

# Atomic-Layer-Deposited ultrathin films of vanadium pentoxide crystalline nanoflakes with controllable thickness and optical band-gap

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**Abstract:** Thin films of crystalline  $V_2O_5$  nanoflakes were prepared through atomic layer deposition (ALD) process. The film thickness was verified to play a critical role in determining structural morphology, optical bandgap and Raman vibration of crystalline  $V_2O_5$  thin films. Two optical bandgaps observed at about 2.8 eV and 2.4 eV result from two growth stages during ALD preparation. We expect that these results help to understand the growth control of ultrathin films and their functional devices.

**Key words:** atomic layer deposition, vanadium pentoxide, thin films

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## 厚度与带隙可调、原子层沉积的超薄五氧化二钒纳米晶薄膜

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**摘要:** 利用原子层沉积方法制备  $V_2O_5$  纳米片晶薄膜. 薄膜厚度可以被精确控制, 并对纳米晶  $V_2O_5$  薄膜的结构形貌、光学带隙和拉曼振动有显著影响. 原子层沉积过程中  $V_2O_5$  薄膜生长的两个阶段导致薄膜具有两个光学带隙, 这将有助于理解超薄薄膜生长与功能应用.

**关键词:** 原子层沉积; 五氧化二钒; 薄膜

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## Introduction

Two-dimensional layered transition metal oxides have attracted considerable attention due to their intriguing chemical activity, distinctive electrical and optical properties. The layered nanostructures have shown promises in facilitating functional devices including optical<sup>[1]</sup> and electronic devices<sup>[2]</sup>, sensors<sup>[3]</sup>, and rechargeable lithium batteries<sup>[4,5]</sup>. Among these layered transition metal oxides, layered  $V_2O_5$  materials have been widely investigated due to its low cost, nature abundance and high charge storage capability. Some investigations have recently demonstrated  $V_2O_5$ -based electrochromism<sup>[6-9]</sup>, photocatalysis<sup>[10-12]</sup>, sensor<sup>[13]</sup>, and energy storage<sup>[14-18]</sup>.

Crystalline  $V_2O_5$  thin films play a vital role in improving the performances of functional electrodes<sup>[19, 20]</sup>. In order to optimize electronic or ionic conductivity, the nanoscale thicknesses of  $V_2O_5$  films are controlled to improve ionic diffusion<sup>[21]</sup> and electronic transportation<sup>[22, 23]</sup>. In general, the film thicknesses of layered  $V_2O_5$  materials intensely affect their optical and electrical properties. It is known that atomic layer deposition (ALD) is an excellent technique to precisely control the thickness of functional films on complicated configurations<sup>[24]</sup>. In this paper, we present the preparation of crystalline  $V_2O_5$  thin films with controllable thickness of several nanometers through an ALD process at low temperature. The investigations focus on the relationships between the film thickness and surface morphology, optical bandgap and pho-

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non vibration of crystalline  $V_2O_5$  thin films. Meanwhile, the distinctions between the initial stage and the follow-up stage are discussed to reveal the growth process of  $V_2O_5$  ultrathin films during the ALD process.

## 1 Experiments

$V_2O_5$  thin films were deposited on the silicon or silica substrates by using oxygen plasma and triisopropoxyvanadium oxide (VTOP) as precursors in an ALD reactor (Picosun-R200) at 200°C.  $N_2$  and Ar were used as carrier gases for oxygen plasma and VTOP respectively. When the pressure was 80 Pa in the ALD chamber, the pulse time of VTOP and oxygen plasma were 0.7 s and 1.5 s respectively.

The surface morphologies were examined by atomic force microscopy (AFM, NT-AIST) in a tapping mode. Scanning electron microscopy (SEM) images were obtained on FEI Sirion200, and transmission electron microscopy (TEM) images were obtained on JOEL JEM-2100F performed at an accelerated voltage of 200 kV. X-ray photoelectron spectroscopy (XPS; Axis Ultra DLD) was used for elements analysis. X-ray diffraction (XRD) patterns were collected on Bruker D8 with a monochromatic source of Cu-K $\alpha$  radiation ( $\lambda = 0.15418$  nm) at 1.6 kW (40 kV, 40 mA). Raman spectra were obtained using TII Tokyo-Nanofinder 30 at room temperature. Spectroscopic ellipsometry (SE) was performed on J. A. Woollam-M2000X-FB-300XTF at an incident angle of 65°. UV-Vis-NIR absorption spectra were measured using PerkinElmer-Lambda 950 spectroscopy with an integration sphere accessory.

## 2 Results and discussions

Fig. 1 shows SEM and AFM images of  $V_2O_5$  thin films deposited with different thicknesses by controlling ALD cycle numbers on silicon substrates. The representative surface morphology is related to the film thicknesses with different ALD cycle numbers. When the thickness is less than 1500 ALD cycles, the films are with smooth surface, as shown in Fig. 1a and 1b. As the films thickness increases, the small grains grow up and finally evolve into orthorhombic shaped crystallites (Fig. 1c-e). The crystal nucleation and growth may determine the surface morphologies of  $V_2O_5$  thin films due to the formation of  $V_2O_5$  nanoflakes during ALD procedure<sup>[25, 26]</sup>. The use of  $O_2$  plasma leads to the crystalline nanoplates with sharp-shaped edges.

Stoichiometric ratio is critical for vanadium oxides to determine the crystalline structure and morphology. Thus, XPS and XRD were utilized to investigate the compositions and structures of thin films with different thicknesses (Fig. 2). The Binding energies at about 517.24 and 524.68 eV result from  $2p_{3/2}$  and  $2p_{1/2}$  of vanadium element. There is an energy difference of nearly 7.4 eV between V  $2p_{3/2}$  and  $2p_{1/2}$ , which implies the existence of  $V^{+5}$  oxidation state<sup>[27]</sup>. In addition, the  $V^{+4}$  oxidation state is not detected in XPS. All these results suggest the formation of  $V_2O_5$  thin films during ALD procedure. For two thin films with thicknesses of 500 and 2000 ALD cy-

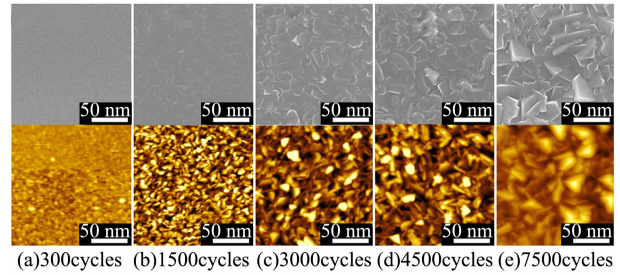


Fig. 1 Top-view SEM (up) and AFM (down) images of  $V_2O_5$  thin films with varying thicknesses of (a) 300, (b) 1500, (c) 3000, (d) 4500, (e) 7500 ALD cycles

图1 不同厚度五氧化二钒的SEM(上部)和AFM(下部)表面形貌:(a)300循环(b)1500循环(c)3000循环(d)4500循环(e)7500循环

cles, the band energies of V 2p are almost the same. However, except for O 1s peak at 530.36 eV, there is an obvious peak at a binding energy of  $\sim 532.1$  eV (Figure 2b) in the case of the films with a thickness of 500 ALD cycles. This result may be related to  $H_2O$  molecules absorbed in the films, and the thinner films are easily affected by the adsorbed molecules.

Fig. 3 shows Raman spectra of the lattice variations in  $V_2O_5$  thin films with different thicknesses. For all the  $V_2O_5$  films with different thicknesses by varying ALD cycles, Raman shifts at 102, 145, 197, 284, 405, 484, and  $700\text{ cm}^{-1}$  are in agreement with those previously reported<sup>[28]</sup>. The intensity of Raman peak rises as the thickness of  $V_2O_5$  film increases (Fig. 3a). The Raman shifts at 303 and  $526\text{ cm}^{-1}$  are indistinguishable because the Si substrates have obvious Raman shifts at about 300 and  $520\text{ cm}^{-1}$ . Raman shifts at  $145\text{ cm}^{-1}$  are the most obvious and come from a mixture of  $B_{1g}$  and  $B_{3g}$  vibrations. The  $B_{1g}$  and  $B_{3g}$  modes correspond to the shear and rotation motion of V-O<sub>B</sub> bond ladders, respectively (Fig. 3e)<sup>[28]</sup>. This bending mode is strongly associated with the layered structure<sup>[29]</sup>. There is a slight blue-shift at  $145\text{ cm}^{-1}$  when the thickness of  $V_2O_5$  thin film decreases from 1500 cycles to 300 cycles, which indicates the presence of a little strain occurred in the initial growth stage of the layered thin films (Fig. 3b). This observed shift may result from small compression strain along c-axis in the layered  $V_2O_5$  structure reported elsewhere<sup>[30, 31]</sup>. No detectable changes are found in Fig. 3c and 3d.

As mentioned above, for thin films with varied thicknesses, the surface morphology and XPS spectra are different. Thus, spectroscopic ellipsometry (SE), cross-sectional SEM and TEM measurements are further performed to understand the essential growth process of  $V_2O_5$  ultrathin films. Fig. 4a shows the relationship between the thickness and ALD cycle number. The thicknesses are obtained from SE and SEM measurements. As shown in Fig. 2f, there is a silica layer with about 2 nm between silicon substrate and  $V_2O_5$  thin film. Thus, a three-layer model is used in SE data and analysis. Moreover, the surface roughness is also considered in SE a-

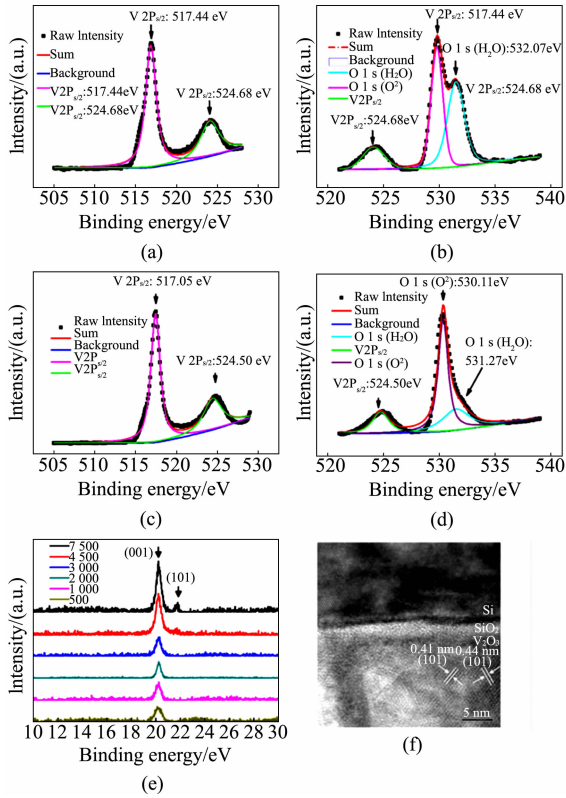


Fig. 2 (a, b)  $V 2p_{3/2}$  and  $O 1s$  peaks of XPS spectra for  $V_2O_5$  films with a thickness of 500 ALD cycles, respectively, and (c, d) with a thickness of 2000 ALD cycles. (e) XRD patterns of  $V_2O_5$  films with varying thickness; (f) TEM image of  $V_2O_5$  films with a thickness of 2000 ALD cycles

图 2 (a, b) 500 循环五氧化二钒薄膜的  $V 2p_{3/2}$  和  $O 1s$  的 XPS 谱, (c, d) 2000 循环五氧化二钒薄膜的  $V 2p_{3/2}$  和  $O 1s$  的 XPS 谱, (e) 不同厚度五氧化二钒的 XRD 谱图 (f) 2000ALD 循环五氧化二钒薄膜的 TEM 图

analysis and can be obtained from the RMS roughness in AFM measurements. For example, for the  $V_2O_5$  film prepared with 7500 ALD cycles, we can use SE simulation to obtain a thickness of 215 nm, which is also verified by cross-sectional SEM measurements. Fig. 4a suggests the effective thickness increases linearly with ALD cycles. In the range from 1500 to 7500 cycles, the growth rate is about  $0.3 \text{ \AA}/\text{cycle}$ . However, the growth rate is about  $0.2 \text{ \AA}/\text{cycle}$  in the range from 100 to 1500 cycles. The ratio of Rms roughness to thickness remains relatively stable although the Rms roughness increases with the increasing ALD cycles, as shown in Fig. 4b. The variation trends of both growth rate and surface morphology indicate that there are two growth stages during the formation of  $V_2O_5$  thin films, which may be understood with the Stranski-Krastanov growth mode<sup>[32]</sup>. Once the film thickness is up to a critical thickness, the growth modes change from the layer growth to the island-based growth. In the initial layer growth stage, the  $V_2O_5$  films are very thin and almost parallel to surface by the reaction between ALD precursors and substrates. The growth is self-

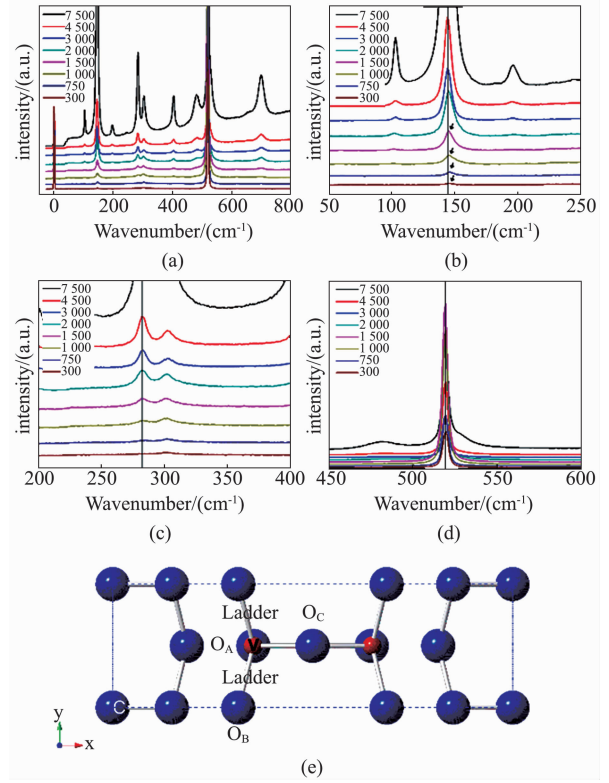


Fig. 3 (a) Raman spectra of  $V_2O_5$  films with various thicknesses; (b, c, d) a zoom at the ranges of  $50 \sim 250 \text{ cm}^{-1}$ ,  $200 \sim 400 \text{ cm}^{-1}$  and  $450 \sim 600 \text{ cm}^{-1}$ , respectively. (e) Schematic molecular structure of  $V_2O_5$

图 3 (a) 不同厚度五氧化二钒薄膜的拉曼图谱, (b, c, d)  $50 \sim 250 \text{ cm}^{-1}$ ,  $200 \sim 400 \text{ cm}^{-1}$  和  $450 \sim 600 \text{ cm}^{-1}$  的局部放大, (e) 五氧化二钒的分子结构模型

limiting on the substrate, which results in a low surface roughness. Once the  $V_2O_5$  ultrathin films form, the precursor molecules are easily adsorbed and selectively reacted with  $V_2O_5$  nanoflakes, which leads to an improved growth rate and an island-based growth mode.

Fig. 5 displays Tauc plots derived from SE and UV-Vis-NIR absorption spectra. The  $V_2O_5$  ultrathin film has an indirect bandgap, and the bandgap red-shifts from 2.66 eV to about 2.40 eV when the thickness increases from 50 to 300 cycles (Fig. 5a). Furthermore, Figure 3c suggests a bandgap of 2.36 eV for the thin film with a thickness of 500 ALD cycles prepared on quartz substrate, which is in agreement with SE simulation. The red-shift can be attributed to the quantum size effect. Once the thicknesses increase up to more than 1000 ALD cycles, two bandgaps are identified as shown in Fig. 3b and 3d. One stays at 2.40 eV and the other varies from 2.73 eV to 2.90 eV. The formation of two bandgaps might result from the two growth stages mentioned above. In the layer growth stage the bandgap decreases with the thickness increasing. Then, the thin film subsequently grows up and another bandgap appears while the primary bandgap of the initial ultrathin film remains. Small grains form in the thin films during the island-based growth stage, which leads to a larger bandgap. All these results

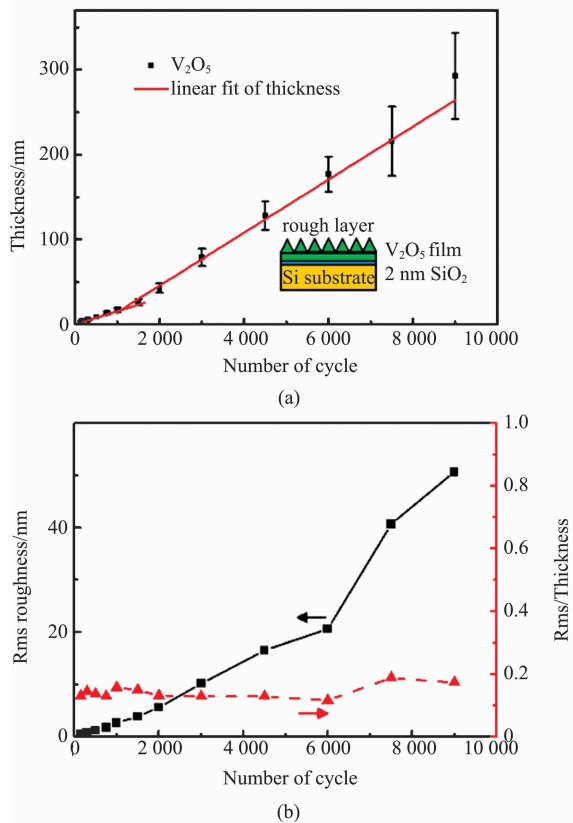


Fig. 4 (a) Relationship between thickness and ALD cycle numbers, the inset shows the schematic structure of thin film deposited on the Si substrate. (b) Plots for the relationships between Roughness and roughness-thickness ratio and ALD cycle numbers

图 4 (a) 厚度和 ALD 循环数的关系, 插入图为硅衬底上五氧化二钒薄膜的模型结构示意图, (b) 表面粗糙度及粗糙度膜厚比值与 ALD 循环数的关系图

suggest that the optical properties of  $V_2O_5$  thin film could be adjusted by the control of film thickness during an ALD process.

### 3 Conclusions

We have successfully prepared  $V_2O_5$  thin films with a thickness down to 5 nm by using ALD process. All investigations on surface morphology, optical bandgap and Raman vibration confirm that the film thickness of crystalline  $V_2O_5$  nanoflakes can be precisely controlled. Meanwhile, the two observed optical bandgaps result from two growth stages during the ALD preparation of crystalline  $V_2O_5$  thin films. We expect that these results will enable us to understand the growth control of ultra-thin films and their applications in functional optical and electrochromic devices.

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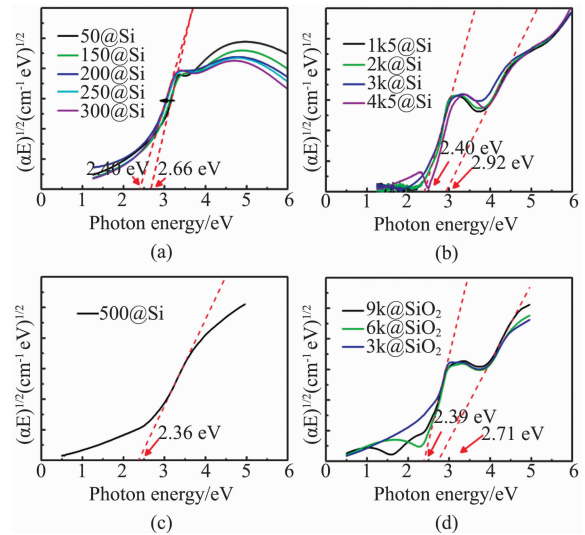


Fig. 5 Tauc plots of  $V_2O_5$  films with various thicknesses of (a) 50 to 300 ALD cycles on Si substrates; (b) 1500 to 4500 ALD cycles on Si substrates; (c) 500 ALD cycles on quartz substrates; (d) 3000, 6000 and 9000 ALD cycles on quartz substrates

图 5 不同厚度五氧化二钒薄膜的能带吸收关系图: (a) 硅衬底上 50 到 300 ALD 循环的薄膜, (b) 硅衬底上 1500 到 4500 ALD 循环的薄膜, (c) 石英衬底上 500 循环 ALD 的薄膜, (d) 石英衬底上 3000 到 9000 ALD 循环的薄膜

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