some examples that demonstrate the performance of FEBID/FIBID FEBID, i.e. precision and controllability of the printed nanostructure.

From the standpoint of metal 3D manufacturing in nano- and micro-scale, the DIW, EHD printing, LIFT and FEBID/FIBID techniques are considered to be at the verge of transitioning from development to application (Hirt, *et al.*, 2017). However, we emphasize that no single technique can presently provide a complete solution of combining high resolution, high purity and intricate geometry.

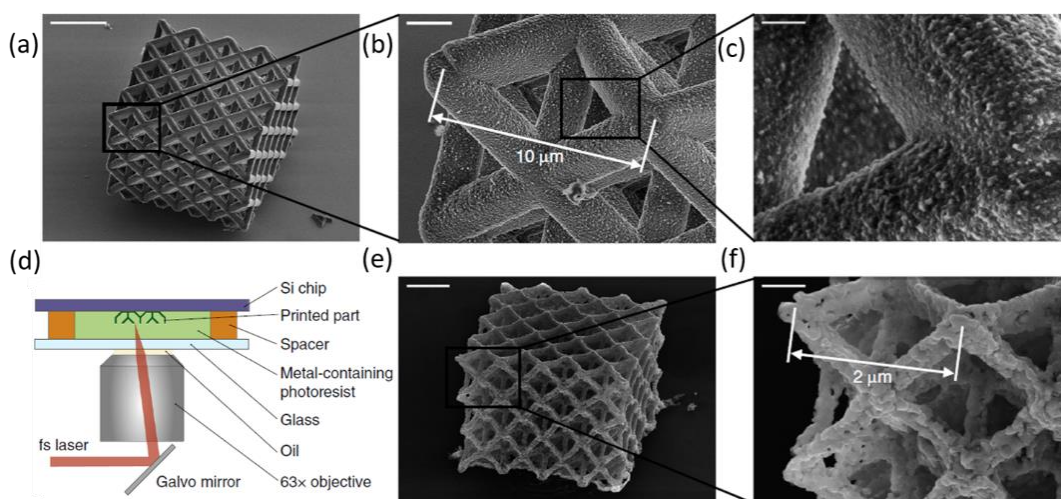


Fig. 14 Process for additive manufacturing of metals with nanoscale features and SEM characterization of the fabricated scaffold. (d) Schematic of two-photon lithography process used to sculpt the scaffold. SEM images of (a)–(c) a representative octet lattice made out of a nickel-containing polymer at different magnifications and (e), (f) a nickel nanolattice after pyrolysis. Magnifications in (b) and (e) (scale bars 2 μm) and also (c) and (f) (scale bars 500 nm) are identical. Scale bar is 15 μm for (a) (Vyatskikh, *et al.*, 2018)

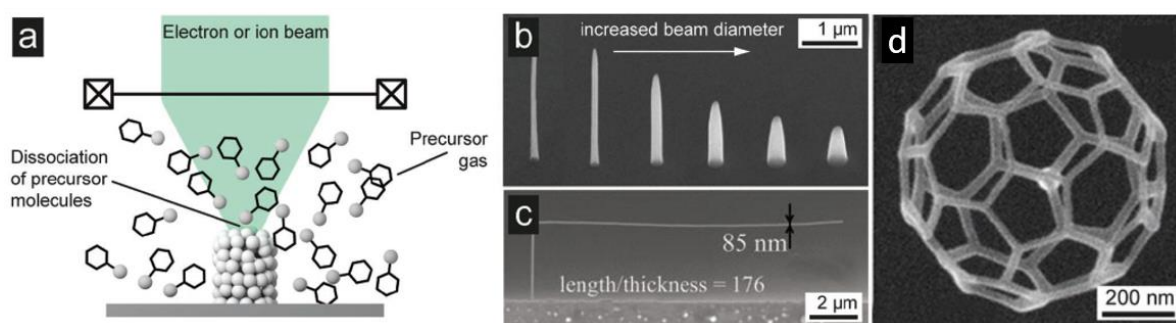


Fig. 15 Focused electron/ion beam induced deposition (FEBID and FIBID) (a) Schematic showing principle of FEBID and FIBID, SEM image of (b) Pt rods deposited with different beam diameters.(Plank, *et al.*, 2008) (c) Horizontally grown DLC wire by feedback of monitored secondary electron current signal to maintain stable growth angle.(Guo, *et al.*, 2013) d) Pt icosahedron fabricated with simulation-guided FEBID.(Fowlkes, *et al.*, 2016)

2.2.3 Ceramic materials

In comparison to plastic and metal processing, the ceramic industry has been slow in implementing additive manufacturing due to the difficulties with shape, a limited resolution and the quality of the produced parts (Schwentenwein and Homa, 2015). However, as global trends point toward customized design and manufacturing and increasing needs for complex nano- and microscale fabrication, it is inevitable that additive manufacturing technologies are also promising for ceramic materials.

Selective laser sintering (SLS) and selective laser melting (SLM) are powder-based direct sintering processes whereby the individual particles are connected using thermal energy by means of a laser. However, significant internal stresses can be induced by high temperature gradients, which result in a rough surface morphology.

On the other hand, indirect methods require a combined binder and ceramic particle mixture and a post-treatment of printed ceramic parts. This is applicable to many polymer-based additive manufacturing processes such as fused deposition modelling (FDM), stereolithography and digital light processing (DLP). (Schwentenwein and Homa, 2015) Upon thermal treatment of the printed object, the organic components are removed and the inorganic particles are sintered to form a high-purity ceramic architecture. Schwentenwein et al. (Schwentenwein and Homa, 2015) demonstrated high density printed alumina ceramic parts which have mechanical strength comparable to conventionally produced alumina parts prepared using a photocurable ceramic suspension with light-emitting diodes (LEDs) combined with a digital micromirror device. The process is known as lithography-based ceramic manufacturing (LCM) because it is based on selective curing by a dynamic mask exposure process which enables printed dense microscale features.

Transparent fused silica is a highly challenging ceramic to produce by additive manufacturing. Using stereolithography 3D printing, Kotz et al. successfully produced high quality printed silica glass parts at a resolution of a few tens of micrometres (Kotz, *et al.*, 2017). A photocurable silica nanocomposite was 3D printed and converted to high-quality fused silica glass by heat treatment, as illustrated in Fig. 16. The printed fused silica glass was non-porous, with a similar level of optical transparency to that of commercial glass, and had a smooth surface with a roughness of only a few nm. Furthermore, by doping with metal salts the colour of the glass can be modified (Kotz, *et al.*, 2017).

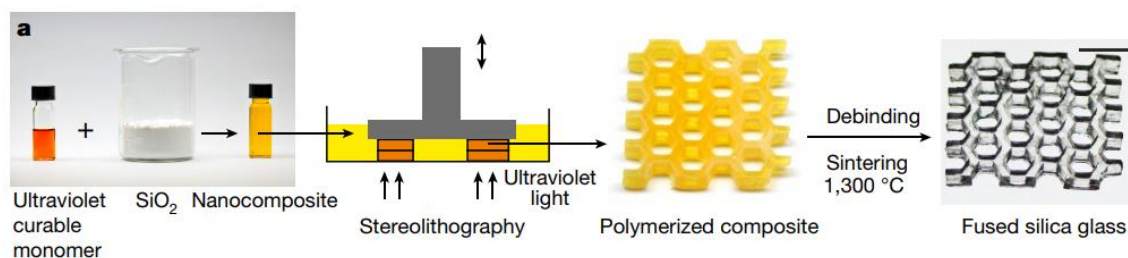


Fig. 16. 3D printing of fused silica glass. Ultraviolet-curable monomer mixed with amorphous silica nanopowder is structured in a stereolithography system. The resulting polymerized composite is turned into fused silica glass through thermal de-binding and sintering (scale bar, 7 mm) (Kotz, *et al.*, 2017)

2.3 Digital control and automation

In order to realize complex 3D structure in micro- or sub micro-scale, digital control and automation are essential, forming a platform to build on using additive manufacturing. A wide range of CAD models can be built, which are then transferred to STL (standard transform language) format to provide cross-section data for each layer of 3D model by digital slicing of the CAD drawing. From there, the tool path (i.e., printer nozzle or laser trajectory) is determined and the automated printing process can begin.

Examination and analysis of performance results for a wide range of compositional and structural parameters is crucial to achieve design optimization and simulation. The complex icosahedron structure of a previously shown Pt nanostructure built by FEBID (Fig. 15 (d)) was possible to fabricate using the simulation-guided processing.(Fowlkes, *et al.*, 2016) The simulation predicts beam-deposit interactions, including spatial distribution of local precursor concentrations, and ‘guides’ the beam accordingly to compensate for such effects. Furthermore, simulation-guided fabrication is considered beneficial for solving other important challenges of additive manufacturing (DebRoy, *et al.*, 2018). For example, shrinkage and deformation of 3D printed structure after annealing are common issues for many metal and ceramic materials. Simulation can help to predict the possible volume loss and geometrical distortion. This result can be used to optimize the original design to compensate for the impact of thermal treatment (DebRoy, *et al.*, 2018).

Similarly to using optical monitoring for precise feedback control, the measured secondary electron current can also be used for achieving a precisely controlled growth angle of the deposit in the FIBID process (Bret, *et al.*, 2004; Rack, *et al.*, 2007). This real time monitoring and feedback control by optical/electrical measurement can be instrumental for improving the stability of the process and the quality of printed film or 3D architecture in the plasma printing system.

The use of simulations and precise control can therefore drastically improve the quality of the product in terms of composition and structure, producing stable, complex 3D structures in a variety of materials.

2.4 Limitations for applications in micro- and nano-scale manufacturing

The surface textures of additively produced materials commonly require a surface finishing process to improve the as-is printed surface characteristics. These include stair-like steps, powder adhesion, fill patterns from extrusion or beam-based systems, and witness marks from support material removal can be improved by choosing a fine filament in FDM; however, this significantly increases the build time. Powder adhesion is a common issue in binding, powder bed and powder feed processes (Bahnini, *et al.*, 2018). To combat this, machining, shot peening and grinding have been widely used to achieve greater surface quality. Chemical processes can also be used, such as electro-polishing. Generally, however, these surface treatment methods are not applicable for microscale structuring. Objects sculpted by additive manufacturing processes generally have inferior mechanical strength due to the intrinsic layer-

by-layer formation and lack of crosslinking between different layers as well as between the printed structure and the substrate. Residual stresses and micropores often exist in the printed metallic parts, which therefore require post thermal processing to improve their mechanical performance and provide the desired microstructure. This is especially problematic when the structure is required to be built on thermally weak materials or thin film-like substrates.

When sculpting composite materials or multi-material 3D objects the necessity of post-processing, especially at high temperature, can limit the choice of materials and processing techniques. Intermittent *in situ* annealing by laser has been proposed to enhance process efficiency and improve the distortion of the sculpted object by removing the thermal post-processing step (Lewis, *et al.*, 2015; Stanford, *et al.*, 2015). However, this intermittent annealing was time-consuming and caused a change in the operating pressure due to the reactive gas supply, which interrupted the deposition process. Pre- and post-treatments in additive manufacturing are considered inevitable to a certain extent, hence research has mostly focussed on creating more facile post-processing methods instead of removing this step completely.

The low reduction rate for certain metals deposition in FEBID/FIBID (Hirt, *et al.*, 2017) is also an area that requires further research. While Au, Co, Fe and Ge could be reduced almost completely, other metals such as Ir, Mo and Ti show low reduction levels of under 35% (Botman, *et al.*, 2009).

In contrast to FEBID/FIBID, which can only provide electrons or ions, plasma can provide a diverse range of chemically reactive species. The dependence on the precursor materials is not as dominant when using plasma and it may be possible to increase the metal reduction level even in composite material printing. Many conventional additive manufacturing techniques require a separate process for each different material printed in one object. Plasma CVD systems with precursor solution have the potential to overcome this, achieving complex micro- or nano-fabrication in one step, either by mixing two (or more) different liquid precursors or by installing two (or more) plasma jet systems next to each other.

In the following section, we shall discuss in more detail the mechanisms of plasma processing to understand the benefits of plasmas and how plasma processes can potentially help overcome the limitations and current issues of additive manufacturing.

3. Relevant plasma-specific effects

The obvious benefits of using low-temperature plasmas in printing technology are the diversity of reactive species that exist and the multiple options for controlling them. Depending on electrode configuration, operation pressure, electron energy and density, as well as the reactive gas environment, the chemical and physical properties of the plasma can be manipulated using different controls. Surface functionalization, etching, deposition and on-surface reactions are all examples of material modification that can be carried out by controlling the plasma properties.

3.1 Surface functionalization

A successful example of the commercialization of atmospheric-pressure plasma processes is the plasma surface modification of a wide range of polymeric substrates. Usually, the highly hydrophobic properties of polymeric materials result in poor wetting and adhesion properties, which often cause difficulties in various manufacturing processes; for example, chemical wet processing, printing, deposition and bonding with a different film or layer. Plasma surface modifications can be used to significantly increase hydrophilicity, even under high speed Roll-to-Roll (R2R) operating conditions. Atmospheric-pressure plasmas generate high density electrons, ions and metastable species, including reactive oxygen and nitrogen-containing reactive radicals even in inert gas discharge that impinge onto the substrate due to its exposure to ambient air during the process. Atmospheric pressure plasma jets have been used as an effective tool for introducing a chemical change on substrate materials with hydrophilic functional groups such as $=O$, $-OH$, $-O$, or $-NH$, both in the presence and absence of an additional oxygen source such as water vapor, air or pure oxygen gas.

Figure 17 shows an example of surface modification used to increase hydrophilicity or hydrophobicity, depending on the reactant gas condition. In Fig. 17, hydrophobicity is obtained by depositing SiO_x layers using an atomized organosilicon precursor such as TEOS (Tetraethyl orthosilicate) or HMDSO (Hexamethyldisiloxane) (Verhoeven, *et al.*, 2014). Changing the reactant gas to a perfluorocarbon compound, for example C_xF_y or NF_3 , can even create a superhydrophobic surface by replacing the C-H functional groups containing C-F bonds (Satulu, *et al.*, 2016). This surface modification function is important in printing applications as it enables selective modification using a microplasma jet or pin-to-plate discharge configuration so that the surface energy and adhesion properties can be modified locally, improving the resolution and durability of the printing process as shown in Fig. 17 (g).

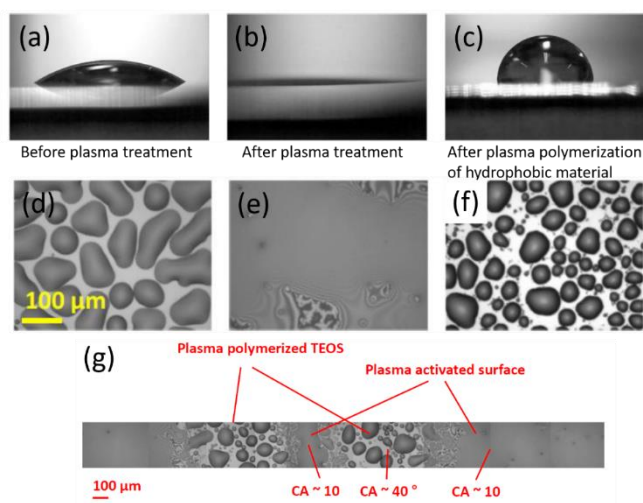


Fig. 17. Contact angle and surface wetting properties modified by the plasma treatment: Contact angle of water (a) before plasma treatment (b) after plasma treatment by air (c) hydrophobic organosilicon film coated surface (d) – (f) corresponding wetting properties of surface (a)–(c) respectively (g) selective modification in $\sim 200 \mu m$ line width of polymeric surface by atmospheric pressure plasma. (Verhoeven, *et al.*, 2014) In panel (g), CA stands for contact angle.

Yick et al. demonstrated the highly confined surface modification of vertically aligned carbon nanotubes (VA-CNTs) for gold nanoparticle deposition using a microplasma jet, which can be beneficial for selective patterning or printing with micrometre precision. This localized plasma treatment enabled the production of a microfluidic strip that effectively confined Au nanoparticles to a 100 μm feature on the VA-CNT support (Yick, *et al.*, 2013). The deposited nanogold was found throughout the entire depth (2.5 μm) of the microfluidic strip, which indicates a 3D Au–CNT hybrid formation over the entire nanotube length. In additive manufacturing processes or decorative depositions on various substrates or 3D objects, this uniform and selective surface modification can be a versatile assisting tool and processing technique.

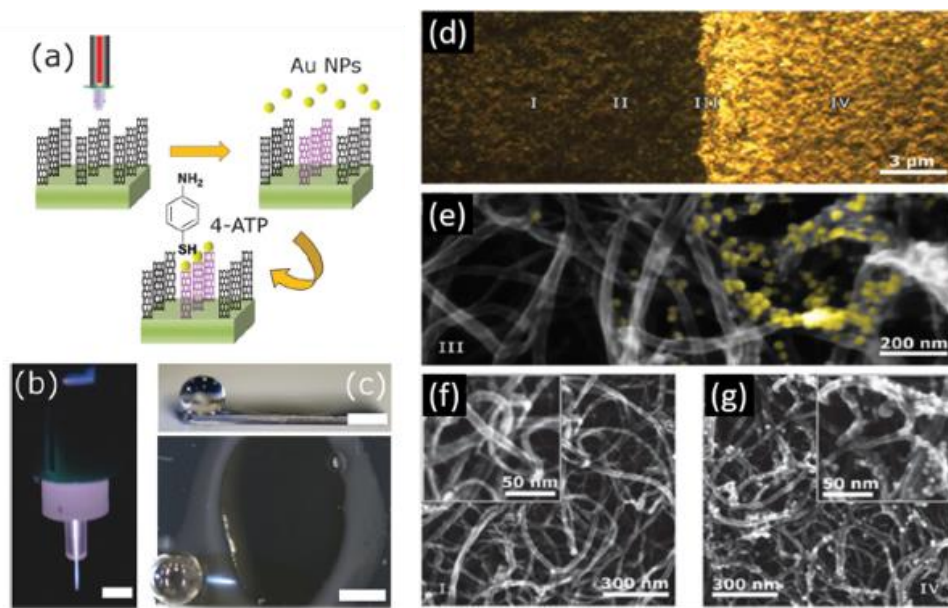


Fig. 18. Plasma surface modifications to confine Au nanoparticles (Au NPs) within microscopic line features: (a) Schematic of the surface modification and the nanogold deposition procedure. The microfluidic strips are highlighted in pink. (b) A photo showing atmospheric pressure plasma jet and the plasma plume's size. (c) Change in surface wettability of the VA-CNTs in the plasma-modified (hydrophilic) and the pristine area (hydrophobic). Scale bars are 2 mm in (b) and (c). (d) Optical micrograph at the interfacial region of Au and CNT (e) SEM image depicting the interfacial region (III). False color is applied to highlight Au nanoparticles. SEM micrographs of the pristine (f) and the Au NP decorated (g) sides of the VA-CNTs, corresponding to areas (I) and (IV) in (a), respectively. Insets in (c) and (d) are the corresponding high-resolution SEM images (Yick, *et al.*, 2013)

3.2 Etching

Plasma etching is one of the most common processes whereby plasma-generated chemical species react with the surface atoms and remove them by forming volatile products. Due to the highly collisional environment at atmospheric pressure, a significant physical sputtering effect by high energy ions, as in low-pressure plasmas, cannot be expected. However, in many polymer materials such as polyethylene (PE), polypropylene (PP), polycarbonate (PC) and poly(ether ether ketone) (PEEK) etc., it has been shown by measurement of the mass loss and etch profiles after the plasma exposure that high etch rates of 50 – 300 nm/s are possible (Fricke, *et al.*, 2011). The experiments showed that reactive oxygen species

play an important role in the polymer etching process (Fricke, *et al.*, 2011). The etching typically leads to chemical changes on the surface and is very effective in changing the morphology of the substrate. Indeed, depending on the process conditions, one can sculpt micro- or nano- structures of different dimensionalities (1D-3D).

Plasma etching at atmospheric pressure is not limited to polymer materials. Thomson *et al.* showed that etching by atmospheric-pressure DBD plasmas is an effective method for controlling the ZnO surface morphology (Thomson, *et al.*, 2013). Surface morphology is a critical parameter that determines the optical scattering properties of transparent conducting oxides in photovoltaic thin films. The reactant gases were found to have a significant effect on the etching process; in particular, water introduction has been shown to be important for the efficient etching of ZnO. It was proposed that a two-stage reaction process occurs, with intermediate products (e.g., a hydrate or partial hydrate) reacting faster with oxygen.

Using hydrogen as a reactant gas, copper (Cu) dry etching with a maximum etching rate of 500 nm/min was demonstrated at close to atmospheric pressure (13.3 kPa, 100 Torr). The application for this is the direct Cu wiring of integrated circuits. As shown in Fig. 19, around the apex of a fine pipe electrode a localized hydrogen plasma is generated. Cu etching was observed only when the process gas contained hydrogen, with hydrogen concentration significantly influencing the etching rate. It was also possible to etch Si and SiO₂ and at the same operating conditions the etching rates were 100 nm/min and 50 nm/min, respectively; hence this etching process is suitable for the Cu wiring application on an SiO₂ layer. In addition to the examples of plasma etching at atmospheric-pressure introduced here, many others have been reported in literature and should be carefully considered in the development of the plasma-assisted printing processes.

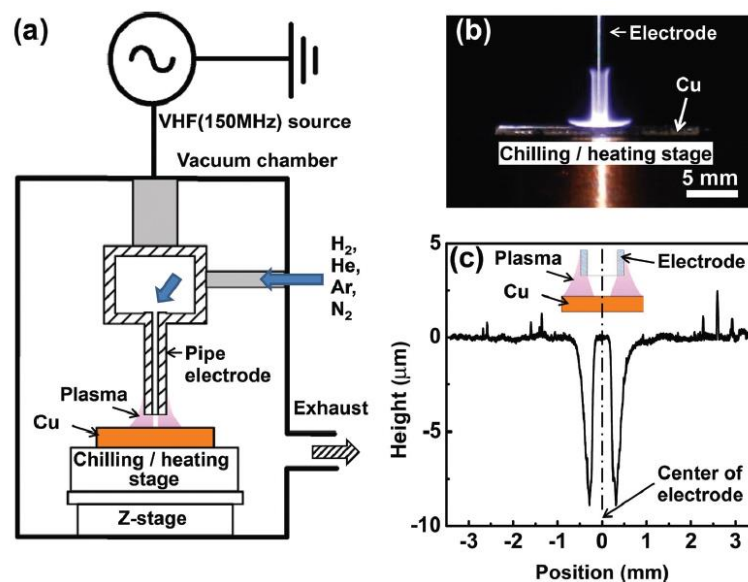


Fig. 19. (a) Schematic of the experimental setup used in Fricke *et al.* (Fricke, *et al.*, 2011) (b) Aspect of the pure hydrogen plasma generated at a VHF input power of 100W, hydrogen pressure of 13.3 kPa, and gas flow rate of 10 slm. The electrode outer diameter is 1mm (c) Typical surface profile of the etched Cu sample and the positional relation between the electrode and the etching profile. The x-axis offset coincides with the center axis of the electrode (Fricke, *et al.*, 2011)

3.3 Deposition

Deposition is directly related to 2D and 3D printing processes as it builds up layers and architects them in three dimensions. *Plasma polymerization* can be understood as an extension of deposition by crosslinking monomers into larger molecules through the build-up of layers. The influence of different species (ions, neutral species, and photons) on the polymerization process within the plasma discharge was studied by examining the crosslinking properties. This was carried out by using a carefully designed setup with different optical and metal windows to deconvolute the effects of the various plasma species on the surface nanomechanical properties of polyethylene at low pressure ($p \sim 500$ mTorr). It was shown that surface crosslinking is primarily due to the simultaneous effects of neutral species and vacuum ultraviolet (VUV) photons, whilst the ion bombardment effect is secondary. The intensity of VUV may differ at atmospheric-pressure conditions. However, predominantly high-density neutral radicals are commonly available in atmospheric-pressure plasma discharges and are expected to have an important role in polymer based additive manufacturing processes, improving the mechanical properties and accelerating polymerization.

It is also possible to obtain metallic film or particles with non-thermal atmospheric pressure plasmas by *physical deposition* processes (Fig. 20) (Shimizu, *et al.*, 2009; Lazea-Stoyanova, *et al.*, 2015; Shimizu, 2017). Lazea-Stoyanova *et al.* demonstrated the synthesis of nanometre-sized, well-distributed copper particles at atmospheric pressure using a radiofrequency (RF) low-temperature argon plasma jet (Lazea-Stoyanova, *et al.*, 2015). The copper bulk material was used as a powered electrode and evaporation target simultaneously. The Cu particles were condensed and collected onto Si substrates. Similarly, Shimizu *et al.* successfully demonstrated nanogold particle generation and deposition using 100 μm diameter gold wire with 450 MHz ultrahigh-frequency (UHF) driven microplasma jet (Shimizu, *et al.*, 2009). The pulse repetition rate was varied from 50 to 100 Hz whilst the duty cycle was kept constant at 5 %, providing the non-thermal process with a time averaged UHF power as low as 0.8 W (Shimizu, *et al.*, 2009). The effective physical deposition and control over the size of the nanogold particles is closely related to the H_2 volume ratio, as the concentration of H_2 affects the plasma discharge and changes the concentration of atomic hydrogen, as well as the heat flux density. Considering the atomic hydrogen-metal interaction model (Ohno and Uda, 1989), it was concluded that hydrogen atoms first dissolve into the molten Au wire and the exothermic hydrogen recombination reaction then enhances the gold wire etching. The nanogold film eventually forms with a uniform size distribution through the condensation from the Au vapour.

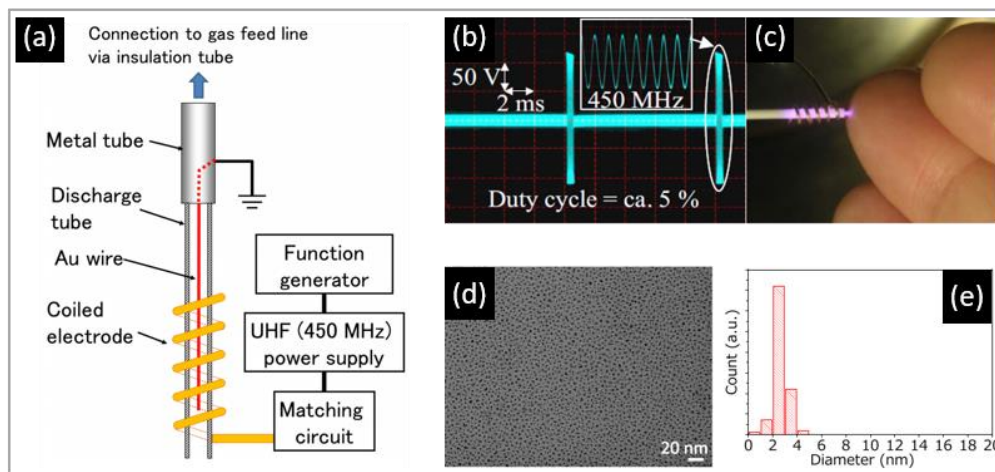


Fig. 20. (a) Schematic of the microplasma setup used for nanogold synthesis (Shimizu, 2017) (b) Pulse-modulated UHF voltage waveform effective for lowering gas temperature (c) Micro plasma jet touching finger without thermal/electrical shock (Shimizu, *et al.*, 2009) (d) SEM image showing finely controlled Au deposition (e) Estimated size distribution (Shimizu, 2017)

In comparison to physical deposition, atmospheric pressure PECVD (AP-PECVD) shows more flexibility in the range of materials that can be treated and the possibility of producing composite materials with varying composition. SiO_x deposition as a barrier coating for electronics, solar cells and food packaging has been intensively investigated and successfully commercialized. Furthermore, metal embedded SiO_x films produced by high-temperature air plasmas are effective for antibacterial applications. For example, hexamethyldisiloxane (HMDSO) was used as a primary precursor and solutions with silver nitrate AgNO_3 , zinc nitrate $\text{Zn}(\text{NO}_3)_2$ and copper nitrate $\text{Cu}(\text{NO}_3)_2$, in a 1:1 volume mixture of isopropanol and water, were sprayed into the plasma as secondary precursors (Zimmermann, *et al.*, 2011; Jager, *et al.*, 2019). After separation from the precursor molecules, Ag, Zn and Cu elements bind to oxygen, thus forming metal oxides that are subsequently incorporated into the SiO_x film.

Amongst the range of target materials that can be produced using the AP-PECVD technique, it is worthwhile to mention the growth of carbon nanotubes (CNT) from a C_2H_2 gas precursor (Kyung, *et al.*, 2006). The deposition was performed at 400°C using dielectric barrier discharge with $\text{He}/\text{C}_2\text{H}_2$, N_2 and NH_3 addition. Interestingly, it was found that pre-treatment of the Ni catalyst can modify the surface morphology, enhancing carbon diffusion during CNT formation. However, this process requires substantial heating, which limits its applicability for plasma printing unless special arrangements are made to print on sufficiently heated substrates.

In practice, chemical and physical deposition occur simultaneously in many cases, especially in atmospheric pressure plasma processing. In the example of deposition of gold (Shimizu, *et al.*, 2009), the unique chemical environment of rich atomic hydrogen enabled the evaporation of solid gold without any additional thermal energy source or sputtering aid. It is important to stress this benefit of the plasma process, as the synergy between the chemical and physical effects may enable the fruition of several processes of interest in additive manufacturing technology.

3.4 On-surface reactions

As an extension of surface modification, in recent years there has been active research into on-surface plasma reactions to provide innovative and practical solutions to various material processing issues. The reductive writing of metal cations with 30 μm linewidth scale shows a plasma jet to be an alternative pathway to conventional lithography (Lee, *et al.*, 2011). On the metal precursor coated surface, microscale plasma treated and effectively reduced the metal ions to crystalline metal nanoparticles. As shown in Fig. 21, metal reduction was not observed with a control sample that was treated through the grounded mesh. This result proves the importance of electrons in the plasma reduction of metal cations, as suggested by electrochemical analysis of the plasma-liquid systems (Richmonds and Sankaran, 2008; Richmonds, *et al.*, 2011). A wide range of metals (Ag, Au, Pt, Ir, and Ru) were tested with different polymeric systems such as polyvinyl alcohol (PVA) and poly(methyl methacrylate) (PMMA). This approach provides a way to fabricate patterned conductive films for applications ranging from plasmonics to catalysis, at low cost and high throughput (Lee, *et al.*, 2011).

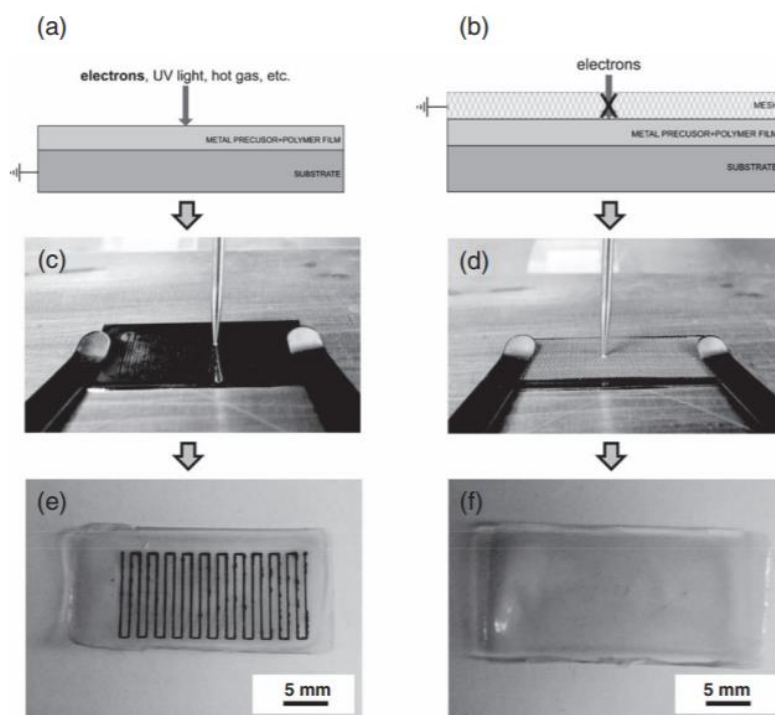


Fig. 21 Control experiment to demonstrate the crucial role of electrons in the reduction process: schematics of plasma reduction by plasma jet (a) without mesh and (b) with grounded mesh to filter out electron interaction; photos of plasma treatment (c) without mesh and (d) showing reduced metal pattern related to (a) and (c); (f) no reduction shown treated through grounded mesh to reveal importance of electrons in metal reduction process (Lee, *et al.*, 2011)

The dehalogenation reaction (Fig. 22) (Hartl, *et al.*, 2019) is another interesting example of plasma on-surface reactions. Similarly to Lee *et al.* (Lee, *et al.*, 2011), N_2 - plasma was applied to a liquid source sprayed surface. In this application, atmospheric-pressure plasma treatment is thought to be promoting a metal-catalysed surface reaction that cleaves carbon-halogen (C-X) bonds in the presence of a transition metal (Hartl, *et al.*, 2019). Due to the low dissociation energy of the C-X bond, catalytic

cleaving may happen through the plasma-assisted Ullmann reaction, where copper acts as a catalyst for C–X bond dissociation and induces subsequent aryl–aryl coupling.(Ullmann and Bielecki, 1901) In this study, a liquid phase small-molecule precursor, 1,2,4-trichlorobenzene (TCB), was converted to a solid, thin-film product in a process carried at ambient conditions. The deposited total energy was found to be a determining factor for C–Cl bond scission, with different plasma parameters leading to variations in film morphologies, roughness and feature size.

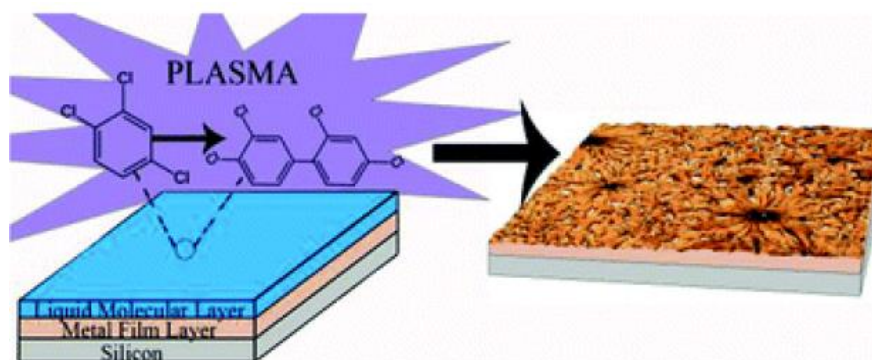


Fig. 22 Schematic to represent the plasma-catalytic dehalogenation process (Hartl, *et al.*, 2019)

Plasma has the unique benefit of non-equilibrium characteristics, containing high energy electrons at low gas temperature whilst also containing high-density reactive radicals. This makes it highly suitable for assisting conventional printing technology, which often faces challenges in dealing with various non-traditional substrates or 3D objects. Plasma techniques, such as surface modification, etching, deposition of new film and various on-surface chemical reactions, can potentially provide a breakthrough and enable the realization of new concepts in 2D and 3D printing applications where current techniques fall short. In section 4, we discuss the examples of plasma applications for developing the micro and nanostructure of materials based on these elemental plasma reactions, before exploring plasma enhanced 3D printing with micro and nanoscale features in section 5.

4 Plasma nanotechnology for printing

Plasma nanotechnology for the fabrication of intricate nanostructures is gaining momentum in certain areas of fundamental research and potential industrial applications and even becoming standard practice in some existing industrial applications. A range of plasma sources and processes have been successfully adapted for the fabrication of a multitude of nanoscale structures, features and devices. In this section, we focus on processes which can be combined with conventional printer systems (inkjet or a 3D printing unit), as well as the possibility for direct one-step plasma printing from a precursor solution. The processes discussed include nanoparticle synthesis, nano-ink production and deposition of nanoparticle in a flow, and feature printing.

4.1 Nanoparticle production

Metallurgical reduction, fuming, atomization, electrolysis, grinding, the sol-gel method, precipitation and spray drying are typical conventional processes used for producing metallic powder (Young and Pfender, 1985). Plasma-based fine metal powder synthesis methods are preferred because they provide numerous particle types at high throughput and with good process control (Forster, *et al.*, 2012; Lazea-Stoyanova, *et al.*, 2015).

4.1.1 Thermal plasma synthesis of nano-sized powders

The plasma synthesis of nano-powders by thermal plasma is one of the most rapidly emerging areas of plasma technology. The precursor material is vaporized within the plasma discharge, followed by rapid quenching of the vapour cloud to produce fine powder, as shown in Fig. 23 (a) (Gonzalez, *et al.*, 2008; Guo, *et al.*, 2010; Shigeta and Murphy, 2011; Kim and Kim, 2019). The process is not limited by the precursor form, as powder, a wire, a suspension, a liquid or even a gaseous precursor can be used. It is also possible to generate exfoliated nanomaterials from multilayered bulk materials, as in Fig. 23 (b) (Kim and Kim, 2019). The process has been successfully applied for the synthesis of metallic nanoparticles and nanoceramic powders such as Al, Cu, Ni and Si and nano-alumina, nano-zinc oxide, nano-glass, and BN nanotubes (BNNTs). It is possible to produce nanopowders with an identical chemical composition to the feed stock, or to significantly modify the chemistry with a fine control of morphology and size distribution through the additional process parameter control, which is not possible in the combustion process (Boulos, 2016; Kim and Kim, 2019).

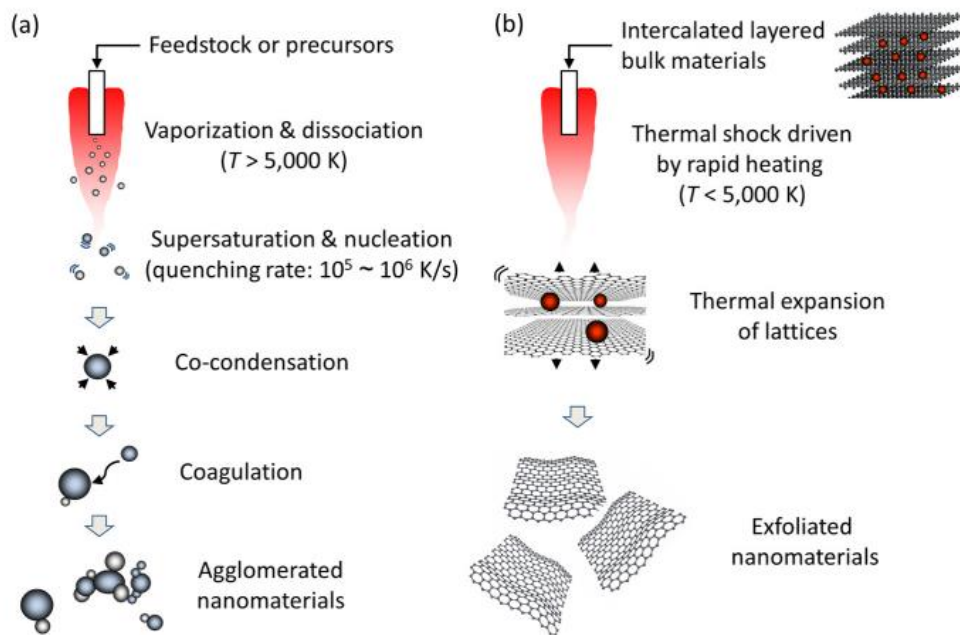


Fig. 23 Nanofabrication using thermal plasma jets. (a) Bottom-up (b) Top-down approaches (Kim and Kim, 2019)

Some of the most common thermal plasma sources for nanopowder synthesis are shown in Fig. 24. The transferred DC arc plasma source, as shown in Fig. 24(a), uses the electrode as an evaporation target and can easily generate nanometre scale metal powder. However, issues with erosion result in the need for frequent replacement of the electrode, making the process non-continuous. Contrary to this, the non-transferred DC plasma torch can support continuous synthesis by using target materials injected near the exit of the plasma plume, as shown in Fig. 24 (b). However, most precursors cannot be fully evaporated within this short length of the plasma flame. In addition to the obvious advantage of electrodeless discharge, inductively coupled (ICP) type RF torch can sustain a much longer plasma discharge length, which enhances the evaporation rate (Seo and Hong, 2012).

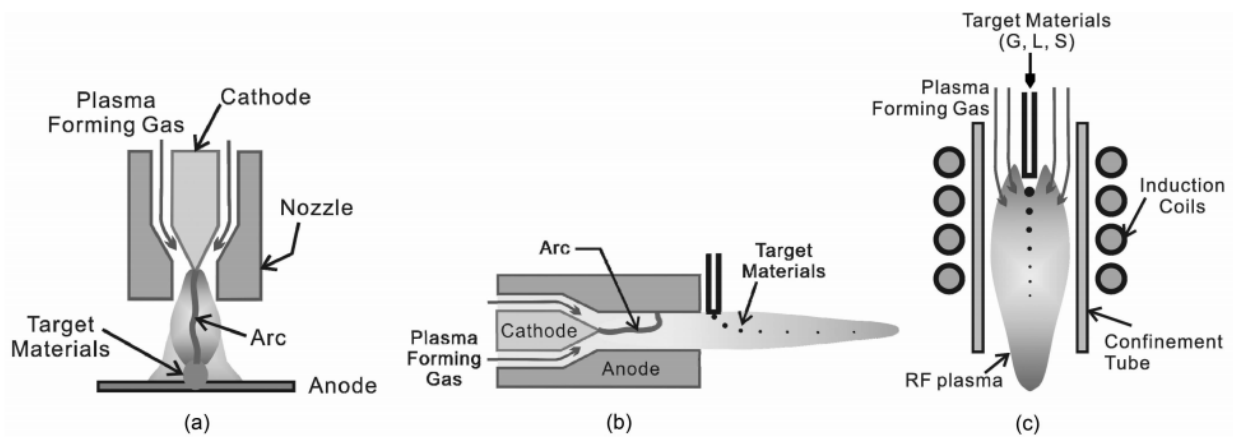


Fig. 24 Schematic diagrams of the typical thermal plasma torches available for synthesis of nano-sized powders. (a) Transferred DC Plasma Torch (b) Non-transferred DC Plasma Torch (c) RF Plasma Torch (Seo and Hong, 2012)

One noteworthy point in nanopowder synthesis using thermal plasma is the dependence of a specific reaction on the Gibbs free energy (ΔG). Mostly, the oxidation reactions are spontaneous, having highly negative ΔG values for the entire operating temperature range. However, some carbonization, nitridation, and boronisation processes with positive ΔG require a specific temperature range to be sustained to maximize efficiency (Kim and Kim, 2019).

Recent advances in nanomaterials synthesis may herald new directions for thermal plasma applications. (Kim and Kim, 2019) Figure 25 (a) illustrates how the new composite materials, namely the high-entropy alloys (HEAs), are distinguishable from typical phase-separated alloys. These alloys react differently in the ammonia oxidation process. The HEA of five dissimilar metals (Pt, Pd, Rh, Ru and Ce) is produced by a two-step carbothermal shock (CTS) method that employs flash heating and cooling (maximum temperature of ~ 2000 K in shock duration of ~ 55 ms) of metal precursors on an oxygenated carbon support. The rapid heating and quenching of the CTS technique enabled uniform diffusion of individual elements and avoided phase separation, which is a fundamental problem in multi-component particle synthesis. Considering the similar characteristics of rapid heating and quenching in thermal plasma, we articulate this new area where thermal plasmas has strong potential to contribute (Yao, *et al.*, 2018).

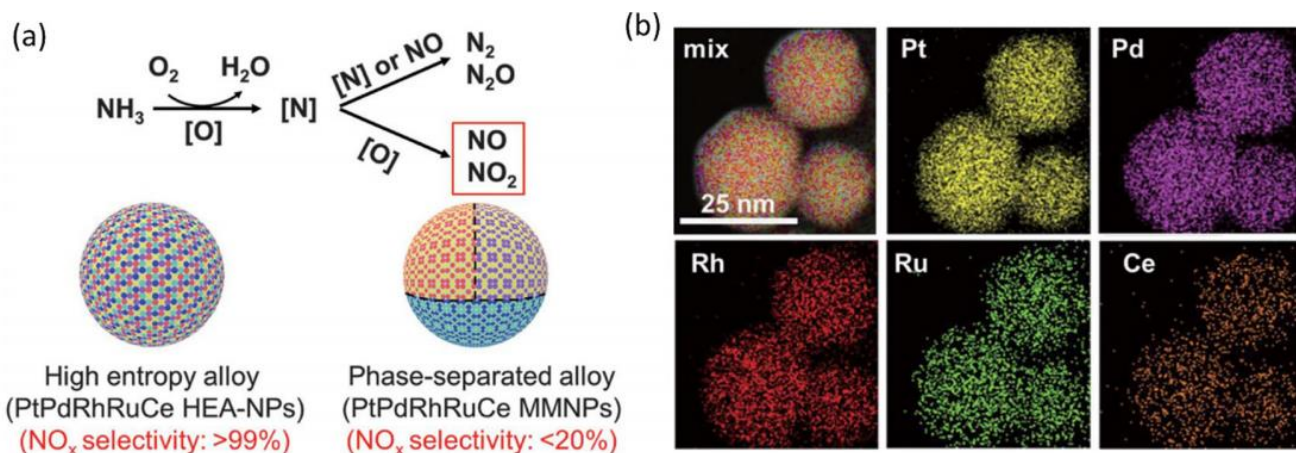


Fig. 25. Synthesis of high-entropy-alloy nanoparticles (HEA-NPs). (a) Schematic comparison of a phase-separated heterostructure and a high-entropy-alloy structure for ammonia oxidation process (b) Quinary HEA-NPs (PtPdRhRuCe) synthesized by a carbothermal shock method (Yao, *et al.*, 2018)

4.1.2 Plasma synthesis of nanoparticles

Non-thermal plasma is a viable option for production of highly functional elemental, alloy and doped nanoparticles. A recent extensive review on this topic includes the fundamental principles, materials and applications (Kortshagen, *et al.*, 2016). Low-temperature, low-pressure plasma processes have been effective for the synthesis of a range of high quality semiconductor nanocrystals (Mangolini, *et al.*, 2005; Hunter, *et al.*, 2017). Through the recombination of electrons and ions at the particle surface, heat is released to enable the particles to reach high temperatures. This is highly desirable for the synthesis of nanometer-sized silicon (Mangolini, *et al.*, 2005). The high energy, highly mobile electrons in the plasma prevent agglomeration by producing negatively charged particles (Matsoukas and Russell, 1995; Kortshagen and Bhandarkar, 1999), whilst also producing the core/shell heterostructure of the nanoparticles (Hunter, *et al.*, 2017).

However, as in many plasma applications, the effort to shift the nanoparticle synthesis to atmospheric pressures is important for lowering cost and making the process attractive for industrial applications (Kramer, *et al.*, 2015). Using kinetic modelling, it has been predicted that much higher plasma densities are required for nanocrystal synthesis at atmospheric pressure due to the strong quenching effect. (Kramer, *et al.*, 2015) Among the numerous examples of nanoparticle synthesis using non-thermal plasma, it is worthwhile to highlight the work related to core/shell heterostructure nanocomposite synthesis, production of metal nanoclusters and in-flight nanogold synthesis from solution precursors.

An all-gas-phase process using low-pressure RF plasma was developed for the synthesis of Ge/Si quantum dots (QDs) (Hunter, *et al.*, 2017), which is usually carried out using wet chemistry approaches. Figure 26 (a) shows the schematic of the plasma reactor for Ge/Si QDs and illustrates the procedure at each different stage. The primary gas feedstock, consisting of GeCl_4 , Ar and H_2 , is supplied to the chamber and electron impact dissociation occurs in region ①, initiating nanoparticle nucleation. In region ②, Ge QDs grow predominantly by agglomeration. In region ③, Ge QD growth and

crystallization continues by surface deposition of radical species and energetic surface reactions (Kramer, *et al.*, 2014). Silane precursor is injected to region ④, where it undergoes decomposition followed by heterogeneous surface growth of Si onto Ge QD cores. The synthesized Ge/Si core/shell QDs exit the plasma reactor and are collected from the gas phase via impact onto the substrate in region ⑤.

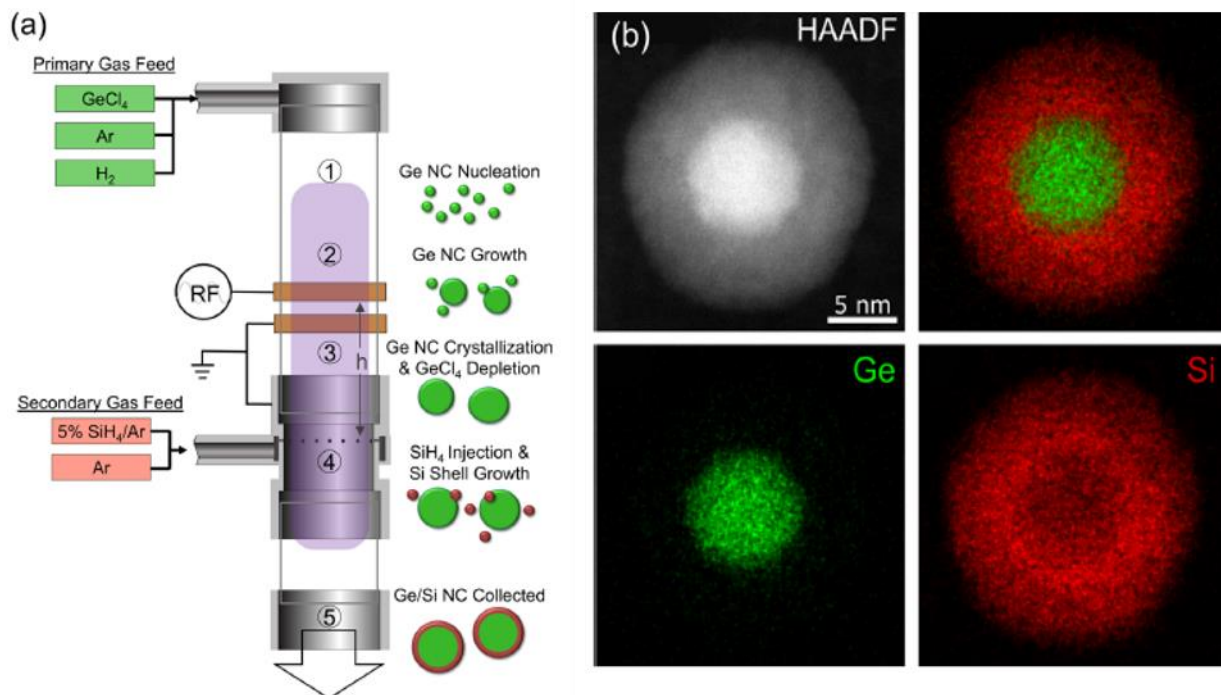


Fig. 26 (a) Schematic of the flow-through non-thermal plasma reactor used for the synthesis of core/shell semiconductor QDs. Numbers 1–5 on the diagram indicate the stages of core/shell QD growth (b) High-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) images of a single Ge/Si core/shell QD. The HAADF image shows a Ge core (bright) surrounded by a Si shell. STEM–EDX elemental maps of Ge (green) and Si (red) show elemental segregation to the core and shell, respectively, as viewed in the projection (Hunter, *et al.*, 2017).

Microplasma at atmospheric pressure has been utilized as an effective tool for the synthesis of a wide range of nanoparticles. Kumar *et al.* have demonstrated the generation of very small Ni nanoclusters using an atmospheric pressure microplasma jet. A powder form of Bis(cyclopentadienyl) nickel (II) [Ni(Cp)₂, nickelocene] was used as a Ni precursor. Atomic force microscopy (AFM) and ion mobility spectrometry (IMS) were used to independently characterize the produced nanoclusters. A combination of bulk matter approximations and calculations based upon structural models were used to correlate the AFM and IMS measurements to the number of atoms in the produced nanoclusters. The analysis results suggested that nanoclusters with less than 10² atoms are likely to be produced using this microplasma nanoparticle synthesis technique (Kumar, *et al.*, 2014).

Maquire *et al.* introduced continuous in-flight synthesis of gold nanoparticles using a solution precursor with an atmospheric-pressure RF plasma source (Fig. 27). Based on liquid droplet irradiation with ultralow (<0.1 eV) energy electrons, well-controlled Au nanoparticles (4.4 ± 1.5 nm) were synthesized in less than 150 μs. Nanoparticle synthesis rates within the droplets were estimated to be

many orders of magnitude greater than radiolysis, electron beam irradiation, and colloidal chemical synthesis where reaction times vary from seconds to hours (Maguire, *et al.*, 2017). The significantly enhanced nanogold particle synthesis rates are considered to be due to high-density electrons, electrons and nanoparticle confinement at the surface of the droplet, as well as the picolitre-sized volume of the droplets which act as microreactors. This approach is promising, with considerable potential for scale-up of the synthesis and, more importantly, for continuous on-demand delivery of high-quality nanoparticles directly to their point of use without additional collection, recovery, and purification steps (Maguire, *et al.*, 2017)

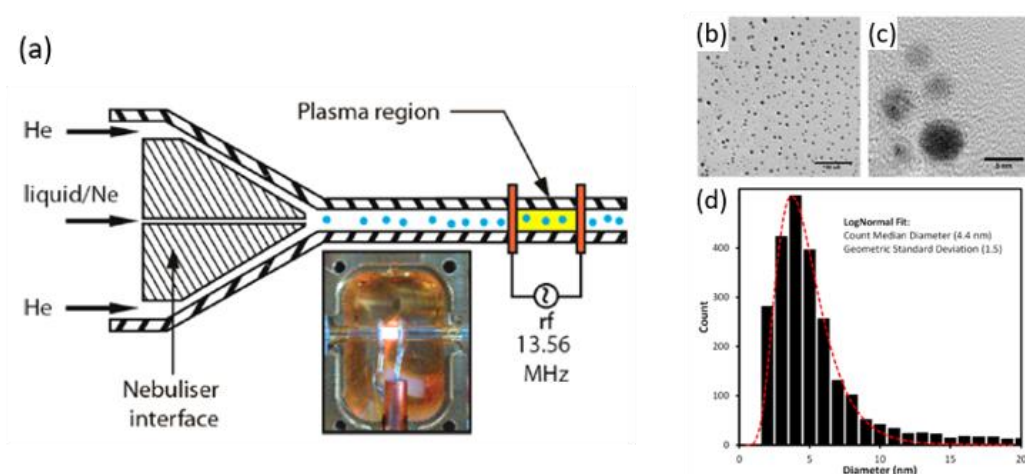


Fig. 27 (a) Schematic representation of the droplet-plasma micro-reactor (inset) Image of ignited plasma region. TEM images of (b) Au nanoparticle onto TEM grid direct from plasma and (c) a typical nanoparticle with diameter 4.5 nm (d) The particle size distribution obtained from a sample of 2100 nanoparticles, showing a mean diameter of 4.4 nm (Maguire, *et al.*, 2017)

4.1.3 Plasma-assisted synthesis of nanocrystal inks

After the synthesis of nanoparticles, production of a stable colloidal ink typically requires complex transfer steps, followed by functionalization in liquid phase. Using non-thermal plasma, Mangolini *et al.* demonstrated the production of silicon nanocrystal inks by combining nanoparticle synthesis and grafting of polymeric material, resulting in a product that can be readily dispersed in organic solvents to form stable silicon ink (Fig. 28) (Mangolini and Kortshagen, 2007). The process requires two different low-pressure RF plasma systems operating at high and low power densities. The nano-silicon crystals are synthesized in the first reactor from an Ar/SiH₄/H₂ supply at high power density (50 W/6.4 mm discharge tube) using RF plasma. The gas flow transports the particles into the second parallel plate plasma region (< 10 W/ gap distance between the electrodes of ~5 cm) where they undergo functionalization from the liquid organic precursor. This two-zone plasma system enhances the optical properties of the product. This methodology of immediate grafting of organic molecules after nanoparticle synthesis could be potentially extended to atmospheric-pressure plasma operation.

Exploration of different materials could also be used to optimise this plasma assisted nanoparticle synthesis process.

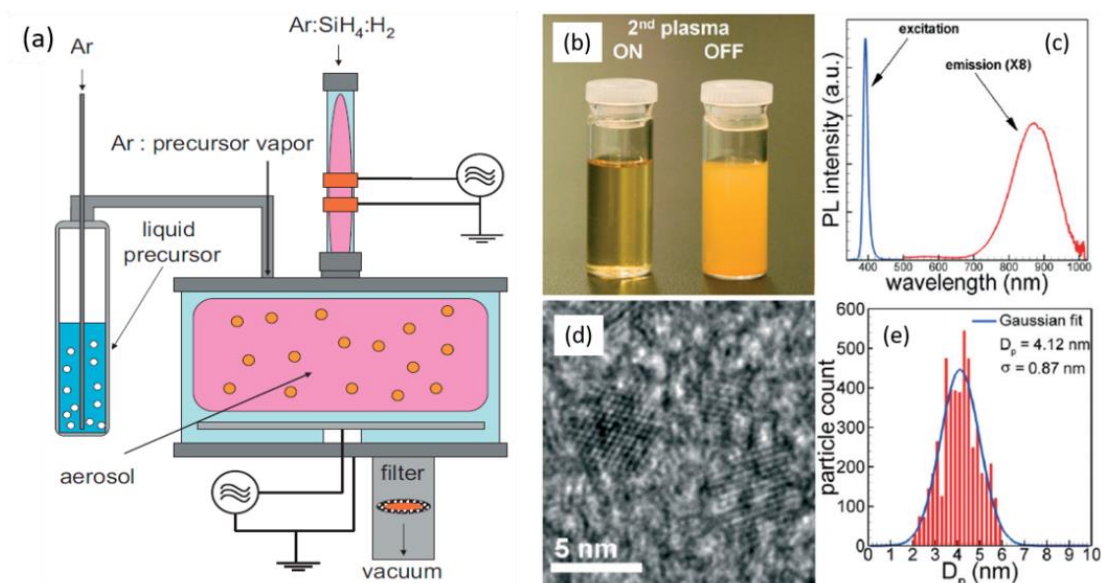


Fig. 28 (a) Schematic of the experimental setup for the synthesis and in-flight functionalization of silicon nanocrystals (b) Photograph of the produced silicon nanocrystals dispersed in toluene, with (left) and without (right) the second plasma (c) Photoluminescence spectra (red) of the in-flight grafted sample shown in (b) with the UV excitation peak (blue) (d) Bright field TEM image of particles redispersed in toluene after in-flight treatment and drop-casting onto a thin film carbon grid (e) Particle size distribution was estimated TEM images (Mangolini and Kortshagen, 2007)

4.2 Deposition of nanoparticles in a flow

Beyond particle synthesis, using plasma printing to build a nanostructured film or material in a controlled manner is important considering the opportunities it could bring in nanoelectronics, nanophotonics, catalysis, sensing and energy applications. Mackie et al. presented a substrate-independent method for the plasma deposition of nanostructured, crystalline metal oxides using an organometallic precursor and micro-hollow cathode plasma discharge, as shown in Fig. 29. The use of a remote ring anode made it possible to deliver a highly directed flux of the precursor species to the substrate, depositing them in a conformal fashion regardless of the type of substrate (Mackie, *et al.*, 2016; Mackie and Gordon, 2017). Figure 30 shows similar “agave-like” nanowire morphologies of CuO film on various substrates including conducting, insulating, flexible, patterned and fiber-based substrates, to demonstrate the versatility of this technique (Mackie, *et al.*, 2016). The deposition process was performed at 15 – 20 Torr, which is a relatively high pressure in comparison to the nominal vacuum processes while raster scanning the substrate allowed uniform deposition over larger areas. This is a viable technique that enables deposition of a variety of functional nanostructured metal oxides on a range of surfaces.

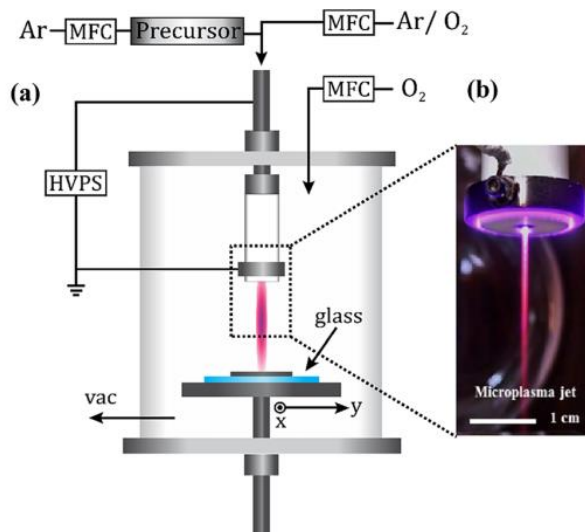


Fig. 29 (a) Schematic of the microplasma deposition system. Nickelocene, ferrocene, copper(II) acetylacetonate ($\text{Cu}(\text{acac})_2$), and copper(II) hexafluoroacetylacetonate hydrate ($\text{Cu}(\text{hfac})_2 \times \text{H}_2\text{O}$) were used as a precursor for NiO , $\alpha\text{-Fe}_2\text{O}_3$ and CuO respectively. HVPS indicates high voltage power supply (b) Photo of Ar microplasma jet at 15 Torr/8.5 mA (Mackie, *et al.*, 2016)

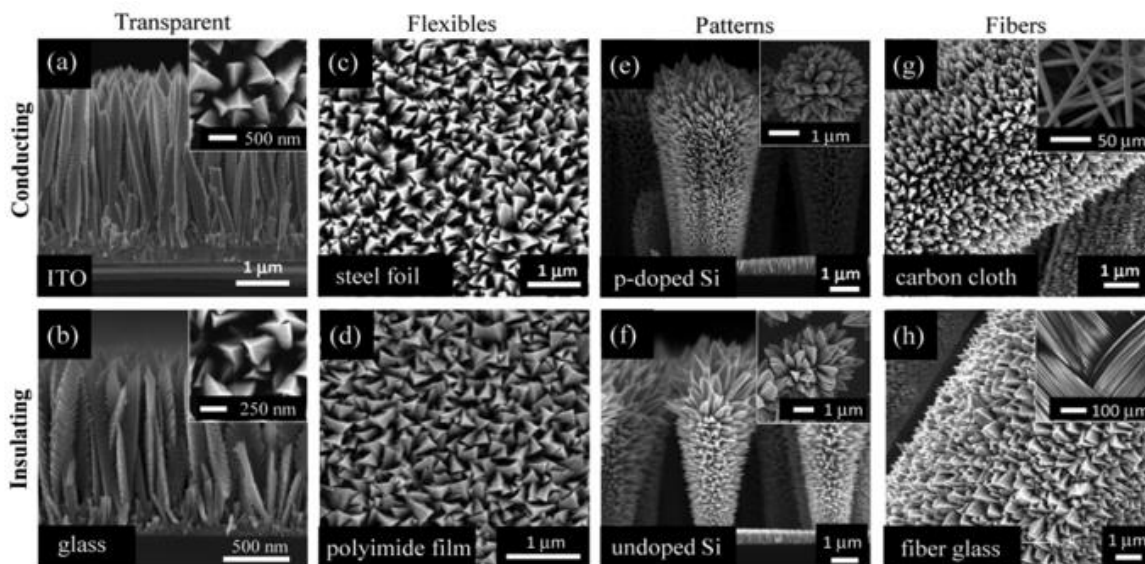


Fig. 30 Microplasma spray deposition of CuO at 20 Torr, 8.5 mA with $\text{Ar}:\text{O}_2=3:1$ on various, unheated substrates (a) ITO, (b) glass, (c) stainless steel foil, (d) Kapton polyimide film, (e) conducting Si micropillars, (f) undoped (insulating) Si micropillars, (g) carbon paper, and (h) fiberglass cloth. Insets show top-down images of the oxide films at various length scales (Mackie, *et al.*, 2016; Mackie and Gordon, 2017)

Anthony *et al.* demonstrated an all-gas-phase approach for the fabrication of nanocrystal-based light-emitting devices. Silicon nanocrystals synthesis, surface functionalization, and deposition were all performed in a single reactor. Si nanoparticles were synthesized at approximately 180 Pa in the upper part of the reactor where RF power is supplied, as shown in Fig. 31 (a). A mixture of silane and helium (5:95) was introduced for Si nanoparticle synthesis by controlled Ar flow. The diameter of the Si nanoparticles could be varied, and thus the corresponding size-dependent emission wavelength. Downstream of the plasma, in the afterglow zone of Fig. 31 (b), the synthesized Si nanoparticles were

exposed to a precursor vapor of 1-dodecene, similar to the dual-plasma-zone process discussed above. A slit-shaped polytetrafluoroethylene (PTFE) orifice shown in section (c) of Fig. 31 plays an important role of yielding a high pressure drop, which accelerates the gas flow to supersonic velocity and results in dense nanocrystal Si film formation, as shown in Fig. 31 (e) and (f). We emphasize that controlled deposition of dense and functional nanocrystal films may be suitable for application in electronic devices.

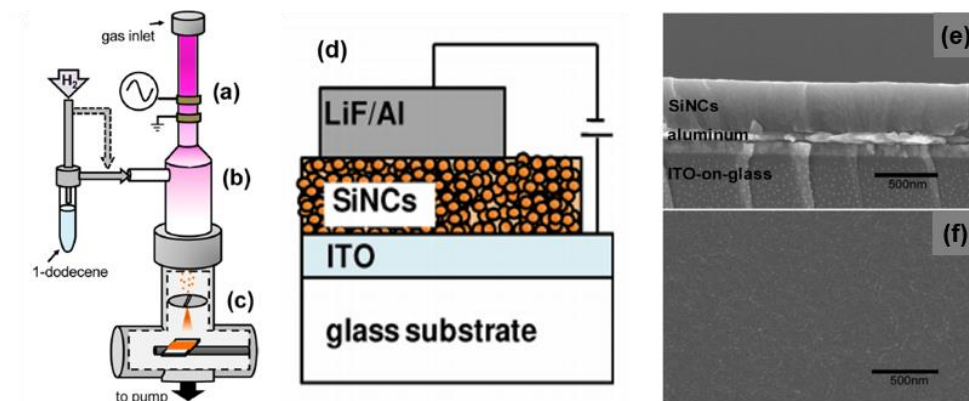


Fig. 31 Illustration of the all-gas-phase Si nanocrystal synthesis (a) Functionalization (b) Impaction (c) Scheme (d) The film structure of nanocrystal silicon-based light emitting devices. SEM images of Si nanocrystal films (e) Cross-sectional image of a SiNC layer deposited on aluminum-coated ITO-on-glass (f) Top-down view of the same SiNC film, demonstrating layer uniformity (Anthony, *et al.*, 2012)

4.3 On-surface reactions

On-surface reactions, the working principle and mechanisms of which were briefly discussed in section 3.4, can be an important element of ‘plasma-assisted’ printing of both 2D and 3D structures. It is therefore worthwhile to highlight what we can achieve from this regarding the plasma printing process. Figure 32 illustrates the patterned post-reduction of a metal precursor/polymer mixture using atmospheric-pressure micro plasma (Lee, *et al.*, 2011). The substrate is spin coated by the precursor/polymer mixture in the first step. Scanning microplasma treats the substrate selectively: only the plasma treated area will have reduced metallic material and the rest of the (unreduced) material is easily removed by the solvent. Free-standing, transparent, flexible films of patterned metal nanoparticles were fabricated by this plasma post-reduction technique (Lee, *et al.*, 2011).

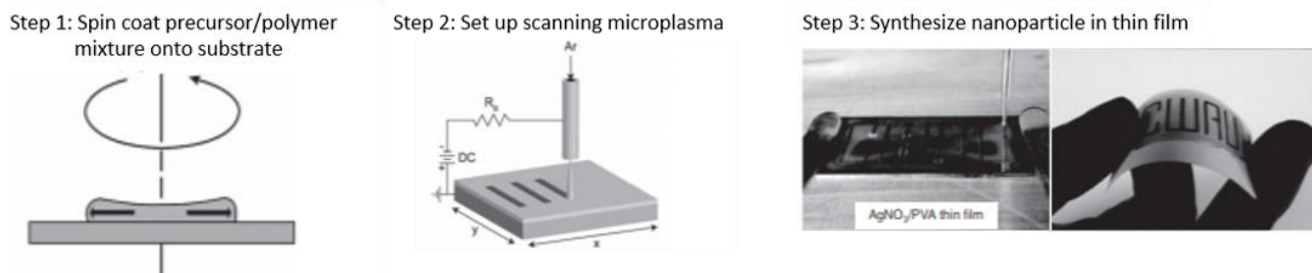


Fig. 32 Illustration of synthesis of metal nanoparticles with polymers. Free-standing, transparent, flexible films of patterned metal nanoparticles were fabricated by step1: spin-coating and drying the solution of metal salt precursor and polymer onto a Si substrate, step 2: exposing to a rastered microplasma, and step3: removing from the substrate (Lee, *et al.*, 2011)

It was recently reported that plasma–liquid interactions become visible due to immediate changes in the surface morphology of the catalytic metal as the reaction progresses through the formation of a solid-phase product from a liquid-phase precursor. The optical microscope images in Figure 33 show tailored morphologies through a catalyzed reaction of small molecules 1,2,4-trichlorobenzene (TCB), as was briefly mentioned in section 3.4. It is thought that the monomer molecule was dissociated by the plasma into reactive fragments, which then crosslinked to form an oligomeric/polymeric product that formed a thin film on the substrate (Hartl, *et al.*, 2019). The surface wettability was influenced by the plasma parameters, with high-dose plasmas leading to a strongly hydrophobic surface with water contact angles up to 130°. The deposited film had a lower stoichiometric Cl content than the intact molecule, which suggests that the plasma-assisted catalytic dehalogenation of TCB had indeed occurred. However, when the applied voltages were below 20 kV or the substrate was a non-catalytic reference (Si) material, the formation of a dehalogenated film was not achieved.

This plasma-liquid interaction on catalytic surfaces is not limited to dehalogenation, but may also be applicable to numerous other chemical reactions, providing an alternate pathway to impart specific functionality for the printed substrate or 3D object through dynamic exchange of the molecules in contact with the surface of the catalyst at low temperatures.

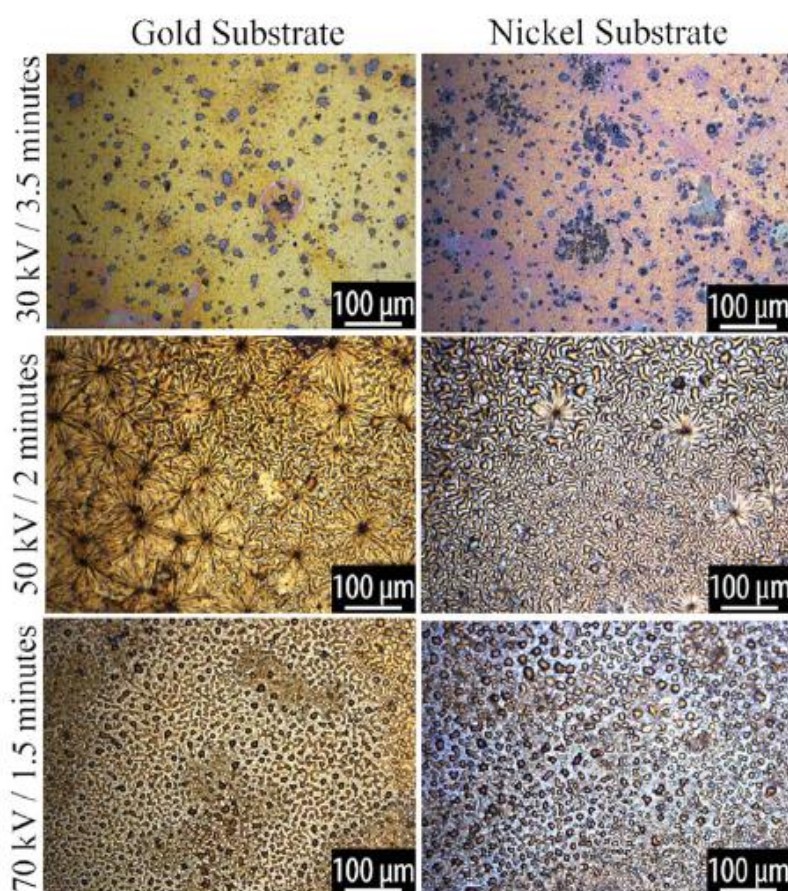


Fig. 33 Optical microscopy of plasma reacted TCB on Au and Ni susbtrates at varing power and duration but constant total dose. Voltage and time parameters are shown along the vertical axis of the figure (Hartl, *et al.*, 2019)

4.4 Feature printing

Recently, Sui et al. (Sui, *et al.*, 2019) reported a new approach for plasma assisted inkjet printing with particle-free inks composed of inorganic metal salts. The possibility of printing a range of metals was demonstrated, including both noble and non-noble metals such as Au, Ag, Pd, and Pt and Cu, Pb, Bi, and Sn, using a single-step process that involves high density RF plasma conversion of the printed structure at low pressure (650 mTorr). These inorganic salt-based inks consist of readily available halide or polyatomic ionic compounds, which are fully soluble and stable in water or organic solutions. More importantly, they are compatible with many ink additives, such as viscosity modifiers and surfactants, that may be essential for tuning the jettability and wettability of various substrates. The printed precursors are then converted into metallic structures by reaction with energetic plasma species, such as argon ions and high-energy electrons, without excessive substrate heating. Direct fabrication on a wide range of polymeric substrates, such as polyethylene terephthalate (PET), cellophane tape, polyvinyl alcohol (PVA), and polyethylene (PE) as shown in Fig. 34, is therefore possible. The printed metallic structures are polycrystalline, porous, and have resistivity ranging from $2\times$ to $10\times$ that of their respective bulk metal, which is a highly desirable electric characteristic, making the printed materials suitable for sensor applications that require large surface areas. A Bi-based trace Pb sensor, an Au-based amyloid- β 42 sensor, and an Au-based strain gauge were fabricated with proven enhanced sensitivity compared to a reference prepared with conventional methods for representative chemical, biological and mechanical sensor applications.

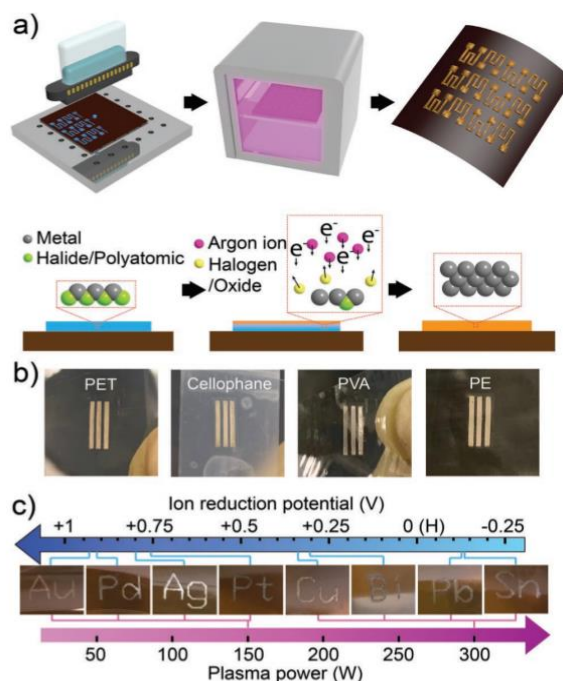


Fig. 34 (a) Process flow diagram for printing metals from inorganic metal salt-based inks by inkjet printing and plasma conversion (top) and a corresponding proposed mechanism (bottom) (b) Photos of printed and plasma-converted Ag on PET, cellophane tape, PVA, and PE substrates (c) Photos of printed and plasma-converted Au, Pd, Ag, Pt, Cu, Bi, Pb, and Sn films on polyimide organized by the reduction potentials of the corresponding inorganic metal salt precursor and plasma power required for conversion (Sui, *et al.*, 2019)

4.5 Direct plasma printing from solution precursor

A direct plasma jet printing process of graphene oxide with *in situ* reduction was recently introduced. Considering the benefit of graphene for flexible electronics and energy storage devices embedded into clothing or other flexible surfaces, a method for the direct printing of graphene from graphene oxide has drawn significant research interest. Graphene oxide is considered a suitable precursor material as it is highly processable in solution, robust and can be made conductive. Low-temperature He-H₂ plasma enabled the *in situ* reduction of a highly acidic graphene oxide suspension (pH < 2) during the deposition. X-ray Photoelectron Spectroscopy (XPS) results confirmed that the reaction intermediates and the concentration of oxygen functionalities bonded to graphene oxide were reduced significantly by this single step plasma jet printing approach. Both near edge X-ray absorption fine structure (NEXAF) and Raman spectroscopies confirmed a recovery of the *sp*² feature of graphene by this plasma jetting process in a reductive gas environment as shown in Fig. 35. Moreover, these printed graphene oxide films revealed enhanced conductivity by Electrostatic force microscopy (EFM) measurement. Hence, this technique has strong potential for printing conducting patterns of graphene oxide for a range of large-scale applications.

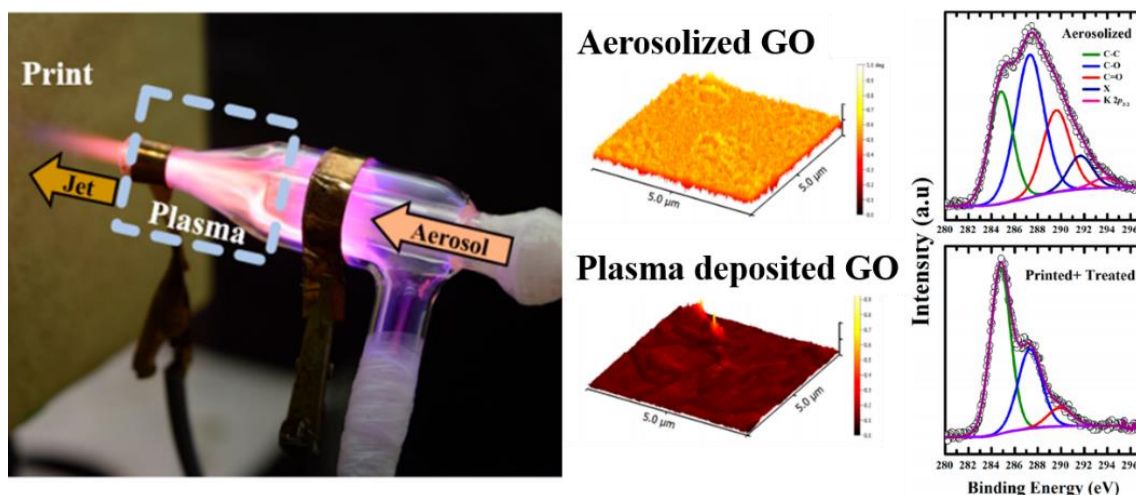


Fig. 35 (Left) a photo of plasma jet with aerosol supply of graphene oxide solution and EFM (Middle) phase map and XPS analysis results (Right) of aerosolized and plasma printed graphene oxide (GO) (Dey, *et al.*, 2018)

A direct one-step nano-gold printing process using an HAuCl₄ solution precursor with an atmospheric-pressure plasma jet was recently demonstrated (Hong, *et al.*, 2019). The system consists of an atmospheric-pressure plasma jet, atomized precursor supply and table-top 3D micro-positioning system for patterning, as shown in Fig. 36. Atomized droplets of the solution are reduced to gold nanoparticles in the plasma and deposited on a range of different substrates. The gold film has minimal Cl content and its structure can be controlled by the deposition time, from nanometer-size particles to a dense film that fully covers the substrate. The applicability of the nano-gold film as a SERS (surface-enhanced Raman scattering) platform is demonstrated by sensing 10⁻⁵ M of Rhodamine B on an Si and

paper substrate. This gold production process is readily applicable to various solution-based metal precursors, providing an easily approachable method for composite material synthesis and printing.

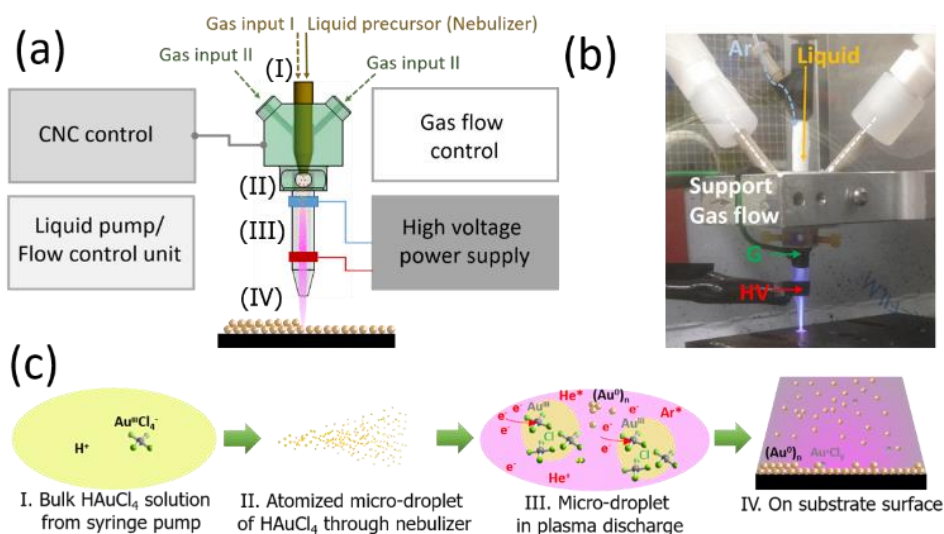


Fig. 36 (a) Schematic of the plasma printing system with atmospheric-pressure plasma jet (b) Photo of discharge showing liquid and gas input with nebulizer (c) Illustration of plasma reduction process in different parts of the system of Fig. 1(a), where Au^xCl_y indicates possible unreduced ionic gold in the discharge and on the surface e.g. $\text{Au}^{\text{III}}\text{Cl}_4^-$ or $\text{Au}^{\text{I}}\text{Cl}_2^-$ (Hong, *et al.*, 2019)

The representative plasma sources and processes discussed above are suitable for a ranges of application and can be utilized to improve the current printing technologies. In the following section 5, we will review research related to 3D structure morphing using plasma processing.

5. Plasma combined with 3D printing

Combining 3D printing with plasmas is promising to advance current printing technologies. This non-equilibrium media provides numerous effects of the plasma-excited species, which may lead to new processes with enhanced capabilities at low process temperatures.

5.1. Plasma-aided localized CVD

The use of plasma in additive manufacturing began with the simple localized technique of plasma-enhanced chemical vapour deposition (PECVD), with pioneering works by Holländer *et al.* (Hollander and Abhinandan, 2003; Abhinandan and Hollander, 2004; Silmy, *et al.*, 2005) and Shimizu *et al.* (Shimizu, *et al.*, 2003), but has since expanded. The traditional role of plasmas in CVD was to lower the deposition temperature, thus increasing the range of substrates that could be employed. In this process, the localization of the deposition can be ensured through the use of capillaries; by pulling a glass tube under controlled conditions (*i.e.* under dry nitrogen, as glass is sensitive to water, and with an accurate control of the temperature – usually satisfied by laser heating), it is relatively easy to reach reproducible aperture sizes around several hundreds of nanometres, leading to unprecedented resolution (Boileau, *et al.*, 2016).

The main advantages of these processes are:

- Possible control of the size of the basic pattern;
- The possibility to deposit any material if liquid (with a sufficiently high partial pressure) or gaseous precursors are available; and
- The simple assembly of multiple materials.

This has been successfully carried out by localized plasma modification and combined atomic layer deposition (ALD) process using μ PlasmaPrinting system developed by Innophysics (Verkuijlen, *et al.*, 2014; Mameli, *et al.*, 2017). However, this process also has certain drawbacks, the most common of which are:

- A relatively low writing speed (the highest possible deposition rate by CVD is around hundreds of micrometres per second (Hollander and Abhinandan, 2003), leading to very porous structures);
- The need for precise control of the surrounding atmosphere, which is required to limit the presence of impurities (air and water to be removed for exact control of the deposit composition); and
- A limited efficiency for treatment of large areas

At a given resolution, the writing speed is comparable to those of laser treatments or ink-jet printers but much lower than Digital Light Processing, where a digital light projector is used to flash a single image of each layer. The use of the many capillaries in parallel improves the global writing efficiency, but at the expense of a complicated control of gas injection.

To compete with conventional 3D printing solutions, plasma sources must operate at atmospheric pressure. However, this condition is not mandatory and low-pressure plasma sources might also be of interest for specific reactive gases that are highly sensitive to impurities, like silane or Trimethylaluminium for instance. Many plasma sources have already been tested as shown in Fig. 37. The use of microplasmas, whereby plasma is ignited inside a capillary tube as shown in Fig. 37 (a), is commonly believed to fail for diameters below the Debye's length, which is around 30-50 μm at atmospheric pressure. However, good results have been reported despite this. (Motrescu and Nagatsu, 2016) Plasmas could be detected outside the capillaries (beyond the tip) with diameters as small as 100 nm. It is unclear whether the plasma was also created upstream, inside the capillary (which could be explained by an increase in the gas pressure and the subsequent lowering of the Debye's length) or if the capillary served as propagation structure for the pulsed electric field and provided outer excitation of the plasma. Nonetheless, this approach likely creates microscopic plasmas and not nanoscale plasmas.

Consequently, the use of a capillary to direct the flux of a reactive gas onto a surface for plasma activation is by far more convenient than creating the plasma inside the capillary. Indeed, separating the plasma unit and the precursor delivery system suppresses clogging at the tip by disabling deposition inside the capillary. Ideally, the plasma unit and the precursor supply should be separated. When the plasma formation and the injection of the precursor are dealt with separately, the choice of plasma source

must be done according to the deposit that is sought. The use of electric arcs as plasma sources as shown in Fig. 37 (b) is very challenging for a PECVD process. Maurau et al.(Maurau, *et al.*, 2013) used titanium tetraisopropoxide (TTIP) as titanium precursor, introducing it in the afterglow of a blown arc generated with a 100 kHz sinusoidal voltage. The dissipated power was kept constant at 600 W, with a 30 L min⁻¹ flow rate of nitrogen and 5 μ L min⁻¹ of precursor. A localized coating was obtained, characterized by a very large porosity and TiO₂-nanocrystallites embedded in an amorphous phase. Using such a high power to coat an area of a few millimetres in diameter requires the system to be operated at a very low precursor flow rate, even when the precursor is introduced downstream of the arc. However, even with this compensation, the formation of powders is still an issue.

The majority of early systems employed in plasma-activated 3D printing used microwave plasmas operated at atmospheric pressure as shown in Fig. 38 (c) (Belmonte, *et al.*, 2011b; Boileau, *et al.*, 2016), most likely due to its high power density and the ease with which they can be implemented as electrodeless configuration. Microwave plasmas provide an important heating element, which hinders the use of labile substrates but favours the synthesis of dense materials that are contaminant-free due to the absence of electrodes. Dielectric barrier discharges are usually preferred for labile substrates; however, the electrode arrangement must be adapted to create access to the capillary. As shown in Fig. 37 (d) after μ PlasmaPrinting system of Innophysics, a plasma head configuration with multi-pin electrode array offers a unique advantage. It has a 24-electrode-array printing head that enables localized atomic layer deposition (ALD)-like deposition at atmospheric pressure and argon shielding curtains that allow the system to be operated in a confined atmosphere.

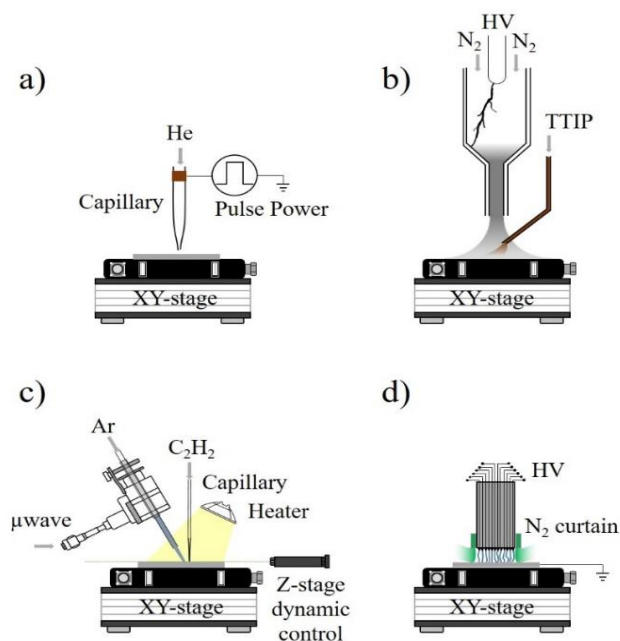


Fig. 37 Examples of plasma-aided 3D printers. (a) Plasma-bullet type system with nanometric aperture capillary(Motrescu and Nagatsu, 2016) (b) Blown arc system with downstream injection of precursor such as titanium isopropoxide (TTIP) for TiO_x film deposition (Maurau, *et al.*, 2013) (c) Surfatron-type microwave plasma with nanometric aperture capillary after Boileau et al.(Boileau, *et al.*, 2016) (d) Multi-pin- electrode array plasma head with nitrogen curtain aside after μ PlasmaPrinting system of Innophysics (Mameli, *et al.*, 2017)

High quality films with a high level of control in terms of thickness, opto-electrical properties and patterning resolution, can thus be synthesized.

Surface barrier discharges are suited to plasma-aided 3D printing as they offer easy access to the activated surface. However, the flux of ions parallel to the deposition surface creates an ion wind that affects the shape of the elementary deposition spot, *i.e.* the coating obtained in a static mode, changing it from a circle to a sort of ellipse. It was shown that the development of an ion wind can be attributed to the electrohydrodynamic force per unit volume that appears in electric surface discharge due to momentum transfer from charged particles to neutral particles (Boeuf, *et al.*, 2007). This observation, also made on a pulsed plasma jet (Park, *et al.*, 2018), raises the more general question of the influence of the flow pattern on the shape of the elementary spot. This aspect was discussed earlier (Belmonte, *et al.*, 2011a; Belmonte, *et al.*, 2011b) who stated that when a separate capillary is used to inject the precursor, the mixing of this precursor with the plasma gas plays a key role in the deposition process. Indeed, the simultaneous presence of both kinds of species on the surface of the substrate is needed for deposition. Since laminar diffusion is very limited at atmospheric pressure, plasma species have to penetrate the jet that contains the precursor by another mechanism in this configuration. Numerical calculations of the flow pattern indicate that species coming from the plasma have a mean velocity that is at least one order of magnitude lower than that of the precursor. Consequently, the transport of species from the plasma to the surface, followed by the species mixing with the precursor, is ensured by the slight turbulence caused by the high velocity of the jet flow. This conclusion provides a fundamental condition to guarantee the simultaneous presence of reactive species on the substrate. Of course, turbulence must not be too strong as this can cause recirculation of the flow, long residence times and powder synthesis. (Belmonte, *et al.*, 2011a; Belmonte, *et al.*, 2011b)

The control of the distance H from the tip of the capillary to the surface is crucial but challenging to monitor. The fluid flow of a jet exiting a capillary of diameter D and impinging a wall on a spot submitted to thermal gradients is very complex (Grenson, *et al.*, 2016; van Hout, *et al.*, 2018). Most important is the formation of a stagnation region around the point where the jet impinges the surface, the size of which increases when H/D increases, typically beyond $H/D > 3$. (Wienand, *et al.*, 2017) In this region, deposition is very fast due to the accumulation of precursors, and so it is recommended to operate the process at the capillary-to-surface distance that verifies $H/D < 3$.

During PECVD, surface buckling and roughness can occur; thus height adjustment is necessary to combat these issues. This issue is the same as when using hollow cantilevers employed in AFM/STM devices for localized fluid dispensing, where tips must be located very close to the surface to scan it. (Ghatkesar, *et al.*, 2014) This method is well mastered today but is costly and is therefore only worthwhile if the object to build is large enough with respect to the resolution. For a capillary of, say, 100 nm, not only must the X and Y movements be controlled at the nanometre scale, but also Z as shown

in Fig. 38 (c). (Boileau, *et al.*, 2016) This kind of stage usually combines two systems: one mechanical for fast moves and one piezoelectric for slow moves, and it must be coupled to an accurate measurement of the height and therefore to an active vibration isolation table.

Another difficulty arises if the substrate conductivity changes under the plasma flow by deposition of an insulating material, making the deposition area drift in an uncontrolled manner. For this reason, resorting to remote plasmas removes the constraint of operating with charges. The flow of the remote plasma must be chosen weak enough with respect to the precursor so as to keep the shape of the precursor jet while enabling transport of metastable species within the jet to activate deposition.

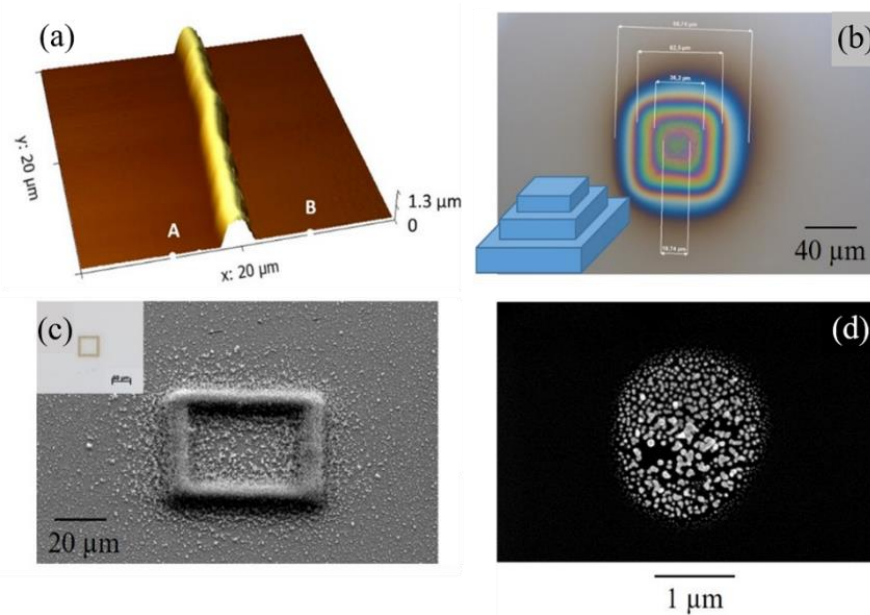


Fig. 38 (a) AFM image of a hydrogenated carbon line deposited with C_2H_2 (Boileau, *et al.*, 2016) (b) Gold spot deposited in a static mode at 623K with $HAuCl_4$ (c) SiO_x rectangular pattern deposited with SiH_4 (d) Top view of a hydrogenated carbon pyramid deposited with C_2H_2 . (b)-(d) are from unpublished works by Thierry Belmonte and colleagues.

Importantly, materials deposited by plasma-aided 3D printing are sensitive to deposition conditions. Plasma-aided 3D printing must overcome the same issues as thermal CVD: dewetting, adhesion, development of stress in the films, etc. Fig. 38(a) shows a hydrogenated carbon line obtained with C_2H_2 using an argon microwave plasma (Boileau, *et al.*, 2016). The presence of air traces in the surrounding atmosphere is useful in this situation as it results in a sharp and well-defined pattern. The oxidizing species completely prevent the presence of splats around the line by suppressing the synthesis of particles in the gas phase, enabling only surface deposition processes.

Consequently, building 3D objects like pyramids gives excellent results, as shown in Fig. 38 (d). The top-view optical image of the pyramid illustrates the sharpness of each step edge whereas the colour, due to interference, shows the high reproducibility of the step thickness from one step to another. On the contrary, in Fig. 38 (c), a rectangular pattern produced with SiH_4 in nitrogen containing 0.1% of O_2 and 2% of relative humidity shows many splats. Fourier Transform Infrared (FTIR) spectroscopy measurements indicate the presence of both Si-H and Si-O bonds, demonstrating a partial oxidation of

the silicon coating. Although it is hard to observe in Fig. 38 (c), the thickness of the line is greater at each corner of the rectangle. This is because the capillary spends longer at spots where one line ends and another one starts. This factor has to be accounted for maintaining the constant height. In Fig. 38(b), deposition of a gold spot was realized from a solution of $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ (8×10^{-4} M). Below 300°C , gold crystallites in the pattern correspond to AuClO . Above this temperature, they are reduced into metallic gold. The dewetting of the coating on the silicon wafer underneath is clearly visible and is a common observation if no accommodation layer is grown in-between. This behaviour can be easily prevented by increasing the surface wettability using a plasma pre-treatment, which gives better printing performance.

We note that it is difficult to form some specific shapes straightforwardly. For example, to build a candy cane-like shape, the opposite parts of the U-shape handle have to be connected together to fill the gap between the two branches; following this, connectors must be removed.

5.2. Nanoparticles assembly

As CVD-based processes are efficient for producing nanoparticles, plasmas can also be used as non-equilibrium sources of nanoparticles for *in situ* production as we discussed above in section 4.1. This avoids the need for *ex situ* handling of hazardous nanoparticles. Powder sintering is usually performed either locally by a laser in 3D-printing processes or generally in volume by thermal annealing. In these cases, using plasma to assemble nanoparticles into larger 3D structures is not particularly beneficial. However, direct plasma synthesis of nanoparticles from precursor materials or in-flight functionalization before assembly or 3D structure formation is ideal to maximize the benefit of plasma processing.

A promising process in this field (Hafiz, *et al.*, 2004; Hafiz, *et al.*, 2006a; Hafiz, *et al.*, 2006b; Girshick, 2008) is based on an idea that is widely applied in the synthesis of nanoparticles: a high-pressure chamber is used to generate nanoparticles that are ballistically extracted through a nozzle (Hafiz, *et al.*, 2006b) or a set of aerodynamic lenses (Girshick, 2008) into a low-pressure chamber as shown in Fig. 39 (a). Figure 39 (b) shows a possible nanostructured patterning system using this hypersonic plasma particle deposition technique. It was demonstrated that it is possible to control and focus nanoparticles (< 30 nm) into tightly collimated beams using the optimized aerodynamic lens systems (Wang, *et al.*, 2005a; Wang, *et al.*, 2005b). Figure 39 (c) is a sketch of a possible nanoparticle-based manufacturing assembly line, in which various types of micro- and nano-scale structures are constructed from focused nanoparticle beams using arrays of aerodynamic lenses.

Figure 40(a) shows that it is possible to obtain a high aspect ratio nanostructured tower-like structure using the lens system. Michelakaki *et al.* (Michelakaki, *et al.*, 2018) demonstrated that the porosity of the film prepared by this nanoparticle beam cluster technique can be significantly improved by applying voltage on the substrate. A fine pattern as small as 400 nm linewidth was realized by adapting lithography process as shown in Fig. 40 (b). However, the deposited film inevitably contains porous

structure even at the highest applied voltage (4.5 kV) condition. It is considered that even with hypersonic impact (estimated around 1700 m/s), the energy released by collision is insufficient to fuse the nanoparticles. High temperatures are subsequently required to form dense structures, which may limit the application of these sources. This strategy has been applied to produce an instant coating of SiC by heating the substrate above 750°C (Hafiz, *et al.*, 2006b). The resolution of the printed pattern is determined by the width of the nanoparticles beam, which is a few tens of μm .

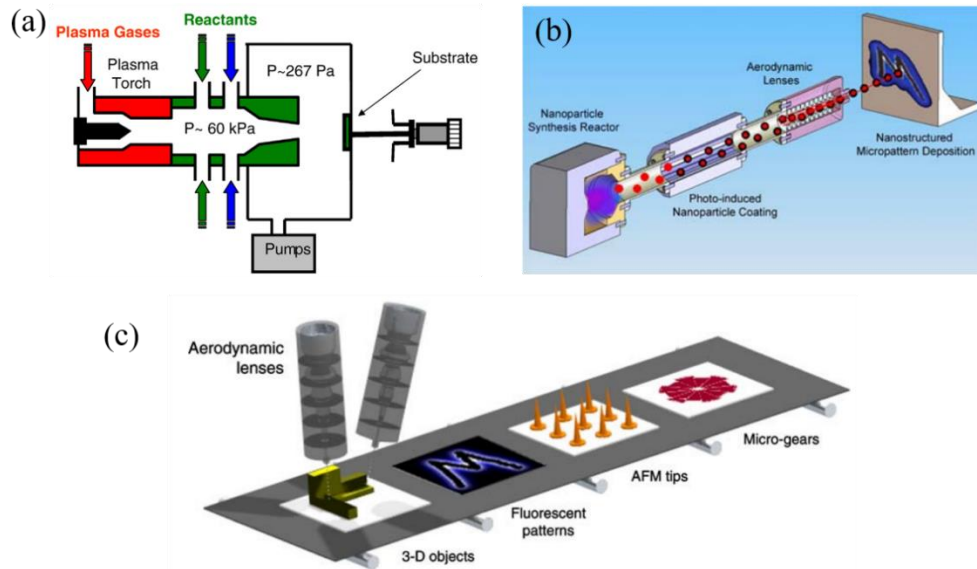


Fig. 39 (a) Schematic of hypersonic plasma particle deposition system (Hafiz, *et al.*, 2006a) (b) Illustration of hypothetical integrated nanoparticle process stream, including nanoparticle synthesis, nanoparticle coating by photo-CVD, nanoparticle focusing by aerodynamic lenses, and deposition of nanoparticles to form micropatterns (Girshick, 2008) (c) Illustration of conceptual nanoparticle-based manufacturing assembly line, in which various types of MEMS-scale objects are constructed from focused nanoparticle beams using arrays of aerodynamic lenses (Girshick, 2008)

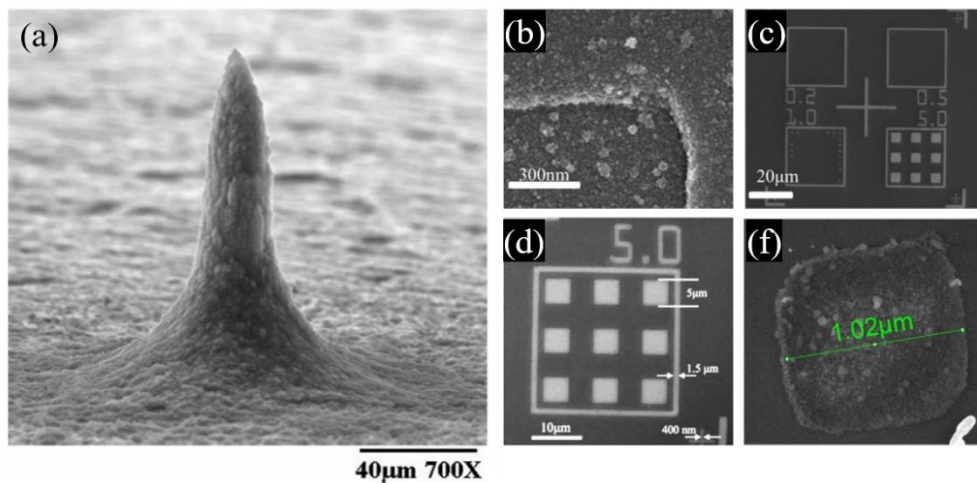


Fig. 40 SEM image (a) Tower-like 3D structure formation of SiC nanoparticles deposited by focused particle beam on a stationary substrate (Hafiz, *et al.*, 2006b), 3D patterns composed of Hf nanoparticles deposited at a substrate voltage of $V_s = 4.5 \text{ kV}$: (b) The high-magnification image shows the granular structure of these patterns; (c,d) the areas that appear white are structures formed from the nanoparticles. Large square structures of 5 μm size were successfully constructed (e) Cohesive structure with 1 μm side found at a random position on the substrate (Michelakaki, *et al.*, 2018).

The use of plasma to produce nanoparticles that can be assembled into 3D structures has been exploited utilizing a nanoxerographic process (Barry, *et al.*, 2003; Barry, *et al.*, 2005). In this process, localized deposits are formed from nanoparticles that are deflected by a high voltage applied on a patterned nano-electrode. The plasma is used only as a monodisperse source of charged nanoparticles. The resolution is spectacular, the deposited patterns having a characteristic dimension of about 100 nm.

Although plasmas has been successfully employed for the *in situ* production of nanoparticles in 3D printing processes, there are still obstacles to overcome in this area; in each of the discussed examples, monodisperse distributions of nanoparticles at given mean size are still required (Hafiz, *et al.*, 2006b).

5.3. Plasma treatment of 3D objects

There are several options for atmospheric-pressure plasma processing of 3D objects. The plasmas can either be mounted directly on a 3D printer head or used as a post-printing technique to clean, functionalize or reduce the roughness of surfaces. A few representative examples are discussed below.

The Flashfuse filament technology is a 3D printing solution commercialized by Essentium, Inc.(Essentium) as we briefly introduced in section 1. The polymer or composite to be deposited is introduced through a nozzle in the centre of a conductor disk positioned above the grounded 3D printed part to be built. Within the gap between the two surfaces, an air plasma is ignited. The whole system is heated to improve the interface exchange mechanisms. Therefore, the issue of delamination, which usually plagues pieces formed by conventional fused deposition discussed in section 2, is solved. Filaments, made of Polyether Ether Ketone (PEEK) or carbon fibre reinforced polymers for instance, are coated with conductive carbon nanotubes to weld each printed cross section to the previously printed layers. By addressing the bonding of the polymer chains at the nanoscale level, larger heat affected zones are created, thereby improving the strength of the cross-sectional welds and the resulting strength of the parts. A result of this is that polymer chains can better migrate and entangle. This technology was mentioned in relation to building the solar cell umbrellas that are expected to provide long-term photovoltaic power on Mars (Adams, *et al.*, 2018).

Another example is the PDD® Plasma technology (Piezoelectric Direct Discharge) by Relyon Plasma which can be integrated directly into 3D printers (Plasma). A voltage supply of 15 V is needed for the compact plasma module (piezobrush® PZ2), which typically has a power consumption of less than 15 W. In typical applications on the basis of conventional FDM technology, this simple system can change the surface wettability of the deposited polymer, leading to the improvements in the gluing or painting steps.

As we discussed earlier, it is also possible to directly deposit other nanomaterials such as carbon nanotubes or nanogold (Gandhiraman, *et al.*, 2016)(Hong, *et al.*, 2019). A low-temperature plasma jet

with emulsion of nanoparticle or solution-based precursor supply can be effective to coat 3-D objects and flexible surfaces, such as paper or cloth, with a functional nanomaterial or even composite.

Plasma-modified polymers used as a scaffold for biomedical applications have been intensively studied (Cools, *et al.*, 2018; Joshy, *et al.*, 2019). Indeed, surface modification using plasmas is a highly exploited and productive technique in tissue engineering and regenerative medicine. Plasma modification is known to improve surface properties and thereby enhance the biocompatibility of the materials involved. As we earlier mentioned in section 1, Fraunhofer Institute for Applied Polymer Research (IAP) and Thin Films (IST) developed a plasma jet technique for 3D Printing customized bone implants, which couples both aspects – plasma treatment and 3D printing – in a single solution (Fraunhofer, 2018). The device blows a cold jet of plasma containing reactive groups directly onto the 3D printed layers. Amino groups present in the precursor bond with the surface and ensure adhesion of bone cells to the substrate. No chemical pre-treatment with solvents is required for the coating, so the procedure is both cost-effective and environmentally friendly. Wang *et al.* (Wang, *et al.*, 2016b) showed that cold atmospheric plasmas modify the nanoscale roughness and chemical composition of a 3D printed poly-lactic-acid scaffold surface, thus promoting both osteoblast (bone forming cells) and mesenchymal stem cell attachment and proliferation.

Perelaer *et al.* (Perelaer, *et al.*, 2012) developed a mixed system combining plasma and microwave flash sintering to sinter an inkjet-printed and tailored silver nanoparticle formulation. The processing temperature remains well below the glass transition temperature of the used polymer substrate. This approach leads to highly conductive features, which are compatible with R2R but also with 3D-printing production.

6. Outlook and Conclusion

It is evident that low-temperature, atmospheric-pressure plasmas can play an impactful role in additive manufacturing and other areas of advanced digital manufacturing. The recent parallel development of both 3D printing and atmospheric plasma technologies offers the feasibility to combine the approaches with a view to producing new and highly controlled materials, structures, and devices.

The diverse nature of the physics and chemistry of plasmas and their multi-modal effects means that there is a vast array of operational variables which need to be examined and optimised for specific AM applications. Compounding this complexity is the continued evolution of multi-material additive manufacturing (MM-AM) which offers complex multi-component products. The behaviour of different materials as they interface with both each other and the plasma during the printing process needs to be understood. As outlined in this paper, there are numerous integration approaches for combining the technologies which need to be classified and ideally standardised for plasma printing. Integration of plasma sources need to not only take into consideration the ease of combining the printing process with

the discharge but to also the plasma induced effects which may influence the process such as gas flow effects, reactive chemistry with the surrounding atmosphere, and other factors.

The ability to effectively scale the process is a key challenge, particularly where large volumes or dimensions are sought with features that are set to a micro or indeed nano-scale level. The time-scale of these processes should also be considered and matched with the induced reactions.

Given the scales and resolution required by advanced manufacturing, effective control of the plasma-printing process is required. Non-invasive and real-time plasma diagnostic techniques could be integrated within the plasma-printing process to provide both mechanistic insights, process tuning and control. The potential to link high resolution imaging of the printing process to plasma data would provide useful information to drive technology evolution in this space.

It is clear from the rapidly emerging literature and indeed the few commercial offerings on the market that the integration of non-thermal plasmas with printing processes is an active area of research with a wide range of potential end applications. The combination of plasma and 3D printing has the potential to transform processes across a wide range of industries, creating products that significantly enhance both science and society.

Many limitations of current processes can be overcome by combining plasma with 3D printing. However, these techniques are still in their early days and require further research to improve and expand their use. This is why we sincerely hope that our effort to critically examine the recent progress and potential of atmospheric-pressure plasmas, and plasma nanotechnology in particular, to help advance the rapidly developing digital manufacturing – the technology which is poised to sustain the next industrial revolution in the digital manufacturing age – will stimulate information exchange and collaborative efforts among the scientists and engineers normally working in disparate fields.

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REFERENCES

- Press release. Fraunhofer Institute for Applied Polymer Research IAP and Thin Films IST
<https://www.fraunhofer.de/content/dam/zv/en/press-media/2018/december/research-news/rn12-2018-IST-precisely-fitting-bone-implants-from-the-printer.pdf>
- L. Abhinandan, and A. Hollander, Localized deposition of hydrocarbon using plasma activated chemical vapour deposition, *Thin Solid Films* **457**, 241-245 (2004)
- G. Adams, J. Banks, C. Frazier, U. Toodi, and M. Lagoudas, 2018, in (Texas A&M University).
- J. Alaman, R. Alicante, J. I. Pena, and C. Sanchez-Somolinos, Inkjet Printing of Functional Materials for Optical and Photonic Applications, *Materials* **9**, 910 (2016)
- R. J. Anthony, K. Y. Cheng, Z. C. Holman, R. J. Holmes, and U. R. Kortshagen, An All-Gas-Phase Approach for the Fabrication of Silicon Nanocrystal Light-Emitting Devices, *Nano Lett.* **12**, 2822-2825 (2012)
- I. Bahnini, M. Rivette, A. Rechia, A. Siadat, and A. Elmesbahi, Additive manufacturing technology: the status, applications, and prospects, *Int J Adv Manuf Tech* **97**, 147-161 (2018)
- C. R. Barry, J. Gu, and H. O. Jacobs, Charging process and coulomb-force-directed printing of nanoparticles with sub-100-nm lateral resolution, *Nano Lett.* **5**, 2078-2084 (2005)
- C. R. Barry, N. Z. Lwin, W. Zheng, and H. O. Jacobs, Printing nanoparticle building blocks from the gas phase using nanoxerography, *Appl. Phys. Lett.* **83**, 5527-5529 (2003)
- T. Belmonte, T. Gries, R. P. Cardoso, G. Arnoult, F. Kosior, and G. Henrion, Chemical vapour deposition enhanced by atmospheric microwave plasmas: a large-scale industrial process or the next nanomanufacturing tool?, *Plasma Sources Sci T* **20**, 024004 (2011)
- T. Belmonte, G. Henrion, and T. Gries, Nonequilibrium Atmospheric Plasma Deposition, *J. Therm. Spray Technol.* **20**, 744-759 (2011)
- J. P. Boeuf, Y. Lagmich, T. Unfer, T. Callegari, and L. C. Pitchford, Electrohydrodynamic force in dielectric barrier discharge plasma actuators, *Journal of Physics D-Applied Physics* **40**, 652-662 (2007)
- A. Boileau, T. Gries, C. Noel, R. P. Cardoso, and T. Belmonte, Sub-micro a-C:H patterning of silicon surfaces assisted by atmospheric-pressure plasma-enhanced chemical vapor deposition, *Journal of Physics D-Applied Physics* **49**, 445306 (2016)
- A. Botman, J. J. L. Mulders, and C. W. Hagen, Creating pure nanostructures from electron-beam-induced deposition using purification techniques: a technology perspective, *Nanotechnology* **20**, 372001 (2009)
- M. I. Boulou, The Role of Transport Phenomena and Modeling in the Development of Thermal Plasma Technology, *Plasma Chem. Plasma Process.* **36**, 3-28 (2016)
- T. Bret, I. Utke, C. Gaillard, and P. Hoffmann, Periodic structure formation by focused electron-beam-induced deposition, *Journal of Vacuum Science & Technology B* **22**, 2504-2510 (2004)
- D. Chakravarty, C. S. Tiwary, C. F. Woellner, S. Radhakrishnan, S. Vinod, S. Ozden, P. A. D. Autreto, S. Bhowmick, S. Asif, S. A. Mani, *et al.*, 3D Porous Graphene by Low-Temperature Plasma Welding for Bone Implants, *Advanced Materials* **28**, 8959-8967 (2016)
- K. Cheng, M. H. Yang, W. W. W. Chiu, C. Y. Huang, J. Chang, T. F. Ying, and Y. Yang, Ink-jet printing, self-assembled polyelectrolytes, and electroless plating: Low cost fabrication of circuits on a flexible substrate at room temperature, *Macromol. Rapid Commun.* **26**, 247-264 (2005)
- J. W. Choi, E. MacDonald, and R. Wicker, Multi-material microstereolithography, *Int J Adv Manuf Tech* **49**, 543-551 (2010)
- L. Chong, S. Ramakrishna, and S. Singh, A review of digital manufacturing-based hybrid additive manufacturing processes, *Int J Adv Manuf Tech* **95**, 2281-2300 (2018)
- A. Clausen, F. W. Wang, J. S. Jensen, O. Sigmund, and J. A. Lewis, Topology Optimized Architectures with Programmable Poisson's Ratio over Large Deformations, *Advanced Materials* **27**, 5523-5527 (2015)
- P. Cools, C. Mota, I. Lorenzo-Moldero, R. Ghobeira, N. De Geyter, L. Moroni, and R. Morent, Acrylic Acid Plasma Coated 3D Scaffolds for Cartilage tissue engineering applications, *Sci. Rep.* **8**, 3830 (2018)
- E. A. Corbin, L. J. Millet, J. H. Pikul, C. L. Johnson, J. G. Georgiadis, W. P. King, and R. Bashir, Micromechanical properties of hydrogels measured with MEMS resonant sensors, *Biomed. Microdevices* **15**, 311-319 (2013)
- R. d'Agostino, P. Favia, C. Oehr, and M. R. Wertheimer, Low-temperature plasma processing of materials: Past, present, and future, *Plasma Process Polym* **2**, 7-15 (2005)
- B. J. de Gans, P. C. Duineveld, and U. S. Schubert, Inkjet printing of polymers: State of the art and future developments, *Advanced Materials* **16**, 203-213 (2004)

- T. DebRoy, H. L. Wei, J. S. Zuback, T. Mukherjee, J. W. Elmer, J. O. Milewski, A. M. Beese, A. Wilson-Heid, A. De, and W. Zhang, Additive manufacturing of metallic components - Process, structure and properties, *Prog. Mater. Sci.* **92**, 112-224 (2018)
- A. Dey, S. Krishnamurthy, J. Bowen, D. Nordlund, M. Meyyappan, and R. P. Gandhiraman, Plasma Jet Printing and in Situ Reduction of Highly Acidic Graphene Oxide, *ACS Nano* **12**, 5473-5481 (2018)
- Essentium, <http://essentium3d.com>
- H. Exner, M. Horn, A. Streek, P. Regenfass, F. Ullmann, and R. Ebert, Laser Micro Sintering - A new method to generate metal and ceramic parts of high resolution with sub-micrometer powder, *Virtual and Physical Prototyping* **3**, 3-11 (2008)
- R. D. Farahani, M. Dube, and D. Theriault, Three-Dimensional Printing of Multifunctional Nanocomposites: Manufacturing Techniques and Applications, *Advanced Materials* **28**, 5794-5821 (2016)
- S. Felton, M. Tolley, E. Demaine, D. Rus, and R. Wood, A method for building self-folding machines, *Science* **345**, 644-646 (2014)
- S. R. Forrest, The path to ubiquitous and low-cost organic electronic appliances on plastic, *Nature* **428**, 911-918 (2004)
- H. Forster, C. Wolfrum, and W. Peukert, Experimental study of metal nanoparticle synthesis by an arc evaporation/condensation process, *J. Nanopart. Res.* **14**, 926 (2012)
- J. D. Fowlkes, R. Winkler, B. B. Lewis, M. G. Stanford, H. Plank, and P. D. Rack, Simulation-Guided 3D Nanomanufacturing via Focused Electron Beam Induced Deposition, *Acs Nano* **10**, 6163-6172 (2016)
- K. Fricke, H. Steffen, T. von Woedtke, K. Schroder, and K. D. Weltmann, High Rate Etching of Polymers by Means of an Atmospheric Pressure Plasma Jet, *Plasma Process Polym* **8**, 51-58 (2011)
- A. Frutiger, J. T. Muth, D. M. Vogt, Y. Menguc, A. Campo, A. D. Valentine, C. J. Walsh, and J. A. Lewis, Capacitive Soft Strain Sensors via Multicore-Shell Fiber Printing, *Advanced Materials* **27**, 2440-2446 (2015)
- P. Galliker, J. Schneider, H. Eghlidi, S. Kress, V. Sandoghdar, and D. Poulikakos, Direct printing of nanostructures by electrostatic autofocussing of ink nanodroplets, *Nat Commun* **3**, 890 (2012)
- R. P. Gandhiraman, E. Singh, D. C. Diaz-Cartagena, D. Nordlund, J. Koehne, and M. Meyyappan, Plasma jet printing for flexible substrates, *Appl. Phys. Lett.* **108**, 123103 (2016)
- M. Gavagnin, H. D. Wanzenboeck, S. Wachter, M. M. Shawrav, A. Persson, K. Gunnarsson, P. Svedlinth, M. Stoger-Pollach, and E. Bertagnolli, Free-Standing Magnetic Nanopillars for 3D Nanomagnet Logic, *Acs Appl Mater Inter* **6**, 20254-20260 (2014)
- H. C. George, T. A. Orlova, A. O. Orlov, and G. L. Snider, Novel method for fabrication of nanoscale single-electron transistors: Electron beam induced deposition of Pt and atomic layer deposition of tunnel barriers, *Journal of Vacuum Science & Technology B* **29**, 06FB01 (2011)
- M. K. Ghatkesar, H. H. P. Garza, F. Heuck, and U. Staufer, Scanning Probe Microscope-Based Fluid Dispensing, *Micromachines* **5**, 954-1001 (2014)
- S. L. Girshick, Aerosol processing for nanomanufacturing, *J. Nanopart. Res.* **10**, 935-945 (2008)
- A. S. Gladman, E. A. Matsumoto, R. G. Nuzzo, L. Mahadevan, and J. A. Lewis, Biomimetic 4D printing, *Nat Mater* **15**, 413 (2016)
- N. Y. M. Gonzalez, M. El Morsli, and P. Proulx, Production of Nanoparticles in Thermal Plasmas: A Model Including Evaporation, Nucleation, Condensation, and Fractal Aggregation, *J. Therm. Spray Technol.* **17**, 533-550 (2008)
- P. Grenson, O. Leon, P. Reulet, and B. Aupoix, Investigation of an impinging heated jet for a small nozzle-to-plate distance and high Reynolds number: An extensive experimental approach, *Int. J. Heat Mass Transfer* **102**, 801-815 (2016)
- D. J. Guo, R. Kometani, S. Warisawa, and S. Ishihara, Growth of ultra-long free-space-nanowire by the real-time feedback control of the scanning speed on focused-ion-beam chemical vapor deposition, *Journal of Vacuum Science & Technology B* **31**, 061601 (2013)
- J. Y. Guo, X. B. Fan, R. Dolbec, S. W. Xue, J. Jurewicz, and M. Boulos, Development of Nanopowder Synthesis Using Induction Plasma, *Plasma Sci Technol* **12**, 188-199 (2010)
- J. Hafiz, R. Mukherjee, X. Wang, J. V. R. Heberlein, P. H. McMurry, and S. L. Girshick, Analysis of nanostructured coatings synthesized by ballistic impaction of nanoparticles, *Thin Solid Films* **515**, 1147-1151 (2006)
- J. Hafiz, R. Mukherjee, X. Wang, P. H. McMurry, J. V. R. Heberlein, and S. L. Girshick, Hypersonic plasma particle deposition - A hybrid between plasma spraying and vapor deposition, *J. Therm. Spray Technol.* **15**, 822-826 (2006)

- J. Hafiz, X. Wang, R. Mukherjee, W. Mook, C. R. Perrey, J. Deneen, J. V. R. Heberlein, P. H. McMurry, W. W. Gerberich, C. B. Carter, *et al.*, Hypersonic plasma particle deposition of Si-Ti-N nanostructured coatings, *Surf Coat Tech* **188**, 364-370 (2004)
- H. Hartl, Y. R. Guo, K. Ostrikov, Y. B. Xian, J. Zheng, X. G. Li, K. E. Fairfull-Smith, and J. MacLeod, Film formation from plasma-enabled surface-catalyzed dehalogenative coupling of a small organic molecule, *Rsc Adv* **9**, 2848-2856 (2019)
- D. Herzog, V. Seyda, E. Wycisk, and C. Emmelmann, Additive manufacturing of metals, *Acta Mater.* **117**, 371-392 (2016)
- L. Hirt, A. Reiser, R. Spolenak, and T. Zambelli, Additive Manufacturing of Metal Structures at the Micrometer Scale, *Advanced Materials* **29**, 1604211 (2017)
- A. Hollander, and L. Abhinandan, Localized deposition by mu-jet-CVD, *Surf Coat Tech* **174**, 1175-1177 (2003)
- J. Hong, S. Yick, E. Chow, A. Murdock, J. Fang, D. H. Seo, A. Wolff, Z. Han, T. van der Laan, A. Bendavid, *et al.*, Direct plasma printing of nano-gold from an inorganic precursor, *J Mater Chem C* **7**, 6369 (2019)
- J. L. Hu, H. P. Meng, G. Q. Li, and S. I. Ibekwe, A review of stimuli-responsive polymers for smart textile applications, *Smart Mater. Struct.* **21**, 053001 (2012)
- K. I. Hunter, J. T. Held, K. A. Mkhoyan, and U. R. Kortshagen, Nonthermal Plasma Synthesis of Core/Shell Quantum Dots: Strained Ge/Si Nanocrystals, *Acs Appl Mater Inter* **9**, 8263-8270 (2017)
- Innophysics, <http://www.innophysics.nl/index.php/projects/plasmaprint-ald>
- E. Jager, J. Schmidt, A. Pfuch, S. Spange, O. Beier, N. Jager, O. Jantschner, R. Daniel, and C. Mitterer, Antibacterial Silicon Oxide Thin Films Doped with Zinc and Copper Grown by Atmospheric Pressure Plasma Chemical Vapor Deposition, *Nanomaterials-Basel* **9**, 255 (2019)
- P. I. John, *Plasma Sciences and the Creation of Wealth* (Tata McGraw Hill Education, New York City, 2005)
- K. S. Joshy, S. Snigdha, and S. Thomas, *Plasma Modified Polymeric Materials for Scaffolding of Bone Tissue Engineering* (Elsevier, Amsterdam, 2019)
- K. S. Kim, and T. H. Kim, Nanofabrication by thermal plasma jets: From nanoparticles to low-dimensional nanomaterials, *J. Appl. Phys.* **125**, 070901 (2019)
- D. B. Kolesky, R. L. Truby, A. S. Gladman, T. A. Busbee, K. A. Homan, and J. A. Lewis, 3D Bioprinting of Vascularized, Heterogeneous Cell-Laden Tissue Constructs, *Advanced Materials* **26**, 3124-3130 (2014)
- H. W. P. Koops, O. E. Hoinkis, M. E. W. Honsberg, R. Schmidt, R. Blum, G. Bottger, A. Kuligk, C. Liguda, and M. Eich, Two-dimensional photonic crystals produced by additive nanolithography with electron beam-induced deposition act as filters in the infrared, *Microelectron. Eng.* **57-8**, 995-1001 (2001)
- U. Kortshagen, and U. Bhandarkar, Modeling of particulate coagulation in low pressure plasmas, *Phys Rev E* **60**, 887-898 (1999)
- U. R. Kortshagen, R. M. Sankaran, R. N. Pereira, S. L. Girshick, J. J. Wu, and E. S. Aydil, Nonthermal Plasma Synthesis of Nanocrystals: Fundamental Principles, Materials, and Applications, *Chem. Rev.* **116**, 11061-11127 (2016)
- F. Kotz, K. Arnold, W. Bauer, D. Schild, N. Keller, K. Sachsenheimer, T. M. Nargang, C. Richter, D. Helmer, and B. E. Rapp, Three-dimensional printing of transparent fused silica glass, *Nature* **544**, 337 (2017)
- N. J. Kramer, R. J. Anthony, M. Mamunuru, E. S. Aydil, and U. R. Kortshagen, Plasma-induced crystallization of silicon nanoparticles, *Journal of Physics D-Applied Physics* **47**, 075202 (2014)
- N. J. Kramer, E. S. Aydil, and U. R. Kortshagen, Requirements for plasma synthesis of nanocrystals at atmospheric pressures, *Journal of Physics D-Applied Physics* **48**, 035205 (2015)
- A. Kumar, S. Kang, C. Larriba-Andaluz, H. Ouyang, C. J. Hogan, and R. M. Sankaran, Ligand-free Ni nanocluster formation at atmospheric pressure via rapid quenching in a microplasma process, *Nanotechnology* **25**, (2014)
- S. Kyung, Y. Lee, C. Kim, J. Lee, and G. Yeom, Deposition of carbon nanotubes by capillary-type atmospheric pressure PECVD, *Thin Solid Films* **506**, 268-273 (2006)
- A. Lazea-Stoyanova, A. Vlad, A. M. Vlaicu, V. S. Teodorescu, and G. Dinescu, Synthesis of Copper Particles by Non-thermal Atmospheric Pressure Plasma Jet, *Plasma Process Polym* **12**, 705-709 (2015)
- H. H. Lee, K. S. Chou, and K. C. Huang, Inkjet printing of nanosized silver colloids, *Nanotechnology* **16**, 2436-2441 (2005)
- S. W. Lee, D. Liang, X. P. A. Gao, and R. M. Sankaran, Direct Writing of Metal Nanoparticles by Localized Plasma Electrochemical Reduction of Metal Cations in Polymer Films, *Adv. Funct. Mater.* **21**, 2155-2161 (2011)
- B. B. Lewis, M. G. Stanford, J. D. Fowlkes, K. Lester, H. Plank, and P. D. Rack, Electron-stimulated purification of platinum nanostructures grown via focused electron beam induced deposition, *Beilstein J Nanotech* **6**, 907-918 (2015)
- J. A. Lewis, Direct ink writing of 3D functional materials, *Adv. Funct. Mater.* **16**, 2193-2204 (2006)

- M. M. Ling, and Z. N. Bao, Thin film deposition, patterning, and printing in organic thin film transistors, *Chem. Mater.* **16**, 4824-4840 (2004)
- Y. Liu, J. K. Boyles, J. Genzer, and M. D. Dickey, Self-folding of polymer sheets using local light absorption, *Soft Matter* **8**, 1764-1769 (2012)
- M. F. Mabrook, C. Pearson, A. S. Jombert, D. A. Zeze, and M. C. Petty, The morphology, electrical conductivity and vapour sensing ability of inkjet-printed thin films of single-wall carbon nanotubes, *Carbon* **47**, 752-757 (2009)
- K. Mackie, and M. Gordon, Microplasma-based deposition of functional nanomaterials for energy storage applications, *Abstracts of Papers of The American Chemical Society* **253**, Apr 2, (2017)
- K. E. Mackie, A. C. Pebley, M. M. Butala, J. P. Zhang, G. D. Stucky, and M. J. Gordon, Microplasmas for direct, substrate-independent deposition of nanostructured metal oxides, *Appl. Phys. Lett.* **109**, 033110 (2016)
- S. Magdassi, A. Bassa, Y. Vinetsky, and A. Kamyshny, Silver nanoparticles as pigments for water-based ink-jet inks, *Chem. Mater.* **15**, 2208-2217 (2003)
- P. Maguire, D. Rutherford, M. Macias-Montero, C. Mahony, C. Kelsey, M. Tweedie, F. Perez-Martin, H. McQuaid, D. Diver, and D. Mariottit, Continuous In-Flight Synthesis for On-Demand Delivery of Ligand-Free Colloidal Gold Nanoparticles, *Nano Lett.* **17**, 1336-1343 (2017)
- R. M. Mahamood, *Laser Metal Deposition Process of Metals, Alloys, and Composite Materials* (Springer, International, 2018)
- A. Mameli, Y. H. Kuang, M. Aghaee, C. K. Ande, B. Karasulu, M. Creatore, A. J. M. Mackus, W. M. M. Kessels, and F. Roozeboom, Area-Selective Atomic Layer Deposition of $\text{In}_2\text{O}_3\text{:H}$ Using a μ -Plasma Printer for Local Area Activation, *Chem. Mater.* **29**, 921-925 (2017)
- L. Mangolini, and U. Kortshagen, Plasma-assisted synthesis of silicon nanocrystal inks, *Advanced Materials* **19**, 2513-2519 (2007)
- L. Mangolini, E. Thimsen, and U. Kortshagen, High-yield plasma synthesis of luminescent silicon nanocrystals, *Nano Lett.* **5**, 655-659 (2005)
- Y. Q. Mao, K. Yu, M. S. Isakov, J. T. Wu, M. L. Dunn, and H. J. Qi, Sequential Self-Folding Structures by 3D Printed Digital Shape Memory Polymers, *Sci. Rep.* **5**, 13616 (2015)
- T. Matsoukas, and M. Russell, Particle Charging in Low-Pressure Plasmas, *J. Appl. Phys.* **77**, 4285-4292 (1995)
- R. Maurau, N. D. Boscher, S. Olivier, S. Bulou, T. Belmonte, J. Dutroncy, T. Sindzingre, and P. Choquet, Atmospheric pressure, low temperature deposition of photocatalytic TiO_x thin films with a blown arc discharge, *Surf Coat Tech* **232**, 159-165 (2013)
- I. Michelakaki, N. Boukos, D. A. Dragatogiannis, S. Stathopoulos, C. A. Charitidis, and D. Tsoukalas, Synthesis of hafnium nanoparticles and hafnium nanoparticle films by gas condensation and energetic deposition, *Beilstein J Nanotech* **9**, 1868-1880 (2018)
- S. Y. Min, T. S. Kim, B. J. Kim, H. Cho, Y. Y. Noh, H. Yang, J. H. Cho, and T. W. Lee, Large-scale organic nanowire lithography and electronics, *Nat Commun* **4**, 1773 (2013)
- S. Mohr, and O. Khan, 3D Printing and Its Disruptive Impacts on Supply Chains of the Future, *Technol Innov Manag* 20-25 (2015)
- I. Motrescu, and M. Nagatsu, Nanocapillary Atmospheric Pressure Plasma Jet: A Tool for Ultrafine Maskless Surface Modification at Atmospheric Pressure, *Acs Appl Mater Inter* **8**, 12528-12533 (2016)
- K. Murakami, and M. Takai, Nano electron source fabricated by beam-induced deposition and its unique feature, *Microelectron. Eng.* **132**, 74-82 (2015)
- L. E. Murr, S. M. Gaytan, D. A. Ramirez, E. Martinez, J. Hernandez, K. N. Amato, P. W. Shindo, F. R. Medina, and R. B. Wicker, Metal Fabrication by Additive Manufacturing Using Laser and Electron Beam Melting Technologies, *Journal of Materials Science & Technology* **28**, 1-14 (2012)
- T. J. Ober, D. Foresti, and J. A. Lewis, Active mixing of complex fluids at the microscale, *Proc. Natl. Acad. Sci. U.S.A.* **112**, 12293-12298 (2015)
- S. Ohno, and M. Uda, Preparation for Ultrafine Particles of Fe-Ni, Fe-Cu and Fe-Si Alloys by Hydrogen Plasma-Metal Reaction, *J. Jpn. Inst. Met.* **53**, 946-952 (1989)
- R. Parashkov, E. Becker, T. Riedl, H. H. Johannes, and W. Kowalsky, Large area electronics using printing, methods, *Proceedings of the Ieee* **93**, 1321-1329 (2005)
- J.-U. Park, M. Hardy, S. J. Kang, K. Barton, K. Adair, D. k. Mukhopadhyay, C. Y. Lee, M. S. Strano, A. G. Alleyne, J. G. Georgiadis, *et al.*, High-resolution electrohydrodynamic jet printing, *Nat Mater* **6**, 782 (2007)
- S. Park, U. Cvelbar, W. Choe, and S. Y. Moon, The creation of electric wind due to the electrohydrodynamic force, *Nat Commun* **9**, 371 (2018)
- J. Perelaer, R. Jani, M. Grouchko, A. Kamyshny, S. Magdassi, and U. S. Schubert, Plasma and Microwave Flash Sintering of a Tailored Silver Nanoparticle Ink, Yielding 60% Bulk Conductivity on Cost- Effective Polymer Foils, *Advanced Materials* **24**, 3993-3998 (2012)

- H. Plank, C. Gspan, M. Dienstleder, G. Kothleitner, and F. Hofer, The influence of beam defocus on volume growth rates for electron beam induced platinum deposition, *Nanotechnology* **19**, 485302 (2008)
- Relyon Plasma, <https://www.relyon-plasma.com/maximum-surface-quality-in-3d-printing/?lang=en>.
- P. D. Rack, J. D. Fowlkes, and S. J. Randolph, In situ probing of the growth and morphology in electron-beam-induced deposited nanostructures, *Nanotechnology* **18**, 465602 (2007)
- C. L. Randall, E. Gultepe, and D. H. Gracias, Self-folding devices and materials for biomedical applications, *Trends Biotechnol.* **30**, 138-146 (2012)
- C. Richmonds, and R. M. Sankaran, Plasma-liquid electrochemistry: Rapid synthesis of colloidal metal nanoparticles by microplasma reduction of aqueous cations, *Appl. Phys. Lett.* **93**, 131501 (2008)
- C. Richmonds, M. Witzke, B. Bartling, S. W. Lee, J. Wainright, C. C. Liu, and R. M. Sankaran, Electron-Transfer Reactions at the Plasma-Liquid Interface, *J. Am. Chem. Soc.* **133**, 17582-17585 (2011)
- P. Richner, S. J. P. Kress, D. J. Norris, and D. Poulikakos, Charge effects and nanoparticle pattern formation in electrohydrodynamic NanoDrip printing of colloids, *Nanoscale* **8**, 6028-6034 (2016)
- S. Sanaur, A. Whalley, B. Alameddine, M. Carnes, and C. Nuckolls, Jet-printed electrodes and semiconducting oligomers for elaboration of organic thin-film transistors, *Org. Electron.* **7**, 423-427 (2006)
- V. Satulu, M. D. Ionita, S. Vizireanu, B. Mitu, and G. Dinescu, Plasma Processing with Fluorine Chemistry for Modification of Surfaces Wettability, *Molecules* **21**, 1711 (2016)
- J. Schneider, P. Rohner, D. Thureja, M. Schmid, P. Galliker, and D. Poulikakos, Electrohydrodynamic NanoDrip Printing of High Aspect Ratio Metal Grid Transparent Electrodes, *Adv. Funct. Mater.* **26**, 833-840 (2016)
- M. Schwentenwein, and J. Homa, Additive Manufacturing of Dense Alumina Ceramics, *Int J Appl Ceram Tec* **12**, 1-7 (2015)
- C. W. Sele, T. von Werne, R. H. Friend, and H. Sirringhaus, Lithography-free, self-aligned inkjet printing with sub-hundred-nanometer resolution, *Advanced Materials* **17**, 997-1001 (2005)
- J. H. Seo, and B. G. Hong, Thermal Plasma Synthesis of Nano-Sized Powders, *Nucl Eng Technol* **44**, 9-20 (2012)
- S. K. Seol, D. Kim, S. Lee, J. H. Kim, W. S. Chang, and J. T. Kim, Electrodeposition-based 3D Printing of Metallic Microarchitectures with Controlled Internal Structures, *Small* **11**, 3896-3902 (2015)
- M. Shigeta, and A. B. Murphy, Thermal plasmas for nanofabrication, *Journal of Physics D-Applied Physics* **44**, 174025 (2011)
- Y. Shimizu, Diameter control of gold nanoparticles synthesized in gas phase using atmospheric-pressure H₂/Ar plasma jet and gold wire as the nanoparticle source: Control by varying the H₂/Ar mixture ratio, *AIP Adv* **7**, 015316 (2017)
- Y. Shimizu, K. Kawaguchi, T. Sasaki, and N. Koshizaki, Generation of room-temperature atmospheric H₂/Ar microplasma jet driven with pulse-modulated ultrahigh frequency and its application to gold nanoparticle preparation, *Appl. Phys. Lett.* **94**, 191504 (2009)
- Y. Shimizu, T. Sasaki, T. Ito, K. Terashima, and N. Koshizaki, Fabrication of spherical carbon via UHF inductively coupled microplasma CVD, *Journal of Physics D-Applied Physics* **36**, 2940-2944 (2003)
- K. Silmy, A. Hollander, A. Dillmann, and J. Thomel, Micro-jet plasma CVD with HMDSO/O₂, *Surf Coat Tech* **200**, 368-371 (2005)
- H. Sirringhaus, T. Kawase, R. H. Friend, T. Shimoda, M. Inbasekaran, W. Wu, and E. P. Woo, High-resolution inkjet printing of all-polymer transistor circuits, *Science* **290**, 2123-2126 (2000)
- M. A. Skylar-Scott, S. Gunasekaran, and J. A. Lewis, Laser-assisted direct ink writing of planar and 3D metal architectures, *Proc. Natl. Acad. Sci. U.S.A.* **113**, 6137-6142 (2016)
- M. G. Stanford, B. B. Lewis, J. H. Noh, J. D. Fowlkes, and P. D. Rack, Inert Gas Enhanced Laser-Assisted Purification of Platinum Electron-Beam-Induced Deposits, *Acs Appl Mater Inter* **7**, 19579-19588 (2015)
- A. R. Studart, Additive manufacturing of biologically-inspired materials, *Chem. Soc. Rev.* **45**, 359-376 (2016)
- N. Stutzmann, R. H. Friend, and H. Sirringhaus, Self-aligned, vertical-channel, polymer field-effect transistors, *Science* **299**, 1881-1884 (2003)
- Y. Sui, Y. Dai, C. C. Liu, R. M. Sankaran, and C. A. ZormanSui, A New Class of Low-Temperature Plasma-Activated, Inorganic Salt-Based Particle-Free Inks for Inkjet Printing Metals, *Advanced Materials Technology* 1900119 (2019)
- J. B. Szczech, C. M. Megaridis, D. R. Gamota, and J. Zhang, Fine-line conductor manufacturing using drop-on-demand PZT printing technology, *Ieee T Electron Pack* **25**, 26-33 (2002)
- T. Takai, H. Nakao, and F. Iwata, Three-dimensional microfabrication using local electrophoresis deposition and a laser trapping technique, *Opt. Express* **22**, 28109-28117 (2014)
- V. Tasco, M. Esposito, F. Todisco, A. Benedetti, M. Cuscuna, D. Sanvitto, and A. Passaseo, Three-dimensional nanohelices for chiral photonics, *Appl Phys A* **122**, 280 (2016)
- EMPA Materials Science and Technology, <http://www.empa.ch>
- Tekna, <http://www.tekna.com>

- M. Thomas, J. Borris, A. Dohse, M. Eichler, A. Hinze, K. Lachmann, K. Nagel, and C. P. Klages, Plasma Printing and Related Techniques - Patterning of Surfaces Using Microplasmas at Atmospheric Pressure, *Plasma Process Polym* **9**, 1086-1103 (2012)
- M. Thomson, J. L. Hodgkinson, and D. W. Sheel, Control of zinc oxide surface structure using combined atmospheric pressure-based CVD growth and plasma etching, *Surf Coat Tech* **230**, 190-195 (2013)
- S. Tibbits, 4d Printing: Multi-Material Shape Change, *Archit Design* **84**, 116-121 (2014)
- R. L. Truby, and J. A. Lewis, Printing soft matter in three dimensions, *Nature* **540**, 371-378 (2016)
- F. Ullmann, and J. Bielecki, Synthesis in the Biphenyl series. (I. Announcement), *Ber Dtsch Chem Ges* **34**, 2174-2185 (1901)
- M. Vaezi, H. Seitz, and S. F. Yang, A review on 3D micro-additive manufacturing technologies, *Int J Adv Manuf Tech* **67**, 1721-1754 (2013)
- R. van Hout, V. Rinsky, and Y. G. Grobman, Experimental study of a round jet impinging on a flat surface: Flow field and vortex characteristics in the wall jet, *Int. J. Heat Fluid Flow* **70**, 41-58 (2018)
- P. Verhoeven, A. Stevens, J. P. Schalken, M. Soltani, and A. Mäntysalo, Digital Printing with Micro Plasmas and its Effects on Surface Wettability, 28th International-Conference on Surface Modification Technologies, Tampere, 421-431 (2014)
- R. O. F. Verkuijlen, M. H. A. van Dongen, A. A. E. Stevens, J. van Geldrop, and J. P. C. Bernards, Surface modification of polycarbonate and polyethylene naphthalate foils by UV-ozone treatment and mu Plasma printing, *Appl. Surf. Sci.* **290**, 381-387 (2014)
- A. Vyatskikh, S. Delalande, A. Kudo, X. Zhang, C. M. Portela, and J. R. Greer, Additive manufacturing of 3D nano-architected metals, *Nat Commun* **9**, 593 (2018)
- D. Z. Wang, W. Zha, L. Feng, Q. Ma, X. M. Liu, N. Yang, Z. Xu, X. J. Zhao, J. S. Liang, T. Q. Ren, *et al.*, Electrohydrodynamic jet printing and a preliminary electrochemistry test of graphene micro-scale electrodes, *J Micromech Microeng* **26**, 045010 (2016)
- J. Z. Wang, J. Gu, F. Zenhausem, and H. Sirringhaus, Low-cost fabrication of submicron all polymer field effect transistors, *Appl. Phys. Lett.* **88**, (2006)
- M. Wang, P. Favi, X. Q. Cheng, N. H. Golshan, K. S. Ziemer, M. Keidar, and T. J. Webster, Cold atmospheric plasma (CAP) surface nanomodified 3D printed polylactic acid (PLA) scaffolds for bone regeneration, *Acta Biomater.* **46**, 256-265 (2016)
- X. L. Wang, A. Gidwani, S. L. Girshick, and P. H. McMurry, Aerodynamic focusing of nanoparticles: II. Numerical simulation of particle motion through aerodynamic lenses, *Aerosol Sci. Technol.* **39**, 624-636 (2005a)
- X. L. Wang, F. E. Kruis, and P. H. McMurry, Aerodynamic focusing of nanoparticles: I. Guidelines for designing aerodynamic lenses for nanoparticles, *Aerosol Sci. Technol.* **39**, 611-623 (2005)
- T. Wei, J. Ruan, Z. J. Fan, G. H. Luo, and F. Wei, Preparation of a carbon nanotube film by ink-jet printing, *Carbon* **45**, 2712-2716 (2007)
- K. D. Weltmann, J. F. Kolb, M. Holub, D. Uhrlandt, M. Simek, K. Ostrikov, S. Hamaguchi, U. Cvelbar, M. Cernak, B. Locke, *et al.*, The future for plasma science and technology, *Plasma Process Polym* **16**, 133502 (2019)
- J. Wienand, A. Riedelsheimer, and B. Weigand, Numerical study of a turbulent impinging jet for different jet-to-plate distances using two-equation turbulence models, *Eur J Mech B-Fluid* **61**, 210-217 (2017)
- J. Xu, C. Zhong, and C. Fu, Novel method for printing high-quality metal wires, *SPIE News* (2017) <http://doi.org/10.1117/2.1200712.0969>
- Y. G. Yao, Z. N. Huang, P. F. Xie, S. D. Lacey, R. J. Jacob, H. Xie, F. J. Chen, A. M. Nie, T. C. Pu, M. Rehwoldt, *et al.*, Carbothermal shock synthesis of high-entropy-alloy nanoparticles, *Science* **359**, 1489-1494 (2018)
- S. Yick, Z. J. Han, and K. Ostrikov, Atmospheric microplasma-functionalized 3D microfluidic strips within dense carbon nanotube arrays confine Au nanodots for SERS sensing, *Chem. Commun.* **49**, 2861-2863 (2013)
- Z. Yin, Y. Huang, Y. Duan, and H. Zhang, *Electrohydrodynamic direct-writing for flexible electronic manufacturing* (Springer, Singapore, 2018)
- R. M. Young, and E. Pfender, Generation and Behavior of Fine Particles in Thermal Plasmas - a Review, *Plasma Chem. Plasma Process.* **5**, 1-37 (1985)
- I. Zein, D. W. Hutmacher, K. C. Tan, and S. H. Teoh, Fused deposition modeling of novel scaffold architectures for tissue engineering applications, *Biomaterials* **23**, 1169-1185 (2002)
- R. Zimmermann, A. Pfuch, K. Horn, J. Weisser, A. Heft, M. Roder, R. Linke, M. Schnabelrauch, and A. Schimanski, An Approach to Create Silver Containing Antibacterial Coatings by Use of Atmospheric Pressure Plasma Chemical Vapour Deposition (APCVD) and Combustion Chemical Vapour Deposition (CCVD) in an Economic Way, *Plasma Process Polym* **8**, 295-304 (2011)