

High-Performance Waveguide-Integrated Bi₂O₂Se Photodetector for Si Photonic Integrated Circuits

Jianghong Wu,[#] Maoliang Wei,[#] Jianglong Mu,[#] Hui Ma, Chuyu Zhong, Yuting Ye, Chunlei Sun, Bo Tang, Lichun Wang, Junying Li, Xiaomin Xu, Bilu Liu, Lan Li,* and Hongtao Lin*



applied to grow Bi_2O_2Se onto mica, and our developed polycarbonate/polydimethylsiloxane-assisted transfer method enables the clean and intact transfer of Bi_2O_2Se on top of a silicon waveguide. We demonstrated the Bi_2O_2Se/Si waveguide integrated photodetector with a small dark current of 72.9 nA, high responsivity of 3.5 A·W⁻¹, fast rise/decay times of 22/78 ns, and low noise-equivalent power of 15.1 pW·Hz^{-0.5} at an applied voltage of 2 V in the O-band for transverse electric modes. Additionally, a microring resonator is designed for enhancing light-matter interaction, resulting in a wavelength-sensitive photodetector with reduced dark current (15.3 nA at 2 V) and more than a 3-fold enhancement in responsivity at the resonance wavelength, which is suitable for spectrally resolved applications. These results promote the integration of Bi_2O_2Se with a silicon photonic platform and are expected to accelerate the future use of integrated photodetectors in spectroscopy, sensing, and communication applications.

KEYWORDS: Bi₂O₂Se, waveguide-integrated photodetector, silicon photonics, near-infrared, low-dimensional materials

ear-infrared (NIR) photodetectors are a significant component for various applications such as optical communication, sensing, spectroscopy, and so on.^{1–3} Over the past decade, atomically thin low-dimensional materials have drawn considerable attention for NIR photodetectors because of their excellent characteristics such as high carrier mobility, intriguing optical absorption, tunable optoelectronic parameters, and dangling-bond-free surface.^{4–10} Recently, an air-stable material named Bi₂O₂Se, which can be obtained with high quality and large size by chemical vapor deposition (CVD) and vapor solid deposition, has been increasingly used in electronic and optoelectronic devices.^{11–16} With an indirect bandgap of 0.8 eV and a high Hall mobility of ~29 000 cm²·V⁻¹·s⁻¹ at 1.9 K,¹⁷ Bi₂O₂Se has emerged as a promising material for NIR photodetectors.^{18–21}

Up to now, free-space NIR photodetectors based on Bi_2O_2Se have already been reported. For instance, the Bi_2O_2Se -based phototransistor shows a broadband photoresponse ranging from 360 to 1800 nm.²² Although this photodetector's responsivity is up to 22.12 A·W⁻¹ at 1550 nm under the excitation of ultralow optical power (nW), its rise/decay times are 44/181 ms, limiting its application in high-

speed detection. Heterostructures of Bi₂O₂Se with other lowdimensional materials have been constructed to optimize the performance. For example, a photodetector based on PbSe quantum dots/Bi₂O₂Se demonstrates a high responsivity at the wavelength of 2 μ m.²³ PbSe quantum dots work as photoactive materials at 2 μ m, while Bi₂O₂Se serves as the conductive channel. The photogating effect in such a device leads to a high responsivity but slow rise/decay times that are usually more than several seconds. Additionally, a photodetector based on WSe₂/Bi₂O₂Se heterostructure demonstrates a self-powered photovoltaic response with fast rise/decay times of 2.4/2.6 μ s at 532 nm, but its responsivity is ultralow in the telewavelength range.²⁴

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Figure 1. (A) Optical micrograph of Bi_2O_2Se flakes transferred onto PDMS. (B) Raman spectra of Bi_2O_2Se (about 100 nm thick) on mica and a SiO₂ substrate. (C) Scanning electron microscope (SEM) image of Bi_2O_2Se on SiO₂ substrate and its corresponding elemental maps analysis. Scale bar: 1 μ m. (D) High-resolution transmission electron microscope (TEM) image of as-grown Bi_2O_2Se . The inset is the selected area electron diffraction (SAED) pattern of a Bi_2O_2Se flake.

The trade-off between the responsivity and rise/decay times in photodetectors can be overcome by a waveguide-integrated architecture, benefiting from the orthogonal directions in the path of photogenerated carriers and light propagation. Indeed, waveguide-integrated photodetectors not only possess the potential to achieve high-speed detection but also provide superior planar integration capability for on-chip applications. Moreover, a high signal-to-noise ratio could be achieved due to reduced detection volume compared with their free-space counterparts.²⁵ Among a wide variety of low-dimensional materials, graphene is the most commonly used material for waveguide-integrated photodetectors.²⁶⁻³³ However, the characteristic of zero bandgap in graphene brings about a large dark current, low photocurrent-to-dark current ratio, and high noise in photodetectors. Therefore, low-dimensional semiconductors with a narrow bandgap such as black phosphorus (BP) and MoTe₂ are alternative candidates for Si waveguide-integrated photodetectors, 3^{4-38} but the low stability of BP prevents it from constructing high-performance photodetectors in air. $MoTe_2$ is air-stable, but a bandgap of 1.04 eV leads to limited absorption when the operational wavelength is larger than 1.26 μ m.³⁹⁻⁴¹ Given its air stability, high carrier mobility, and narrow bandgap, Bi₂O₂Se is attractive to be integrated with a Si waveguide to realize a high-performance NIR photodetector. However, there are currently no reports on the waveguideintegrated photodetector based on Bi₂O₂Se. Transferring Bi₂O₂Se from f-mica to a Si waveguide is one key point for obtaining a high-performance waveguide-integrated photodetector. At present, wet transfer with the assistance of a

polymer film such as poly(methyl methacrylate) (PMMA) is the most common strategy to peel off Bi_2O_2Se from f-mica.^{12,23} But this approach is cumbersome and not suitable for preparing a waveguide-integrated photodetector. On the contrary, the dry transfer process facilitates the alignment of low-dimensional materials with the underlying nanostructured waveguides.⁴²

In this paper, we demonstrate both the waveguide-integrated and microring resonator (MRR)-integrated photodetectors based on Bi₂O₂Se. First, we synthesize Bi₂O₂Se on an f-mica substrate by the CVD method, and the dry transfer method assisted by polydimethylsiloxane (PDMS) or polycarbonate (PC)/PDMS is used to transfer Bi₂O₂Se from f-mica to the Si waveguide and MRR. Clean Bi₂O₂Se flakes are transferred on top of the Si waveguide, where light propagating in the waveguide can be evanescently coupled to Bi₂O₂Se. The waveguide-integrated photodetector shows a small dark current of 72.9 nA, high responsivity of 3.5 A·W⁻¹, fast rise/ decay times of 22/78 ns, and low noise-equivalent power (NEP) of 15.1 pW·Hz^{-0.5} at an applied voltage of 2 V in the Oband. The MRR-integrated photodetector shows a wavelengthsensitive photoresponse. The photocurrent increases about 3fold at the resonance wavelength, which is induced by the enhanced light-matter interaction within the MRR structure. Owing to the thinner thickness and shorter length of Bi₂O₂Se that has been applied on the MRR, the MRR-integrated photodetector exhibits a reduced dark current (15.3 nA at 2 V) and comparable responsivity (1.83 $A \cdot W^{-1}$).



Figure 2. (A) Diagram of the waveguide-integrated Bi_2O_2Se photodetector. (B) Optical image of a typical integrated Bi_2O_2Se photodetector on an unbalanced MZI. (C) Transmission spectra of the unbalanced MZI with and without Bi_2O_2Se (73.6 nm thick).

RESULTS AND DISCUSSION

Bi₂O₂Se, possessing a layered structure, consists of covalently bonded $[Bi_2O_2]_n^{2n+}$ and $[Se]_n^{2n-}$ layers that stack alternately in a tetragonal unit cell with weak electrostatic force, which is similar to f-mica. We use f-mica as a substrate for growing Bi_2O_2Se in this work (see details in Methods). The size of the obtained Bi2O2Se flakes, with square or rectangular shapes, ranges from several microns to more than 20 μ m. We transfer Bi_2O_2Se onto the SiO₂ substrate with the assistance of PDMS, and the optical micrograph of Bi₂O₂Se on PDMS is shown in Figure 1A, in which no cracks have been observed in Bi₂O₂Se. Raman spectra of Bi₂O₂Se on mica and a SiO₂ substrate are illustrated in Figure 1B, in which a characteristic A_{1g} peak at about 159 cm⁻¹ is observed before and after transfer. The characteristic peak of f-mica is not shown due to the limited penetration depth of Raman light sources into a thick Bi₂O₂Se flake. The energy-dispersive X-ray spectroscopy (EDS) mappings of Bi₂O₂Se on a silicon-on-insulator (SOI) substrate (Figure 1C) reveal the uniform distribution of Bi, Se, and O elements. As for the image of O element distribution in Figure 1C, the SiO_2 substrate possesses a high percentage of O atoms compared with Bi₂O₂Se, leading to a bright color. A shallow dark line in the SiO₂ substrate is the bottom Si waveguide (Figure 1C). We transfer Bi₂O₂Se onto a Cu grid for TEM characterization. The high-magnification TEM image and the SAED pattern shown in Figure 1D demonstrate the singlecrystal nature of our Bi₂O₂Se flakes. These measurements in Figure 1 indicate that the CVD-grown Bi₂O₂Se exhibits high quality, and the applied transfer methods do not damage the quality of Bi₂O₂Se.

The diagram of the waveguide-integrated Bi₂O₂Se photodetector is conceptually illustrated in Figure 2A, which consists of a couple of grating couplers, a single-mode waveguide, multimode interference (MMI), a pair of Au pads, and a Bi₂O₂Se flake. The grating couplers in our experiment only support TE (transverse electric) modes, and all results of optoelectronic response in this article correspond to TE modes as well. The passive optical devices are fabricated on 220 nm SOI wafers with multiproject wafer (MPW) processes involved, and the rib Si waveguide with a width of 397 nm and an etching depth of 151 nm (Figure S1) is planarized by backfilling with SiO_2 to protect Bi_2O_2Se from cracking at the edge of the Si waveguide. A pair of Au pads is located around one arm of an unbalanced Mach-Zehnder interferometer (MZI) with a suitable distance, avoiding optical loss and ensuring an effective collection of the photogenerated carriers. Bi₂O₂Se is well aligned to the gap between two Au pads on top of the waveguide by the PDMS-assisted transfer method. A detailed flowchart of the fabrication processes is illustrated in Figure S2. The optical image shown in Figure 2B represents a

typical integrated photodetector where Bi_2O_2Se can absorb photons in the waveguide *via* evanescent coupling.

Optical absorption is a significant factor of photodetectors, which is affected by the thickness of photoactive materials and the degree of overlap between photoactive materials and the optical field. To determine the absorption coefficient of the Bi₂O₂Se/Si hybrid waveguide, we transfer Bi₂O₂Se onto one arm of the unbalanced MZI. As for devices based on the straight waveguide, the power difference between the output signal with and without photoactive materials is an approach to obtain the absorption coefficient. However, the practical coupling efficiency of the same grating coupler may be different in two operations, leading to a measurement error. This can be avoided by assessing the optical absorption information from analyzing the extinction ratio (ER) of the unbalanced MZI in the transmission spectra.⁴³ In an unbalanced MZI, a pair of MMIs works as a 3 dB beam splitter and combiner, respectively. Six serially cascaded MMIs (Figure S3A) are designed to better investigate and quantify the MMI's performance. The measured result shown in Figure S3B demonstrates that the beam splitter could equally allocate half of the input power to each output port of the 1×2 MMI. As shown in the inset of Figure S3B, the blue and red points are the measured power of two output ports of the sixth MMI. The two sets of data points overlap well, indicating the outstanding power equilibrium of the designed 1×2 MMI. The interference occurs when light passes through the beam combiner. Since the extinction ratio is more than 30 dB (before transferring Bi₂O₂Se), the propagation loss difference between the two arms could be neglected. Thereby the output power of an unbalanced MZI can be expressed as

$$P_{\rm out} = \left(\frac{P_{\rm in}}{4}\right) [e^{-\alpha l_1} + e^{-\alpha l_2} + 2e^{-\alpha (l_1 + l_2)/2} \cos(2\pi n_{\rm eff} \Delta L/\lambda)]$$

where $P_{\rm in}$ is the input power, α is the absorption coefficient of the Bi₂O₂Se/Si hybrid waveguide, and l_1 and l_2 are the lengths of the Bi₂O₂Se/Si hybrid waveguide on two arms, respectively. In our experiment, Bi₂O₂Se integrates with only one arm; thereby, l_2 is zero. ΔL is their length difference, λ is the wavelength of the incident light, and $n_{\rm eff}$ is the effective refractive index. Figure 2C shows the interferometric transmission spectra from which the peak position and ER of interference fringes of the unbalanced MZI are acquired accordingly. And there are apparent changes in both the ER and peak position upon the applying of Bi₂O₂Se on the MZI owing to its influence on optical absorption and the effective refractive index. In fact, the ER can be expressed as⁴³



Figure 3. (A) Current–voltage (IV) curves of a Bi₂O₂Se photodetector in the dark and under illumination under different optical power at 1260 nm. (B) Energy band diagram of a Bi₂O₂Se photodetector at equilibrium, under illumination, and under an externally applied voltage. (C) Measured photocurrent (left-hand axis) and calculated responsivity (right-hand axis) of a Bi₂O₂Se photodetector under different optical power at 1260 nm at 2 V. (D) Calculated responsivity (left-hand axis) and external quantum efficiency (right-hand axis) of a Bi₂O₂Se photodetector under different applied voltages and an input power of 2.04 μ W at 1260 nm. (E) Normalized photocurrent-to-dark-current ratio (NPDR) under different applied voltages and an input power of 2.04 μ W at 1260 nm. (F) Spectral photoresponse of a Bi₂O₂Se photodetector at a wavelength ranging from 1260 to 1310 nm. The thickness of Bi₂O₂Se in the photodetector characterized is about 142.5 nm.

$$ER = \frac{P_{max}}{P_{min}} = \frac{e^{-\alpha l_1} + 1 + 2e^{-\alpha l_1/2}}{e^{-\alpha l_1} + 1 - 2e^{-\alpha l_1/2}} = \left(\frac{1 + e^{-\alpha l_1/2}}{1 - e^{-\alpha l_1/2}}\right)^2$$

Thus the α of the Bi₂O₂Se/Si hybrid waveguide is calculated to be 0.07 dB/ μ m with the Bi₂O₂Se thickness of 73.6 nm. When the thickness of Bi₂O₂Se is 38.9 nm, the calculated α is about 0.03 dB/ μ m (Figure S4), indicating that designing a lighttrapping structure to enhance optical absorption is necessary for a photodetector based on Bi₂O₂Se flakes. Furthermore, the SiO₂ layer being about 79.4 nm between Bi₂O₂Se and the Si waveguide degrades the α in our device, which is consistent with the results of the graphene/Si waveguide structure.⁴³

The static photoresponse of the straight waveguideintegrated photodetectors based on Bi₂O₂Se with a thickness of 142.5 nm (Figure S5A) is characterized by a continuous wave laser, and the optical and SEM images of this photodetector are shown in Figure S6. Figure 3A illustrates the measured IV curves of an integrated photodetector under the applied voltage from -2 to 2 V in the dark and under illumination at the wavelength of 1260 nm. The dark current of this photodetector is about 72.9 nA at an applied voltage of 2 V. The current increases to 3.1 μ A under illumination with an optical power of 2.04 μ W, and the current rises gradually with the increasing optical power, indicating that Bi₂O₂Se is suitable for a photodetector in the O-band. The nonlinear behavior shown in the IV curves can be attributed to Schottky contact between Bi₂O₂Se and a metal. In our devices, Au pads are deposited on the flat substrate before the transfer of Bi₂O₂Se, which can reduce Fermi-level pinning and chemical disorder at the interface of Au and $Bi_2O_2Se^{.44}$ In this case, the Schottky barrier depends mainly on the work function of Au. The energy band diagram of the Au/Bi₂O₂Se/Au photodetector is shown in Figure 3B. The work function of Au is larger than the electron affinity of n-Bi2O2Se; thereby electrons in Bi2O2Se drift to Au and result in an upward energy band bending at the

interface. This bending energy band suppresses the carrier transport and brings about a small dark current.

We then characterized the photoresponse under different optical power ranging from 0.14 to 1809.07 μ W, and the measured photocurrent and calculated responsivity $(R = I_{\rm pb}/$ $P_{\rm in}$) are shown in Figure 3C. $I_{\rm ph}$ and $P_{\rm in}$ are the photocurrent and input optical power launched into the photodetector, respectively. Responsivity decreases from 3.5 A·W⁻¹ to 40 mA· W^{-1} when the optical power increases from 0.14 μW to 1809.07 μ W. This nonlinear relationship between photocurrent and optical power probably results from trap states in Bi₂O₂Se, which leads to photocurrent gain under weak optical power. On the contrary, trap states are filled under high optical power, resulting in lower responsivity. We calculated the responsivity under the applied voltage ranging from -2 to 2 V, as shown in Figure 3D. The responsivity increases with the rising applied voltage, and it reaches up to 1.47 A·W⁻¹ at an applied voltage of 2 V when the input power is 2.04 μ W. The corresponding external quantum efficiency (EQE) can be calculated by EQE = $R \times hc/(q\lambda)$, where h, c, q, and λ are Planck's constant, the speed of light in a vacuum, elementary electron charge, and operational wavelength, respectively. EQE possesses the same tendency with responsivity, and it increases to 144.33% at an applied voltage of 2 V. This tendency may be attributed to the higher tunneling current induced by the lower barrier height of the Au/Bi₂O₂Se junction. And more efficient separation-collection of photoinduced carriers can be obtained at a larger applied voltage. The NPDR ((NPDR = $(I_{\rm p}/I_{\rm d})/P_{\rm in} = (I_{\rm p}/P_{\rm in})/I_{\rm d} = R/I_{\rm d})$ is estimated to be 3 \times 10⁴ mW^{-1} at the applied voltage of -2 V when the input power is 2.04 μ W in Figure 3E, which is several orders of magnitude higher than that of graphene, BP, and even a Ge waveguide-integrated photodetector.^{36,45,46} This high NPDR is attributed to both a small dark current and large responsivity, and it can be optimized through tuning the carrier concentration to a minimum by electrostatic doping. Some graphene photo-



Figure 4. (A) Dynamic photoresponse of a Bi_2O_2Se photodetector under the excitation of the modulated laser at 1310 nm. (B) 1/f noise spectra of a Bi_2O_2Se photodetector under different applied voltages. (C) Spectral noise-equivalent power (NEP) at a wavelength ranging from 1260 to 1310 nm under an applied voltage of 2 V; left-hand axis corresponds to the calculated NEP limited by 1/f noise, and the right-hand axis corresponds to the calculated NEP limited NEP limited by shot and thermal noise. (D) Summary of the device performance for some waveguide-integrated photodetectors based on low-dimensional materials. The blue arrow represents the better performance. The thickness of Bi_2O_2Se in the photodetector is about 142.5 nm.

detectors possess zero dark current in the self-driven mode, but responsivity is weak in this situation. Therefore, the Bi_2O_2Se photodetector shows obvious advantages in terms of both responsivity and dark current.

We measured the spectral photoresponse as well, and the corresponding results are illustrated in Figure 3F. The responsivity fluctuates slightly when the wavelength increases from 1260 nm to 1310 nm, which can be attributed to the smaller gain at larger optical power and more efficient photon-to-electron transition at shorter wavelengths in the O-band considering Bi₂O₂Se's indirect bandgap of 0.8 eV.⁴⁷ At a wavelength of 1330 nm, the responsivity of the proposed photodetector is 1.75 A·W⁻¹. In our design, the adopted grating couplers have limited operation optical bandwidth. Taking advantage of the edge coupling method, we can replace the grating couplers with end-face coupling waveguides to ensure the functionality of the Bi₂O₂Se-based photodetector with a broader optical bandwidth.

The dynamic response (Figure 4A) is measured under the excitation of the modulated laser, and a magnified plot of one response cycle is shown in (Figure S7). The rise and decay times are 22 and 78 ns, respectively, benefiting from the fast carrier mobility of Bi₂O₂Se and the short channel length between two Au pads. Furthermore, the noise-equivalent power (NEP) characterizing the detection limit is another significant parameter for a photodetector. The total current noise of a photodetector mainly includes the 1/f noise, shot noise, and thermal noise. Hooge's empirical relationship is usually used for analyzing 1/f noise, which can be expressed as⁴⁸ $S_i = Ai^{\alpha}/f^{\beta}$, where *i*, *f*, and *A* are the channel current, the frequency, and noise amplitude, respectively. Figure 4B is the

measured 1/f noise spectral density, demonstrating that our photodetectors possess typical characteristics of 1/f noise. The measured 1/f noise decreases from 2.9 × 10^{-19} Å²·Hz⁻¹ at a frequency of 1 Hz to 5.6 \times 10⁻²⁴A²·Hz⁻¹ at 100 kHz at an applied voltage of 2.0 V. A higher external applied voltage brings about an optimal responsivity but leads to a higher 1/fnoise. The shot noise can be expressed as $S_s = 2qi_d\Delta f_t^{49}$ where $i_{\rm d}$ and Δf are the dark current and bandwidth, respectively. The dark current of the proposed Bi2O2Se-based waveguideintegrated photodetector is 72.9 nA at an applied voltage of 2 V, and the calculated shot noise is $2.3 \times 10^{-26} \text{ A}^2 \cdot \text{Hz}^{-1}$. The thermal noise is calculated to be $6.0 \times 10^{-28} \text{ A}^2 \cdot \text{Hz}^{-1}$ by $S_t =$ $4kT\Delta f/R_{st}^{49}$ where k, t, and R_{s} are the Boltzmann constant, the temperature (about 300 K in our experiment), and the device's resistance. Therefore, 1/f noise is the main noise at low frequency, and the NEP is estimated to be 15.1 pW·Hz^{-0.5} at a frequency and wavelength of 1000 Hz and 1260 nm, respectively. On the contrary, the shot noise is dominant at a higher frequency, and NEP is estimated to be 0.1 $pW \cdot Hz^{-0.5}$. The spectral NEP at the wavelength ranging from 1260 to 1310 nm is shown in Figure 4C, indicating a small detectable power within this wavelength range.

As shown in Figure 4D, we list several reported waveguideintegrated photodetectors based on low-dimensional materials. ${}^{34,37,39,40,50-53}$ The proposed photodetector based on Bi₂O₂Se not only possesses a low dark current but also has high responsivity at the O band, exhibiting potential applications in the fields of spectroscopy, sensing, and communication as high-performance photodetectors integrated onto silicon photonic platforms.



Figure 5. (A) Schematic of the MRR-enhanced photodetector showing Bi_2O_2Se integrated with a planarized microring. (B) Transmission spectra of the MRR with and without Bi_2O_2Se . Inset: Zoom-in of the spectra. (C) Photocurrent at resonance and off-resonance wavelength. The inset is the photocurrent ratio at resonance and off-resonance wavelength. (D) Spectral response of the Bi_2O_2Se -based MRR-enhanced photodetector. (E) *IV* curves of the Bi_2O_2Se -based MRR-enhanced photodetector in the dark and under illumination with different optical power at resonance wavelength. (F) Calculated responsivity (left-hand axis) and external quantum efficiency (right-hand axis) of a Bi_2O_2Se photodetector under different applied voltages and an optical power of 0.86 μ W. The thickness of Bi_2O_2Se in the photodetector is about 63 nm.

Furthermore, Bi₂O₂Se is applied onto a silicon MRR to achieve a high-performance integrated photodetector. On the one hand, the enhanced light-matter interaction by a MRR makes it possible to realize quantum efficiency comparable to a thicker and longer Bi2O2Se photodetector on the straight waveguide.^{37,54,55} On the other hand, such a cavity-enhanced photodetector is wavelength sensitive, which plays an important role in spectrally resolved applications such as wavelength division multiplexing systems.^{56,57} Figure 5A is a schematic of the MRR-enhanced photodetector, indicating a pair of Au pads close to the MRR and Bi₂O₂Se on top of the Au pads and the MRR. Furthermore, FIB-SEM of this photodetector is shown in Figure S1B,C. Typical transmission spectra of an MRR are shown inFigure 5B. The transmission spectra of an MRR with different gaps (from 200 to 400 nm) are measured (Figure S8A), from which the gap under the critical coupling condition is confirmed to be less than 300 nm. The cavity loss increases after introducing Bi₂O₂Se, so it is difficult to observe the resonance peak in the spectra of the Bi₂O₂Se photodetector integrated with an undercoupled MRR (Figure S8B). Thereby, overcoupled MRRs are used to ensure resonance peaks with high extinctions upon applying Bi₂O₂Se onto a silicon MRR. We choose the MRR with a gap of 200 nm for the photodetector, and the corresponding transmission spectra without and with Bi₂O₂Se are shown in Figure 5B. The quality factor of the MRR decreases from 3.5×10^4 to $2.4 \times$ 10^4 , which can be attributed to the additional optical loss that Bi_2O_2Se brings to the cavity. And for the MRR with Bi_2O_2Se , the number of times that a resonance photon passes through the cavity is calculated to be 754 (the detailed process is shown in Supporting Information 9), and the photon circulation brings about an enhanced photocurrent at resonance wavelength. The thickness of the Bi₂O₂Se flake in this photodetector is about 63.0 nm (Figure S5B), and the optical and SEM images of this photodetector are shown in Figure S6. Moreover, the *n* value of Bi_2O_2Se is estimated to be 4.079 from the resonance peak shift after transferring Bi₂O₂Se, and the simulated modal intensity for fundamental TE modes in the

Bi₂O₂Se/Si waveguide is shown in Figure S9. Shown in Figure 5C are the IV curves of the photodetector at resonance and offresonance wavelength. The inset is the ratio between current at resonance and off-resonance wavelengths, indicating a maximum enhancement of 3.4 (at an applied voltage of -2V) in photocurrent at resonance wavelength. This photodetector shows a 5-fold decrease in the dark current (15.3 nA@2 V) compared with the straight waveguide-integrated photodetector, which is partially attributed to the thinner thickness. The spectral response of the MRR-enhanced photodetector at a wavelength ranging from 1260 to 1265 nm is shown in Figure 5D, where the wavelength-variation photoresponse is observed, and the maximum photocurrent is achieved at the resonance wavelength at an applied voltage of 2 V. The photocurrent exhibits more than a 3-fold enhancement at resonance wavelength, comparing with that at off-resonance wavelength. IV curves of an MRR-enhanced photodetector under different power at the resonance wavelength are shown in Figure 5E. The current increases gradually with the increasing optical power. The calculated responsivity and EQE at different applied voltage are shown in Figure 5F. And a responsivity of 1.83 A·W⁻¹ and EQE of 179.6% at the resonance wavelength (at an applied voltage of 2 V) are obtained. All in all, this MRR-enhanced photodetector shows an advantage for optimizing a photodetector, and the wavelength-sensitive characteristics have the potential in a wavelength division multiplexing system for on-chip spectroscopy, sensing, and communication applications.

CONCLUSIONS

In conclusion, we demonstrated the silicon waveguide and MRR-integrated Bi_2O_2Se photodetectors operating in the Oband. Bi_2O_2Se interacts with the light propagating along the waveguide through evanescent field coupling in this type of photodetector. The optical absorption of the Bi_2O_2Se/Si hybrid waveguide is measured by an unbalanced Mach– Zehnder interferometer. As a result, we showed its broadband operation capability in the O-band with a responsivity of 3.5 A. W^{-1} at 1260 nm and an applied voltage of 2 V, a dark current of 72.9 nA, and rise/decay times of 22/78 ns, respectively. Moreover, we also analyzed noise characteristics and found that 1/*f* noise is the dominant noise when the frequency is below 10⁵ Hz, and the corresponding NEP is about 15.1 pW Hz^{-0.5}. The photodetection is further enhanced by integrating Bi₂O₂Se with MRR cavities to increase the optical absorption. The MRR-enhanced photodetector shows a 3-fold improvement in photocurrent at the resonance wavelength comparing with that at the off-resonance wavelength, suitable for devices with a small volume of photoactive materials. These results promote the integration of Bi₂O₂Se photodetectors on a silicon photonic platform and hold great promise for extensive applications in spectroscopy, sensing, and communication.

METHODS

Synthesis of Bi₂O₂Se. Bi₂O₂Se nanoflakes were grown by lowpressure CVD (LPCVD) processes in the one-inch diameter quartz tube (length of 120 cm) located inside a horizontal furnace. A specific mass ratio of Bi₂O₃ (purchased from Macklin, 99.99%) and Bi₂Se₃ (purchased from Aladdin, 99.99%) as reaction precursors was put into two quartz boats, respectively. Bi2O3 powder was sited at the hot center of the furnace, while Bi₂Se₃ powder was placed upstream at a distance of 10 cm, and the freshly cleaved mica substrates (1 cm \times 1 cm) were placed at 10-12 cm downstream. The quartz tube was evacuated using a mechanical pump in all growth processes to avoid the ambient environment. High-purity argon (Ar) was used as the carrier gas to transport precursors' vapor from upstream to downstream. The furnace was heated to the goal temperature of 700 °C with a heating rate of 30-35 °C·min⁻¹ and kept for 1–10 min. Meanwhile, the flow rate of Ar was 200 standard cubic centimeters per minute (sccm), and the growth pressure in the quartz tube was about 2 Torr. Finally, the furnace cooled naturally to ambient temperature.

Materials Characterization and Transfer. Raman spectra (WITec, Alpha300R), AFM (Oxford Instruments, Cypher ES), TEM (Thermo Fisher Scientific, Talos F200X G2), and SEM (Hitachi, Regulus 8230) were used to evaluate the quality and characteristics of the obtained Bi2O2Se. The flexible PDMS was attached and pressed to make a strong adhesion between Bi₂O₂Se flakes and PDMS. Next, PDMS along with Bi2O2Se was peeled off from the f-mica with the help of the water droplet (5–10 μ L). After that, Bi₂O₂Se was aligned with the Au electrode and waveguide under a microscope, and PDMS was removed after pressing Bi₂O₂Se onto the Au electrode and being heated for 10 min (60 °C). Finally, Bi₂O₂Se was transferred onto the Au electrode upon the removal of PDMS. However, there were still lots of Bi₂O₂Se flakes on the f-mica after repeating these steps several times, but PMDS-assisted transfer was not efficient enough at this stage. We proposed a PC-assisted approach to peel off Bi₂O₂Se from the f-mica more efficiently. PC (dissolved in chloroform) was dissolved in chloroform (PC/ chloroform = 6/94) to obtain a transparent solution. This transparent solution was spin-coated onto the surface of the mica that was fixed on a glass slide, which was heated for 60 s (90 °C). With the assistance of a water droplet (5–10 μ L), PC along with Bi₂O₂Se could be peeled off from f-mica, and this exploited PC film was fixed on a blue tape. Afterward, PDMS was attached and pressed onto PC, transferring Bi₂O₂Se from PC to PDMS. The next step was the same as the PDMS-assisted transfer.

Device Fabrication. The passive Si photonic devices were fabricated with standard 220-nm-SOI MPW processes in IMECAS. On top of the device is the buried SiO₂ layer with a 2 μ m thickness, which was chemically and mechanically polished to a flat surface. Metal pads in the devices were patterned by electron-beam lithography (Raith, Voyager) followed by deposition of metal pads (Ti/Au, 5/50 nm) by electron-beam evaporation (ULvac Vacuum Technology (Suzhou) Co., Ltd., Ei-SZ) and a standard lift-off process.

After depositing the metal pads, Bi_2O_2Se was transferred with the help of a PC/PDMS film.

Device Measurement. Electrical measurements were collected by a semiconductor analyzer (PDA, FS-Pro). Static photoresponse measurements (Figure S10) were done under the excitation of a tunable semiconductor laser (Santec Corporation, TSL-550) with tunable wavelength ranging from 1260 to 1620 nm. A variable optical attenuator (Thorlab, V1550A) was used to tune the power of the incident light. A polarization controller was applied to tune the polarization of the incident light and improve the coupling efficiency of the grating coupler. Dynamic photoresponse measurements (Figure S11) were under the excitation of a modulated laser (1310 nm) driven by a signal generator (SDG6032X-E), and the output signal was obtained by an oscilloscope (SDS5104X).

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsnano.1c04359.

Cross-section SEM images of the rib waveguide and Bi_2O_2Se photodetector; detailed flowchart of the fabrication processes; optical image and optical performance of cascading MMIs; optical absorption of Bi_2O_2Se ; thickness of Bi_2O_2Se ; optical and SEM images for photodetector; dynamic optoelectronic response of the Bi_2O_2Se photodetector; transmission spectra of the microring resonator; the relationship between performance enhancement and the cavity Q; the simulated modal intensity for the Bi_2O_2Se photodetector; static photoresponse measurement setup; dynamic photoresponse measurement setup; (PDF)

AUTHOR INFORMATION

Corresponding Authors

- Hongtao Lin State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China; orcid.org/0000-0001-7432-3644; Email: hometown@zju.edu.cn
- Lan Li Key Laboratory of 3D Micro/Nano Fabrication and Characterization of Zhejiang Province, School of Engineering, Westlake University, Hangzhou 310024, China; Institute of Advanced Technology, Westlake Institute for Advanced Study, Hangzhou 310024, China; orcid.org/0000-0002-9097-9157; Email: lilan@westlake.edu.cn

Authors

- Jianghong Wu State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China; Key Laboratory of 3D Micro/Nano Fabrication and Characterization of Zhejiang Province, School of Engineering, Westlake University, Hangzhou 310024, China; Institute of Advanced Technology, Westlake Institute for Advanced Study, Hangzhou 310024, China
- **Maoliang Wei** State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China
- Jianglong Mu Shenzhen Geim Graphene Center, Tsinghua-Berkeley Shenzhen Institute and Tsinghua Shenzhen International Graduate School, Tsinghua University, Shenzhen, Guangdong 518055, China

- Hui Ma State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China
- **Chuyu Zhong** State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China
- Yuting Ye Key Laboratory of 3D Micro/Nano Fabrication and Characterization of Zhejiang Province, School of Engineering, Westlake University, Hangzhou 310024, China; Institute of Advanced Technology, Westlake Institute for Advanced Study, Hangzhou 310024, China
- Chunlei Sun Key Laboratory of 3D Micro/Nano Fabrication and Characterization of Zhejiang Province, School of Engineering, Westlake University, Hangzhou 310024, China; Institute of Advanced Technology, Westlake Institute for Advanced Study, Hangzhou 310024, China
- Bo Tang Institute of Microelectronics, Chinese Academic Society, Beijing 100029, China
- Lichun Wang State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China
- Junying Li State Key Laboratory of Modern Optical Instrumentation, College of Information Science and Electronic Engineering, Zhejiang University, Hangzhou 310027, China
- Xiaomin Xu Shenzhen Geim Graphene Center, Tsinghua-Berkeley Shenzhen Institute and Tsinghua Shenzhen International Graduate School, Tsinghua University, Shenzhen, Guangdong 518055, China
- Bilu Liu Shenzhen Geim Graphene Center, Tsinghua-Berkeley Shenzhen Institute and Tsinghua Shenzhen International Graduate School, Tsinghua University, Shenzhen, Guangdong 518055, China; Orcid.org/0000-0002-7274-5752

Complete contact information is available at: https://pubs.acs.org/10.1021/acsnano.1c04359

Author Contributions

H.L., L.L, and J.W. conceived the project. X.X., B.L., and J.M. grew Bi_2O_2Se . J.W., M.W., H.M., C.Z., Y.Y., L.W., J.L., L.W., and B.T. fabricated these devices. C.S. and M.W. designed the passive optical devices. J.W. performed optical and optoelectronic measurements. H.L., L.L, and J.W. analyzed the data and wrote the manuscript. All authors commented on the manuscript.

Author Contributions

[#]J.W., M.W., and J.M. contributed equally to this work.

Notes

The authors declare no competing financial interest.

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