



Full length article

Layered Al₂O₃-SiO₂ and Al₂O₃-Ta₂O₅ thin-film composites for high dielectric strength, deposited by pulsed direct current and radio frequency magnetron sputteringBenjamin V.T. Hanby^{a,*}, Bryan W. Stuart^a, Miquel Gimeno-Fabra^a, Jonathan Moffat^b, Chris Gerada^c, David M. Grant^{a,*}^a Advanced Materials Research Group, Faculty of Engineering, University of Nottingham, NG7 2RD, UK^b Oxford Instruments, Asylum Research, High Wycombe HP12 3SE, UK^c Department of Electrical and Electronic Engineering, University of Nottingham, NG7 2RD, UK

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ABSTRACT

Multilayer thin films have the potential to act as high dielectric strength insulation for wire and microelectronics. In this study, films consisting of 2, 4 or 8 layers, composed of Al₂O₃ with SiO₂ or Ta₂O₅, were prepared *via* pulsed direct current and radio frequency magnetron sputtering to a thickness of between 152 and 236 nm. The dielectric strengths of all films exceeded the 310 Vμm⁻¹ achieved for PDC Al₂O₃. Maximum dielectric strengths were obtained for four layer composites; Al₂O₃-SiO₂-Al₂O₃-SiO₂ (466 Vμm⁻¹) and Al₂O₃-Ta₂O₅-Al₂O₃-Ta₂O₅ (513 Vμm⁻¹), each containing two PDC-Al₂O₃ and two RF-SiO₂/Ta₂O₅ layers. Whilst the average dielectric strength was higher in the Ta₂O₅ composites, they suffered from higher leakage prior to breakdown with *ca.* 6.5 nA compared to *ca.* 0.1 nA for SiO₂ composites. The mechanical properties of the composites were poorer due to increased intrinsic coating stress. Samples exhibited complete interfacial delamination with maximum coating adhesion strengths of 22 and 25 MPa. The variance resulted from larger coefficient of thermal expansion for Ta₂O₅ compared to SiO₂. Sputtered composites of Al₂O₃ and either SiO₂ or Ta₂O₅ had high breakdown strength with reasonable adhesion and could be suitable for insulating copper conductors in the aerospace and automotive industries.

1. Introduction

Highly insulating ceramic films are of great interest as dielectric layers in microelectronics for thin-film capacitors and field effect transistors [1,2]. Furthermore, thin high-strength dielectric films offer a route to improve packing densities in wire insulation and are desirable in aviation and automotive applications, where a drive for electrification is pushing for the improvement of all aspects within electrical systems. As such, lightweight, thermally stable and high breakdown strength insulation is required for windings in motors and actuators [3]. An airbus A3800–800 contains *ca.* 470 km of wire with a total weight of 5700 kg. The electrical insulation commonly consists of PTFE/polyimide/PTFE composites, because of its arc track resistance, compared to polyimide alone and maximum operating temperature of *ca.* 260 °C [4,5]. With the further evolution of more electrical aircraft the amount of wire used is bound to increase, whereby improvements in dielectric strength/ thermal stability of the insulation could be used to reduce the

weight and space taken up by the total wire [3].

Physical Vapour Deposition (PVD) techniques such as magnetron sputtering have been shown to be capable of producing alumina films with dielectric strength as high as 1.5 kVμm⁻¹, such as those produced by *CARRIER* et al. when using special power supplies to pulse the Direct Current (PDC) power and quench arc events [6–8]. Further to these desirable electrical properties PDC alumina films are considered a suitable base material for electrically insulating multilayer composites as they have been shown to adhere well to copper, exhibiting pull off adhesion values of up to 56 MPa and have deposition rates up to 76 nmh⁻¹ with sample rotation [8]. Sputtering is also attractive because the thin films it deposits can be tailored to prevent cracking and delamination, namely by monitoring thickness and mitigating thermal expansion variation and high internal stress [9,10].

Multilayer materials have been used as a route to combine desirable material properties, such as in ceramic capacitors where multilayers improve electrical properties including dielectric constant and

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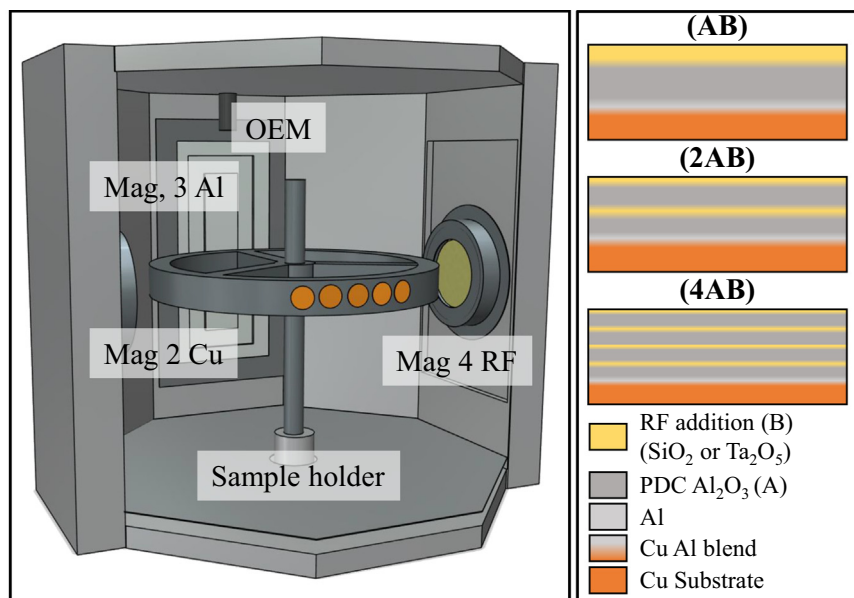


Fig. 1. TEER UDP-650 front on configuration with targets in Mag 2 (RF Cu), Mag 3 (PDC Al) and Mag 4 (RF SiO₂ or RF Ta₂O₅). The OEM is located above mag 3 and is used to control plasma intensity of the Al target used to prepare multilayer PDC alumina materials with a blended copper aluminium interlayer. On the right is a summary of the nomenclature used to refer to the Al₂O₃-Ta₂O₅ and Al₂O₃-SiO₂ composites produced in this study (where A refers to PDC alumina and B refers to the RF addition SiO₂ or Ta₂O₅).

dielectric strength, by introducing barriers to current flow and defect filling [11–13]. It is important to consider barrier layer thickness, as shown by *Chu and Shen* where increasing the number of layers from 174 to 800 in TiN/TiBN films was shown to result in a 50% reduction in hardness and a 79% reduction in adhesive scratch failure [14]. Film doping is another potential route to improving the dielectric strength of sputtered ceramics; for instance a 10 mol% lanthanum inclusion was shown to increase the dielectric strength of sol-gel Al₂O₃ films from 310 to 393 V μ m⁻¹ and reduce leakage current three fold, as shown by *Zou et al.* [15]. However, multilayers are more appropriate because of the higher level of consistency attainable, because of issues with complex preferential sputtering and target erosion during sputtering from compound and composite targets when doping [16]. Preferential sputtering is caused by the varying sputtering yields of elements in a compound target, which results in a coating composition which differs from the target [17].

SiO₂ is an attractive option for use in multilayer materials and it has been sputtered using various power supply options from Si or SiO₂ targets. Films deposited using reactive radio frequency (RF) sputtering by *Jun et al.* achieved good dielectric strengths up to 570 V μ m⁻¹, higher than that of sputtered alumina films produced in previous work [18,19]. However, these RF sputtered films suffer from low deposition rate, for example ca. 300 nmh⁻¹ without sample rotation, making them impractical from a fabrication standpoint. Multilayer films consisting of sputtered Al₂O₃ and SiO₂ produced by *Martínez-Perdiguero et al.* showed a two fold increase in dielectric strength when comparing the electrical properties to those of Al₂O₃ alone [20]. Also displaying a maximum dielectric strength of > 160 V μ m⁻¹ for an annealed 8 layer structure. Improvements in this case were attributed to structure stabilisation. Al₂O₃-SiO₂ multilayer materials have also been studied for gas diffusion barriers as produced by atomic layer deposition and sputtering [13,21]. These multilayers have found such application because of subsequent layers ability to close pinhole defects. In the case of electrical insulating films this effect would also be beneficial.

Ta₂O₅ is another ceramic which has often been utilised in the electronics industry, as a material with a very high dielectric constant of up to 23, therefore, it is a highly attractive material for thin film capacitors among other applications [22]. Ta₂O₅-thin films have been frequently deposited using various techniques as discussed by *Chaneliere et al.* [23]. Deposition of films using RF sputtering has been able to produce materials with high dielectric strength, such as 400 V μ m⁻¹ achieved by *Oehlein et al.* and 600 V μ m⁻¹ for ca. 60 nm films deposited

by *Seki et al.* [24,25]. Ta₂O₅ Al₂O₃ nano laminates have been produced using ALD by *Smith et al.* [26]. These ALD composites were produced as films for capacitors and to have a high dielectric constant between 8.9 and 23.9 for pure Al₂O₃ and Ta₂O₅ respectively. Their work showed that the layered films exhibited lower leakage current values than Al₂O₃ alone over certain applied voltage ranges.

Dielectric properties have been reported for layered magnetron sputtered Al₂O₃-SiO₂ composites. However, reports are lacking for layered magnetron sputtered Al₂O₃-Ta₂O₅ composites and further elucidation is required to fully understand the mechanisms by which incorporating multilayers improves the dielectric strength in both material systems [20,26]. A more substantial assessment of the effect of the number of layers and the layer thickness would also aid in the future design of high performance ceramic multilayers to increase the dielectric strength and adhesion strength. Further to this it is noted that AFM breakdown methods have not been used to measure the properties of composite materials with such thickness, which is usually limited to ultra-thin films and it is important to show the methods applicability [27–29].

The aim of this study is to utilise the potential of multi-layered ceramic films, based upon pulsed direct current sputtered Al₂O₃ with either RF sputtered Ta₂O₅ or SiO₂ to improve the dielectric strength of the single layer Al₂O₃ films as previously deposited [8]. The combination of PDC and RF sputtering is proposed in order to utilise the high quality of RF films and the higher deposition rates of PDC deposition whilst also facilitating blending to prevent cohesive failure and limiting the requirements for multiple optical emission monitoring (OEM) systems. All of this is carried out in order to produce the next generation of thermally stable, high breakdown strength insulators, which maintain adhesive properties to copper substrates.

2. Methodology

2.1. Coating and sample preparation

Cu disk substrates of 5 and 10 mm diameter with a thickness of 2 mm were punched from Cu sheet (Cu-CW004A-H065 of 99.9% purity) and were polished using sequentially higher grit silicon carbide grinding and polishing papers (240 to 4000 Grit) with water as a lubricant. Followed by colloidal silica polishing (particle size of 0.06 μ m) using an MD-Chem polishing pad (Struers®). Samples were then ultrasonicated for 10 min each in acetone and ethanol. Coatings were

Table 1

Alumina coating parameters for the multilayer composite films. Note the time used for RF stages (Ta_2O_5 or SiO_2) excluded the 15 min ramp stages prior and post layer formation. The initial PDC layer was followed with either RF Ta_2O_5 or RF SiO_2 depending on the type of composite being produced.

Layer config.	PDC Al_2O_3 per layer (min)	RF Ta_2O_5 per layer (min)	RF SiO_2 per layer (min)	RF target power (W)	Ar Gas Flow (sccm)	Pulse Freq. (kHz)	Duty Cycle (%)
AB	133.3	30.9	155.4	100	140	150	40
2AB	66.7	15.0	77.2	100	140	150	40
4AB	33.3	7.2	38.6	100	140	150	40

deposited in a TEER UDP-650 magnetron sputtering rig (see Fig. 1 for sputtering rig layout and magnetron placement). Al targets were pre-cleaned until the optical emissions monitoring (OEM) plasma intensity returned to that of the un-poisoned target. Optical emission monitoring and optimisation has been discussed previously and is used to control the flow of reactive gas, in order to deposit stoichiometric materials [8]. The optical emissions monitoring system monitors the intensity of the plasma and is calibrated when sputtering in 100% argon, the reactive gas is then added until the intensity is decreased to a desired set point. An OEM set point of 23% was used to deposit Al_2O_3 from an Al target in this study. The Cu target (mag 2 – Fig. 1) was pre-cleaned for 10 min at 60 W. RF targets (mag 4 - Fig. 1) were pre-cleaned for a total of 15 min with a 5 min power ramp up stage, so as not to damage the target.

Films were sputtered to a total thickness of ca. 200 nm with a varying number of layers. Alternate layers of pulsed DC Al_2O_3 (allocated A) and RF SiO_2 or RF Ta_2O_5 (the added RF layer is allocated B) were deposited so as to form AB, ABAB (2AB) and ABABABAB (4AB) composites as seen in Fig. 1 using deposition parameters as shown in Table 1.

Al_2O_3 layers were sputtered using pulsed direct current sputtering according to films as produced by in a previous study, with a blended Cu/Al interlayer [8]. The Al interlayer was deposited using PDC power at 4 A as opposed to 6 A DC as used in the previous study. This was primarily done to achieve stable arc free deposition. Sputtering times were used as in Table 1 in order to deposit a film of ca. 200 nm total thickness, containing ca. 150 nm of pulsed DC Al_2O_3 . Note that RF magnetrons were allowed to ramp for 15 min prior to the time given in Table 1 and also allowed 15 min to ramp to 0 following each layer, creating the blended interfaces.

2.2. Characterisation

2.2.1. Fourier transform infrared spectroscopy

Fourier Transform Infrared Spectroscopy (FTIR) was carried out using a Bruker Tensor FTIR instrument. An Attenuated Total Reflectance (ATR) attachment with a diamond mirror and ZnSe lens was used to measure sample IR absorbance between 500 and 1500 cm^{-1} wavenumbers. A resolution of 4 cm^{-1} was used.

2.2.2. XRD

X-ray Diffraction (XRD) was carried out using a Bruker D8 Advanced diffractometer. A glancing angle of 1.2° was applied with a step size of 0.04° and a counting time of 20 s, scanning between 2θ values of 15 and 80° . Diffraction patterns were analysed using Bruker's EVA software.

2.2.3. XPS

X-ray photoelectron spectroscopy (XPS) was used to determine the elemental composition of the top surface (ca. 5 nm) of the material [30]. Samples were analysed in an ESCA lab mark II with an Al K α non-monochromatic X-ray source at $< 10^{-7}$ Torr. A survey spectra was collected from two scans between 0 and 1200 eV with a step size of 1.0 eV and a dwell time of 0.2 s. Five high resolution scans of O_{1S}, C_{1S} and either Ta_{4F} or Si_{2P} were averaged with a step size of 0.2 eV and a dwell time of 0.4 s. Charge shift correction was applied to all spectra

through calibration with the C_{1S} peak using the ideal value of 284.8 eV.

2.2.4. SEM

Coated copper samples were mounted in resin perpendicular to the normal to obtain polished cross section as in Section 2.1. Samples were ultra-sonicated and iridium coated (ca. 8 nm) using a Q150R Plus coater. Samples thickness and structure were determined using A JEOL 7100F Field Emission Gun Scanning Electron Microscope (FEG-SEM) operating at 15 kV, with a working distance of 10 mm.

2.3. Mechanical testing

2.3.1. Pull off testing

In accordance with ASTM-D4541–09 DFD E1100S epoxy was applied to a cleaned 2.8 mm diameter stub and pressed onto a 10 mm diameter coated Cu disk. The stub was pressed down to remove any air and then cured on a heating plate for 1 h at 140°C . Once cured any excess glue from around the stub was removed with the supplied cylindrical cutting tool. The samples were tested with a DFD instruments P.A.T. handy pull off adhesion unit and adhesion strength in MPa was recorded. A Nikon LV100ND light microscope was then used to determine the failure mechanism. Eight measurements were carried out for each sample type. A FEI Quanta 650 microscope equipped with an Oxford Instruments X-Max -150 EDX Detector was used for elemental mapping of the pull off stubs.

2.4. Electrical testing

2.4.1. AFM dielectric breakdown measurements

Atomic Force Microscopy (AFM) was utilised to measure the dielectric breakdown of the films as deposited onto copper disks. This was carried using an Oxford Instruments Cypher S equipped with a high voltage module and an Asylum Research ASYELEC.01 Ti/Ir coated tip [31]. Measurements were carried out by generating a $30 \times 30\ \mu\text{m}$ topographic image (in tapping mode) and then a grid of 4×4 locations was used on top of the image. At each location a voltage ramp (0–150 V at a rate of 37.5 Vs^{-1}) was applied to the copper substrate until a sharp increase in current was detected through the probe.

3. Results

3.1. Structural characterisation

FTIR spectra for RF sputtered SiO_2 shows peaks resulting from Si–O bonds at 1151 and 807 cm^{-1} . RF Ta_2O_5 spectra displayed a broad absorbance band with peaks at 745 and 890 cm^{-1} . Multilayer composites contained a spectra resulting from the superimposition of these peaks with an Al–O LO phonon vibration peak at ca. 900 cm^{-1} . The SiO_2 composites display an additional peak at 1229 cm^{-1} . All peaks for RF films and composites have been identified in Fig. 2a. Diffraction patterns (see Fig. 2b) showed the X-ray amorphous nature of all of the films, suggested by broad diffraction associated with an amorphous hump, which resulted from contributions from both the film and amorphous glass substrate. In the case of RF Ta_2O_5 , two broad

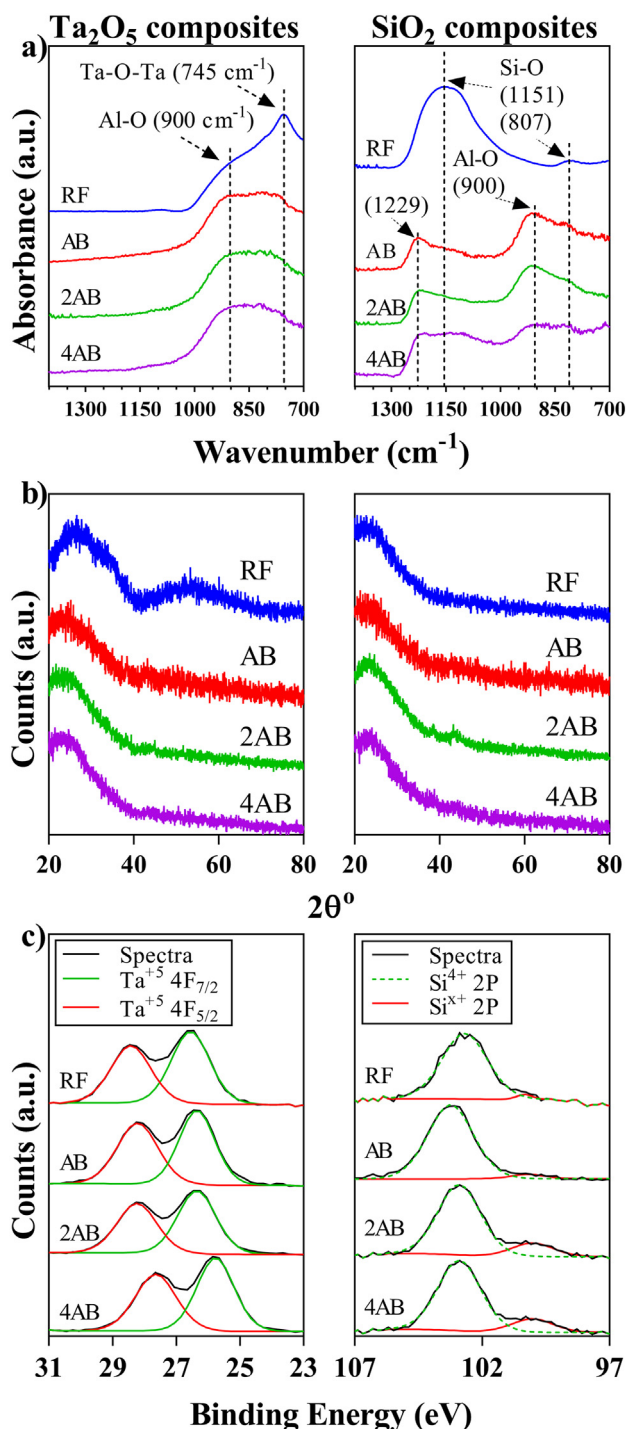


Fig. 2. a) FTIR spectra for Ta₂O₅ and SiO₂ composites as deposited onto copper foil. The main absorbance bands and their assignments have been indicated. b) shows XRD diffraction patterns for Ta₂O₅ and SiO₂ composites as deposited onto glass coverslips, indicating the films amorphous nature. c) XPS peak deconvolution for Ta₂O₅ and SiO₂ composites (as deposited onto Cu disks, identity of the peaks fitted are indicated).

amorphous humps can be seen which correlate to amorphous Ta₂O₅.

XPS spectra gave the stoichiometry and binding energies of the top 5 nm layer of the films. Binding energies of 26.3 ± 0.1 eV were observed for Ta_{4F} photoelectrons, whereas binding energies of 103.2 ± 0.1 eV were observed for Si_{2P} photoelectrons with a shoulder at 100.5 ± 0.1 eV, see Fig. 2c and Table 2. The amount of non-stoichiometric silicon calculated from the Si peak was seen to

Table 2

XPS Ta_{4F} and Si_{2P} binding energies, elemental ratios and oxidation states for composite films as deposited onto Cu disks.

Layer config.		Ta ⁵⁺ _{4F} (eV)	Ta ⁰ _{4F} (eV)	Ta ⁰ %	Ta:O ratio
Ta ₂ O ₅	RF	26.3	N/A	N/A	0.39
	AB	26.2	N/A	N/A	0.35
	2AB	26.3	N/A	N/A	0.39
	4AB	26.2	N/A	N/A	0.40
Layer config.		Si ⁴⁺ _{2P} (eV)	Si ¹⁺ _{2P} (eV)	Si ¹⁺ %	Si:O ratio
SiO ₂	RF	103.2	100.8	3.5	0.43
	AB	103.2	100.2	1.9	0.62
	2AB	103.2	100.2	4.4	0.58
	4AB	103.2	100.3	16.1	0.56

increase with an increasing number of layers. Yielding a transition from 1.9 to 16.1% between AB and 4AB composites. The Si:O ratios for the films varied between 0.43 and 0.62 where the ideal Si:O is 0.5. No peaks were found in the Ta₂O₅ composite spectra resulting from Ta⁰ and the Ta:O ratios were generally consistent between, 0.35 and 0.4 where the ideal ratio is 0.4.

Deconvolution of the O_{1S} peak for the SiO₂ composites gave information as to the binding energy and composition of the Si species, see Table 3 for the oxygen binding energies and percentages determined by deconvolution.

Cross-sectional SEM imaging was used to show film thickness for use in the dielectric breakdown strength calculations. Layers could not be distinguished in most cases due to limiting resolution of the instrument and similar appearance, however, using backscattered imaging for Ta₂O₅ composites some layer structure could be seen in AB and 2AB films (the light areas can be seen resulting from the high molecular weight ca. 181 g mol⁻¹ Ta atoms) as seen along with sample thickness in Fig. 3 and Table 4. All films appeared to be predominately featureless. The layers in SiO₂ composite films were not clearly observed in backscattered or secondary electron mode (see Fig. 3).

3.2. Electrical testing

Dielectric breakdown measurements revealed a range of breakdowns for the multilayer materials. The highest dielectric strength was observed in both the case of Ta₂O₅ and SiO₂ composites where the 2AB motif was used, at 513 and 466 μm^{-1} respectively. The lowest dielectric strength was seen in AB films at 332 and 350 μm^{-1} for SiO₂ and Ta₂O₅ respectively.

The standard error in the voltage measurements was at least four times lower for Ta₂O₅ films as seen in Table 4, the error in AB films was also highest for each set of materials as seen in Table 4 and Fig. 4. The maximum leakage current prior to breakdown was determined using the current applied field plots and was shown to be up to ca. 60 times larger in the Ta₂O₅ composites, with maximum leakages of ca. 6.48, 3.13 and 2.35 nA (AB, 2AB, 4AB) whilst leakage prior to breakdown was < 0.1 nA in all Al₂O₃-SiO₂ composite films. Oscillations in the current leakage prior to breakdown were also seen in Ta₂O₅ composites but not in SiO₂ composites, see Fig. 4b.

Table 3

Binding energies and contribution of Si in the O_{1S} high resolution spectra from SiO₂ composites as deposited onto Cu disks.

Layer config.	Si ⁴⁺		Shoulder 1		Shoulder 2	
	(eV)	%	(eV)	%	(eV)	%
RF	532.6	68.8	534.6	29.1	531.6	2.1
AB	532.5	93.9	534.5	3.4	530.2	2.6
2AB	532.6	91.2	534.5	5.1	530.8	3.7
4AB	532.56	93.6	534.5	4.4	530.7	2.2

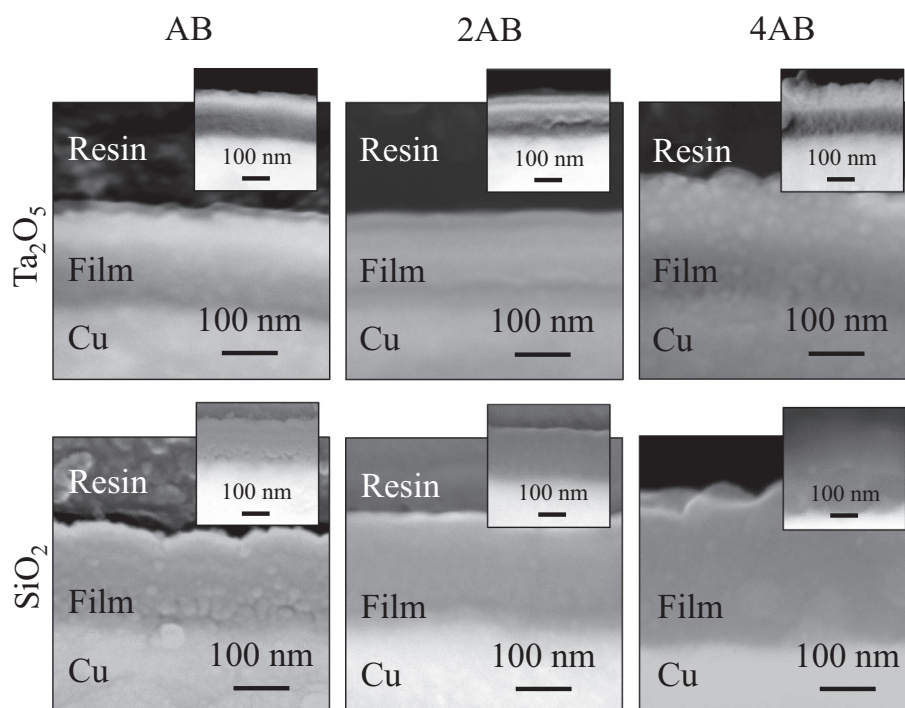


Fig. 3. Secondary and back scattered (inset) Cross sectional SEM of composite films Ta_2O_5 (top) and SiO_2 (bottom) as deposited onto Cu disks and mounted in resin. Average film thickness are displayed in Table 4.

Table 4

Breakdown voltage and dielectric breakdown for Ta_2O_5 and SiO_2 multilayer composites as deposited onto Cu disks. Layer structure and overall film thickness has been included (and was used to calculate the dielectric strength).

	Layer config.	Thickness (nm)	Breakdown AVG. (V)	Error (V)	N	Dielectric Strength ($\text{V}\mu\text{m}^{-1}$)
SiO_2	AB	191 ± 4	63.4	30.4	16	332
	2AB	182 ± 4	84.9	15.6	16	466
	4AB	236 ± 11	98.3	16.5	16	416
Ta_2O_5	AB	$203. \pm 6$	71.0	7.5	16	350
	2AB	152 ± 4	78.0	2.1	16	513
	4AB	210 ± 9	84.6	4.1	16	403

3.3. Mechanical testing

Pull off testing carried out on the multilayer composite films showed that the adhesive strength varied between 11.4 ± 1.8 and 34.3 ± 4.4 MPa with failure occurring through complete interfacial delamination and partial interfacial delamination between the coating and the substrate for all of the films. Average pull off values and failure mechanisms for each film are given in Table 5.

It can be seen that Ta_2O_5 composites produced with an AB layer configuration had a larger adhesion strength than the other Ta_2O_5 composites by ca. 1.5 times. Whilst the highest adhesion seen in SiO_2 composites was for the 4AB layer structure. EDX of the pull off stubs revealed a layer of copper on tested samples indicating that failure occurred at the initial Cu–Cu interface (see Fig. 5).

4. Discussion

The deposition of amorphous thin SiO_2 and Ta_2O_5 composites using RF power was shown to be possible with FTIR (showing bonding peaks for each component of the films) and XPS which confirmed the expected bonding configurations and desired compositional stoichiometry

for the surface layers of the composites (see, Fig. 2). These materials were combined with PDC Al_2O_3 , as deposited previously with a view to improving the dielectric strength above $310 \text{ V}\mu\text{m}^{-1}$ [8]. RF sputtered SiO_2 and Ta_2O_5 both have the potential to have high breakdown strength as reported in the literature (higher than that achieved for pulsed DC alumina alone 570 and $600 \text{ V}\mu\text{m}^{-1}$ respectively) making them ideal candidates for the multicomponent films [19,25]. The use of multi-layers is attractive because of the potential for suppression of charge transfer as explored by Mackey et al. for extruded polymer films [32]. The changes in mechanical and electrical properties will be discussed in the following section.

4.1. Structural

For the composite films multiple infrared absorption peaks associated with the composites constituent layers were seen, see Fig. 2. For Ta_2O_5 composites a broad absorbance peak resulted from Al–O phonon vibrations combined with Ta_2O_5 peaks at ca. 785 cm^{-1} . The Ta_2O_5 peak decreased in intensity from 0.97 to 0.92 a.u. with respect to the peak at ca. 820 cm^{-1} as the Ta_2O_5 RF surface layer decreased in thickness. The decreasing layer thickness resulting in a diminished absorption peak, Ta–O–Ta peaks were similarly observed by Mannequin et al. in electron beam evaporated and sputtered films [22].

The FTIR spectra of SiO_2 composites had overlapping Al–O and Si–O peaks, similar combinations of peaks - resulting from Al_2O_3 and SiO_2 vibrations and Al–O–Si linkages - were seen in heat treated Al_2O_3 - SiO_2 composite aerogels produced by Wu et al. [33]. Hence assignment of the peak at 1229 cm^{-1} was to either Al–O–Al bonding or Al–O–Si bonding. This could suggest Al–O–Si bonding at the Al_2O_3 , SiO_2 interfaces, which may lead to improved adhesion [34].

Discrete layers were too thin to be imaged using SEM techniques, as seen in Fig. 3. The XRD pattern for RF Ta_2O_5 was not replicated in multi-layer Ta_2O_5 films, this was because the sample used to measure the RF properties was deposited to 774 nm whilst the Ta_2O_5 layers in the composites had a total and maximum thickness of 50 nm which resulted in the single amorphous hump, due to a lack of instrument

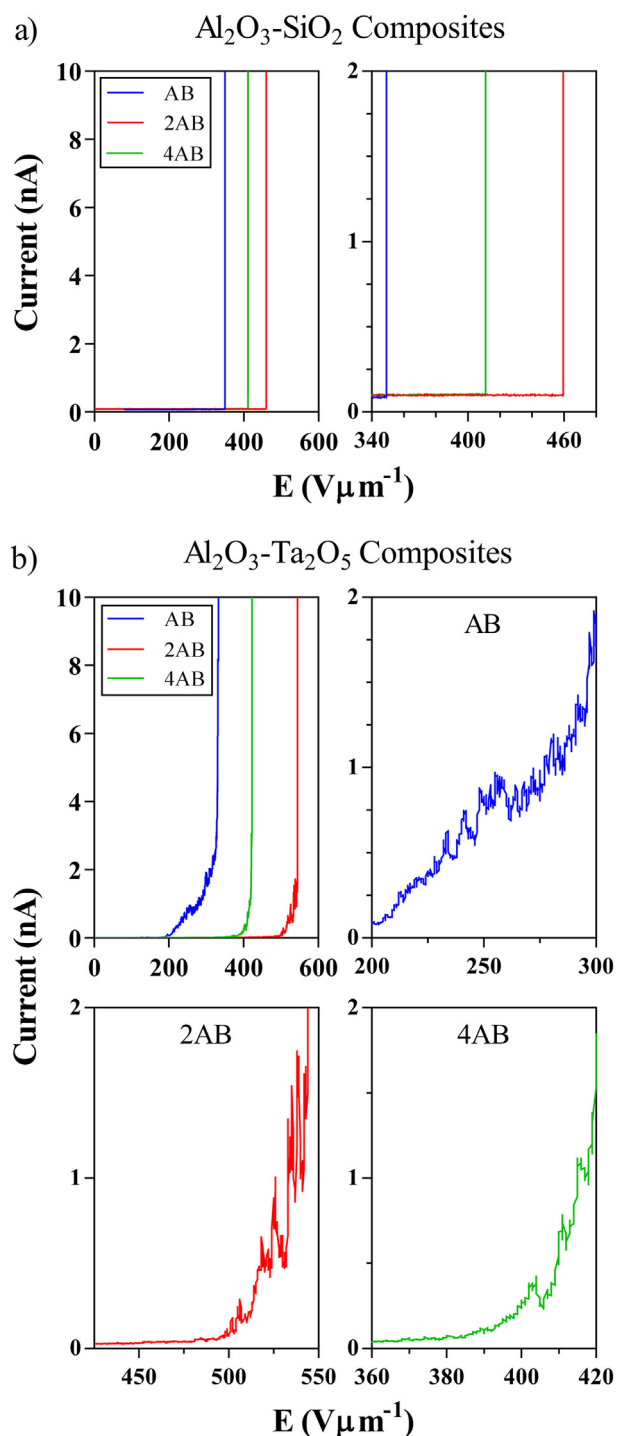


Fig. 4. AFM measurement of current, applied field plots indicating the onset of breakdown and current leakage prior to breakdown. a) Plots for SiO_2 composites, including a high resolution graph showing the low leakage current (< 0.1 nA) prior to breakdown. b) Plots for Ta_2O_5 composites which includes high resolution graphs showing current oscillation associated with charge trapping and de-trapping, prior to breakdown. For both materials the layer configuration is indicated in the key.

sensitivity.

All the RF layers used for the 4AB composites had a thickness in excess of the XPS penetration depth such that all the photo electrons originated from the surface of the top layers of Ta_2O_5 or SiO_2 . For the Al_2O_3 - SiO_2 composites XPS determined that the level of Si^{1+} increased in every instance where the number of layers was increased, hence the

Table 5

Average adhesive pull off strength (MPa) and failure mechanism for Ta_2O_5 and SiO_2 multilayer films as deposited onto Cu disks.

	Sample	Average failure strength (MPa)	Complete interfacial failure	Partial interfacial failure	Failure in Adhesive
Ta_2O_5	AB	24.6 ± 3.1	4	3	0
	2AB	16.0 ± 2.2	8	0	0
	4AB	15.3 ± 2.8	8	0	0
SiO_2	AB	22.0 ± 3.4	6	2	0
	2AB	11.4 ± 1.8	8	0	0
	4AB	34.3 ± 4.4	1	7	0

layers decreased in thickness. Deconvolution of the O_{1S} peaks (not shown) was used to determine the identity of the Si^{1+} species. This analysis showed that the Si^{1+} shoulders were made up of contributions from surface hydroxides at 534.5 ± 0.1 eV accounting for between 2.1 and 3.7% and SiO_{2-x} species at 530.6 ± 0.2 eV accounting for between 3.4 and 5.1%. The amount attributed to SiO_{2-x} species was lower than the 29.1% seen for RF films [35]. This improvement – with respect to the level of SiO_{2-x} in RF films – is attributed to target conditioning during reactive PDC Al_2O_3 sputtering as well as the decreased sputtering time [36]. These parameters combined to decrease the effect of preferential sputtering on the film composition. This, however, cannot account for the large increase in the Si^{1+} shoulder seen for 4AB films. The lack of Al_{2P} photo electrons in the survey spectra rules out the influence of compound material at the boundaries (such as Al-O-Si as suggested by FTIR) on the XPS results, thus the amount of Si^{1+} in the films. This is therefore likely a result of the increase in total sputtering time (when considering the multiple blending stages required for the 4AB films) resulting in preferential sputtering effects, which increased the amount of compositional defects.

Conversely, the Al_2O_3 - Ta_2O_5 films emitted only Ta^{5+} photoelectrons and had stoichiometry which varied between a Ta:O ratio of 0.35 and 0.4, as seen in Table 2. Thus it is likely that conditioning stages during PDC sputtering was sufficient to produce stoichiometric Ta_2O_5 films. Deconvolution of the O_{1S} peak in this case yielded two peaks similarly to work by Mannequin et al. who showed the possibility of the peak resulting from O-H, oxygen vacancies or O-C from contaminants or surface species [22,37].

4.2. Electrical characterisation

Dielectric breakdown measurements of the composite films showed a significant improvement in the dielectric strength of 2AB and 4AB composites, resulting in a minimum increase of 93 ± 37 $\text{V}\mu\text{m}^{-1}$ compared to PDC Al_2O_3 alone, whilst the difference between the AB and PDC films was insignificant. The improvements in 2AB and 4AB composites are likely a result of barrier properties of the inserted layers, whereby they inhibit the flow of current through defect sites within the films, meaning that charge transit through the thickness of the material occurs at a higher applied field. Barrier effects have been studied for solid polymer dielectrics by Gefle et al. who discussed a mechanism by which material interfaces prevent charge propagation across films improving the dielectric strength [12]. Other work which implemented clay modification of polymers was shown to improve dielectric strength by up to 17.2% [38,39]. The improvement was attributed to the suppression of charge injection and migration, which resulted in the inhibition of an electron cascade. Interestingly, bi-axially orientated polypropylene only showed marginal improvements (1.1%) because of its intrinsic dielectric strength. Mattox also noted that multilayers can be useful for decreasing the number of pinholes resulting from abnormalities during growth [40].

Taking this research into account it can be seen that incorporating

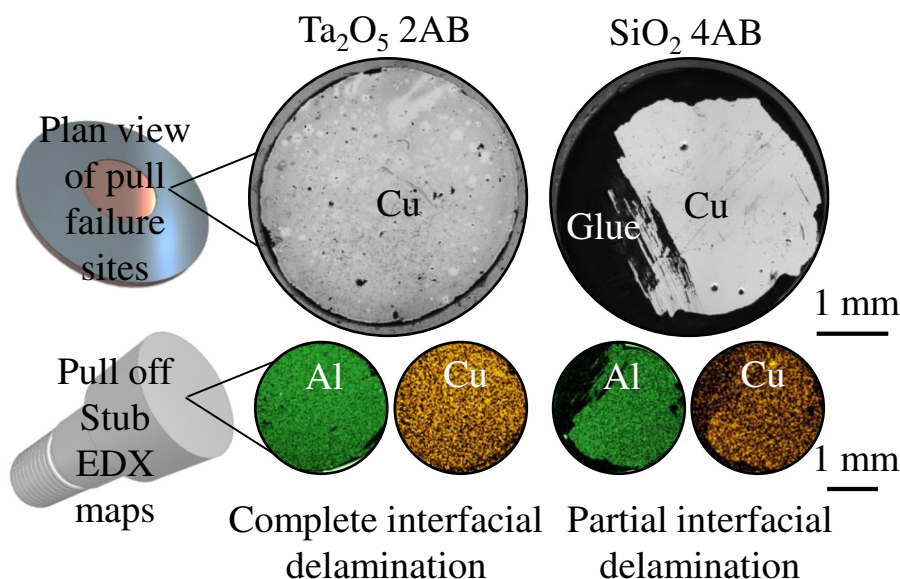


Fig. 5. Example images of complete and partial delamination sites, for Ta₂O₅ 2AB (left) and SiO₂ 4AB (right) respectively. Below each light micrograph are the EDX maps of the pull off stub for Cu and Al, which indicate that the failure was at the Cu substrate interface.

barrier layers with a higher dielectric strength relative to the base material has the potential to increase the breakdown strength and lower leakage current. It is suggested that the mechanisms for the improvement in SiO₂ and Ta₂O₅ composites include: capping structural and compositional defects, prevention of charge injection and disrupting the growth of breakdown channels. The work by Ren et al. showed materials with sufficient intrinsic breakdown strength were not affected by layering, thus the RF deposition method is seen as key to producing films with high enough breakdown voltage to act as sufficient barrier layers in the PDC Al₂O₃.

In ceramic multilayers produced by Martinez-Perdiguero et al. a two times improvement in the dielectric strength of sputtered SiO₂-Al₂O₃ composite coatings was noted, giving a maximum of 80 V μm^{-1} . This condition was met when transitioning from single layer Al₂O₃ into 2,4 and 8 layer composite materials [20]. Improvements of this kind could be the result of added layers masking structural or compositional defects, such as in the Al₂O₃-SiO₂ gas diffusion barrier films, produced using ALD by Dameron et al. [13]. AFM is a powerful tool which has the capability of generating dielectric breakdown measurements and current voltage data which can be used to determine leakage current. This method allows high spatial resolution in a well-defined array avoiding the effect of making multiple measurements too close to one another and allowing the avoidance of extrinsic defects. AFM has been used to measure the dielectric strength of thin films by Ganesan et al. and even to perform layer by layer breakdown testing on BN by Hattori et al. [41,42]. However, AFM dielectric breakdown measurements have not been widely used for multilayer ceramic films past the BN films mentioned above.

Ta₂O₅ composites had a higher maximum dielectric strength of 513 \pm 18 V μm^{-1} than the SiO₂ composites where the maximum was 466 \pm 86 V μm^{-1} (see Fig. 4). There was, however, a large difference in the conduction prior to breakdown with higher current leakage and large oscillations seen in Ta₂O₅ composites resulting from charge trapping and de-trapping. The higher the leakage current the worse the dielectric is at preventing current flow; which is the primary purpose for these films. Larger levels of leakage current are symptomatic of defects in the film, which are capable of facilitating charge transfer. These oscillations caused by charge trapping and de-trapping, which can be caused by intrinsic or electrical stress induced defects, were noticeably more severe in Ta₂O₅ AB composites. Such oscillations were also seen in reactively sputtered Al₂O₃ films as measured by AFM

techniques [8,42]. The maximum current flow prior to breakdown was ca. 6 and 0.1 nA for Ta₂O₅ and SiO₂ composites respectively. This shows that the RF SiO₂ acted as a superior barrier layer than the RF Ta₂O₅. The lack of observable oscillations in the SiO₂ composites suggests that the SiO₂ layers contained fewer defects capable of trapping charge and were not susceptible to stress induced leakage current, which could also cause charge trapping and de-trapping [42].

The Ta₂O₅ and SiO₂ 2AB layer composites produced the films with the highest dielectric strengths. This is likely due to a compromise between layer thickness and frequency of the barrier layers; given that films were all deposited to < 236 nm. The barrier frequency in AB films was insufficient and the thickness of the RF layer in 4AB films was too thin to offer as effective barrier effects (the RF layer is half as thick as the 2AB sample). Such layer dependence was seen in HfO₂ Al₂O₃ composites produced by Park et al. where composites with 7 and 5 layers had enhanced breakdown characteristics compared to 3 layer materials; this phenomena was attributed the Al₂O₃ layers blocking the current between grain boundaries in the HfO₂ layers [43]. The poorer electrical properties in the SiO₂ 4AB composites could also be the result of apparent increased amount of sub Si⁴⁺ silicon from 1.9 to 15.8% at the surface of the SiO₂ composite films as identified in the XPS in Fig. 6.

4.3. Mechanical characterisation

The pull off adhesion strengths of all of the composites were shown to be on average 2.4 times lower than that of the solely PDC alumina films which had an average failure strength of 55.7 \pm 2.9 MPa [8]. The failure also shifted from predominantly partial interfacial failure to complete interfacial failure (in this study, see Fig. 5). The substrates in both cases were treated using the same methods, thus the decrease must be ascribed to the use of multilayers, due to the use of dissimilar materials in the films. It could therefore be the case that the implementation of multilayers resulted in a higher level of intrinsic stress, which would result in this decrease in adhesion. Intrinsic stress in PVD thin films originate from either variation in thermal expansion between the substrate and the films or from growth induced stresses produced during condensation [44]. Knotek et al. have explored the effect of PVD deposition techniques on intrinsic stress, showing the dramatic effect of processing parameters, achieving a 1 GPa reduction of growth induced compressive stress by applying substrate heating [44]. For instance a two fold increase in film adhesion strength was seen when decreasing

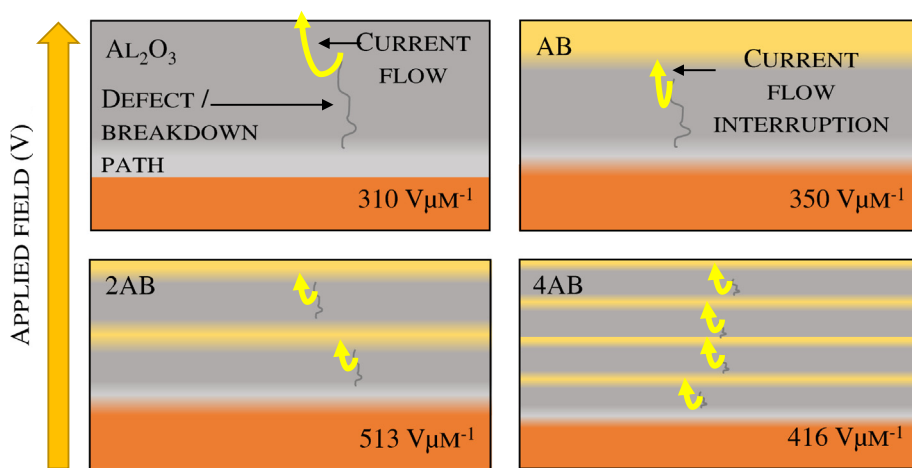


Fig. 6. Effect of multilayers in capping defects and improving barrier properties, by resisting current flow and disrupting the breakdown path. Defects are shown to be passivated in films with multilayer configuration and the dielectric strength is highest for the 2AB films. The maximum dielectric strength for each layer configuration is given in the bottom right of each illustration.

the intrinsic stress from 4.84×10^{-9} to 0.21×10^{-9} Pa by chopping in *ca.* 180–200 nm CeO_2 films deposited by *Patil et al.*, who also showed the same effect in ZnS films [45,46]. The use of interlayers also increased intrinsic stress in multilayer BaTiO_3/Ni ceramic capacitors, analysed by *Park et al.* [11,47]. An increase in the BaTiO_3/Ni layer frequency increased the compressive in plane stress, which reached a maximum at < 135 MPa compressive stress. Complete interfacial failure at the Cu–Cu interface occurred at all test sites in the current study, which indicates that the blending stages were sufficient to mitigate cohesive failure between the sputtered layers. Blending stages have been used extensively for improving the adhesion of diamond like carbon films. For instance, a number of graded layers (such as TiC) can be used to avoid detachment and high interfacial stresses. This is achieved through a gradation of the thermal expansion coefficient (8.6×10^{-6} to 6.2×10^{-6} to $1.0 \times 10^{-6} \text{ }^\circ\text{C}^{-1}$ for Ti, TiC and DLC respectively). Gradation of this kind can also decrease lattice mismatches, such as in the Ti–TiC–DLC system by transitioning through a number of intermediate mixed structures from the dominant hcp phase of Ti to fcc for TiC [48–50]. It was also shown by *Pei et al.* that graded interfaces in ZrCu/Cu multi-layered films improved mechanical properties (elastic modulus, tensile yield stress and tensile elongation increased by a minimum of 1.3 times for 8 to 10 layer films), compared to sharp interfaces because of the reduced stress and mismatches at the interfaces [51].

There was no significant difference in the average adhesion strength of the Ta_2O_5 and SiO_2 composites in the case of the AB and 2AB conformers, with pull off adhesion averages of 20.3 ± 3.8 and 16.7 ± 4.4 MPa for Ta_2O_5 and SiO_2 composites respectively. SiO_2 4AB composites did, however, have superior maximum pull off adhesion over the equivalent Ta_2O_5 composites, as the average pull off adhesion attests: 34.3 ± 4.4 and 15.3 ± 2.8 MPa respectively. The coefficients of thermal expansion were reported to be 5.4×10^{-6} , 5.1×10^{-7} and $3.6 \times 10^{-6} \text{ }^\circ\text{C}^{-1}$ for sputtered Al_2O_3 , SiO_2 and Ta_2O_5 respectively [52]. Thus the decrease in properties when using Ta_2O_5 in the 4AB composites was a direct result of the larger thermal expansion coefficient; which led to an increase in internal stress within the composites upon cooling from the processing temperature. Generally a decrease in properties was seen when transitioning to 2AB and 4AB films from AB films (Table 5) suggesting an increase in internal stress with an increasing number of layers and interfaces.

Combining PDC Al_2O_3 with RF sputtered multilayers has been shown to be beneficial when regarding the dielectric strength of thin films, through a combination of defect capping and disruption of breakdown paths. Whilst the adhesive properties suffered, development of a more comprehensive interlayer system could allow these coatings to be used as insulation for wires or micro electrical components.

5. Conclusions

AFM dielectric breakdown data displayed in this work has shown the electrical breakdown potential for multilayer films consisting of pulsed DC sputtered Al_2O_3 and a RF sputtered SiO_2 or Ta_2O_5 layers to have improved breakdown strength compared to those of pulsed DC Al_2O_3 alone. This was attributed to barrier effects and the requirement for charge carriers to travel across layer interfaces. The best layer configuration was seen to be the 2AB motif, which offered the best compromise between layer thickness and number of layers. Leakage current in SiO_2 composites was a maximum of 60 times lower than that of the Ta_2O_5 composites prior to breakdown. SiO_2 was shown to act as a better barrier layer preventing the flow of current more effectively prior to breakdown. The mechanical testing showed that the blending stages used during film production were sufficient to prevent cohesive failure between the various layers. However, the use of the multilayer systems was shown to decrease the adhesive properties of the films when compared to PDC Al_2O_3 alone by a minimum of two times with respect to pull off adhesion.

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