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A parallel four-quadrant sensing method utilizing a specially designed gold nanodot array is created for sensing virus-like particles with a sub-diffraction limit size (~ 100 nm) in a wide-field image. Direct label-free sensing of viruses using multiple four-quadrant sensing channels in parallel in a wide-field view enables the possibility of high-throughput onsite screening of viruses.

Infections from viruses and other pathogens are a major threat to human health and even to the survival of the human race. The outbreak of SARS-CoV-2 has resulted in millions of deaths worldwide.¹ Rapid detection techniques with high throughput and sensitivity which allow onsite operation with minimal requirements for sample preparation and the test environment are essential for the prevention and control of pandemics.²⁻⁴

Conventional methods for virus detection include polymerase chain reaction (PCR), serological analysis, and imaging technologies such as computerized tomography (CT) scans.⁵⁻⁹ Various diagnostic sensors have been developed to support standard SARS-CoV-2 diagnostic techniques, which are mainly based on antigenic probes such as whole virus particles and surface antigens (capsids, envelopes, and nucleocapsid proteins) and deliver qualitative outputs (*i.e.*, positive or negative). However, these methods necessitate skilled operators and

Label-free sensing of virus-like particles below the sub-diffraction limit by wide-field photon state parametric imaging of a gold nanodot array†

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require hours or even days to perform the analysis. Meanwhile, they are also restricted by the small number of antibodies produced during different phases of infection.^{10,11}

Compared with contemporary virus identification methods, optical imaging is a promising method for fast virus screening,^{12,13} which includes a visual inspection of cell cultures, fluorescence-based readouts and indirect detection *via* refractive index changes and other mechanisms.¹⁴⁻¹⁶ Challenges faced in direct optical imaging of viruses include the diffraction limit size of particles and the need for sophisticated labeling.^{17,18}

In order to improve the sensitivity of optical imaging, research efforts have been made to develop various labeling and signal amplification methods for virus imaging and sensing. The utilization of nanoparticles, including quantum dots, magnetic particles, polymer nanoparticles, and noble metal nanoparticles, has led to significant enhancement in the detection sensitivity of viruses.¹⁹⁻²⁴ Among various types of nanoparticles, metallic nanoparticles show the distinct feature of localized surface plasmon resonance (LSPR), induced by the interaction of the free-electron cloud of metal nanoparticles with the incident electromagnetic field. Gold nanoparticles have attracted much attention due to their easy modification with biomolecules, allowing tight attachment of the aforementioned antibody or nucleic acid.²⁵⁻²⁷ These strategies mainly rely on exploiting the spectral response of gold nanoparticles decorated with virus-specific antigens, which needs relatively complicated sample preparation. Moreover, their single-channel operation manner limits their application to high-throughput virus screening.

In our recent studies, gold nanoparticles (AuNPs) were used to amplify the scattering signal of viruses in photon state parametric imaging, which suggestss the possibility of wide-field sensing of virus particles.²⁸ It is found that antigen-decorated AuNPs attach to virus particles and enhance the optical scattering signal due to localized plasmons. However, stray AuNPs would randomly aggregate with pathogens, forming different morphologies at random locations in the field of view of the

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microscope. The random distribution of AuNPs degrades the connection efficiency of viruses to antigens, and also means that the virus-specific signal is obscured by noise scattered from other locations. Furthermore, similar sample preparation is required to that in the gold spectrum-based techniques.

To overcome the limitations of current techniques which will be difficult to use in onsite virus screening, in this paper, a virus-like particle sensing method is proposed that operates in a parallel manner with wide-field optical imaging. This is achieved by utilizing grouped gold nanodots as sensing channels in a specifically designed gold nanodot matrix on a Si substrate. A well-regulated optical distribution is formed in the field of view by the scattering of each dot in the matrix. The gold nanodots also demonstrate high efficiency in capturing the virus, allowing label-free sensing of virus-like particles under the sub-diffraction limit with a simple optical setup. In addition, sample preparation processes are minimized and can be used easily in a drop-and-observe manner, which makes the technique attractive for large-scale onsite virus screening. It should be noted that the process of optical scattering in our method is independent of labels, which provides various label-free applications like water impurity detection or concentration and particle size analysis for a known sample.

The layout of the gold nanodot array is shown in Fig. 1. The fabrication process of the gold nanodot array is given as follows (more details in the ESI and Fig. S2[†]): The surface of the silicon substrate was first cleaned with acetone and isopropyl alcohol (IPA). After that, a 200 nm thick polymethyl methacrylate (PMMA) photoresist layer was spun on the substrate. Then the gold nanodot array was defined using electron beam lithography (EBL) and such a pattern was subsequently transferred onto PMMA by development. Next, a 10 nm thick titanium layer for enhanced adhesion and a 70 nm thick gold layer were deposited successively by employing the electron beam evaporation method. Followed by lift-off and cleaning procedures, the desired gold nanodot array was finally realized. Here, the

diameter of a single gold disk is 110 nm, with a height of 70 nm. To avoid light coupling between adjacent nanodots, the spacing between adjacent disks in the horizontal and vertical directions was 1 μ m. The gold nanodot array with a spacing of 0.5 μ m was also fabricated and tested as a reference result to confirm the influence of light coupling. Since every group of four neighboring nanodots in the array will be utilized as a sensing channel for virus detection, their detailed geometry plays an important role in determining the image quality and detection sensitivity. The nanodots were made as round disks with dimensions close to the size of the virus-like particles. The circular shape of the disks guarantees an equal probability of virus attachment in any direction around the nanodot. The diameter of the gold disk was chosen to enhance photon scattering at the wavelength of 554 nm, which lies in the spectral range of the camera with a relatively high quantum efficiency.

The photon state parametric images of the sample, *i.e.*, $\sin \delta$ and ϕ , were obtained using the polarization indirect microscopic imaging (PIMI) technique previously developed by our group (see the ESI[†]).^{29,30} The experimental system is based on a conventional reflection microscope (Olympus BX51), where a 554 nm filter is placed in front of the light source. A highly sensitive CCD (PiA2400-17gm, Basler) with 5 million pixels was used in combination with a 100 \times objective lens, making it possible to obtain images in which the pixel size corresponds to 34.5 nm.

The schematic of the PIMI system is shown in Fig. 2. In our PIMI measurement, the light reflected by the sample was passed through a quarter-wave plate and a linear analyzer was placed in front of the image plane with their fast axes oriented at 45° and 90° respectively relative to the *x*-axis.

The incident polarized light was rotated from 0° to 180° in steps of 18°, and the resulting 10 images were collected and fitted to a sine curve at every pixel (more details in the ESI and Fig. S1[†]), according to the formula:

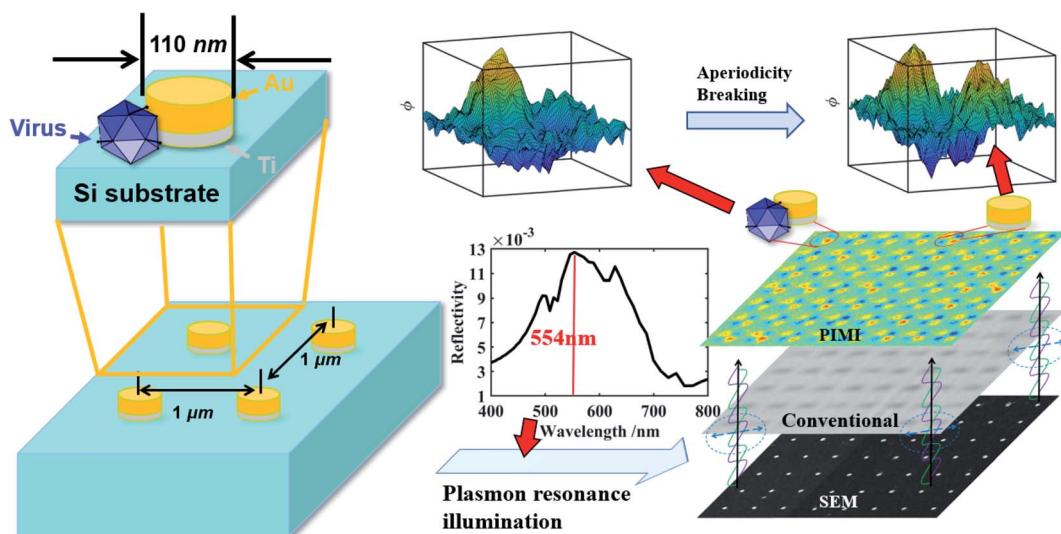


Fig. 1 Schematic of virus-like particle detection and dimensions of gold nanodot arrays on a silicon substrate.



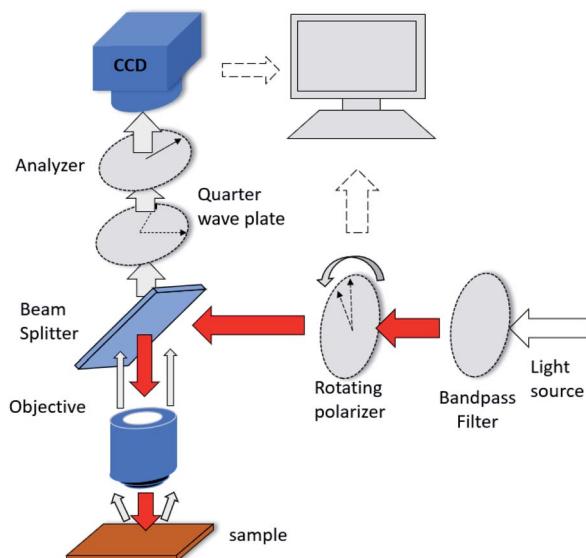


Fig. 2 Diagram of measurement using the PIMI system.

$$I_i = 1/2 \times I_0[1 + \sin \delta \sin 2(\theta_i - \phi)]. \quad (1)$$

where I_i is the i th output intensity, I_0 is the background intensity and θ_i is the i th rotating angle. After curve fitting, the two indirect photon parameters, $\sin \delta$ and ϕ were obtained which respectively represent the phase difference between the two orthogonal polarization components of the scattered light and the polarization ellipse orientation angle of the beam reflected from the sample.

The polarization states of photons scattered by deep sub-wavelength structures or particles heavily depend on the coupling between the incident photons and the optical anisotropy of the sample at the nanoscale. Calculations of the polarization states of scattered photons in the far-field demonstrate that parametric images can have a resolving power beyond the diffraction limit of conventional microscopy. This advantage was exploited in this work to determine whether virus-like particles were attached to the nanodot array. The lower right part of Fig. 1 displays images from a scanning electron microscope (SEM), conventional microscope and PIMI of the same region on the gold nanodot array. An intact virus particle typically has a dimension of 100 nm, close to the size of a single nanodot and below the optical diffraction limit. As shown in Fig. 1, the point spread function of conventional microscopy smeared out any structural features of the nanodots, making it impossible to recover information about possible virus-like particles that may have attached to the gold nanodots. The PIMI images in the upper right part of Fig. 1, however, show that the scattering photon state pattern of the nanodots exhibits high sensitivity to the attachment of virus particles.

The virus (obtained from Hanbio Technology Co., Ltd), was dissolved in phosphate buffer saline (PBS). After being diluted to a concentration of 1 ng mL^{-1} , a microdroplet of around $1 \mu\text{L}$ was dropped directly on the gold nanodot array. Virus-like particles in the droplet randomly attach to the gold disk due

to Brownian motion and surface adsorption, which breaks the periodic structure of nanodots and results in a local variation of the scattering photon state distribution in the PIMI images. To quantitatively evaluate the influence of virus particles attached the gold nanodisk, the finite difference time domain (FDTD) method was used to simulate PIMI images for the experimental configuration. For comparison, every simulation image is down-sampled to the same pixel density as the experimental image.

Every four neighboring nanodots were grouped as a four-quadrant sensing channel and the purpose of such a four-quadrant sensing configuration is to provide a self-reference of the scattering pattern in each group, suppressing the effects of minor variations in morphology between individual nanodots in the quantitative analysis. Furthermore, this four-quadrant sensing channel will provide a mechanism for fast confirmation of the virus particle location in the whole sensing array, by repetitively analyzing each four-nanodot lattice, as shown in later parts.

For the gold nanodots with an interval of $1 \mu\text{m}$, Fig. 3 shows the comparison between the four bare disks and four gold disks with a virus particle located between them. When the virus particle is in the vicinity of the gold disk, the symmetry of the localized surface plasmon resonance (LSPR) of the gold disk and the symmetry of the scattered photon state distribution are both broken, which is shown clearly in Fig. 3b. This change is barely visible in the conventional microscopy images (Fig. 3a) because the attached virus results in only a small variation in the number of scattered photons (and hence intensity) and cannot be distinguished from the variation caused by background impurities on the substrate or irregularities in the shape of the gold nanodots. On the other hand, polarization state distribution of the scattered photons was drastically changed by the attached virus, not only due to the symmetry breaking of the morphological structure of the gold nanodisk, but also due to the plasmonic emission conditions which are associated with collective electron oscillation in the gold disk. The disturbance of the plasmonic emission further enhances the asymmetry of the photon state distribution induced by the structure of the gold disk–virus combination. As shown in Fig. 3b and c, the simulation and experimental PIMI images $\sin \delta$ and ϕ show a significant variation due to the disturbance of the structural periodicity of the nanodots.

Characteristic curves were plotted along the trajectory shown in Fig. 3b and c, to show the detailed features of the PIMI images. In each figure, regularly appearing peaks in the curve of bare gold disks were disturbed by the attached virus, as indicated by the difference between the red and blue curves in Fig. 3d. In the characteristic curve along certain directions, the difference was significant due to the drastic change of the scattering photon states caused by the virus, *e.g.*, the disappearance of the third peak in the red curve of 5 in Fig. 3d. This change of the characteristic curve can also be utilized to identify which nanodot was attached to the virus and its attaching direction, *i.e.*, from the upper left corner of the attached nanodot.

To quantify the aperiodicity caused by virus, two characteristic parameters were defined to describe the aperiodicity of



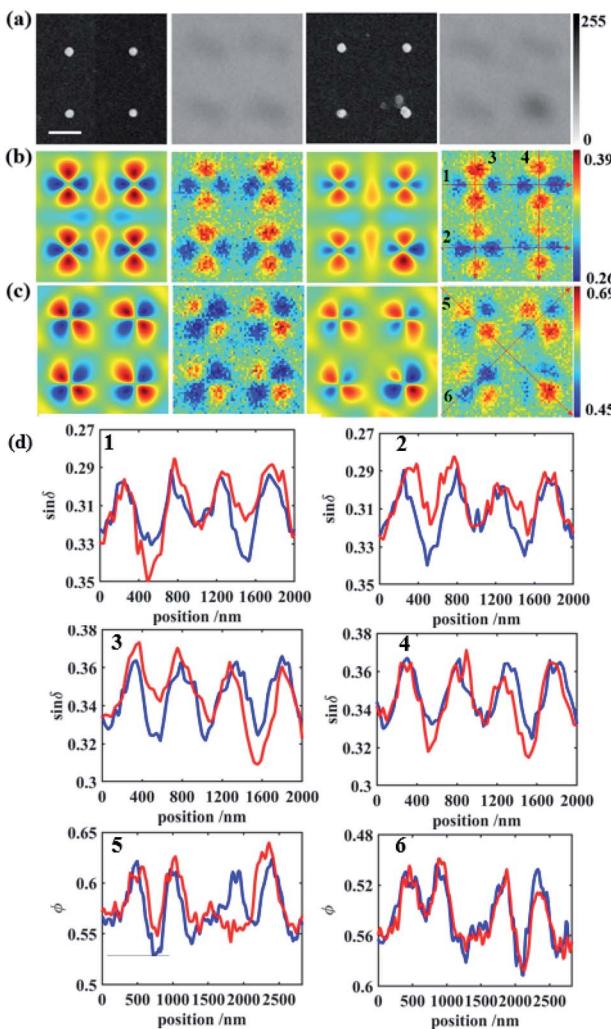


Fig. 3 (a) From left to right: SEM and conventional images of bare gold disks and SEM and conventional images of deposited gold disks, scale bar: 500 nm, (b) from left to right: simulation result of bare gold disks, the experimental result of bare gold disks, the simulation result of deposited gold disks, and the experimental result of deposited gold disks from PIMI $\sin \delta$ images, (c) from left to right: simulation result of bare gold disks, the experimental result of bare gold disks, the simulation result of deposited gold disks, and the experimental result of deposited gold disks from PIMI ϕ images, (d) comparison of characteristic curves of bare and deposited gold disks following the six lines labeled in (b) and (c). Blue curves represent bare gold and red curves represent gold attached to the virus.

$\sin \delta$ and ϕ at every nanodisk site. As shown in Fig. 4a, the photon state distribution at each site is split into four parts, which correspond to the four peaks of the scattering light distribution. We sum the pixel intensities in the four parts and define as follows:

$$Ap_{\sin \delta} = A/B + B/A + C/D + D/C \quad (2)$$

where $Ap_{\sin \delta}$ represents the aperiodicity of the $\sin \delta$ image, and A, B, C , and D represent the sum of the pixel intensities in the four parts. Ap_{ϕ} represents the aperiodicity factor of the ϕ image, and shares the same formula as that in eqn (2) for $Ap_{\sin \delta}$. For a 6

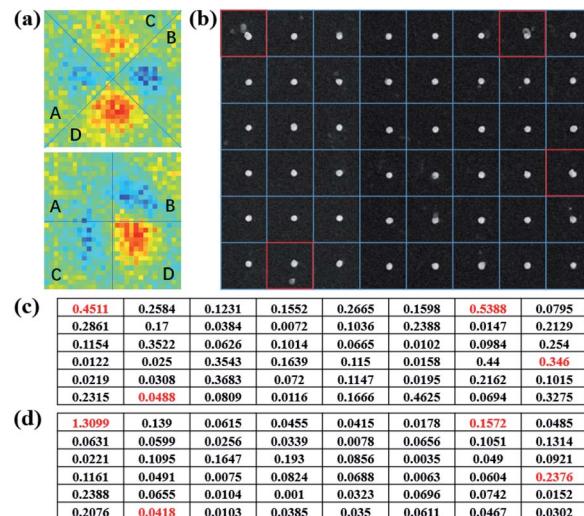


Fig. 4 (a) Partition of PIMI $\sin \delta$ (top) and ϕ images (bottom). (b) Separation of every site in the SEM image. (c) $Ap_{\sin \delta}$ on the array surface. (d) Ap_{ϕ} on the array surface.

$\times 8$ array, $Ap_{\sin \delta}$ and Ap_{ϕ} are shown in Fig. 4c and d, respectively. We can therefore make a preliminary connection between a gold nanodot and virus and values of $Ap_{\sin \delta}$, and Ap_{ϕ} with a higher value being associated with a higher probability of the presence of the virus.

However, although there was a virus present in the square of row 6, and column 2, its $Ap_{\sin \delta}$ and Ap_{ϕ} values were relatively low, *i.e.*, 0.0488 and 0.0418, respectively. This is because the virus was too far away from the gold nanodot to modify the scattered photon distribution. The broken LSPR caused by virus-like particles only occurs at a distance smaller than half wavelength, which means that the period of gold nanodots determined the ratio of the detection area. In this paper, a series of distances between the virus and gold nanodots are simulated to determine the detection radius of nanodots, which is written as R_{det} . After evaluating the values of the corresponding aperiodicity, R_{det} is estimated to be around 150 nm (ESI and Fig. S5†).

Here, we can also calculate the detection limit of the virus concentration. If N viruses are randomly distributed on the substrate and at least one virus is detected by the gold array, the minimum number of viruses should obey the inequality below:

$$N > \frac{-3}{\lg \left(1 - \frac{\pi R_{\text{det}}^2}{L^2} \right)} \quad (3)$$

Here L is the interval of gold nanodots. We assume that a sample volume of 150 μL is reasonable for a real test. When we designed L as 1 μL , the calculated detection limit of the concentration is 96 copies/150 μL . With a smaller L , the detection limit of concentration will decrease to 21 copies/150 μL . The limit is very competitive compared with that of conventional PCR methods, in which the common value is from 10^2 – 10^3 copies/1 mL (more details in the ESI†).

However, the denser gold nanodots will also lead to the overlap and merging of LSPR between two nanodots, which

causes more complex conditions. An interval of 1 μm is the acceptable distance to distinguish the scattering field of two neighboring gold nanodots. To clarify the influence of the interval distance of the nanodisk, another gold nanodot array with a 0.5 μm interval is fabricated, and in such an interval, the light coupling between adjacent dots could not be ignored (Fig. S4†). In this interval, the PIMI ϕ image is rambling and cannot be used to accurately distinguish the virus-like particles. The scattering field of δ from neighboring nanodots merged into one and every four nanodots were grouped as a four-quadrant sensor. As shown in Fig. 5, a virus-like particle will always fall within the sensing area of a certain group of four nanodots and affect its scattering distribution. To find a parameter to define and quantify the aperiodicity in the whole field, such as the condition of 1 μm interval, is difficult to achieve. Here four other characteristic curves are utilized to quantify the influence of virus particles in the vicinity of the gold nanodots in the same region in Fig. 5b and c.

The characteristic curves for $\sin \delta$ images before and after deposition of the virus are shown in Fig. 5d. In general, the amplitude of characteristic curves decreases, due to the reduction of polarization characteristics caused by the virus-like particles. Characteristic curves before deposition mainly possess regular peaks with the same amplitude. After deposition, however, curves crossing the center of the disk connected with the virus generate a height difference among peaks. It is clear that in the curve of 1 and 3 in Fig. 5d, the virus raises the local aperiodicity and hardly influences the peaks in curves 2 and 4 in Fig. 5d. The results of the gold nanodots with an interval of 0.5 μm verify the detection feasibility at a higher ratio

of the detection area. However, we can hardly conduct a quantitative analysis at every site and obtain similar tables of aperiodicity in Fig. 4 due to interaction within adjacent gold nanodots.

Conclusions

In conclusion, we have demonstrated a method for label-free sensing of virus-like particles using a wide-field optical imaging system with a specially designed gold nanodot array. The scattering spatial spectra of photon states was found to be highly sensitive to the presence of virus particles attached to or close to gold nanodots. The presence and location of virus-like particles below the diffraction limit can be sensed by quantitative evaluation of the scattering pattern. This technique has the advantages of requiring minimal sample preparation, purification, and labeling, and eliminates the need for stringent and time-consuming image processing. Furthermore, the gold nanodot array provides a platform with the potential of high throughput parallel sensing in wide-field images, employing each nanodot group in the field of view to act as a sensing channel for viruses. Gold nanoarrays on a silicon substrate can be fabricated using the standard CMOS foundry process or nanoimprint technology, which makes it easy to be fabricated on a large scale at a relatively low cost. Hopefully, this can be developed to handle the complex detection process for infectious diseases. We did not give a complete system with a set judging condition or detection rate. This work currently only confirms the single-particle level detection ability of our method. Also, it should be noted that we used samples with known adenovirus particles to prove the capability of this method as a fast onsite screening technique before detailed analysis of the sample using lab-based techniques such as PCR. In future work, antibody modification of the nanodots will be employed to achieve virus-specific capturing and sensing, allowing accurate screening of samples with unknown particles and contaminations with a similar size and morphology as the target virus particles.

Author contributions

Xuefeng Liu, Jichuan Xiong and Ming Sun designed and oversaw the project. Shengwei Ye and Xiao Sun fabricated the nanodot arrays with assistance from Lianping Hou and John H. Marsh. Weiping Liu designed the nanodot array with assistance from Bin Ni and Bin Xu. Xiao Jin and Heng Zhang performed the measurement of PIMI images and analysis of the experimental results. Heng Zhang prepared the virus particles with the assistance of Xiaofeng Li, Shanhui Li, and Lei Dong. Xiao Jin and Heng Zhang wrote the manuscript with the help of Jamie Jiangmin Hou. All the authors contributed to the discussion and revision of the manuscript.

Conflicts of interest

There are no conflicts to declare.

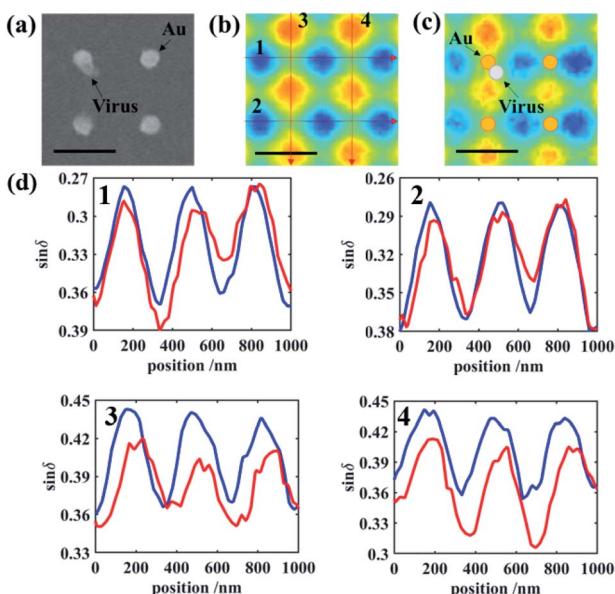


Fig. 5 (a) SEM image of a four-quadrant sensor with a virus, (b) $\sin \delta$ image of the four-quadrant sensor before deposition, (c) $\sin \delta$ image of the four-quadrant sensor after deposition, and the scale bar for (a)–(c) is 0.5 μm , (d) four characteristic curves labeled in (b). Blue curves represent bare gold and red curves represent gold attached to the virus.



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