Carbon-based materials for stable, cheaper and large-scale processable perovskite solar cells

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Almost ten years after their first use in the photovoltaic (PV) field, perovskite solar cells (PSCs) are now hybrid devices that, in addition to having reached silicon performance, can accelerate the energy transition and boost the use of abundant elements for their manufacturing process. However, noble metals (in particular gold) represent the most typically used sources for back electrode fabrication, and this issue has been intensively considered by the research community in the last five years. This review shows how the most promising solution, considering also the need to develop a large-scale production process, is based on the use of carbon-based materials for the preparation of back electrodes. Graphite, carbon black, graphene and carbon nanotubes (CNTs) have been proposed, functionalized and characterized, leading to laboratory-scale solar cells and modules capable of providing excellent efficiencies and ensuring stability greater than those of gold-based devices. Strengthened by these results and its hydrophobizing properties, carbon has also started to be used as an electron transporting material (ETM), with excellent results on both rigid and flexible substrates. This review discusses the major advances and the updated state-of-the-art in the carbon-based PSC scenario, keeping a solid trajectory where the accessibility, low cost, high electrical conductivity, chemical stability and controllable porosity of carbon are highlighted and exploited in the design of upscalable hybrid solar cells.

1 Introduction

Among all PV fields, PSCs have shown the highest increase in terms of power conversion efficiency (PCE), passing from 3.8% to 25.2% in just a few years. Perovskites are materials with the generic formula ABX₃, where A is an organic larger cation, situated in the eight corners of the unit cell, with cubo-octahedral coordination, B is a metal cation located at the body center, and X represents an anion that binds them both. This kind of material has attracted much attention, due to its superb characteristics, such as strong and broad optical absorption from the visible to the near-infrared (NIR) range, long carrier lifetime, long diffusion length (from 100 nm to 1 µm), a direct optical band-gap of 1.5 eV and low exciton binding energy. In addition, these semiconductors offer good versatility and processability from solution, being used as light absorber and charge transporting materials. One of the most commonly used perovskites is the hybrid organic–inorganic compound, CH₃NH₃PbI₃. One of the main drawbacks toward the commercialization of PSCs is the use of noble metals, such as gold and silver, as back electrodes, thus limiting their large-scale application. Gold and
silver are expensive rare metals and their deposition requires a highly energy-consuming vacuum evaporation method.\textsuperscript{22} In addition, silver and gold electrodes may deteriorate cell performance,\textsuperscript{23} due to migration of halogen atoms from the perovskite phase, with formation of silver and gold halides.\textsuperscript{24}–\textsuperscript{26}

The most common carbon-based materials used in the PSC field are graphite/amorphous carbon, graphene, and CNTs. They represent a suitable solution to substitute noble metals, due to their low cost, high conductivity, eventual low-temperature processing (100 °C)\textsuperscript{1,27,28} and work function close to that of gold (5.0 and 5.1 eV, respectively).\textsuperscript{29,30} Another advantage of carbon-based components reflects on perovskite degradation in the presence of water, that leads to the formation of the hydrated phases \(\text{CH}_3\text{NH}_3\text{PbI}_2\cdot\text{H}_2\text{O}\) and \((\text{CH}_3\text{NH}_3)_4\text{PbI}_6\cdot2\text{H}_2\text{O}\);\textsuperscript{31} while the formation of the monohydrate phase is reversible, the dihydrate phase degrades irreversibly to yellow \(\text{PbI}_2\) and \(\text{CH}_3\text{NH}_3\text{I}\).\textsuperscript{32} Conversely, carbon-based materials can improve PSC stability due to their highly hydrophobic nature that leads to the prevention of moisture penetration in the perovskite layer, even without carrying out the sealing process.\textsuperscript{24}

Nonetheless, carbon-based materials also present many drawbacks, associated especially with their processability on a large scale. For example, graphene, the 2D layer of graphite, is a very versatile material with metal-like properties; however, it is difficult to be obtained by mechanical or chemical exfoliation on a large scale. The dispersion of graphene layers from graphite requires a large amount of organic solvents, such as dimethylformamide; on the other hand, the epitaxial growth on SiC by vacuum graphitization has the disadvantages of high temperature (1300 °C) and high costs of the monocristalline SiC sheet.\textsuperscript{33,34}

Chemical vapour deposition (CVD) is more applicable on a large scale, but has hardly any control on film thickness and requires expensive substrates (Ni, Cu, Pd, Pt).\textsuperscript{35} CNTs are composed of graphitic layers rolled up to form a cylinder. For this reason, in each tube, an elevated charge-transfer resistance is created.\textsuperscript{34,36} The problem is partially solved by the use of multi-walled CNTs (MWCNTs), which consist of multiple CNTs stacked together.

In some cases reported in this review, graphite and amorphous carbon require a high processing temperature. They form a rigid structure that impedes the penetration of the perovskite precursor solution. For this reason, the use of carbon black (CB) is necessary to break the oriented and impermeable graphite layer. In addition, in most cases, the small CB nanoparticles are obtained by the incomplete combustion of fossil fuels.

This review focuses on the current trends and major achievements obtained from hundreds of research teams working on carbon-based PSCs.\textsuperscript{37} From our analysis of literature databases, it clearly emerges that graphite and CB represent the most credible realities if the target is – in a short time – a printable and industrially scalable solar energy conversion technology. We will show how the strategies for functionalizing and engineering the interfaces between carbon and perovskites or carbon and hole transporting materials (HTMs) strongly influence the cell PCE and stability. Finally, we will offer an overview of the state of the art of alternative carbon-based materials, such as CNTs and graphene, that show interesting aspects when low temperature processes or flexible device fabrications are envisaged. The achievement of stable, cheap and reproducible PSCs will be fundamental in view of their worldwide diffusion in the energy scenario, as well as integration with energy storage technologies.\textsuperscript{38}–\textsuperscript{47}

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2 Graphite- and carbon black-based back electrodes

Generally speaking, a PSC is based on a perovskite layer sandwiched between an ETM and a HTM. The ETM usually consists of TiO₂ or another n-type semiconductor anode, while the back electrode is deposited on top of the HTM. For efficient charge extraction, the valence band (VB) and the conduction band of the perovskite should lie below the highest occupied molecular orbital (HOMO) of the HTM and under the lowest unoccupied molecular orbital (LUMO) of the ETM, respectively. After light absorption, in the perovskite layer an electron–hole pair is generated. Electrons are collected by the ETM and migrate towards the anode side, while holes are collected by the HTM and migrate towards the back electrode. The most common organic HTMs are expensive and require an inert atmosphere for the deposition process. They are also subjected to halide and metal ion migration, resulting in the degradation of the cell. In addition, carbon-based materials are processable, highly conductive, thermally inert and hydrophobic. It needs to be doped with lithium bis[trifluoromethanesulfonyl]imide (LiTFSI), a hygroscopic and deliquescent salt that allows the penetration of water molecules, leading to the decomposition of the perovskite. Luckily, perovskite materials show a large carrier path and PSCs can also work without the HTM. In this architecture, an insulating layer, such as Al₂O₃ or ZrO₂, is necessary to keep the front electrode separated from the back one.

To date, the best performances have been obtained with gold as the back electrode. However, the noble element may be replaced by carbon-based materials, such as graphite, graphene, CNTs and so on. In fact, carbon has a suitable work function of 5.0 eV, very close to that of gold (5.1 eV). In addition, carbon-based materials are processable, highly conductive, thermally inert and hydrophobic. Carbon-based back electrodes represent a concrete and indispensable strategy towards the large-scale industrial production of PSCs. Fig. 1 shows that it took 4 years, from the first reported perovskite-based PVs (2009), to see the first articles published on this topic. However, it must also be specified that all articles on PSCs published before 2012 were focused on devices using liquid electrolytes. Carbon-based devices appeared approximately only one year after the first solid-state PSC based on spiro-OMeTAD. As well known, the unexpected efficiency improvement that spiro-OMeTAD promoted is sometimes considered the true starting point of the PSC topic. To date, publications on carbon-based PSCs count for about 10% of the overall PSC articles, maintaining an upward trend after 2015.

Among the most known compounds, graphite is a soft material, composed of a hexagonal lattice layer, with high electrical conductivity and excellent thermal stability. On the other hand, CB is an amorphous nanopowder obtained by incomplete combustion of carbonaceous materials. Even though it shows lower conductivity than graphite, carbon black is used to distort and break the preferential orientation of graphite flakes. Indeed, a full graphite layer would not ease the infiltration of the perovskite solution. Furthermore, it must also be stressed that the ratio between carbon black and graphite is relevant not only in terms of electrical conductivity; conversely, it also has an influence on the processability of the carbon paste, in terms of viscosity, adhesion to the other cell layers and infiltration degree of the perovskite.

Carbon-based materials may be processed by two main deposition techniques. The first process requires temperatures in the range 400–500 °C. A mesoporous carbon layer is deposited by doctor-blade or screen-printing techniques on the top of an insulating layer, and subsequently sintered. The insulator has the role of preventing the contact between the front and the back electrode, thus avoiding photocurrent leakage. To the best of our knowledge, three oxides have been successfully used as insulating layers in the PSC field: ZrO₂, Al₂O₃ and TiO₂. The latter can be used both as a compact layer and as a mesoscopic layer, in combination with ZrO₂ and Al₂O₃. Generally speaking, ZrO₂ is more efficient as the insulating layer with respect to Al₂O₃. In fact, its larger pores facilitate the infiltration of the perovskite phase and its contact with the anodic material. The effect of the thickness of the insulating layer has been investigated by Liu et al. and Barichello et al. Optimal thicknesses of 1 μm and 1.8 μm for ZrO₂ and Al₂O₃, respectively, were found. Below these values, most of the generated photoelectrodes recombined with holes, leading to a high energy loss. On the other hand, the thickness of the insulating layer should not be higher than the limit of carrier diffusion length. More details on this topic can be found in a recently published review article.

The perovskite phase is formed later, since it is not stable over 120 °C, by drop-casting a precursor solution on top of the carbon layer. The perovskite precursor solution must penetrate the entire thickness of the carbon layer, to ensure a contact with the anode. Mesoscopic cells, where a good interface between carbon and the perovskite is established, are obtained by this strategy. However, the penetration is somehow difficult, may require the use of additives and is not applicable to flexible
substrates. The other strategy is a layer-by-layer deposition,\textsuperscript{58} where a perovskite phase is formed by a one- or two-step method and carbon is doctor-bladed or screen-printed on top of CH\textsubscript{3}NH\textsubscript{3}PbI\textsubscript{3} or HTM. The carbon layer may also be deposited on another substrate and then transferred onto the cell. These depositions are simpler, do not require high temperature and may be suitable also for flexible devices, but the interface usually suffers from poor contact.\textsuperscript{84–86} Planar architectures with and without HTMs are obtained.

The most significant and state-of-the-art outcomes on these two families of processing strategies of graphite- and carbon black-based back electrodes are reviewed in the forthcoming sub-sections.

### 2.1 High-temperature processed front electrodes

In 2013, Ku et al. reported the first use of carbon black/graphite as a back electrode in a PSC.\textsuperscript{29} Exploiting the ambipolar charge transfer properties of perovskites, a HTM was not used. Instead, an insulating ZrO\textsubscript{2} layer was put on top of the TiO\textsubscript{2} layer to prevent contact between the front and back electrodes. To further improve the device performances, spheroidal graphite with better conductivity and favorable morphology for pore filling was used in the carbon composite electrode. As evidenced by cross-sectional scanning electron microscopy (SEM) images, shown in Fig. 2, a more effective pore filling was ensured by the smooth structure of spheroidal graphite. In fact, the PCE was improved from 4.08% to 6.64%. The stability dropped only by 2% from the initial value after 70 days, confirming the positive role of the hydrophobic carbon layer in preventing the perovskite phase from the penetration of water molecules.

The influence of both carbon layer thickness and graphite size on the cell PCE was investigated by Zhang et al. in 2015.\textsuperscript{87} In fact, if the carbon layer is too thick, the penetration of the perovskite precursor solution is hindered. As a consequence, a poor contact between carbon and the perovskite phase is established. However, too thin carbon layers do not provide sufficient conductivity. The carbon layer thickness ranged from 5 to 15 \( \mu \)m and it was found that the optimal thickness was 9 \( \mu \)m, with a short-circuit current density \( J_{sc} \) of 18.06 mA cm\textsuperscript{-2} and a remarkable PCE of 11.63%. On the other hand, graphite flakes with different sizes were added to the carbon paste, to further enhance the conductivity of the carbon back electrode. Dimensions of graphite flakes ranged from 0.5 to 8 \( \mu \)m; it was found that the 8 \( \mu \)m-large sample ensured the best perovskite infiltration, having the biggest pore size. In addition, it had the lower square resistance.

Different results were obtained by Raminafshar et al. in 2018.\textsuperscript{88} They investigated the effect of TiO\textsubscript{2}, ZrO\textsubscript{2} and carbon thickness in HTM-free, high-temperature processed PSCs. Concerning the back electrode, the carbon layer should be thin enough to ensure perovskite infiltration; however, too thin layers are mechanically fragile and can crack, with a drop in the conductivity. Four thickness values were tested: 6.5, 15, 25 and 54 \( \mu \)m. On the other hand, the thicknesses of TiO\textsubscript{2} and ZrO\textsubscript{2} were kept constant at 0.4 and 1.7 \( \mu \)m, respectively. As reported in Table 1, the PCE improved with carbon thickness. Only when the layer became too thick (54 \( \mu \)m) the PCE dropped due to the poor penetration of the perovskite phase. The best carbon thickness was 25 \( \mu \)m, with an open-circuit voltage \( V_{oc} \) of 0.88 V, a \( J_{sc} \) of 21.4 mA cm\textsuperscript{-2}, a fill factor (FF) of 0.57 and a PCE of 10.7%.

It is matter of debate in the scientific community if the carbon layer behaves simply like an electrode ohmic contact or has HTM properties. Indeed, carbon shows a higher work function than other commonly used metals, such as silver and aluminium, and its Fermi level \((-5.0\text{ eV})\) is close to the VB maximum of CH\textsubscript{3}NH\textsubscript{3}PbI\textsubscript{3} \((-5.4\text{ eV})\). This makes the hole extraction at the perovskite/carbon interface possible and paves the way to HTM-free PSCs. In the last six months, the origins of subgap tail states and \( V_{oc} \) were investigated by Du et al., and the main outcome was that the cell voltage was more affected by the perovskite crystallinity than by other factors.\textsuperscript{89} Gelmetti et al. studied the energy alignment and recombination in PSCs.

![Fig. 2](image-url)  
Cross-sectional SEM images of (A) a spheroidal graphite-based cell and (B) a flake/bulk graphite-based device. (C) Long-term stability of PSCs at room temperature in the dark. Adapted and reprinted with permission from ref. 29.

### Table 1: PV parameters of carbon-based PSCs fabricated with back electrodes bearing different thicknesses. Adapted and reprinted with permission from ref. 88

| Thickness (\( \mu \)m) | Series resistance (\( \Omega \)) | \( V_{oc} \) (V) | \( J_{sc} \) (mA cm\textsuperscript{-2}) | FF | PCE (%) |
|-----------------------|-------------------------------|----------------|---------------------------------|----|--------|
| 6.5                   | 169                           | 0.85           | 0.33                            | 4.6 |
| 15                    | 120                           | 0.88           | 12.7                            | 0.54 | 5.9    |
| 25                    | 56                            | 0.88           | 21.4                            | 0.57 | 10.7   |
| 54                    | 41                            | 0.67           | 14.3                            | 0.45 | 4.3    |
by choosing HTMs with close chemical and physical properties; however, once they were deposited on the perovskite layer, different performance characteristics were obtained with respect to those predicted. In particular, upon deposition on top of the perovskite, an important change in the energy level position was detected. From some recent reports and communications to relevant conferences, it has often been hypothesized that the HOMO and VB matching between the HTM and hybrid perovskite is almost irrelevant to achieve high $V_{oc}$ in PSCs. This is quite a disruptive topic, especially when considering that just a couple of years ago the lower $V_{oc}$ of carbon-based HTM-free PSCs was motivated by the absence of a standard HTM compound in the cell. Indeed, if $V_{oc}$ is determined by the difference between the electron quasi-Fermi level (e.g., ETM/TiO$_2$ interaction) and the hole quasi-Fermi level (e.g., perovskite/HTM interaction), after removing the HTM the latter level should be determined by the perovskite/carbon interaction, which will lift the level position due to the higher Fermi level of carbon than the HOMO of common HTMs. As a result, $V_{oc}$ of carbon-based HTM-free PSCs was supposed to be lower than that of HTM-based cells and lowering the Fermi level of carbon electrodes was considered as a milestone to increase the cell potential. Even if the scope of this review is not that of investigating the origins of $V_{oc}$ in PSCs, the reader is invited to consider this point when checking the solar cell parameters of carbon-based HTM-free devices.

Another important parameter to consider is the temperature of the annealing process. In fact, it can influence both morphologic and electrical properties of the carbon electrode. In 2019, Mishra et al. studied the effect of annealing temperature in carbon electrodes deposited by the screen-printing of commercial carbon pastes. It was found out that temperatures below 300 °C did not lead to working PSCs. In fact, high-temperature sintering ensured a better contact between particles, thus allowing a faster charge transfer and higher conductivity. The vaporization of organic solvents was facilitated by the use of high temperature. In this way, a highly mesoporous structure, with pore filling ability, was created. On the other hand, highly resistive electrodes, with poor perovskite infiltration, were formed at temperatures lower than 300 °C. The samples treated at 350 and 400 °C achieved a remarkable PCE of 8.4% and 12.4%, respectively. SEM images showed an increased uniformity and a defect-free morphology in high-temperature treated carbon films. Raman spectra underlined the high graphitization degree and the lack of defect regions.

Besides thermal treatment, another efficient solution to improve the charge extraction ability of the carbon back electrode was proposed by Tian et al. in 2018. They suggested that a fine regulation of the oxygen content in the carbon black, used to fabricate the back electrode, may be exploited for tuning the work function of carbon and increasing the surface area and the contact with the perovskite phase. Oxygen can be considered as a dopant, since it increases the work function of the material; it can be present as C–OH, COOH and C=O groups on the surface, enlarging the interlayer distance through electrostatic repulsion, thus increasing the surface area. Two samples, with a high (oxygen rich carbon – ORC) and low (oxygen deficient carbon – ODC) content of oxygen-containing groups, were prepared. Brunauer–Emmett–Teller (BET) analysis highlighted the increased surface area of ORC with respect to ODC (186.4 vs. 112.5 m$^2$ g$^{-1}$, respectively). Cross-sectional SEM of ODC showed particles with surface defects and pin-holes, whereas ORC was present in the form of particles entrapped in the perovskite phase, thus ensuring an effective contact with the active material. The improved properties of ORC reflected also on the PV results. In fact, $V_{oc}$ increased from 0.88 to 0.98 V, $J_{sc}$ from 22.98 to 23.20 mA cm$^{-2}$, FF from 67.2 to 69.1 and PCE from 13.59 to 15.70%.

A highly interconnected carbon can also be obtained starting from natural biomass, treated at high temperatures. In fact, natural organisms are rich in interconnected biomolecules with high molecular weight, and the dense and interconnected structure can be maintained even after the thermal treatment. For example, in 2018 Mali et al. reported, in an interesting work, a carbon material derived from an Aloe Vera plant (AV-C), used as a back electrode for PSCs. The Aloe Vera gel was extracted from the leaves and dried under sunlight, forming a black powder. Then, the product was washed with HCl and treated at very high temperature (1000 °C). The resulting material presented highly interconnected nanoparticles with dimensions of 40–45 nm, as confirmed by SEM and transmission electron microscopy (TEM) techniques, and a graphitic structure with sp$^2$ carbon, as demonstrated by X-raydiffraction (XRD) and Raman analysis (see Fig. 3). When used in PSCs with ZrO$_2$ as an insulator, AV-C provided a PCE of 12.58% and quite good stability: indeed, it retained 85% of the initial efficiency after 1000 h.

High-temperature processed carbon may suffer from poor contact with the perovskite phase in mesoscopic PSCs. In fact, bulk graphite has a large grain size, with small specific surface area. However, to ensure a good contact with the perovskite phase, a material with large surface area and interconnected channels is required. For this reason, malleable ultrathin graphite (UG) was used instead of bulk graphite (BG) for the first time by Duan et al. as a cathode material for PSCs. UG was prepared by mechanical exfoliation of BG and mixed with carbon black. The latter is a fine carbonaceous powder, with small particles, that breaks the extremely oriented and impermeable graphite layer, improving infiltration phenomena. As confirmed by SEM images shown in Fig. 4A–D, although particle dimensions of BG and UG were similar, UG had a more malleable morphology, better suited for the perovskite phase, and that ensured an effective pore filling of the precursor solution. In addition, the specific surface area of UG (202.24 m$^2$ g$^{-1}$) was remarkably higher than that of BG (69.01 m$^2$ g$^{-1}$). A UG-based cell resulted mostly in higher FF ($V_{oc}$ = 901 mV, $J_{sc}$ = 22.97 mA cm$^{-2}$, FF = 0.68, PCE = 14.07%) with respect to that of the BG-based one ($V_{oc}$ = 893 mV, $J_{sc}$ = 22.89 mA cm$^{-2}$, FF = 0.62, PCE = 12.63%), measured with reverse scan direction.

Another strategy for improving the contact of carbon with perovskites is the use of a volatile, pore forming agent in carbon pastes. When the carbon paste is deposited and thermally treated, this agent decomposes and leaves pores that facilitate the penetration of the perovskite precursor solution into the
mesoporous structure. In 2018, Tao et al. used different contents of polystyrene spheres (PSs) as pore forming agents in carbon pastes, in order to obtain a highly porous carbon back electrode for PSCs. Fig. 4E shows the pristine material, deposited by screen-printing. Fig. 4F shows the PSs–10% carbon paste before annealing, with PSs clearly visible on the surface. After the thermal treatment, PSs were removed, leaving a highly porous material, as shown in Fig. 4G. The weight ratio of PSs with respect to the carbon paste had an effect on the PCE of the resulting devices. In fact, the optimal percentage was 10%, with a $V_{oc}$ of 782 mV, a $J_{sc}$ of 9.32 mA cm$^{-2}$, a FF of 0.56 and a PCE of 4.10%; these values were improved with respect to those of the pristine material ($V_{oc} = 704$ mV, $J_{sc}$ of 8.36 mA cm$^{-2}$, FF = 0.55 and PCE = 3.36%). A further increase in the PSs content brought to a reduced PCE of 3.13%, mainly due to the decreased conductivity of the carbon electrode.

One of the main issues with carbon-based PSCs is the high resistance of the cell and the uncontrolled crystallization of the perovskite phase in the porous stack. An attempt to reduce the sheet resistance and ameliorate interfaces between printed stacks was made by Hashmi et al. in 2017. They reported a humidity-assisted thermal exposure (HTE) as a post-treatment method for PSCs. After the pore filling of the substrate with the perovskite precursor solution (one-step deposition method), the cell was exposed to 70% relative humidity (RH) for 115 h. This treatment resulted in a drastic enhancement of the preliminarily measured PV parameters. Also, the hysteresis effect was reduced. This beneficial effect may be ascribed to...
two main factors: first of all, the sheet resistance of the carbon layer, measured by the four-probe sheet resistance method, was found to drastically reduce after the HTE treatment. Water molecules probably increased the interconnectivity between carbon nanoparticles (NPs) and perovskite crystals. Secondly, the hydrophobic carbon impeded the fast intrusion of water drops, that would have led to the decomposition of the perovskite, and allowed only the penetration of gaseous water molecules. Perovskite growth under humidity was already demonstrated by You et al.99 Larger crystals, with an optimal contact with both anode and carbon layer, were obtained. One batch of devices was stored at 40 °C under full sun illumination and retained 92% of its initial efficiency after 350 h, confirming the irrereplaceable role of hydroscopic carbon in preventing moisture penetration.

A suitable strategy to further enhance PSC efficiency is the replacement of a methylammonium (MA) cation100 with 5-amino valeric acid (5-AVA)99 or formamidinium (HC(NH$_2$)$_2$) ensuring a better interfacial contact with the anode.102 Mei reported an increase in PCE from 7.2% to 11.6% by replacing MA with 5-AVA99 while Santhosh reported an average PCE of 6.47% for the same cell (see Table 2).102 The lower PCE was ascribed to the poor FF, due to a non-optimal assembly of the cell. On the other hand, the FA cation forms a perovskite phase with a band-gap of 1.47 eV, lower than that of MAPbI$_3$ (1.55 eV). FAPbI$_3$ presents great charge transport ability and mean free path.126 This was ascribed to the fact that high-temperature treated lead-based perovskites, both in mixed-cation perovskites117,118 and in fully inorganic CsPbI$_3$ perovskites,119,120 Despite a remarkable initial PCE of 10%,121 inorganic CsPbI$_3$ underwent an irreversible transformation toward a non-perovskite phase. Therefore, the stability was seriously compromised121,122. However, the thermal stability can be increased by fully replacing iodide with bromide.123,124 In fact, CsPbBr$_3$ is stable till 580 °C and presents great charge transport ability and mean free path.126 This perovskite was used in planar PSCs and showed a remarkable stability for more than three months in humid air.122 For the first time, in 2018 Poli et al. demonstrated that CsPbBr$_3$ was able to work as an active material also in mesoscopic PSCs.125 The mesoporous carbon layer was annealed at different temperatures, ranging from 100 to 400 °C. Samples annealed below 400 °C still presented traces of organic binder present in carbon pastes. Only for the sample annealed at 400 °C a pure perovskite phase was found. The PV parameter that was mostly influenced by temperature was the $V_{oc}$. In fact, the presence of other species can truly influence the recombination rate.127,128 The optimal annealing temperature was 400 °C, and the corresponding devices showed an average $V_{oc}$ of 1.27 V, a $J_{sc}$ of 6.29 mA cm$^{-2}$, a FF of 73.5 and a PCE of 4.77% (see Table 2). For comparison, an analogous planar architecture was also reported, which presented a lower PCE (3.67%), even if the $V_{oc}$ was higher (1.33 V) as shown in Table 2. This was ascribed to the fact that high-temperature treated mesoporous carbon presented high roughness, which allowed the permeation of the precursor solution and ensured a more effective contact with the perovskite phase, with respect to the planar carbon layer.

TiO$_2$ is the most used anodic material not only for PSCs, but also for dye-sensitized solar cells (DSSCs).139–141 It presents several advantages, such as low cost, low toxicity, adequate band-gap and high electron collection efficiency. Some efforts have been made to further ameliorate its performance in

### Table 2

| Perovskite          | $V_{oc}$ (mV) | $J_{sc}$ (mA cm$^{-2}$) | FF | PCE (%) | Ref. |
|---------------------|--------------|-------------------------|----|---------|------|
| MAPbI$_3$           | 835          | 13.9                    | 0.61 | 7.2     | 99   |
| (5-AVA)$_2$[MA]$_2$PbI$_4$ | 843          | 21.3                    | 0.65 | 11.6    | 99   |
| (5-AVA)$_2$[MA]$_2$PbI$_4$ | 838          | 22.8                    | 0.66 | 12.84   | 99   |
| (5-AVA)$_2$[MA]$_2$PbI$_4$ | 830          | 15.6                    | 0.49 | 6.47    | 102  |
| MAPbI$_3$           | 928          | 17.5                    | 0.70 | 11.4    | 100  |
| FAPbI$_3$           | 901          | 18.4                    | 0.72 | 11.9    | 100  |
| (FA)$_2$[MA]$_2$PbI$_4$ | 921          | 20.9                    | 0.67 | 12.9    | 100  |
| MAPbI$_3$           | 900          | 11.4                   | 0.60 | 6.30    | 103  |
| MAPbI$_3$           | 900          | 16.12                  | 0.60 | 8.68    | 103  |
| CsPbBr$_3$ (mesoscopic) | 1270        | 6.29                    | 0.60 | 4.77    | 125  |
| CsPbBr$_3$ (planar) | 1330         | 5.09                    | 0.55 | 3.67    | 125  |
carbon-based PSCs. In 2014, Rong et al. synthesized TiO$_2$ nanosheets (NSs) by a hydrothermal route in hydrofuran solution, and compared the resulting PCE with that of TiO$_2$ NPs.\textsuperscript{142} The structure of TiO$_2$ NSs ensured a higher number of (001) facets, with respect to that observed for TiO$_2$ NPs, due to the elongated geometry. It was found that the (001) anatase facet was slightly more reactive than the (101) counterpart because of the larger number of titanium atoms without coordination.\textsuperscript{143} In addition, the (001) facet presented a higher charge density, that may strengthen the interaction with the active absorbing material. Consequently, electron injection was facilitated.\textsuperscript{144,145} As shown in J–V curves (see Fig. 6A), $V_{oc}$, $J_{sc}$ and PCE of the NSs-based device were higher than those of the NPs-based one (868 mV, 20.1 mA cm$^{-2}$ and 10.64% vs. 839 mV, 13.7 mA cm$^{-2}$ and 7.36%); a TEM image of NSs is shown in Fig. 6B.

Similar results were obtained by Zhang et al. in 2018.\textsuperscript{146} They built two types of 1 mm-thick anodes, one based on TiO$_2$ NSs (see Fig. 6D) and the other one based on TiO$_2$ NPs. The PCE of the NS-based device shown in Fig. 6C was almost twice that of the NP-based one (8.11% vs. 4.37%, respectively), confirming that the (001) anatase facet was more reactive and suitable for perovskite adhesion and electron injection. In addition, they varied the TiO$_2$ layer thickness from 0.3 to 1.4 mm, and found that the best results were provided by 1 mm-thick TiO$_2$. In fact, recombination decreased by increasing the layer thickness, as confirmed by electrochemical impedance spectroscopy (EIS) analysis.\textsuperscript{147} However, a too thick TiO$_2$ layer caused high series resistance.

Another strategy used to ameliorate the anode performances is the surface modification of TiO$_2$. In 2015, Liu et al. reported a HTM-free PSC with an organic silane self-assembled between TiO$_2$ and the perovskite.\textsuperscript{148} The silanization of the anode was achieved by dipping a high-temperature processed TiO$_2$/ZrO$_2$/carbon substrate and subsequent hydrolysis. This extra layer drastically reduced recombination processes at the interphase between the perovskite and TiO$_2$, as confirmed by the drastically increased recombination resistance in the EIS spectrum. PCE values were determined as a function of silane treatment time. Efficiencies of 9.71%, 11.10%, 12.41%, 12.77% and 10.44% were reached with treatment times of 0, 0.5, 1, 4 and 24 h, respectively. Therefore, the PCE increased with treatment time, till a maximum of 4 h. After that, an excess of silane on the TiO$_2$ surface caused poor infiltration of the perovskite precursor solution.

In order to replace expensive HTMs, many p-type inorganic semiconductors have been reported. Among them, NiO showed several advantages, such as low cost, stability, wide band-gap and high hole mobility.\textsuperscript{149} In 2015, Cao et al. used NiO as the HTM and reported a TiO$_2$/Al$_2$O$_3$/NiO/carbon architecture with infiltrated perovskite precursor solution, as shown in Fig. 7A.\textsuperscript{150} All layers were deposited by the screen-printing method, ensuring a reproducible and scalable procedure. NiO successfully increased the PV efficiency. As a matter of fact, the PCE
increased from 11.20% (for the NiO-free PSC) to 15.03% (for the NiO-containing PSC). It was suggested that NiO limited charge recombination and ameliorated hole collection efficiency toward the carbon electrode.

A similar architecture was reported in the same year by Xu et al.\textsuperscript{151} ZrO\textsubscript{2} was used as both spacer and insulator to keep the TiO\textsubscript{2} anode and NiO semiconductor separated. The resulting TiO\textsubscript{2}/ZrO\textsubscript{2}/NiO/carbon(MAPbI\textsubscript{3}) architecture presented a p–i–n configuration as shown in Fig. 7B and C. Besides enhancing charge collection at the interface with the carbon back electrode, NiO successfully acted also as an electron blocking layer, thus reducing charge recombination. As a matter of fact, the device with this architecture attained remarkably high photovoltaic performances, with a $V_{oc}$ of 917 mV, a $J_{sc}$ of 21.36 mA cm\textsuperscript{-2}, an FF of 0.76, an average PCE of 13.7% and a maximum PCE of 14.9%. Long-term stability of the device was confirmed, since at room temperature and at 40% RH it retained 93% of the initial PCE after 1000 h of aging under dark conditions. The device was not sealed; in fact, hydrophobic carbon can prevent atmospheric moisture from penetrating the perovskite layer.

Charge transport in NiO can be attributed to positively-charged nickel vacancies. By increasing the temperature, the removal of some nickel(ii) cations results in increasing the number of oxygen atoms surrounding the remaining nickel atoms. For this reason, some nickel(ii) are oxidized to nickel(iii) and conductivity is attributed to hole hopping from the Ni\textsuperscript{3+} sites to Ni\textsuperscript{2+} ones. To further enhance charge mobility, NiO can be treated with ultraviolet (UV)/O\textsubscript{3} and annealed at high temperature. The UV/O\textsubscript{3} treatment generates NiOOH species on
the surface of nickel oxide. Hydroxyl groups are then removed by thermal treatment. Ni$_2$O$_3$ with a low transport resistance and a high charge mobility is formed. This treatment has been used by Behrouznejad et al. to provide a mesoscopic PSC with a NiO/carbon back electrode.$^{152}$ When a 450 nm-thick Al$_2$O$_3$ space layer was used, an average PCE of 11.28% (with best device having a PCE of 12.12%) was achieved. However, a drop in PCE of 18.2% was observed after three months.

Aside from NiO, other transition metal oxides were taken into account as the HTM for PSCs.$^{153}$ However, it was found out that only NiO had a positive effect on PV parameters. In fact, MoO$_3$, Co$_2$O$_3$, and CuO did not ensure an efficient interaction with the perovskite phase and the PCEs of the corresponding devices were quite lower than those of NiO-based devices: 0.82%, 6.93%, 6.08% and 13.94%, respectively.

Besides using NiO as the HTM, heteroatomic doping of graphite may be a suitable strategy for modifying the work function of carbon electrode and facilitating charge extraction.$^{154,155}$ In fact, the tiny difference between the work function of carbon and the valence band of perovskite causes a carrier extraction barrier at the interphase, limiting PV performances. Carbon atoms in the graphiteic layers can be easily replaced by electron-deficient boron by high-temperature thermal treatment with B$_4$C, as shown in Fig. 8A.$^{156,157}$ After this treatment, the work function of the carbon electrode was improved by the incorporation of the electron-deficient boron atom (5.10 vs. 4.81 eV of pure graphite, PG). In addition, boron-doped graphite (BdG) showed a lower sheet resistance (16 vs. 35 Ω sq$^{-1}$ of the PG). This was attributed to a better graphite layer packing and higher graphitization degree, as confirmed by XRD patterns. BdG-based devices showed improved $V_{oc}$, FF and PCE with respect to PG-based cells ($940$ mV, 0.63 and 13.6% vs. 900 mV, 0.60 and 12.4%, respectively). The increased FF and reduced hysteresis effect were ascribed to the more favorable band alignment in BdG-cells.$^{157}$

Other alternatives to NiO are p-type spinel oxides that are promising HTMs due to their excellent hole mobility. The first use of p-type Co$_3$O$_4$ as the HTM in carbon-based PSCs was reported in 2018 by Bashir et al.$^{158}$ Co$_3$O$_4$ has a cubic structure with high spin Co(II) in the tetrahedral sites and low spin Co(III) in the octahedral sites. In addition, the Co$_3$O$_4$ VB (5.3 eV) matches well with that of perovskite (5.4 eV), favoring efficient hole extraction. The effective contact between the perovskite and cobalt oxide and the improved hole extraction at the carbon electrode were demonstrated by the negligible hysteresis effect and by a high PCE of 13.27% ($V_{oc}$ = 0.88 V, $J_{sc}$ = 23.43 mA cm$^{-2}$, FF = 0.64).

Despite their dark appearance, carbon-based materials can also be used as transparent electrodes, necessary for semitransparent solar cells.$^{159}$ An ideal transparent electrode should exhibit high transparency, low resistance and charge collection efficiency. In this view, a thick layer of carbon-based materials may be the ideal candidate. Graphene and CNTs have already been successfully applied as back electrodes in semitransparent PSCs.$^{160,161}$ However, in most cases, they are still prepared by

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**Fig. 8** (A) Preparation of BdG starting from graphite and B$_4$C. (B) Schematic diagrams of carbon back electrodes with different carbon grids (blue represents carbon), and (C) optical images of as-fabricated semitransparent PSCs. Adapted and reprinted with permission from ref. 157 and 162.
non-scalable procedures, such as transferring and lamination. For these reasons, in 2017 Li et al. reported for the first time a fully printable carbon-based PSC with a TiO$_2$(5-AVA)$_{0.05}$(MA)$_{0.95}$PbI$_3$/Al$_2$O$_3$/carbon architecture.

The carbon layer was deposited by the screen-printing method, using several grids with different geometries, as reported in Fig. 8B and C. For comparison, a carbon-based PSC without grid lines was also fabricated (device 0). Device 1 showed the best performances, even when compared to device 0, with a $V_{oc}$ of 0.87 V, a $J_{sc}$ of 21.07 mA cm$^{-2}$, a FF of 58.9% and a PCE of 10.80% (Table 3). This result was due to the fact that the smaller grid lines of device 1 ensured a better pore filling of the perovskite precursor solution. The decreased PCE with increasing number of grid lines was ascribed to the increase in series resistance of the electrode. To solve this issue, MWCNTs were coated on top of device 6. MWCNTs enhanced conductivity by creating a charge transport network. In fact, the PCE dramatically increased from 5.08% to 8.04%.

Further issues for the realization of scalable high-temperature processed carbon-based PSCs are (i) mechanical infiltration of the perovskite solution, in order to have a fully reproducible and printable fabrication method; (ii) long-term stability under UV-light soaking; (iii) high-temperature thermal stability; and (iv) enlargement of the active area of the cell. The manual infiltration of the perovskite precursor solution onto the carbon layer limits reproducibility and is also time-consuming. For this reason, Hashmi et al. reported in 2017 a suitable strategy for the inkjet infiltration of the perovskite ink in HTM-free carbon-based PSCs. The precursor ink was loaded into a disposable cartridge and deposited by an inkjet printer on the high-temperature processed carbon electrode. 5-AVA iodide was added in the precursor ink to slow down perovskite crystal growth, thus preventing the inkjet printer cartridge from clogging and ensuring a better filling through the carbon layer. In particular, 10 × 10 cm$^2$ fluorine-doped tin oxide (FTO)-glass substrates having 18 individual cells of HTM-free triple layer PSCs were fabricated, as shown in Fig. 9A. Photovoltaic parameters showed improved reproducibility, with average values of 0.845 V ($V_{oc}$), 15.2 mA cm$^{-2}$ ($J_{sc}$), 63.5% (FF) and 8.15% (PCE). PCE of the devices stored under dark conditions was found to initially increase by 13% in the first three weeks. This was ascribed to a further curing of the perovskite phase. Then, the PCE restored to its initial value and remained stable for 1046 h. On the other hand, the stability under sunlight soaking conditions at 35 °C was also determined. In this case, after 1046 h, the drop in PCE was only 4.5%. The sluggish degradation of the perovskite, with formation of PbI$_2$, was observed and monitored by the camera imaging technique.

UV light exposure has been identified as one of the key issues for the evaluation of PSCs’ long-term stability. For this reason, the same cells obtained by inkjet printing and stored without encapsulation were subjected to intense 1.5 sun UV light illumination in an electronic weather chamber, at 45% RH and 40 °C. In the first 250 h, the cell ameliorated its performances in both $J_{sc}$ (+17%) and PCE (8.6%); this behavior may be ascribed to the additional curing of perovskite crystals under UV light. After this time, a slow degradation of cell
performance started. Part of the dark-brown perovskite layer became yellow after 751 h, due to the formation of PbI₂ and the disappearance of HI and CH₃NH₂. The JSC reduced by 25% and the PCE by 28%. Some batches were sealed with epoxy glue to prevent contact with moisture. Epoxy did not dissolve any phase and did not penetrate into the carbon layer over the photo-active area. In these cells, the average JSC and PCE were maintained after 1002 h. XRD demonstrated that PbI₂ reflections appeared only in epoxy-free aged samples. PL measurements demonstrated that the carrier lifetimes of batch 1 (without epoxy) and batch 2 (with epoxy) were 21.6 ns and 4.1 ns, respectively. This means that the epoxied samples were more likely to accelerate light-induced charge separation, thus restricting charge recombination, which correlated well with the results of aging tests.

Thermal stability is necessary for outdoor application of PSCs in many warm areas of the Earth, for example desert regions. However, the perovskite undergoes a phase transition from tetragonal to cubic at 54 °C. For these reasons, Baranwal et al. reported a three-layer printable HTM-free PSC with carbon as the back electrode, able to keep stability till 100 °C. Despite the hydrophobic nature of this carbon component, owing to its porous nature, sealing of the device was necessary under high-temperature conditions. Sealing was conducted with a UV-cured gel and a glass cover, in two different configurations (over-sealed and side-sealed, as shown in Fig. 9B and C). Sealed and unsealed devices were kept at 100 °C before measuring J-V curves. Both the unsealed and over-sealed devices lost 20% of their initial efficiency in the first 30 h. In contrast, side-sealed devices did not degrade during thermal tests and their PCE remained constant over 1500 h (see Fig. 9D), suggesting that sealing is a necessary protection against moisture, but it could also cause internal decomposition in the over-sealed configuration.

Finally, the enlargement of the active area of the cell is a desirable goal for the real application of a PV solar cell. However, this task is not trivial when a thermal treatment, which may cause the cracking or the bending of the glass, is involved. Both high-temperature treated TiO₂ and carbon may suffer from poor uniformity and compactness. In 2014, De Rossi et al. reported the first example of A4-size carbon-based PSC, with an active area of 198 cm². They optimized the printing process with the method of edge registration to avoid short circuit and damage to the layers. The PSC was composed of 22 units, with dimensions of 5 x 180 mm² and spaced by 6 mm, as shown in Fig. 10. A 1.5 mm-wide ZrO₂ layer and a low-temperature-treated TiO₂ layer were used as the insulator and blocking layer, respectively. A 10 μm-thick carbon layer was deposited by the doctor-blade technique in order to minimize the series resistance between adjacent units. With this large-area device, the highest PCE was 3.2%.

2.2 Low-temperature processed back electrodes

High-temperature thermal treatment of carbon-based electrodes is a time and energy-consuming method, that may limit large-scale manufacturing of PSCs. In addition, high-temperature processed carbon cannot be applied in plastic and flexible devices.

In 2014, Wei, Zhou and Zhang reported the first examples of low-temperature processed carbon electrodes for PSCs. In particular, the innovation introduced by Wei was the use of an ink containing both carbon and CH₃NH₃I in isopropanol, that was directly printed on FTO/TiO₂/PbI₂, as shown in Fig. 11A. In this way, the deposition of the carbon layer and the formation of perovskite phase occurred simultaneously, ensuring an efficient interfacial contact between CH₃NH₃I/PbI₂ and carbon. For comparison, another cell was prepared by the “traditional” route; after the deposition of the carbon layer onto the FTO/TiO₂/PbI₂ substrate, the cell was soaked into the CH₃NH₃I solution, as confirmed in the SEM images shown in Fig. 11B and C. Recombination resistance (Rrec) was estimated by fitting EIS data. It was confirmed that charge recombination was several times lower in the cell obtained with the CH₃NH₃I/polymer ink. Consequently, PV parameters improved: Voc increased from 0.90 to 0.95 V, JSC from 15.00 to 17.20 mA cm⁻², FF from 0.63 to 0.71 and PCE from 8.51 to 11.60%. Finally, devices maintained about 90% of their initial PCE after 12 days of dark storage without sealing.

Zhou et al. investigated for the first time the influence of carbon layer thickness on the performance of low-temperature processed PSCs. Carbon particles were dispersed in chlorobenzene and deposited by the doctor-blade technique, varying the thickness in the range 2–40 μm. It was found that the PCE of devices increased with carbon layer thickness till a maximum of 20 μm. 2.1 μm-thick carbon presented a high sheet resistance
of 160 Ω, therefore the corresponding cell exhibited a poor PCE of 2.25%. On the other hand, the best PCE of 6.90% was obtained with a carbon layer thickness of 20.6 μm. Even though a clear explanation for this structure–performance correlation was not provided by the authors, cell stability was confirmed for up to 2446 h. Such a good result under long-term testing was ascribed to the presence of hydrophobic carbon, which impeded the penetration of moisture, and to the absence of noble metals, which could corrode CH₃NH₃PbI₃.

The intrinsic resistivity of graphite is high. However, it can be reduced by introducing carbon black NPs (CBNPs) and nano-graphite powder or by ameliorating carbon morphology. In literature studies, PV parameters were influenced by the relative content of CBNPs with respect to graphite. In fact, the addition of CBNPs in the ratios 5:1 and 3:1 dramatically increased the PCE, as shown in Table 4 (entries I–III). This improvement was attributed to the reduced sheet resistance of the carbon material, that changed from 1.11 to 0.69 Ω cm (after having been pressed at 0.25 MPa). However, a further increase of CBNPs (ratio 2:1) slightly diminished cell performances (see Table 4, entry IV); in fact, an excess of CBNPs brought about a stronger recombination.

A similar strategy was followed by Liu and coworkers. Besides CBNPs, they also explored the effect of nano-graphite powder with different particle dimensions of 40 and 400 nm. The introduction of nano-graphite powder made the carbon film more compact and improved the interfacial contact. Thus, PV performances ameliorated after the introduction of nano-graphite powder. Table 4 (entries VI and VII) reveals that cells containing smaller nano-graphite powder (40 nm) perform better than cells containing larger graphite powder (400 nm). Larger particles decreased the contact between grains, establishing an overpotential at the interphase. However, increasing the content of 40 nm-graphite powder did not lead to a further increase of PCE (see Table 4, entry VIII). In fact, too much nano-graphite powder could have led to higher resistance and stronger recombination.

It has to be noted that, even though the cell architectures are identical, PCE values obtained by Wei are remarkably higher than those obtained by Liu (13.53% vs. 4.47%, respectively), as shown in Table 4 (entries III and V). In fact, the latter researcher fabricated the devices under uncontrolled high humidity conditions, which may have deteriorated the perovskite layer, thus limiting the PCE.

The poor interface contact between carbon and the perovskite phase is the main efficiency-limiting factor of low-temperature carbon-based PSCs with respect to gold-based PSCs. In fact, commercial carbon pastes contain solvents that, during evaporation, can create bulges and pinholes in the material, and also the perovskite layer may present holes that are not filled with the carbon material. A suitable solution to overcome this issue was proposed by Zhang et al. in 2018. They fabricated a self-adhesive carbon film, processed at room temperature by the solvent exchange method. A carbon electrode, namely C2, was doctor-bladed on a glass, soaked in ethanol and dried. It exhibited a macroporous, flexible morphology, with a thickness of 60 μm. It was removed from the glass and pressed onto the perovskite layer. The adhesion was ensured by its compressibility: in fact, the thickness was reduced to 31.9 μm, allowing an optimal contact at the interphase. Furthermore, after being pressed, its sheet resistance decreased from 424 to 58 Ω sq⁻¹. For comparison, a carbon electrode, namely, C1, with a similar thickness was created by the traditional route at 100 °C, without being soaked in ethanol and transferred from another substrate. As expected, the PCE of the C2-cell was dramatically higher than that reached by the C1-based cell (19.2% vs. 15.2%, see Table 4, entry IX).
Another suitable solution to overcome the poor contact between the perovskite phase and carbon back electrode was proposed by Cheng et al. in 2016. As demonstrated by the SEM image shown in Fig. 11D, the CH$_3$NH$_3$PbI$_3$ layer had a rough surface, with particles of different dimensions. On the other hand, the carbon material was composed of nano-sized carbon black and micro-sized graphite flakes. The latter, due to the large dimension, did not provide an effective contact with perovskite crystals, thus reducing the PCE to 9.8% (see Table 4, entry X). To solve this issue, MWCNTs were used. CNTs consist of a graphene layer bent and joined in one direction, to form a hollow cylinder. MWCNTs are composed of more nested CNTs, aligned together. Similar to other graphene-based materials, CNTs exhibit high electrical conductivity and thermal stability. MWCNTs with a diameter of 55 nm were added that can act as recombination centers. On the other hand, DEA-covered films were compact and homogeneous and fully covered the FTO substrate. A remarkable PCE of 11.92% was observed when a Ti(OC$_4$H$_9$)$_4$ : DEA = 1 : 0.75 ratio was used. Besides carbon, TiO$_2$ and insulating ZrO$_2$ also require high-temperature sintering and are not suitable for flexible applications, thus hindering the fabrication of PSCs. ZnO is a suitable n-type semiconductor, which requires only low-temperature processing (120 °C) and can be deposited by spin coating. Since its conduction band is at a lower energy with respect to that of the perovskite, electron injection is possible between excited CH$_3$NH$_3$PbI$_3$ and ZnO as shown in Fig. 12A. Therefore, it can replace the role of TiO$_2$. In 2015, Zhou et al. reported the first example of a TiO$_2$- and HTM-free carbon-based PSC with the simple architecture ZnO/CH$_3$NH$_3$PbI$_3$/carbon. The thickness

![Image](https://example.com/fig12)

**Table 4** Comparison between PV parameters, obtained under 1 sun illumination, for PSCs fabricated by varying the composition of the carbon-based electrode

| Entry | Carbon electrode composition | $V_{oc}$ (V) | $J_{sc}$ (mA cm$^{-2}$) | FF | PCE (%) | Ref. |
|-------|-----------------------------|--------------|-------------------------|----|---------|------|
| I     | Graphite                    | 0.97         | 18.8                    | 0.57 | 10.27   | 176  |
| II    | Graphite/CBNPs 5:1         | 0.98         | 20.0                    | 0.58 | 11.43   | 176  |
| III   | Graphite/CBNPs 3:1         | 1.00         | 21.3                    | 0.63 | 13.53   | 176  |
| IV    | Graphite/CBNPs 2:1         | 0.97         | 20.6                    | 0.62 | 12.47   | 176  |
| V     | Graphite/CBNPs 3:1         | 0.79         | 13.0                    | 0.43 | 4.47    | 177  |
| VI    | Graphite/CBNPs/nanographite powder (40 nm) 1:1:2 | 0.82 | 14.9 | 0.43 | 5.31 | 177 |
| VII   | Graphite/CBNPs/nanographite powder (40 nm) 1:1:2 | 0.84 | 16.8 | 0.43 | 6.16 | 177 |
| VIII  | Graphite/CBNPs/nanographite powder (40 nm) 1:1:5 | 0.87 | 11.0 | 0.40 | 3.84 | 177 |
| IX    | Self-adhesive carbon film   | 1.08         | 23.3                    | 0.76 | 19.2    | 178  |
| X     | Carbon                      | 0.87         | 20.2                    | 0.56 | 9.8     | 179  |
| XI    | Carbon/MWCNT                | 0.93         | 21.3                    | 0.59 | 11.6    | 179  |

*Indicates cells fabricated under high humidity conditions.
of the ZnO layer had a strong influence on PSC efficiency. In fact, a too thin layer (25 nm) was not completely covered by the active material and presented many holes. Its PCE and $V_{oc}$ were limited to 0.90% and 0.47 V, respectively. On the other hand, a too thick ZnO layer (89 nm) showed a high series resistance and an increased recombination rate. In fact, PCE reached a small value of 3.55%, even though its $V_{oc}$ of 0.80 V was the highest found in this kind of system. The optimal ZnO thickness was found to be 55 nm ($V_{oc}$ of 0.74 V, $J_{sc}$ of 20.68 mA cm$^{-2}$, FF of 0.46 and PCE of 7.05% in the forward scan).

Another ZnO-based low-temperature processed PSC was reported by Jin et al. in 2015.183 In this work, all films were deposited at room temperature. An electrochemically exfoliated graphite layer was used as the HTM to improve the extraction efficiency between the perovskite and the carbon back electrode. The obtained graphene had an excellent hole mobility of $\approx 310$ cm$^2$ V$^{-1}$ s$^{-1}$. The atomic force microscopy (AFM) image (Fig. 12B) showed that the thickness of graphene sheets was lower than 10 nm, corresponding to 30 atomic layers and the majority of graphite was in the form of NPs. Overall, the carbon-based back electrode was quite compact and completely covered the perovskite film. An average PCE of 7.2% was reached with this architecture.

SnO$_2$ is a high band-gap semiconductor ($E_g = 3.6$ eV), with high charge mobility and without a UV photocatalytic effect.184 It has already been employed to replace mesoporous TiO$_2$ as the anode material for PSCs, due to its low-temperature processability.185 Metal doping of SnO$_2$ further increases its electronic properties, making it more suitable for application in the PSC field. In particular zinc-doping shifts the work function of SnO$_2$ from $-4.28$ to $-4.19$ eV, resulting in a more favorable band alignment, as shown in Fig. 12C. Ye et al. reported in 2019 an entirely low-temperature processed carbon-based PSC with a Zn–SnO$_2$ ETM, as shown in Fig. 12D.186 FF and PCE were strongly improved, with respect to the undoped SnO$_2$-based cell. In particular, PCE increased from 15.31% to 17.78%. The more favorable band alignment generated a stronger electron injection from the excited active material to the ETM.

Even if not necessary, the HTM plays the important role of helping in hole extraction and ameliorating surface contact between the perovskite phase and carbon electrode. The most common organic HTMs are spiro-OMeTAD, which is expensive and needs to be doped with LiTFSI to improve conductivity. The latter is a hygroscopic salt that threatens the stability of the perovskite layer.187 Many efforts have been made for the replacement of spiro-OMeTAD with polymeric188,189, organic24,190,191 and inorganic HTMs.192–194 In this framework, poly(3-ethyl-3-hexylthiophene) (P3HT) is an intrinsic semiconductor with high stability (up to 350 °C in air) and low cost. Mashhoun et al. showed in 2018 that it can efficiently work as the HTM in carbon-based PSCs.188 Since the carbon back electrode is usually deposited from carbon pastes, the solvent content in the suspension should not be harmful for the underlying layer. For this reason, the effect of the nature of the solvent was investigated. In particular, the lowest and the highest PCEs were found with chlorobenzene and toluene, respectively (see Table 5, entries I–III). PV performances ameliorated by decreasing the solubility of P3HT in the solvent, but a clear explanation was not given by the authors. Further interface engineering with TaWO$_x$ NPs ameliorated the contact between P3HT and carbon, thus enhancing the PCE from 5.04% to 11.58%, as shown in Table 5 (entry IV).

| Entry | Solvent   | P3HT solubility (mg mL$^{-1}$) | $V_{oc}$ (V) | $J_{sc}$ (mA cm$^{-2}$) | FF (%) | PCE (%) |
|-------|-----------|-------------------------------|-------------|------------------------|--------|---------|
| I     | Chlorobenzene | 15.9                         | 0.545       | 6.27                   | 0.36   | 1.27    |
| II    | Xylene    | 2.7                          | 0.662       | 13.46                  | 0.40   | 3.45    |
| III   | Toluene   | 0.7                          | 0.764       | 15.86                  | 0.42   | 5.04    |
| IV    | Toluene$^a$ | 0.7                          | 1.012       | 16.35                  | 0.70   | 11.58   |

$^a$ Indicates a cell with TaWO$_x$-doped P3HT.

Smaller molecule-based HTMs have the advantages of facile synthesis and tunable electronic properties. In addition, in most cases, they do not require the use of a dopant to increase conductivity. Among organic HTMs, copper phthalocyanine (CuPC) is noteworthy.195 It is a strong conjugated planar molecule, with a tendency to form π–π stacking in the film state. Its p-type semiconductor character, low cost, ease of preparation, good thermal stability and long charge diffusion length make it the ideal candidate as the HTM in PSCs.196–198 Zhang et al. used for the first time CuPc as the HTM in carbon-based PSCs.190 Results underlined the importance of CuPc, since every PV parameter improved with respect to the HTM-free cell. In particular, the PCE increased from 9.0 to 16.1%. For comparison, an analogous cell with spiro-OMeTAD as the HTM was also reported. It showed an efficiency of 15.0%, that was still lower than that of the CuPc-cell. In addition, the device with CuPc was more stable: PCE dropped by only 8.5% in 600 h, as shown in Fig. 13A.

Another n-delocalized small molecular HTM was reported by Zhang et al. in 2015.19 Triazatruxene (TAT) contains three indole units combined by one benzene. Due to its electron rich aromatic structure, it has a strong charge carrier ability and has already been employed as the HTM in the field of organic PVs.199,200 Its derivative 5,10,15-triphenyl-5H-diiindolo[3,2-a:3',2'-c] carbazole (TPDI) shows the same properties and, in addition, exhibits good thermal stability and compatible band alignment with both carbon and CH$_3$NH$_2$PbI$_3$. When both TPDI and spiro-OMeTAD were doped with LiTFSI, TPDI-based cells showed comparable PCE with respect to spiro-OMeTAD-based ones.
the HTM/perovskite interface. A novel fluorinated HTM was reported by Ren et al., i.e. BDT2MeDPA, composed of difluorobenzene, benzo[1,2-b:4,5-b']dithiophene (BDT) and 4,4'-dimethoxydiphenylamine (DPA) groups. Electro-donating BDT presented excellent charge transport properties and facilitated the synthetic process. For comparison, the fluorine-free counterpart (BDT0MeDPA) was also used as the HTM. PV performance remarkably increased with the use of the novel HTM. When BDT0FMeDPA was used, $J_{sc}$ shifted from 12.5 to 17.2 mA cm$^{-2}$ and the PCE increased from 8.4% to 11.3%, while $V_{oc}$ maintained a constant value of 0.98 V. BDT2MeDPA-based cells showed a further increase in $J_{sc}$ (21.4 mA cm$^{-2}$) and PCE (14.5%), that were slightly lower than those obtained with spiro-OmEATAD (PCE = 15.2%). PL quenching was stronger in the BDT2FMeDPA/perovskite than in the BDT0FMeDPA/perovskite, underlining the well matched energy levels and efficient charge extraction process.

Inorganic p-type HTMs are more thermally stable with respect to molecular ones. The most commonly used are CuS, CuSCN and NiO. CuS has already been investigated in DSSCs, and was reported as the HTM for PSCs for the first time in 2018. Hu et al. fabricated a carbon-based PSC where small particles of CuS, obtained by a low-temperature precipitation process, were added to the carbon paste to increase the hole-extraction ability of the cathodic hybrid material. The resulting carbon electrode had a thickness of 10 μm and CuS presented its nanostructure form below 100 nm, as shown by SEM image in Fig. 13B. The EDX spectrum evidenced the homogeneous distribution of CuS into the carbon electrode.

CuS was added in the carbon paste in the ratio of 0.5–2 wt%. $J$-$V$ curves evidenced that a small CuS-doping increased by 21% the performances of the carbon-based PSCs. The average PCE shifted from 8.41% for the undoped device to 9.32% for the 0.5%-doped one and 10.22% for the 1%-doped one, highlighting the increased hole collection ability of the doped electrode. On the other hand, efficiency reduced when the amount of CuS exceeded 1 wt%, due to the smaller conductivity of CuS with respect to graphite. For these reasons, the optimal amount of CuS was 1 wt%.

Another copper-based inorganic HTM was reported by Lv et al. in 2018. CuSCN is an inorganic p-type semiconductor, with low cost and high stability. It was deposited by spin coating under the perovskite layer with a thickness of 300 nm and established a good contact with both the absorber material and carbon back electrode. It caused a strong emission quenching in the PL spectrum of the perovskite, highlighting its effective contact and suitable band alignment with CH$_3$NH$_3$PbI$_3$. The effect of CuSCN on the PV performance was remarkable: $V_{oc}$ shifted from 0.72 to 0.78 V, $J_{sc}$ from 17.32 to 19.58 mA cm$^{-2}$, FF from 0.53 to 0.59 and PCE from 6.61% to 9.01%. This result was confirmed by Mashhoun, who found a PCE of 8.59% for CuSCN-based devices.

As already said, NiO has been largely used as the HTM in high-temperature processed carbon-based PSCs. Its properties of hole conduction and electron blocking effect have been exploited to suppress recombination and increase both $V_{oc}$ and $J_{sc}$.
However, in high-temperature processed PSCs the poor pore filling causes an inhomogeneous crystallization of the perovskite and this issue is worsened when NiO is used as the HTM. In 2017, Peiris et al. reported for the first time a mesoscopic PSC, with NiO as the HTM and low-temperature processed carbon as the back electrode.\textsuperscript{194} The perovskite was deposited on the FTO/TiO\textsubscript{2}/ZrO\textsubscript{2}/NiO substrate by a modified one-step method in the presence of N\textsubscript{2} flow. Subsequently, the carbon film was doctor-bladed on top of NiO: this strategy solved the problem of poor crystallization of CH\textsubscript{3}NH\textsubscript{3}PbI\textsubscript{3}. The PCE was almost doubled up from 5.96\% to 10.35\% by the presence of NiO. A lower charge transfer resistance was measured by EIS in the NiO-based device, confirming the more effective charge extraction and transfer. In addition, recombination resistance was stronger in NiO-based cells, suggesting that charge recombination was lower.

Another suitable strategy for replacing time and energy-consuming sintering is the NIR technology, that has already been used to thermally treat TiO\textsubscript{2} for DSSC application.\textsuperscript{203} NIR wavelengths are absorbed by a substrate, and converted into heat, causing the rapid warming of the substrate.\textsuperscript{204} If the substrate is transparent in the NIR range, a metal can be exploited to transfer heat. This method is very fast and allows energy saving. In 2017, Baker et al. reported the use of NIR technology for rapid thermal treatment (12.5 s) in mesoporous carbon-based PSCs.\textsuperscript{205} Both TiO\textsubscript{2} and ZrO\textsubscript{2} are transparent in the NIR range, so, when they were deposited and treated with NIR, radiation reached the FTO. The heat generated was enough to burn out all the organic additives of the two pastes. On the other hand, carbon has a high absorption coefficient in the NIR range; so, it absorbed all the radiation that, in this way, did not reach the underlying layer. Heat was generated only in the carbon layer, allowing the evaporation of solvents and volatilization of organic compounds present within the carbon paste. Consequently, the as-formed mesoporous layer was successfully infiltrated with the perovskite precursor solution. To the best of our knowledge, this is the only example of mesoscopic carbon-based PSCs obtained without thermal treatment. The champion device reached a $J_{SC}$ of 20 mA cm\textsuperscript{-2} and a stabilized PCE of 11\%.

As concluding remarks, a comparison of sheet resistance and thickness of carbon-based back electrodes prepared following different recipes, along with the best PCE values of the resulting devices, is given in Table 6 (for carbon-infiltrated perovskites) and Table 7 (for layered carbon electrodes).

Within the comparison between high and low temperature processes for the fabrication of carbon-based PSCs, it should be stressed that this distinction just refers to the preparation of cell components and device assembly. Indeed, many carbon electrode recipes contain graphite flakes that quite always come from synthetic sources, the latter requiring very high temperature treatments. Therefore, a life cycle assessment (LCA) study of

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|c|c|}
\hline
Paste composition & Deposition technique & Thermal treatment & Sheet resistance ($\Omega$ cm$^{-2}$) & Thickness (\textmu m) & PCE (%) \\
\hline
CB/flaky graphite & Blade coating & 400 °C for 30 min & 10.49 & 10 & 4.08 29 \\
CB/spheroidal graphite & Blade coating & 400 °C for 30 min & 8.02 & 10 & 6.64 29 \\
2 g CB (particle size 30 nm), 6 g graphite powder, 1 g ZrO\textsubscript{2} (particle size 20 nm), 1 g hydroxypropyl cellulose in 30 mL terpineol & Blade coating & 400 °C for 30 min & 11.47 & 9 & 11.63 87 \\
9 g graphite powder (particle dimension < 20 \textmu m), 3 g CB, 1 g ZrO\textsubscript{2} (particle dimension < 100 nm), 15 g ethylcellulose in 18 g terpineol & Screen-printing & 400 °C for 30 min & 56 & 25 & 10.7 88 \\
5 g graphite, 1 g CB (particle size 30 nm), 1 g ZrO\textsubscript{2} (particle size 20 nm), terpineol (30 mL) & Printing & 400 °C for 30 min & 5–20 & 5–16 & 12.63 94 \\
UG (obtained by exfoliation of BG), CB, hydroxypropyl cellulose, terpineol & Printing & 400 °C for 30 min & 5–25 & 5–16 & 14.01 94 \\
Carbon material [graphite : CB = 3 : 7], hydroxypropyl cellulose, terpineol & Screen-printing & 400 °C for 30 min & 23 & 11 & 15.70 92 \\
4 g graphite, 1 g CB, 0.6 g ZrO\textsubscript{2} in ethanol & Screen-printing & 400 °C for 60 min & 30.5 & Not reported & Not reported 13.7 151 \\
3.8 g graphite, 0.95 g CB, 0.57 g ZrO\textsubscript{2}, 0.28 g polystyrene spheres in ethanol & Screen-printing & 400 °C for 60 min & 56.7 & Not reported & Not reported 3.36 95 \\
3.6 g graphite, 0.9 g CB, 0.54 g ZrO\textsubscript{2}, 0.56 g polystyrene spheres in ethanol & Screen-printing & 400 °C for 60 min & 68.3 & Not reported & Not reported 3.87 95 \\
2.8 g graphite, 0.7 g CB, 0.42 g ZrO\textsubscript{2}, 1.68 g polystyrene spheres in ethanol & Screen-printing & 400 °C for 60 min & 180.7 & Not reported & Not reported 3.13 95 \\
6.5 g graphite, 2 g CB, 1 g ZrO\textsubscript{2}, 1 g hydroxypropyl cellulose, 30 mL terpineol & Screen-printing & 400 °C for 30 min & 35 & 11 & 12.4 157 \\
6.5 g BdG, 2 g CB, 1 g ZrO\textsubscript{2}, 1 g hydroxypropyl cellulose, 30 mL terpineol & Screen-printing & 400 °C for 30 min & 16 & 9.6 & 13.6 157 \\
CB (5 wt%), Al\textsubscript{2}O\textsubscript{3} (5 wt%), graphite (10 wt%), ethyl cellulose (20 wt%), terpineol (60 wt%) & Screen-printing & 125 °C for 10 min, 325 °C for 10 min, 375 °C for 10 min, 400 °C for 30 min & Not reported & 4–5 & 12.12 152 \\
2 g CB (particle size 30 nm), 6.5 g graphite, 1 g ZrO\textsubscript{2} (particle size 20 nm), 1 g hydroxypropyl cellulose, 30 mL terpineol & Screen-printing & 400 °C for 30 min & 8.98 & 10 & 14.15 153 \\
\hline
\end{tabular}
\caption{Comparison of sheet resistance and thickness of carbon-based back electrodes prepared following different recipes; the best PCE values of the resulting devices are also listed. The table refers to carbon-infiltrated perovskites.}
\end{table}
carbon-based PSCs (out of the scope of this review) should also consider the preparation of the graphite part of the electrodes, independently if the device assembly contains a high temperature step or not. A LCA study of a pre-industrial production process of large-area perovskite modules based on a carbon stack architecture was recently proposed by Alberola-Borras et al.\(^{207}\)

The perovskite layer was found to be the one with the greatest impact, mainly due to the energy consumed in the preparation and annealing of the precursor solution, rather than the lead content. As regards the back electrode, the inventory of the photovoltaic module included input/output relative to carbon, nitrocellulose, terpineol, screen-printing, annealing at 400 °C for 30 min, emissions and transportation burden by a lorry. No relevant mention was present concerning the source of carbon, therefore we further highlight the necessity to consider this point in future LCA approaches. Of course, if the carbon source comes from biomass or biosourced matter, the outcome of the LCA analysis will be markedly affected.

### 3 Graphene-based back electrodes

Among carbon-based materials, graphene, a 2D dimensional sheet of sp\(^2\) carbon atoms packed into a honeycomb crystal lattice, is surely noteworthy. Graphene presents several remarkable properties, such as high surface area, high charge mobility, both electrical and thermal conductivity, large mechanical flexibility and transmittance (it absorbs only 2.3% of the incident radiation from infrared, IR, to visible range).\(^{208}\) Due to its unique characteristics, it has been reported for catalysis,\(^{209}\) sensors,\(^{210}\) and optoelectronic devices (diodes, displays, solar cells, etc.).\(^{211,212}\) Concerning PSCs, many reviews summarize the applications of graphene.\(^{208,213–216}\) In particular, its high flexibility and transmittance make it suitable for numerous applications in flexible, portable and inverted PSCs.\(^{161,217,218}\)

Both planar and mesoscopic normal architectures require the use of high-temperature processed TiO\(_2\) to form a compact film. This issue is not only time- and energy-consuming, but also impedes the fabrication of plastic and flexible devices. On the other hand, in inverted architectures, the device is illuminated from the back electrode side. Photogenerated electrons and holes are collected by the ETM and HTM and reach the aluminum anode and the carbon electrode, respectively. Consequently, the latter needs to be highly transparent in both the visible and IR range. For this reason, graphene is the ideal candidate.\(^{208}\) Finally, the use of a transparent conductive oxide (TCO) is not necessary; indeed, plastic and transparent materials can be used, allowing the fabrication of flexible devices.

The first example of the use of graphene as the transparent electrode in inverted PSCs was reported by You et al. in 2015.\(^{161}\) Graphene was synthesized by the CVD method on copper foils, then coated with poly(methyl methacrylate) (PMMA) and poly(dimethyldisiloxane) (PDMS). The resulting graphene/PMMA/PDMS sample was peeled off from the substrate and deposited onto a perovskite/HTM layer. Poly(3,4-ethylenedioxythiophene)–poly(styrenesulfonate) (PEDOT:PSS) was used to increase the

### Table 7

| Paste composition | Deposition technique | Thermal treatment | Sheet resistance (Ω cm\(^{-2}\)) | Thickness (μm) | PCE (%) | Ref. |
|-------------------|---------------------|-------------------|-------------------------------|---------------|---------|------|
| 15 mg mL\(^{-1}\) CB in 2-propanol | Screen-printing | 100 °C for 60 min | Not reported | Not reported | 7.55 | 174 |
| 15 mg mL\(^{-1}\) CB, 10 mg mL\(^{-1}\) CH\(_3\)NH\(_3\)I in 2-propanol | Screen-printing | 100 °C for 60 min | Not reported | 40 | 10.03 | 174 |
| 20 wt% polyvinyl acetate, 80 wt% carbon material (graphite : CB = 1 : 0) | Doctor blade | 85 °C for 15 s | 1.11 | 40 | 10.27 | 176 |
| 20 wt% polyvinyl acetate, 80 wt% carbon material (graphite : CB = 5 : 1) | Doctor blade | 85 °C for 15 s | 0.75 | 40 | 11.43 | 176 |
| 20 wt% polyvinyl acetate, 80 wt% carbon material (graphite : CB = 3 : 1) | Doctor blade | 85 °C for 15 s | 0.69 | 40 | 13.53 | 176 |
| 20 wt% polyvinyl acetate, 80 wt% carbon material (graphite : CB = 2 : 1) | Doctor blade | 85 °C for 15 s | 0.61 | 40 | 12.47 | 176 |
| Commercial carbon paste (CC, Shenzhen DongDaLai Chemical Co., Ltd) | Doctor blade | 100 °C for 30 min | 6.4 | 10 | 16.1 | 190 |
| Graphite flakes (10 μm) : CB (40 nm) = 3 : 1 in ethyl acetate | Doctor blade | Not reported | 17 | 65 | 4.47 | 177 |
| Graphite flakes (10 μm) : CB (40 nm) : nano-graphite powder (400 nm) = 1 : 1 : 2 in ethyl acetate | Doctor blade | Not reported | 70 | 65 | 5.31 | 177 |
| Graphite flakes (10 μm) : CB (40 nm) : nano-graphite powder (400 nm) = 1 : 1 : 2 in ethyl acetate | Doctor blade | Not reported | 100 | 65 | 6.16 | 177 |
| Graphite, CB, PMMA ( binder), Y-037 ( solvent) | Doctor blade | 100 °C for 15 min | Not reported | 10.9 | 10.22 | 192 |
| Commercial carbon paste (Guangzhou Saidi Technology Co., Ltd) | Doctor blade | 100 °C (until the residual solvent is totally evaporated) | 34.5 | 15.2 | 178 |
| The layer was soaked in ethanol to remove the residual solvent | Printing | Not reported | Not reported | Not reported | 17.78 | 186 |
| Commercial carbon paste (Shenzhen DongDaLai Chemical Co., Ltd) | Blade coating | 100 °C for 15 min | Not reported | 11.2 | 11.92 | 206 |
hole extraction ability of graphene and ameliorate the contact with the perovskite. Furthermore, the addition of α-sorbitol to PEDOT:PSS improved the contact of the graphene layer with the HTM and the perovskite, and increased the PCE from 4.13% to 12.37% when the devices were illuminated from the FTO side. However, the PCE was almost the same when the graphene side was illuminated: 4.37% and 12.02% for devices without and with α-sorbitol, respectively. The PCE was strictly dependent on the number of graphene layers. The device with two layers showed the best PV parameters, with an efficiency of 12.37% vs. 9.18% for the cell assembled with the single layer. However, further increasing the number of graphene layers led to a worsening of PCE (11.45% and 11.27% for the triple and four-fold layers, respectively). In fact, a high number of layers decreased the transmittance of the electrode, resulting in a lower number of photons that reached the perovskite phase.

The use of the inverted architecture also allows the elimination of expensive, brittle and rigid TCO substrates. In 2016, Sung et al. reported an inverted cell, using graphene as the transparent cathode.217 MoO₃ was added to graphene to make its surface more hydrophilic, thus increasing the contact with the PEDOT:PSS used as the HTM, as shown in Fig. 14A–F. In addition, MoO₃ elevated the graphene work function from 4.23 to 4.71 eV, ameliorating the band alignment with the HTM. The best performing device showed a PCE of 16.1%, with a $V_{oc}$ of 1.03 V, a $J_{sc}$ of 21.9 mA cm⁻² and a FF of 0.72. The beneficial effect of MoO₃ was confirmed also by Yoon et al.,218 who reported the same architecture of Sung,217 but with superflexible polyethylene naphthalate (PEN) as the substrate. Devices with graphene/MoO₃ as the back electrode showed an average PCE of 15.0% ($V_{oc} = 0.99$ V, $J_{sc} = 21.0$ mA cm⁻², FF = 0.72), highlighting the role of MoO₃ in ensuring a better ohmic contact with the HTM.

A successful strategy for increasing the conductivity of carbon materials is nitrogen doping. In fact, the addition of lone pair electrons of nitrogen atoms can facilitate charge transfer. The nitrogen-doping of graphene for PSC application was performed by Zhu et al. in 2018.219 Nitrogen-doped graphene framework (NGF) was used as the cathode material and ensured a good network for charge transfer. PV parameters confirmed the increased properties of the NGF with respect to undoped samples. In fact, the PCE shifted from 8.98% to 10.32%, for pristine and NGF-based devices, respectively.

Finally, in a noteworthy work, Lou et al. used only carbon-based materials as electrodes for the fabrication of PSCs, as shown in Fig. 14G.220 In particular, graphene was chosen as the anode and was deposited on a poly(ethylene terephthalate) (PET) flexible substrate, due to its high transparency and flexibility; cross-stacking CNTs were used as the back electrode material, due to their high electrical conductivity. The optimum number of graphene layers was two, as reported by You.161 This double layer device demonstrated a $V_{oc}$ of 0.89 V, a $J_{sc}$ of 20.25 mA cm⁻², a FF of 0.65 and a PCE of 11.9%, and photographs of its components are shown in Fig. 14H and I.

4 Carbon nanotube-based back electrodes

CNTs are hollow cylinders resulting from the rolling up of a graphene layer. From their discovery in 1991,221 CNTs have attracted much attention in electronic and optoelectronic applications, due to their excellent charge transport ability, large hydrophobicity and high stability.222 CNTs are p-type semiconductors, with work functions ranging from −4.95 to −5.05 eV, suitable for working as back electrodes in PSCs. Their application in the PSC field has already been reviewed.75,221 Li et al. reported the first example of the use of CNTs as a back electrode in PSCs.160 They adopted an interfacial engineering method to improve the surface contact between CH₃NH₃PbI₃ and CNTs: drops of toluene were added on the top of CNT film after the deposition onto the perovskite phase. Toluene did not dissolve any phase, but wetted the surface of both films, driving CNTs toward CH₃NH₃PbI₃. PL quenching confirmed the strong adhesion through van der Waals interactions between the two films. As a result, this cell was able to convert solar radiation with a PCE of 6.87%.

Device performance can be dramatically increased by using MWCNTs,223 consisting of several CNTs stacked together providing a continuous and homogeneous film, strongly attached to the perovskite surface. This represents a strong advantage over other carbon-based materials, such as carbon black, which forms a loosely packed film easy to be cracked. Graphite forms...
a uniform layer too, but the larger particles do not lead to a perfect contact with the perovskite phase. In a HTM-free device, the use of MWCNTs as a back electrode provided a PCE of 10.30%,223

Zheng et al. made a further effort toward the optimization of MWCTNs for PSC application,224 starting from the idea that the work function of MWCTNs (~4.46 eV) was still too high compared with the VB of the perovskite. As already mentioned before,157 the boron-doping of carbon-based materials is a suitable strategy to reduce their work function. Boron-doped MWCTNs were obtained by the reaction of MWCTNs with boric acid and subsequent thermal treatment, as shown in Fig. 15A. The product showed the same structure of MWCTNs, suggesting that boron can replace carbon without any change in the graphitic structure, and the thermal annealing reduced the number of defects. MWCT- and boron-doped MWCTN-based devices attained average PCE values of 10.70% and 14.60%, respectively. The dramatic increase in PCE was ascribed to the ameliorated band alignment and improved charge extraction ability of boron-doped samples, as depicted in the scheme shown in Fig. 15B.

Like graphene,225 CNTs may also be exploited for the fabrication of transparent electrodes. In addition, their mechanical properties permit the use of flexible substrates, such as PET. In 2017 Jeon et al. demonstrated the use of CNTs as both front and back electrode materials.226 In these devices, the transparency of the electrode is fundamental to allow light reach the active material. In addition [6,6]-phenyl-C61-butyric acid methyl ester (PC61BM) and PEDOT:PSS were used as the ETM and HTM, respectively. This architecture (shown in Fig. 15C and D) was found to work well in both glass and PET-based flexible substrates, with a PCE of 7.3% and 7.1%, respectively.

As in the case of graphite and carbon black back electrodes, organic227–229 and inorganic149 HTMs have been exploited to further increase the hole extraction ability of CNTs.230,231 Aitola et al. reported a CNT-based PSC with spiro-OMeTAD as the HTM.228 An average PCE of 14.3% was achieved, still lower than that obtained with gold as the back electrode (18.4%). However, CNTs strongly increased the stability of the device at relatively high temperatures (60 °C): while the gold-based device degraded dramatically, exhibiting a loss of 20% in just 8 h, the CNT-based device was highly stable (loss of 0.04% in the same time). The migration of gold atoms and the penetration of air moisture were responsible for the degradation of the perovskite layer. On the other hand, these phenomena were suppressed by the use of hydrophobic CNTs.

P3HT is another suitable hydrophobic HTM for PSCs. It has already been used for graphite, carbon black and graphene electrodes.188,189 When used with CNTs, it acts as a strong crosslinker that binds together CNTs.227 Therefore, a uniform morphology and a network for charge transfer are obtained by using P3HT as an organic/polymeric HTM. With an optimized concentration of 2 mg mL⁻¹ in the precursor solution, an average PCE of 12.54% was obtained (Voc = 0.90 V, Jsc = 22.21 mA cm⁻², FF = 0.63).227 Below this concentration, the amount of P3HT was not enough to connect CNTs. On the other hand, above this concentration the excess of P3HT blocked the conductive pathway, increasing recombination phenomena.

A similar strategy involves the use of poly(ethylenimine) (PEI) in the CNT film.239 PEI played the important role of ameliorating the interface between carbon and PC61BM, the latter being used as an electron acceptor in inverted FTO/NiO/CH3NH3PbI3/PC61BM/CNTs:PEI architectures. The beneficial effect of PEI was demonstrated by the dramatic increase in PCE, i.e. 10.8% vs. 3.3% for the PEI-free device. This result was ascribed to the better contact and decreased energy barrier between CNTs and PC61BM.

Last, a p-type NiO semiconductor was demonstrated to work as a HTM in carbon-based PSCs.150–152,194 In 2017, Liu et al. reported the use of NiO as the HTM in fully printable CNT-based devices.149 The energy level of the NiO/CNTs composite matched well with that of the perovskite, ensuring a good hole extraction rate. The addition of NiO to the carbon material allowed the reduction of the thickness of the electrode (1.8 μm), while keeping the electrical conductivity constant. As a matter of fact, this device attained an average PCE of 12.7% (Voc = 0.945 V, Jsc = 20.7 mA cm⁻², FF = 0.64), while the corresponding device with 1.8 μm-thick graphite achieved a PCE of only 6.2%.

Fig. 15 (A) Schematic structure of MWCTNs before and after boron-doping and thermal treatment; (B) schematic diagram of the stronger hole extraction and transport capability of boron-MWCNTs (right) with respect to pristine MWCTNs (left) and thermally treated MWCTNs (center). (C) Schematic architecture of a CNTs/PC61BM/CH3NH3PbI3/PEDOT:PSS/CNTs device and (D) digital picture of a PET-based device, highlighting its remarkable flexibility. Adapted and reprinted with permission from ref. 224 and 226.
5 Stability of carbon-based PSCs

This review has clearly highlighted that using noble metals is problematic and expensive, but the main drawback for the commercialization of PSCs is the poor long-term stability of the devices. Being carbon electrode-based devices the best performing architectures in terms of stability, this advantage will be further stressed in this section. Stability is a widely discussed topic in the PSC framework, and the authors of this review agree with the general opinion that tests carried out under maximum power point (MPP) conditions possess wider significance than the vast majority of shelf stability tests reported in the literature. Therefore, even though it was surprisingly noted that the vast majority of research groups proposed shelf stability tests for their carbon-based devices, this section highlights the most significant results achieved under MPP conditions.

Bashir et al. investigated the stability of a HTM-free device fabricated with a thin layer of copper-doped nickel oxide (Cu:NiO\textsubscript{x}) NPs, which helped in improving the photocurrent and reducing the recombination resistance.\textsuperscript{325} MPP tracking was performed and the output current density was monitored under continuous illumination with a constant applied bias of 0.62 V. The current remained above 20 mA cm\textsuperscript{-2} for more than 50 h, and the device showed no sign of degradation for 60 h (see Fig. 16A). Moreover, a slight increase in current was observed during the experiment, and this was attributed to the interaction of the perovskite layer with ambient humidity. These researchers also fabricated a monolithic perovskite module on a 100 cm\textsuperscript{2} glass substrate (active area = 70 cm\textsuperscript{2}) and the champion device showed an impressive PCE of 12.1%.

Almost all of the carbon-based PSCs reported so far have been fabricated using TiO\textsubscript{2} as the ETM, the latter requiring a high-temperature sintering process. As an alternative, Zhou et al. proposed solution-processed hexamethonium bromide (HMB)-doped C\textsubscript{60} as the ETM, achieving a high PCE of over 16% without hysteresis.\textsuperscript{326} The remarkable stability of the resulting PSCs allows them to maintain nearly 90% of the initial performance after continuous operation for 338 h at MPP under 1 sun illumination in ambient air and with a 420 nm cut-off UV filter (see Fig. 16B).

Coupling carbon-based electrodes with all-air processing would truly lead to an easily upscalable PSC fabrication procedure. This goal was achieved by Zhou et al., who prepared MWCNT-incorporated FA\textsubscript{0.5}MA\textsubscript{1.5-x}Pb\textsubscript{0.5}Br\textsubscript{x-y} perovskite films in ambient air, where hydrophobic MWCNTs mediated the crystallization of the perovskite and prevented the erosion by moisture.\textsuperscript{327} With the low annealing temperature (< 150 °C) adopted, a PCE of 16.23% was achieved and the photostability of the devices without encapsulation was studied by MPP tracking under continuous 1 sun illumination at a RH of 30–80%. Cells fabricated with 0.5% MWCNTs maintained 94.9% of the initial PCE after 500 h, while pristine devices retained only 23.4% after 200 h, as shown in Fig. 16C.

The self-adhesive macroporous carbon electrodes proposed by Zhang et al. and previously described in this review were subjected to an aging test under a N\textsubscript{2} atmosphere, full AM1.5 sun-equivalent and UV-free conditions at the MPP.\textsuperscript{328} Gold-based PSCs suffered from much faster degradation due to the chemical reaction between the perovskite and gold, retaining just 22% of the initial PCE after 80 h (see Fig. 17A). Conversely, carbon-based
devices exhibited a short rapid degradation and subsequent recovery, due to the improvement of the back contact between the electrode and spiro-OMeTAD; this phenomenon was attributed to an interface passivation caused by ion migration under illumination and an electric field. Carbon-based cells retained over 94% of the initial PCE, indicating greatly improved stability with respect to their gold-based counterparts.

Meng et al. fabricated a carbon-based PSC without a HTM and using C_{60} as the ETM, reaching a PCE of 15.38% without hysteresis.\textsuperscript{328} Besides improved electron extraction, suppressed charge recombination and reduced sub-bandgap states at the C_{60}/MAPbI_{3} interface, the device showed moisture and ion migration resistance. As shown in Fig. 17B, the long-term stability was tested under continuous 1 sun illumination (with a 420 nm UV filter) at the MPP, and the devices assembled without any encapsulation retained 95% of their initial performance after 180 h in air, with a RH of 40–60%.

Since the commonly used MAPbI_{3} photoabsorber is ambient-unstable and incompatible with the low-cost mass-production of carbon-based PSCs, Zhou et al. fabricated (EA)\textsubscript{2}(MA)\textsubscript{n}/PbI\textsubscript{n+1} films (n = 20, 10, 6) and assembled ITO/C\textsubscript{60}/(EA)\textsubscript{2}(MA)\textsubscript{n}/PbI\textsubscript{n+1}/C devices with PCE exceeding 11.88\%.\textsuperscript{60} The photostability was assessed without encapsulation, by MPP tracking under continuous 1 sun illumination (without a cut-off UV filter) in air with a RH of 30–80%. The PCE was retained at 91.8% after 308 h, while that of the reference cell fabricated with MAPbI\textsubscript{3} dropped to 66.5% of the initial value after 83 h, as shown in Fig. 17C.

The inorganic interlayer of spinel Co\textsubscript{3}O\textsubscript{4} used by Bashir et al. to suppress charge recombination and extract holes efficiently, and described in the previous sections of this review, was further investigated in aging tests at the MPP.\textsuperscript{158} A slight increase in J\textsubscript{sc} was observed (see Fig. 18A), attributed to the interaction of the perovskite layer with ambient humidity, and the perovskite modules showed a degradation lower than 10% after continuous light soaking for 140 h.

Excess PbI\textsubscript{2} is commonly considered as a way to boost the PCE in standard PSCs, and Kapoor et al. checked this issue in fully printable carbon-based devices.\textsuperscript{328} An improvement in V\textsubscript{oc} was detected, while other parameters remained unchanged. Continuous simulated solar illumination under ambient conditions (25 °C and 65% RH) and the MPP was carried out for 68 h. Fig. 18B shows that no decrease of current was detected for the standard device, whereas the current reduced by 10% for the device containing 15% excess PbI\textsubscript{2}.

A 140 h MPP tracking of gold- and single-walled CNT-contacted devices under illumination at 1 sun in a N\textsubscript{2} atmosphere was carried out by Aitola et al.\textsuperscript{228} The cells were kept at 20 °C for the first 14 h, and the gold-based cell showed pronounced changes; then, a stabilization period occurred, which was not seen in high temperature experiments, where gold-based devices underwent an irreversible degradation process. Noteworthy, the CNT-based device exhibited only a small PCE drop in this region, as shown in Fig. 18C.

Carbon cloth and carbon fibers, widely used in DSSCs and batteries, replaced gold in a batch of PSCs proposed by Gholipour et al.\textsuperscript{330} The optimized device configuration led to a PCE of 14.8%. Long-term stability was studied under a N\textsubscript{2} atmosphere, 85 °C, constant 1 sun illumination and MPP tracking. While the gold-based cell suffered from rapid degradation, retaining less than 20% of the initial PCE after 20 h and less than 10% after 65 h, the carbon-based cell retained close to half of the starting
PCE after 115 h (see Fig. 18D). This slight degradation of the carbon-based cell was attributed to chemical reactions between spiro-OMeTAD$^+$ and migrating I$, hampering the conductivity of the HTM and thus the solar cell performance.

Several articles among those mentioned in this section focused on aging tests based on external degradation agents (e.g., moisture, oxygen, etc.), but no insight into the intrinsic degradation vector from the perovskite itself (releasing reactive gases) or studies of the true chemical inertness properties of carbon layers were provided, along with comparison with reference molecular/polymer-based HTMs. For sure it can be stated that, in the absence of HTMs, 4-tert-butylpyridine and organic lithium salt additives (typically used to dope spiro-OMeTAD) will not diffuse into the perovskite layer or absorb water traces; this surely improves the intrinsic stability of carbon-based devices. As a further advantage, carbon is not prone to formation of metal halides as many metal electrodes used in the PSC field do, leading to high resistivity and iodide deficiency in the perovskite layer. Moreover, carbon has not been demonstrated to penetrate the semiconductor material, while it is well known in the PSC community that Au diffuses through the perovskite layer. To further strengthen the solidity of carbon-based back electrodes in HTM-free devices, in situ analytical studies upon aging, highlighting the intrinsic stability of cell components, are required.

6 Conclusions

This review has summarized the most significant outcomes on PSC fabrication, characterization and scale up when carbon-based back electrodes are used. These platforms, prepared with graphite, carbon black, graphene or CNTs, exhibited significantly high stability, especially due to the high water resistance property of carbon. Research groups have proposed several processing routes, from scalable screen-printing approaches to inkjet printing, slot-die coating, and electrodeposition. The main trend in the literature scenario highlights that the PCE values of carbon electrode-based PSCs are lower than those of their gold-based counterparts, and the main reason lies in the poor contact at the perovskite/electrode interface; as secondary issues, resistance of carbon electrodes and charge recombination phenomena also contribute to lower $J_{sc}$ and FF values.

As a solution, current trends highlighted in this review show that interfacial engineering of carbon-based PSCs is becoming a fruitful research field, where scientists are working to decrease the heterogeneity in (opto)electronic properties of perovskites and carbon films and passivating their defects, thus boosting charge transfer and reducing recombination.

A further challenge in the long (but in rapid evolution) path towards gold-free PSCs will be the investigation behind the
markedly improved stability of carbon-based devices. Indeed, even though many groups have carried out long-term stability experiments, not much is understood on the relationship between the singular aging of internal interfaces and components, and to what extent they affect the overall device stability against different external stress sources.

**Conflicts of interest**

There are no conflicts to declare.

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