A simple growth method for Nb$_2$O$_5$ films and their optical properties

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A simple method for the synthesis of Nb$_2$O$_5$ films of thicknesses ranging from tens to several hundreds of nanometers on amorphous silicon dioxide or quartz substrates is presented. Nb$_2$O$_5$ films were formed by annealing the sputter deposited Nb films under an Ar flow and without oxygen plasma in a quartz tube within a furnace at 850 °C. The structural, compositional, optical, and vibrational properties were characterized by grazing incidence X-ray diffraction, transmission electron microscopy, X-ray photoelectron spectroscopy, ultraviolet visible spectroscopy, and Raman scattering. Each of the Nb$_2$O$_5$ films is polycrystalline with an orthorhombic crystal structure. We observed vibrational modes including longitudinal optical, transverse optical, and triply degenerate modes, and measured the indirect optical band gap to be ~3.65 eV. The transmittance spectrum of the ~20 nm thick Nb$_2$O$_5$ film shows over 90% transmittance below the band gap energy in the visible wavelength range and decreases to less than 20% in the ultraviolet regime. The optical properties of the films in the UV-vis range show potential applications as UV detectors.

1. Introduction

Niobium oxides have attracted attention among researchers for their rich charge states, phases, morphologies, and associated properties. The number of charge states of niobium ranges from +2 to +5 for NbO, NbO$_2$ and Nb$_2$O$_5$, respectively. These suboxides can be conductors, n-type semiconductors, or insulators. Nb$_2$O$_5$ has the highest heat of formation (−1899.54 ± 8.37 kJ mol$^{-1}$) compared with NbO$_2$ (−792.87 ± 6.28 kJ mol$^{-1}$) and NbO (−407.94 ± 12.55 kJ mol$^{-1}$) and is the most stable once it is formed. Nb$_2$O$_5$ exists in various phases including the pseudohexagonal (TT-phase), orthorhombic (T-phase), and monoclinic (H-phase). Various physical or chemical processing methods have been developed to prepare Nb$_2$O$_5$ thin films and nanostructures. Examples of Nb$_2$O$_5$ thin film preparations are reactive sputtering of a Nb target in an Ar–O$_2$ plasma on SiO$_2$/Si substrates or in an atmosphere composed of oxygen and argon, pulsed laser deposition of Nb on Si followed by exposure to atmosphere, reactive RF magnetron sputtering of a Nb$_2$O$_5$ target on glass substrates, atomic layer deposition, and sol–gel dip coating. The extended sol–gel method was also used to form nanocrystalline Nb$_2$O$_5$ films. Other nanostructures synthesized by chemical processes in the form of nanowires, nanorods, nanotubes, nanobelts, nanopores, nanopowders, and nanohollow fibers have been reported. A wide range of potential applications has been demonstrated: catalysis, gas sensors, electrochromic windows, photodetectors, electron field emitters, and solid oxide fuel cells.

In this work we report a simple method to fabricate ultrathin Nb$_2$O$_5$ films by post annealing the magnetron sputtered Nb films. We study their properties including structure, chemical composition, optical transmittance and reflection, optical bandgap, surface morphology, and vibrational modes. The characterization techniques used are grazing incidence X-ray diffraction (GIXRD), transmission electron microscopy (TEM), X-ray photoelectron spectroscopy (XPS), ultraviolet visible (UV-VIS) spectroscopy, atomic force microscopy (AFM), X-ray specular reflectivity (XRR), and Raman scattering.

The advantages of our ultrathin Nb$_2$O$_5$ film growth method are (1) the oxide film thickness can be controlled by the initial thickness of the Nb film and the oxide film is continuous, (2) there is no need to use the oxygen plasma, pulsed laser, or chemical precursors, but it only needs an Ar flow in the quartz tube in a furnace, and (3) the optical properties including transmittance, reflection and optical bandgap energy of 3.65 eV are suitable for UV detection.

2. Experimental

Ultrathin Nb films of thicknesses 5, 10, 100, and 300 nm were deposited by DC magnetron sputter deposition on 300 nm thick...
SiO₂ covered Si (001) substrates (of resistivity 0.001–0.005 ohm cm) and quartz slides (Structure Probe, Inc.). The base pressure of the sputter deposition chamber was ~7.5 × 10⁻⁷ Torr. The Ar gas flow rate was controlled at 2.01 standard cubic centimeters per minute (sccm) and the working pressure of Ar was set to 2.3 mTorr. The power was set to 50 W. The Nb target (99.95% purity) was a 3 inch round disk and the distance from the target surface to the center of the substrate was ~15 cm. The deposition rate was ~3.5 nm min⁻¹. This deposition rate was calibrated from the cross sectional image obtained from the scanning electron microscopy (SEM).

The Nb ultrathin films on SiO₂/Si and quartz substrates were then placed in a quartz boat (100 × 45 × 20 mm) positioned at the center of a 30 inch long quartz tube with outside and inside diameters of 59 and 55 mm, respectively. This quartz tube was placed inside a one zone furnace and then purged with an ultrahigh purity Ar gas (99.999%) at a flow rate of 950 sccm. After 20 min of Ar purging, the Ar flow rate was reduced to 200 sccm while the quartz boat was gradually heated up from room temperature to 850 °C in the furnace over about 80 min. The temperature was measured using a type K thermocouple with an accuracy of ±5 °C. The boats were kept at 850 °C for 30 min before the furnace filament current was reduced to zero over 5–6 min. The quartz tube then cooled down to room temperature in about 120 min. Ar gas purging was continuous during the entire heating up and cooling down processes. The temperature ramping was managed using a Eurotherm 2116 PID programmable temperature controller attached to the furnace. The furnace and Ar gas handling lines were placed inside a venting hood. Fig. 1(a) shows a schematic of the sputter deposited Nb film on SiO₂/Si. For optical measurements the Nb films were sputter deposited on quartz substrates. Fig. 1(b) shows a TEM cross-sectional view of a Nb₂O₅ film synthesized from a 300 nm thick Nb film grown on a SiO₂/Si (001) substrate that was annealed at 850 °C for 30 min inside the quartz tube furnace. The uniform contrast areas near the bottom left are Si and SiO₂. The area to the right of SiO₂ is the Nb₂O₅ film followed by a thin layer of epoxy (M-bond) and a Pt layer. The 300 ± 5 nm thickness of the SiO₂ layer is in agreement with the vendor’s thickness specification. The thickness of the Nb₂O₅ film is 800 ± 20 nm. This oxide layer is a result of the oxidation of the 300 nm thick Nb film. Many grains with sizes on the order of hundreds nm to microns are seen.

Grazing incidence X-ray diffraction (GIXRD) theta–2theta (θ–2θ) scans were collected using a PANalytical X’Pert Pro diffractometer to characterize the atomic structure and average grain size of the Nb₂O₅ films. A parallel X-ray beam generated from a parabolic mirror with a wavelength of 1.5405 Å was incident on the sample at a fixed 1.5° grazing angle (ω). A point detector with a 0.013° scanning step size and a 0.25 s counting time at each step, was used to collect X-ray diffraction scans. During the GIXRD scan, the ω was fixed at 1.5° while the 2θ was varied. The detector scanned over 2θ in the scattering plane while the source and sample stage were fixed. The detector was scanned in the scattering plane that contains the incident X-ray and the surface normal.

For the TEM sample, a thin layer of epoxy (M-bond) was applied to the surface of the Nb₂O₅ film. A sacrificial Pt layer was deposited on the surface of the epoxy layer for focused ion beam (FIB)/scanning electron microscopy (SEM) sample preparation. The sample was then thinned by FIB milling using energetic gallium ions. The thinned sample was mounted on a liquid nitrogen cooled sample stage to minimize the sample's mechanical drifting. A 200 keV energy electron beam was used to obtain the TEM (JEOL 2100 F) bright field images, high resolution images, and electron diffraction patterns.

X-ray photoelectron spectroscopy (XPS) was used to determine the near surface composition. The nearly monochromatic X-ray Mg K-alpha source (PHI model 04-151) had a primary energy of 1253.6 eV. The X-ray source was operated at 270 W with a 10 kV high-voltage bias and an emission current of 27 mA. A double pass cylindrical mirror energy analyzer (PHI model 15-255G) in a high vacuum chamber was used to detect the emitted electrons from the sample surface. A high-resolution spectrum was averaged from 40 repeated scans at a passing energy of 50 eV. The energy resolution of the analyzer at this passing energy is about 0.7 eV. The energy step size (0.08 eV) used to scan Nb and O peaks is smaller than 0.7 eV in order to provide a good relative resolution, defined as \( R = \Delta E/E_o \), where \( \Delta E \) is the passing energy and \( E_o \) is the kinetic energy of a peak position.⁷⁷

UV-visible reflectance and transmission spectra were collected to determine the optical band gap of the ultrathin Nb₂O₅ films on quartz. The spectra were obtained from a Perkin-Elmer Lambda 950 UV/VIS Spectrometer. The transmission spectra were obtained with the slides held normal to the incident light beam in air and were corrected for the quartz background. Specular reflectance spectra were obtained at a near-normal incidence angle of 6°.

The AFM surface morphology images were collected using a PSI XE100 in non-contact mode. The HiRES-C14/CR-AU AFM tip used has a tip radius of ~1 nm, a force constant of 5 N m⁻¹ and a resonant frequency of 160 kHz.

X-ray reflectivity (XRR) scan obtained for the Nb₂O₅ films grown on quartz were used to determine thickness, surface roughness and density. XRR scans were taken with a Bruker D8 Discover X-ray Diffractometer using a 2-circle \( \theta/2\theta \) goniometer, a centric Eulerian cradle, and a sealed copper tube X-ray source.
(Cu Kα, λ = 1.54 Å) operated at 40 kV. A 40 mm Gobel collimating mirror and a four-position rotary absorber were used for the incident beam with attenuations of approximately 1 : 1, 1 : 10, 1 : 100 and 1 : 1000. Slits of 1.0 mm and 0.2 mm were used before and after the rotary absorber, respectively. A knife edge was employed to optimize the measurements. Data were collected at grazing incidence angles ranging from 0 to 5° at 0.01° increments and repeated scans were collected and averaged until the desired signal to noise level was reached.

Raman spectra that provide vibrational mode information were collected using a Witec Alpha 300 confocal Raman microscope. The laser wavelength and power used for the measurements of the Raman spectra were 532.1 nm and 17 mW, respectively. A 100× objective lens produced a spot size of ~721 nm (=1.22 × wavelength/numerical aperture of 0.9 for the 100× objective lens). The spectral resolution and step size used in Raman scattering data collection were each 0.02 cm⁻¹.

3. Results

3.1. Structural property

3.1.1. Grazing incidence X-ray diffraction. Fig. 2 shows GIXRD θ/2θ scans of the 300 nm thick SiO₂/Si (001) substrate, 5 nm thick Nb on SiO₂/Si (001), and the synthesized Nb₂O₅ films from 5, 10, 100, and 300 nm thick Nb on SiO₂/Si (001). The intensity from SiO₂/Si (001) is mostly flat over the 2θ range with the exception of a low intensity bump near 22.8° that is fromreported short range (~1 nm) order in SiO₂ measured by X-ray diffraction. For the 5 nm Nb, film there is a peak at 38.6° from the (110) planes of Nb. See Table 1 for Bragg peak positions. The Nb film has a bcc structure and the (110) is the lowest surface energy plane. According to the structure zone model, the surface atoms’ mobility is high in zone II and atoms form the lowest surface energy plane. (Note that our GIXRD-measured plane corresponding to the (110) peak is not exactly parallel to the surface. ~17.8° from the substrate normal = θ, the grazing incident angle of 1.5°). For the film synthesized from the 5 nm thick Nb film, the (110) peak disappears but two new peaks appear at 2θ = 22.6° and 46.1°. These are identified as (001) and (002) reflections of Nb₂O₅ using lattice constant c = 3.930 Å. Similar to the film synthesized from the 5 nm thick Nb film, a GIXRD scan from the film synthesized from the 10 nm thick Nb film has (001) and (002) peaks with slightly higher intensities. For the films synthesized from the 100 nm and 300 nm Nb films, two additional peaks appear at 28.4° and 36.6°. When the GIXRD 2θ scan range is greater than 50°, two peaks at 50.9° and 55.1° are also observed. These four additional peak positions are in agreement with the calculated peak positions of the (180), (181), (380) and (182) planes using the lattice constants a = 6.175 Å, b = 29.175 Å and c = 3.930 Å, and the angles α = β = γ = 90°. The above results show that the films synthesized from all thickness of Nb films have a Nb₂O₅ orthorhombic phase with a Pham space group. The [001] direction has a lattice constant c. For the films synthesized from the 100 nm and 300 nm thick Nb films there are more orientations than just [001]. These additional Nb₂O₅ orientations are (180), (181), (380), and (182). For the film synthesized from the 300 nm thick Nb film there is a small (110) peak at 38.58°.

From the full-width-at-half-maximum (FWHM) of a peak intensity profile one can estimate the average ordered crystallite size using the Scherrer formula. That is, the average crystallite size = kλ/(β cos θ), where λ is a shape factor of about 0.9, λ is the wavelength of the X-ray, β is the FWHM in radians, and θ is the Bragg angle in degrees. Note that the size obtained from the Scherrer formula is smaller than or equal to the actual grain size. This is because there are lattice imperfections such as stacking faults and dislocations that can contribute to the peak broadening in addition to the finite crystallite size. Also, there is a small broadening due to instrumental effects. From the FWHM of the (110) peak, the average crystallite size of the thinnest pure Nb film is estimated to be 5.5 nm. This is consistent with the intended 5 nm thick sputtered Nb film. The average crystallite sizes of the Nb₂O₅ films estimated from the (001) peaks of the synthesized films from 5, 10, 100, and 300 nm thick Nb films are about 12.5 ± 0.2, 18.6 ± 0.2, 30.8 ± 0.4, and 42.6 ± 0.4 nm, respectively. The crystallite size increases as the starting Nb film thickness increases. The increased crystallite sizes of the Nb₂O₅ films relative to the original smooth Nb films (see Fig. 6(a) and (b) later) result from the synthesized Nb₂O₅ films having a 3D island structure with a height distribution (see Fig. 6(c) and (d) later), rather than being smooth and continuous films. The crystallite size extracted from the small (110) peak of Nb from the Nb₂O₅ film synthesized from the 300 nm thick Nb film is about 14.8 ± 0.2 nm.

3.1.2. High resolution TEM image and diffraction pattern. Fig. 3(a) and (b) show a high resolution TEM image of the 800 ± 20 nm thick Nb₂O₅ film and a selected area TEM diffraction pattern (DP) of the same film, respectively. The zone axis is along the [100], or a direction. Line scans in Fig. 3(a) show the spacing between adjacent atoms along the [001], or c direction.
measured the reciprocal length \( b \) measured from the spots in the DP shown in Fig. 3(b). We next.

To determine the unit length along the \([010]\), or \(b\) direction seen in Fig. 3(b). Note the lower resolution bright field TEM image shown in Fig. 1(b) does not reveal these stacking faults. The grain size seen in TEM is much larger than the crystallite size determined from the FWHM of XRD peak.

The lengths of the reciprocal unit vectors \( c^* \) and \( b^* \) are measured from the spots in the DP shown in Fig. 3(b). We measured the reciprocal length \( b^* \) between the 2\(^{nd}\) order spots in the positive and negative \( b^* \) directions, or 4\(b^*\), and then divided by 4 (the 000 spot is too strong and the TEM beam stopper blocks the spot). The measured values are \( c^* = 2.64 \pm 0.01 \text{ nm}^{-1} \) and \( b^* = 0.357 \pm 0.005 \text{ nm}^{-1} \). The inverses of \( c^* \) and \( b^* \) correspond to real space spacings \( c \approx 0.378 \text{ nm} \) and \( b \approx 2.80 \text{ nm} \), respectively. These are in reasonable agreement with the measured real space unit lengths \( c \) and \( b \) shown in Fig. 3(a) as well lattice constants obtained from X-ray diffraction. Miller indices of the stronger diffraction spots are labeled. All reciprocal lattice vector lengths \( G(hkl) \) match well to the real space spacings \( d_{hkl} \) through \( d_{hkl} = 1/ G(hkl) \). The fast Fourier transform of the real space image shown in Fig. 3(a) using ImageJ software gives a DP in agreement with the observed DP. The real space high resolution image in Fig. 3(a) shows that many stacking faults exist along the \( b \) direction. The stacking faults give raise to the elongated diffraction spots along the \( b^* \) direction seen in Fig. 3(b). Note the lower resolution bright field TEM image shown in Fig. 1(b) does not reveal these stacking faults.

Other high resolution TEM images of the Nb\(_2\)O\(_5\) film and the corresponding selected area TEM DPs were also measured from a grain with the zone axis along the [001], or \(c\) direction so that the [100], or \(a\) direction is parallel to the high resolution image. The measured unit vector length, \( a \), from the real space high resolution TEM image and TEM DP are consistent with the value of 0.6168 nm determined from the (180) peak in Section 3.1.1.

Table 1  Bragg peak and average ordered crystallite size from XRD and film’s roughness parameters including surface width, correlation length and roughness exponent from AFM

<table>
<thead>
<tr>
<th>Sample (nm)</th>
<th>GIXRD Orientation (XRD Bragg peak)</th>
<th>Ave. ordered crystalline size (nm)</th>
<th>AFM Surface width ( \omega ) (nm)</th>
<th>Correlation length ( \xi ) (nm)</th>
<th>Roughness exponent ( \alpha )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nb(_2)O(_5) synthesized films from 300 nm Nb</td>
<td>5 nm Nb 110 (38.6°)</td>
<td>5.5 ± 0.1</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Nb(_2)O(_5) synthesized films from 300 nm Nb</td>
<td>5 nm Nb 001 (22.6°), 002 (46.1°)</td>
<td>12.5 ± 0.2</td>
<td>3.6 ± 0.2</td>
<td>84 ± 2</td>
<td>0.78</td>
</tr>
<tr>
<td>Nb(_2)O(_5) synthesized films from 300 nm Nb</td>
<td>10 nm Nb 001 (22.6°), 002 (46.1°)</td>
<td>18.6 ± 0.4</td>
<td>3.8 ± 0.2</td>
<td>91 ± 2</td>
<td>0.72</td>
</tr>
<tr>
<td>Nb(_2)O(_5) synthesized films from 300 nm Nb</td>
<td>300 nm Nb 001 (22.6°), 002 (46.1°), 180 (28.4°), 181 (36.6°), 380 (50.9°), 182 (55.1°)</td>
<td>42.6 ± 0.4</td>
<td>17.4 ± 0.3</td>
<td>193 ± 7</td>
<td>0.61</td>
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</table>

Fig. 3  (a) High resolution real space TEM image (scale bar is 2 nm) and (b) TEM diffraction pattern (scale bar is 1 nm\(^{-1}\)) of Nb\(_2\)O\(_5\) film synthesized from 300 nm thick Nb grown on 300 nm thick SiO\(_2\)/Si (001) and annealed at 850 °C for 30 min.
3.2. Composition of Nb and O in Nb₂O₅ film

Fig. 4 shows an XPS spectrum of the sample synthesized from the 5 nm thick Nb film. Six peaks including Nb 3p1/2, Nb 3p3/2, Nb 3d5/2, O 1s, and C 1s in the binding energy range from 150 to 550 eV are observed. The inset shows Nb 3d5/2 and Nb 3d3/2 peaks collected in a smaller energy step size from 190 to 215 eV. Quantitative composition of Nb and O near the surface can be calculated from the ratio of the Nb 3d5/2 peak area divided by the Nb sensitivity factor ($S_{\text{Nb}} = 1$) to the O 1s peak area divided by the O sensitivity factor ($S_{\text{O}} = 0.63$). The XPS background intensity was first subtracted from the peak area using a Shirley background subtraction and then the peak area was fitted using a Gaussian intensity distribution. The peak area and fitting error were obtained from the Gaussian fit. The peak positions for Nb 3d5/2, Nb 3d3/2, and O 1s are 207.2, 209.9 and 530.2 eV, respectively. Table 2 lists XPS peak positions measured from the Nb film and Nb₂O₅ films synthesized from the 5, 10 and 300 nm thick Nb films. Our measured binding energy of 207.2 eV from Nb 3d5/2 peak of synthesized Nb₂O₅ films is close to the binding energy 207.4 ± 1.14 eV of major oxide species reported from pure Nb₂O₅. For comparison, we measured the Nb 3d5/2 peak at 202.2 eV from a pure Nb film. We observed a shift of 5.2 eV in the Nb 3d5/2 peak position for Nb₂O₅ films from that of the Nb film. This 5.2 eV shift is in agreement with the literature reported shift. Our measured 202.2 eV binding energy for Nb 3d5/2 from the Nb film is close to 202.1 ± 0.79 eV recorded in the XPS handbook. The measured binding energy of 530.2 eV from O 1s peak is also close to the O 1s binding energy of 530.4 ± 1.35 eV recorded in the XPS handbook.

The O 1s and Nb 3d5/2 peak areas are 603 and 361 in arbitrary units, respectively. The ratio of O/Nb = (area of O 1s/S_o)/(area of Nb 3d5/2/S_Nb). The calculated O to Nb ratio is 5.3/2.0. The O used in the calculated ratio of O to Nb is estimated from all detected oxygen in the surface. In addition to O that bonds to Nb, there also exists about 10% C in the surface. This means that part of the total detected O signal may come from CO or CO₂ or H₂O near the surface. Thus our calculated ratio of O to Nb is an overestimate. We expect the actual O to Nb ratio to be less than 5.3/2.0. Similarly, the O to Nb ratio has also been obtained from films synthesized from 10 and 300 nm thick Nb films (these spectra are not presented here). See Table 2 for the ratio of O to Nb from three film thickness. The ratios are each close to 5/2.

When the photon energy increases, it can penetrate deeper into the surface of the Nb₂O₅ film. It has been shown that energy resolved XPS with increasing photon energy (100–1600 eV) can reveal the nature of the suboxides. Starting at 750 eV, the suboxides of Nb including NbO₂, NbO and Nb₂O (binding energy between 203 to 209 eV) exist as a transition zone between Nb₂O₅ and the Nb metal (binding energy 202.5 eV). The X-ray photon energy of 1254 eV that we used can reveal the existence of the Nb suboxides in the transition zone if they exist. The fact that our XPS spectra do not reveal suboxides of Nb, but only Nb₂O₅ implies that the Nb has been fully consumed to form Nb₂O₅ and any suboxides have been converted to Nb₂O₅ after the synthesis. Therefore, there are neither detectable suboxides of Nb nor pure Nb near the surface in our sample synthesized from the 5 nm thick Nb film. Another study also supports that the outermost Nb oxide layer is always Nb₂O₅. A similar XPS
spectrum shows that no suboxide exists in the sample synthesized from the 10 nm thick Nb film. In contrast, for the film synthesized from the 300 nm thick Nb film under the same condition of 850°C annealing for 30 min, there is pure Nb near the surface. This implies that within the annealing temperature and time, the oxidation process has not consumed all the Nb in the 300 nm thick Nb film and converted it to Nb$_2$O$_5$. This is why a small (110) peak from pure Nb was observed in XRD.

### 3.3. Optical bandgap of Nb$_2$O$_5$ film on quartz measured from UV-visible absorption

Fig. 5(a) shows the UV-vis reflectance and transmission spectra of the Nb$_2$O$_5$ film synthesized from the 10 nm thick Nb film on quartz. These spectra have been corrected for the quartz background measured from an annealed quartz substrate without the Nb film. A small bump exists in the reflectance spectrum around a wavelength of 300 nm. The transmittance spectrum shows a high transmittance in the visible wavelength range, which decreases in the UV wavelength range. The resulting absorptivity coefficient, $\alpha^*$ of the Nb$_2$O$_5$ film (Fig. 5(b)) shows a sharp onset around 4 eV. The optical bandgap of the Nb$_2$O$_5$ film is determined from the experimental UV-vis absorption spectra via the Tauc method by extrapolating the linear portion of the $(\alpha h\nu)^{1/2}$ vs. $h\nu$ plot to the intercept, where $\alpha$ is the absorption coefficient, $h\nu$ is the photon energy, and the exponent 1/2 is for an indirect bandgap. Fig. 5(c) shows the Tauc analysis for the Nb$_2$O$_5$ film synthesized from the 10 nm thick Nb film on quartz which gives an optical bandgap of 3.65 ± 0.02 eV (=340 nm).

The bandgap for bulk Nb$_2$O$_5$ is about 3.4 eV. Experimentally measured bandgaps for films range from 3 to over 4 eV depending on the structural phase, grain size, and morphology.

![Atomic force microscopy images](image_url)\[Fig. 6\] Atomic force microscopy images (5 μm × 5 μm) and histograms of (a) 5 nm thick Nb film and (b) 10 nm thick Nb film on SiO$_2$/Si (001) before synthesis. AFM images (1 μm × 1 μm) of Nb$_2$O$_5$/SiO$_2$/Si (001) synthesized from (c) 5 nm thick Nb film and (d) 10 nm thick Nb film.

### Table 2  Measured binding energies of Nb in Nb film and Nb in Nb$_2$O$_5$ films from XPS. The "—" means no peak

<table>
<thead>
<tr>
<th>Samples</th>
<th>Binding energy of Nb in Nb film (eV)</th>
<th>Binding energy of major species in Nb$_2$O$_5$ films (eV)</th>
<th>Ratio of O to Nb</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Nb 3d$_{3/2}$</td>
<td>Nb 3d$_{3/2}$</td>
<td>Nb 3d$_{3/2}$</td>
</tr>
<tr>
<td>Nb$_2$O$_5$ synthesized from</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5 nm Nb</td>
<td>202.2</td>
<td>205.0</td>
<td>—</td>
</tr>
<tr>
<td>5 nm Nb</td>
<td>—</td>
<td>—</td>
<td>207.2</td>
</tr>
<tr>
<td>10 nm Nb</td>
<td>—</td>
<td>—</td>
<td>207.2</td>
</tr>
<tr>
<td>300 nm Nb</td>
<td>—</td>
<td>204.9</td>
<td>207.2</td>
</tr>
</tbody>
</table>
of the Nb2O5 film as well as the exponent used in the Tauc plot.\textsuperscript{8,23} Our measured bandgap of 3.65 eV is close to that measured from single crystalline Nb2O5 nanorods grown by hydrothermal technique\textsuperscript{18} using UV-vis (note that the exponent of \(a\)\textsubscript{ph} used in that work was 2 instead of 1/2). Our value is also comparable to the value of 3.7 eV determined for pulsed laser deposited Nb film exposed in air and measured by ultraviolet photoelectron spectroscopy.\textsuperscript{3} Density functional theory (DFT) calculations of Nb2O5 indicate that Nb2O5 is an indirect bandgap semiconductor with a bandgap of 2.55 eV. The underestimation by DFT calculations is typical due to the discontinuity of exchange correlation energy.\textsuperscript{5}

The reflectance and transmission spectra of Nb2O5 film synthesized from 10 nm thick Nb film shown in Fig. 5 do not have the interference feature as compared with those of the few hundreds of nm thick Nb2O5 samples grown on Si substrate or glass substrates reported in previous work.\textsuperscript{3,4,5} Unlike their thicker films, our film has a much smaller dimension than the UV-vis wavelength used in the measurement and the wave interference from the film and substrate interface does not occur.

### 3.4. Film morphology and roughness

#### 3.4.1. Atomic force microscopy

Fig. 6 shows AFM images and height histograms of the 5 and 10 nm thick Nb films on SiO\textsubscript{2}/Si (001), and the Nb2O5 films synthesized from the 5 and 10 nm thick Nb films on SiO\textsubscript{2}/Si (001) substrates. The 5 and 10 nm Nb films shown in Fig. 6(a) and (b) are smooth and almost featureless with root mean square (RMS) roughnesses in the sub-nm range (~0.6 nm). In contrast, the synthesized Nb2O5 films shown in Fig. 6(c) and (d) have island-like features. The height histograms shown below the AFM images range from 0 to 25 nm with a peak around 12 nm and 0 to 30 nm with a peak around 17 nm for the Nb2O5 films synthesized from the 5 and 10 nm thick Nb films, respectively.

We analyzed the two-dimensional height–height correlation function, \(H(r)\) from these AFM images to extract the roughness parameters including the vertical surface width, \(\omega\) (or RMS roughness), lateral correlation length, \(\xi\), and roughness exponent, \(\alpha\).\textsuperscript{39–41} \(H(r)\) is defined as \(H(r) = \langle [h(r) - h(0)]^2 \rangle\), where \(h(r)\) is the surface height at position \(r\) and \(h(0)\) is the surface height at a reference position \(r = 0\). Surface width, \(\omega\) is the square root of \(\langle [h(r) - h(0)]^2 \rangle\), where \(h\) is the average height of the surface. \(\omega\) describes the surface roughness at large distances. The lateral correlation length, \(\xi\) describes the “wavelength” of the surface fluctuations. For large \(r\), the surface height fluctuations should not be correlated, and \(H(r) = 2\omega^2\).

For small \(r\), the surface height–height correlation has the asymptotic form \(H(r) \sim r^{2\alpha}\), where \(\alpha\) is the local roughness exponent. \(\alpha\) describes how "wiggly" the surface is and has a value \(0 < \alpha \leq 1\), with 1 being the smoothest.

To perform the height–height correlation function analysis, one needs to be aware of the sampling induced oscillation.\textsuperscript{42} If the sampling size \(L\) is sufficiently larger than the correlation length \(\xi\), then the oscillation will diminish when \((\xi/L)^{d/2} \ll 1\), where \(d\) is a dimension, and when a large number of images from a rough surface are averaged. If the oscillations do not diminish under these conditions, then the surface is a mounded rough surface. Table 1 lists the height–height correlation function analyses of ten AFM images from each of these Nb2O5 films synthesized from the 5, 10 and 300 nm thick Nb films. Table 1 indicates that the vertical surface width \(\omega\) increases from 3.6 ± 0.2 nm to 17.4 ± 0.3 nm, and the lateral correlation length \(\xi\) increases from 84 ± 2 nm to 193 ± 7 nm. The local roughness parameter \(\alpha\) decreases from 0.78 to 0.61 and becomes more wiggly as the film thickness increases. The 1 × 1 \(\mu\)m sampling size \(L\) of the ten AFM images and the ~90 nm size of \(\xi\), which gives \((\xi/L)^{d/2} \approx 0.09\) for \(d = 2\). After averaging 10 images, the height–height correlation function still has obvious oscillations with regular separation. This is consistent with results obtained for a rough surface with mounds.\textsuperscript{43} These mounds are seen as islands in the AFM images shown in Fig. 6(c) and (d).

From Table 1, the lateral correlation length \(\xi\) extracted from AFM is larger than the average grain size obtained from GIXRD. This implies that there is more than one grain in each small AFM image. The model was fitted to the data using the

![Fig. 7 X-ray specular reflectivity curve (black filled circles) and fitted model (red curve) vs. momentum transfer \(q\) for the Nb2O5 film synthesized from 10 nm thick Nb film on quartz substrate.](image-url)
thickness of the niobium oxide film, and the roughness and density of both the film and substrate as variables.

The Nb₂O₅ film thickness was determined to be 22.57 ± 0.09 nm with a surface roughness of 2.2 ± 0.3 nm from the XRR curve. This thickness is higher than the histogram shown in Fig. 6(b) probably due to the AFM tip size effect when imaging the depth of closely spaced small features. The density was determined to be 5.34 ± 0.07 g cm⁻³, which agrees, approximately, with the orthorhombic density of 5.24 g cm⁻³ (ref. 32) and is slightly higher than the T′ and T phases of Nb₂O₃, of 4.99 and 5.00 g cm⁻³, respectively. Using the density of the 10 nm Nb film thickness, and considering the stoichiometric ratio in mass change by converting Nb to Nb₂O₅ (2 Nb + 2.5 O² → Nb₂O₅), we can estimate the expected density of the 22 nm thick Nb₂O₅ film to be 5.43 g cm⁻³.

3.5. Vibrational modes of Nb₂O₅ films from Raman spectra

Fig. 8 shows three Raman spectra: (a) the reference spectrum of a bare SiO₂ covered Si (001) substrate after 850 °C annealing for 30 min; (b) the spectrum of the Nb₂O₅ film synthesized from the 5 nm thick Nb film; (c) the difference between the Nb₂O₅ film synthesized from the 5 nm thick Nb film and the reference spectrum. The reference spectrum in Fig. 8(a) shows two Si peaks, a sharp peak around 521 cm⁻¹ and a broad peak around 972 cm⁻¹. Both Si peaks also show up with high intensity in the Raman spectrum of the Nb₂O₅ film shown in Fig. 8(b). For the 532 nm laser wavelength that we used in our experiment, the optical penetration depth is about 0.7 μm and the photon energy is 2.33 eV, which is higher than the Si bandgap of 1.11 eV. Raman resonances occur at 521 cm⁻¹ and 972 cm⁻¹ as shown in Fig. 8(a). In order to remove the observed enhancement effect in Raman intensity from the Si (001) substrate when a 532 nm incident laser wavelength is used, we subtracted the reference spectrum shown in Fig. 8(a) from the spectrum shown in Fig. 8(b). The difference spectrum shown in Fig. 8(c) has five distinct Raman peaks at 238, 304, 647, 695, and 993 cm⁻¹.

Among the five peaks, the full-width-at-half-maximum (FWHM) of the Raman peaks in the difference spectrum Fig. 8(c) varies from sharp to broad. The broad peak may contain more than one peak. Using Origin peak fit software we have fitted the broad peaks and obtained extra peaks with the coefficient of determination (goodness of fit) = 0.992. The positions of peaks in the Raman difference spectra from our Nb₂O₅ films synthesized...
Vibrational mode & Raman shift range (cm\(^{-1}\)) & Raman shift of film synthesized from 5 nm Nb & 10 nm Nb & 300 nm Nb & Monoclinic Nb\(_2\)O\(_5\) powders\(^{47}\) nanowires on Nb foil in oxygen plasma\(^{10}\) \\
\(T_{2u}\) & 200–300 & 217.65 & 217.63 & 220.58 & 207 (M) \\
 & & 238.70 & 238.71 & 238.71 & 235 (S) \\
 & & 304.29 & 304.25 & 268.81 & 261 (S) \\
 & & 311.96 & 312.14 & 301.31 & 305 (M) \\
 & & 324.23 & 324.22 & 359.27 & \\
\(T_{2g}\) & 350–560 & 470.51 & 470.51 & 470.51 & 347 (W) \\
 & & & & & 471 (W) \\
TO mode \(E_g\) & 600–700 & 647.65 & 647.65 & 647.24 & 647 (S) \\
 & & 666.88 & 666.85 & 682.63 & 659 (S) \\
 & & 695.72 & 695.75 & & 674 (S) \\
LO mode \(A_{1g}\) & 700–1000 & 722.79 & 725.23 & 707.47 & 760 (W) \\
 & & & & & 846 (W) \\
 & & & & & 897 (W) \\
 & & 962.65 & 962.68 & 962.21 & \\
 & & 992.62 & 993.54 & 993.84 & 992 (S) \\

from the 5, 10, and 300 nm Nb films are listed in columns 3, 4 and 5 in Table 3. For a comparison, Nb\(_2\)O\(_5\) powder samples\(^{47}\) and Nb\(_2\)O\(_5\) nanowires\(^{47}\) are also listed in the last column in the same table. Note that the phase of Nb\(_2\)O\(_5\) in these works is monoclinic \((a = 2.13 \text{ Å}, b = 3.81 \text{ Å}, c = 1.95 \text{ Å})\) and not orthorhombic. The laser wavelength used in Raman scattering is 514.5 nm. \(^{47}\)

The laser, the substrate has no effect on the Raman spectrum. The theoretical modeling is needed to understand the observed difference in Raman shifts shown in Table 3.

Table 3 lists the vibrational modes and their ranges of Raman shifts in columns 1 and 2, respectively. In the 700–1000 cm\(^{-1}\) range, the Raman peak around 993 cm\(^{-1}\) is a longitudinal optical (LO), \(A_{1g}\) mode using Mulliken symbols. \(^{11,14}\) This vibration is most prominent in oxide systems. It is a symmetric single stretching vibration mode (symbol A) of Nb and O double bond (Nb=O) relative to the principle rotational axis (symbol 1) and an inversion through the center of the molecule (symbol G). This stretching mode has a short bond and is also supported by the DFT calculation result described previously. \(^{48}\) This peak is also sensitive to ordering. The peak intensity is either low or zero if the structure is disordered. \(^{48}\) The FWHM of this peak is \(\sim 20 \text{ cm}^{-1}\), which is relatively narrow, implying a reasonably ordered film. In the 700–1000 cm\(^{-1}\) range there are two other peaks at 962 and 722 cm\(^{-1}\). These are single Nb and O (Nb=O) stretching bonds associated with NbO\(_6\) and NbO\(_4\). In the range of 600–700 cm\(^{-1}\) we observed three transverse optical (TO), \(E_g\) (symbol E is doubly degenerated modes) modes. In the range of 200–300 cm\(^{-1}\) we observed \(T_{2u}\) modes (T is triply degenerated modes, 2 and \(\mu\) are anti-symmetric to a rotation axis and an inversion, respectively). However, for the Nb\(_2\)O\(_5\) films synthesized from 5 and 10 nm thick Nb films, we did not observe \(T_{2g}\) modes in the range of 350–550 cm\(^{-1}\).

4. Discussions

4.1. Formation of Nb\(_2\)O\(_5\) films in Ar flow

Previous ultraviolet photoemission spectroscopy studies of oxygen adsorption on Nb (110) surface at 300 K in ultrahigh vacuum showed that oxygen molecules dissociated and chemisorbed on the Nb (110) surface. NbO and NbO\(_2\) formed initially as protective layers and then Nb\(_2\)O\(_5\) formed a heavy oxidation. \(^{39}\) In our case, the sputtered Nb films deposited on SiO\(_2\)/Si (001) were heated up to 850 °C (1123 K) in a furnace and the oxygen came from the exogenous oxygen in the SiO\(_2\) and as impurity in the Ar flow. Based on the heat of formation of Nb\(_2\)O\(_5\) as compared to that of SiO\(_2\), Nb is not an effective reactant towards SiO\(_2\). \(^{31}\) When Nb was deposited on the SiO\(_2\) surface, the high heat of formation of Nb\(_2\)O\(_5\) \((-1899.54 \text{ kJ mol}^{-1}\)) was higher than that of SiO\(_2\) \((-879.48 \pm 2.93 \text{ kJ mol}^{-1}\)). Assume the following reaction: \(4\text{Nb} + 3\text{SiO}_2 \rightarrow 2\text{Nb}_2\text{O}_5 + 5\text{Si}\). Then \(\Delta H_{\text{reaction}} = 2(-1899.54 \text{ kJ mol}^{-1}) - 5(-879.48 \text{ kJ mol}^{-1}) = +598.32 \text{ kJ mol}^{-1}\). The positive heat of reaction means

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this reaction is endothermic and therefore the reduction of SiO2 cannot be the origin of oxygen for the formation of the Nb2O5 films. The TEM cross section shows that the thickness of SiO2 is about 300 nm. The oxygen must come from external sources such as impurities in the Ar gas supply. During our synthesis of Nb2O5, there was no external oxygen flow or oxygen plasma. The fact that Nb2O5 was formed after 850 °C annealing for 30 min under Ar flow in a furnace implies there were oxygen supplies in the Ar flow. One of the major oxygen containing chemicals in the air is the oxygen molecule. Its density is about 21% of air density (2.504 x 10^{19} atoms per cm^3) or 5.24 x 10^{19} atoms per cm^3 under 1 atm at 20 °C. Other minor oxygen containing chemicals are water molecules (1–5%) and CO2 molecules (0.03%). All these oxygen containing molecules are available for oxidation reaction. The Ar flow reduces the amount of air flow and therefore the oxygen supply from the air is reduced. In an ultrahigh vacuum environment, the Nb2O5 film can only grow to a thickness of ≤2 nm in an ultrahigh vacuum furnace at 1850 °C for 2 hours because of the much reduced amounts of oxygen and water.\(^{26}\) We have also synthesized a Nb2O5 film from the 10 nm Nb film in air without Ar flow. The XRD of this sample shows many additional peaks (not shown here). In contrast, the sample prepared under Ar flow shows only the (001) and (002) XRD peaks (see Fig. 2).

The growth of Nb oxides in the presence of O2, H2O and H2O2 is described by Cabrera Mott theory.\(^{29}\) Once the initial Nb oxide is formed, the anion (O\(^{–}\)) diffusion in the oxide film is significant compared with cation (Nb\(^{3+}\)) diffusion and the new oxide layers are initiated at the oxide–metal interface. The oxidation steps are Nb → NbO → NbO + NbO2 → Nb2O5. The oxygen reacted with the Nb film to form Nb2O5 at the interface between Nb and oxygen in the Ar flow ambient. After all the Nb atoms at or near the interface were consumed, the oxygen reacted with Nb atoms located further away from the interface. The formation of Nb2O5 was supported by our XRD, TEM and XPS results described above.

4.2. Dominant [001] growth orientation in Nb2O5 films

Recent ab initio density functional theory calculation (DFT) of monolayer oxygen adsorption on Nb (110) using linear augmented plane wave method (LAPW) indicates that oxygen atoms form stronger bonds with Nb atoms on the (110) surface in the [001] direction rather in the [110] direction. The stronger bond shortens the bond length between Nb and O in the [001] direction. The shorter bond results in a structural relaxation in neighboring Nb atoms.\(^{49}\) This DFT result qualitatively agrees with our XRD finding that the oxygen reacts with [110] surface of Nb and that the dominant oxide orientation is the [001] direction. Previous work shows that the unit cell of the orthorhombic Nb2O5 has 16.8 Nb ions and 42 oxygen atoms.\(^{50}\) Eight Nb ions are in distorted octahedral positions, and eight Nb ions are in pentagonal bipyramidal positions. Both groups are in a sheet parallel to the [001] plane. These 16 Nb ions are surrounded by six or seven oxygen atoms. The remaining 0.8 Nb ion is positioned at the interstitial nine-coordinated site in the unit cell.\(^{51}\) The adjacent spacing among Nb atoms is not uniform in the [010] or b direction. This is consistent with the high resolution TEM image along the b direction shown in Fig. 3(a).

5. Summary

We report the annealing of Nb films grown on amorphous substrates and show that optically smooth orthorhombic Nb2O5 films can be formed at 850 °C in a short time under Ar flow in a furnace. This takes advantage of the high heat of formation of Nb2O5. The oxide film thickness is two to three times thicker than that of the starting Nb film as shown by TEM and X-ray reflectivity measurements. Therefore the thickness of Nb2O5 film can be controlled by altering the starting Nb film thickness.

The composition ratio of Nb/O determined by XPS is close to 2/5. The optical bandgap is ~3.65 eV. The low optical transmittance in the UV wavelength range and high transmittance in the visible wavelength range indicate potential applications in UV detection, photocatalysis and electrochromic devices. Although we have not measured the mechanical properties of our synthesized Nb2O5 films, other work shows the Nb2O5 films is scratch resistance and hydrophobic.\(^{5}\) Our simple growth method can be scaled up and can have potential realistic applications.

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Notes and references