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Optical Whispering-Gallery-Mode Microbubble Sensors

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Abstract: Whispering-gallery-mode (WGM) microbubble resonators are ideal optical sensors due to their high quality factor, small mode volume, high optical energy density, and geometry/design/structure (i.e., hollow microfluidic channels). When used in combination with microfluidic technologies, WGM microbubble resonators can be applied in chemical and biological sensing due to strong light-matter interactions. The detection of ultra-low concentrations over a large dynamic range is possible due to their high sensitivity, which has significance for environmental monitoring and applications in life-science. Furthermore, WGM microbubble resonators have also been widely used for physical sensing, such as to detect changes in temperature, stress, pressure, flow rate, magnetic field and ultrasound. In this article, we systematically review and summarize the sensing mechanisms, fabrication and packing methods, and various applications of optofluidic WGM microbubble resonators. The challenges of rapid production and practical applications of WGM microbubble resonators are also discussed.

Keywords: whispering-gallery-mode microcavity; optical microbubble resonator; sensors; microfluidic technology

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1. Introduction

The invention of lasers in the 1960s promoted the rapid development of different types of optical sensors. Optical sensors have had a significant impact based on their ability to directly convert detected changes into the information of the light intensity, wavelength, polarization and phase with a high sensitivity and a fast response time. Various types of optical sensors, such as surface plasmon resonance (SPR) [1,2], planar waveguides [3,4], optical fiber sensors [5,6], optical interferometers [7,8] and optical microcavities [9,10] have been studied. Optical microcavities are miniature optical resonators at the micrometer or sub-micrometer scale, which confine the photons to a small space through the reflection or diffraction effects and are flexible for the integration in lab-on-chip systems.

Optical microcavities can be categorized according to the constraint mechanism of the light as Fabry-Pérot (F-P) microcavities [11–14], photonic crystal microcavities [15–20] and whispering-gallery-mode (WGM) [21–24] microcavities, as shown in Figure 1. Among these, WGM microcavities have been widely used in lasers, sensors, nonlinear optics, optomechanical and optical trapping due to their high quality factor (Q, as high as 10^{11}), extremely small mode volumes, and high optical energy density [9,25–34]. WGM was discovered in the field of acoustics: during the twentieth century, Lord Rayleigh discovered that sound waves could propagate inside the corridor in St. Paul's Cathedral [35]. The same phenomenon was discovered in the field of optics when light meets the conditions of total internal reflection and constructive interference on a closed concave surface, as shown in Figure 1c.

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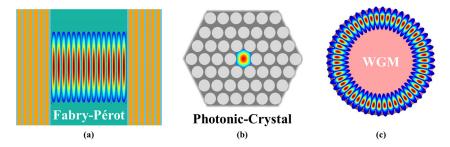


Figure 1. Schematic of the **(a)** Fabry–Pérot, **(b)** photonic crystal, and **(c)** WGM optical microcavities.

The optical WGM microcavities studied thus far include microrings [36-40], microtoroids [41-46], microdisks [47-52], microcylinders [53-55], microcapillaries [56-58], microspheres [59-62], microbottles and microbubbles [63-73], as shown in Figure 2. For different microcavity morphologies, the microcapillaries, microbottles and microbubbles have attracted attentions due to their natural optofluidic channels, which was always combined with the microfluidic technology for analyte delivery [74-83]. In 2005, the optical characteristics of microbottle resonators were studied theoretically by Louyer et al., who demonstrated that the microbottle resonators sustain modes with enhanced field strength due to the confinement of light between two turning points. This results in the superior optical characteristics of the microbottle resonators compared with the microsphere resonators [84,85]. Five years later, the first microbubble resonator was fabricated with a silica microcapillary by Sumetsky et al. [86]. In the same year, they also demonstrated that the tunable bandwidth of a silica microbubble resonator was approximately two times of its azimuthal free spectral range (FSR) [87]. The favorable optical characteristics of the microbubble resonators, such as their high Q-factor and small mode volume, have made them a focus of research in the studying of various sensors.

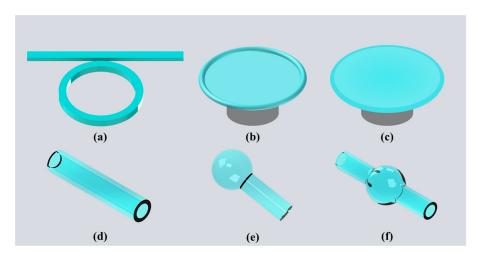


Figure 2. Types of WGM optical microcavities: (a) microring cavity; (b) microtoroid cavity; (c) microdisk cavity; (d) microcapillary cavity; (e) microsphere cavity; (f) microbubble cavity.

In this paper, we systematically review and summarize the information on optofluidic microbubble resonators, including their optical characteristics and sensing mechanisms, the fabrication and packaging methods and various sensing applications. Firstly, different sensing mechanisms of the microbubble resonators are introduced in Section 2. We summarize the methods for fabricating and packaging the microbubble resonators in Section 3. In Section 4, we provide an overview of the microbubble resonators, including their physical, chemical to biological sensing applications. Finally, a summary of the research status of the microbubble resonators and the further improvement outlooks are provided in Section 5.

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2. Optical Characteristics and Sensing Mechanism of a Microbubble Resonator

2.1. Optical Characteristics

The optical characteristics of a microbubble resonator are typically described according to their Q-factor, photon lifetime (τ), FSR and mode volume [88]. The Q-factor is related to the energy attenuation or the ability to constrain energy in the microcavity, which can be calculated as follows:

$$Q = \omega_0 \frac{U}{-dU/dt} = \frac{\omega_0}{\Delta \omega} = \frac{\lambda_0}{\Delta \lambda},$$
 (1)

where ω_0 is the central frequency of a resonance mode, U is the energy inside the microcavity, -dU/dt is the energy attenuation per unit of time, $\Delta\omega$ is the spectral linewidth for frequency, λ_0 is the central wavelength of a resonance mode, and $\Delta\lambda$ is the spectral linewidth for wavelength, which can be written as follows:

$$\Delta\omega = -2\pi \frac{c}{\lambda_0^2} \Delta\lambda,\tag{2}$$

where c is the speed of light in vacuum. The photon lifetime is defined as the time scale when the light intensity decays to 1/e of its initial value, which is caused by a depletion in the cavity. The photon lifetime can be written as follows:

$$\tau = \frac{Q}{\omega_0}.$$
 (3)

The photon lifetime increases with the increasing of *Q*-factor, which indicates a stronger interaction between the light and the analytes. The FSR is defined as the wavelength interval between two adjacent resonance modes, which can be written as follows:

$$FSR = \lambda_m - \lambda_{m-1} = \frac{\lambda_m^2}{2\pi n_{eff} R},$$
(4)

where n_{eff} is the effective refractive index (RI), and R is the radius of the microbubble.

The mode volume is the spatial distribution of a resonance mode in the cavity, which can be written as follows:

$$V = \frac{\oint \varepsilon(\vec{r}) |E(\vec{r})|^2 dV}{\max(\varepsilon(\vec{r}) |E(\vec{r})|^2)},$$
(5)

where $\varepsilon(r)$ is the spatial dielectric constant, E(r) is the spatial electric field strength, and V is the volume of the microcavity. The energy density increases as the mode volume decreases. Therefore, a smaller mode volume can lead to a stronger interaction between the light and the analytes.

2.2. Sensing Mechanism

Due to the extension of WGM fields into the surrounding analytes, any changes occurring around the evanescent field can lead to the changes of the resonance mode. Therefore, the sensing mechanism of WGM microbubble resonators are divided into the wavelength shift [89,90], mode-broadening [91,92], and mode-splitting [46,55,64].

2.2.1. Wavelength Shift

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WGM microbubble resonators are constituted of an inner air core and an outer spherical shell, which provides a fluidic channel for analyte delivery. Due to the total internal reflection, the light propagates along the azimuth of the microbubble wall, and it will resonate stably when it meets the constructive interference. The resonance wavelength λ can be written as follows [84]:

$$\lambda = \frac{2\pi n_{eff} R}{\sqrt{m^2 + m(2q+1)\Delta kR}},\tag{6}$$

where m and q represents the azimuth quantum number and axial quantum number, respectively, and Δk is the microbubble curvature parameter, which increases with the increases in the microbubble radius ($\Delta k = \frac{2}{L} \sqrt{(\frac{R}{R_a})^2 - 1}$) [84,88], as shown in Figure 3. For a zero-order axial mode that propagates along the microbubble equator (q = 0), due to $m^2 \gg m\Delta kR$, the resonance wavelength can be approximated as follows:

$$\lambda = \frac{2\pi n_{\text{eff}} R}{m}.$$
 (7)

The resonance wavelength is related to the effective RI and the microbubble radius, and it increases when either the effective RI or the microbubble radius increase. Therefore, it is possible to monitor the environmental disturbances by converting it into the changes of the RI and microbubble radius, such as concentration, temperature, pressure, and stress.

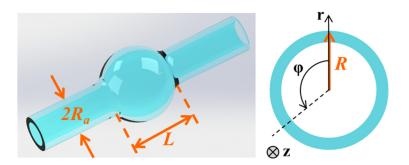


Figure 3. Schematic of an optical WGM microbubble resonator.

The effective RI changes with the changing of the surrounding medium, resulting in a wavelength shift. For different concentrations of the surrounding medium, the resonance wavelengths will also differ. Generally, the resonance wavelength red shifts with the increasing of the medium concentration, as shown in Figure 4a. However, the sensitivity of the microbubble resonator is low due to most of the mode field is constrained inside the microbubble wall. To achieve a higher sensitivity, the microbubble is usually corroded with hydrofluoric acid (HF), or a higher order radial mode is chosen for sensing [89,90,93]. Furthermore, the effective RI changes with the changing of the temperature [94,95] and magnetic field [96,97] due to the thermo-optical effect and magneto-optical effect, respectively. Changing of geometric shapes can also lead to a wavelength shift. For example, the microbubble size is easily changed by the pressure, flow rate, stress, tension, and thermal expansion [98–101]. By monitoring the wavelength shift, the changes in the external physical parameters are easily determined, as shown in Figure 4b. Note that the wall thickness of a microbubble resonator is dependent on its practical applications: a thin wall is always chosen for the detection of media inside the microbubble, while a thick wall is chosen for physical sensing, such as the temperature.

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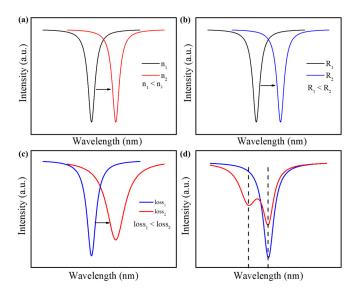


Figure 4. Sensing mechanism of the microbubble resonators: (a) wavelength shift with RI; (b) wavelength shift with microbubble radius; (c) mode-broadening; (d) mode-splitting.

2.2.2. Mode-Broadening

The spectral linewidth of a resonance mode is related to the *Q*-factor, as shown in Equation (1). Variations in the optical loss will result in different values of the *Q*-factor. Generally, the total *Q*-factor of a microbubble resonator can be written as follows [88]:

$$\frac{1}{Q_{Total}} = \frac{1}{Q_{Material}} + \frac{1}{Q_{Scattering}} + \frac{1}{Q_{Surface}} + \frac{1}{Q_{Radiation}} + \frac{1}{Q_{External}},$$
(8)

where $1/Q_{Material}$ is the absorption loss of the material; $1/Q_{Scattering}$ is the scattering loss caused by the surface roughness and other artificially introduced scattering points; $1/Q_{Surface}$ is the unclean surface loss; $1/Q_{Radiation}$ is the radiation loss due to the evanescent field leaks outside the microbubble wall; and $1/Q_{External}$ is the coupling loss due to the coupling between the cavity and the external coupling device. The spectral linewidth increases with increasing the cavity loss, such as the scattering loss and radiation loss [91,92], as shown in Figure 4c. Different from the wavelength shift, mode-broadening is not affected by the disruptions in external environment due to its self-referenced sensing mechanism. Therefore, it has smaller detection limits for the small molecules and single nanoparticles. However, the additional loss of the surrounding medium can significantly damage the microbubble surface and preclude its reuse.

2.2.3. Mode-Splitting

WGM microcavity has a degenerate state between clockwise- and counterclockwise-propagating resonance modes [23]. When a small molecule or nanoparticle enters the optical evanescent field, coupling between the counterpropagating modes is introduced due to the node and antinode of each mode at the molecules or nanoparticles location. Therefore, the eigenstates are transformed into two orthogonal standing waves in the cavity, as shown in Figure 4d. The size and number of scatters, such as the nanoparticles, can be determined by measuring the frequency-splitting [46,55,64]. Note that the two modes experience the same thermal noise. Therefore, disturbances in the surrounding environment and thermal fluctuations are eliminated by the self-referencing of the two modes. Compared with the mode-broadening, a cavity of the ultra-high *Q*-factor is required to differentiate the two modes in the splitting spectrum.

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3. Fabrication and Packaging Methods of a Microbubble Resonator

3.1. Fabrication Methods of a Microbubble Resonator

The fabrication of microbubbles is based on the fuse-and-blow method [86,102–105]. For melting of the silica microcapillaries, different methods, such as the electrode-discharge heating [102,103], oxyhydrogen-flame heating [104,105], and carbon dioxide laser heating [106], have been proposed. Differently from the carbon dioxide laser heating and oxyhydrogen-flame heating, the electrode-discharge heating is accomplished using a portable, low-cost fiber fusion splicer. A microbubble is fabricated through the following four steps: (I) sealing one end of the silica microcapillary via a high temperature while the other is left open to allow air to be pushed into the microcapillary, as shown in Figure 5(a2); (II) removing the outer polymer coating of the silica microcapillary at the intended location and then cleaning it until there is no residue, as shown in Figure 5(a3); (III) heating the position where the coating has been removed and pressing air into the silica microcapillary at the same time, as shown in Figure 5(a4); (IV) repeating the process of step (III) until a symmetrical and high-optical-quality microbubble has been fabricated, as shown in Figure 5(a5),(a6),b.

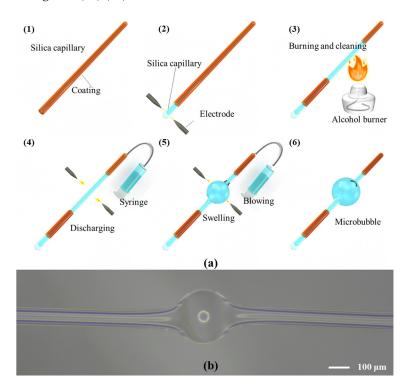


Figure 5. Fabrication of the microbubble resonators: (a) process for fabrication a microbubble resonator by the electrode discharge heating. (1) Silica microcapillary; (2) sealing one end of the silica microcapillary; (3) burning and cleaning the polymer coating; (4–6) fuse-and-blow process for microbubble fabrication; (b) optical microscopy image of a microbubble resonator.

To reduce the wall thickness of the microbubble to meet special experimental requirements, different methods have been proposed. One method involves corroding a silica microcapillary with HF solution before fabrication [89]. The wall thickness of a silica microcapillary can be controlled via the corrosion time. However, the internal surface smoothness of the microbubble may be reduced due to the uneven corrosion. Therefore, a new method by melting and tapering the silica microcapillary was proposed. The wall thickness of the silica microcapillary and its outer diameter were controlled by the tapering time [105].

In addition to the microbubble fabrication with a silica microcapillary, other new materials have also been used. For example, a gas microbubble was generated by locally

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heating a fiber tip, where a gold nanofilm had been coated [107]. To further improve the sensitivity, additional materials have been used for the microbubble fabrication, such as polymethylmethacrylate (PMMA)-based optical microbubbles for temperature sensing [108] and lead-silicate-based optical microbubbles for nonlinear optical effects [109], etc. [110–112].

3.2. Packaging Process of a Microbubble Resonator

The optical coupling between a microbubble resonator and the tapered fiber is easily affected by surrounding perturbations, such as the mechanical vibration and airflow. Therefore, noise levels limit its practical application as well as result in the poor detection of weak signals. To improve the stability while maintaining a high Q-factor for a long working period, specific packing schemes have been proposed [89-91,93]. As shown in Figure 6a, a glass scaffold was fabricated for a coupling system. Firstly, the tapered fiber was fixed on the glass scaffold, and two ports were further fixed with a low-RI polymer (e.g., MY 133) to reduce the vibration of the tapered fiber, as shown in Figure 6b,c. Then, the gap between the tapered fiber and the microbubble resonator was precisely controlled by 5D stages until the optimal coupling condition was realized, which was monitored through the transmission spectrum. Thirdly, the two ports of the microbubble were also fixed to the glass scaffold to reduce the coupling position change between the tapered fiber and the microbubble resonator, as shown in Figure 6d. Finally, the optical coupling region was wrapped with a low-RI polymer via the UV exposure, which further improved the stability of the coupling system, as shown in Figure 6e. A coverslip was placed on the glass scaffold to insulate the low-RI polymer from oxygen and accelerate its solidification, as shown in Figure 6f. Various coupling methods, such as the waveguide coupling, prism coupling, and fiber-tip coupling have been proposed for the WGM microcavity. However, the tapered fiber has typically been used for the microbubble resonators due to its high efficiency and ease of integration [113–116].

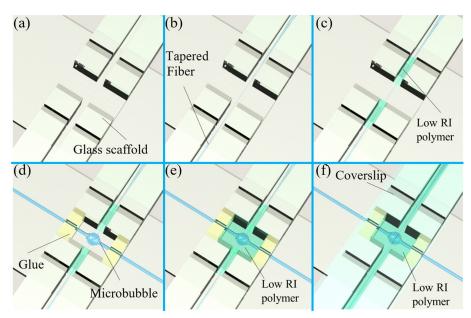


Figure 6. Packing process of a microbubble resonator: (a) a glass scaffold for holding the coupling system; (b,c) fixation of the tapered fiber; (d) fixation of the microbubble resonator; (e) wrapping the optical coupling region with low-RI polymer; (f) placement of a coverslip on the glass scaffold.

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4. Sensing Applications of Microbubble Resonators

4.1. Physical Sensing

4.1.1. Temperature Sensing

Due to the thermo-optical effect and thermal expansion, the RI and geometric shape are easily affected by temperature. Furthermore, the optical characteristics of a high Qfactor and a small mode volume imply a high sensitivity for the temperature sensing [94]. The thermal drift of two different cores of the microbubble, such as water and ethanol, were measured by Ward et al. in 2013 [95]. They found that the liquid core with a negative thermo-optical coefficient material can realize a high sensitivity of 100 GHz/K while effectively reducing the thermal drift. Furthermore, microbubbles based on polymethyl methacrylate (PMMA) material were proposed by He et al. in 2018 [108], as shown in Figure 7a. Different microbubble sizes were fabricated using a volume-controllable pipette, and a sensitivity of 39 pm/°C for a temperature range of 25 °C to 80 °C was obtained. To address the issues of the microbubble structure fragility and solution evaporation, a stable and encapsulated quasi-droplet microbubble was used for temperature sensing by Chen et al. in 2018 [117]. A high sensitivity of 205.3 pm/°C was experimentally achieved, as shown in Figure 7b. Optical WGM barcode technology based on the simultaneous measurement of multiple WGM modes was proposed by Liao et al. in 2021 [118] for temperature sensing, as shown in Figure 8a,b. Compared with monitoring the relative wavelength drift of a specific mode, the dynamic detection range was further increased, and the temperature was directly derived by searching the spectrum in the database. This method achieved a detection limit of 0.002 °C. Compared with other optical temperature sensors, the thermal drift can be reduced by introducing materials with negative thermo-optic coefficients inside the WGM microbubble resonators. Therefore, a high sensitivity and large dynamic detection ranges for the temperature sensing can be realized using WGM microbubble resonators.

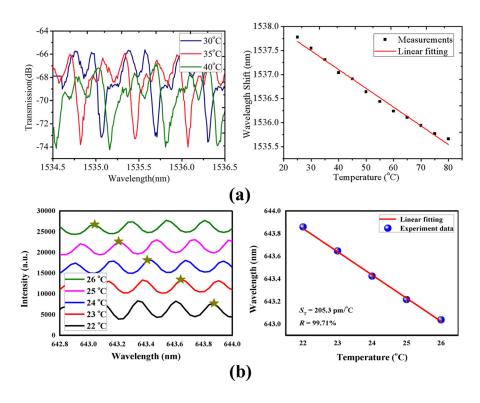


Figure 7. Temperature sensing with WGM microbubble resonators based on: (a) PMMA microbubbles (reprinted with permission from [108], © IOP Science), (b) encapsulated quasi-droplet microbubbles (reprinted with permission from [117], © CC BY license).

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4.1.2. Bulk RI and Liquid-Concentration Sensing

As the distribution of an evanescent field penetrates to the microbubble core, the changing of RI and concentrations can result in wavelength shifts. Therefore, bulk RI and liquid-concentration sensing can be accomplished through a combination of microfluidic technologies. Lab-on-fiber sensing technology based on gas microbubbles was proposed by Zhang et al. in 2018 [107], in which they locally heated a fiber tip with a gold nanomembrane, as shown in Figure 9a. The growth rate of the air microbubbles is dependent on the surrounding solution. For the evaporation of a solution, the growth rate of a microbubble decreases as the concentration increases due to the increase in boiling point. In addition, chemical decomposition can also lead to the generation of air microbubbles. The growth rate usually increases when the concentration increases, which is due to the increasing of decomposition rate. The dynamic detection range of the sucrose solutions from 0.5% to 50% and five orders of the magnitude (10-5~1 M) for hydrogen peroxide were realized. Such air microbubbles can also be repeatedly reused. However, they are easily affected by surrounding mechanical disturbances. To reduce the effect of environmental perturbance, a packaged microbubble tapered-fiber coupling system was proposed by Tang et al. in 2016 [89], as shown in Figure 9b. A type of low-RI polymer was used as the cladding layer. They found such a sensing system maintained a high Q-factor for a significant length of time and had a high signal-to-noise ratio. Furthermore, bulk-RI sensitivity of 18.8 nm/RIU and a detection limit of 5.4×10^{-5} RIU were experimentally achieved, as shown in Figure 9c. In 2018, this same research group proposed a new packaged microbubble tapered-fiber coupling system. The signal-to-noise ratio and stability were further increased, and the detection of a low concentrations of small molecules was realized [90]. The microfluidic channel in microbubble resonators provides advantages when used for RI-based and liquid-concentration-based sensing. In addition, ultra-low detection limits can be realized due to the high Q-factor. However, only a small part of the WGM field extends into the surrounding analytes, resulting in a low sensitivity of the RI sensing.

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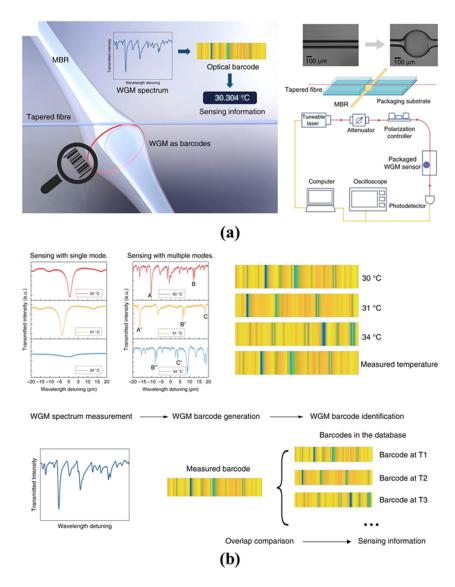


Figure 8. Temperature sensing with WGM microbubble resonators based on optical WGM barcode technology (reprinted with permission from [118], © CC BY license): (a) the sensing system of optical WGM barcode technology; (b) sensing principle of the optical WGM barcode technology.

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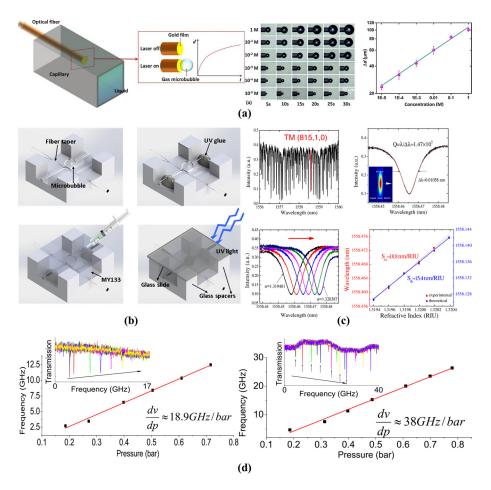


Figure 9. Liquid concentration and pressure sensing with the microbubble resonators: (a) a lab-on-fiber microbubble for liquid-concentration sensing (reprinted with permission from [107], © Elsevier); (b,c) a packaged microbubble for the liquid RI sensing (reprinted with permission from [89], © Optica, formerly the Optical Society of America); (d) pressure sensing with the microbubble resonators (reprinted with permission from [106], © Optica, formerly the Optical Society of America).

4.1.3. Pressure and Flow Rate Sensing

Due to the hollow microfluidic channel, an accurate measurement of the pressure and flow rate can be realized through a combination with microfluidic technologies. For different pressures inside the microbubble, the geometric shape is different [119]. By changing the pressure inside a microbubble, the resonance mode was tuned over hundreds of GHz [98]. To further improve the sensitivity, an ultra-thin microbubble wall (500 nm) was used for aerostatic pressure sensing by Yang et al. in 2016 [106], as shown in Figure 9d. A high sensitivity of 19 GHz/bar at 1.55 μ m, 38 GHz/bar at 0.78 μ m and a resolution of 0.17 mbar were realized. In addition, the microbubble wall thickness was precisely measured with a combination of pressure sensing by Lu et al. in 2016 [99]. The minimum measurement uncertainty of the microbubble wall thickness was approximately 0.02 μ m. The advantages of such pressure sensors are their simple structures and their low detection limits. However, the sensitivity is limited due to the large Young's modulus of the silica material. Therefore, new materials are needed for the fabrication of microbubble resonators aimed at pressure or flow rate sensing.

The pressure inside the microbubble decreases with increasing of velocity. Therefore, a microbubble resonator can also be used for the flow rate sensing. As shown in Figure 10a, a flow rate sensor whereby detection is based on the Bernoulli effect was proposed and experimentally proven by Chen et al. in 2019 [100]. By combination with a packaged microbubble resonator, a dynamic detection range of 10 μ L/min to 200 μ L/min and

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sensitivity of 0.0196 pm/(μ L/min) were realized. To further improve the sensitivity, a higher order radial mode and a thinner microbubble wall were applied for the flow rate sensing by Wang et al. in 2021 [101]. A high sensitivity of 0.079 pm/(μ L/min) and a dynamic detection range of 0 μ L/min to 200 μ L/min were realized. Moreover, the microbubble resonators were also used for air-coupled ultrasound sensing [120]. By the combination with an optical frequency comb, a femtometer resolution of the resonance wavelength shift and sub-microsecond response time were realized to trace the ultrasound pressure, as shown in Figure 10b. The noise equivalent pressure was 4.4 mPa/ μ Hz. Compared with other flow rate sensors, microbubble resonators provide a simple and flexible structure while maintaining a high sensitivity.

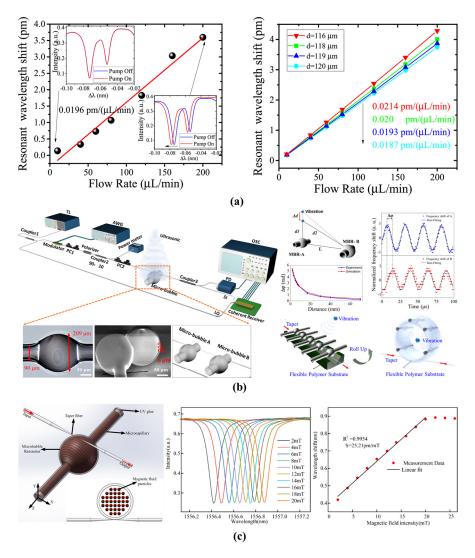


Figure 10. (a) A packaged microbubble resonator for flow rate sensing (reprinted with permission from [100] © Optica, formerly the Optical Society of America); (b) ultrasound sensing of a microbubble resonator (reprinted with permission from [120] © Chinese Laser Press); (c) magnetic field sensing of a magnetic-fluid-filled microbubble resonator (reprinted with permission from [97] © Elsevier).

4.1.4. Magnetic-Field Sensing

By coating with magnetic material, the microbubble resonators have been used for magnetic field sensing due to the magneto-optical effect [121]. Terfenol-D with a high magnetostriction coefficient was fixed to one side of the microbubble resonator [96]. When the strength of the surrounding magnetic field was altered, the longitudinal length and RI

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of microbubble were changed. A sensitivity of 0.081 pm/mT and a dynamic detection range of 0.41 mT to 21.8 mT were realized. To further increase the sensitivity, magnetic-fluid-filled microbubble resonators were proposed by Liu et al. in 2021 [97], as shown in Figure 10c. A high sensitivity of 25.21 pm/mT was realized. Due to their high *Q*-factor, small mode volume, and high optical energy density, microbubble resonators can provide the high sensitivity required to sense the magnetic-fields. Compared with other magnetic-field sensors, the microbubble resonators have the advantages of its small size and having high accuracy for magnetic-filed sensing.

4.2. Chemical Sensing

4.2.1. Gas Sensing

By modifying the inner surface of the microbubble with special materials, the detection of low concentrations of chemical gases becomes possible, especially for harmful gases. Due to the high Q-factor and the hollow microfluidic channel of the microbubble resonator, real-time and high sensitivity gas sensors were realized. As shown in Figure 11a, an approach that extended electron-photon interaction to electron-phonon-photon interaction in a graphene-deposited microbubble was proposed by Yao et al. in 2017 [122]. Ultra-sensitive and rapid detection of ammonia gas was realized with a noise equivalent limit of 1 ppb and a dynamic detection range of more than five orders of magnitude, as shown in Figure 11b. In addition, a carbon dioxide (CO₂) gas sensor developed by coating polyhexamethylenebiguanide (PHMB) inside the microbubble surface was proposed by Li et al. in 2019 [123], as shown in Figure 11c. When the CO₂ gas is delivered into the microbubble resonator, it interacts with the PHMB molecular layers, resulting in a blue shift in resonance wavelength. A sensitivity of 0.46 pm/ppm and a detection limit of 50 ppm in a range of 200 ppm to 700 ppm with a good selectivity was realized. Furthermore, a self-assembled polydimethylsiloxane (PDMS) microbubble was proposed for the detection of ethanol gas [111]. When the ethanol gas concentrations changes, the volume and RI of PDMS change accordingly, resulting in a resonance wavelength shift, as shown in Figure 11d. A sensitivity of 36.24 pm/ppm and a dynamic range of 4.19 ppm to 272.35 ppm were realized. However, those gas sensors were dependent on the internal surface coating and can only detect one medium.

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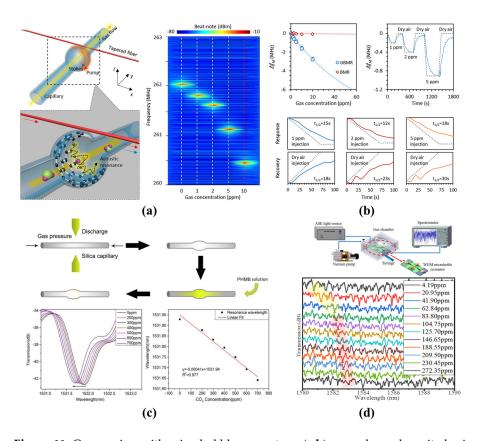


Figure 11. Gas sensing with microbubble resonators: (**a**,**b**) a graphene-deposited microbubble for the ammonia gas sensing (reprinted with permission from [122], © ACS Publications); (**c**) a PHMB-deposited microbubble for the CO₂ gas sensing (reprinted with permission from [123], © Optica, formerly the Optical Society of America); (**d**) a PDMS-deposited microbubble for the ethanol gas sensing (reprinted with permission from [111], © Elsevier).

To overcome the difficulty of surface coating and allow the detection of different gases at the same time, a method based on pressure-induced geometric deformation and gas-molecules-induced heat dissipation in the microbubble resonator was proposed by Peng et al. in 2020 [124]. When the flow rate increases, the resonance frequency red-shifts while the thermal effect decreases due to the heat dissipation. The different thermal conductivities of He, N₂, and CO₂ gases allowed them to be simultaneously distinguished. Using a combination of microbubble resonators, different chemical gases were detected with high sensitivity and low detection limits. In addition, the sensor size was further decreased. However, the complicated coating process has limited its practical application.

4.2.2. Ion and pH Sensing

The ultra-sensitive and real-time specificity detection of lead ions was proposed by Fu et al. in 2020 [125], as shown in Figure 12a. Preprocessing with the piranha solution and poly-L-lysine solutions results in the inner surface of the microbubbles being positively charged. The GR-5 DNAzyme and substrate strands were then successfully modified by electrostatic adsorption. When the lead ions solutions are delivered into the microbubble, they cleave the substrate strands, resulting in blue shifts of the resonance modes, as shown in Figure 12b. A detection limit of 15 fM and a dynamic detection range of 0.1 pM to 100 pM were realized. In addition, a pH-sensitive polymer, N-isopropylacrylamide (polyNIPA), was coated onto the inner surface for pH sensing by Stoian et al. in 2019 [126], as shown in Figure 12c,d. When changing the pH inside the microbubble, the polymer particles changed from a shrunken to a swollen state, resulting in a change in the RI. A response time of 10~15 s and a resolution of 0.06 pH were realized. The

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drawback of such a method is the large size of the polyNIPA particles, and it was not adapted to the thin-film-related theoretical models.

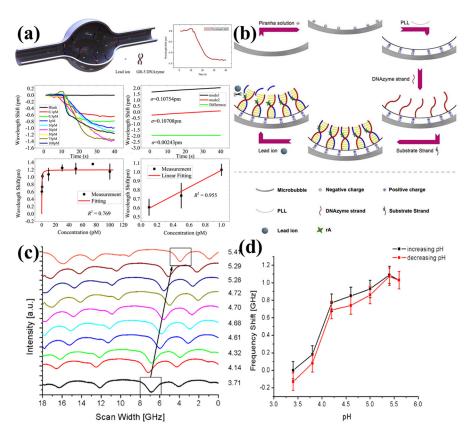


Figure 12. Sensing of (a,b) lead-ions concentrations with the microbubble resonators (reprinted with permission from [125], © Elsevier) and (c,d) pH values with the microbubble resonators (reprinted with permission from [126], © Elsevier).

4.2.3. Hydrogel Phase Sensing

Real-time monitoring of the internal structural changes of hydrogel was proposed by Yang et al. in 2020 [91,92,127]. The phase transition from hydrophilic to hydrophobic with changing of temperatures was studied, as shown in Figure 13a. By simultaneously monitoring the resonance wavelength shift, spectral linewidth broadening and microscope imaging, RI increase and light scattering enhancement were observed during the transition from hydrophilic to hydrophobic, as shown in Figure 13b. Compared with other methods, rapid and real-time monitoring of the phase transition dynamics of certain materials was achieved via the microbubble resonators. In addition, the cost was further decreased.

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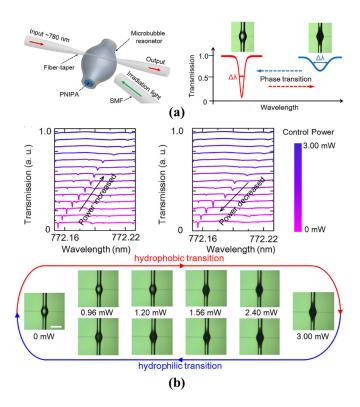


Figure 13. Sensing of the hydrogel phase transition between hydrophobic and hydrophilic (reprinted with permission from [92], © The Chinese Laser Press): (a) schematic of the hydrogel sensing system with a microbubble resonator; (b) resonance wavelength shift, spectral linewidth broadening, and scattering enhancement from hydrophilic to hydrophobic.

4.3. Biosensing

Due to their favorable optical characteristics and microfluidic channel, microbubble resonators have been used for the biosensing, such as biotin and protein. In 2013, a self-referencing optofluidic microbubble resonator was studied and used for bovine albumin (BSA) detection [64]. The noise was further suppressed to 0.029 pm by monitoring two splitting modes. A detection limit of 0.5 pg/mL was realized. Furthermore, a method of theoretical analysis for the sensitivity and the detection limit of biosensing was proposed by Barucci et al. in 2016 [128]. By decreasing the microbubble wall thickness or choosing a higher order radial mode, the sensitivity and detection limit for biosensing were further increased. As shown in Figure 14a,c, D-biotins of different concentrations could be detected with the microbubble resonators [90]. A detection limit of 0.41 pM was realized. However, the noise limited its further measurement of ultra-low concentrations of biomolecules. To further reduce the noise, an external referencing microbubble resonator was proposed by Guo et al. in 2019 [93]. An ultra-low concentration of 1 fg/mL for BSA and D-biotin were reported, as shown in Figure 14b,d. The drawback of this method is its complicated referencing systems.

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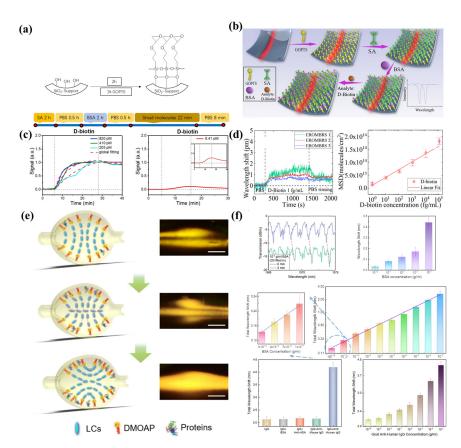


Figure 14. Biomolecule sensing with microbubble resonators: (**a,c**) D-biotin sensing of packaged microbubble resonators (reprinted with permission from [90], © CC BY license); (**b,d**) external referencing microbubble resonators for the D-biotin sensing (reprinted with permission from [93], © Optica, formerly the Optical Society of America). (**e,f**) biomolecule sensing of microbubble resonators with an amplification of the liquid crystals (reprinted with permission from [129], © CC BY license).

Furthermore, ultrasensitive biomolecule detection employing the amplification of liquid crystals (LC) was proposed by Wang et al. in 2021 [129], as shown in Figure 14e,f. The orientation of the LC molecules changed with the increasing of biomolecule concentrations, resulting in the amplification of the wavelength shift. Different biomolecules were detected at the concentrations of 2 fM. In addition, a single DNA molecule with 8 kDa was detected by the plasmonic-enhanced interface mode in the microbubble resonators [130]. However, the above methods can only detect one specific biomolecule. To realize the simultaneous detection of multiple biomolecules, a method based on selectively immobilizing antibodies on a specific microbubble was proposed by Berneschi et al. in 2016 [131]. As shown in Figure 15a,b, by the combination with the photochemical activation method, multiple antibodies were modified on different microbubbles for parallel antigen detection. Using a combination of microbubble resonators, an ultra-low detection limit of biosensing was realized with a miniature sensor system. However, the internal surface modification had low efficiency, which would need to be further improved for practical application.

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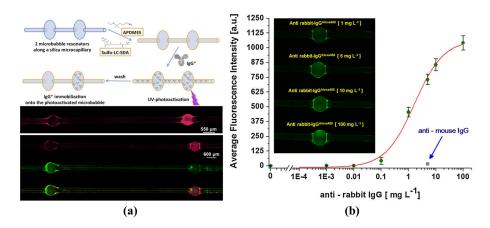


Figure 15. (**a**,**b**) Parallel biomolecule detection based on a photochemical activation method with microbubble resonators (reprinted with permission from [131], © Elsevier).

5. Summary and Outlook

Here, we summarize the recent developments in the use of WGM microbubble resonators, including the sensing principles, fabrication methods, as well as various physical, chemical, and biological sensing applications, as shown in Table 1. Due to their high *Q*-factor, small mode volume, high optical energy density and hollow microfluidic channel, microbubble resonators possess advantages for the sensing of small changes. Miniaturized sensors have been fabricated using a combination of microfluidic technologies and can be used for the detection of temperature, stress, strain, pressure, flow rate, RI, ultrasound, gas, ions, and biomolecules, as shown in Figure 16. The different types of microbubble sensors used indicate their broad potential research value and future practical applications.

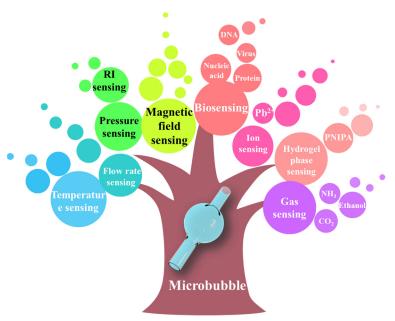


Figure 16. Overview of the application of microbubble resonators in sensing.

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Table 1. Summary of different application of the microbubble resonators.

Application		Q-Factor	Fabrication Material	Sensitivity	Dynamic Detection Ranges	Detection Limit
	[95]	106	Glass capillary	100 GHz/K	-	8.5 mK
Temperature Sensing	[108]	1.6×10^4	PMMA	39 pm/°C	25 °C-80 °C	-
	[117]	5.8×10^{3}	Silica capillary	205 pm/°C	22 °C-26 °C	-
	[118]	106–107	Silica capillary	205 pm/°C	~65 °C	$0.002~^{\circ}\mathrm{C}$
Liquid- concentration Sensing	[89]	5.8 × 10 ⁶	Silica capillary	18.8 nm/RIU	-	5.4 × 10 ⁻⁵ RIU
	[93]	1.8×10^{6}	Silica capillary	11.4 nm/RIU	-	5.5×10^{-5} RIU
	[98]	$10^3 - 10^7$	Glass capillary	-1.1 GHz/bar	-	-
Pressure Sensing	[99]	-	Silica microcapillary	6.21 GHz/bar	-	0.02 μm
	[106]	5×10^{7}	Silica Capillary	38 GHz/bar	-	0.17 mbar
	[119]	1.5×10^{6}	Silica capillary	51.2 pm/bar	-	-
Flow Rate Sensing	[100]	10 v 105	Ciliaa aanillamu	0.0196	10 μL/min-	
	[100]	1.0 ^ 10°	Silica capillary	$pm/(\mu L/min)$	200 μL/min	-
	[101]	1.7×10^{5}	Silica capillary	0.079	$0 \mu L/min-200$	
				$pm/(\mu L/min)$	μL/min	-
Magnetic Sensing	[96]	2.1 × 10 ⁴	Hollow Fiber	0.08 pm/mT	0.41 mT-21.8 mT	-
	[97]	4.0 × 10 ⁶	Microcapillary (Magnetic Fluid)	25.2 pm/mT	2 mT–20 mT	-
Gas Sensing	[111]	2.2 × 10 ⁴	PDMS	36.24 pm/ppm	4.2 ppm–272.4 ppm	-
	[122]	4.4×10^{4}	Silica capillary (Graphene)	200 kHz/ppm	1 1010100 4000	1 ppb
	[123]	1.1×10^{5}	Silica capillary (PHMB)	0.46 pm/ppm	200 ppm–700 ppm	50 ppm
Ion Sensing	[125]	4.7×10^{4}	Silica capillary	265.2 nm/RIU	0.1 pM - 100	15 fM
pH Sensing	[126]	-	Silica capillary (polyNIPA)	33 nm/RIU	pH 3.4–pH 5.6	pH 0.06
Hydrogel Sensing	[92]	9.1 × 10 ⁷	Silica capillary	-	Hydrophobic -hydrophilic	-
Biosensing	[64]	3.0×10^{5}	Silica capillary	-	0 ng/mL–20 ng/mL	2 pg/mL
	[90]	3.7×10^{5}	Silica capillary	-	-	0.41 pM
	[93]	1.8 × 10 ⁶	Silica capillary	11.4 nm/RIU	1 fg/mL–100 pg/mL	1 fg/mL
	[129]	1.4×10^4	Silica capillary (5CB liquid crystal)	-	10 ⁻¹² g/Ml–10 ⁻³ g/mL	1.92 fM
	[130]	8.6×10^{8}	Microcapillary	~0.9 fm/(pg.cm ⁻²)	-	0.3 pg/cm ²
	[131]	3×10^{7}	Silica microcapillary	-	10 ⁻³ mg/L-10 ² mg/L	0.015 mg/L

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However, there remain technical challenges to overcome regarding their production and practical application. Firstly, microbubble resonators with three-dimensional structures are difficult to prepare quickly and in batches using the conventional semiconductor processing technology. Therefore, it is possible to further optimize the fabrication method based on an automated process. Secondly, it can be difficult to miniaturize a tunable laser while maintaining a high stability. The large size of a tunable laser is not flexible for integration and commercialization. Thirdly, it is important to maintain the same performance of the sensor outside the laboratory environment for applicability in real-world measurements. Therefore, it will be necessary to further improve the stability and signal-to-noise ratio of microbubble sensors. In addition, new data processing and analysis techniques are required to fully utilize the multiple WGMs formed. The level of sensitivity and the dynamic measurement range of the sensor will also require further improvement. Among the potential analysis methods developed for WGM microbubble sensors thus far, the barcode technology may be a feasible solution.

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