Article

Influence of Temperature on Vibrational Frequency of Graphene Sheet Used as Nano-Scale Sensing

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Abstract: In this study, the vibrational properties of single- and double-layer graphene sheets (GSs) with attached nanoparticles are analyzed based on the nonlocal elasticity theory. The potential applications of atomic-scale mass sensing are presented using GSs with simply supported boundary condition. The frequency equation for GSs with an attached nanoparticle is derived to investigate the vibration frequency of the GSs under thermal environment. Using the proposed model, the relationship between the frequency shifts of graphene-based mass sensor and the attached nanoparticles is obtained. The nonlocal effect and the temperature dependence on the variation of frequency shifts with the attached nanomass and the positions on the GS are investigated and discussed in detail. The obtained results show that the nanomass can be easily detected by using GS resonator which provides a highly sensitive nanomechanical element in sensor systems. The vibrational frequency shift of GS increases with increasing the temperature dependence. The double-layer GSs (DLGSs) have higher sensitivity than the single-layer GSs (SLGSs) due to high frequency shifts.

Keywords: graphene sheet; sensor; nonlocal elasticity theory

1. Introduction

Graphene sheets (GSs) are an excellent two-dimensional (2D) material, and have drawn attention in recent years. Due to the unique properties of small size, super mechanical properties, and high vibration frequency, graphene sheets have been considered an ideal structure element for fabricating nano electromechanical systems (NEMS) [1–3]. Novel graphene-based nanostructures are believed to have tremendous potential applications for developing sensors of various types such as magnetic and electric field sensors, strain and nanoscale mass sensors [3–7]. The mechanism of using GSs as nanoscale mass sensing is based on the fact that the resonant frequency of GSs is shifted upward when a nanoparticle is loaded onto the GSs and the shift frequency is sensitive to the attached nanomass. One relevant application of the GSs is given by carbon nanotubes (CNTs) which can be obtained by rolling a graphene sheet into a cylinder. Due to their mechanical and physical properties, CNTs are applied as ultrahigh-frequency nano-mechanical resonators in a large number of NEMS such as sensors, oscillators and charge detectors [8–14].

At present, the elastic continuum theory and model have been regarded as an effective way to investigate the vibration behavior of GSs [15,16]. The vibration analysis of multilayered graphene sheets (MLGSs) using a continuum model has been reported by Kitipornchai et al. [15]. They indicated that the vibration modes of the graphene sheets that were associated with the resonant frequencies...
were quite different due to the influence of the Van der Waals (vdW) interaction between any two nanosheets. The lowest natural frequency of a MLGS was independent of the vdW interaction, but all of the other higher natural frequencies depended largely on the vdW interaction. Natsuki’s group has investigated the vibrational characteristics of embedded double-layer graphene nanoribbons (DLGNRs) based on the nonlocal elasticity theory [16]. The results showed that the vibrational frequency of DLGNRs had a significant difference between the in-phase mode (IPM) and the anti-phase mode (APM). The nonlocal parameter affected the natural frequencies, especially for higher modes of vibration. Moreover, the natural frequencies of the APM were less sensitive to the vibration modes than those of the IPM. The operation of a NEMS mass sensor relied on monitoring how the resonance frequency of a nanomechanical resonator changed when an additional nanomass was adsorbed onto its surface [17–20]. Many studies have shown that GSs exhibit ultrahigh frequency ranges up to the terahertz order because of the inherent characteristics of the materials [15,16,21–23]. These inherent characteristics led to a higher sensitivity to atomic-scale mass detection, whereby the attached nanomass can easily cause a shift in the resonant frequency of the GS resonator.

Ultrasonic vibration and attenuation are the important properties for design and performance of the sensor devices. Graphene sheets appear to be excellent element materials for structures of nanomechanical resonators because they can generate terahertz frequency vibration. Some studies have been carried out on utilizing the molecular dynamics (MD) method [24,25], molecular structural mechanics [26] or continuum mechanics [20,27–30] to investigate the applications of single-layer graphene sheets (SLGSs) for NEMS mass sensors. However, there is very little investigation of the resonant frequency of double-layer graphene sheets (DLGSs) used as mass sensors and how temperature affects the vibration frequency of GSs [27].

In this paper, an analytical method based on the nonlocal elasticity theory [31–33] is introduced to predict the vibrational property of GSs as nanoscale sensing. The potential applications using single- and double-layer GSs with a simply supported boundary condition are illustrated. To investigate the vibration frequency of GSs under a thermal environment, the frequency equation in GSs for attaching a nanoparticle is derived. Using the present theory, the relationship between the frequency shifts of the graphene-based mass sensor and the attached nanoparticles is investigated in detail.

2. Theoretical Approach

2.1. Governing Equation Based on Nonlocal Elasticity

We present a continuum elastic model to analyze the vibration frequency of the nanomechanical mass sensor when temperature effects are considered. As shown in Figure 1, a nanoparticle \( m_c \) is attached to a simply supported graphene sheet (GS) in an arbitrary position \( (x_0, y_0) \), thereby inducing a vibrational frequency shift due to the attached mass. The origin is taken at a corner of the mid-plane of the nanoplate. The coordinates \( x \) and \( y \) are taken along the length \( L_a \) and the width \( L_b \) of the GSs, respectively. The coordinate \( z \) is taken along the thickness \( h \) direction of the GSs.

![Figure 1. Schematic illustration showing graphene sheet with an attached mass.](image)
In the analysis, the vibration frequency of GSs is analyzed based on nonlocal elasticity theory. For the nonlocal linear elastic solids, Hook’s law of nanoplates can be expressed as

\[
\begin{align*}
\sigma_{xx} &= (\epsilon_0 a)^2 \nabla^2 \sigma_{xx} = \frac{E}{1-\nu^2} (\epsilon_{xx} + \nu \epsilon_{yy}) - \frac{E a \Delta T}{1-\nu}, \\
\sigma_{yy} &= (\epsilon_0 a)^2 \nabla^2 \sigma_{yy} = \frac{E}{1-\nu^2} (\epsilon_{yy} + \nu \epsilon_{xx}) - \frac{E a \Delta T}{1-\nu}, \\
\tau_{xy} &= (\epsilon_0 a)^2 \nabla^2 \tau_{xy} = G \gamma_{xy}
\end{align*}
\]

(1)

where \(\sigma\) and \(\tau\) are the normal stress and the shear stress. \(\epsilon\) and \(\gamma\) are the axial strain and shear strain. \(E\), \(G\) and \(\nu\) are the tensile, shear elastic modulus and Poisson’s ratio, respectively. \(a\) is the coefficient of thermal expansion, and \(\Delta T\) is the temperature change. The parameter \(\epsilon_0 a\) is the nonlocal parameter appropriate to GSs, where \(\epsilon_0\) is a constant appropriate to the material and has to be determined for each material independently by experiment or atomistic simulation. \(a\) is the internal characteristic length of a carbon-carbon bond (e.g., lattice parameter, granular size, distance between atoms’ bond) [13,16].

The expressions between the strain and the displacement are [34]

\[
\begin{align*}
\epsilon_{xx} &= -\frac{1}{2} \frac{\partial^2 w}{\partial x^2}, \\
\epsilon_{yy} &= -\frac{1}{2} \frac{\partial^2 w}{\partial y^2}, \\
\gamma_{xy} &= -2\nu \frac{\partial^2 w}{\partial x \partial y}
\end{align*}
\]

(2)

where \(w\) is the transverse displacement.

The bending and shear moments can be defined by

\[
\begin{align*}
M_{xx} &= \int_{-h/2}^{h/2} z \sigma_{xx} dz \\
M_{yy} &= \int_{-h/2}^{h/2} z \sigma_{yy} dz \\
M_{xy} &= \int_{-h/2}^{h/2} z \tau_{xy} dz
\end{align*}
\]

(3)

Substituting Equation (1) into Equation (3) and using Equation (2), we obtain

\[
\begin{align*}
M_{xx} &= (\epsilon_0 a)^2 \left( \frac{\partial^2 M_{xx}}{\partial x^2} + \frac{\partial^2 M_{xx}}{\partial y^2} \right) = -D \left( \frac{\partial^2 w}{\partial x^2} + \nu \frac{\partial^2 w}{\partial y^2} \right) \\
M_{yy} &= (\epsilon_0 a)^2 \left( \frac{\partial^2 M_{yy}}{\partial x^2} + \frac{\partial^2 M_{yy}}{\partial y^2} \right) = -D \left( \frac{\partial^2 w}{\partial y^2} + \nu \frac{\partial^2 w}{\partial x^2} \right) \\
M_{xy} &= (\epsilon_0 a)^2 \left( \frac{\partial^2 M_{xy}}{\partial x^2} + \frac{\partial^2 M_{xy}}{\partial y^2} \right) = -D (1 - \nu) \frac{\partial^2 w}{\partial x \partial y}
\end{align*}
\]

(4)

and

\[
D = \frac{E h^3}{12(1-\nu^2)}
\]

(5)

According to the elastic theory of thin plates, the governing equation of the vibration for GSs with the external load is given in

\[
0 = \frac{\partial^2 M_{xx}}{\partial x^2} + 2 \frac{\partial^2 M_{xy}}{\partial x \partial y} + \frac{\partial^2 M_{yy}}{\partial y^2} + N^T \left( \frac{\partial^2 w}{\partial x^2} + \frac{\partial^2 w}{\partial y^2} \right) = \left[ \rho h + m_c \delta(x-x_0) \delta(y-y_0) \right] \frac{\partial^2 w}{\partial t^2}
\]

(6)

where \(m_c\) is an attached nanoparticle mass, \(h\) is the thickness of GSs, \(t\) is the time, \(\delta\) is the Dirac delta function, and \(N^T\) is the load caused by the temperature change given by

\[
N^T = \frac{E a \Delta T h}{1-\nu}
\]

(7)

Substituting Equation (4) into Equation (6), we yield

\[
0 = \frac{\partial^2 M_{xx}}{\partial x^2} + 2 \frac{\partial^2 M_{xy}}{\partial x \partial y} + \frac{\partial^2 M_{yy}}{\partial y^2} + N^T \left( \frac{\partial^2 w}{\partial x^2} + \frac{\partial^2 w}{\partial y^2} \right) = \left[ \rho h + m_c \delta(x-x_0) \delta(y-y_0) \right] \frac{\partial^2 w}{\partial t^2}
\]

(8)
The bi-harmonic operator is given by

\[ \nabla^4 = \frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2}, \quad \nabla^4 = \frac{\partial^4}{\partial x^4} + 2 \frac{\partial^4}{\partial x^2 \partial y^2} + \frac{\partial^4}{\partial y^4} \]  

(9)

2.2. Dynamic Analysis for Single-Layer Graphene Sheet (SLGSs)

We can introduce the following substitution to obtain the vibration frequency of the governing equations for SLGSs

\[ w(x, y, t) = Y(x, y) e^{i\omega t} \]  

(10)

where \( Y(x, y) \) is the shape function of deflection, and \( \omega \) is the resonant frequency of the GSs.

Substituting Equation (10) into Equation (8), the governing equation is written as

\[ D \nabla^4 Y - \omega^2 \left[ 1 - (\epsilon_0 a)^2 \nabla^2 \right] \left[ \rho h + m_c \delta(x - x_0) \delta(y - y_0) \right] Y = N! \left[ 1 - (\epsilon_0 a)^2 \nabla^2 \right] \nabla^2 Y \]  

(11)

To obtain the solution of Equation (11), we consider GSs subjected to a simply supported condition. For the simply supported GSs, we have

\[ \frac{\partial^2 w(x, y, t)}{\partial x^2} = \frac{\partial^2 w(x, y, t)}{\partial y^2} = 0, \text{ on } x = 0, \text{ } x = L_a \text{ and } y = 0, \text{ } x = L_b \]  

(12)

To satisfy the above set of boundary conditions, therefore, the shape function of deflection in Equation (10) can be expressed as:

\[ Y(x, y) = A_{mn} \sin \frac{m \pi x}{L_a} \sin \frac{n \pi y}{L_b} \]  

(13)

where \( A_{mn} \) is the vibration amplitude of oscillation. \( m \) and \( n \) indicate the mode numbers.

Substituting Equation (13) into Equation (11), we get

\[ D \pi^4 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right)^2 \sin \frac{m \pi x}{L_a} \sin \frac{n \pi y}{L_b} - \omega^2 \left[ 1 + \pi^2 (\epsilon_0 a)^2 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right) \right] \left[ \rho h + m_c \delta(x - x_0) \delta(y - y_0) \right] \sin \frac{m \pi x}{L_a} \sin \frac{n \pi y}{L_b} + N! \left[ 1 + \pi^2 (\epsilon_0 a)^2 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right) \right] \pi^2 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right) \sin \frac{m \pi x}{L_a} \sin \frac{n \pi y}{L_b} = 0 \]  

(14)

Multiplying Equation (14) by \( \sin \frac{m \pi x}{L_a} \sin \frac{n \pi y}{L_b} \) and integrating over the whole region with respect to \( x \) and \( y \) with the limits \( x = 0 \) to \( L_a \) and \( y = 0 \) to \( L_b \), we obtain the following frequency equation after simplifications,

\[ \omega^2 = \frac{D \pi^4 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right)^2 + N! \left[ 1 + \pi^2 (\epsilon_0 a)^2 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right) \right] \pi^2 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right)}{1 + \pi^2 (\epsilon_0 a)^2 \left( \frac{m^2}{L_a^2} + \frac{n^2}{L_b^2} \right)} \left( \rho h + \frac{4}{L_b} m_c \sin^2 \frac{m \pi x}{L_a} \sin^2 \frac{n \pi y}{L_b} \right) \]  

(15)

where \( \xi = x_0 / L_a, \eta = y_0 / L_b \) is the non-dimensional position of the attached nanoparticle. According to the Equation (15), the natural frequency of GSs can be given as \( f = \omega / 2\pi \).
2.3. Dynamic Analysis for Double-Layer Graphene Sheet (DLGSs)

In the analysis for DLGSs with attached nanoparticles, the governing equations are given by the two coupled equations

\[
D \nabla^4 w_1 + \left[ ph + m_c \delta(x - \xi) \delta(x - \eta) \right] \frac{\partial^2 w_1}{\partial t^2} - N^2 \nabla^2 w_1 = c(w_2 - w_1) \\
D \nabla^4 w_2 + ph \frac{\partial^2 w_2}{\partial t^2} - N^2 \nabla^2 w_1 = c(w_1 - w_2)
\]

(16)

where \( w_j(x, y, t), j = 1, 2 \), are the flexural deflections of the upper sheet \((j = 1)\) and lower sheet \((j = 2)\). \( c \) is the vdW interaction coefficient between the upper and lower layers, which can be obtained from the Lennard-Jones pair potential \([20]\), given as:

\[
c = b \left( \frac{4\sqrt{3}}{3w_0} \right)^2 2^{4/3} \left( \frac{\sigma}{\varepsilon} \right)^8 \left[ \frac{3003\pi}{256} \sum_{k=0}^{5} \frac{(-1)^k}{2k+1} \left( \frac{5}{k} \right) \left( \frac{c}{a} \right)^6 \frac{1}{(z_1 - z_2)^2} \right]
\]

(17)

where \( \varepsilon = 2.968 \text{ meV} \) and \( \sigma = 0.34 \text{ nm} \) are parameters chosen to fit the physical properties of GSs. \( z_j = z_j/a_i, (j = 1, 2) \), where \( z_j \) is the coordinate of the \( j \)th layer in the direction of thickness with the origin at the midplane of the GSs, and \( a = 1.42 \text{ nm} \) is the C–C bond length.

To obtain the vibration frequency for the governing equations Equation (16), we can introduce the following substitution

\[
w_j(x, y, t) = Y_j(x, y)e^{i\omega t}, \quad j = 1, 2
\]

(18)

where \( Y_j(x, y), j = 1, 2 \) is the shape function of deflection in the upper and the lower sheets, whereas \( \omega \) is the resonant frequency of the DLGS sensor.

Substituting Equation (18) into Equation (16), the coupled governing equations of the vibration in DLGSs are written in the following matrix form:

\[
\begin{bmatrix}
D \nabla^4 - N^2 \nabla^2 + c - \mu(x, y, t)\omega^2 \\
-c
\end{bmatrix}
\begin{bmatrix}
Y_1 \\
Y_2
\end{bmatrix}
= \begin{bmatrix}
0 \\
0
\end{bmatrix}
\]

(19)

and

\[
\mu(x, y) = ph + m_c \delta(x - x_0) \delta(y - y_0)
\]

(20)

Algebraic manipulation of Equation (20) reduces it to a single equation, which is

\[
\nabla^8 Y - \frac{2N^2}{D^2} \nabla^6 Y + \left[ \frac{2c - \rho \omega^2 - \mu(x, y, t)\omega^2}{D} + \left( \frac{N^2}{D} \right)^2 \right] \nabla^4 Y + \frac{\rho \delta n(x, y, t)\omega^4 - \mu(x, y, t)\omega^4}{D^2} Y = 0
\]

(21)

where \( Y = Y_1 \), or \( Y = Y_2 \).

For simply supported boundary condition, we can obtain the following frequency equation based on the same method with single-layer GSs.

\[
r_0\omega^4 - r_1\omega^2 + r_2 = 0
\]

(22)

and

\[
r_0 = \left( \frac{ph}{D^2} \right)^2 + \frac{4ph}{L_t d_t D^2} m_c \sin^2 m \pi \xi \sin^2 n \pi \eta
\]
\[
r_1 = \left( \frac{\pi^2}{L_t^2} \left( \frac{w_0^2}{L_x^2} + \frac{w_0^2}{L_y^2} \right)^2 + \frac{N^2}{D^2} \pi^2 \left( \frac{w_0^2}{L_x^2} + \frac{w_0^2}{L_y^2} \right) + \frac{c}{D} \right) \left( 2ph + \frac{4n \pi \xi m_t \sin^2 m \pi \xi \sin^2 n \pi \eta \right)
\]
\[
r_2 = \pi^2 \left( \frac{w_0^2}{L_x^2} + \frac{w_0^2}{L_y^2} \right)^4 + \frac{2N^2}{D^2} \pi^2 \left( \frac{w_0^2}{L_x^2} + \frac{w_0^2}{L_y^2} \right)^3 + \frac{2D + (N^2)^2}{D^2} \pi^4 \left( \frac{w_0^2}{L_x^2} + \frac{w_0^2}{L_y^2} \right)^2 + \frac{2N^2}{D^2} \pi^2 \left( \frac{w_0^2}{L_x^2} + \frac{w_0^2}{L_y^2} \right)
\]

(23)
The solution of Equation (22) yields the sought natural frequency of DLGSs. The anti-phase mode of the sought natural frequency, in which the deflection of the upper and lower layers occurs in the opposite direction, is obtained from:

$$\omega^2 = \frac{r_1 + \sqrt{r_2^2 - 4r_0^2}}{2r_0}$$  \hspace{1cm} (24)

3. Results and Discussion

In this simulation, GSs used as the nanomechanical resonator are considered to be simply supported. The vibration mode takes the fundamental frequency $m = n = 1$, and the anti-phase vibration for the DLGSs, in which the deflection of the upper and lower layers occurs in the opposite direction. In order to investigate the vibrational behavior of a GS mass sensor, the geometrical dimensions and the material constants are given as follows: The effective thickness $h$ of each layer of GS was 0.127 nm. The aspect ratio of GSs is 20, which is defined as the side length to the thickness. The Young’s modulus $E$ and the density $\rho$ of GSs were 2.81 TPa and 2300 kg/m$^3$ [35]. The coefficient of thermal expansion of GSs was $2.2 \times 10^{-6}$/K [2].

The Young’s modulus and the thickness of GSs from the literature [35] are given in Table 1, and the frequency shifts of GSs with attached nanomass of 100 zg (1 zg = $10^{-24}$ kg are obtained based on the proposed analytical mode.

<table>
<thead>
<tr>
<th>Authors</th>
<th>Young’s Modulus ($E$ (TPa))</th>
<th>Thickness ($t$ (nm))</th>
<th>Frequency Shift * ($\Delta \omega_1$ (GHz))</th>
<th>Frequency Shift ** ($\Delta \omega_2$ (GHz))</th>
<th>$\Delta \omega_2 - \Delta \omega_1$</th>
<th>Variation Error</th>
</tr>
</thead>
<tbody>
<tr>
<td>Van Lier et al.</td>
<td>1.11</td>
<td>0.34</td>
<td>113.0</td>
<td>115.11</td>
<td>2.21%</td>
<td></td>
</tr>
<tr>
<td>Zhao et al.</td>
<td>106.8</td>
<td>0.33</td>
<td>106.8</td>
<td>109.25</td>
<td>2.29%</td>
<td></td>
</tr>
<tr>
<td>Natsuki et al.</td>
<td>1.06</td>
<td>0.34</td>
<td>110.4</td>
<td>112.87</td>
<td>2.24%</td>
<td></td>
</tr>
<tr>
<td>Meo et al.</td>
<td>0.945</td>
<td>0.34</td>
<td>104.2</td>
<td>106.57</td>
<td>2.24%</td>
<td></td>
</tr>
<tr>
<td>Duan et al.</td>
<td>6.88</td>
<td>0.052</td>
<td>68.44</td>
<td>117.14</td>
<td>7.11%</td>
<td></td>
</tr>
<tr>
<td>Shi et al.</td>
<td>2.81</td>
<td>0.172</td>
<td>113.8</td>
<td>123.47</td>
<td>8.45%</td>
<td></td>
</tr>
</tbody>
</table>

* The frequency shift at room temperature; ** The frequency shift at temperature of 5 K.

Here, the frequency shift is defined as the difference between the natural frequency of a GS with and without attached nanoparticles, that is $\Delta \omega = \Delta \omega(\omega) - \Delta \omega(\omega + m_c)$. It can be observed that the frequency shifts of GSs with attached nanoparticles are larger for the temperature dependence. The variation errors of frequency shifts are about 2% when the temperature change is only 5 K, especially if the error value increases up to 8.45 % at the Young’s modulus (6.88 TPa) and there is a small thickness of around 0.052 nm of the GS. This suggests that the temperature dependence of the frequency shift is affected largely by the Young’s modulus and the thickness of GSs.

Figure 2 shows the location dependence of attached nanoparticles on the frequency shift of the SLGSs as a function of the attached mass under the temperature difference of 100 K. The attached mass is located at the center ($\zeta$, $\eta$) = (0.5, 0.5) of the GSs; near to the corner ($\zeta$, $\eta$) = (0.25, 0.25); and near to the edge ($\zeta$, $\eta$) = (0.5, 0.25). The frequency shift value of GSs increases with increasing attached mass because the attached nanoparticles increase the overall mass of the GS resonator. It can be deduced from Figure 2 that the location influence of the nanoparticles on the frequency shift is significant within a mass range from 10 to 1000 zg. The value of frequency shifts of GSs at the center load is larger than those away from the center location. This suggests that the sensitivity of the GS mass sensor is high when nanoparticles are located at the center of GSs.
The frequency shift value of GSs increases with increasing nonlocal parameter, and increases with the temperature difference. Figure 4 shows vibration frequency shift of GSs are investigated for an attached mass of 100 zg. In this simulation, the relationship between frequency shift and temperature difference is greater for a small mass than for a large mass. It is evident that the effect of temperature difference on the frequency shift is greater for a small mass than for a large mass.

As shown in Figure 3, the effects of temperature difference and nonlocal parameter on the vibration frequency shift of GSs are investigated for an attached mass of 100 zg. In this simulation, we assume that the nonlocal parameter \( (\xi, \eta) \) is 0, 2.0 and 4.0 nm [13]. It is found that the nonlocal parameter affects the vibrational frequency shift of GSs. The value of the frequency shift decreases with increasing nonlocal parameter, and increases with the temperature difference. Figure 4 shows the relationship between frequency shift and temperature difference. It is evident that the effect of temperature difference on the frequency shift is greater for a small mass than for a large mass.
When the total mass attached to the GSs is less than 1.0 zg.

According to this simulation, the relationship between the frequency shift and the attached mass can be well represented by the exponential function $\Delta \omega = \alpha m_e^\beta$, which has the double logarithmic linear relationship. The values of the parameters $a$ and $\beta$ can be determined easily by fitting the simulated curve and to obtain high accuracy calibration between frequency shift and attached mass. A logarithmically linear relationship is found between the frequency shift and the attached mass when the total mass attached to the GSs is less than 1.0 zg.

Figure 4. Variation of the vibration frequency shift with the temperature difference for different nanomass sizes ($e_{\vartheta a} = 2.0$ nm, $\xi = \eta = 0.5$).

Figure 5 shows a comparison of the frequency shift between SLGSs and DLGSs with the temperature difference of 100 K. The frequency shift of the GS resonator increases with increasing mass of nanoparticles. The DLGSs used as a nanomechanical resonator can provide higher sensitivity than the SLGSs due to large change in the vibrational frequency shift. It can be found that the frequency shift versus the attached mass show a nearly logarithmic linear relationship within a certain range. According to this simulation, the relationship between the frequency shift ($\Delta \omega$) and the attached mass ($m_e$) can be well represented by the exponential function $\Delta \omega = \alpha m_e^\beta$, which has the double logarithmic linear relationship. The values of the parameters $a$ and $\beta$ can be determined easily by fitting the simulated curve and to obtain high accuracy calibration between frequency shift and attached mass. A logarithmically linear relationship is found between the frequency shift and the attached mass when the total mass attached to the GSs is less than 1.0 zg.

Figure 5. Comparison of the frequency shift between DLGSs and SLGSs with attached nanoparticle for the temperature difference of $\Delta T = 100$ K ($e_{\vartheta a} = 0$ nm, $\xi = \eta = 0.5$).
4. Conclusions

In this study, we propose an analytical model to investigate the vibration frequency of simply supported graphene sheets (GSs) using the nonlocal elasticity theory. The relationship between the vibrational frequency of GSs and the attached mass is derived analytically, and the influences of the position of the nanoparticles and the mass on the frequency shifts of GSs are analyzed in detail. Because the mass of a GS is quite small, the frequency shifts can easily change when a relatively small mass of a nanoparticle is added to the GS. Based on the simulation, the results indicate the resolution of a mass sensor could achieve an order of 100 zg. The result demonstrates that the temperature and the nonlocal parameter affect the vibration frequency shift of the GSs. Moreover, the double-layer graphene sheets (DLGSs) used as a nanomechanical mass sensor provide higher sensitivity than the SLGSs. In this simulation, the results show that the Young’s modulus and the thickness of the GS affect temperature dependence of frequency shifts, especially the error value which increases for high Young’s modulus and small thickness. It is evident that the variation errors of frequency shifts are about 2% when the temperature change is only 5 K. The simulation technology and the parameters of investigation, such as temperature, nonlocal coefficient, geometric and material parameters, are thus very useful in designing GSs used as nano-scale sensing.

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