Numerical Study on Mass Sensitivity of Magnetoelastic Biosensors with Concentrated Mass Load under Different Resonance Modes

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1. Introduction

With the advantages of wireless in detection, easiness in operation, and low cost, magnetoelastic biosensors as an important type of acoustic wave devices have been widely used for the detection of pathogens like Listeria monocytogenes [1, 2], Escherichia coli [1, 2], Staphylococcus aureus [2], Salmonella typhimurium [2–7], Bacillus anthracis spores [7, 8], Staphylococcus epidermidis [9], and other chemicals [10, 11]. A magnetoelastic biosensor is typically made of a free-standing rectangular magnetoelastic strip immobilized with bioreceptors on its surface. Due to magnetostriction effect, a magnetoelastic sensor can be excited to vibrate by applying an alternative magnetic field. The working principle of a magnetoelastic biosensor is based on the shift in resonance frequency due to the mass change caused by the specific binding of bioreceptor target on the sensor surface. Therefore, a magnetoelastic biosensor is actually a mass-detection device. For such a device, a key parameter to evaluate its performance is mass sensitivity ($S_m$) which is defined as the change in the resonance frequency per unit mass load. Mass sensitivity can be well determined if the targets (i.e., mass load) are uniformly immobilized on the whole sensor surface [1]. This kind of immobilization mode has been widely used for magnetoelastic biosensors in various applications [1–11]. However, recent study has shown that such immobilization mode is not the best way to obtain the optimized mass sensitivity due to the location dependence of mass sensitivity [12–14].

Ramasamy et al. [12] experimentally found that the resonance frequency of a magnetoelastic sensor changed with the concentrated mass position. Based on the simulation results, equations for determining the maximum and minimum mass sensitivity were established. Li and Cheng [13] did the experiments for studying the resonance frequency of a magnetoelastic sensor attached with uniform mass load at different locations under different resonance modes. The experimental results indicated that the resonance frequency not only changed with mass load position but also changed...
Table 1: The properties of the sensor material and selected loading positions for this study [15, 16].

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Unit</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Young's modulus</td>
<td>( E )</td>
<td>GPa</td>
</tr>
<tr>
<td>Density</td>
<td>( \rho )</td>
<td>kg/m(^3)</td>
</tr>
<tr>
<td>Poisson’s ratio</td>
<td>( \nu )</td>
<td>—</td>
</tr>
<tr>
<td>Length</td>
<td>( l )</td>
<td>mm</td>
</tr>
<tr>
<td>Width</td>
<td>( w )</td>
<td>mm</td>
</tr>
<tr>
<td>Thickness</td>
<td>( h )</td>
<td>( \mu ) m</td>
</tr>
<tr>
<td>Loading position</td>
<td>( x_i/l )</td>
<td>—</td>
</tr>
</tbody>
</table>

...with resonance modes. They also found that when the mass load was attached on the node of a magnetoelastic sensor, the resonance frequency did not change. That is, mass sensitivity is zero when mass is loaded on the node. In our previous work, a vibration governing equation of a magnetoelastic sensor attached with a concentrated mass for the first resonance mode was established based on mechanical vibration theory [14]. The node position, resonance frequency, and thus mass sensitivity with a given loading position were mathematically calculated. In this work, the behavior of mass sensitivity of a magnetoelastic sensor under higher resonance modes was further investigated and the magnitude of load mass on mass sensitivity with given loading positions were mathematically calculated. In this study, the loading positions are listed in Table 1. The sensor size of this size can work well for real biodetection.

2. Materials and Methods

Figure 1 shows a magnetoelastic biosensor with a concentrated mass attached on its surface. Here, the commercially available soft magnetic alloy Metglas\textsuperscript{\textregistered} 2826 MB is selected as the sensor material and its material properties as well as the loading positions are listed in Table 1. The sensor size is 1 mm \( \times \) 0.2 mm \( \times \) 15 \( \mu \)m which is the same as that in the published paper regarding magnetoelastic biosensors [8]. The experimental results have proved that the magnetoelastic sensor of this size can work well for real biodetection.

3. Theory and Calculation

In real applications for biodetection, an alternative magnetic field is applied along the length direction of a magnetoelastic sensor, so that the sensor is driven to vibrate longitudinally (i.e., vibrating along its length direction). In this study, the sensor is assumed to vibrate longitudinally along \( x \)-axis, and its kinetic energy (\( T \)) and potential energy (\( V \)) can be expressed as follows:

\[
T = \frac{1}{2} \int_0^l \rho_s A_s \left( \frac{\partial u(x,t)}{\partial t} \right)^2 dx + \frac{1}{2} m_k \left( \frac{\partial u(x,t)}{\partial t} \right)^2_{x=x_i}, \quad (1)
\]

\[
V = \frac{1}{2} \int_0^l E \left( \frac{\partial u(x,t)}{\partial x} \right)^2 dx, \quad (2)
\]

where \( \rho, E, \nu \), and \( A_s \) represent the density, Young’s modulus, Poisson’s ratio, and cross-sectional area \( (w \times h) \) of the sensor, respectively, \( m_k \) represents the loading mass, \( u(x, t) = \varphi_n(x)q(t) \), where \( u(x, t) = [u_1(x, t), u_2(x, t), \ldots, u_n(x, t)] \), \( u_n(x, t) \) is the displacement of point \( x \) at time \( t \) for the \( n \)-th order resonance, \( \varphi_n(x) = [\varphi_1(x), \varphi_2(x), \ldots, \varphi_n(x)] \), where \( \varphi_n(x) \) is the mode shape function for the \( n \)-th order resonance, \( q(t) \) is the generalized coordinate which is a \( n \times n \) matrix, and \( u_n(x, t) = \varphi_1(x)q_1(t) + \varphi_2(x)q_2(t) + \cdots + \varphi_n(x)q_n(t) \). Theoretically, the potential energy \( V \) in (2) consists of the potential energy of the sensor itself and that of biomolecules which depend on Young’s modulus of the sensor material and biomolecules, respectively. Considering that Young’s modulus of biomolecules is greatly less than that of the sensor material, the contribution of biomolecules to the potential energy is neglected in this study.

Equations (1) and (2) can be converted into

\[
T = \frac{1}{2} \dot{q}^T (M_0 + M_1) \dot{q} = \frac{1}{2} \dot{q}^T M \dot{q},
\]

\[
V = \frac{1}{2} q^T K q. \quad (3)
\]

where \( K \) is the stiffness matrix in which the element \( K_{ij} = \int_0^l (E/(1-\nu))A_s \varphi_i'(x)\varphi_j'(x) dx \), \( M_0 \) is the mass matrix for the sensor in which the element \( M_{0ij} = \int_0^l \rho_s A_s \varphi_i(x)\varphi_j(x) dx \), \( M_1 \) is the mass matrix for the concentrated mass in which the element \( M_{1ij} = m_i \varphi_i(x)\varphi_j(x) \), and \( M \) is the total mass matrix in which the element \( M_{ij} = M_{0ij} + M_{1ij} \).

It is known that \( \varphi_n = \cos(n \pi x/l) \) \( (n = 1, 2, 3, \ldots) \) satisfies the boundary conditions of the sensor and thus \( \varphi_n \) is selected as the shape function for the \( n \)-th order resonance.
By applying Lagrange’s equation \((d/dt)(\partial L/\partial \dot{q}) - (\partial L/\partial q) = 0\) (where \(L = T - V\)), the governing vibration equation of the sensor is derived as

\[
(K - \omega_n^2 M) \dot{q} = 0. \tag{4}
\]

By solving (4), the generalized eigenvalue \(\omega_n\) and \(q(t)\) are obtained. The \(n\)th-order resonance frequency is then obtained by \(f_n = \omega_n/2\pi\). The mass sensitivity \(S_{m,n}\) for the \(n\)th-order resonance is determined by

\[
S_{m,n} = \frac{f_{n,0} - f_{n,m}}{m^n}, \tag{5}
\]

where \(f_{n,0}\) and \(f_{n,m}\) represent the \(n\)th-order resonance frequency of the sensor without and with mass load. The point displacement for the \(n\)th-order resonance is obtained by \(u_n(x, t) = \phi_1(x)q_{m,1}(t) + \phi_2(x)q_{m,2}(t) + \cdots + \phi_n(x)q_{m,n}(t)\). All the calculations are done by MATLAB software.

4. Results and Discussion

From the above equations, we can see that the resonance frequency \(f_n\) and mass sensitivity are dependent on the total mass matrix \(M\). Since \(M\) is the function of loading position \(x\) and concentrated mass \(m_\xi\), different loading positions and loading mass would result in different mass sensitivities. Figure 2(a) shows mass sensitivity \((S_m)\) as the function of loading position \((x_\xi)\) under different order resonance modes. Clearly, all the data curves exhibit periodic variation and there are \(n + 1\) loading positions \((x/l = m/n, \text{where } n = 1, 2, 3,\ldots\text{ and } m = 0, 1, 2,\ldots, n)\) corresponding to the maximum \((S_{m,\text{max}})\) and \(n\) loading positions \((x/l = (2m - 1)/2n, \text{where } n = 1, 2, 3,\ldots\text{ and } m = 1, 2,\ldots, n)\) corresponding to the minimum \((S_m = 0)\) for the \(n\)th-order resonance. In addition, it is found that the data points in Figure 2(a) are well fitted by \(S_m = A \cos^2 t\) (A is a constant) as presented by the dotted lines. That is, mass sensitivity is the function of only one independent parameter \(t\) and is linearly proportional to \(\cos^2 t\). Interestingly, it is found that \(u^2\) as the function of \(x_\xi\) as shown in Figure 2(b) shares the same variation trend as that of the curve in Figure 2(a) for the same order resonance mode. Therefore, it is believed that \(|\cos t|\) is linearly proportional to \(|u|\). In other words, \(S_m\) is linearly proportional to \(u^2\) at the point where the mass is loaded (i.e., \(S_m \propto u^2\)).

When the concentrated mass is uniformly loaded on the whole sensor surface, the mass sensitivity \((S_{m,u})\) can be determined by the well-known equation [1]:

\[
S_{m,u} = \frac{f_n}{2m_s}, \tag{6}
\]

where \(f_n\) is the \(n\)th-order resonance frequency which is expressed as [15]

\[
f_n = \frac{n}{2l} \sqrt{\frac{E}{\rho (1 - \nu)}}. \tag{7}
\]

Figure 3 compares the plots of \(S_{m,u}\) and the maximum mass sensitivity \(S_{m,\text{max}}\) in Figure 2(a) as the function of resonance order. Clearly, both curves show a linear relationship. In addition, it is found that \(S_{m,\text{max}}/S_{m,u} = 2\) for the same resonance mode. Based on the aforementioned theory (i.e., \(S_m \propto u^2\)), \(S_{m,u}\) should be related to \(u^2\) of all the points with mass load but we do not know what the relationship is. Now we assume the concentrated mass \((m_\xi = m_s/1000)\) is evenly distributed on 101 positions \((i.e., x_\xi/l = 0, 0.01, 0.02,\ldots, 1)\) on the sensor. We find that \(u_{\text{max}}^2/(\sum_{i=1}^{101} u_i^2/101)\) is very close to that of \(S_{m,\text{max}}/S_{m,u} = 2\). Therefore, it is believed that mass sensitivity for uniform mass load is the average contribution of \(u^2\) from all the loading points; that is, \(S_{m,u} \propto \sum_{i=1}^{n} u_i^2 / n\).
To investigate the effect of magnitude of mass load on mass sensitivity, mass sensitivities under six different loading masses (i.e., \( m_{c0} = m_s/1000 \), \( m_{c1} = m_s/900 \), ... , \( m_{c5} = m_s/500 \)) are calculated and compared. Here \( S_{m0}, S_{m1}, ..., S_{m5} \) are the mass sensitivities corresponding to the above six loading masses, respectively, and \( \Delta S_m = S_{m,n} - S_{m,0} \) and \( \Delta M = m_{c,n} - m_{c0} \) \((n = 1, 2, 3, ..., 5)\). Figure 4 shows the plot of \( \Delta S_m \) as the function of \( \Delta M \) for the first-order resonance.

It is found that \( \Delta S_m \) can be either negative or positive or zero depending on the loading position. When the loading position satisfies \( 0 \leq x_c/l < 0.3 \) or \( 0.7 < x_c/l \leq 1 \), \( \Delta S_m \) is negative and decreases as the loading mass increases which indicates that \( S_m \) decreases with loading mass increasing. The opposite trend is observed when \( 0.3 < x_c/l < 0.5 \) or \( 0.5 < x_c/l \leq 0.7 \). When the concentrated mass is loaded at the nodal point (i.e., \( x_c/l = 0.5 \)), \( \Delta S_m \) keeps zero no matter how the mass is changed. The above behaviors again can be confirmed by \( S_m \propto u^2 \). Here we compare \( u^2 \) at \( x/l = 0.2 \) \((0 \leq x_c/l < 0.3)\) and \( x/l = 0.4 \) \((0.3 < x_c/l < 0.5)\) where two different masses \( (m_c = m_s/2 \text{ and } m_c = m_s) \) are loaded on the two positions, respectively, as shown in Figure 5. We can see that \( u^2 \) at the loading position \( x_c/l = 0.2 \) decreases as \( m_c \) increases while \( u^2 \) increases as \( m_c \) increases at the loading position \( x_c/l = 0.4 \) which agrees with \( S_m \propto u^2 \). For the mass loaded at the nodal point \( (x_c/l = 0.5)\), the plots of \( u^2 \) versus \( x/l \) are exactly the same as that of the one for the sensor without loading mass.

That is, \( u^2 \) at \( x/l = 0.5 \) is always zero no matter how the load mass changes at \( x/l = 0.5 \) which again is consistent with \( S_m \propto u^2 \).
5. Implications

The theoretical results provide important guidance on the immobilization patterns of bioreceptors and targets on a magnetoelastic sensor for biodetection, particularly for single-target detection. For example, a single bacterial cell can be treated as a concentrated mass load. In such case, the position of the cell attached on the sensor would be very critical to mass sensitivity. In the worst case, that is, the cell attached on the node, there would be no frequency shift and thus a false result would be obtained. Based on the simulation results, we can conclude that the maximum mass sensitivity for the first resonance mode is obtained when the cell is attached on either end of the sensor. That means that the bioreceptor should be immobilized on the two ends of the sensor to ensure that the single target is attached on either end.

6. Conclusion

In this work, we established a governing equation of a magnetoelastic sensor attached with a concentrated mass under different resonance modes. Based on the results and analysis, several conclusions were drawn as follows:

(1) Mass sensitivity as the function of loading position $x_c$ can be well fitted by $S_m = A \cos^2 t$ and there are $n + 1$ loading positions corresponding to the maximum and $n$ loading positions corresponding to the minimum ($S_m = 0$) for the $n$th-order resonance.

(2) For the first resonance mode, when the mass load position $x_c$ satisfies $0 \leq x_c/l \leq 0.3$ or $0.7 < x_c/l \leq 1$, mass sensitivity decreases as loading mass increases while the opposite trend is obtained when $0.3 < x_c/l < 0.5$ or $0.5 < x_c/l \leq 0.7$. Mass sensitivity does not change when concentrated mass is loaded at the nodal point ($x_c/l = 0.5$).

(3) Mass sensitivity comes from the average contribution of square of displacement ($\mu^2$) of all the loading positions; that is, $S_{mu} \propto \sum_{i=1}^{n} \mu_i^2 / n$.

Competing Interests

The authors declare no conflict of interests.

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References


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