Comparison of chemical stability and corrosion resistance of group IV metal oxide films formed by thermal and plasma-enhanced atomic layer deposition

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The wide applications of ultrathin group IV metal oxide films (TiO$_2$, ZrO$_2$, and HfO$_2$) probably expose materials to potentially reactive etchants and solvents, appealing for extraordinary chemical stability and corrosion resistance property. In this paper, TiO$_2$ ultrathin films were deposited on Si at 200 °C while ZrO$_2$ and HfO$_2$ were grown at 250 °C to fit their growth temperature window, by thermal atomic layer deposition (TALD) and plasma-enhanced ALD (PEALD). A variety of chemical liquid media including 1 mol/L H$_2$SO$_4$, 1 mol/L HCl, 1 mol/L KOH, 1 mol/L KCl, and 18 MΩ deionized water were used to test and compare chemical stability of all these as-deposited group IV metal oxides thin films, as well as post-annealed samples at various temperatures. Among these metal oxides, TALD/PEALD HfO$_2$ ultrathin films exhibit the best chemical stability and anti-corrosion property without any change in thickness after long time immersion into acidic, alkaline and neutral solutions. As-deposited TALD ZrO$_2$ ultrathin films have slow etch rate of 1.06 nm/day in 1 mol/L HCl, however other PEALD ZrO$_2$ ultrathin films and annealed TALD ones show better anti-acid stability, indicating the role of introduction of plasma O$_2$ in PEALD and post-thermal treatment. As-deposited TiO$_2$ ultrathin films by TALD and PEALD are found to be etched slowly in acidic solutions, but the PEALD can decrease the etching rate of TiO$_2$ by ~41%. After post-annealing, TiO$_2$ ultrathin films have satisfactory corrosion resistance, which is ascribed to the crystallization transition from amorphous to anatase phase and the formation of 5% Si-doped TiO$_2$ ultrathin layers on sample surfaces, i.e. Ti-silicate. ZrO$_2$ and TiO$_2$ ultrathin films show excellent corrosion endurance property in basic and neutral solutions. Simultaneously, 304 stainless steel coated with PEALD-HfO$_2$ is found to have a lower corrosion rate than that with TALD-HfO$_2$ by means of electrochemical measurement. The pre-treatment of plasma H$_2$ to 304 stainless steel can effectively reduce interfacial impurities and porosity of overlayers with significantly enhanced corrosion endurance. Above all, the chemical stability and anti-corrosion properties of IV group metal oxide coatings can be improved by using PEALD technique, post-annealing process and plasma H$_2$ pre-treatment to substrates.
uniformity, excellent three-dimensional conformality, simple and precise control in film thickness, dense and pinhole-free films, and low processing temperature. Compared with the conventional thermal ALD (TALD), plasma-enhanced ALD (PEALD) is an energy-assisted method for fabrication of thin films, where plasma species are utilized as reactive gas during one step of the cyclic deposition process. It can produce some virtues over the TALD route, including more choice of materials and precursors, substrate temperature at room temperature, improved film density and quality with high purity and little defects, etc. Therefore, extensive and intensive studies on ALD-derived ultrathin films as protective coatings and anti-corrosion barriers are needed greatly. In consideration of better film quality, PEALD may be the more suitable candidate to obtain the ultrathin corrosion-resistant coatings with long-term stability in severe environments.

So far, some researches on surface passivation and chemical protection by thermal ALD-derived thin films have been performed, the majority of which focus on several common metal oxides like alumina and titania formed by TALD and their stability in various environments, especially in aqueous solutions for photoelectrochemical water splitting and protection of stainless steel or other metal substrates. For example, Abdulagatov et al. deposited Al2O3 and TiO2 on Cu and found that only Al2O3 or TiO2 TALD films were insufficient to prevent copper from corrosion whereas the introduction of TiO2 capping layer on Al2O3 films was more resilient to dissolution in water. Strandwitz et al. have demonstrated TALD-derived HfO2 thin films can be used to stabilize n-Si photoelectrodes in chemically reactive conditions. The chemical stability of TALD Al2O3 and TiO2 films in different acidic, basic and neutral media, and the influence of post-deposition thermal treatment on different samples have been discussed. These valuable results are helpful to expansive applications of organic electronic devices, stabilization of semiconductor photoanodes for water oxidation and many other fields.

In this paper, the chemical stability of as-deposited and post-annealed TiO2, ZrO2 and HfO2 thin films derived from TALD and PEALD in various chemical liquid media, including 1 mol/L H2SO4, 1 mol/L HCl, 1 mol/L KOH, 1 mol/L KCl, and pure water, has been investigated and compared systematically. A series of analytical methods were used to characterize the change of thickness, chemical composition, structure and morphology of different samples. It is found that TALD/PEALD HfO2 films exhibit the best chemical stability and anti-corrosion property without any change in thickness after long time immersion into various chemical solutions. The corrosion endurance of HfO2 coating for 304 stainless steel (SS) in 1 mol/L KCl solution has been studied by in-situ electrochemical characterization.

Results and Discussion

Composition, morphology and structure. Chemical composition of all as-deposited TALD and PEALD samples on Si was examined by XPS, as seen in Fig. 1. The strong doublet peaks of Ti 2p (a), Zr 3d (b) and Hf 4f (c) with respective splitting energy of 5.7 eV, 2.4 eV, and 1.7 eV, and the strong O 1s signal at 529.83 eV (d, only showing TALD and PEALD HfO2) verify the formation of TiO2, ZrO2 and HfO2 films by TALD and PEALD. Herein the weak O 1s peak at 531.73 eV might come from the surface adsorption oxygen of HfO2 films.

After 600 °C anneal, there is no obvious change in XPS spectra of ZrO2 and HfO2 films on Si. However, it is found that after 450 °C and 900 °C thermal treatment, the Si 2p signal appears on the surface of TiO2 films/ Si. After anneal, both samples exhibit Si 2p peaks at 101.98 eV, assigned to the chemical bond of Ti-O-Si. After 450 °C anneal, both samples exhibit Si 2p peaks at 101.98 eV, assigned to the chemical bond of Ti-O-Si. When further raising the anneal temperature to 900 °C, the Si 2p peaks shift to higher binding energy of 102.48 eV, indicating that more Si from substrate diffuses onto TiO2 film surface and produce more Ti-silicate.

The XPS composition data attest that the atomic ratio of Si: Ti in the film surface is 0.25:1.00 at 450 °C and 0.31:1.00 at 900 °C for TALD samples, and in PEALD samples surface, the atomic ratio of Si: Ti is 0.28:1.00 at 450 °C and 0.55:1.00 at 900 °C, respectively. The content of Si element on the surface of annealed TiO2 films is rather high, so XPS depth analysis on as-deposited, 450 °C and 900 °C annealed TALD and PEALD-TiO2 films surface. No Si 2p signal is detected on the surface of as-deposited TALD and PEALD samples. After 450 °C anneal, both samples exhibit Si 2p peaks at 101.98 eV, assigned to the chemical bond of Ti-O-Si. When further raising the anneal temperature to 900 °C, the Si 2p peaks shift to higher binding energy of 102.48 eV, indicating that more Si from substrate diffuses onto TiO2 film surface and produce more Ti-silicate.

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Atomic percent content of different elements was recorded every 60 s and the etch rate is around 0.56 Å/s. There is hardly Si signal for as-deposited TiO2 film surface, but ~5 at% Si can be clearly observed on the annealed TiO2 film surface. After the first etching process, Si content decreases to zero sharply with increased Ti and O contents, whose ratio is among 1:8–2:1, in basic agreement with the stoichiometric TiO2. When it comes to 420 s, Si starts to emerge and increase gradually, and Ti and O contents become low. The interfacial diffusion layer of (TiO2)m(SiO2)n is formed between TiO2 films and Si substrate because TiO2 film is thermodynamically unstable on Si substrate, which easily diffuses into the film and reacts with TiO2 during thermal treatment, resulting in formation of Ti-silicate.

In addition, with the increase of anneal temperature, the thickness of interfacial diffusion layer becomes larger, which can be verified by the following thickness measurement.

FTIR spectra were also utilized to obtain the chemical group information of TALD-samples before and after thermal anneal, as recorded in Fig. 3. As-deposited sample show Ti-O bond at 614 cm−1 with Ti-Cl bond at 811 cm−1, C-Cl bond at 739 cm−1 and C-OH bond at 1108 cm−1, indicating the chlorine residue in film from TiCl4 precursor and surface carbon contamination due to the lower deposition temperature of 200 °C. Besides Si-O bond, 450 °C- and 900 °C-annealed TiO2 films still display Ti-O-Si and Si-O modes of 1024 cm−1 and 1148 cm−1 with the disappearance of the Ti-Cl, C-Cl and C-OH bonds, confirming the Si element diffusion into films and removal of Cl residue and organic component after high temperature anneal.

SEM and AFM were conducted to characterize the film surface morphologies. Besides, the film thickness can be determined using spectroscopic ellipsometry and cross-section SEM images. The thicknesses of as-deposited and annealed TALD and PEALD TiO2, ZrO2 and HfO2 films from spectroscopic ellipsometry before chemical test are summarized in Table 1. Evidently ZrO2 and HfO2 films become thinner after anneal, which can be ascribed
to remove of oxygen vacancy and densification of films. However, TiO₂ films become thicker after thermal treatment, especially for 900 °C samples with remarkable thickness increase. This result is also verified by cross-section FESEM images of TALD TiO₂ films on Si substrate, as seen in Fig. 4.

Figure 5 is AFM images of as-deposited and annealed TALD and PEALD TiO₂ films. Thanks to the advantage of ALD technique, large area uniformity, the 200 °C as-deposited and 450 °C annealed films are of little roughness with root-mean-square (RMS) value of 0.2 nm, especially for samples formed by TALD. It can be seen from Fig. 5(c,f), after 3-hour 900 °C post annealing in N₂ atmosphere, surface morphology of TALD- and PEALD-TiO₂ films have greatly changed with small particles growing and gathering, the surface roughness increased to 1.8 nm and 3.1 nm respectively. Normally, thermal treatment at high temperature can enlarge grain size in the films, resulting in better crystallinity. Further discussion on structure of samples will be done according to XRD and XPS data.

Similarly, AFM images of as-deposited and annealed TALD ZrO₂ and HfO₂ films are presented in Fig. 6. After 600 °C annealing, the ZrO₂ films have relatively flat surfaces (RMS: 1.42 nm) without easily recognized grains, indicating poor crystallinity. The GIXRD pattern in Fig. 7(b) also proves this. Whereas 600 °C annealed HfO₂ films contain quite a few 20~30 nm particles with enhanced RMS of 2.56 nm. The XRD result in Fig. 7(c) reveals that stable HfO₂ monoclinic phase has been formed.

Due to thinner thickness of metal oxide films deposited by ALD technique, GIXRD with the help of synchrotron radiation was introduced to perform phase analysis. Figure 7 illustrates the GIXRD patterns of as-deposited and annealed TALD TiO₂, ZrO₂ and HfO₂ ultrathin films. As-deposited TiO₂ samples at 200 °C in Fig. 7(a) are amorphous without any diffraction peaks. 450 °C annealed samples show relatively stronger (101) peak at 25.2° and weak peaks of (004), (200), (105) and (211), which can be assigned to the anatase phase. In addition, a wide peak at 27.5° corresponds to the slight rutile phase. So the mixture of main anatase and slight rutile phases exists in 450 °C TiO₂ films. Nevertheless, after 900 °C anneal, the TiO₂ film crystallinity becomes worse, as confirmed by the reduced (101) peak intensity and some disappeared weak peaks. It has been reported that TiO₂ forms the anatase phase when thermal-treated between 300 and 500 °C, and then turns to the rutile structure above 800 °C anneal27. Herein the evident phase discrepancy can be attributed to the severe Si diffusion between TiO₂ ultrathin films and Si substrate, leading to the interfacial diffusion layer and the poor crystalline of TiO₂. Meanwhile, Si-doped TiO₂ films also prevent the transform from anatase to rutile phase. The XRD results are in good agreement with the mentioned-above XPS and FTIR analyses.

Comparing GIXRD patterns of ZrO₂ and HfO₂ films after 10-minute 600 °C RTA (Fig. 7(bc)), obviously HfO₂ films have changed from amorphous to monoclinic phase with sharp peaks. The calculated grain size is 23.3 nm.
by Schereer equation, in consistent with AFM observation. While under the same anneal condition, the crystallinity of ZrO₂ films is not as good as that of HfO₂ and a weak and broad diffusion peak at 30.5° implies the trend of crystallinity.

**Chemical stability in chemical liquid media.** To study anti-corrosion property of group IV metal oxides, we immersed a subset of as-deposited and annealed TALD and PEALD TiO₂ films in various chemical liquid media with a range of pH value, including acid solutions (1 mol/L H₂SO₄, 1 mol/L HCl), an alkaline solution (1 mol/L KOH) and neutral solutions (1 mol/L KCl, 18MΩ pure water). Film thickness was examined using spectroscopic ellipsometry after regular intervals. Figure 8 illustrates the thickness change dependence on immersion time for...
all as-deposited and post-annealed TALD/PEALD-TiO₂ films in various chemical environments, so as to evaluate the influence of anneal and deposition method on etching resistance of TiO₂ films.

From the point of view of film thickness variation, TiO₂ thin films in alkaline and neutral solutions of 1 mol/L KOH and 1 mol/L KCl are very stable, and the film thickness has hardly changed, even if as-deposited ones at 200 °C without post-anneal, as indicated in Fig. 8(c,d). In Fig. 8(e), as-deposited and 450 °C annealed TiO₂ samples in 18 MΩ water show a little slight thickness increase due to the hydration role; after 900 °C post-anneal, the film thickness basically keeps constant.

However, as-deposited TiO₂ thin films are unstable in acidic solutions with slow dissolution during the immersion duration, resulting in the visual change of the films in reflection color. The etch rates in 1 mol/L H₂SO₄ and 1 mol/L HCl in Fig. 8(a,b) are summarized in Table 2.

Evidently H₂SO₄ aqueous solution is more corrosive to as-deposited TiO₂ films than HCl solution at the same molar concentration, because sulfuric acid is a binary strong acid, which can ionize twice amount of H⁺ of single hydrochloric acid. Moreover, the etch rate of as-deposited PEALD samples in acid drops by about 41% than that of as-deposited TALD ones, which can be ascribed to the fact that PEALD-derived samples have enhanced film density and lower impurity concentration due to the O₂ plasma’s higher surface reactivity than only thermal energy-assisted process alone²⁸.

After annealed 450 °C and 900 °C for 3 h, the chemical stability of TALD and PEALD TiO₂ ultrathin films on Si is improved significantly with nearly unchanged thickness in strong acidic solutions over test period of 20 days in Fig. 8(ab). As described above in Figs 2 and 7(a), the XPS and XRD results confirm that the longer time annealing at 450 °C and 900 °C for 3 h causes the Si diffusion into TiO₂ films to form the amorphous Ti-silicate containing partial crystalline anatase TiO₂ main phase. The amorphous Ti-silicate and crystalline TiO₂ exhibit better anti-acid corrosion performance than amorphous TiO₂. Especially, the formation of 5% Si-doped TiO₂ ultrathin layers on sample surfaces in Fig. 2(c,d) also plays an important role in enhancing the acid-endurance.

The thickness variation dependence on immersion time for all as-deposited and post-annealed TALD/PEALD-ZrO₂ films in various chemical environments are recorded in Fig. 9. Except for as-deposited TALD ZrO₂ films dissolving in 1 mol/L HCl with the etch rate of 1.06 nm/day, all other ZrO₂ samples exhibit better stability in different aqueous environments during the whole 20-day test, including annealed TALD and all PEALD samples. Herein, the influence of PEALD is much more significant without any etching than TiO₂ case for O₂ plasma contains more reactive radical and ionic species, beneficial to the corrosion-endurance ZrO₂ films’ growth. Meanwhile annealing at high temperature may remove the residual impurities in films, leading to densification and crystallization trend of ZrO₂ films. This produces positive effects on improving anti-corrosion property of ZrO₂ films, as indicated in Fig. 9(b).

Figure 10 displays the anti-etching characteristics of all TALD and PEALD HfO₂ films in various chemical liquid media. Among three kinds of group IV metal oxides, HfO₂ films may be the best candidate as barrier layer in caustic environments, because after the long-time immersion in any acidic, alkaline and neutral solutions, all the HfO₂ samples show negligible change in thickness, no matter from TALD and PEALD or before and after anneal. So, we chose HfO₂ films as one candidate to further examine the corrosion resistance by electrochemical measurements.

Electrochemical result. We deposited 300 cycles HfO₂ films (~36 nm-thick) onto 304 stainless steels (304 SS) by TALD and PEALD. It was reported that the anti-corrosion properties of ALD Al₂O₃ thin films on steel could be improved by the H₂–Ar plasma pre-treatment²⁹. In our experiment, 304 SS was treated by in-situ 50 s H₂–Ar plasma pulse before HfO₂ deposition so as to study the effect of the H₂ pre-treatment.

The polarization curves and Bode plots for the EIS data measured at OCP for the bare and coated 304 SS in 1 mol/L KCl are shown in Fig. 11. Only a peak at the corrosion potential can be observed in the polarization curves of each sample, indicating the HfO₂ coating films are electrochemically inert³⁰. Compared with the bare 304 SS, the corrosion potential (Ecorr) of TALD- and PEALD-HfO₂ coated 304 SS with and without H₂ plasma pretreatments, shifts to more negative potential, leading to improved corrosion resistance. Corrosion current density (Icorr) is another key factor to evaluate the corrosion reaction kinetics, which is usually proportional to the

| Table 1. Thicknesses of as-deposited and annealed TiO₂, ZrO₂ and HfO₂ films prepared by TALD and PEALD from spectroscopic ellipsometry. |
|---|---|---|---|---|---|---|
| TIO₂ (1000 cycles) | TALD | PEALD | TALD | PEALD | TALD | PEALD |
| as dep. | post-anneal | as dep. | post-anneal | as dep. | post-anneal | as dep. | post-anneal |
| 200°C | 450°C | 900°C | 200°C | 450°C | 900°C | 200°C | 450°C | 900°C |
| 44.0 nm | 49.6 nm | 57.9 nm | 17.5 nm | 18.1 nm | 33.8 nm |
| ZrO₂ (500 cycles) | TALD | PEALD | TALD | PEALD | TALD | PEALD |
| 250°C-as dep. | 600°C-anneal | 250°C-as dep. | 600°C-anneal | 250°C-as dep. | 600°C-anneal |
| 38.1 nm | 34.8 nm | 40.1 nm | 37.5 nm |
| HfO₂ (300 cycles) | TALD | PEALD | TALD | PEALD |
| 250°C-as dep. | 600°C-anneal | 250°C-as dep. | 600°C-anneal |
| 37.9 nm | 35.4 nm | 35.8 nm | 32.2 nm |
corrosion rate and can be obtained by Tafel fit analysis. Additionally, polarization resistance (R_p) and porosity (P) are calculated and presented in Table 3, using Eqs 1 and 2.

The HfO_2 coatings lead to a marked decrease in corrosion current density (I_corr) from 1.09 × 10^{-7} A/cm² for bare 304 SS to ~10^{-10} A/cm² with three orders of drop. Accordingly, polarization resistance (R_p) increases by three orders of magnitude. The porosity (P) represents the nominal uncoated surface fraction of the 304 SS substrate based on the ratio of polarization resistance without and with HfO_2 coatings. The smaller the porosity (P) is, the better the corrosion resistance is. Among all samples, PEALD-HfO_2 coated 304 SS with H_2 plasma pretreatments exhibits the most encouraging anti-corrosion property with most negative potential of −0.48 V, lowest corrosion current density of 5.25 × 10^{-11} A/cm², largest polarization resistance of 3.43 × 10^8 Ω · cm² and smallest porosity of 0.01. This can be ascribed to the combination role of the H_2 plasma pre-treatment and the high O_2 plasma activity during PEALD. The removal of organic contamination and improvement of adhesion to the 304 SS caused by the H_2 plasma pre-treatment are beneficial to anti-corrosion property.

Electrochemical impedance spectra (Bode plots) of different samples are presented in Fig. 11(b). The HfO_2 coating properties are evaluated from low frequency to high frequency. Compared to the bare 304 SS, all HfO_2 coated 304 SS samples show a clear increase in the global impedance. Moreover, PEALD HfO_2 coated 304 SS samples maintain a higher phase angle in a wide frequency range from 10^{-2} Hz to 10^5 Hz, giving better corrosion protection. Above all, the electrochemical results obtained from the LSV and the EIS confirm that PEALD HfO_2 coating onto H_2 plasma pretreated 304 SS has optimal corrosion-endurance.
Conclusion

TiO2, ZrO2 and HfO2 thin films were deposited on p-type Si (100) with nature oxide layer by the technique of TALD and PEALD. A variety of chemical liquid media including 1 mol/L H2SO4, 1 mol/L HCl, 1 mol/L KOH, 1 mol/L KCl, and 18 MΩ deionized water were used to test and compare chemical stability of all these as-deposited group IV metal oxides thin films, as well as post-annealed samples. In acidic solutions, TALD/PEALD HfO2 films, whether annealed or not, exhibit the best chemical stability without any change in thickness after long time immersion. Except as-deposited TALD-ZrO2 films with slowly etch rate of 1.06 nm/day in 1 mol/L HCl, all other ZrO2 films show better anti-acid stability, indicating the role of introduction of plasma O2 in PEALD and post-thermal treatment. As-deposited TiO2 films are relatively unstable in acidic solutions of H2SO4 and HCl, where the etch rate is 1.39 nm/day and 0.82 nm/day for TALD ones, and 0.81 nm/day and 0.48 nm/day for PEALD ones, respectively. The etch rate of PEALD samples in acid solutions drops by about 41%, compared to TALD ones. After post-annealing, TiO2 films have good corrosion endurance, which is ascribed to the crystallization transition from amorphous to anatase main phase and the formation of 5% Si-doped TiO2 ultrathin layers on sample surfaces. In basic and neutral solutions, HfO2, ZrO2, and TiO2 all show excellent corrosion resistance property with negligible change in thickness. Simultaneously, compared with bare 304 SS, 304 SS with HfO2 coating shows enhanced anti-corrosion property, and 36 nm PEALD-HfO2 is found to have a lower corrosion rate than TALD-HfO2 ones, which is ascribed to the improved film quality when activated O2 plasma is used. The pre-treatment of plasma H2 to substrate of SS can reduce interfacial impurities and porosity of overlayers with increased corrosion resistance characteristics. Electrochemical analysis yields an exponential decay of the corrosion current density by four orders of magnitude. Above all, using PEALD technique, post-annealing process and plasma H2 pre-treatment can improve the chemical stability and corrosion resistance of group IV metal oxide coatings. As a result, HfO2, ZrO2, and TiO2 ultrathin films derived from TALD and PEALD allows various protective applications in several commonly-used chemical liquid environments.

Materials and Methods

Preparation of TiO2, ZrO2 and HfO2 films by TALD and PEALD. TiO2 films were deposited at 200 °C, ZrO2 and HfO2 films were deposited at 250 °C, in a commercial Picosun SUNALE™ R-200 advanced PEALD system, which can be operated under PEALD and TALD modes. P-type Si (100) substrates (1 ~ 10 Ω · cm) were ultrasonically cleaned by acetone, alcohol and deionized water in turn without removing native oxide. TiCl4, Zr[N(C2H5)CH3]4 (TEMAZ) and Hf[N(C2H5)CH3]4 (TEMAH) as metal precursors were used at room temperature, 150 °C and 155 °C, respectively. H2O and O2 plasma were adopted as oxygen source for deposition of metal.
oxide by TALD and PEALD, respectively, where remote plasma power and O2 gas flow rate were 2,500 W and 160 sccm. For TALD-derived metal oxide thin films, all precursors’ pulse time was 0.1 s, followed by a 4 s N2 purge step to remove extra precursors and by-products. For PEALD-derived samples, the pulse and purge time of metal precursor kept unchanged, but pulse and purge time of plasma O2 was extended to 13.5 s and 10 s, respectively.

All as-deposited samples were also annealed at high temperature to evaluate the effect of thermal treatment on chemical stability. Both TALD- and PEALD-TiO2 films were heated in tube furnace at 450 °C and 900 °C for 3 h under N2 atmosphere, while annealed ZrO2 and HfO2 films were made in O2 environment at 600 °C for 10 min by rapid thermal annealing (RTA) to promote crystallization.

As-deposited and post-annealed samples were immersed in various chemical liquid media for various time, including 1 mol/L H2SO4, 1 mol/L HCl, 1 mol/L KOH, 1 mol/L KCl, and 18 MΩ deionized water. All the samples were stored in acid/alkaline-resistant boxes at room temperature in dark while control samples were placed in air.

| Solution | TALD-TiO2 | PEALD-TiO2 |
|----------|-----------|------------|
|          | as dep.  | post-anneal| as dep. | post-anneal |
| H2SO4    | 1.39 nm/day | — | 0.81 nm/day | — |
| HCl      | 0.82 nm/day | — | 0.48 nm/day | — |

Table 2. Etch rates of TALD- and PEALD-TiO2 films in 1 mol/L H2SO4 and 1 mol/L HCl.

Figure 7. GIXRD patterns of as-deposited and annealed TALD TiO2, ZrO2 and HfO2 ultrathin films.

Figure 8. The thickness variation vs. immersion time for all as-deposited and post-annealed TALD/PEALD TiO2 films in various chemical liquid media: (a) 1 mol/L H2SO4, (b) 1 mol/L HCl, (c) 1 mol/L KOH, (d) 1 mol/L KCl and (e) 18 MΩ water.

![Diagram of GIXRD patterns](image1)

![Diagram of thickness variation vs. immersion time](image2)
Characterization. The physical thickness of films (before and after immersion) was estimated by spectroscopic ellipsometry (Sopra GES-5) at an incidence angle of 75° and with wavelengths between 300 nm to 800 nm at increments of 10 nm. The data were then fit using the Cauchy (ε) model for TiO$_2$, ZrO$_2$, and HfO$_2$ films to acquire thickness values. A fixed native SiO$_2$ layer (1–2 nm) was considered in the model at the Si-metal oxide interface using the software of WinElli. X-ray photoelectron spectroscopy (XPS, Thermo Fisher K-Alpha, USA) was explored to characterize the chemical state and component of samples, using a monochromatic Al Kα source ($hν$ = 1,486.6 eV) for excitation of photoelectrons and binding energy scale was calibrated using C 1 s peak at 284.6 eV. Fourier transform infrared spectroscopy (FTIR) was used to obtain the chemical group information of the thin films using a pristine Si substrate as a reference. Crystallinity and phase structures of the films were analyzed by grazing incidence X-ray diffraction (GIXRD, Shanghai Synchrotron Radiation Facility, BL14B1) system with the light source energy of 10 k eV and grazing incidence of 0.12° due to the extremely thin thickness of the films. The surface topography was recorded by atomic force microscopy (AFM, Cypher, USA) and field emission scanning electron microscopy (FESEM, Zeiss Ultra 55, German). The cross-section FESEM image was also recorded to corroborate the thickness of samples.

Electrochemical measurements were performed by CHI660E electrochemical workstation and three-electrode tests where bare or coated 304 SS was used as the working electrode, a platinum wire as the counter electrode, saturated Ag/AgCl solution as a reference, and 1 mol/L KCl as the electrolyte and corrosion solution.

Table 3. Parameters related to the polarization curves in Fig. 11(a) for untreated bare 304 SS and TALD- and PEALD-HfO$_2$ coated 304 SS with and without H$_2$ plasma pretreatments.

|                        | H$_2$ plasma pre-treatment | $E_{corr}$ (V) | $I_{corr}$ (A·cm$^{-2}$) | $R_p$ (Ω·cm$^2$) | $P$ (%) |
|------------------------|----------------------------|----------------|--------------------------|------------------|---------|
| Bare 304 SS            | No                         | -0.17          | 1.09E-07                 | 3.49E+05         | 100     |
| TALD-HfO$_2$           | No                         | -0.18          | 3.92E-10                 | 2.77E+07         | 0.95    |
|                         | Yes                        | -0.24          | 3.55E-10                 | 4.73E+07         | 0.24    |
| PEALD-HfO$_2$          | No                         | -0.26          | 1.78E-10                 | 9.29E+07         | 0.15    |
|                         | Yes                        | -0.48          | 5.25E-11                 | 3.43E+08         | 0.01    |

Figure 9. The thickness variation vs. time for all 250°C as-deposited, 600°C-annealed TALD/PEALD-ZrO$_2$ films in various chemical liquid media: (a) 1 mol/L H$_2$SO$_4$, (b) 1 mol/L HCl, (c) 1 mol/L KOH and (d) 18 MΩ water.
rubber was used to encapsulate the surface of the sample. The reverse side of the sample was sealed after the extraction of the Al electrode ears equipped with the electrochemical workstation. Only a square area of about 1 × 1 cm² was left on the front for testing. Figure 12 shows the experimental setup for electrochemical measurements including electrochemical workstation, three-electrode test and sample package picture.

During measurement, the open circuit potential (OCP) was detected for one hour to attain a steady potential, then the electrochemical impedance spectroscopy (EIS) measurements were conducted in a frequency range of 10⁻² to 10⁵ Hz at OCP. The linear sweep voltammetry (LSV) data were collected at a rate of 20 mV/min and the corrosion potentials, corrosion currents, polarization resistances and porosities were calculated based on the polarization curves using the Stern-Geary equation. 

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![Figure 10](image1.png)

**Figure 10.** The thickness variation vs. time for all 250 °C as-deposited, 600 °C-annealed TALD/PEALD-HfO₂ films in various chemical liquid media: (a) 1 mol/L H₂SO₄, (b) 1 mol/L HCl, (c) 1 mol/L KOH and (d) 18 MΩ water.

![Figure 11](image2.png)

**Figure 11.** (a) Polarization curves and (b) Bode plots of untreated bare 304 SS and TALD- and PEALD-HfO₂ coated 304 SS with and without H₂ plasma pretreatments.
where $b_a$ and $b_c$ are the slopes of the anodic and cathodic branches of the Tafel plot, respectively. The electrochemical porosity of the films deposited onto SS was obtained by comparing $R_p$ of coated and bare SS:\(^{29}\)

$$P = \frac{R_{p\text{–bare}}}{R_{p\text{–ALD}}} \times 100\%$$  

(2)

References

1. George, S. M. Atomic layer deposition: an overview. Chem. Rev. 110, 111–131 (2010).
2. Lim, J. W., Yun, S. J. & Lee, J. H. Characteristics of TiO$_2$ films prepared by ALD with and without plasma. Electrochim. Solid-State Lett. 7, F73–F76 (2004).
3. Aarik, J. et al. Phase transformations in hafnium dioxide thin films grown by atomic layer deposition at high temperatures. Appl. Surf. Sci. 173, 15–21 (2001).
4. Niinistö, J., Kukli, K., Heikkilä, M., Ritala, M. & Leskelä, M. Atomic layer deposition of high-k oxides of the group 4 metals for memory applications. Adv. Eng. Mater. 11, 223–234 (2009).
5. Zhang, W. et al. Bipolar Resistive Switching Characteristics of HfO$_2$/TiO$_2$/HfO$_2$ Trilayer-Structure RRAM Devices on Pt and TiN-Coated Substrates Fabricated by Atomic Layer Deposition. Nanoscale Res. Lett. 12, 393–1–11 (2017).
6. Cao, Y. Q. et al. Improved thermal stability and electrical properties of atomic layer deposited HfO$_2$/AlN high-k gate dielectric stacks on GaAs. J. Vac. Sci. Technol. A 33, 01A136–1–6 (2015).
7. Puurunen, R. L. Surface chemistry of atomic layer deposition: A case study for the trimethylaluminium/water process. J. Appl. Phys. 97, 121301 (2015).
8. Miikkulainen, V., Leskela, M., Ritola, M. & Puurunen, R. L. Crystallinity of inorganic films grown by atomic layer deposition: Overview and general trends. J. Appl. Phys. 113, 021301 (2013).
9. Profijt, H. B., Potts, S. E., De Sanden, M. C. & Kessels, W. M. Plasma-assisted atomic layer deposition: basics, opportunities, and challenges. J. Vac. Sci. Technol. A 29, 050801 (2011).
10. Kim, H. Characteristics and applications of plasma enhanced-atomic layer deposition. Thin Solid Films 519, 6639–6644 (2011).
11. Potts, S. E. et al. Ultra-thin aluminium oxide films deposited by plasma-enhanced atomic layer deposition for corrosion protection. J. Electrochem. Soc. 158, C132–C138 (2011).
12. Potts, S. E. et al. Low temperature plasma-enhanced atomic layer deposition of metal oxide thin films. J. Electrochem. Soc. 157, P66–P74 (2010).
13. Matero, R. et al. Atomic layer deposited thin films for corrosion protection. J. Phys. IV 9, 493–499 (1999).
14. Hu, S. et al. Amorphous TiO$_2$ coatings stabilize Si, GaAs, and GaP p–n junctions and GaAs p–n photoanodes for efficient water oxidation. Science 344, 1005–1009 (2014).
15. Marin, E., Lanzutti, A., Guzman, L. & Fedrizzi, L. Corrosion protection of AISI 316 stainless steel by ALD alumina/titania nanometric coatings. J. Coat. Technol. Res. 8, 655–659 (2011).
16. Shan, C. X., Hou, X. H. & Choy, K. L. Corrosion resistance of TiO$_2$ films grown on stainless steel by atomic layer deposition. Surf. Coat. Technol. 202, 2399–2402 (2008).
17. Mihashemihaghighi, S. et al. Corrosion protection of Aluminium by Ultra-thin Atomic Layer Deposited Alumina Coatings. Corros. Sci. 106, 16–24 (2016).
18. Abdulagatov, A. I. et al. Al$_2$O$_3$ and TiO$_2$ Atomic Layer Deposition on Copper for Water Corrosion Resistance. ACS Appl. Mat. Interfaces 3, 4593–4601 (2011).
19. Strandwitz, N. C. et al. Photoelectrochemical behavior of n-type Si (100) electrodes coated with thin films of manganese oxide grown by atomic layer deposition. J. Phys. Chem. C 117, 4931–4936 (2013).
20. Correa, G. C., Bao, B. & Strandwitz, N. C. Chemical stability of titania and alumina thin films formed by atomic layer deposition. ACS Appl. Mat. Interfaces 7, 14816–14821 (2015).
21. Kim, L. H. et al. Al$_2$O$_3$/TiO$_2$ nanolaminate thin film encapsulation for organic thin film transistors via plasma-enhanced atomic layer deposition. ACS Appl. Mat. Interfaces 6, 6731–6738 (2014).
22. Chen, Y. W. et al. Atomic layer-deposited tunnel oxide stabilizes silicon photoanodes for water oxidation. Nat. mater. 10, 539–544 (2011).
23. Klein, T. M. et al. Evidence of aluminum silicate formation during chemical vapor deposition of amorphous Al$_2$O$_3$ thin films on Si(100). Appl. Phys. Lett. 75, 4001 (1999).
24. Zhang, W. Q., Huang, L. Y., Li, A. D. & Wu, D. Hf$_x$Zr$_{1-x}$O$_2$ films chemical vapor deposited from a single source precursor of anhydrous Hf$_x$Zr$_{1-x}$(NO$_3$)$_3$. J. Cryst. Growth 346, 0–16 (2012).
25. Wilk, G. D., Wallace, R. M. & Anthony, J. M. Hafnium and zirconium silicates for advanced gate dielectrics. J. Appl. Phys. 87, 484–492 (2000).
26. Gilmer, D. C. et al. Low Temperature CVD of Crystalline Titanium Dioxide Films Using Tetranitratotitanium(IV). Chem. Vap. Deposition 4, 9–11 (1998).
27. Lim, G. T. & Kim, D. Characteristics of TiO2 films prepared by chemical vapor deposition using tetrakis-dimethyl-amido-titanium and water. Thin Solid Films 498, 254–258 (2006).
28. Yun, S. J., Lim, J. W. & Lee, J. H. Effect of plasma on characteristics of zirconium oxide films deposited by plasma-enhanced atomic layer deposition. Electrochem. Solid-State Lett. 8, F47–F50 (2005).
29. Harkonen, E. et al. Hydrogen–argon plasma pre-treatment for improving the anti-corrosion properties of thin Al2O3 films deposited using atomic layer deposition on steel. Thin Solid Films 534, 384–393 (2013).
30. Daubert, J. S. et al. Corrosion protection of copper using Al2O3, TiO2, ZnO, HfO2, and ZrO2 atomic layer deposition. ACS Appl. Mater. Interfaces 9, 4192–4201 (2017).
31. Díaz, B. et al. Low-temperature atomic layer deposition of Al2O3 thin coatings for corrosion protection of steel: Surface and electrochemical analysis. Corros. Sci. 53, 2168–2175 (2011).

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Author Contributions
M.L. and Z.X.J. carried out the sample fabrication and characterization of SEM and XRD, W.Z. and Y.H.B. conducted the XPS, AFM and FTIR tests. M.L. and Y.Q.C. conducted the chemical stability test and electrochemical measurement. W.M.L., D.W. and A.D.L. participated in the discussion of results. M.L. drafted the manuscript. A.D.L. supervised the whole work and revised the manuscript. All authors critically read and commented on the manuscript.

Additional Information
Competing Interests: The authors declare no competing interests.

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