Graphene-Based Strategies in the Diagnosis of Viral Diseases

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The occurrence of new pandemic viruses, such as SARS-CoV-1 (2003), MERS-CoV (2012) and SARS-CoV-2 (2019) has indicated an urgent need for diagnostic tools able to reliably identify infected individuals and to determine if the infection is in the acute phase. Although nanotechnology based on graphene has been poorly applied for the rapid diagnosis of viral diseases, the extraordinary properties of graphene have been recently exploited for the diagnosis of COVID-19. Novel graphene-based field-effect transistor (GFET) biosensors were developed for the quantitative detection of viral RNA and viral spike protein. The fabrication of COVID-19 FET sensor for spike protein recognition is based on the integration of the SARS-CoV-2 spike antibody with graphene, whereas the GFET sensor for viral RNA recognition exploited the CRISPR/Cas biotechnology.

Direct methods, exploiting graphene nanotechnology, for the rapid virus detection, have been only marginally investigated in the past, and no critical discussion has been reported in successive literature reviews.[1][2]

This attitude was unchanging even during SARS-CoV-1 emergency that was responsible for the 2003 severe acute respiratory syndrome (SARS) infection in Asia, causing about 8000 cases and 774 deaths, also during the Middle East respiratory distress syndrome (MERS) of 2013, which affected Saudi Arabia causing close to 858 deaths.[3][4][5][6] Advances in nanotechnology have begun to play an important role in viral detection, to improve the detection limit, operational simplicity of viral diagnostics.[2]

A coplanar-gate graphene field-effect transistors (GFETs)[8] have been proposed for the detection of HIV-1 (human immunodeficiency virus 1) and MLV (murine leukemia virus) viruses using antibodies of vesicular stomatitis Indiana virus (VSV) as biorecognition element. VSV antibodies are immobilized on the G layer using 1-pyrenebutanoic acid succinimidyl ester (PASE). PASE binds G by π-π interactions, anchoring the antibody’s primary amine groups by the opposite succinimidyl group. The formation of the virus-antibody complex leads to a downward shift of the Dirac point voltage, regardless of the types of detected viruses. The proposed platform has worked in a wide range of concentrations (from 47.8 aM to 10.55 nM), but the lack of virus specificity appears the main limitation of this strategy.

An surface plasmon resonance (SPR) sensor based on an polyamidoamine-functionalized rGO(composite, with monoclonal antibodies immobilized on self-assembled dithiobis (succinimidyl undecanoate, DSU) for the detection/quantification of Dengue virus (DENV), has been recently described.[2]

The specificity and the sensitivity of the sensor have been achieved by anchoring a stable biorecognition element (antibodies (IgM) against Dengue type 2 envelope proteins) on the gold surface of the sensor. The specific binding of antibody-DENV 2 E-protein allows a significant change in the angle of the reflectivity minimum that is correlated to Dengue virus detection. The proposed sensor has shown a sensitive and selective response towards DENV 2 E-proteins compared to DENV 1 E-proteins and ZIKV (Zika virus) E-proteins. Although no G materials have been integrated into the above-described sensor[2], the criteria used for its fabrication were included in this review since the strategy could be extended to other viruses, and the performance of SPR noble metal could be improved in the presence of G.[9]

Differently from the past, the current sanitary pandemic emergency caused by the new type of coronavirus (SARS-CoV-2) is characterized by global effort to identify biomarkers that predict the severity of COVID-19 patients and to develop diagnostic tools for the rapid detection of SARS-CoV-2 infection[10].

Currently, nucleic acid testing on respiratory specimens is the reference gold standard method for the
diagnosis of COVID-19 infected patients. The test requires a series of laboratory procedures: (a) viral RNA extraction; (b) addition to a master mix containing nuclease-free water, reverse primers, a fluorophore-quencher probe, and a reaction mix (i.e., polymerase, reverse transcriptase, magnesium, nucleotides, and additives); (c) loading of extracted RNA/master mix into a PCR thermocycler; (d) several cycles at settled temperature. During the RT-PCR cycles, the cleavage of the fluorophore-quencher probe generates a fluorescent signal detected and recorded in real-time.

RT-PCR uses respiratory samples to genetically detect SARS-CoV-2; some data have suggested that 20–34% of COVID-19 patients resulted negative in the test despite being infected. This variance in the sensitivity could be mainly attributed to low viral load (i.e., patients tested in the early stage of the viral disease). Other RT-PCR issues include the time consuming and expensive analysis and the technical expertise in carrying out the test.

Other technologies, such as point-of-care technologies and serologic immunoassays, are rapidly emerging to address these deficiencies.

Analytic methods to assess prior infection and immunity to SARS-CoV-2 by antibody identification are essential for epidemiologic studies, although sensibility and specificity of the tests currently available in the market remain undefined. Cross-reactivity of antibody to non-SARS-CoV-2 coronavirus proteins is the main issue of these serologic tests. The development of an antigen detection test could take advantage of progress in the production of monoclonal antibodies against the nucleocapsid protein of SARS-CoV-2. The global effort to increase SARS-CoV-2 testing capacity takes advantage of the most recent advances in chemistry, molecular biology, genome technology, and nanotechnology. Several projects are ongoing in this direction, and some results are already reported in the literature.

The detection of SARS-CoV-2 in respiratory samples has been achieved by LSPR biosensor, combining the photothermal effect and plasmonic sensing transduction for SARS-CoV-2 viral nucleic acid.

A field-effect transistor (FET)-based biosensing device for detecting SARS-CoV-2 spike protein (S) in clinical samples was reported by Seo et al. Antibodies against S protein were anchored to the graphene sheet (external coating of FET) by 1-pyrenebutanoic acid succinimidyl ester (PBASE, Figure 1).

Figure 1. Coronavirus disease 2019 (COVID-19) field-effect transistor (FET)-sensor. Graphene is selected as sensing material and is decorated with the SARS-CoV-2 spike antibody using 1-pyrenebutanoic acid succinimidyl ester (PBASE) as interfacing molecule and probe linker. Reprinted with permission from reference, Copyright © 2020, American Chemical Society. Further permissions related to the material excerpted should be directed to the ACS.

The performance of the sensor is determined using antigen protein, cultured virus, and nasopharyngeal swab specimens from COVID-19 patients. The device could detect S protein at concentrations of 1 fg/mL in PBS and 100 fg/mL in the clinical transport medium, and it could distinguish the SARS-CoV-2 antigen protein from those of MERS-CoV. The successful fabrication of a COVID-19 FET sensor based on the integration of the SARS-CoV-2 spike antibody with graphene suggests the key role of G for diagnostic scope.
Specifically, the functionalization of G with diverse functional molecules could be the key element to tailor its properties and to obtain advanced diagnostic tools for the SARS-CoV-2 diagnosis. Meanwhile, for the revision of this manuscript, some works dealing with sensors for COVID-19 diagnosis based on graphene are reported in the literature, and, although further researches are undoubtedly necessary, the leading role of G in the world’s fight against COVID-19 is clearly coming out.

In summary, the biomolecules till now used to target SARS-CoV-2 includes the viral RNA, the viral spike proteins, and the specific immunoglobulins produced by the host immune system. The biosensing community is actively working to improve portability, time, and cost of PCR-based SARS-CoV-2 detection, as well as to create manufacturable PCR-based microfluidic devices. Recently, also the gene-editing technology (CRISPR/Cas) has been developed to overcome the issues of PCR-based systems. Two different detection modes have been proposed in CRISPR technology, i.e., binding- or cleavage-based. The sensor is developed by immobilization on a graphene-based field-effect transistor (GFET) of Cas9 with a sgRNA, specific to the target sequence of SARS COV-2; the electrical signal originated by the binding of the target nucleic acid by the Cas9-sgRNA complex is recorded via a simple handheld device without amplification.

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