Abstract: Electrochemical sensors and biosensors have attracted considerable attention for the sensitive detection of a variety of biological and pharmaceutical compounds. Since the discovery of carbon-based nanomaterials, including carbon nanotubes, C₆₀ and graphene, they have garnered tremendous interest for their potential in the design of high-performance electrochemical sensor platforms due to their exceptional thermal, mechanical, electronic, and catalytic properties. Carbon nanomaterial-based electrochemical sensors have been employed for the detection of various analytes with rapid electron transfer kinetics. This feature article focuses on the recent design and use of carbon nanomaterials, primarily single-walled carbon nanotubes (SWCNTs), reduced graphene oxide (rGO), SWCNTs-rGO, Au nanoparticle-rGO nanocomposites, and buckypaper as sensing materials for the electrochemical detection of some representative biological and pharmaceutical compounds such as methylglyoxal, acetaminophen, valacyclovir, β-nicotinamide adenine dinucleotide hydrate (NADH), and glucose. Furthermore, the electrochemical performance of SWCNTs, rGO, and SWCNT-rGO for the detection of acetaminophen and valacyclovir was comparatively studied, revealing that SWCNT-rGO nanocomposites possess excellent electrocatalytic activity in comparison to individual SWCNT and rGO platforms. The sensitive, reliable and rapid analysis of critical disease biomarkers and globally emerging pharmaceutical compounds at carbon nanomaterials based electrochemical sensor platforms may enable an extensive range of applications in preemptive medical diagnostics.
Keywords: SWCNTs; graphene; buckypaper; methylglyoxal; NADH; glucose; acetaminophen; valacyclovir

1. Introduction

Owing to the advantages of instrumental simplicity, moderate cost, and portability, electroanalytical techniques provide a powerful sensing strategy, superior to various traditional analytical methods [1]. Assorted electroanalytical techniques have been employed for the development of electrochemical sensors and biosensors for the detection and quantification of myriad biomarker species, chemical compounds, and minerals in environmental and biological samples [2–4]. The development of active electrocatalysts plays a key role in the design of efficient, reliable, stable, and innovative sensing devices [5]. Numerous electrochemical sensor platforms have been developed for the detection and quantification of medically and pharmaceutically important compounds such as glucose, methylglyoxal, nicotinamide adenine dinucleotide (NADH), acetaminophen, and valacyclovir. Moreover, recent developments in the field of nanotechnology and materials science have paved the way for the synthesis of many new materials with desired morphologies and unique physicochemical properties [6].

Carbon-based nanomaterials such as carbon nanotubes, graphene, buckypaper, and nanohybrids among various other types of engineered nanomaterials have received enormous attention due to their promising sensor applications, which are most widely used in the electrochemical sensing of various compounds [7–9]. Carbon nanomaterials offer diverse advantages with their unique properties, such as a high surface-to-volume ratio, high electrical conductivity, chemical stability, biocompatibility, and robust mechanical strength [10–12]. Carbon possesses the capacity to hybridize into sp, sp² and sp³ configurations with narrow gaps between their 2s and 2p electron shells. These unique properties are responsible for enabling the design of versatile carbon-based nanomaterials for the sensitive detection of biological compounds [13–15]. Carbon nanotubes may be comprised of a single graphitic layer, or multiple coaxial layers, resulting in the formation of single-walled carbon nanotubes (SWCNTs) and multiple-walled carbon nanotubes (MWCNTs). SWNTs are seamlessly wrapped into cylindrical tubes, having diameters of between 0.4 nm and 2.5 nm. These materials offer excellent physical and chemical properties that enable a wide range of applications in biomedicine [16].

Methylglyoxal is a predictor in type 2 diabetes mellitus of intima-media thickening, vascular stiffening, and the elevation of systolic blood pressure, suggesting its clinical usefulness as a biomarker for diabetic macroangiopathy [17]. In view of the prominence of methylglyoxal in clinical applications, there are many methods reported for its determination. The use of SWCNTs as a sensing material dramatically decreases the overpotential, which makes a promising carbon nanomaterial for the detection of variety of biological compounds. Chen and co-workers have demonstrated a SWCNT based sensor platform for the quantitative determination of methylglyoxal in 0.1 M PBS (pH 7.4), showing that the sensor system was highly specific and sensitive for the detection of plasma levels of methylglyoxal in healthy individuals and diabetic patients [18].
Valacyclovir is the drug of choice for the treatment of herpes zoster and cold sores, and is an L-valyl ester and prodrug of the antiviral drug acyclovir. Subsequent to absorption, valacyclovir is rapidly and almost completely hydrolyzed to acyclovir and L-valine, which is an essential amino acid. It is also effective for the suppression or treatment of genital herpes in immunocompetent individuals. SWCNTs have been utilized to modify GCE for the sensitive detection of valacyclovir [19].

Graphene is a much newer member in the family of carbon materials in comparison to fullerenes. Graphene is an atomically thin film that consists of hexagonally arranged carbon atoms with sp² hybridization in two dimensions [15]. The combination of attributes such as a high surface area, enhanced mobility of charge carriers, and high stability makes graphene an ideal platform for the anchoring of metal nanoparticles for electrochemical sensing applications [20–22]. NADH/NAD⁺ is an important co-enzyme couple that plays a significant role in energy production/consumption within the cells of living organisms, which participates in a variety of enzymatic reactions involving more than 300 dehydrogenases [23]. Acetaminophen is a widely used analgesic (pain reliever) and fever reducer, and is considered to be very safe when administered at recommended dosages; however, it causes hepatotoxicity at higher doses [24]. Recently, graphene and Au nanoparticle-rGO nanocomposite based electrochemical sensors have been fabricated by Chen and co-workers [25,26] via an electrochemical method for the sensitive detection of NADH and acetaminophen.

Buckypaper is a thin film (5–25 µm) comprised of lateral arranged networks of nanotubes, which constitute a promising platform for the fabrication of highly concentrated and aligned nanotube-reinforced composites. Recently, much attention has been given to macroscopic assemblies of carbon nanotubes, which include carbon buckypapers, in order to exploit the characteristic properties of single carbon nanotubes at macroscopic scales [27]. Buckypapers and composites thereof are innovative materials with intriguing physical and chemical properties that have strong potential for pharmacological and prosthetic applications. Buckypaper has been incorporated into the design of a mediator-free glucose sensor with high sensitivity, stability, selectivity, and reproducibility [28].

This feature article highlights some recent development of electrochemical sensor platforms employing SWCNTs, rGO, SWCNT-rGO nanocomposites, Au nanoparticle-rGO nanocomposites, and buckypapers as sensing materials for the sensitive electrochemical detection of methylglyoxal, acetaminophen, valacyclovir, NADH and glucose with a main focus of the work carried out in the authors’ group. The conceptualization, design, optimization, and electrocatalytic activity of carbon based nanomaterials and nanocomposites for the detection of various important biological and pharmaceutical compounds are also described to demonstrate the promising medical applications of the carbon nanomaterial based electrochemical platforms.

2. Experimental

2.1. Chemicals and Reagents

All chemicals used in experiments, including methylglyoxal, acetaminophen, valacyclovir, glucose oxidase (GOx) (EC 1.1.3.4), horseradish peroxidase (HRP), β-D-glucose, NADH, graphene oxide (4 mg/mL), SWCNTs and chitosan, were purchased from Sigma-Aldrich (St. Louis, MO, USA). The buckypaper employed in this study was produced by the High-Performance Materials Institute at
Florida State University (Tallahassee, FL, USA), with a thickness of 0.035 mm. All experiments were performed in a 0.1 M phosphate buffer solution (PBS) prepared with Na₂HPO₄ and NaH₂PO₄. Titanium foils (99.2%) were purchased from Alfa Aesar (Ward Hill, MA, USA). All chemicals were used as received without further purification. All solutions were prepared with pure water produced by a Nanopure® water purification system. All solutions were freshly prepared and used within 24 h.

2.2. Fabrication of Electrochemical Sensors

Prior to modification, a glassy carbon electrode (GCE) was polished with 0.05 μm alumina powders, then sonicated for 3 min in the pure water, and finally allowed to dry at room temperature [26]. Similarly, titanium plates (1.25 cm × 0.8 cm × 0.5 mm) were ultrasonically cleaned in acetone, ethanol and water [28]. For the fabrication of the SWCNT/GCE, a 0.5 mg/mL suspension was prepared by dispersing 0.5 mg SWNTs in 1.0 mL N,N-dimethylformamide via ultrasonic agitation. The GCE surface was then coated with a known volume (40 μL) of this suspension and allowed to dry at room temperature [18,19]. For the rGO/GCE, a 4 mg/mL GO solution was added to a 0.1 M (pH 9) phosphate buffer solution (PBS) to dilute a 0.3 mg/mL GO colloidal dispersion followed by 30 min of sonication. The GO suspension within the electrochemical cell was deoxygenated using Ar gas for 15 min. Cyclic voltammetric deposition was performed in the GO suspension (0.3 mg/mL) with an electrode potential range between 0.5 to −1.5 V at scan rate of 10 mV/s using a three-electrode system with a bare GCE as the working electrode. Five reduction cycles were employed in this study to deposit rGO onto the GCE surface [26]. The Au nanoparticle-rGO/GCE was fabricated by casting a 5 μL mixture of GO (0.5 mg/mL), AuCl₃ (10 mM), and Nafion (0.5%) on the cleaned GC electrode, which was then allowed to air dry.

Scheme 1. Schematic diagram of the preparation of the buckypaper-based electrochemical biosensor. (a) a titanium plate sputtered with a thin layer of gold; (b) gold interaction with chitosan; (c) attachment of buckypaper to the gold surface via chitosan; (d) activation of the buckypaper; (e) immobilization of enzymes; (f) the fabricated biosensor. All figures are reproduced with permission from [28]. Copyright 2011 Elsevier.

The in situ formation of Au nanoparticle-rGO sheets on the GCE was achieved in 0.1 M H₂SO₄ by applying a potential of −1.0 V (vs. Ag/AgCl) for 500 s [25]. The SWCNT-rGO nanohybrid/GCE was
fabricated by casting 40 μL (0.5 mg/mL) of each batch of SWCNTs prepared in N,N-dimethylformamide and GO suspension (4 mg/mL) on the GCE surface, followed by air drying and electrochemical reduction of GO through continuous cycles under a potential range between 0.5 and −1.5 V for five cycles. Details on the fabrication of the buckypaper based biosensor are presented in Scheme 1 [28]. The titanium electrode for glucose sensing was prepared by attaching a portion of the buckypaper to a gold coated titanium plate, which was facilitated by 5 μL of 2 mg/mL chitosan. The buckypaper surface was then activated through cyclic voltammetry with a potential range between −0.8 V and 0.4 V at a scan rate of 10 mV/s in a phosphate buffer solution (PBS) followed by 90 s amperometric scanning at a potential of 1.5 V in order to generate carboxylic groups. The activated surface was then enzymatically immobilized via the casting of a mixed solution comprised of 20 μL of 5 mg/mL GOx, 10 μL of 2 mg/mL HRP and 10 μL of 2 mg/mL chitosan onto the buckypaper.

2.3. Apparatus

All electrochemical experiments, including cyclic voltammetry (CV), differential pulse voltammetry (DPV), square wave voltammetry (SWV) and amperometry were conducted with a CHI 660D electrochemical workstation (CH Instruments Inc., Austin, TX, USA) using a conventional three-electrode system that consisted of a platinum wire counter electrode, a 3 M KCl saturated Ag/AgCl reference electrode, and a carbon nanomaterial based working electrode. A field-emission scanning electron microscope (FE-SEM; Hitachi SU-70, Japan) was employed to characterize the surface morphologies of the carbon nanomaterials and their nanocomposites, which were deposited on the surfaces of the electrode. All experiments were performed at room temperature and the electrode potentials quoted are versus an Ag/AgCl electrode.

3. Results and Discussion

3.1. Electrochemical Sensing of Methylglyoxal, Acetaminophen and Valacyclovir at SWCNTs

SEM was utilized to study the surface characteristics and morphology of the fabricated SWCNT-modified GCE. Figure 1A depicts a SEM image of the SWCNTs on the GCE surface, which were uniformly covered with an interlinked nanoporous network structure. The prepared SWCNT/GCE was tested for the detection of methylglyoxal, acetaminophen, and valacyclovir. Figure 1B shows the cyclic voltammograms (CVs) of the SWCNT/GCE recorded in 0.1 M PBS (at pH 7.4) at a scan rate of 20 mV/s in the absence (Curve a) and presence of 10 μM methylglyoxal (Curve b). A single well-defined reduction peak of methylglyoxal was observed at −0.864 V (Curve b) compared with the CV recorded in 0.1 M PBS (Curve a). The absence of anodic peaks in the reverse scan revealed that the reduction of methylglyoxal in this particular system was irreversible in nature.

Figure 1C depicts a typical reversible behavior of 50 μM acetaminophen at the SWCNT/GCE 0.1 M PBS at pH 7.4 under a scan rate of 20 mV/s. A distinct oxidation peak at 0.372 V and reduction peak at 0.355 V obtained during the measurement of CV revealed its reversible behavior. As shown in Figure 1D, the CV of 50 μM valacyclovir in 0.1 M PBS (pH 7.4) at the scan rate of 20 mV/s generated distinct broad oxidation peak at 0.978 V, whereas no obvious reduction peak was observed during reverse scanning, indicating the irreversible catalytic behavior of valacyclovir.
Figure 1. (A) FE-SEM images of the SWCNTs on the GCE surface. CV performance recorded at the SWCNT/GCE, in the absence (red solid line), and in the presence of (blue solid line) (B) 10 µM methylglyoxal, (C) 50 µM acetaminophen, and (D) 50 µM valacyclovir, in 0.1 M PBS (pH 7.4) at a scan rate of 20 mV/s. Figure 1B reproduced with permission from [18]. Copyright 2013 Elsevier.

Figure 2. Responses of the SWCNTs/GCE recorded in 0.1 M PBS (pH 7.4) (A) and (B) SWV responses of different methylglyoxal concentrations (0.1 µM to 100 µM) (C) DPV responses of acetaminophen (5 nM to 80 µM) and (D) DPV responses of valacyclovir (5–55 nM). Figure 2A,B reproduced with permission from [18]. Copyright 2013 Elsevier. Figure 2D reproduced with permission from [19]. Copyright 2013 Royal Society of Chemistry.
Figure 2A,B illustrate the responses of various concentrations of methylglyoxal through square wave voltammetry (SWV), where the peak current was increased linearly with the successive addition of different concentrations of methylglyoxal. Figure 2A depicts a linear increase in the peak current responses with increased methylglyoxal concentrations in the range 0.1–2.0 μM, with a strong linear association (R² value of 0.99). Similarly, Figure 2B shows the responses of the higher concentration range of (20–100 μM). Different concentrations of acetaminophen were studied through DPV as presented in Figure 2C, where the peak current increased linearly with increasing concentrations from 5 nM to 80 μM with a linear association (R²) value of 0.97 and a calculated limit of detection of 4.3 nM. Figure 2D shows the DPV responses of different valacyclovir concentrations varied from 5 nM to 55 nM with a limit of detection of 1.8 nM.

Besides the aforementioned three examples, carbon nanotubes and carbon nanotube composite based electrochemical sensing platforms have been widely explored for the detection of various biological and pharmaceutical compounds including glucose [29–31], lactate [32], dopamine [33–35], rutine [36], human serum albumin [37], DNAs [38], epinephrine [39,40], cholesterol [41,42], methimazole [43], sumatriptan [44], and paracetamol [45]. The electrochemical methods, the linear range and the detection limit of those studies are compared in Table 1.

**Table 1.** List of electrochemical sensors and biosensors based on carbon nanomaterials and their composites for the detection of biological and pharmaceutical compounds.

| Carbon Based Nanomaterials | Analytes          | Methods      | Linear Range          | Detection Limit | Reference |
|----------------------------|-------------------|--------------|-----------------------|-----------------|----------|
| SWNT-Nafion-GOX            | Glucose           | Amperometry  | 2 mM to 20 mM         | -               | 29       |
| SWNT-GoX                   | Glucose           | Amperometry  | Up to 40 mM           | -               | 30       |
| Pt-Nafion-SWCNTs-GOX       | Glucose           | Amperometry  | 0.5 μM to 5 mM        | 0.5 μM          | 31       |
| SWNT-mineral-oil paste     | Lactate           | Amperometry  | Up to 7.0 mM          | 0.3 mM          | 32       |
| Nafion-SWNT                | Dopamine          | DPV          | 0.02 μM to 6.0 μM     | 5.00 nM         | 33       |
| SWNT polymer composite     | Dopamine          | CV           | 16 nM to 600 μM       | 8 nM            | 34       |
| SWCNTs                     | Dopamine          | DPV          | 3 μM to 200 μM        | 48 nM           | 35       |
| SWCNTs                     | Rutine            | CV           | 20 nM to 5 μM         | 10 nM           | 36       |
| SWCNTs                     | Human serum albumin | CV         | 0.075 nM to 7.5 nM    | 75 pM           | 37       |
| SWCNTs                     | DNAs              | DPV          | 5 μM to 30 μM         | 1.43 μM         | 38       |
| MWCNTs-Nafion              | Epinephrine       | CV and DPV   | 0.06 mM to 0.24 mM    | 0.02 mM         | 39       |
| MWCNTs                     | Epinephrine       | LSV          | 50 nM to 10 μM        | 10 nM           | 40       |
| MWCNTs                     | Cholesterol       | Amperometry  | Up to 6.0 mM          | 0.2 mM          | 41       |
| MWCNTs                     | Cholesterol       | Amperometry  | 100 mg/dL to 400 mg/dL| -               | 42       |
| MWCNTs                     | Methimazole       | Amperometry  | 0.074 μM to 63.5 μM   | 0.056 μM        | 43       |
| MWCNTs-silver nanoparticles| Sumatriptan       | CV           | 80 nM to 100 μM       | 40 nM           | 44       |
| MWCNTs                     | Paracetamol       | ASV          | 0.01 μM to 20 μM      | 10 nM           | 45       |
| SWCNTs                     | Methylglyoxal     | SWV          | 0.1 μM to 100 μM      | -               | 18       |
| SWCNTs                     | Valacyclovir      | DPV          | 5 nM to 55 nM         | 1.8 nM          | 19       |
| SWCNTs                     | Acetaminophen     | DPV          | 5 nM to 80 μM         | 4.3 nM          | This work|
| CuO-graphene               | Glucose           | Amperometry  | 1 μM to 8 mM          | 1 μM            | 46       |
| CuNPs/graphene             | Glucose           | Amperometry  | 0.5 μM to 4.5 mM      | 0.5 μM          | 47       |
Table 1. Cont.

| Carbon Based Nanomaterials | Analytes      | Methods   | Linear Range       | Detection Limit | Reference |
|----------------------------|--------------|-----------|--------------------|-----------------|-----------|
| Graphene-ppy               | Glucose      | Amperometry | -                  | 3 µM            | 48        |
| Graphene-Pt                | Ascorbic acid| DPV       | 0.03 µM to 8.13 µM | 0.03 µM         | 49        |
| Graphene                   | Norepinephrine| Amperometry | 0.04 µM to 100 µM  | 0.84 nM         | 50        |
| Reduced GO                 | NADH         | Amperometry | 10 µM to 600 µM    | 0.33 µM         | 51        |
| Graphene-Au nanorod        | NADH         | Amperometry | 5 µM to 337 µM     | 1.5 µM          | 52        |
| Au-TiO2/graphene           | NADH         | Amperometry | 10 µM to 240 µM    | 0.2 µM          | 53        |
| Graphene-TiO2              | NADH         | Amperometry | 10 nM to 2 mM      | 3 × 10⁻⁹ M      | 54        |
| AuNPs-rGO                  | NADH         | Amperometry | 50 nM to 500 µM    | 1.13 nM         | 25        |
| Nitrogen doped Graphene    | Uric acid    | DPV       | 0.1 µM to 20 µM    | 0.045 µM        | 55        |
| Graphene                   | Uric acid    | Amperometry | 0.19 µM to 49.68 µM| 0.132 µM        | 56        |
| Nafion-AgNPs-rGO           | Uric acid    | LSV       | 10 µM to 800 µM    | 8.2 µM          | 57        |
| Pt-rGO                     | Uric acid    | DPV       | 10 µM to 130 µM    | 0.45 µM         | 59        |
| ERGO                       | Serotonin    | DPV       | 5 µM to 300 µM     | 0.11 µM         | 59        |
| ERG/ NiO₂/ NiO             | Acetaminophen| DPV      | 0.04 µM to 100 µM  | 0.02 µM         | 60        |
| Graphene-chitosan          | Acetaminophen| DPV      | 1 µM to 100 µM     | 0.3 µM          | 61        |
| Graphene                   | Acetaminophen| SWV      | 0.1 µM to 20 µM    | 0.032 µM        | 62        |
| rGO                        | Acetaminophen| DPV      | 5 nM to 800 µM     | 2.13 nM         | 26        |
| SWCNTs-GNS                 | Acetaminophen| DPV      | 0.05 µM to 64.5 µM | 0.038 µM        | 63        |
| MWCNT-graphene nanosheets  | Acetaminophen| DPV      | 0.8 µM to 110 µM   | 0.1 µM          | 64        |
| MWCNT/GO                   | Acetaminophen| DPV      | 0.5 µM to 400 µM   | 47 nM           | 65        |
| SWCNTs-rGO                 | Acetaminophen| DPV      | 5 nM to 80 µM      | 1.4 nM          | This work |
| MWCNT/GO                   | Dopamine     | DPV       | 0.2 µM to 400 µM   | 22 nM           | 65        |
| MWCNT/GONR                 | Dopamine     | DPV       | 0.15 µM to 12.15 µM| 0.08 µM         | 66        |
| Buckypaper-SWCNTs          | Glucose      | Amperometry | 0 nM to 10 mM     | 0.022 mM        | 70        |
| Buckypaper-GOx-HRP         | Glucose      | Amperometry | Up to 9 mM        | 0.01 mM         | 28        |

3.2. Electrochemical Sensing of Acetaminophen and NADH at rGO and Au Nanoparticle-rGO Nanocomposites

Figure 3A displays a FE-SEM image of rGO sheets deposited on GCE through an electrochemical technique, where a characteristic rippled-like structure indicated that the GCE surface was uniformly coated with graphene. The electrochemical behavior of 50 µM acetaminophen was studied through CV at rGO/GCE in 0.1 M PBS at pH 7.4 at the scan rate of 20 mV/s (Figure 3B). The appearance of the redox peaks indicated a reversible electrode process. The performance of rGO/GCE toward different concentrations of acetaminophen was studied through differential pulse voltammetry (DPV). Figure 3C shows the linear increments of current responses upon the successive addition of acetaminophen concentrations from 5 µM to 800 µM with an excellent correlation coefficient ($R^2 = 0.996$). The analytical performance of the rGO/GCE for the lowest acetaminophen concentrations were evaluated through an amperometric technique under a constant electrode potential of 0.5 V. A rapid increment of current was observed upon the successive addition of 5.0 nM, 0.2 µM, and 2.0 µM acetaminophen solutions in 20 mL of 0.1 M PBS (Figure 4A). The calibration plot of current vs acetaminophen concentration gave a strong linear association with a correlation coefficient of $R^2 = 0.986$ (Figure 4B).
The limit of detection (LOD) was calculated to be 2.1 nM using 3s/b, where s is the standard deviation of the blank and b is the slope of the calibration curve.

Figure 3. (A) FE-SEM images of electrochemically reduced graphene oxide (rGO) on the GCE surface; (B) CVs recorded at the rGO/GCE in the absence (red solid line), and presence of 50 µM acetaminophen (blue solid line) in 0.1 M PBS (pH 7.4) at a scan rate of 20 mV/s; (C) DPVs of the rGO/GCE recorded in 0.1 M PBS (pH 7.4) containing different acetaminophen concentrations (5 µM to 800 µM). Figure 3A and 3B are reproduced with permission from [26]. Copyright 2015 Elsevier.

Figure 5A shows a typical FE-SEM image of the Au nanoparticle/rGO nanocomposite electrode. Au nanoparticles with an average size of 8.2 nm were uniformly distributed across the rGO sheet. Figure 5B compares the CVs of the rGO/GCE (blue) and Au nanoparticle-rGO/GCE (red) recorded in 0.1 M PBS (pH 7.2) containing 1 mM NADH at the scan rate of 20 mV/s. A broad anodic peak appeared at 0.54 V for NADH at the rGO/GCE whereas a large well-defined anodic oxidation peak appeared at the Au nanoparticle-rGO/GCE, with a ~2.3 fold increase of the current density. Figure 5C displays amperometric i-t curves under an applied potential of 0.55 V for the sensitive detection of NADH using an Au nanoparticle-rGO/GCE platform. The calibration plot of current vs. NADH concentrations are presented in Figure 5D, which shows two linear concentration ranges: one from 50 nM to 50 µM with a correlation coefficient of $R^2 = 0.987$, and the other from 50 µM to 500 µM with a correlation coefficient of $R^2 = 0.997$. The limit of detection was calculated to be 1.13 nM. Such a low detection limit was far improved from those of most existing reports with graphene and their composite materials for the electrochemical detection of NADH. Moreover, a variety of electrochemical sensors were
prepared with graphene and its nanocomposites for the detection of glucose \cite{46–48}, ascorbic acid \cite{49}, norepinephrine \cite{50}, NADH \cite{51–54}, uric acid \cite{55–58}, serotonin \cite{59}, and acetaminophen \cite{60–62}. The performance of these electrochemical sensors is also compared in Table 1.

**Figure 4.** (A) Amperometric responses recorded at rGO/GCE in 0.1 M PBS under various acetaminophen concentrations (5 nM–4 µM), $E_{\text{app}}$: 0.5 V and (B) the calibration plot of current vs acetaminophen concentrations. All figures are reproduced with permission from \cite{26}. Copyright 2015 Elsevier.

**Figure 5.** (A) FE-SEM images of the AuNP-rGO on the GCE surface. (B) CVs recorded at the rGO/GCE (blue), and AuNPs-rGO/GCE (red) recorded in 1 mM NADH. (C) Amperometric i-t curve recorded at the Au-rGO/GCE in 0.1 M PBS under various NADH concentrations (50 nM–0.5 mM), $E_{\text{app}}$: 0.55 V and (D) the calibration plot of current density vs. NADH concentration. All figures are reproduced with permission from \cite{25}. Copyright 2015 Elsevier.
3.3. Electrochemical Sensing of Acetaminophen and Valacyclovir at SWCNTs-rGO Nanocomposites

Figure 6A depicts a FE-SEM image of the SWCNT-rGO nanocomposite, in which graphene films were uniformly dispersed throughout the SWCNTs. The electrochemical behaviors of the SWCNT-rGO/GCE were investigated through the CV response of 50 µM acetaminophen in 0.1 M PBS (pH 7.4) at the scan rate of 20 mV/s. As shown in Figure 6B, a distinct well-defined reversible peak (blue solid line) was observed, which can be attributed to the oxidation and reduction of acetaminophen. The current response observed through the oxidation and reduction of 50 µM acetaminophen at the SWCNT-rGO/GCE was much higher than that of the responses obtained through individual SWCNT (Figure 1C) and rGO (Figure 3B) platforms. Figure 6C shows the typical valacyclovir response at the SWCNTs-rGO/GCE with a distinct oxidation peak (blue solid line), having no obvious reduction peak during reverse scanning. DPV responses of different concentrations of acetaminophen at the SWCNT-rGO/GCE are presented in Figure 6D. The current responses were increased with increasing concentrations of acetaminophen (from 5 nM to 80 µM), and shows a linear association (R² value of 0.97) between current vs. acetaminophen concentrations. The low detection limit of 1.4 nM was achieved at the SWCNT-rGO nanocomposite platform through DPV, which is much lower than that of the value achieved through individual SWCNT and rGO platforms. Some other carbon nanotube-graphene nanocomposite based electrochemical sensing platforms for the detection of acetaminophen [63–65] and dopamine [65, 66] are summarized in Table 1.

![Figure 6. (A) FE-SEM images of the SWCNTs-rGO nanocomposite on the GCE surface. CV performance recorded at the SWCNT-rGO nanocomposite/GCE in the absence (red solid line), and presence of (blue solid line) (B) 50 µM acetaminophen, (C) 50 µM valacyclovir in 0.1 M PBS (pH 7.4) at a scan rate of 20 mV/s. (D) DPVs of the SWCNT-rGO nanocomposite/GCE recorded in 0.1 M PBS (pH 7.4) containing different acetaminophen concentrations (5 nM to 80 µM).]
3.4. Electrochemical Sensing of Glucose at Buckypaper

The morphology and composition of the buckypaper were characterized using FE-SEM. Figure 7A shows a typical SEM image of the buckypaper film, which reveals that the SWNTs were arranged in bundles that formed networks. The biosensor composed of Au/BP/GOx-HRP on a titanium surface was studied for its glucose response. Figure 7B shows the CV responses of Au/BP/GOx-HRP/Ti prior to (a) and following (b) the addition of 10 mM glucose in 0.1 M PBS (pH 7.4) at the scan rate of 10 mV/s. The CV curves revealed a decreased oxidation peak, as well as a significantly increased reduction peak, showing a strong response to glucose oxidation. The increase of the reduction peak might have been due to the H$_2$O$_2$ that was generated through the oxidation of glucose via the immobilized GOx [67–69]. The amperometric response of the Au/BP/GOx-HRP/Ti biosensor at an optimized potential under physiological pH 7.4 in 0.1 M PBS upon the successive addition of 1 mM glucose is shown in Figure 7C. For comparison, the amperometric response of the Ti/Au/BP electrode is also included in Figure 7C (Curve ii). No obvious current response was observed at the non-enzymatic electrode, while a strong response was achieved through the Au/BP/GOx-HRP/Ti biosensor (Figure 7C(i)). Figure 7D illustrates the corresponding current vs. different glucose concentration with an excellent linear correlation coefficient of $R^2 = 0.993$. The limit of detection for the detection of glucose through this fabricated biosensor was estimated to be 0.01 mM. The detection limit obtained through this study was much lower than that of previously reported for the buckypaper based detection of glucose [70].

![Figure 7.](image-url)
3.5. Electrocatalytic Performance of SWCNTs, rGO and SWCNTs-rGO for Acetaminophen and Valacyclovir: Comparative Analysis

To demonstrate the electrocatalytic behavior of three carbon nanomaterial-based platform nanocomposites (SWCNT, rGO, and SWCNT-rGO), the responses to acetaminophen and valacyclovir were compared. As seen in Figure 1C, Figure 3B, and Figure 6B, the electrocatalytic behavior of 50 µM acetaminophen at the SWCNT, rGO and SWCNT-rGO nanocomposite electrodes revealed significant changes in oxidation-reduction peak current responses. In the present study, the lowest oxidation peak current was observed with the rGO platform, whereas a six fold increase in the oxidation current response was observed at the SWCNT platform under identical experimental conditions. The electrocatalytic oxidation current response was greatly improved at the SWCNT-rGO nanocomposite, which exhibited an over six times higher current response than that of SWCNT platform, and was 12 times higher than that of the rGO. The peak to peak separation (ΔEp) was estimated to be 75 mV, 45 mV, and 30 mV at the rGO, SWCNT and SWCT-rGO nanocomposites, respectively. Decreasing ΔEp values, in the order rGO > SWCT > SWCT-rGO observed through these platforms reflected a more rapid electron transfer reaction with efficient electrocatalytic oxidation. A similar pattern was observed for the oxidation of 50 µM valacyclovir through the SWCNT (Figure 1D) and SWCT-rGO nanohybrid (Figure 6C), where we observed an almost 21% increment of oxidation current response with the SWCT-rGO nanocomposite. In general, the structure and properties of carbon materials, such as surface functional groups, graphitization structure, charge density and surface area, have a significant impact on their electrocatalytic activities. The high crystallinity and one-dimensional (1D) structure of CNTs endow high conductivity and high surface areas. Graphene, in addition to possessing similar stable physical properties as CNTs, has larger edge planes with high charge densities. Further, the heterogeneity of carbon materials with edge planes may better stabilize and augment electrocatalytic activity [71]. Due to strong van der Waals forces, 1D CNTs and two-dimensional (2D) rGO sheets are both inclined to aggregate, or stack with each other in forming a three-dimensional (3D) CNT-rGO nanocomposite network with synergetic properties [72]. As a result, this hybrid material provides exceptional characteristics with a higher nominal edge density per unit area than individual CNTs and rGO nanosheets, thereby fully utilizing it capacities for charge density, surface area, and available active sites for catalytic reactions. In fact, the integration of CNT and rGO to comprise nanocomposite materials is a quite intriguing strategy for enhancing the dispersion of CNT and rGO, toward exploiting the distinct advantages of both CNT and rGO in obtaining a robust and efficacious electronic and thermally conductive 3D network [73].

4. Conclusions

Carbon-based nanomaterials and composites thereof were employed in the design of highly sensitive electrochemical sensors for the detection of biologically and pharmaceutically important compounds. Carbon and its derivatives possess excellent electrocatalytic properties for the modified sensors, such as enhanced detection sensitivity, electrocatalytic effects, high conductivity, and reduced fouling. These superior attributes endow SWCNTs, graphene, and buckypaper nanocomposites with great advantages for enhanced sensing and biosensing applications. In the present feature article,
important biomolecules such as methylglyoxal, glucose, and NADH, as well as pharmaceutical molecules such as acetaminophen and valacyclovir were selected to establish and verify the sensing performance of carbon nanomaterials based electrochemical platforms. Carbon based nanomaterials provide an excellent electrocatalytic platform for the sensitive detection of these compounds. Further demonstrations with acetaminophen and valacyclovir revealed that SWCNT-rGO nanocomposites possess excellent electrocatalytic activity in comparison to individual SWCNTs and rGO platforms. Among the carbon based nanomaterials tested in this study, SWCNT-rGO nanocomposites were shown to be an excellent platform for use in the core architectures of future sensor/biosensor designs. Carbon nanomaterials and nanohybrids thereof may play a critical role in the future development of advanced point of care diagnostics.

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Author Contributions

B.-R.A. and A.C. conceived the original idea, approach and design of experiments. B.-R.A and M.G. conducted the experiments and wrote the manuscript. A.C. supervised the project and reviewed the paper.

Conflict of Interest

The authors declare no conflict of interest.

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