Article

Investigating the Feasibility of Preparing Metal–Ceramic Multi-Layered Composites Using Only the Aerosol-Deposition Technique

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Abstract: The preparation of metal–ceramic layered composites remains a challenge due to the incompatibilities of the materials at the high temperatures of the co-firing process. For densification, the ceramic thick-film materials must be subjected to high-temperature annealing (usually above 900 °C), which can increase the production costs and limit the use of substrate or co-sintering materials with a low oxidation resistance and a low melting point, such as metals. To overcome these problems, the feasibility of preparing dense, defect-free, metal–ceramic multilayers with a room-temperature-based method should be investigated. In this study, we have shown that the preparation of ceramic–metal Al2O3/Al/Al2O3/Gd multilayers using aerosol deposition (AD) is feasible and represents a simple, reliable and cost-effective approach to substrate functionalisation and protection. Scanning electron microscopy of the multilayers showed that all the layers have a dense, defect-free microstructure and good intra-layer connectivity. The top Al2O3 dielectric layer provides excellent electrical resistance (i.e., $7.7 \times 10^{12} \ \Omega \cdot m$), which is required for reliable electric field applications.

Keywords: aerosol deposition; multilayers; interdigitated aluminium electrodes; alumina insulating layers

1. Introduction

Thick-film multilayer technology is of great value in the electronics industry. It enables the development and design of a wide variety of products, such as microsystems, electronic circuit boards and micro-electromechanical systems. The fabrication of conventional thick-film multilayers involves well-developed technologies, i.e., tape-casting and screen-printing, followed by a co-firing process at elevated temperatures [1]. The co-firing process poses many problems in the fabrication of metal–ceramic multilayers. First, the mismatch of firing temperatures significantly limits the choice of compatible materials, as metals require much lower temperatures than ceramics. Second, a high-temperature process facilitates metal oxidation and diffusion between the layers, which can lead to functional degradation, posing major challenges in material selection [2]. On the other hand, a high-temperature firing process can be avoided by using aerosol deposition (AD), which is a room-temperature spray-coating method for producing dense, micrometre-thick films. It requires dry powders of approximately micrometre-sized particles that are mixed with a carrier gas to form an aerosol [3]. In the deposition process, the aerosol jet hits the substrate with a high kinetic energy under vacuum conditions. The AD method is often referred to as a simple and rapid deposition method capable of producing very dense films without adding any external thermal energy to the aerosol or deposited films [4]. The high density of
the deposited materials is achieved by the hammering effect of powder particles colliding with the surface of the substrate, fragmenting and re-bonding [4–6]. This deposition mechanism is referred to as a room-temperature, impact-consolidation mechanism [6]. AD is a unique approach to the deposition of ceramic coatings at room temperature, which means the vast majority of research has been conducted on ceramic coatings, such as simple oxides (Al₂O₃ [7], TiO₂ [8], Y₂O₃ [9]), perovskites (Pb(Zr,Ti)O₃ [10], BaTiO₃ [11]) and others (AlN [12], hydroxyapatite [13]). However, AD is not limited to ceramics; metals, glasses or polymers can also be deposited. The deposition of metal layers is often associated with other spray-coating techniques, such as cold spray, which requires heating of the carrier gas (up to 700 °C). In contrast to cold spray, the AD process is much more advantageous for the deposition of metal layers, since the deposition at room temperature avoids deleterious oxidation, decomposition and thermal shock to the coating and the substrate [14]. However, the AD of metal films is still poorly understood, since in the AD community research is mainly focused on ceramic deposition. The deposition of metals poses a great challenge because the modification of powder parameters (e.g., particle size distribution, morphology and agglomeration state) is much more challenging for metals compared to ceramic materials. For example, ceramic powders can be coarsened very easily by partial sintering, which only requires heating the powder at ambient atmosphere. In addition, the brittleness of ceramic powders allows straightforward particle size reduction and de-agglomeration in a ball milling process. On the other hand, metal powders are much more susceptible to oxidation during heat treatment and their ductile behaviour significantly complicates the milling process. The first dense metal films prepared using AD at room temperature were reported less than 10 years ago [15]. The development of metal films by AD is still in its early stages. Thus far, AD has been used to deposit metal films such as Cu [16,17], Ag [15,18], Fe [19,20] and Fe-based amorphous alloys [21]. In this study, we have deposited metal layers of Al, which has not yet been reported.

The AD method is mostly used for the fabrication of single layers [4]. Although AD is considered as an additive technology, only a few multilayers have been demonstrated thus far. There are few reports involving the fabrication of multilayers using a combination of different deposition methods. For example, functional ceramic layers are deposited by AD and conductive metal electrode layers are deposited by physical vapour deposition [22,23]. Such an approach can pose difficulties because the metal layers deposited via physical vapour deposition are very thin and their properties degrade after an additional AD of ceramic layers on top. The impact of the ceramic powder on the metal layer is particularly problematic, leading to roughening of the metal–ceramic interface and a potential connectivity loss of the metal layer. There are few reports dealing with the fabrication of multilayers using the AD method. Simple structures consist of ceramic–ceramic [24] or metal–metal multilayers [20]. Thus far, only Leupold et al. [19] have fabricated a ceramic–metal multilayer structure. In this report, multiple thick-film layers of Al₂O₃ and Fe were interchangeably deposited, but the microstructural analysis again revealed the connectivity loss of the metal layers. It is obvious that using the AD method to build a metal–ceramic multilayer without structural defects is still a challenge.

In this investigation, we look at the possibility of fabricating a laminated metal–ceramic composite, i.e., a multilayer with a good intra-layer connectivity, using only the AD method. Ceramic alumina (Al₂O₃) and metallic aluminium (Al) powders were selected for the deposition due to their low price, high abundance and because they are one of the most commonly used dielectric and electrically conductive materials, respectively. Al₂O₃ is often used as an insulating and protective coating, while Al is used as an electrode material. In our previous report we showed that an Al₂O₃ thick film deposited on a Gd substrate provides an excellent electrical insulating layer [25]. Based on this, we used AD to build a metal–ceramic multilayer structure with an inter-digitated electrode layer to add functionality to the system. The deposited Al₂O₃/Al/Al₂O₃ multilayers on the Gd substrate provide an excellent starting point for the development of future electrowetting-on-dielectric devices, such as those used as thermal switches in the magnetocaloric cooling
process [26,27] or potentially in any other solid-state, fluidic or mechanical thermal control devices [28,29]. In this study, we have shown that multilayer fabrication using AD is feasible and represents a simple, reliable and cost-effective approach to add functionality and protection to existing substrates.

2. Materials and Methods

In the AD, a raw Al powder (99.96 %, 2 HPC, Toyal Europe, Accous, France) and an Al$_2$O$_3$ powder (99.8 %, A 16 SG, Almatis, Ludwigshafen, Germany) were used. The Al$_2$O$_3$ powder was thermally treated in a chamber furnace (Custom-made, Terna, Ljubljana, Slovenia) at 1150°C for 1 h (with 5 K-min$^{-1}$ heating and cooling rates) and additionally milled in a planetary ball mill (PM400, Retsch, Haan, Germany) at 200 min$^{-1}$ for 5 h using Al$_2$O$_3$ milling jar and yttria-stabilised-zirconia milling balls with diameters of 3 mm in iso-propanol as a liquid medium. The thermal treatment and the subsequent ball milling of the Al$_2$O$_3$ powder are necessary to achieve suitable particle size distribution for efficient AD [25]. In the case of Al, the as-received powder already resulted in successful film deposition. Therefore, no further powder modification was required. Prior to the AD, both powders were sieved through an 80-micrometre mesh and vacuum dried for 12 h at 100°C and at 10 mbar. The AD apparatus was provided by InVerTec e.V., Bayreuth, Germany. Commercial gadolinium foils (Metall Rare Earth Limited, Hong Kong) were used as the substrate material. The process parameters used during the AD are shown in Table 1. The number of scans was adjusted to achieve the desired film thickness.

Table 1. Process parameters used during the AD.

| Process Parameters | Al$_2$O$_3$ Powder | Al Powder |
|--------------------|--------------------|-----------|
| Carrier gas species | N$_2$               |           |
| Nozzle geometry (slit size) | (0.5 × 10) mm$^2$ |           |
| Distance between nozzle and substrate | 5 mm |           |
| Sweep speed | 5 mm·s$^{-1}$ |           |
| Gas flow rate | 4 L·min$^{-1}$ | 2 L·min$^{-1}$ |
| Pressure in aerosol chamber | 180 mbar | 120 mbar |
| Pressure in deposition chamber | 2 mbar | 1 mbar |

Particle size analyses of the raw Al powder and treated Al$_2$O$_3$ powders were performed using a light-scattering laser granulometer (S3500, Microtrac, York, PA, USA) with isopropanol as the medium. Scanning electron microscopy (SEM) and energy dispersive spectroscopy (EDS) analyses were performed using a field-emission scanning electron microscope (FE-SEM, JSM-7600F, JEOL, Tokyo, Japan) equipped with an energy dispersive X-ray spectrometer (Inca Oxford 350 EDS SSD, Oxford Instruments, Abingdon, UK). For the SEM powder analyses, Al$_2$O$_3$ and Al powders were deposited on carbon tape. For the cross-sectional analysis of the multilayers, the samples were cut, mounted in epoxy resin, ground and fine-polished with a colloidal silica suspension. Prior to the SEM analyses, all the samples were coated with a 3-nanometre-thick carbon layer using a Precision Etching and Coating System (PECS 682, Gatan, Pleasanton, CA, USA).

The X-ray diffraction (XRD) analysis was performed using a high-resolution diffractometer (X’Pert PRO, PANalytical, Almelo, The Netherlands) with Cu–K$_\alpha_1$ radiation. Diffraction patterns were recorded in the BraggBrentano geometry with a 100-channel X'Celerator detector in a 2θ range 10–120° with a step of 0.017° and an integration time of 100 s per step. The software X’Pert HighScore Plus 2.1, PANalytical was used to analyse the XRD patterns and to estimate the penetration depth of X-rays in the multilayer samples during XRD analysis. In the case of the multilayer sample, the X-rays penetrate all
deposited layers (top Al₂O₃ layer, middle Al layer and bottom Al₂O₃ layer) and reach the Gd substrate, since the penetration depth is higher than the thickness of deposited layers. The total thickness of deposited layers is ~8 µm; while the calculated penetration depth for Al and Al₂O₃ at 2θ of 10° is ~30 µm and the value even increases with increasing 2θ angle.

The Topas R package (version 2.1, Bruker AXS GmbH, Karlsruhe, Germany) was employed for the Rietveld refinement and the Fundamental Parameters Approach (FPA) was used for line-profile fitting of all samples [30]. The FPA uses the geometrical properties of the diffraction experiment to build up the instrumental linewidth from first principles. It, thus, allows an explicit determination of the sample-dependent, line-broadening contributions to the peak profile, which are dominated by the microstrain and the broadening of the crystallite size [31].

The topography of the layers was analysed using atomic force microscopy (AFM) and contact stylus profilometry. The 20-micrometre line scans were acquired using an atomic force microscope (Jupiter XR, Asylum Research, Santa Barbara, CA, USA) in AC air topography mode. A Si tip on a Si/Al cantilever with a diameter of ~7 nm (AC240TS-R3, Asylum Research, Santa Barbara, CA, USA) was used for scanning. A contact stylus profilometer (DektakXT, Bruker, Karlsruhe, Germany) was used to measure 2-millimetre line scans. Then, the root-mean-square surface roughness (Rq) was determined from the roughness profile obtained after high-pass filtering of the primary profile with a cut-off wavelength of 0.08 mm.

Silver electrodes with a diameter of 0.75 mm were deposited on the top Al₂O₃ layer before electrical characterisation. Current density–electric field (J–E) measurements were performed using a Keithley 237 high-voltage-source measurement unit (Keithley Instruments, Cleveland, OH, USA). A step-like electric field in the range ±75 kV·cm⁻¹ was applied between Al layer and silver electrodes. The electrical resistivity was determined from the slope of the J–E curve, assuming Ohm’s law.

3. Results

We prepared an Al₂O₃/Al/Al₂O₃ multilayer structure on the surface of a magnetocaloric gadolinium element. A schematic representation and a photograph of the multilayer structure are shown in Figure 1a,b, respectively. First, the Gd substrate was almost completely covered with an Al₂O₃ layer, followed by the deposition of an Al layer. Then, the Al₂O₃ layer was deposited on top of the Al layer, keeping certain areas free to allow for electrical connections (i.e., placement of the contact wires). The first Al₂O₃ layer protects the Gd substrate from the environment and prevents corrosion and mechanical damage. The second Al layer is an electrically conductive electrode layer covered by the third Al₂O₃ layer, which electrically insulates the Al surface and completes the multilayer structure with an embedded electrode.
The Al₂O₃ and Al powders used in the deposition were analysed using laser granulometry and SEM. Laser granulometry shows a monomodal (Figure 2a) and a multi-modal (Figure 2b) particle size distribution of the Al₂O₃ and Al powders, respectively. The particle size range of the two powders is between 0.1 and 20 µm, and the median particle size ($d_{50}$) of the Al₂O₃ and Al powders is 0.6 and 1.5 µm, respectively. The most abundant particle size fraction of the Al₂O₃ powder is represented by the peak at 0.6 µm, while the Al powder contains three peaks at 0.2, 0.5 and 3 µm. Both powders have an acceptable particle size range for the AD. In the literature, particles with sizes between a few hundred nm and a few µm are generally considered suitable for deposition and produce dense thick films with good adhesion [4,6]. According to the SEM analysis, the Al powder exhibits round particles, often with a perfect circular shape (Figure 2d). In contrast, the Al₂O₃ particles are irregularly shaped with smooth edges (Figure 2c).

XRD analysis (Figure 3) was performed on the Al₂O₃ and Al powders, on the Gd substrate and on the multilayer sample after deposition. The XRD patterns of the Al₂O₃...
and Al powders contained only Al₂O₃ (JCPDS 46-1212) and Al reflections (JCPDS 89-2769), respectively. Therefore, no significant powder contamination in the ball milling process of the Al₂O₃ powder was detected. The very sharp peaks indicate large crystallites (>100 nm) and no microstrain in both powders. As expected, the XRD pattern of the Gd substrate also exhibits sharp Gd reflections (JCPDS 89-2924) with no impurities. After deposition, the XRD pattern of the multilayer (purple) shows the reflections of the Al layer, the Al₂O₃ layer and the Gd substrate. None of the three phases underwent a phase transformation. In the multilayer, the Gd reflections (marked with a red cross) did not undergo any peak shift or change in the peak shape. Only the intensity of the Gd decreased, since the substrate was covered by Al₂O₃ and Al layers. After the deposition, the multilayer sample exhibits peak broadening of the Al₂O₃ (marked with a grey asterisk) and Al (marked with blue dash) reflections, indicating a decrease in the crystallite size and/or an increase in the microstrain due to the fragmentation of the colliding powder particles in the AD process [4,32]. To quantitatively evaluate the crystallite size and microstrain in the Al₂O₃ and Al deposited layers, we performed a Rietveld refinement. The calculated crystallite size and microstrain in the Al₂O₃ layers are 22 ± 4 nm and 0.46 ± 0.14%, respectively. On the other hand, the crystallites in the Al layers are larger, i.e., 98 ± 8 nm, while the microstrain is almost insignificant, i.e., 0.04 ± 0.01%. In conclusion, the XRD results show that the deposition of Al₂O₃ leads to a much larger decrease in crystallite size and an increase in microstrain compared to Al.

![X-ray diffraction (XRD) patterns of the Al₂O₃ powder (grey), Al powder (blue), Gd substrate (red) and prepared multilayer sample (purple).](image)

Figure 3. X-ray diffraction (XRD) patterns of the Al₂O₃ powder (grey), Al powder (blue), Gd substrate (red) and prepared multilayer sample (purple).

Previously, it was reported that a significant reduction in the crystallite size and an increase in the microstrain after the deposition of an oxide powder are necessary conditions for successful film deposition [33]. According to our XRD analysis, the same hypothesis can be valid for the AD of ceramic Al₂O₃, but not completely for the AD of metallic Al. Both Al₂O₃ and Al powders formed consolidated layers, but only the ceramic Al₂O₃ powder obtained significant peak broadening, indicating that the cracking and fragmentation of particles predominate in the deposition mechanism. On the other hand, the deposition mechanism of ductile metals is different from that of ceramics. The less intense XRD peak broadening of Al indicates that plastic deformation is more prevalent in the deposition mechanism of metals.

The SEM and EDS analyses of the multilayer structure in cross-section are shown in Figure 4. All the deposited layers are very dense without any visible pores (Figure 4a). The thickness of the deposited Al₂O₃, Al and Al₂O₃ layers (bottom-up) is 5.0, 1.7 and 1.6 µm, respectively. Apparently, the roughness of the interfaces increases with the number of increasing layers. However, a sufficient layer thickness ensures good connectivity of the deposited layers. The interface between the layers is well defined (enlarged SEM images in
Figure 4b,c), indicating good adhesion without any reactions. An additional adhesion test revealed no peeling off or delamination of the deposited layers (Supplementary material: Figure S1). The EDS map scans (Figure 4d–f) of the multilayer show a typical elemental distribution of the layers, which confirms no reaction between the layers.

The surface roughness of the deposited layers including the Gd substrate was evaluated using AFM and contact profilometry. The AFM map scans and the contact-profilometry line scans are shown in Figure 5a,b, respectively. A comparison of the root-mean-square surface roughness ($R_q$) between the two measurement methods is shown in Figure 5c. As expected, the values obtained with AFM are lower than those obtained with contact profilometry due to the hundred-times-smaller scanned area (20-micrometre lines in the case of the AFM and 2-millimetre lines in the case of the contact profilometry). However, the same trend is observed for both methods. The $R_q$ is lowest for the Gd substrate and increases with each deposited layer. The $R_q$ of the top Al$_2$O$_3$ layer is about 200 nm.

To test the electrical insulation of the upper Al$_2$O$_3$ layer, measurements of the current density ($J$) versus the electric field ($E$) were performed (Figure 6). The upper Al$_2$O$_3$ layer withstands high electric fields (75 kV·cm$^{-1}$) without an electrical breakdown. The calculated electrical resistivity at room temperature is very high ($7.7 \times 10^{12}$ $\Omega$·m) and corresponds to the resistivity of the commercially available Al$_2$O$_3$ ceramics for electrical insulation [34,35].

Figure 4. (a) Scanning electron microscopy (SEM) analysis of the multilayer structure in cross-section. Panels (b,c) show the magnified areas marked in (a). Energy dispersive spectroscopy (EDS) elemental mapping images are showing the distribution of (d) Al, (e) Gd and (f) O.
Figure 5. (a) Atomic force microscopy (AFM) map scans and (b) contact-profilometer line scans of the deposited layers, including the Gd substrate. The corresponding Rq values of the two methods, AFM (cyan) and contact profilometry (dark blue), are shown in (c).

Figure 6. The current density (J) vs. electric field (E) curve (grey) of the top Al2O3 layer and its linear approximation (blue).

4. Conclusions

We used the AD method to prepare an Al2O3/Al/Al2O3 multilayer composite on a Gd substrate. The complete room-temperature processing of the AD enabled the integration of metallic and ceramic materials that are otherwise incompatible at high temperatures. Inexpensive commercial powders with appropriate micrometre-sized particles were used for the successful film deposition. The SEM analysis revealed a dense multilayer with a defect-free microstructure and good intra-layer connectivity. In addition, the top Al2O3
dielectric layer provides excellent electrical resistance, which is required for reliable electric field application. In summary, we have shown that the fabrication of ceramic–metal multilayers using AD is feasible and represents a simple, reliable and cost-effective approach to functionalise and protect existing substrates. For example, the deposited $\text{Al}_2\text{O}_3/\text{Al}/\text{Al}_2\text{O}_3$ multilayers on the Gd substrate provide an excellent starting point for the development of future electrowetting-on-dielectric devices.

**Supplementary Materials:** The following are available online at https://www.mdpi.com/article/10.3390/ma14164548/s1, Figure S1: Images of the sample surface (top $\text{Al}_2\text{O}_3$ layer) taken with an optical light microscope. (a) before and (b) after the adhesion test.

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**Data Availability Statement:** The data presented in this study are available upon request from the corresponding author.

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**References**

1. Kosec, M.; Kuscer, D.; Holc, J.; Pardo, L.; Ricote, J. Processing of Ferroelectric Ceramic Thick Films. In *Multifunctional Polycrystalline Ferroelectric Materials*; Springer: Dordrecht, The Netherlands, 2011.

2. Pan, M.-J.; Randall, C. A brief introduction to ceramic capacitors. IEEE Electr. Insul. Mag. 2010, 26, 44–50. [CrossRef]

3. Schubert, M.; Hanft, D.; Nazarenus, T.; Exner, J.; Nieke, P.; Gloses, P.; Leupold, N.; Kita, J.; Moos, R. Powder aerosol deposition method—Novel applications in the field of sensing and energy technology. Funct. Mater. Lett. 2019, 12, 1930005. [CrossRef]

4. Hanft, D.; Exner, J.; Schubert, M.; Stöcker, T.; Fuierer, P.; Moos, R. An overview of the Aerosol Deposition method: Process fundamentals and new trends in materials applications. J. Ceram. Sci. Technol. 2015, 6, 147–181. [CrossRef]

5. Lee, D.-W.; Kim, H.-J.; Kim, Y.-H.; Yun, Y.-H.; Nam, S.-M. Growth Process of Alpha-$\text{Al}_2\text{O}_3$ Ceramic Films on Metal Substrates Fabricated at Room Temperature by Aerosol Deposition. J. Am. Ceram. Soc. 2011, 94, 3131–3138. [CrossRef]

6. Akedo, J. Room Temperature Impact Consolidation (RTIC) of Fine Ceramic Powder by Aerosol Deposition Method and Applications to Microdevices. J. Therm. Spray Technol. 2008, 17, 181–198. [CrossRef]

7. Saunders, R.; Johnson, S.D.; Schwer, D.; Patterson, E.A.; Ryou, H.; Gorzkoowski, E.P. A Self-Consistent Scheme for Understanding Particle Impact and Adhesion in the Aerosol Deposition Process. J. Therm. Spray Technol. 2021, 30, 523–541. [CrossRef]

8. Ryu, J.; Park, D.-S.; Hahn, B.-D.; Choi, J.-J.; Yoon, W.-H.; Kim, K.-Y.; Yun, H.-S. Photocatalytic TiO$_2$ thick films by aerosol-deposition: From micron-sized particles to nano-grained thin film at room temperature. Appl. Catal. B Environ. 2008, 83, 1–7. [CrossRef]

9. Lee, B.-K.; Park, D.-S.; Yoon, W.-H.; Ryu, J.-H.; Hahn, B.-D.; Choi, J.-J. Microstructure and Properties of Yttria Film Prepared by Aerosol Deposition. J. Korean Ceram. Soc. 2009, 46, 441–446. [CrossRef]

10. Baba, S.; Tsuda, H.; Akedo, J. Thickness dependence of electrical properties of PZT films deposited on metal substrates by laser-assisted aerosol deposition. IEEE Trans. Ultrason. Ferroelectr. Freq. Control 2008, 55, 1009–1016. [CrossRef]

11. Khansur, N.H.; Eckstein, U.; Benker, L.; Deisinger, U.; Merle, B.; Webber, K.G. Room temperature deposition of functional ceramic films on low-cost metal substrate. Ceram. Int. 2018, 44, 16295–16301. [CrossRef]

12. Heo, Y.J.; Kim, H.T.; Kim, K.J.; Nahm, S.; Yoon, Y.J.; Kim, J. Enhanced heat transfer by room temperature deposition of AlN film on aluminum for a light emitting diode package. Appl. Therm. Eng. 2013, 50, 799–804. [CrossRef]

13. Park, D.-S.; Kim, I.-S.; Kim, H.; Chou, A.H.K.; Hahn, B.-D.; Li, L.-H.; Hwang, S.-J. Improved biocompatibility of hydroxyapatite thin film prepared by aerosol deposition. J. Biomed. Mater. Res. Part B Appl. Biomater. 2010, 94, 353–358. [CrossRef]

14. Papyrin, A. Cold Spray Technology. Adv. Mater. Proc. 2001, 159, 49–51.

15. Kim, Y.-H.; Lee, J.-W.; Kim, H.-J.; Yun, Y.-H.; Nam, S.-M. Silver metallization for microwave device using aerosol deposition. Ceram. Int. 2012, 38, S201–S204. [CrossRef]

16. Lee, D.-W.; Kwon, O.-Y.; Cho, W.-J.; Song, J.-K.; Kim, Y.-N. Characteristics and Mechanism of Cu Films Fabricated at Room Temperature by Aerosol Deposition. Nanoscale Res. Lett. 2016, 11, 1–8. [CrossRef] [PubMed]
17. Khansur, N.H.; Eckstein, U.; Ursic, H.; Sadl, M.; Brehl, M.; Martin, A.; Riess, K.; de Ligny, D.; Webber, K.G. Enhanced Electromechanical Response and Thermal Stability of 0.93(Na_{1/2}Bi_{1/2})TiO_3-0.07BaTiO_3 Through Aerosol Deposition of Base Metal Electrodes. *Adv. Mater. Interfaces* 2021, 8, 2100309. [CrossRef]

18. Cho, M.-Y.; Lee, D.-W.; Kim, I.-S.; Lee, W.-H.; Yoo, J.-W.; Ko, P.-J.; Koo, S.-M.; Choi, Y.-K.; Oh, J.-M. Formation of silver films for advanced electrical properties by using aerosol deposition process. *Ipn. J. Appl. Phys.* 2018, 57, 11UF05. [CrossRef]

19. Leupold, N.; Denneler, S.; Rieger, G.; Moos, R. Powder Treatment for Increased Thickness of Iron Coatings Produced by the Powder Aerosol Deposition Method and Formation of Iron–Alumina Multilayer Structures. *J. Therm. Spray Technol.* 2020, 30, 480–487. [CrossRef]

20. Sugimoto, S.; Chan, V.; Noguchi, M.; Tezuka, N.; Inomata, K.; Akedo, J. Preparation of Fe/Ni–Zn–Cu ferrite stacked films by aerosol deposition method. *J. Magn. Magn. Mater.* 2007, 310, 2549–2551. [CrossRef]

21. Kwon, J.; Park, H.; Lee, I.; Lee, C. Effect of gas flow rate on deposition behavior of Fe-based amorphous alloys in vacuum kinetic spray process. *Surf. Coat. Technol.* 2014, 259, 585–593. [CrossRef]

22. Imanaka, Y.; Akedo, J. Embedded capacitor technology using aerosol deposition. *Int. J. Appl. Ceram. Technol.* 2010, 7, 23–32. [CrossRef]

23. Miyoshi, T. Preparation of multilayer piezoelectric device by aerosol deposition using a novel detachment method. *J. Ceram. Soc. Jpn.* 2009, 117, 899–903. [CrossRef]

24. Kambale, R.C.; Han, G.; Yoon, W.-H.; Kim, J.-W.; Choi, J.-J.; Hahn, B.-D.; Park, D.-S.; Ahn, C.-W.; Ryu, J. Reliability of ferroelectric multilayer PZT thick films fabricated by aerosol deposition. *Ferroelectrics* 2014, 470, 183–193. [CrossRef]

25. Sadl, M.; Tomc, U.; Prah, U.; Ursic, H. Protective Alumina Coatings Prepared by Aerosol Deposition on Magnetocaloric Gadolinium Elements. *Inf. MIDEM J. Microelectron. Electron. Compon. Mater.* 2019, 49, 177–182. [CrossRef]

26. Cha, G.; Kim, C.-J.; Ju, Y.S. Thermal conductance switching based on the actuation of liquid droplets through the electrowetting on dielectric (EWOD) phenomenon. *Appl. Therm. Eng.* 2016, 98, 189–195. [CrossRef]

27. Klinar, K.; Kitanoivski, A. Thermal control elements for caloric energy conversion. In *Renewable and Sustainable Energy Reviews*; Elsevier: Amsterdam, The Netherlands, 2020; Volume 118, p. 109571. [CrossRef]

28. Swoboda, T.; Klinar, K.; Yalamarthi, A.S.; Kitanoivski, A.; Rojo, M.M. Solid-State Thermal Control Devices. In *Advanced Electronic Materials*; Blackwell Publishing Ltd.: Hoboken, NJ, USA, 2021; Volume 7, p. 2000625. [CrossRef]

29. Klinar, K.; Swoboda, T.; Rojo, M.M.; Kitanoivski, A. Fluidic and Mechanical Thermal Control Devices. In *Advanced Electronic Materials*; Blackwell Publishing Ltd.: Hoboken, NJ, USA, 2021; Volume 7, p. 2000623. [CrossRef]

30. Cheary, R.W.; Coelho, A. A fundamental parameters approach to X-ray line-profile fitting. *J. Appl. Crystallogr.* 1992, 25, 109–121. [CrossRef]

31. Klug, H.P.; Alexander, L.E. *X-Ray Diffraction Procedures: For Polycrystalline and Amorphous Materials*, 2nd ed.; John Wiley & Sons: New York, NY, USA, 1974.

32. Akedo, J. Aerosol deposition of ceramic thick films at room temperature: Densification mechanism of ceramic layers. *J. Am. Ceram. Soc.* 2006, 89, 1834–1839. [CrossRef]

33. Exner, J.; Schubert, M.; Hanft, D.; Kita, J.; Moos, R. How to treat powders for the room temperature aerosol deposition method to avoid porous, low strength ceramic films. *J. Eur. Ceram. Soc.* 2019, 39, 592–600. [CrossRef]

34. Barsoum, M.W. *Fundamentals of Ceramics*; CRC Press: Boca Raton, FL, USA, 2019.

35. Kingery, W.D.; Bowen, H.K.; Uhlmann, D.R. *Introduction to Ceramics*; Wiley: London, UK, 1976.