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## Detection of microorganisms using terahertz metamaterials

## SUBJECT AREAS:

METAMATERIALS

TERAHERTZ OPTICS

SUB-WAVELENGTH OPTICS

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Microorganisms such as fungi and bacteria cause many human diseases and therefore rapid and accurate identification of these substances is essential for effective treatment and prevention of further infections. In particular, contemporary microbial detection technique is limited by the low detection speed which usually extends over a couple of days. Here we demonstrate that metamaterials operating in the terahertz frequency range shows promising potential for use in fabricating the highly sensitive and selective microbial sensors that are capable of high-speed on-site detection of microorganisms in both ambient and aqueous environments. We were able to detect extremely small amounts of the microorganisms, because their sizes are on the same scale as the micro-gaps of the terahertz metamaterials. The resonant frequency shift of the metamaterials was investigated in terms of the number density and the dielectric constants of the microorganisms, which was successfully interpreted by the change in the effective dielectric constant of a gap area.

Culture-based detection methods such as polymerase chain reaction (PCR) and fluorescence-based microbial detection systems have been widely used in order to detect and quantify microorganisms<sup>1–3</sup>. However, although the PCR is capable of detecting a wide range of infectious fungi and bacteria, this method is time-consuming and labor-intensive. Conversely, most of fluorescence-based microbial detection systems such as epifluorescence microscopy (EFM), flow cytometry (FC) and solid-phase cytometry (SPC) require fluorescent materials for efficient detection of the microorganisms<sup>4,5</sup>. Optical scattering methods such as light scattering and autofluorescence do not require the dyes; however they suffer from the lack of sensitivity. Therefore, there is an increasing need to develop novel techniques for the effective on-site detection of minute amounts of microbial substances.

Recently, terahertz (THz) spectroscopy has emerged as a promising technique that enables the label-free, non-contact, and non-destructive inspection on the chemical and biological substances<sup>6–15</sup>. In particular, the recently developed, portable THz spectroscopic tools enable on-site detection and identification of these materials with high signal to noise ratio<sup>16,17</sup>. THz frequency range detection of microorganisms such as fungi, bacteria, and viruses has generated a great deal of interests because of its relevance to food and security inspection<sup>18</sup>. In many cases, however, the microorganisms are non-responsive in the THz frequency range since they are mostly transparent to THz waves. In addition the size of a typical microorganism is on the order of  $\sim\lambda/100$ , resulting in low scattering cross-section.

Metamaterials consist of periodically arranged, sub-wavelength metallic elements and exhibit unique electromagnetic properties such as negative refraction<sup>19,20</sup>, sub-diffraction limited focusing<sup>21–23</sup> and cloaking<sup>24,25</sup>. In addition, the metamaterials have gap structures characterized by strongly localized and enhanced fields, enabling sensitive detection of extremely small amounts of chemical and biological substances<sup>26–32</sup>. In particular, metamaterials operating in the THz frequency range have a micro-sized gap<sup>33–39</sup> and can therefore serve as an ideal platform for the sensitive detection of fungi and bacteria, because the size of these microorganisms is compatible with the gap size. Moreover, THz metamaterials are extremely sensitive to the substances near the surface, which is favorable for sensing in aqueous environment since it allows us to use a thin water layer without suffering from the significant loss in the THz wave transmission.

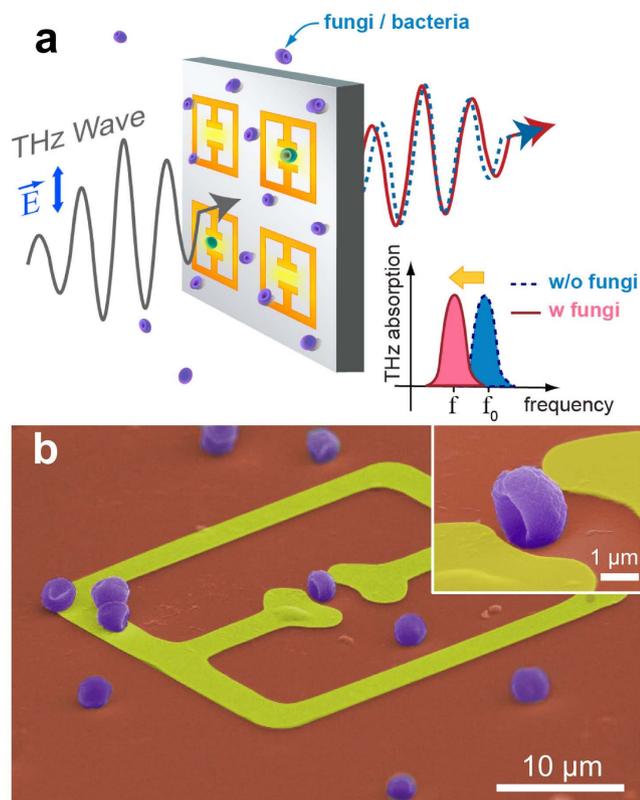
In this work, we performed THz time-domain spectroscopy on metamaterial sensors for the high speed detection of viable and live microorganisms such as molds, yeast cells and bacteria. We measured a frequency



shift in the inductive-capacitive resonance following the deposition of microorganisms with very low surface density. In particular, we functionalize the sensors with the antibody specific to bacteria in aqueous environment, enabling the selective detection. The resonance frequency shift in THz metamaterials was studied as a function of the dielectric constants and the density of fungi which is in accordance of the dielectric constant measurement of individual fungi. Our experimental findings are in a good agreement with the results of simulations.

Schematic presentation of the experiments is shown in Fig. 1a. We measured the change in spectra of the THz radiation transmitted through the metamaterials following the deposition of microorganisms. The metamaterials consist of metallic arrays of a square ring with a micro-gap at the center. Here, the resonance of the transmission is determined both by the capacitance ( $C$ ) of the gap structure and the inductance ( $L$ ) associated with the shape of the square ring. In other words, the resonance of the transmission dip is expressed by  $f_{res} = 1/(2\pi\sqrt{LC})$ . The dielectric microorganisms placed in the gap area will cause a change in the effective dielectric constants of the capacitor, resulting in the shift of the resonant frequency in metamaterials.

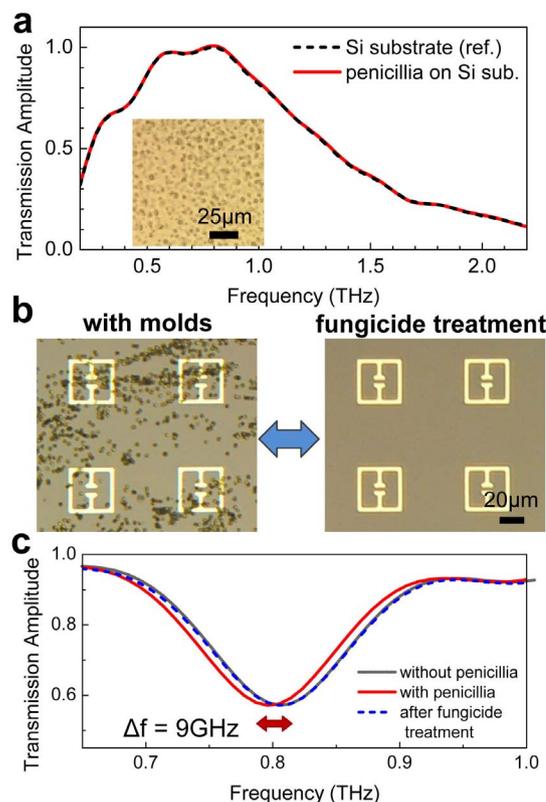
A representative scanning electron microscopy (SEM) image is shown in Fig. 1b, with the *Penicillium chrysogenum* (penicillia) deposited on the metamaterials. The fungi samples were grown by a streaking on medium method followed by the incubation at 37°C for 2 days. Metamaterial patterns were prepared by using a conventional photo-lithography method on an undoped Si substrate, followed by the metal deposition in order to define the arrays of split ring resonator patterns with a linewidth of 4  $\mu\text{m}$  and gap sizes of 2–3  $\mu\text{m}$ . THz transmission spectra were obtained by using the THz time-domain spectroscopic techniques with an acquisition time of 5 sec for each spectrum<sup>40,41</sup>.



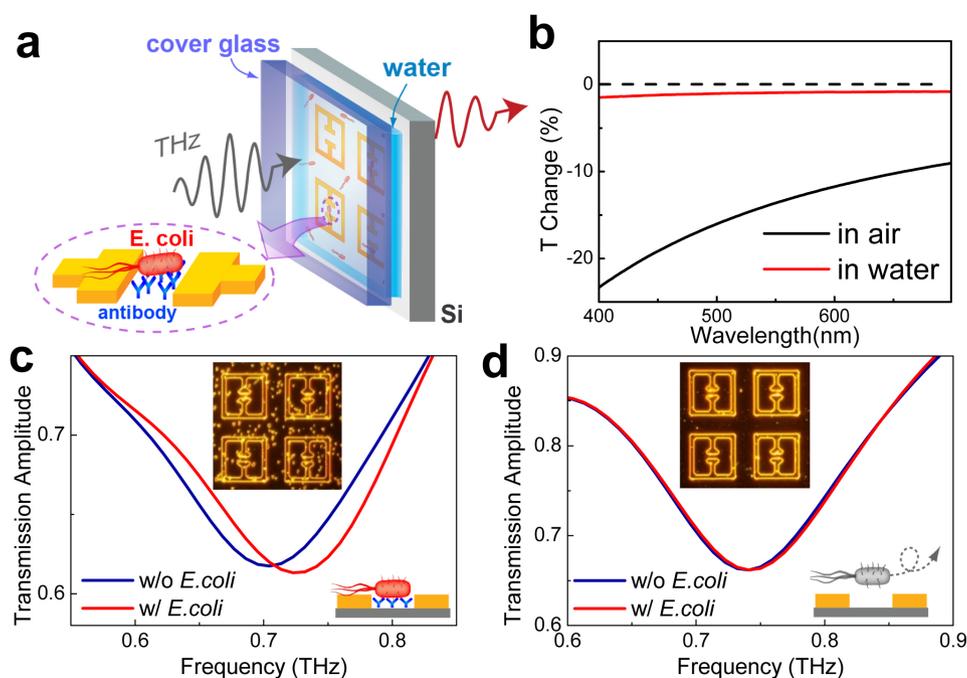
**Figure 1 | Sensing microorganisms using THz metamaterials.** (a) A schematic presentation of THz metamaterials sensing of microorganisms. (b) A color-enhanced SEM image of metamaterials coated by penicillia. (inset) Magnified image of the fungi located in the micro-gap.

We first describe the THz transmission experiments for the low-density microorganisms that were deposited on a plain silicon substrate. Molds and *Saccharomyces cerevisiae* (yeasts) were deposited on the substrate by rubbing a fungi-coated swab on it. Fig. 2a shows the THz transmission amplitude obtained when the penicillia were deposited on the plain substrate at a density of  $0.032/\mu\text{m}^2$ , as shown by the microscope image in the inset of Fig. 2a. No noticeable change in transmission amplitude was found following the deposition of the molds compared to the bare substrate case. Many of the microorganisms such as fungi and bacteria do not have spectral fingerprints in the THz frequency range and thus are basically transparent to the THz waves. Besides, the scattering cross-section of those microbial materials to THz waves is very low, because the typical size of these microorganisms ( $\sim\mu\text{m}$ ) is significantly smaller than the wavelength of THz waves. A similar behavior with respect to THz transmission was observed for other microorganisms, such as yeasts and *Escherichia coli* BL21(DE3) (*E. coli*), plated at comparable or higher densities (See Supplementary Figure S1).

A representative result on the highly sensitive detection of low-density molds by using the THz metamaterials is demonstrated in the following. Fig. 2b shows two microscopic images of our metamaterial pattern (so-called “electrical split ring resonator”), which is an array of a square ring with a micro-gap at the center<sup>33,34</sup>. The transmission dip yields 0.837 THz with a full-width at half maximum (FWHM) of 0.15 THz, which corresponds to the inductive-capacitive resonance for the device shown in the image. The left side image in Fig. 2b shows metamaterials with fungi (penicillia) that were deposited at a density



**Figure 2 | Metamaterial sensing of penicillia.** (a) THz transmission amplitudes, for transmission through a plain Si substrate, with (red solid line) and without (black dashed line) deposition of penicillia. (inset) A microscopic image of penicillia deposited on the Si substrate. (b) Two microscope images of metamaterials patterns on Si substrate, with the deposition of penicillia a (left) and after fungicide treatments (right). (c) THz transmission amplitudes measured before (gray solid line) and after (red solid line) the deposition. Transmission after the fungicide treatment is shown as blue dashed line.



**Figure 3 | Metamaterial sensing of *E. coli* in aqueous environment.** (a) A schematic of selective bacteria (*E. coli*) detection in solution environment. The Si substrate is coated with antibodies specific to *E. coli*. (b) Visible transmittance spectra of the quartz substrate coated by *E. coli* (density of  $0.078 \mu\text{m}^2$ ) both in aqueous (red line) and in ambient (black line) conditions. (c) THz transmission before (blue line) and after (red line) the deposition of *E. coli* on the functionalized metamaterials in aqueous environments. (inset) A corresponding dark-field microscopic image obtained after the deposition of *E. coli*. (d) THz transmission before (blue line) and after (red line) the deposition of *E. coli* on the sensors without the surface functionalization. (inset) A corresponding dark-field microscopic image after the deposition process.

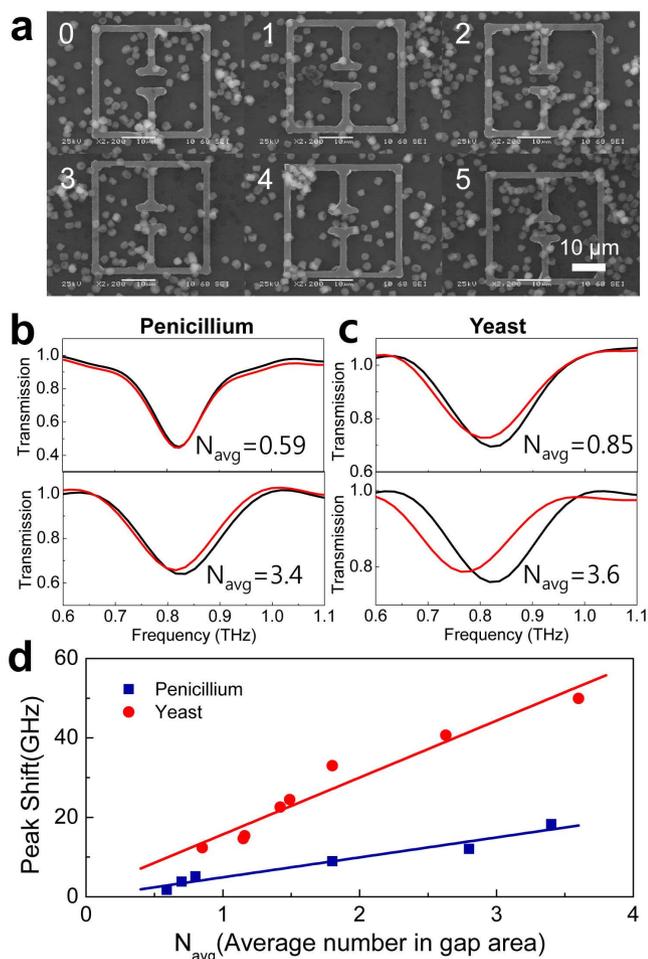
of  $0.090/\mu\text{m}^2$ , and individual molds that are separated from each other can be clearly seen in this figure. As shown in the right side image, the fungi can be removed completely by using a commercial fungicide (sodium hypochlorite solution). Results for the corresponding THz transmission are shown in Fig. 2c. Unlike the results on the transmission through the plain substrate (Fig. 2a), the resonant frequency exhibited a shift towards the red (red solid line) relative to the resonant frequency obtained for clear metamaterial patterns (gray solid line). The amount of the shift was found to be 9 GHz (6% of the FWHM) for the density of penicillia shown in Fig. 2b. Once the molds were cleaned by using the fungicide, the initial resonant frequency was retrieved (blue dashed line), which verifies the reusability of our metamaterial sensors.

The shift in the resonance frequency can be explained simply by the change of the dielectric environment in the gap area that modifies the capacitance according to  $C = \epsilon A/d$ , where  $A$  is the area,  $d$  is the distance, and  $\epsilon$  is the dielectric constant of the capacitor. It is clear that the increase in the dielectric constant due to the presence of the microorganisms results in the red-shift of the resonant frequency. Similar behaviors were also observed for yeasts and different kinds of molds, such as *Neurospora sitophila* (neurospora) and *Aspergillus niger* (niger) (See Supplementary Figure S2). We also performed experiments by using different types of the metamaterial patterns with a gap structure such as single/double split ring resonators and found similar red-shifts following the deposition of fungi; however the electrical split-ring resonator delivers better sensitivity in general (See Supplementary Figure S3).

Our technique can be extended to the highly sensitive detection of bacteria in aqueous environments, which is not accessible by the conventional optical techniques. More importantly, THz metamaterials functionalized with receptors enable the specific sensing of bacteria. A representative result, demonstrating the detection of bacteria (*E. coli*) in a solution environment, is shown in Fig. 3. The *E. coli* were deposited from the solution (100  $\mu\text{g}/\text{ml}$ ) on the Si substrate

with metamaterial patterns functionalized with *E. coli* anti-body<sup>42</sup>, followed by a rinsing with purified water (Fig. 3a). The patterns were enclosed by a cover glass, and the water layer thickness was fixed at 55  $\mu\text{m}$  by using a spacer. This thickness is large enough to cover the small detection volume that is strongly localized at the surface, yet allowing a considerable amount of THz wave transmission. We note that, the *E. coli* are not as clearly resolved in the solution as they are in the air. Fig. 3b shows the optical transmission spectra in the visible range, obtained for transmission through the quartz substrate coated by the *E. coli* (density of  $0.078/\mu\text{m}^2$ ), for both aqueous (red line) and ambient (black line) environments. Transmission through the bare quartz substrate was used as a reference in both cases. The change in the optical transmission reaches only  $\sim 1\%$  in water, which is 10-fold smaller as compared to the air. This strongly suggests that the microbial detection techniques based on the scattering of visible light can be limited for some of the microbial systems in the solution phase.

In contrast, using the THz metamaterials sensors, we can successfully detect the low-density *E. coli*. Shown in Fig. 3c are the results for the THz transmission obtained with (red line) and without (blue line) the *E. coli* on the functionalized substrate. The inset of Fig. 3c shows a dark-field microscope image of metamaterials in water with the *E. coli* (density of  $0.019/\mu\text{m}^2$ ) adsorbed on the substrate. A clear blue-shift of 23 GHz (yielding  $\sim 15\%$  of a FWHM) is observed for the metamaterials with *E. coli* compared to the case of the clean substrate (blue line). The reason that the deposition of *E. coli* in aqueous solution causes a blue shift is likely due to the relatively low dielectric constant of *E. coli* ( $\sim 1.6$  at 1 THz) as compared to that of water ( $\sim 4.2$  at 1 THz)<sup>43</sup> (Also see Supplementary Figure S4). Without the surface functionalization, however, we could not observe a noticeable change in the peak position (Fig. 3b). Therefore it is possible to fabricate the biosensors that are specific to the bacteria of interests. Quite obviously, our scheme can be extended to the selective detection of variety of microorganisms such as fungi<sup>44</sup> and viruses



**Figure 4 | Resonant frequency shift as a function of number density and dielectric constants of fungi.** (a) A series of six SEM images of the metamaterials with the number of penicillia in the gap areas ranging from 0 to 5. (b) THz transmission amplitudes with the penicillia number density of  $N_{avg} = 0.59$  and 3.4, respectively from top to bottom. (c) Same as (b) but with the yeast number density of  $N_{avg} = 0.85$  and 3.6, respectively from top to bottom. (d) A plot of resonant frequency shifts as a function of  $N_{avg}$  for depositions of penicillia (blue boxes) and yeasts (red circles). Solid lines are linear fits to the data.

since the specific binders and their surface functionalization have been widely investigated.

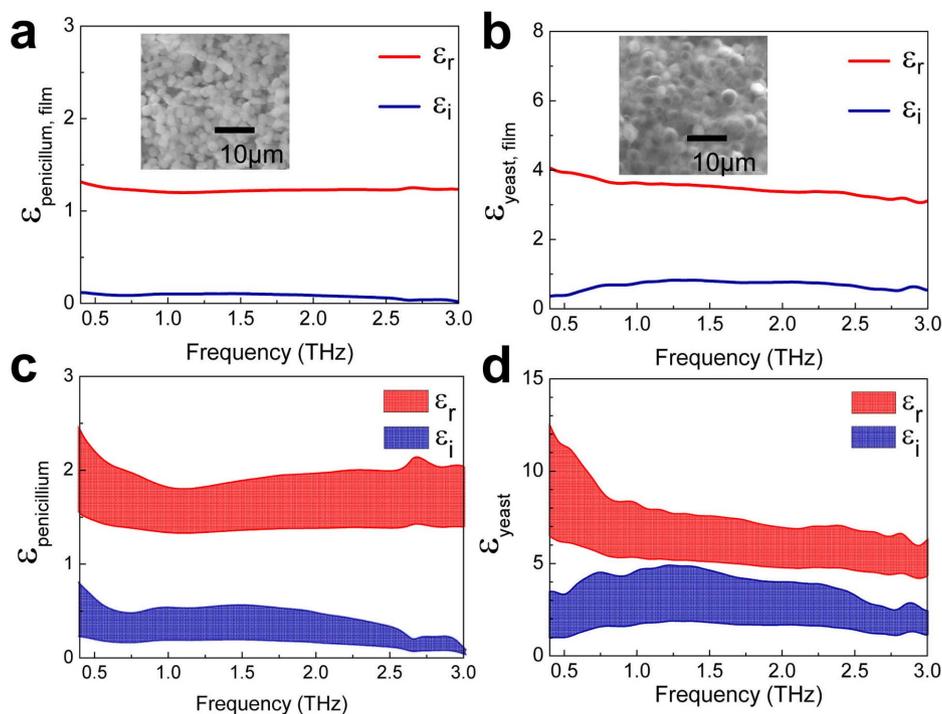
It is evident that the shift in the resonance frequency will strongly depend on the number of fungi and bacteria located in the gap area and on the dielectric constants of the specimens. In Fig. 4, the shift of the resonant frequency is shown as a function of the number density, for different kinds of fungi. The average number of fungi in the gap area was determined by counting them for the entire set of elements ( $10 \times 10$ ) positioned in the spot area of the THz fields ( $\sim 1 \text{ mm}^2$ ). For example, Fig. 4a shows a series of SEM images of metamaterials with the number of penicillia in the gap area ( $\sim 3 \times 10 \mu\text{m}^2$ ) ranging from 0 to 5. Fig. 4b shows the representative data on the resonant frequency shifts, for two different number densities of  $N_{av} = 0.59$  (corresponding to surface density of  $0.020/\mu\text{m}^2$ ) and 3.4 ( $0.113/\mu\text{m}^2$ ), respectively from top to bottom for the penicillia. Similar experimental data are shown in Fig. 4c for the yeast cells with  $N_{av} = 0.85$  ( $0.028/\mu\text{m}^2$ ) and 3.6 ( $0.120/\mu\text{m}^2$ ), respectively. We emphasize that we were able to detect extremely small amounts of fungi, in other words, even in the case when the average number in the gap area was less than unity. A plot of the resonant frequency shift as a function of  $N_{av}$  is shown in Fig. 4d for the penicillia (blue boxes) and yeasts (red

circles), respectively. The magnitude of the resonant frequency shift is gradually increasing with  $N_{av}$  for both fungi types. More importantly, it is very clear that the frequency shift is  $\sim 3$ -fold higher in the case of the yeasts. This is a strong indication that the dielectric constant of the yeast is higher than that of the penicillium.

As mentioned earlier, the resonant frequency shift as a function of  $N_{av}$  and the dielectric constant of the individual fungi ( $\epsilon_f$ ) can be analyzed in terms of the changes in the effective dielectric constant of the capacitor. The modified effective dielectric constant  $\epsilon$  due to the presence of the fungi in the ambient condition can be expressed as  $\epsilon = \epsilon_{eff} + \alpha N(\epsilon_f - 1)$ , where  $\epsilon_{eff}$  is the effective dielectric constant without the deposition of fungi,  $N$  is the number of microorganisms, and  $\alpha$  is the coefficient which is associated with the volume fraction of the fungi. Here,  $\epsilon_{eff}$  in the capacitance (and hence the resonant frequency) is influenced both by the dielectric constants of the substrate and the air<sup>41</sup>, in which we obtained  $\epsilon_{eff} = 6.4$  from simulation results (See Supplementary Figure S5). From the relation  $f = f_0(\epsilon/\epsilon_{eff})^{-\frac{1}{2}}$ , where  $f(f_0)$  is the resonant frequency with (without) the deposition of the fungi, the frequency shift  $\Delta f = f - f_0$  leads to  $\Delta f/f_0 \approx -\frac{1}{2}\alpha N(\epsilon_f - 1)$ . Here we neglect higher order terms in  $\alpha N(\epsilon_f - 1)/\epsilon_{eff}$ , since  $\epsilon_f/\epsilon_{eff}$  is on the order of or less than unity as will be shown below, whereas  $\alpha$  is on the order of 0.01. It is clear from the above equation that, when the density of the microorganisms is low, the frequency shift is linearly dependent on  $\epsilon_f - 1$  as well as on  $N$ .

To verify the effects of the dielectric constants on the resonant frequency shift, we measured the dielectric constants of the fungi ( $\epsilon_f$ ) used in the experiments by preparing thick and dense fungi films. The films were prepared by stacking the large amount of fungi layer by layer on a cellulose membrane. SEM images of the penicillium and yeast films are shown in the insets of Fig. 5a and 5b, respectively. We found from the images that the yeast film consists of very closely packed spheres, whereas that of penicillium is rather loose in general. By measuring both the amplitudes and phases of the transmitted THz pulses, we extracted the complex dielectric constants of the films that consist of the closely packed molds and yeasts. The complex dielectric constants of the thick films ( $\epsilon_{film}$ ) are shown in Fig. 5a and 5b for the penicillium and yeast films, respectively.

The dielectric constants of the fungi can be obtained from those of the films by using the effective medium theory (Maxwell-Garnett model) of  $(\epsilon_{film} - 1)/(\epsilon_{film} + 2) = f(\epsilon_f - 1)/(\epsilon_f + 2)$ , where  $f$  is the volume fraction of the fungi contained in the films<sup>45–47</sup>. Here it is important to estimate the volume fractions reasonably in order to obtain the dielectric constants of the individual fungi accurately. In the case of penicillium film, we extracted  $f$  of 0.28–0.62 by measuring the total mass and volume of the film and from the information on the mass of penicillium (ranging 58–129 pg, depending on the wetting condition)<sup>48</sup>. On the other hand, the volume fraction  $f$  of the yeast film was chosen to be 0.64–0.78, assuming that it consists of the closely packed spheres<sup>49</sup>. Fig. 5c shows the complex dielectric constants of the individual penicillium as extracted from the THz transmission experiments, which shows the upper and lower bounds corresponding to the volume fractions of 0.28 and 0.62, respectively, yielding the real part of the dielectric constants as 1.37–1.94 at 0.8 THz. Similarly, the dielectric constant of yeast is determined to be 5.38–8.54. The dielectric constant of the yeast (which is significantly higher than that of penicillium) is in accordance with the results shown in Fig. 4, in which the resonant frequency shift for a given number density is higher for the yeasts as compared to the penicillia. However, their relative difference in  $\epsilon_f - 1$  is somewhat larger than the value expected from Fig. 4. It is likely that other parameters such as the shape of the individual fungi and the wetting conditions need to be considered further in order to explain the discrepancy, which requires future study. We also measured the dielectric constant of *E. coli* in order to explain the blue shift of resonant peak in a solution environment [Supplementary Figure S4].



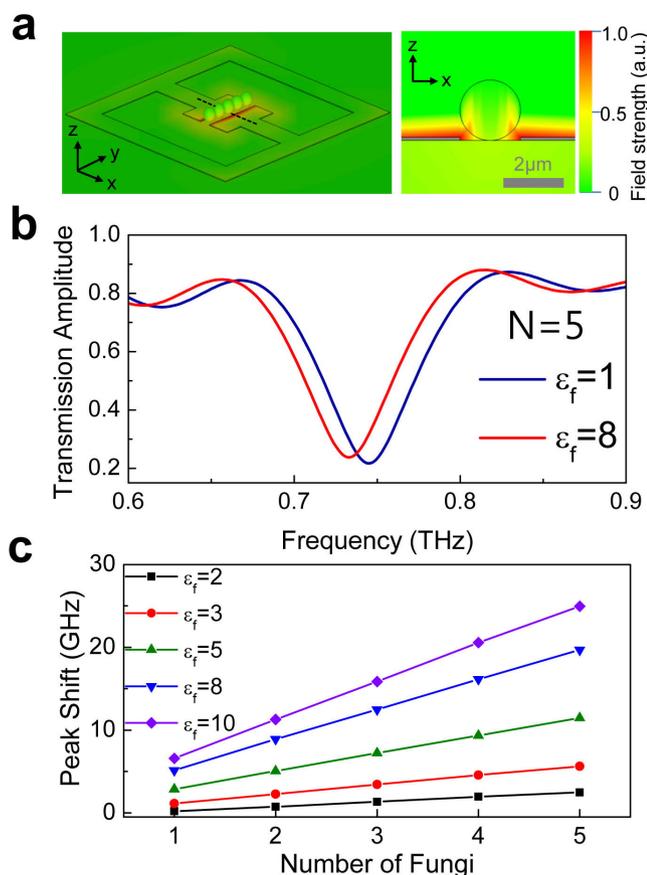
**Figure 5 | Dielectric constant measurements of Fungi.** Plots of frequency-dependent, complex dielectric constants of the penicillium (a) and yeast (b) films, extracted from the THz transmission experiments. Films' thickness were 350 and 125  $\mu\text{m}$ , respectively. Shown together are the dielectric constants of the individual penicillium (c) and yeast (d), extracted from (a) and (b) by using a Maxwell-Garnet theory.

Our experimental findings were analyzed by using the finite-difference time-domain simulations, as shown in Fig. 6. We used the metamaterial patterns (described by gray lines) with the geometric parameters similar to those in our experiments, as shown by the three-dimensional (left) and two-dimensional cross-sectional plots (right) in Fig. 6a. In order to reproduce the effects of the fungi deposition, we added a series of dielectric spheres (which represent the fungi) in the gap areas of the metamaterials. Five dielectric spheres, each with the diameter of 2  $\mu\text{m}$ , were used in this case. The color images overlaid in the patterns are the representative plots showing the electric field strength at the resonant frequency of 0.73 THz. It is clear that the electric fields are strongly localized in the gap area as mentioned before, whereas the fields are strongly modified in the presence of the dielectric spheres. Fig. 6b shows the typical transmission spectra of the simulation results with the dielectric constants of the spheres set to  $\epsilon_f = 1$  (air) and  $\epsilon_f = 8$  (fungi), respectively.

A clear red-shift in the resonant frequency is observed for the case of  $\epsilon_f = 8$  (with fungi) as compared to the case of  $\epsilon_f = 1$  (without fungi), consistent with the experimental findings. Finally, we show a plot of the resonant frequency shift as a function of the number of spheres in the gap areas, for different dielectric constants ranging from 2 to 10 (Fig. 6c). The resonant frequency shift increases linearly with the increasing number of fungi in the gap areas, which is apparently higher for the higher dielectric constants (See Supplementary Figure S6). Results of these simulations successfully reproduce our experimental findings. We also found that the inductive-capacitive resonant peak is not influenced noticeably by the dielectric materials located outside of the gap area (See Supplementary Figure S7). It remains unclear why the magnitude of the resonant frequency shift in the simulation is rather smaller than the one that is obtained in the experiments. The discrepancy is probably because we used photolithographic methods with limited spatial resolution (see Fig. 1b) and possibly is due to approximating the shape of the individual fungi as spheres.

We would like to mention that our technique is sensitive to the surface number density of the fungi and bacteria located at the surface, whereas most of the contemporary techniques refer the volume density<sup>50,51</sup>. In fact, there is a tradeoff between the detection speed and the sensitivity. In our case, the sensitivity depends on the deposition time from a solution in sensing bacteria as shown in Fig. 3. For the detection of *E. coli* in drinking water, the latest culture-based methods report the detection limit of 50 CFU/ml in 3 hr, in which only colony-forming units (CFU) are counted<sup>52</sup>. By simply counting the number density of bacteria in the solution, we can estimate our detection limit to be  $\sim 10^7$  units/ml (including non-colony-forming units) for 1 hr of *E. coli* deposition. Therefore, improving the capture efficiency of these microorganisms at the surface, and near the gap-area in particular, will be essential for improving the sensitivity and the speed of the THz metamaterial sensing. Besides, the sensitivity can be improved further by adopting advanced designs with higher Q-factors<sup>53,54</sup>, by using the substrate of lower refractive index, and by increasing the metal thickness so that it is comparable to the size of the microorganisms.

To conclude, we found that the THz metamaterials can be used to detect various microorganisms such as molds, yeasts, and bacteria. A clear shift in the inductive-capacitive resonant frequency is observed following the deposition of microorganisms, and arises due to the change in the effective dielectric constant in a gap area of the metamaterial patterns. The resonant frequency shift is higher for the larger dielectric constants, which was confirmed by the dielectric constant measurements of individual fungi and by the numerical simulations. Our technique can be applied to various viable and living microorganisms that dwell in aqueous environments, as well as in ambient conditions, because it is highly sensitive to the substances located near the surface, allowing us to use a thin water layer without suffering from the poor transmission in conventional THz spectroscopies. The THz metamaterial sensing is a universal method because it is based on the dielectric sensing, while a selective detection is also possible by functionalizing the substrates with antibodies



**Figure 6 | Results of finite-difference time-domain simulations.** (a) 3D plot of the electric field strength at the resonant frequency of 0.73 THz, when the series of five dielectric spheres (with  $\epsilon_f = 8$ ) are placed in the micro-gap of the metamaterials. Shown on the right is a 2D cross-sectional image taken along the dashed line in the 3D plot. (b) THz transmission spectra for the cases in which the dielectric constant of each of the five spheres are of  $\epsilon_f = 1$  (blue line) and  $\epsilon_f = 8$  (red line), respectively. (c) A plot of resonant frequency shift as a function of the number of spheres, for five different  $\epsilon_f$ 's ranging from 2 to 10.

specific to the target substances. Our approach is likely to constitute an important step toward the fabrication of the highly sensitive biosensors and lab-on-a-chip devices, enabling the high-speed on-site detection of hazardous substances in various environments.

## Methods

**Terahertz time-domain spectroscopy.** The real-time THz transmission amplitudes of the metamaterials with fungi were measured with a conventional THz time-domain spectroscopy (THz-TDS) setup. A femtosecond laser at  $\lambda = 800$  nm is incident on the photoconductive antenna which emits a linearly polarized THz pulse. Subsequently, this THz pulse is focused on the metamaterials with  $\sim 1$  mm spot diameter under ambient conditions. Time traces of the transmitted THz electric field both in amplitude and phase were measured by varying the time delay between the 800 nm probe beam and the THz pulse. THz spectrum is taken by applying a fast Fourier transform (FFT) to the time trace and normalized with respect to the reference. The typical spectral resolution was  $\sim 30$  GHz for a temporal scan range of 33 ps.

**Fabrication of Metamaterial Patterns.** Metamaterial patterns were prepared by using a conventional photo-lithography method on a high-resistivity Si substrate (n-type, resistivity  $> 6000 \Omega \cdot \text{cm}$ , and thickness of 550  $\mu\text{m}$ ). Cr (2 nm) and Au (98 nm) metal films were deposited by an e-beam evaporator in order to define the arrays of split ring resonator patterns with a linewidth of 4  $\mu\text{m}$ , outer dimension of 36  $\mu\text{m} \times 36 \mu\text{m}$  and gap size of 3  $\mu\text{m}$ . The periodicities of array patterns were 86  $\mu\text{m}$  (Fig. 2 and 4) and 50  $\mu\text{m}$  (Fig. 3).

**Preparation of fungi.** *Penicillium chrysogenum* (KACC 40399; penicillia), *Neurospora sitophila* (KACC 40972; neurospora), *Aspergillus niger* (KACC 40280;

niger), and *Saccharomyces cerevisiae* (yeasts) were grown by a streaking on medium method followed by the incubation at 37 °C for 2 days. All the fungi were obtained from the Korean Agricultural Culture Collection (KACC). The culture media used for the penicillia, yeasts, neurospora and niger were V8 juice agar, yeast peptone dextrose, potato dextrose agar, and malt extract agar, respectively. *Escherichia coli* BL21(DE3) (*E. coli*) were grown by a similar method on the medium of luria broth.

**Surface Functionalization and Deposition of *E. coli*.** We begin with surface functionalization by immersing the metamaterials substrate in (3-Aminopropyl)triethoxysilane (APTS):toluene (2 : 1000) solutions for 3 hours. *E. coli* antibodies (Abcam) were immobilized on Si substrate with metamaterial patterns by immersing the substrate in antibody: phosphate buffered saline (PBS) 1 × in 200  $\mu\text{g}/\text{ml}$  for 3 hours. After antibody immobilization, *E. coli* were deposited from the *E. coli*: PBS solution in 100  $\mu\text{g}/\text{ml}$  for 1 hour, followed by a rinsing with distilled water to eliminate remaining surface non-linked *E. coli*. The patterns were enclosed by a quartz cover glass (with a thickness of 170  $\mu\text{m}$ ), and the water layer thickness was fixed at 55  $\mu\text{m}$  by using a spacer.

**Dielectric Measurement of fungi and *E. coli*.** The films for the dielectric measurement were prepared by stacking the large amount of fungi layer by layer on a cellulose membrane (with a pore size of 200 nm and a thickness of 100  $\mu\text{m}$ ), as a result, the thickness of the penicillia and yeasts films reached 350  $\mu\text{m}$  and 125  $\mu\text{m}$ , respectively. The membrane substrate is almost transparent to the THz waves, whose refractive index of the membrane substrate is close to unity in the THz frequency range<sup>55</sup>. By measuring both the amplitudes and phases of the transmitted THz pulses, we extracted the complex dielectric constants of the films, from which we extracted those of the individual fungi by using the Maxwell-Garnett model. We performed the same procedures on the three different films for both fungi and found the similar dielectric constants each time.

**Finite Difference Time-Domain (FDTD) Simulation.** To predict the electromagnetic behavior of the metamaterials, the structure is simulated by using CST MICROWAVE STUDIO.

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## Author contributions

Y.H.A. conceived and designed the experiments. S.J.P. and J.T.H. fabricated metamaterials and performed THz experiments. S.J.C. and H.S.K. provided fungi and bacteria samples. W.K.P. and S.T.H. participated in FDTD analysis. All authors including J.Y.P., S.L., and D.S.K. discussed the results and participated in writing the manuscript.

## Additional information

**Supplementary information** accompanies this paper at <http://www.nature.com/scientificreports>

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