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#### **Research Article**

Ning Liu, Qiang Liu, Yutian Lin, Zhihong Zhu\* and Ken Liu\*

# Second-harmonic generation in NbOI<sub>2</sub>-integrated silicon nitride microdisk resonators

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Abstract: In recent years, two-dimensional (2D) niobium oxide dihalides (e.g., NbOI<sub>2</sub>) have garnered significant research interest in nonlinear photonics due to their prominent second-order nonlinear optical properties. Integrating these materials with high-quality-factor optical microcavities represents a crucial approach for developing high-performance on-chip nonlinear optical devices. This work demonstrates NbOI2-integrated silicon nitride (Si3N4) microdisk resonators that achieve second-harmonic generation under low-power (sub-milliwatt) continuous-wave laser pumping, leveraging the superior second-order nonlinearity of NbOI<sub>2</sub> and the strong optical field confinement capability of Si<sub>3</sub>N<sub>4</sub> microdisks. The conversion efficiency of the device is calculated to be about 0.024 %/W. The intrinsic lack of inversion symmetry in NbOI2 crystals avoids the laborious layer-number-dependent symmetry screening typically required for other 2D materials, while the developed van der Waals transfer technique provides a universal strategy for integrating niobium oxide dihalides with photonic microcavities. This study not only establishes a material-photon co-design strategy for on-chip nonlinear light sources but also lays a critical foundation for advancing quantum photonic chips and on-chip metrology systems.

**Keywords:** second-harmonic generation; niobium oxide diiodides; silicon nitride; microdisks; resonators

liukener@163.com (K. Liu).

https://orcid.org/0000-0003-3199-9323 (Z. Zhu). https://orcid.org/0000-0003-1941-1160 (K. Liu)

**Ning Liu, Qiang Liu and Yutian Lin**, College of Advanced Interdisciplinary Studies & Hunan Provincial Key Laboratory of Novel Nanooptoelectronic Information Materials and Devices and Nanhu Laser Laboratory, National University of Defense Technology, Changsha 410073, China. https://orcid.org/0000-0003-2632-0849 (N. Liu)

#### 1 Introduction

As one of the central processes in second-order nonlinear optical effects, second-harmonic generation (SHG) occupies a critical position in integrated photonics, with its applications extensively permeating cutting-edge fields such as materials science [1]-[3], biomedicine [4], [5], and laser technology [6]-[8]. High-quality-factor (Q factor) whispering gallery mode (WGM) optical microcavities [9], [10], featuring strong optical field confinement capability, significantly enhance light-matter interaction and serve as an ideal platform for the study of integrated nonlinear optics. However, for the widely utilized silicon nitride (Si<sub>2</sub>N<sub>4</sub>) photonic platform, the intrinsic centrosymmetry of the material fundamentally prohibits direct SHG implementation in Si<sub>3</sub>N<sub>4</sub> microdisk resonators. SHG has been demonstrated in symmetric materials through methods such as interfacial effects [11], [12] or optical-induced nonlinearity [13]–[15]. However, achieving efficient SHG under moderate pump power remains a key bottleneck, which limits the expansion of nonlinear functionalities in Si<sub>3</sub>N<sub>4</sub> microcavity platforms.

Two-dimensional (2D) niobium oxide dihalides (NbOX<sub>2</sub>, X = Cl, Br, I) [16], [17] offer novel opportunities for integrated nonlinear photonics due to their unique layered structure and physical properties. First, the intrinsic lack of inversion symmetry in NbOX2 avoids a demanding choice of the layer parity of the 2D materials [18], [19]; Second, via van der Waals (vdW) forces, NbOX<sub>2</sub> enables universal integration with silicon photonic devices (e.g., Si<sub>3</sub>N<sub>4</sub> microcavities) [20], [21], effectively circumventing the lattice mismatch constraints inherent to conventional integration processes. While current studies have confirmed the exceptional SHG performance of NbOX2 in free-space optical configurations [22]-[24], its integration with on-chip optical systems remains unexplored - advancing this direction could propel multifunctional applications of NbOX2 in chip-scale nonlinear optical devices.

In this work, SHG was realized in a fabricated NbOI $_2$ -integrated Si $_3$ N $_4$  microdisk resonator under the pump of a sub-milliwatt continuous-wave (CW) laser and the conversion efficiency was estimated to be 0.024 %/W. The technological breakthrough stems from the following synergistic

<sup>\*</sup>Corresponding authors: Zhihong Zhu and Ken Liu, College of Advanced Interdisciplinary Studies & Hunan Provincial Key Laboratory of Novel Nanooptoelectronic Information Materials and Devices and Nanhu Laser Laboratory, National University of Defense Technology, Changsha 410073, China, E-mail: zzhwcx@163.com (Z. Zhu),

design strategies: (1) Precise transfer of few-layer NbOI<sub>2</sub> onto the edge of the Si<sub>3</sub>N<sub>4</sub> microdisk via a site-specific vdW transfer technique, ensuring interaction between the second-order nonlinear medium and the cavity modes' evanescent field; (2) Overcoming the optical path length limitation imposed by the nanometer-scale thickness of NbOI<sub>2</sub> through the resonance of WGMs. Beyond validating the feasibility of 2D NbOX2 materials for integrated nonlinear photonics, this work establishes a universal strategy for developing sub-milliwatt threshold on-chip nonlinear light source via a co-design strategy that synergizes "material nonlinearity with cavity field enhancement."

### 2 Results and discussion

### 2.1 Optical characterization and environmental stability of 2D NbOI<sub>2</sub>

Layered NbOI<sub>2</sub> nanosheets were prepared from NbOI<sub>2</sub> single crystals using the conventional mechanical exfoliation method (Figure S1). Ultraviolet (UV) - near-infrared (NIR) absorption spectra (Figure 1a) reveals strong absorption for a 100 nm-thick NbOI<sub>2</sub> flake in the visible range (400-700 nm), while the absolute absorption at the targeted pump wavelength band (around 1,550 nm, inset of Figure 1a) and the second harmonic region (near 775 nm) is below 10 %, indicating the material's suitability for subsequent SHG studies. The bandgap values of the 100 nm- and 80 nm-thick NbOI<sub>2</sub> were calculated as 1.74 eV and 1.92 eV, respectively, based on Tauc plot analysis method (Figures 1b and S2) by plotting  $(\alpha h \nu)^{1/2}$  versus  $h \nu$  [25]. These results align with literature reports [26], [27]. Under ambient conditions (relative humidity ~60 %), 30 nm-thick NbOI<sub>2</sub> on a SiO<sub>2</sub>/Si substrate exhibited progressive edge degradation over time (Figures 1c and S3). After 5 weeks, the material fully degraded, with the characteristic Raman peaks of NbOI<sub>2</sub> (located at 104 cm<sup>-1</sup>, 208 cm<sup>-1</sup>, 271 cm<sup>-1</sup>, 608 cm<sup>-1</sup>) disappearing, leaving only the Si substrate peak at 521 cm<sup>-1</sup> (Figure 1d). Correspondingly, the second-order nonlinear optical properties in NbOI2 also disappeared. For the degradation of the NbOX<sub>2</sub> materials, the current general solution strategy is to use a thin layer of hexagonal boron nitride (hBN) for packaging [22], which can significantly improve the stability and power threshold of the materials in air.

The NbOI<sub>2</sub> crystal belongs to the monoclinic *C*2 space group (Figure 1e) with a monolayer thickness of 0.73 nm. Along the a-axis, the layers are bonded via vdW forces, with each NbOI<sub>2</sub> layer composed of NbOI<sub>2</sub> octahedra; along the caxis, NbI<sub>4</sub> chains are interconnected through I atoms, where first-order Peierls distortion induces alternating Nb-Nb

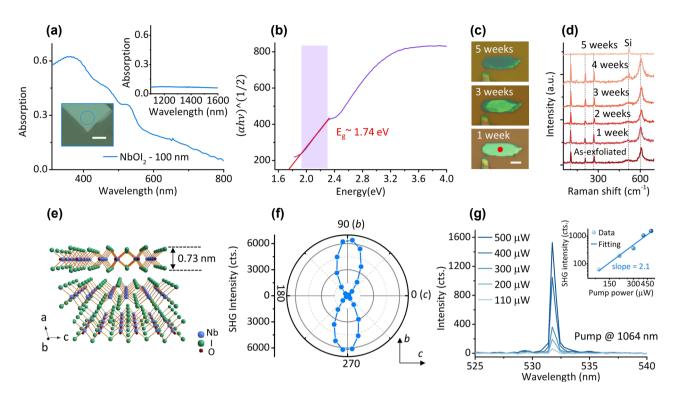
bond lengths (3.17 Å  $\leftrightarrow$  4.35 Å, Figure S4a); along the baxis, the structure is connected via O atoms, and secondorder Peierls distortion causes Nb ions to deviate from the centers of the octahedra (displacement: 0.14 Å, Figure S4b), resulting in strong spontaneous polarization along the baxis. Polarization-resolved SHG measurement revealed a characteristic C2 symmetry pattern – a two-lobed pattern (Figure 1f), with maximum intensity observed when the pump polarization aligns with the b-axis ( $\theta = 90^{\circ}$ ), consistent with theoretical predictions. Under the pump of a 1,064 nm ps pulsed laser, a prominent SHG signal was detected at 532 nm (Figure 1g). The quadratic dependence of SHG intensity on pump power in a log scale (fitted slope: 2.1, inset in Figure 1g) confirms its intrinsic second-order nonlinear optical process.

#### 2.2 Integration of 2D NbOI<sub>2</sub> and Si<sub>3</sub>N<sub>4</sub> microdisk

As shown in Figure 2a, second-order nonlinearity can be endowed to the Si<sub>3</sub>N<sub>4</sub> microdisk by precisely transferring a NbOI₂ flake to the edge of the suspended Si₃N₄ microdisk through the vdW integration technique. When the pump light ( $\omega_p$ , around 1,550 nm) in the NIR band is coupled with the NbOI2-integrated microdisk cavity through a tapered fiber, resonance enhancement occurs at specific wavelengths. The evanescent field of the WGMs interacts with the NbOI2 flake, driving the SHG process. The generated second-harmonic signal ( $\omega_{SHG} = 2\omega_p$ ) is then coupled out from the NbOI<sub>2</sub>-integrated microcavity via the tapered fiber, achieving the extraction of the nonlinear signal.

Figure 2b shows a Si<sub>3</sub>N<sub>4</sub> microdisk used in the experiment. The suspended Si<sub>3</sub>N<sub>4</sub> microdisk was prepared using a hybrid etching process (Figure S5): a 300 nm-thick Si<sub>3</sub>N<sub>4</sub> film was deposited on a Si/SiO<sub>2</sub> (3 µm) substrate by plasmaenhanced chemical vapor deposition (PECVD); UV lithography was employed to pattern the photoresist layer, followed by reactive ion etching (RIE) to transfer the microdisk pattern to the Si<sub>3</sub>N<sub>4</sub> layer; the release of Si<sub>3</sub>N<sub>4</sub> microdisks was achieved using 80 °C KOH solution (1/3 mol/L) to etch the  $SiO_2$  layer (etching rate ~100 nm/h, Figure S6). By precisely controlling the etching duration (29-30 h), a 0-100 nmthick SiO<sub>2</sub> sacrificial layer was retained (Figure S7), ensuring structural stability of the microdisk during subsequent transfer of 2D NbOI<sub>2</sub>.

The vdW integration of 2D NbOI<sub>2</sub> and Si<sub>3</sub>N<sub>4</sub> microdisk was achieved using the multi-step thermal-controlled transfer method illustrated in Figure 2f: a polydimethylsiloxane (PDMS)-polyvinyl alcohol (PVA) stamp was used to pick up the NbOI<sub>2</sub> (Figures 2c and S8a) from the SiO<sub>2</sub>/Si substrate at 60 °C (for 1 min); the picked-up NbOI2 was



**Figure 1:** Optical characterization and environmental stability of 2D NbOI<sub>2</sub>. (a) UV-NIR absorption spectra of a NbOI<sub>2</sub> flake (thickness: 100 nm). The main panel shows absorption in the 300–800 nm range. Top inset: absorption in the telecommunication band (1,100–1,600 nm); bottom inset: optical microscopy (OM) image of the measured sample (blue circle marks the test region; scale bar: 5 μm). (b) Curve of the  $(\alpha h \nu)^{1/2} - h \nu$  relationship based on the Tauc plot method (purple shaded area indicates the linear fitting region), determining a bandgap of 1.74 eV for the 100 nm-thick NbOI<sub>2</sub>. (c) OM images of time-dependent degradation of a NbOI<sub>2</sub> flake (initial thickness 30 nm) under ambient conditions (scale bar: 5 μm). Red dot in the bottom panel marks the Raman test position for Figure 1d. (d) Evolution of Raman spectra for the sample in (c): initial state displays characteristic NbOI<sub>2</sub> peaks (104 cm<sup>-1</sup>, 208 cm<sup>-1</sup>, 271 cm<sup>-1</sup>, 608 cm<sup>-1</sup>); after five weeks, only the Si substrate peak at 521 cm<sup>-1</sup> remains (gray dashed lines indicate the peak positions). (e) 3D schematic of the NbOI<sub>2</sub> crystal structure: the *a*-axis corresponds to the vdW stacking direction with a monolayer thickness of 0.73 nm. (f) Polarization-resolved SHG polar plot: second-harmonic signal intensity exhibits a two-lobed distribution, with maximum response along the *b*-axis (θ = 90°). (g) Pump power-dependent SHG spectra of a NbOI<sub>2</sub> flake under the pump of a picosecond pulsed laser at 1,064 nm. Inset: log plot of the SHG intensity versus pump power, showing a linear fit slope of 2.1, confirming that it is a second-order nonlinear optical process.

then aligned to the microdisk edge using a high-precision nano-manipulation stage; followed by hot-pressing release (for 5 min) of PVA-NbOI<sub>2</sub> onto the microdisk at 145 °C (Figures 2f(iii) and S8b); finally, the PVA was dissolved in deionized water, yielding the NbOI2-integrated Si3N4 microdisk (Figure 2d and e). After transfer, the NbOI<sub>2</sub> adhered to the suspended edge of the Si<sub>3</sub>N<sub>4</sub> microdisk via vdW force. Assuming a monolayer thickness of 0.73 nm for NbOI2, the integrated NbOI2 on the microdisk measures ~29.3 nm in thickness (yellow line in Figure 2c), corresponding to approximately 40 layers. As indicated by prior research [23], within a certain thickness range, the intensity of the second-harmonic signal scales quadratically with the number of layers. The layer-independent broken centrosymmetry in NbOI2 crystals significantly reduces the screening time of 2D materials for the fabrication of the integrated structure.

Achieving SHG in bulk materials requires precise matching of the wavevectors of the pump and the secondharmonic. However, in integrated optical microcavities, the paradigm for phase matching undergoes a transformation: it shifts toward matching the effective mode refractive indices at the fundamental and second-harmonic frequencies. These two approaches are fundamentally equivalent in their underlying principles. In order to satisfy the phase matching condition [28], [29], we precisely designed the geometric parameters of the Si<sub>3</sub>N<sub>4</sub> microdisk resonator. In the simulation, the thickness of the Si<sub>3</sub>N<sub>4</sub> microdisk was set to 300 nm; the refractive index of Si<sub>3</sub>N<sub>4</sub> was set to be 2 and did not vary with wavelength. The pump wavelength is set to 1,550 nm, with the corresponding second-harmonic wavelength at 775 nm. It can be seen from Figure 2g that the transverse electric fundamental mode at the fundamental frequency ( $\omega_p$ -TE $_0$ , blue line) and the transverse

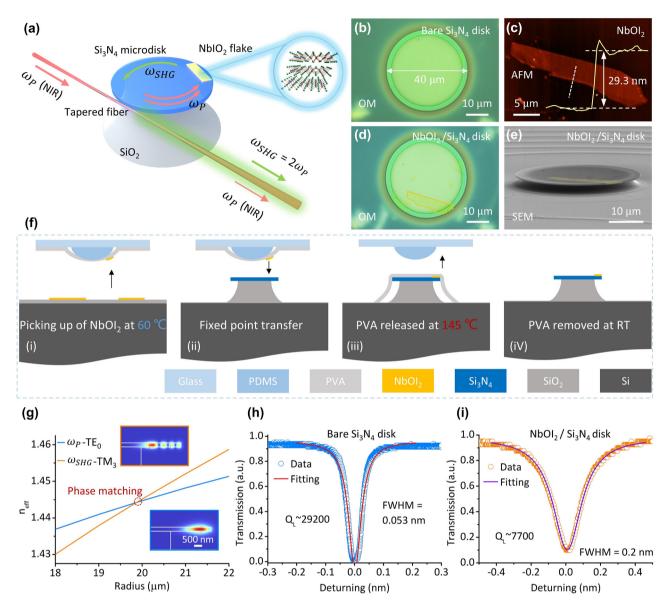


Figure 2: Integration of NbOI<sub>2</sub> and Si<sub>3</sub>N<sub>4</sub> microdisk resonator. (a) Schematic of the SHG process in the NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonator: NIR pump light  $(\omega_p)$  is coupled into the  $Si_3N_4$  microdisk via a tapered fiber. The interaction between NbOI, flake and the evanescent field of the WGMs generates the second-harmonic signal ( $\omega_{\mathsf{SHG}}$ ), which is coupled out through the same tapered fiber. (b) OM image of a bare  $\mathsf{Si}_3\mathsf{N}_4$  microdisk used in the experiment, with a diameter of 40 µm. (c) Atomic force microscopy (AFM) image of the NbOI<sub>2</sub> flake used in the experiment. Inset (yellow line): height profile along the white dashed line reveals a thickness of 29.3 nm. (d) OM image of the NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonator, with the NbOI<sub>2</sub> integration region marked by an orange dashed line. (e) False-colored scanning electron microscopy (SEM) image of the NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk, where NbOI<sub>2</sub> is treated in yellow. (f) Flowchart of the multi-step thermally-controlled transfer process: (i) picking up of NbOI<sub>2</sub> by PDMS-PVA stamp at 60 °C; (ii) transferring the NbOI<sub>2</sub> to the edge of the  $Si_3N_4$  microdisk; (iii) thermally pressurizing and releasing the PVA-NbOI<sub>2</sub> at 145 °C; (iV) removing PVA in deionized water and completing the integration. (g) Effective refractive index ( $n_{\text{eff}}$ ) curves of the TE<sub>0</sub> mode at the fundamental frequency ( $\omega_p$ -TE $_0$ , blue line) and the TM $_3$  mode at the second harmonic frequency ( $\omega_{SHG}$ -TM $_3$ , orange line) as functions of the microdisk diameter. The intersection at a radius of ~20 µm indicates the phase-matching point. Insets: electric field distributions of the TE<sub>0</sub> (fundamental frequency) and TM<sub>3</sub> (second-harmonic frequency) modes. (h, i) Transmission spectra at the resonance wavelengths of the Si $_3$ N $_4$  microdisk before (h) and after (i) the integration of NbOI<sub>2</sub>, with the loaded Q factor ( $Q_1$ ) decreases from 29,200 to 7,700. Circles: experimental data; solid lines: Lorentzian fitting curves.

magnetic third-order mode at the second harmonic frequency ( $\omega_{
m SHG}$ -TM $_3$ , orange line) have an intersection of the effective refractive index ( $n_{\rm eff}$ ) curves at microdisk radius close to 20 µm, i.e., the phase-matching point. At this point, both energy and momentum conservation are satisfied for the SHG process. The bottom and top insets in Figure 2g illustrate the electric field distributions of the TE<sub>0</sub> mode at the fundamental frequency and the TM<sub>2</sub> mode at the secondharmonic frequency, respectively, for a 20 µm-radius Si<sub>3</sub>N<sub>4</sub> microdisk. We have also analyzed the phase-matching condition of the Si<sub>2</sub>N<sub>4</sub> microdisk integrated with a 30 nm-thick NbOI<sub>2</sub> flake (Figure S9). Although the NbOI<sub>2</sub> affects the effective refractive indices of both fundamental and secondharmonic modes, phase matching can still be achieved at a microdisk diameter close to 20 µm. This approach of achieving phase matching by designing the geometry of microcavities is indeed highly versatile and robust, serving as a powerful tool for engineering a wide range of nonlinear processes (Supporting Material Note S1).

Figure 2h and i show the transmission spectra at resonance wavelengths before and after the integration of NbOI<sub>2</sub> with the Si<sub>3</sub>N<sub>4</sub> microdisk, respectively, where Lorentzian fittings of the data (circles) yield the Q values of the microdisk resonator. After the integration of NbOI<sub>2</sub>, the loaded Q value  $(Q_L)$  of the microdisk resonator decreased from 29,200 (Figure 2h) to 7,700 (Figure 2i), primarily due to increased scattering losses at the microdisk edge. Meanwhile, the decrease of the Q value also indicates that the WGMs in the Si<sub>3</sub>N<sub>4</sub> microdisk interacted effectively with the integrated NbOI<sub>2</sub>.

Since high Q factor is essential for high frequency conversion efficiency, after transferring NbOI<sub>2</sub> to the Si<sub>3</sub>N<sub>4</sub> microdisk we grew a certain thickness of Al<sub>2</sub>O<sub>3</sub> by atomic layer deposition (ALD), trying to improve the Q value of the NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk [30]. However, no obvious increase in Q value was found. Future studies should investigate strategies to mitigate the reduction in the microcavity Q-factor, including: (1) Optimizing the integration process: adopting cleaner dry transfer techniques (e.g., direct peeling using PDMS/polycarbonate (PC) films) to minimize contaminants/polymers introduced during transfer; optimizing annealing processes to reduce scattering centers as much as possible. (2) Using hBN encapsulation: covering NbOI<sub>2</sub> with a layer of hBN before integrating it with the Si<sub>3</sub>N<sub>4</sub> microdisk resonator. hBN not only enhances the stability of NbOI2 in ambient conditions but also, due to its atomically flat surface, reduces the strength of interaction between the optical field and defects/edges of NbOI<sub>2</sub>, thereby significantly lowering scattering losses. (3) Optimization of the microdisk design and post-integration cleaning procedures.

### 2.3 SHG in the NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonator

After the integration of NbOI2 and Si3N4 microdisk resonator, the evanescent field of the WGMs in the Si<sub>3</sub>N<sub>4</sub>

microdisk resonator interacts with the NbOI<sub>2</sub> flake, enabling SHG under the pump of a sub-milliwatt CW laser. As indicated in Figure 1c and d, bare NbOI2 gradually degrades in ambient conditions, with a degradation period of approximately five weeks under our laboratory conditions. For our SHG study, all SHG measurements were completed within 1-2 days after device fabrication, ensuring that the optical properties and nonlinear response remained stable during the measurements. Therefore, the reported experimental results were not affected by material degradation. Besides, the challenge of long-term operational stability for the NbOI2-integrated device can be effectively addressed through established 2D material encapsulation techniques, such as hBN and polymethyl methacrylate (PMMA) [22].

Figure 3a illustrates the experimental setup for the measurement, in which the pump light is coupled with the NbOI2-integrated Si3N4 microcavity through a tapered fiber (waist diameter:  $\sim$ 1  $\mu$ m), and the polarization state is adjusted by a polarization controller (PC); when the emission wavelength of the tunable laser matches the resonant wavelength of the NbOI2-integrated Si3N4 microcavity, the optical field inside the cavity is resonantly enhanced, driving SHG in the NbOI<sub>2</sub> flake; the generated second harmonic signal is coupled out through the tapered fiber and detected by a spectrograph for the VIS light; a photodetector (PD, Daheng Optics, DH-GDT-D002N) and an oscilloscope (OWON Technology Inc., VDS1022) are used to record the transmission spectrum for the pump light; the output power for the pump is measured using an optical power meter (Daheng Optics, GCI-080201).

The b-axis of the NbOI<sub>2</sub> crystal (marked with a black arrow) is tangential to the edge of the microdisk, i.e., aligned along the radial direction of the microdisk. In practice, during our experimental testing, by adjusting the PC to ensure excitation of the fundamental TE mode in the microdisk resonator. The pump electric field is aligned parallel to the b-axis of the NbOI<sub>2</sub> crystal (marked with a red arrow). According to Figure 1f, this configuration yields the maximum second-harmonic signal.

Laser frequency scanning was performed in the 1,540-1,560 nm wavelength range. The output raw data from the oscilloscope is shown in Figure S10. The laser (Aunion Tech Co., Ltd) scanned at a speed of 1 nm/s. To obtain the second harmonic signals, we tuned the pump wavelength to the resonant wavelength of the microcavity (1,540.6 nm and 1,551.9 nm). Figure 3b shows the transmission spectrum at the resonant wavelength of 1,540.6 nm, with a Lorentzian-fitted linewidth of  $\sim$ 0.2 nm; Figure 3c displays the SHG spectra under varying pump powers  $(175 \mu W - 350 \mu W)$ , the specific values for the pump power

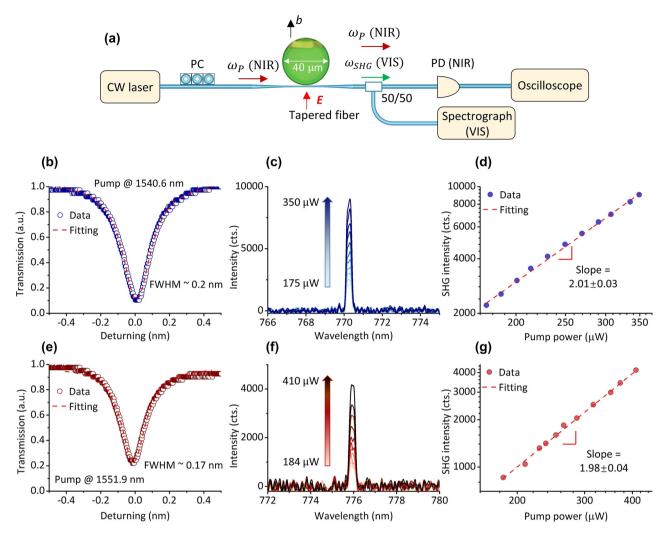


Figure 3: SHG in the NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonator. (a) Schematic of the experimental setup: a tapered fiber is used for the input of the pump  $(\omega_p)$  and the extraction of the second harmonic signal  $(\omega_{SHG})$  from the microdisk; the polarization state of the input light is controlled by a polarization controller (PC), the red arrow indicates the direction of the pump electric field; a photodetector (PD) and oscilloscope record the transmission spectrum of the pump in the NIR band, while a visible (VIS) spectrograph collects the SHG signal. Inset: OM image of the  $NbOI_2$ -integrated  $Si_2N_A$  microdisk resonator, the black arrow indicates the direction of the *b*-axis of the integrated  $NbOI_2$  crystal. (b) Transmission spectrum at the resonant wavelength of 1,540.6 nm, with a Lorentzian-fitted linewidth of 0.2 nm. (c) SHG spectra under varying pump powers. The second harmonic wavelength is locked at 770.3 nm, corresponding to half of the pump wavelength. (d) The relationship between the SHG intensity and the pump power in a log scale. The fitting slope of 2.01 + 0.03 verifies the intrinsic second-order nonlinear process. ( $e-\alpha$ ) SHG characterization at a resonant wavelength of 1,551.9 nm. (e) Transmission spectrum with a Lorentzian-fitted linewidth of 0.17 nm. (f) SHG spectra under different pump powers, with the SHG wavelength located at 776 nm, corresponding to half of the pump wavelength. (g) The relationship between the SHG intensity and the pump power in a log scale. The fitting slope of  $1.98 \pm 0.04$  verifies the intrinsic second-order nonlinear process.

are as follows: 175 μW, 187 μW, 201 μW, 214 μW, 231 μW,  $250 \mu W$ ,  $270 \mu W$ ,  $291 \mu W$ ,  $308 \mu W$ ,  $336 \mu W$  and  $350 \mu W$ ), where the second-harmonic wavelength of 770.3 nm corresponds to half of the pump wavelength; Additionally, the quadratic dependence of the SHG intensity on pump power in a log scale (fitted slope:  $2.01 \pm 0.03$ ) further confirms the second-order nonlinear optical nature of the SHG process (Figure 3d). We also investigated the transmission spectrum at a resonant wavelength of 1,551.9 nm

(Figure 3e), the corresponding SHG spectra under varying pump powers (Figure 3f) and the dependence of the SHG intensity on pump power in a log scale (Figure 3g). In Figure 3f, the pump power gradually increases from 184  $\mu$ W to 410  $\mu$ W, in the following order: 184  $\mu$ W, 210  $\mu$ W, 229  $\mu$ W,  $238 \mu W$ ,  $253 \mu W$ ,  $265 \mu W$ ,  $287 \mu W$ ,  $317 \mu W$ ,  $352 \mu W$ ,  $373 \mu W$ and 410 µW. Due to the resonance enhancement effect of the cavity, SHG process is achieved under the pump of a low-power CW laser.

Under experimental conditions identical to those used for Figure 3, we characterized a bare Si<sub>2</sub>N<sub>4</sub> microdisk resonator. We sequentially tuned the pump wavelength to each resonant wavelength and performed SHG measurements. Although theoretically, cavity enhancement could amplify interface effects and contribute to measurable SHG [12], under our experimental conditions – specifically, a Q-factor on the order of 10<sup>4</sup> and CW laser pumping at sub-milliwatt power levels - we did not detect any second-harmonic signal in the bare Si<sub>3</sub>N<sub>4</sub> microdisk resonator. In contrast, the SHG process was only achieved after integrating the NbOI<sub>2</sub> flake onto the Si<sub>3</sub>N<sub>4</sub> microdisk resonator, which directly confirms that the collected SHG signal originates from NbOI<sub>2</sub>.

Next, we estimate the conversion efficiency of SHG: under the condition where the pump wavelength is 1,540.6 nm, the output power of the tapered fiber is 350  $\mu$ W, which is used as the pump power  $P_{\text{pump}}$ ; For the generated second harmonic signal, considering the absorption loss of single-mode fiber in the communication band for visible light, the diffraction efficiency of the spectrograph, and the quantum efficiency, the estimated power for the generated second harmonic signal  $P_{\rm SHG}$  is 29 pW. Therefore, the estimated conversion efficiency of SHG is:  $\eta_{\rm integrated} = \frac{P_{\rm SHG}}{P_{\rm pump}^2} \times$  $100 \% \approx 0.024 \%/W$ .

Although the conversion efficiency is relatively low, this process is indeed enabled by the cavity field enhancement. The high-Q microresonator (loaded  $Q \sim 7,700$ ) builds up the intracavity pump power significantly, making the SHG process feasible. This is a key advantage of our cavity-integrated approach, allowing us to study SHG process in NbOI<sub>2</sub> without using pulsed lasers.

The spectral data in Figure 1g were obtained using a 100× objective with a collection efficiency of approximately 12 % for the second-harmonic signal at 532 nm. Based on this, the SHG conversion efficiency of pure NbOI<sub>2</sub> under free-space configuration at 400 μW pump power was calculated to be approximately  $\eta_{\text{space},1064}$  =  $2.04 \times 10^{-6}$  %/W. After wavelength scaling to 1,550 nm, we obtain an cavity-enhancement factor of approximately  $\eta_{\rm integrated}/\eta_{\rm space,1550} \approx 25{,}000.$ 

Table 1 summarizes the progress made in integrating 2D materials with high intrinsic  $\chi^{(2)}$  with WGM microcavities for SHG research. In all cases, the pump sources are continuous-wave lasers operating in the communication band;  $\chi_{\rm intrinsic}$  (2) represents the second-order nonlinear polarizability of the 2D material; and the microcavity platforms used include Si<sub>3</sub>N<sub>4</sub> microring resonators (MRR), SiO<sub>2</sub> microsphere resonators (MSR), and Si<sub>3</sub>N<sub>4</sub> microdisk resonators (MDR) used in this work. "Q factor (before)" and "O factor (after)" presents the O factors for the microcavities before and after the integration with 2D materials. The Si<sub>2</sub>N<sub>4</sub> microdisk resonator platform used in our work offers distinct advantages over other resonator architectures: Compared to microring resonators, its fiber coupling configuration allows more flexible and robust critical coupling control under fabrication variations; relative to microspheres, our fully planar platform provides superior integration compatibility with 2D materials and complementary metaloxide-semiconductor (CMOS) compatibility while maintaining high Q-factors. Furthermore, unlike Fabry-Perot cavities that offer high Q but lack mode discreteness [31], or nanoplasmonic systems that provide strong spatial confinement but suffer from broad resonances and high loss [32]. the microdisk resonator platform employed in this work simultaneously combines high-Q temporal confinement, a discrete mode spectrum for phase matching, and a flexible and robust experimental testing configuration, making it an ideal and highly promising platform for nonlinear enhancement studies of 2D materials. Although a direct comparison of effective nonlinear coefficients ( $\chi_{\rm eff}^{(2)}$ ) across different 2D materials-integrated microcavity platforms is infeasible due to the lack of reported values, the excellent secondorder nonlinear optical properties of NbOX2 confirmed by free-space optical studies [22], [23], combined with the high SHG enhancement factor achieved in our microcavity platform, strongly demonstrate that the NbOX<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonator proposed in this work possesses highly competitive effective second-order nonlinearity and significant potential for further optimization.

Table 1: Summary of SHG research by integrating 2D materials with WGM microcavity platforms.

Materials	$\chi_{ m intrinsic}^{(2)}$ (pm/V)	Microcavities	Q factor (before)	Q factor (after)	η (%/W)	Ref.
GaSe	2,400	Si <sub>3</sub> N <sub>4</sub> MRR	2,000	1,800	849	[33]
SnP <sub>2</sub> Se <sub>6</sub>	1,300	Si <sub>3</sub> N <sub>4</sub> MRR	_	1,000	43.2	[34]
NbOBr <sub>2</sub>	91.6	Si <sub>3</sub> N <sub>4</sub> MRR	14,000	24,000	158	[35]
WSe <sub>2</sub>	~10 <sup>3</sup>	SiO <sub>2</sub> MSR	$1 \times 10^{8}$	$5 \times 10^{6}$	$6.6 \times 10^{-4}$	[36]
Surface molecule	210	SiO <sub>2</sub> MSR	$6 \times 10^{7}$	$3 \times 10^{7}$	6.7	[37]
WS <sub>2</sub>	~10 <sup>3</sup>	SiO <sub>2</sub> MSR	_	~10 <sup>6</sup>	$1.08 \times 10^{-5}$	[38]
NbOI <sub>2</sub>	~90	Si₃N₄ MDR	29,200	7,700	0.024	This worl

We can analyze the reasons for the low SHG conversion efficiency of the device in this work from a qualitative perspective: (1) Relatively low Q value of microcavity (for the bare  $Si_3N_4$  microdisk,  $Q \sim 10^4$ ), which can be improved by subsequent ultra-low-loss microcavity design; (2) In the phase-matching analysis in Figure 2g, the second-harmonic mode corresponding to the phase-matching point is the asymmetric TM<sub>3</sub> mode, resulting in a low mode overlap between the fundamental mode and the second-harmonic mode. This can be optimized by the design of the geometric parameters of the microcavity and gradient thickness engineering of NbOI<sub>2</sub>; (3) The absorption of the generated second harmonic signal by the NbOI<sub>2</sub> material itself (Figure 1a) is also a factor contributing to the relatively low calculated  $\eta_{\text{integrated}}$ ; (4) For the pump of light and the extraction of the second harmonic signal, we used the same tapered fiber, which has absorption for the second harmonic signal, also leading to a lower calculated  $\eta_{\mathrm{integrated}}.$  The experimental setup can be optimized in subsequent work by employing two separate tapered fibers for the coupling of the pump and the extraction of the second-harmonic signal, respectively [12]. Optimizing these limiting factors will further improve the SHG conversion efficiency.

We have also theoretically analyzed the maximum achievable SHG conversion efficiency (Supporting Material Note S2), projecting a potential improvement of two orders of magnitude over the current highest reported SHG conversion efficiency listed in Table 1. Beyond cavityinduced SHG enhancement, the intrinsic  $\chi^{(2)}$  of NbOI<sub>2</sub> could be further enhanced via band-edge resonance when the laser energy approaches half the bandgap of the NbOI<sub>2</sub>: for NbOI<sub>2</sub> with a bandgap of  $\sim$ 1.74 eV, this implies that tuning the microdisk resonance to approximately 1,427 nm (corresponding to a photon energy of  $\sim$ 0.87 eV) could theoretically maximize its intrinsic  $\chi^{(2)}$  response. This would enable dualresonance enhancement, combining pump photon energy resonance with optical microcavity resonance, thereby pushing the overall second-order nonlinear response of the system to a new level, further demonstrating the significant application potential of our proposed NbOI2-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonator platform in the field of integrated second-order nonlinear optics.

#### 3 Conclusions

In this study, NbOI<sub>2</sub>-integrated Si<sub>3</sub>N<sub>4</sub> microdisk resonators were successfully fabricated through a vdW integration strategy, and SHG was realized under the pump of a lowpower (~175 µW) CW laser and the SHG conversion efficiency was calculated to be 0.024 %/W. By leveraging the exceptional second-order nonlinear optical properties of 2D NbOI<sub>2</sub> with the strong optical confinement capability of the Si<sub>3</sub>N<sub>4</sub> microdisk, we overcame the optical path length limitation imposed by the nanoscale thickness of 2D materials in traditional free-space optical systems. Furthermore, the site-specific vdW transfer technique, combined with NbOI $_2$ 's intrinsic lack of inversion symmetry (independent of layer parity), greatly shortens the time required for device preparation. In future experiments, polarization-resolved SHG measurements should be conducted to fully characterize the anisotropic nonlinear response within the cavity; meanwhile, electro-optic tunable frequency conversion devices can be developed by leveraging the ferroelectric properties of NbOX2 materials. This work not only establishes a universal strategy for the co-design of 2D materials and microcavities but also opens new avenues for integrated nonlinear photonics in quantum technologies and metrological systems.

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individuals included in this study.

**Author contributions:** ZZ and KL conceived and supervised the project. NL performed the numerical simulations. NL, QL, and YL fabricated the devices. NL and KL built the setup and characterized the devices. NL, ZZ, and KL analyzed the data. NL wrote the manuscript with inputs from all authors. All authors have accepted responsibility for the entire content of this manuscript and consented to its submission to the journal, reviewed all the results and approved the final version of the manuscript.

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