Conductive Gels: Properties and Applications of Nanoelectronics

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Abstract
Conductive gels are a special class of soft materials. They harness the 3D micro/nanostructures of gels with the electrical and optical properties of semiconductors, producing excellent novel attributes like the formation of an intricate network of conducting micro/nanostructures that facilitates the easy movement of charge carriers. Conductive gels encompass interesting properties, like adhesion, porosity, swelling, and good mechanical properties compared to those of bulk conducting polymers. The porous structure of the gels allows the easy diffusion of ions and molecules and the swelling nature provides an effective interface between molecular chains and solution phases, whereas good mechanical properties enable their practical applications. Due to these excellent assets, conductive gels are promising candidates for applications like energy conversion and storage, sensors, medical and biodevices, actuators, superhydrophobic coatings, etc. Conductive gels offer promising applications, e.g., as soft sensors, energy storage, and wearable electronics. Hydrogels with ionic species have some potential in this area. However, they suffer from dehydration due to evaporation when exposed to the air which limits their applications and lifespan. In addition to conductive polymers and organic charge transfer complexes, there is another class of organic matter called “conductive gels” that are used in the organic nanoelectronic industry. The main features of this family of organic materials include controllable photoluminescence, use in photon upconversion technology, and storage of optical energy and its conversion into electricity. Various parameters change the electronic and optical behaviors of these materials, which can be changed by controlling some of the structural and chemical parameters of conductive gels, their electronic and optical behaviors depending on the applications. If the conjugated molecules with π bonds come together spontaneously, in a relative order to form non-covalent bonds, they form a gel-like structure that has photoluminescence properties. The reason for this is the possibility of excitation of highest occupied molecular orbital level electrons of these molecules due to collision of landing photons and their transfer to the lowest unoccupied molecular orbital level. This property can be used in various nanoelectronic applications such as field-effect organic transistors, organic solar cells, and sensors to detect explosives. In this paper, the general introduction of conductive or conjugated gels with π bonds is discussed and some of the physical issues surrounding electron excitation due to incident radiation and the mobility of charge carriers, the position, and role of conductive gels in each of these applications are discussed.

Keywords: Polymer, Π bond, Mobility, Molecular, Receptor molecule, Photoluminescence, Conductive gel, Field-effect transistor, Solar cells, Sensor

Introduction
The organs of some marine animals have structures rich in water and organic matter. For example, mermaid tissues are composed of 1% organic matter in 96% water and are a good example of functional hydrogels.
Chemists and materials engineers have long sought to mimic nature and make materials with the function of natural hydrogels. “Gels” are examples of these materials that can trap very large volumes of solutions in their three-dimensional lattice-like structure. These semisolid viscoelastic materials are used today in various parts of human life, including food and cosmetics. The process of gelation or synthesis of hydrogels is accelerated by the formation of non-covalent bonds (i.e., hydrogen bonds, ionic bonds, π bonds, and van der Waals bonds in low molecular weight organic molecules) [1–3]. “Conjugated polymers with π-bonds” are another class of materials that are widely used in optoelectronic applications. The physical foundations, electronic properties, and nanoelectronic applications of this family of materials have been studied in detail in other articles on the Nano Education site. As mentioned, conductive polymers are often used as main components in organic electronics such as light-emitting diodes (LEDs), organic field-effect transistors (OFETs), and organic solar cells. Organic solar cells (OSCs) are used. The performance and efficiency of these devices are controlled by the degree of crystallinity of these materials in the process of making thin layers. Controlling the morphology and crystallinity of conductive polymers along with limited access to high purity in the synthesis process and low solubility of conjugated polymers in aqueous solutions is one of the problems and challenges facing the organic electronics industry. These challenges often lead to the formation of undesirable arrays of organic and non-uniform molecules of chemical composition and crystal structure. To solve these problems, “supramolecular polymers made of conjugated molecules with π bonds” have been developed [4–6]. The formation of these large molecule polymers under certain conditions leads to the formation of conductive or conjugated gels with π bonds (π gels). In fact, the molecular structure of conductive gels consists of regular arrays of conjugated molecules of various shapes and dimensions [7–11]. Conductive gels, due to the dynamic nature of molecular arrays, have the ability to regulate some electronic properties such as photoluminescence, mobility of charge carriers, and electrical conductivity. In this paper, recent developments in the field of conductive gels and their applications in the field of nanoelectronics will be reviewed [12–17]. Many conjugated organic materials with π bonds have a wide barrier band and exhibit semiconductor behaviors when doped with appropriate materials [18–22]. Tetrathiafulvalene (TTF) and its derivatives are an exception to this rule and, if properly doped, show very high electrical conductivity [23–26]. In recent years, some researchers have sought to make conductive gels based on TTF in order to obtain a very high electrical conductivity [27–30]. The chemical structure of this substance is shown in Fig. 1 (Structure 1) [31]. The current curve according to the voltage of this material, if not doped with a suitable additive, is in the form of a yellow horizontal line, which is shown in Fig. 1b. This curve shows that with any voltage difference, the current flowing through the material is very small. On the other hand, by doping compound I2 at room temperature, the electrical conductivity of these materials is improved. A new solution to increase the conductivity of these materials is to bind gold nanoparticles to the ligands of TTF molecules. This is shown in Fig. 1 as structures 16 and 17. As can be seen, the addition of gold nanoparticles leads to metallic behaviors in TTF-based conductive gels.

![Fig. 1 a Molecular structure of tetrathiafulvalene (TTF)-based conductive gels; b current–voltage diagrams obtained by atomic force microscopy (AFM) for conductive gels in the presence and absence of structures 2 and 3 and doping conditions with I2 [37]](image)
general, in order to be able to use conductive gels in the manufacture of nanoelectronic devices, the method of doping or adding conductive nanoparticles is very helpful [32–36]. As another practical example, large molecules of conductive gels can be produced as nanofibers after doping and their conductivity behaviors in different molecular arrangements can be investigated [37–42]. For example, Fig. 2 shows two different molecular structures in oligothiophene-based conductive gels that are produced as continuous, cohesive fibers. The current–voltage diagram of these nanofibers shows that the molecular structure and chemical composition of large molecules and their self-arrangement strongly affect the electrical resistance of these materials. Therefore, to achieve the highest electrical conductivity in conductive gels, control of the spatial arrangement of molecules is very important [43–45]. Table 1 shows the summary of properties and nanoelectronic applications of conductive gels.

Conductive polymers have been researched over the past few decades owing to their unique ability to provide tunable electrical conductivity and flexibility during processing [46–48]. The conductivity of conductive polymers depends on the molecular structures of the constituent materials, the level of doping, and the ordering of molecular packing. With the rapid emergence of nanoscience and nanotechnology, it is anticipated that conductive polymers with well-defined nanostructures can translate the properties of their bulk forms and exhibit unusual chemical/physical properties because of the confined dimensions of the nanomaterials [49–54].

Conductive polymers with various nanostructures including 0D nanoparticles, [55] 1D nanofibers [56, 57], and 2D nanosheets [58] have been developed and applied in a range of technological areas, such as sensors, electronics, and energy storage and conversion devices. However, the electrical properties of these nanostructured conductive polymers could be weakened by structural defects induced by inhomogeneous aggregation, severe restacking, and poor contacts during processing and assembly [59]. The development of nanostructured conductive polymers with tunable microstructures and controllable chemical/physical properties still remains a challenge [60–62].

Inspired by the chemical/structural features and synthetic approaches of natural gels [64], conductive polymer gels (CPGs) with 3D networked structures were recently developed by cross-linking the conjugated polymer chains using molecules with multiple functional groups [65]. CPGs show the unique features of gel materials: they are dilute cross-linked systems and exhibit no flow when in a steady state. This monolithic structure inherits the conductive properties of the conjugated polymeric chains and generates highly tunable chemical/physical properties derived from its cross-linked network [2, 48, 50–54, 60–63, 66–70], including flexibility, stretchability, ionic conductivity, electrochemical activity,
| Conductive components/nanowires | Materials | Optical property | Application | References |
|--------------------------------|-----------|-----------------|-------------|------------|
| Metallic nanoparticles/nanowires | Ag/Au/Cu nanoparticles | Transparent or semitransparent | Biosensor drug delivery, tissue engineering | [85-87, 98] |
| Carbon-based materials | CNTs | Mostly black or semitransparent | Strain sensor, tissue engineering, biosensor, supercapacitor, drug delivery | [99-116] |
| Carbon-based materials | GO/rGO/graphene | Mostly black or semitransparent | Strain sensor, tissue engineering, biosensor, supercapacitor | [99-113] |
| Conducting polymers | Polyaniline | Mostly black and rarely transparent | Tissue engineering, biosensor, drug delivery, supercapacitor, bioelectrode, strain sensor | [114-139] |
| Conducting polymers | Polypyrrole | Mostly black and rarely transparent | Tissue engineering, biosensor, drug delivery, supercapacitor, bioelectrode, strain sensor | [114-135, 140-143] |
| Conducting polymers | PEDOT:PSS | Mostly black and rarely transparent | Tissue engineering, biosensor, drug delivery, supercapacitor, bioelectrode, strain sensor | [114-135, 144-147] |
| Hybrid | Pt/Ag/(GO + rGO)/SWCNTs + polyaniline | Usually black or semitransparent | Biosensor, battery, fuel cell, supercapacitor, tissue engineering, pressure sensor, strain sensor | [148-163] |
| Hybrid | Au/(Fe + Co)/Fe3O4/CNTs + polypyrrole | Usually black or semitransparent | Biosensor, battery, fuel cell, supercapacitor, tissue engineering, pressure sensor, strain sensor | [148-160, 164-167] |
| Hybrid | Graphene/GO + Ni/Ni(OH)2 | Usually black or semitransparent | Biosensor, battery, fuel cell, supercapacitor, tissue engineering, pressure sensor, strain sensor | [148-160, 168-172] |
| Hybrid | GO + PEDOT:PSS | Usually black or semitransparent | Biosensor, battery, fuel cell, supercapacitor, tissue engineering, pressure sensor, strain sensor | [148-160] |
| Hybrid | GO + CuS + carbon dot | Usually black or semitransparent | Biosensor, battery, fuel cell, supercapacitor, tissue engineering, pressure sensor, strain sensor | [148-160, 173-175] |
| Acids | H2SO4/H3PO4 | Transparent | Supercapacitor | [176-187] |
| Metallic salts | LiCl/Na+ /Ca2+ | Transparent or semitransparent | Supercapacitor, solar cell, nanogenerator, actuator, electronic eel/fish, displays, touch panels, pressure sensor, strain sensor, biosensor | [180-182, 188-221] |
| Metallic salts | Al3+/Fe3+/Tb3+/K+ | Transparent or semitransparent | Supercapacitor, solar cell, nanogenerator, actuator, electronic eel/fish, displays, touch panels, pressure sensor, strain sensor, biosensor | [183-185, 188-221] |
| Metallic salts | Na2S + sulfur + NaOH | Transparent or semitransparent | Supercapacitor, solar cell, nanogenerator, actuator, electronic eel/fish, displays, touch panels, pressure sensor, strain sensor, biosensor | [186-221, 227-230] |
| Ionic liquids | Ionic liquids | 1-Ethyl-3-methylimidazolium chloride | Semitransparent | [222] |
| Electron conductive | Na++ Au nanoparticles | Usually black or opaque | Supercapacitor, biosensor, pressure sensor, battery strain sensor, tissue engineering | [223] |
and so forth. Meanwhile, CPGs have emerged as a unique material platform to develop functional materials by building interpenetrating structures with a secondary polymeric network, loading specific nanoparticles, or serving as a precursor for graphitic carbon frameworks [71].

Conductive hydrogels have drawn significant attention in the field of stretchable/wearable sensors due to their intrinsic stretchability, tunable conductivity, biocompatibility, multi-stimuli sensitivity, and self-healing ability. Recent advancements in hydrogel- and organohydrogel-based sensors, including a novel sensing mechanism, outstanding performance, and broad application scenarios, suggest the great potential of hydrogels for stretchable electronics. However, a systematic summary of hydrogel- and organohydrogel-based sensors in terms of their working principles, unique properties, and promising applications is still lacking. In this spotlight, we present recent advances in hydrogel- and organohydrogel-based stretchable sensors with four main sections: improved stability of hydrogels, fabrication and characterization of organohydrogel, working principles, and performance of different types of sensors. We particularly highlight our recent work on ultrastretchable and high-performance strain, temperature, humidity, and gas sensors based on polyacrylamide/carrageenan, double-network hydrogel and ethylene glycol/glycerol modified organohydrogels obtained via a facile solvent displacement strategy. The organohydrogels display higher stability (drying and freezing tolerances) and sensing performances than corresponding hydrogels. The sensing mechanisms, key factors influencing the performance, and application prospects of these sensors are revealed. Particularly, we find that the hindering effect of polymer networks on the ionic transport is one of the key mechanisms applicable for all four of these kinds of sensors [12].

**Electronic Properties of Conductive Gels**

Chromophore is the part of a molecule that causes color in it. Chromaticity occurs when matter can absorb a certain wavelength of visible light and pass or reflect the rest. A chromophore is a region of a molecule where the energy balance difference between two molecular orbitals is within the energy range of part of the incident beam spectrum [56]. A certain wavelength of light due to a collision with a molecule can be absorbed by its electrons and excite them from the ground state to the excited state. What is very important in conductive gels and affects the efficiency of absorption of light energy by conductive gels is the arrangement and spatial arrangement of chromophores giving or receiving electron with optimal distance and orientation. To date, a wide range of conductive gels have been developed as high-performance scaffolds for energy transfer or conversion. In each case, they have seized it, despite obstacles we can scarcely imagine. In other words, it is generally preferred that the chromophores giving and receiving conductive gels can be arranged on their own and that the electron excitation properties of these materials due to light beam can be optimized by controlling the molecular self-arranging property and increasing the energy conversion efficiency [57]. In other words, by controlling the amount of energy transfer between the donor and electron acceptor molecules in the three-dimensional structure and network like conductive gels, the optical energy absorption or optical behavior of the material can be adjusted and used to make optical emitting devices (LEDs). In order to be able to use conductive gels in the nanoelectronics and organic electronics industries, it is necessary for the three-dimensional structure of these materials to be able to transfer excited electrons at a macroscopic distance. The emergence of this feature requires regular arrangement of electron donor and receiver centers in different directions. In practice, the self-arrangement of donor and receptor molecules does this. Figure 3 shows a diagram of the molecular structure of multi-chromophore conductive gels arranged in self-assembled nanofibers. The core of each nanofiber is composed of an elongated chain of conjugated molecules with π bonds that, with their regular arrangement, lead to the formation of a continuous structure. The gel changes color when exposed to ultraviolet light and its macroscopic appearance changes from dark orange to yellow. The reason for this discoloration is the excitation of electron donor and electron acceptor chromophores by the collision of ultraviolet light and their movement in the three-dimensional structure of the gel. Today, photon upconversion is one of the most important applications of electron excitation in conductive gels. In this phenomenon, several weak and low-energy photons are absorbed by the material due to electron excitation, and in return, a higher-energy photon is emitted by the material. It has been observed that in conductive gels, the electron donor

| Conductive components and ions | Materials | Optical property | Application | References |
|-------------------------------|-----------|----------------|-------------|------------|
| HCl/HClO4/(Na+ + SWCNTs) + polyaniline | Supercapacitor, biosensor, pressure sensor, battery strain sensor, tissue engineering | [224–226] |
centers can be combined in a polymer field containing electron acceptor centers; with a special arrangement, the photon upconversion property in the gel can be achieved. Take Fig. 4, for example. In this form, the conducting gel of 9,10-diphenylanthracene as the electron receptor is doped by Pt (II) octaethylporphyrin (PtOEP)
Fig. 4 Molecular structures: a conductive gels and electron donor centers. b a schematic of the specific spatial arrangement of the two-phase structure of the conductive gel–electron donor centers after doping, the photon upconversion phenomenon is shown schematically; c the photoluminescence spectrum of a conductive gel doped with electron donor centers; and the emission of light output is done in the wavelength range of 400 to 475 nm. In this study, 9,10-diphenylanthracene acted as conductive gels (electron receptor centers) and PtOEP acted as electron donor centers [3]
as the electron donor. As can be seen, the collision of the incident beam leads to the excitation of electrons in the donor centers and the transfer of charge carriers to the recipient centers. Finally, matter emits a higher energy beam for several electrons excited at a lower energy [58]. The photoluminescence spectrum diagram of this conductive gel shows that the output beam has a relatively wide range of wavelengths in the range of 400 to 475 nm.

**Electron Transfer Due to Optical Excitation**

In addition to controllable photoluminescence behavior, one of the main approaches to the development of conjugated gels with π bonds is the conversion of optical energy into electrical energy by electron excitation. In other words, scientists are looking for conductive gels to emit their excited electrons through an electronic circuit, instead of emitting light or exhibiting photon upconversion due to electron stimulation after exposure to the incoming beam. Collected and stored. Observations show that conductive gels are a good option for converting solar energy into electricity, because the electron donor and receiver centers in the three-dimensional structure of these materials are arranged independently and in a controllable arrangement, and the possibility of electron transfer at macroscopic distances [59, 72]. There is: One method of producing conductive gel-based systems to convert solar energy into electricity is to functionalize conductive gels with electron donor–receiver units. What is very important in these gel structures is to prevent the recombination of electrons, which is also done by choosing the right chromophores for giving and receiving electrically charged carriers. Recently, efforts have been made to design and synthesize gels with optimal conversion and storage properties of electric current through visible light absorption. One of these systems is a gel-like structure, which is schematically shown in Fig. 5. In this gel, naphthalimide as electron donor and perylene monoimide as electron receptor are dissolved in hexane/dichloromethane solution and then functionalized with α,β-dihydroxypropyl chains. The functionalization of these compounds leads to the self-assembly of these molecules and the formation of regular molecular structures within the solution at low concentrations (Fig. 5a). As the concentration of the solution increases, non-covalent bonds predominate and the cell turns into a gel (Fig. 5b). It has been observed that at concentrations above a critical limit, a nanostructured gel is formed in which the charge carriers generated by the excitation of visible light are transferred from the naphthalimide unit to the perylene monoimide unit, and thus, the electric current in it flows inside the network structure like a conductive gel. Also, if indole is used as the electron donor, the electron transfer process will be accelerated and the electron path will be from naphthalimide to indole and finally to perylene monoimide, respectively. It should be noted that in biphasic conducting gels, the rate of electron–hole recombination is much lower than the rate of electron production due to light excitation, and the reason for this is the self-assembly arrangement of the large molecules that make up the gel. Such a stable transfer
from charge carriers can be used in water splitting and artificial photosynthesis.

**Mobility of Electric Charge Carriers**

As a general rule, the overall efficiency and performance of organic electronics, such as FETs and heterogeneous solar cells, depend on the mobility of the active charge carriers in the system. Self-assembly materials obtained from conjugated gels with \( \pi \) bonds are good options for making organic electronic devices, because the regular molecular arrangement in the gel structure can contribute to the greater mobility of the electric charge carriers in these structures [73, 74]. However, it is very difficult to measure the mobility of charge carriers in gels without destroying their molecular arrangement. Conductive gels with thienylenevinylene base are among the gels in which electric charge carriers have high mobility. Figure 6a shows two examples of these molecular structures (structures 11a and 11b). The large molecules in these gels self-assemble into continuous fibers. The absorption spectra of these materials show that the mobility of the electrons in the mentioned gels is due to the very regular molecular self-arrangement of these materials and the overlap of the band structure of the electron donor and receiver units. Also, if \( \text{N,N-bis(2,5-di-tert-butylphenyl)3,4,9,10-pyreneledicarboximide (PDI)} \) is used in these gel structures, the mobility of the electrons will increase (Fig. 6b). In fact, it helps to gel more gel on a molecular scale, thereby increasing the mobility of energy carriers. On the other hand, the electron mobility in gel structures is relatively low compared to xerogel structures of the same material (Fig. 6c). It should be noted that if the gel dries at relatively low temperatures, it becomes a very porous solid product called xerogel.

**Sensors for Explosives**

Photoluminescence is one of the most attractive properties of conductive gels. This is done by absorbing colliding photons with organic matter, stimulating electrons at its HOMO level, and transferring them to the LUMO level. Studies show that if the macromolecules of a conductive gel (which are rich in excited electrons) are exposed to electron-deficient analyses, the gel-stimulated electrons are readily transferred to the analytic band structure and photoluminescence properties of these materials they destroy [71, 75]. This interesting phenomenon has recently become the basis for the identification of nitro-aromatic explosives using conductive gels. For example, the presence of TNT explosives in very small amounts (in the range) can be detected using pentajuoroarene functional groups. This is shown in Fig. 7. When these functional groups form a gel-like structure,
the macromolecules assume a relatively orderly arrangement and are expected to exhibit photoluminescence properties. Now, if this gel is placed on a special paper in a thin layer, in the presence of sunlight, it will show a special color. If some of the TNT explosives come in contact with the surface of the paper, the photoluminescence property of the gel is rapidly lost, and the rate of drop of this property is directly related to the amount of explosive. The higher the concentration of the explosive, the lower the intensity of the light emitted by the gel, and the presence of an explosive can be detected by the color change. It should be noted that the reason for the suitability of conductive gels for the detection of TNT explosives is the efficient absorption of these materials into the self-assembled structure of the gel molecules and energy absorption by it.

Applications
Recently, conductive gels have been used in the manufacture of many nanoelectronic devices due to their controllable conductivity. The most important of these applications are organic field-effect transistors (OFETs), organic solar cells, and organic sensors. In the following, the role of these materials in the mentioned applications is described. It should be noted that one of the main applications of conductive gels is the production of hydrogen using water splitting, but since the present article deals with the nanoelectronic applications of conductive gels, this issue will not be discussed [76, 77].

Field-Effect Organic Transistors (OFETs)
Organic semiconductors are preferred over silicon-based semiconductors in many applications, including integrated circuits, sensors, and electronic chips, due to their lightweight and flexibility. The relatively regular arrangement of the large molecules in these materials due to the processing of the solution phase can provide continuous conduction paths for the electrons and cavities (percolation network) and increase the efficiency and electrical efficiency of the devices. Since conductive gels are prepared from solution phase and in which large molecule chromophores have the ability of molecular self-assembly, these materials are a very good option for making active materials in field-effect organic transistors (OFETs) [78, 79]. Before discussing the application of conductive gels in OFETs transistors, it is advisable to discuss the general structure of field-effect capacitors. The most common structure of field-effect transistors often consists of a n- and p-type semiconductor junction with two layers of oxide insulating material (such as silicon oxide) and a conductive layer (non-crystalline metal or silicon). Figure 8 shows a schematic of this structure. As can be seen, these transistors consist of three main parts: The main body is a p-type semiconductor that holds two separate n-type semiconductor parts at a certain distance from each other. One type n region is called the source terminal, and the other is called the drain terminal. The area between the source and discharge terminals is covered by a thin, insulating layer of metal oxide, and the conductive layer is placed on top of the oxide layer. The whole area between the two terminals is called the gate terminal. Due to the nature of n-type semiconductors, the concentration of electrons is higher in the areas near the source terminal and discharge, but there is no current between the two regions [80, 81]. Now, if a positive voltage is applied to the gate terminal and the source terminal is connected to ground and the discharge terminal is connected to a positive potential, the negative electrons tend...
to accumulate in the gate region due to the positive voltage of the gate terminal. If we increase the gate voltage and the positive voltage applied to the discharge source sufficiently, the negative charges accumulated under the metal oxide layer will tend to move toward the gate terminal and the discharge terminal. By doing this, applying a gate voltage and a positive voltage to the discharge terminal causes the electron current to flow from the source terminal to the discharge terminal. In other words, no current of electrons will be established between the two terminals until the gate voltage reaches a critical level. Conversely, if the gate voltage increases exponentially, a path of positive charges in the gate region is created and the intensity of the electron current flux increases. Therefore, the gate terminal acts as a valve for the electron flow tube. It should be noted that if the main body of the field-effect transistor is made of p-type semiconductor, it is called P-FETs and if it is n-type, it is called N-FETs. The same mechanism applies to both types of transistors. In recent years, many attempts have been made to use new materials as the main body or substrate of FETs transistors on which the source and discharge terminals as well as the gate are mounted. Conductive gels are one of these emerging materials. For example, Hong et al. used single nanofibers made of conjugated gels with π bonds as the main body of nanotransistors (Fig. 9). Due to its one-dimensional self-arrangement and chemical structure, this fiber behaves similarly to the p-type semiconductor. The mobility of cavities in this fiber is in the range of 0.48–0.1 cm/s [82]. As shown in Fig. 9, an increase in gate voltage leads to an increase in the current flow from the source terminal to the discharge terminal. In some studies, a thin layer of conductive gels has been used as the main body of OFETs transistors. Figure 5 shows an example of these structures. As can be seen, a field-effect transistor with a downwards gate terminal can be produced using a thin layer of conductive gel with a 23 structure. The important point is that the large molecules of this conductive gel have the property of molecular self-assembly, and this leads to high mobility of charge carriers in this thinner layer. The structure 23 behaves similarly to the n-type semiconductor, and the majority charge carriers are electrons. Also, the current–voltage diagrams in Fig. 10 clearly show that as the gate voltage increases and the voltage difference between the source terminal and the discharge increases, the current passing through the transistor increases exponentially. However, the effect of gate voltage on the current density is greater than the discharge terminal voltage.

Organic Solar Cells

Organic solar cells (OSCs) have relatively low efficiencies in converting solar energy into electrical energy, but very low weight, low cost, and flexibility of organic matter relative to inorganic semiconductors are the focus of energy industries. Has attracted. Has attracted Conductive gels, on the other hand, have been considered in comparison with other semiconductor or non-conductive organic materials for several important reasons for making OSCs [83, 84]:

(a) Ease of processing, coating, production of thin films, and production of nanometer fibers.
(b) The possibility of making heterogeneous connections from chromophores giving or receiving large molecules in meso- or nanodimensions.
(c) Possibility of easy self-assembly of large molecules and engineering of three-dimensional arrangement of electron donor and receiver chromophores to achieve high energy conversion efficiency.

In this section, a sample of organic solar cells based on conductive gels is reviewed. But before entering into this discussion, it is better to introduce the general structure of organic solar cells. Figure 10 schematically shows the layer structure of an organic solar cell. As can be seen,
these cells consist of several main parts: (1) glass, (2) a transparent conductive oxide layer (such as indium-tin oxide or ITO), (3) a conductive transparent polymer (such as PEDOT: PSS), (4) an active layer or absorber, (5) a bonding layer, and (6) a conductive metal layer. In general, the conductive metal layer and the transparent metal oxide layer act as “contact points” to connect to the external circuit. A thin layer of transparent conductive polymer such as PEDOT: PSS is usually used as the cavity conductor material directly above the ITO electrode. What plays the most effective role in building an organic solar cell is the “active layer.” Solar cells are divided into several categories in terms of number of layers and electronic architecture (Fig. 10):

- Organic monolayer solar cells

These solar cells are the simplest type of photovoltaic device in terms of energy storage conversion mechanism. In the structure of these devices, an organic semiconductor is placed between two thin layers of conductive metal. One of these metal layers, such as indium-tin oxide (ITO), has a very high working function, and the other layer, such as aluminum or magnesium, has a relatively lower working function (Fig. 10a). The mechanism of storage of electric charge is such that due to the collision of the landing photon, a large number of excitations (electron–hole pair) are formed in the band structure of the organic semiconductor layer. These electrons begin to move due to the electric field created between the two metal layers due to the difference in their working function [84]. They flow toward the positive metal electrode and the cavity toward the negative electrode. The band structure of this type of solar cell and the path of electrons and holes are shown in Fig. 10a.

- Organic bilayer solar cells

This type of solar cell is made of two separate organic layers with completely different electronegativity...
between conductive metal layers. This difference in electronegativity leads to the creation of an electrostatic field between two metal layers. The semiconductor layer that has a high electronegativity tends to absorb more electrons and the other organic layer tends to give electrons. Therefore, when a landing photon strikes these two semiconductor layers, it leads to the formation of an electron–hole pair at the LUMO level of the band structure of the two materials, and the electrons and holes due to the electrostatic field created between the two layers begin to separate and move. They do it in the opposite direction. In other words, the
layer with high electronegativity acts as the electron acceptor and repulse of the cavity, and the layer with lower electronegativity acts as the electron giver and the absorber of the cavity. In this way, the excitations are separated from each other and placed in holes in the outer circuit by holes and electrons through metal layers. The band structure of this group of solar cells and the path of electron–hole motion are shown in Fig. 11.

- Organic solar cells with mass heterogeneous connections

In organic solar cells, two metal layers (mostly aluminum and ITO) are used as the two points of contact for the external circuit. Also, conductive gels are often used as the organic layer between these two conductive metal layers. This layer is also called the active layer. Recently, some researchers have used organic gels based on conductive polymers as the active layer of organic solar cells [85]. Figure 12 shows the chemical structure of two types of polymer molecules (structures 26a and 26b). These two molecules are similar in molecular structure, but one of them has a CN functional group. The use of these materials as active layers in organic solar cells has been shown to improve the energy storage efficiency of these systems by 1.75%. This efficiency is also increased if heat annealing is used to convert the solution containing these molecules into gel structures in which molecules 26a and 26b are self-arranged. This shows how much the gelling of solutions containing conjugated polymers with π bonds can affect the self-assembly of molecules and energy storage efficiency. Figure 12 shows the increase in efficiency of generated solar cells in the form of stored current density graphs. Molecular structure 29 in Fig. 13 is another example of the application of conductive gels in the fabrication of solar organic cells as the active layer. These large molecules can affect energy conversion efficiency, depending on the number of repetitive units and the degree to which they are self-assembled. As can be seen in Fig. 13, gelling and self-assembly of large molecules significantly increase the slope of changes in the energy conversion efficiency diagram. Efforts to replace
various types of conductive gels in the main body of organic solar cells continue today.

**Conclusion**

a. In the organic nanoelectronics industry, in addition to organic charge transfer complexes and conductive polymers, conjugated gels with π bonds are also used. These materials are large molecules that, similar to organic charge transfer complexes, have electron donor and receptor centers, and, like conductive polymers, are composed of conjugated molecules with π bonds. These large molecular units in gel structures are placed next to each other with non-covalent bonds, increasing the viscosity of the system. One of the main features of conductive gels is the ability to change the optical behavior of these materials by controlling chromophores, viscosity, arrangement of molecular arrays, and doping with other molecules. These materials can also be used in photon upconversion technology. Conductive gels are now used in the manufacture of solar cells to convert solar energy into electricity.

b. In the present paper, the nanoelectronic applications of conductive gels were reviewed. The most important of these applications are sensors for detecting explosives, thin-film organic solar cells, and field-effect nanotransistors. As mentioned, conductive gels can be used as non-intrinsic semiconductors due to the self-arrangement of large molecules and can be used as the active layer in organic solar cells or the main body of semiconductors in field-effect transistors. The basis of these materials is the excitation of HOMO level electrons and their transfer to the LUMO level. The efficiency and performance of this group of materials are strongly related to the kinetics of the process of excitation and transfer of charge carriers due to the collision of landing photons. Conductivity can significantly increase the conductivity of conductive gels. The most important method is to dope or add a specific functional group to the molecular arrays of these materials. More details of the mentioned applications are discussed in the present article.

**Abbreviations**

HOMO: Highest occupied molecular orbital; LUMO: Lowest unoccupied molecular orbital; TTF: Tetrathiafulvalene.

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**Author contributions**

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Ethics approval and consent to participate
I wrote to you in regard to your question about naming some people in my article, I must point out that in some cases, help was sought from people and it was necessary to mention the names of these people in order to maintain professional ethics in research issues.

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