Sub-nanometer Atomic Layer Deposition for Spintronics in Magnetic Tunnel Junctions Based on Graphene Spin-Filtering Membranes

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ABSTRACT We report on the successful integration of low-cost, conformal, and versatile atomic layer deposited (ALD) dielectric in Ni—Al2O3—Co magnetic tunnel junctions (MTJs) where the Ni is coated with a spin-filtering graphene membrane. The ALD tunnel barriers, as thin as 0.6 nm, are grown layer-by-layer in a simple, low-vacuum, ozone-based process, which yields high-quality electron-transport barriers as revealed by tunneling characterization. Even under these relaxed conditions, including air exposure of the interfaces, a significant tunnel magnetoresistance is measured highlighting the robustness of the process. The spin-filtering effect of graphene is enhanced, leading to an almost fully inversed spin polarization for the Ni electrode of 42%. This unlocks the potential of ALD for spintronics with conformal, layer-by-layer control of tunnel barriers in magnetic tunnel junctions toward low-cost fabrication and down-scaling of tunnel resistances.

KEYWORDS: atomic layer deposition · spintronics · spin filter · graphene · dielectrics · magnetic tunnel junction

Atomic layer deposition (ALD) is a key technology for applications requiring ultrathin, uniform, high-quality, and conformal films. Owing to the chemically surface-limited growth mechanisms involved, ALD enables atomic control of the film thickness even for high aspect ratio substrates while being scalable and compatible with other industrial techniques such as roll-to-roll processing.1–3 These advantages are already exploited in a number of applications, including multilayer stacks in thin film electroluminescent flat panel displays,4,5 high-quality dielectrics in dynamic random access memory (DRAM) capacitors,6,7 conformal passivation coatings for organic electronics, e.g., organic light-emitting diodes (OLED),8,9 and conformal functionalization for porous biomaterials and bioelectronic devices.10,11 Furthermore, ALD-grown ultrathin, high-k gate dielectrics were industrially adopted for complementary metal oxide semiconductor (CMOS) logic technologies for high-volume manufacturing as early as the 45 nm node,12 and ALD is particularly well adapted for the high aspect ratio architectures such as FinFET and TriGate transistors used in the current 22 nm node.13,14

To date, despite a number of pioneering attempts15–19 with promising results, the advantages of ALD have not been widely employed in the field of magnetic tunnel junctions for spintronics, which has instead had to rely on more complex physical deposition setups. Spin valves, the active elements in technologies such as hard-drive read-heads and magnetic random-access memories (MRAM),20 would appear to be prime candidates to benefit from the use of ALD.15–17 Spin devices rely heavily on ultrathin films such as nanometer-scale oxide tunnel barriers, whose quality and atomic composition crucially impact the electron’s spin polarization. A key issue has however remained: performing ALD directly...
on metallic spin sources (usually ferromagnets such as Ni, Co, Fe, and their alloys) leads to their oxidation during the fabrication and thus a quenching of their spintronic performances. This has drastically limited the range of processes that can be used to fabricate spin-valve devices. Hence, although promising in terms of thickness and composition control, the application of water- or ozone-based ALD processes to magnetic tunnel junctions has so far been hindered by the associated ferromagnet oxidation.15–17

Here we demonstrate the successful realization of graphene-coated Ni/Al₂O₃/Co magnetic tunnel junctions based on sub-nanometer, ozone-based ALD-grown tunnel barriers. We are thus able to take advantage of the layer-by-layer thickness control, inherent to ALD, to down-scale the thickness and the resistance × area product of tunnel barriers. Measurements of large inverse tunnel magneto-resistance (TMR) signals highlight the almost full reversal of the graphene-coated Ni spin polarization (P = 42%) compared to the max reported value (P = +46% for the highest quality Ni/Al₂O₃ interfaces) and demonstrate the quality of the ALD barrier.

RESULTS AND DISCUSSION

In Figures 1 and 2, we present the technological steps employed to integrate ALD-grown tunnel barriers in vertical spin valves. Ni bottom electrodes are defined by a standard lift-off step, and a subsequent chemical vapor deposition (CVD) process covers these electrodes with a continuous, few-layer graphene coating. This yields a reduced and passivated Ni surface. This direct graphene growth method on patterned ferromagnetic electrodes is readily scalable and avoids graphene transfer or exfoliation steps otherwise typical for the fabrication of graphene devices.15,16 A 1 μm² junction is then defined in the patterned resist (see Figure 1). Next, after exposing the bottom electrode to air, ALD is performed with cycles of ozone and trimethylaluminum (TMA) pulses to grow ultrathin Al₂O₃ layers (see Methods). This fabrication flow results in electrodes fully covered with an Al₂O₃ thin film. We note that the growth of ultrathin dielectrics on sp²-bonded graphene coatings has previously been considered challenging: both the preservation of the pristine graphene’s sp² structure and the homogeneous wetting of the dielectric require careful engineering, which has been particularly discussed for films deposited by evaporation or water-based ALD. In particular, ALD on graphene has been shown to require surface seeding (such as modified ALD cycles, inorganic clusters, or molecular layers such as PTCA) to increase wetting as required for ultralow film thicknesses. Interestingly, the unseeded ozone-based ALD (O₃-ALD) process used here is shown to be nondetrimental to the graphene layer (see Figure 3), while yielding Al₂O₃ films homogeneous enough to be used as tunnel barriers (see Figures 4 and 5).

As ozone is a powerful oxidant, its use in the process might be expected to result in significant oxidation of the graphene coating. We have thus investigated the impact of the whole O₃-ALD growth process on CVD graphene monolayers using Raman spectroscopy, for three different growth temperatures (T): 80, 150, and 200 °C (see Figure 3). At the highest growth temperature of 200 °C, a strong Raman defect-enabled D peak (1350 cm⁻¹) appears in the spectra, with an amplitude even exceeding that of the graphitic G peak (1600 cm⁻¹), indicating a heavily degraded sp² structure.
structure.30 Clearly at this high temperature, ozone reacts with the graphene layer introducing defects and forming functional groups such as epoxides on carbon sites.31,32 Remarkably, however, this reactivity decreases sharply with lower growth temperature. In particular, Raman spectra measured after the 80°C growth of Al2O3 remain unchanged compared to those measured on bare as-grown CVD graphene (see Figure 3): graphene appears to be inert to ozone during the 80°C ALD process. These observations relate to the fact that ozone physisorption occurs on the graphene surface, while further chemical bonding between the oxygen and carbon atoms requires a non-negligible activation energy barrier to be overcome.31,32 In the case of our 80°C O3-ALD process, the kinetics of this reaction are quenched for the exposure times considered. This inert process is thus considered in the following to grow ultrathin Al2O3 films in complete magnetic tunnel junctions.

Given the low wettability of the carbon sp2 structure, another challenge is the formation of continuous, ultrathin, pinhole-free tunnel barriers on graphene. Despite it so far not being achievable by evaporation,33 we previously reported that sputtering led to high-quality Al2O3 films on graphene down to a thickness of 1 nm.34 However, growing Al2O3 films thinner than 1 nm on graphene by sputtering did not yield continuous layers. In particular, AFM characterization of a 0.6 nm tunnel barrier, produced by sputtering a 0.4 nm film of metallic Al that was further oxidized in a 50 Torr O2 atmosphere (as for the high-quality 1 nm tunnel barrier34), reveals a particularly high roughness (RMS > 0.7 nm, even in excess of the targeted nominal film thickness). In fact, at these lower thicknesses the coalescence of Al islands is not achieved following the sputtering process, leading to severely pinholed films. In stark contrast, 0.6 nm of Al2O3 obtained by the O3-ALD process on graphene shows significantly higher homogeneity and no observable pinholes, as confirmed by AFM. The measured root-mean-square (RMS) roughness of 0.25 nm for the 0.6 nm ALD film is comparable to what is achieved for the thicker, high-quality, 1 nm, sputtered films. These characteristics underline that the inherent attributes of ALD are well suited for the needs of spintronics devices in terms of producing conformal, ultrathin, high-quality, films. Indeed, in the following the <1 nm films grown by ALD are shown to be homogeneous enough to act as quality tunnel barriers.

These ultrathin electron-transparent tunnel barriers grown by O2-ALD are then integrated into 1 μm² vertical graphene-coated Ni/Al2O3/Co magnetic tunnel junctions. The tunnel junctions are contacted by a Au-capped ferromagnetic Co top electrode to probe their charge and spin transport properties (as shown schematically in Figure 4a). We first consider their
The reduction of the number of ALD layers grown by ALD as tunnel barriers.36 Thus, the simple and low-cost O3-ALD process presented here remarkably leads to well-wetted tunnel barriers on the bottom electrode with control down to extremely low thicknesses of 0.6 nm, below what is usually achievable by more sophisticated sputtering processes.37,38

We now discuss the spin transport properties of these tunnel devices. In Figure 5, a magneto-transport characterization obtained at 1.4 K is presented. Independent switching of the magnetizations of the top and bottom ferromagnetic electrodes allows the observation of two resistance states, revealing spin-dependent tunneling across the ALD layer. The large magneto-resistance (MR) measured confirms that the graphene passivation layer preserves a spin polarization for the bottom electrode following the ALD step. In contrast to unsuccessful previous studies where ALD was performed directly on top of ferromagnetic metals,15–17 here we show that the graphene layer prevents the ferromagnet from oxidizing while allowing a spin-polarized current to be extracted. We obtain up to MR = (Rap − R0)/Rap = −31% from the device characterization presented in Figure 5, where Rap and R0 are the device resistances in the parallel and antiparallel state, respectively. The MR can be analyzed to extract the spin polarization Ptop and Pbot of the top and bottom interfaces with MR = 2PbotPtop/(1 + PtopPbot), following ref 39. Taking the largest value of the Co/Al2O3 spin polarization we measure in reference devices40 (Ptop = +32%), we derive a large Pbot = −42% for the air-exposed graphene-coated nickel/alumina. This negative sign was expected to arise from the nickel/graphene interface.24,41 Strikingly, this is comparable in amplitude to the maximum reported P = +46% value difficultly achieved for the highest quality Ni/Al2O3 interfaces.21 Hence this corresponds to a practically full inversion of the spin polarization vs graphene-free Ni/Al2O3 spin sources. The ability to achieve high spin signal amplitude from a Ni/Al2O3 source is remarkable when considering how difficult it has been to raise its spin polarization above +10% in state of the art tunneling experiments21,42–44 and that the surface here has been exposed to air in this simple ex situ process. We note that the precleaning inherent to the O2-ALD process used is able to remove contaminants from the air-exposed electrode. Another possible origin of this large MR signal could be the homogeneity of the Al2O3 layers grown by ALD. Indeed, high-quality sputtered Al2O3 tunnel barriers are usually produced by oxidizing a thin metallic Al film. This results in a gradient in the extent of oxidation through the layer while tunnel barriers grown by ALD are oxidized layer by layer, providing a more homogeneous material.12 Following from the work presented herein, the layer-by-layer ALD growth of other tunnel barriers such as MgO can be foreseen. This may ultimately lead to higher spin polarization and drastically reduced resistances.

CONCLUSION

The ability to produce conformal, high-quality dielectrics in a layer-by-layer manner, combined with its notable simplicity and low cost, has led ALD to supersede other deposition techniques in many advanced microelectronics applications. Still, despite several attempts, its use has not been widely adopted for magnetic tunnel junctions where an industry-compatible alternative to sputtering has remained elusive. By demonstrating the fabrication of functional spin valves based on ultrathin and efficient ALD tunnel barriers, we unveil here the potential of ALD for spintronics and more particularly magnetic tunnel junctions. While this...
METHODS

Ni electrodes, 10 μm wide, are defined by e-beam lithography, evaporation of 150 nm of Ni, and subsequent lift-off. Reduction and graphene passivation of the Ni surface are achieved by a CVD process22–24 performed in a custom-built cold-wall reactor whose base pressure is 5 × 10⁻⁷ mbar. Samples are heated in a 1 mbar atmosphere of H₂ to 600 °C at 300 °C/min and annealed for 15 min. The H₂ is removed, and then the samples are exposed to a 10⁻⁷ mbar atmosphere of C₂H₂ at 600 °C for 15 min. Finally, the samples are cooled in a vacuum at ~100 °C/min. This growth results in a few-layer graphene (2–5 layers) coating, which prevents the oxidation of the Ni surface as checked by in situ XPS measurements.24 Complete spin valves are then fabricated using ALD tunnel barriers. Windows of 1 μm² are opened in a UVIII resist layer above the electrode by a second e-beam lithography step. Al₂O₃ layers of different thicknesses (0.6, 0.8, 1 nm) are grown by ALD using a Cambridge Nanotecnah System. The junctions are then contacted by a Co/(15 nm)/Au(80 nm) top electrode sputtered through a shadow mask to prevent bonding pads.

ALD is performed with the growth chamber temperature (T₀) held at 80 °C, low enough to promote adsorption and quench ozone reactivity with graphene, yet high enough to enable complete ALD reactions. Ozone (500 Torr, 20 s purges) and trimethylaluminum (1 Torr, 60 s purges) pulses are sequentially carried out with a growth rate of 1 Å per cycle. Before the growth, the graphene layers are exposed to a 500 Torr ozone atmosphere for 60 s to promote adsorption and subsequent homogeneous nucleation. CVD-like reactions in the chamber during the ALD process are prevented as checked by ellipsometry and breakdown measurements in nm² test structures. The quenching of the reactivity of the complete 80 °C ozone process with the graphene coating is provided by Raman spectroscopy with a Renishaw InVia system using a 532 nm laser. The roughness of the sputtered and ALD-grown tunnel barriers on graphene is characterized using AFM in tapping mode.

Conflict of Interest: The authors declare no competing financial interest.

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REFERENCES AND NOTES


