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Assessment of candidate metallization systems deposited on diamond using nano-indentation and nano-scratching tests

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A B S T R A C T

Mechanical suitability of ohmic contacts among the select metallization systems, deposited on a p-type heavily boron-doped homoepitaxial diamond layer, was evaluated via mechanical tests on the nanoscale. Two candidate metallization systems were considered: Si/Al and Ti/Pt/Au. Metallizations were performed using two different techniques: plasma-enhanced chemical vapour deposition and “lift-off”. Effectiveness of the techniques was assessed via mechanical tests on the microscale and the nanoscale. Nano-indentation experiments were performed to determine the mechanical properties of the layers. Nano-scratching experiments were used to evaluate the mechanical adhesion on the diamond substrate. Scanning electron microscopy was applied for observation of the morphology of the surface and the indent and for detecting defects.

Keywords:

Diamond
Metallization
Deposition
Nano-indentation
Nano-scratching
Power electronics

1. Introduction

Diamond is well known as a promising material for power electronics due to its desirable electrical and thermomechanical properties [1–3]. Nevertheless, many technological issues must be resolved to adequately exploit diamond and to manufacture competitive power electronic devices that can withstand high temperatures of up to 300 °C. This study focuses on diamond metallization systems that play the role of ohmic contacts. The metallization systems must satisfy many conditions, such as having the ability to withstand high voltages and current density, good mechanical properties, and good adhesion on the diamond substrate, among other requirements.

Many researchers are interested in the deposition of ohmic contacts on diamond based on the premise that good mechanical adhesion of the metallization systems is a key requirement for achieving efficient device bonding and power transmission. Refractory metals such as Ti, Mo, Ta, and W are known to form adhering carbide layers on diamond at high temperatures [4–6]. Ti-based contacts are more widely used due to the formation of a TiC layer during annealing, which enables achievement of low contact resistivity and good mechanical adhesion. Moreover, transition metals such as Cr, Ni, and Ni—Cr are widely used in other multi-chip module (MCM) technologies involving Si, AlN, SiC, and alumina substrates [7]. The mechanical and electrical performances of Cr, W, Ni, and Al layers as frameworks have been investigated and

documented in the literature [8]. The ohmic contact metallization systems evaluated in the studies mentioned above were deposited on a diamond sample by the same photolithography process. It was noted that Ni is a suitable contact metallization system for utilization in high power, high temperature, and good mechanical strength diamond Schottky barrier diode applications. Nevertheless, the mechanical adhesion and thermal stability of these species was not investigated. The electrical capability and thermal stability after post-deposition annealing of metallization systems such as C/Ti/Ni, C/Ti/Cr, and C/Ti/W have previously been investigated. Under the relevant deposition conditions of these studies, it was concluded that the adhesion of Ti/Ni to diamond is higher than that of single layer Cu due to the formation of TiC, and that only Cr is inert to AuSn during annealing [9]. Cr also offers better adhesion and enhances the diffusion barrier properties (compared with Ti—W) for Au-based conductor metallization systems on diamond substrates [7]. From the latter observation, it can be deduced that both the die attachment and the diffusion mechanisms are highly important for the thermomechanical stability of the assembly involving diamond. The deposition technique also appeared to be of great importance. The adhesion of sputtered Au/(Ti—W) to diamond was significantly increased when the surface was activated by the sputter etch technique [7]. The electrical performance and intermetallic reaction of a Ti/Pt/Au multilayer after annealing at 800 °C and 900 °C under vacuum was also studied [10]. It was found that Au and Pt inter-diffused, but the Pt barrier was not fully consumed and effectively served as a barrier to Ti diffusion through the Au layer to the contact surface. Notably, some frameworks have been appended to Si/Al based contacts, leading to good electrical characteristics of such systems [11]. However, overall,

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concrete advancements on the study of adhesion of such deposited films on diamond are rare.

Due to the lack of data concerning the mechanical and the adhesion characteristics of metallic films on diamond substrates, herein, we study some candidate metallization systems for use as ohmic contacts on diamond for high temperature power electronics. The ohmic contacts studied here are Ti/Pt/Au and Si/Al systems. The use of a thick Si or Ti layer allows the formation of a SiC or TiC carbide layer, which ensures strong bonding. This latter point is essential if high reliability of the diamond device packaging is required. Surface morphology analyses are performed using scanning electron microscopy (SEM) and the energy dispersive X-ray (EDX) technique to detect surface defects derived from deposition or annealing. Nano-indentation tests with large (XP) and small (DCM) displacements are performed to determine the elastic properties and hardness of the deposited films. Finally, the adhesion of the multilayers on the diamond substrate is evaluated using nano-scratching tests.

2. Experimental procedure

2.1. Deposition of thin films

A1.5 mm thick, heavily boron-doped diamond film deposited on the $3 \times 3 \times 0.5$ mm lb. diamond sample used in this study. The boron density in the film was approximately $3 \times 10^{20} \text{ cm}^{-3}$. The sheet resistance of the sample was measured by four probe measurement to be about $110 \text{ } \Omega/\square$. Various metallization systems were deposited on the diamond sample by vacuum evaporation. For Si/Al metallization, a $0.2 \text{ } \mu\text{m}$ Si layer was first deposited at $475 \text{ } ^\circ\text{C}$ by plasma-enhanced chemical vapour deposition (PECVD). The metallized substrate was then annealed in a rapid thermal processing (RTP) furnace at $1200 \text{ } ^\circ\text{C}$ under vacuum for 10 s prior to cleaning with hydrofluoric acid to remove any oxides on the surface, including the native SiO_2 . The purpose of this operation is to form a silicon carbide layer at the interface between silicon and diamond. This layer was used to help to ensure good adhesion of the layer stack on diamond. Upon completion of this operation, a $0.5 \text{ } \mu\text{m}$ thick Al film was formed by the photolithography technique or by “lift-off”. This step was followed by annealing at $450 \text{ } ^\circ\text{C}$ for 20 min in a tubular passage furnace to improve the adhesion of the metallization system.

On the second diamond substrate, a Ti/Pt/Au multilayer was formed by “lift-off”, where the multilayer comprised respective layers with thicknesses of 0.1, 0.05, and $0.5 \text{ } \mu\text{m}$. Once these films were obtained, the sample was annealed at $450 \text{ } ^\circ\text{C}$ for 30 min. This is the minimum temperature at which Ti and Si form a carbide layer with diamond [11].

2.2. Mechanical characterization

2.2.1. Nano-indentation tests

The mechanical properties of the layers, such as the stiffness and hardness, were determined by using an MTS© nano-indentation device equipped with a Berkovich pyramidal indenter. This type of device can continuously measure the stiffness of the contact via “Continuous Stiffness Measurement”. Nano-indentation tests with small and large displacements were carried out at room temperature ($23 \text{ } ^\circ\text{C}$) depending on the thickness of the deposited films. The test conditions are specified in Table 1. For large displacements, the tests were performed with an MTS XP nano-indenter in dynamic mode, making it possible to obtain the overall Hardness and Young’s Modulus values of the multilayer. However, the DCM nano-indenter was utilized for small displacement (400 nm). The DCM nano-indenter permits determination of the individual properties of the films with quite good precision. The mechanical properties of the layers, such as the hardness and elastic modulus, were determined via the common Oliver-Pharr method [12]. For more details about the nano-indentation technique, one can refer to papers by Alexits or Msolli and co-workers [13–15], for instance.

Table 1

Operating conditions for nano-indentation tests.

Berkovich tip nature	DCM	XP
Load resolution	1 nN	100 nN
Column displacement resolution	0.05 nm	0.1 nm
Number of tests	30	10
Displacement into the surface	400 nm	2 μm
Approach velocity	8 nm/s	8 nm/s
Distance between nano-indents	10 μm	50 μm

2.2.2. Nano-scratching tests

The adhesion of the deposited multilayers was also assessed using nano-scratching tests. The test parameters were adapted depending on the thickness of the metallization layer (see Table 2). Nano-scratching tests were performed on the sample surface by moving a Berkovich tip in three successive steps and measuring the penetration depth. The first step was carried out to determine the surface topography and to correctly set the position of the indenter. The aim of the second step was to scratch the sample surface. The applied load was then increased progressively up to a maximal value and subsequently reduced. Finally, the spring-back and the topography of the residual scratch were measured from a last scan using a small applied load.

2.2.3. Morphology and EDX analysis

Observations were carried out with a field emission gun scanning electron microscope (FEG SEM-7000F from JEOL with the incident electron beam maintained between 10 and 15 kV) to analyse the surface morphology of the films. The analysis was corroborated by chemical analysis with an SDD Bruker X flash energy dispersive X-ray spectrometer (EDX).

3. Study of Si/Al metallization system

3.1. Morphology of the surface

Fig. 1 shows micrographs obtained by scanning electron microscopic (SEM) analysis of the metallization surface at various magnifications ranging from $\times 500$ to $\times 50,000$. The Si/Al multilayer did not have any detectable defects. The Si/Al multilayer appeared relatively dense and homogeneous at low magnification. Observations at higher magnification revealed the presence of clusters of a few microns, evenly distributed on the sample surface (Fig. 1a). The microstructure of the deposited Al comprised equiaxed grains with diameters of 100 nm (Fig. 1b).

EDX spot analyses were carried out on the metallization surface. The overall spectrum attests the presence of two layers of aluminium and silicon in region 3 (Fig. 1a). Indeed, the “interaction pear” of the SEM incident beam has a depth greater than the deposition thickness of aluminium, which also explains the detection of the sub-Si layer. However, the previously observed clusters denoted as region 4 (Fig. 1a) are those of aluminium because the Al content is higher than the Si content.

Table 2

Operating conditions for nano-scratching tests.

Distance between scratches	500 μm
Scratch length	700 μm
Scratching velocity	10 $\mu\text{m/s}$
Topography load	0.1 mN
Scratching load	0.1 mN to 80 mN
Final load	0.1 mN
Number of scratches	2 – 3

Fig. 1. Si/Al surface micrographs obtained by SEM observation at different magnifications. (a) 6000 and (b) $\times 50,000$.

3.2. Mechanical properties of Si/Al system

3.2.1. Nano-indentation with MTS XP indenter

To obtain a scatter in the data, ten nano-indentation tests were performed with an imposed depth of 2 μm . Nano-indentation tests at large displacement (where the maximum load reached 1500 mN) were useful for determining the variation of the mechanical properties of the Si/Al multilayer with the indented depth. Fig. 2a shows the load evolution as a function of the penetration into the surface for all the tests. Good repeatability of the load displacement responses was observed. To facilitate interpretation of the curves, the nature of the different layers that were reached and their thicknesses are shown directly on the graph. Each load curve can be divided into three distinct parts.

The first part appears linear and is located between the loading start and a penetration depth close to 500 nm. It is characterized by the lowest slope for the entire load curve. This is explained by the fact that the stiffness and hardness of Si and Al are much smaller than those of diamond. The second portion corresponds to a transition and ranges between 500 nm and 1300 nm, where the influence of diamond becomes more important. Over 1300 nm, the influence of metallization becomes negligible and the applied load increases sharply due to the very high hardness of diamond.

Fig. 2b shows the average stiffness and hardness curves as functions of the indentation depth. The Young's modulus increased progressively from the start of the response to reach a plateau at 750 GPa from a penetration depth of 1100 nm. The highest dispersion observed at the end of the test and the slight decrease of the load, are due to initiation of cracking of the diamond substrate, which begins from an indentation depth of 1500 nm. This crack is confirmed by the flat portions present on the load displacement curves (see Fig. 2a).

As shown in the same figure, the hardness evolved differently, where it increased very slightly up to a penetration depth of 600 nm. This was followed by a rapid increase above 600 nm to reach a value of 25 GPa. These initial results do not permit determination of the stiffness and hardness of each layer, considering the influence of the

diamond substrate at this stage. It is interesting to note that the influence of the diamond substrate on the stiffness calculation emerges faster (even at small loads) than on the hardness determination. This has previously been observed for other coated substrates [16]. The hardness and stiffness of the diamond substrate appear to be respectively equal to 25 GPa and 750 GPa. Other studies of the mechanical properties reported values for thick diamond coatings deposited on WC substrates close to that found herein [17,18].

The nano-indenters relevant for these curves were observed by scanning electron microscopy (SEM) to confirm the previous analyses. Fig. 3 shows the morphology of the nano-indent and the indented layers. Microcracks were noticeable in the diamond substrate (Fig. 3a). The nano-indent had a notable "barrel-shaped" form on its edges, resulting from incompressible plastic deformation and the up-flow of the aluminium layer around the indenter. This "pile-up" against the indenter faces is characteristic of work-hardened materials such as aluminium. Assessment of the pile-up is important because pile-up leads to differences between the true contact area and the apparent contact area, which is usually observed after indentation. In this case, the indented material appears to be stiffer. The advantage of continuous stiffness measurement (CSM) is that it takes into account the instantaneous variation of the contact area by use of particular values of some adjusting parameters (see Ref. [19] and [20] for the meaning and details of the contact area calculations). During testing, all the layers are crossed, as shown by EDX mapping (see Fig. 3b and EDX profilometry on bottom of the nano-indent; see Fig. 4).

3.2.2. Nano-indentation with DCM indenter

The large displacement nano-indentation tests did not permit determination of the intrinsic mechanical properties of the aluminium layer. Only the properties of the diamond substrate were estimated. Therefore, small displacement nano-indentation tests were carried out with the dynamic control module (DCM) nano-indenter MTS that offers better resolution even at small applied loads due to its lower stiffness.

Fig. 2. Nano-indentation data for the Si/Al bilayer using XP tip. (a) Loading/unloading curves and (b) variation of hardness and Young's modulus against penetration into surface.

Fig. 3. Nano-indent obtained from nano-indentation test on Si/Al metallization system; (a) SEM micrograph of the deformation morphology showing cracks in diamond and (b) EDX mapping showing the composition of the multilayer.

Fig. 5a shows the load evolution with the imposed penetration depth of the indenter (the maximum depth was 400 nm). The load profiles do not show any discontinuities despite the significant dispersion, mainly due to the higher sensitivity of the DCM tip to the surface roughness. The spring-back was very small (<20 nm). This is mainly due to the small yield strength of Al, associated with its significant plastic deformation. Since DCM type indenter is highly sensitive, creep displacements are observed during the holding load segment. Indeed, during the hold, the indenter continues to penetrate into the surface. Subsequently, the area of the indent increases. For the studied metallization, the displacement creep reaches 40 nm at room temperature.

The curves of the average hardness and elastic stiffness module as a function of the penetration depth are shown in Fig. 5b. The stiffness was stabilized at 70 GPa for indentation depths ranging from 40 to 60 nm, which is just one-tenth of the Al film thickness. The influence of Si and diamond was small at this depth. However, over 80 nm, the measured stiffness increased progressively because of the influence of the diamond substrate. Therefore, the stiffness and hardness of the Al layer could be estimated as 70 GPa and 1 GPa respectively. Other studies provide reference values consistent with the modulus values found herein. The hardness values also remained very close to bibliographical results [21].

3.2.3. Nano-scratching tests

The adhesion of the Si/Al bilayer was evaluated via nano-scratching tests according to the experimental conditions described in Table 2. The penetration depth and the dimensionless friction coefficient were determined with respect to the scratch distance (Fig. 6).

The nano-scratching tests provide comparable and reproducible profiles regardless of the location of the preformed scratches (Fig. 7). The behaviour was initially linear with a constant penetration rate of $-2 \text{ nm} \cdot \mu\text{m}^{-1}$. This behaviour is mainly due to the ductile and rather hard Al layer. Once the diamond substrate was reached, an abrupt

change of the penetration rate occurred, but the profile remained linear. For depths below 500 nm, the topographic profile of the final scan matched the penetration profile during scratching. This indicates that the Al layer undergoes substantial plastic deformation as during the indentation tests. Over the stated depth, the topographic profile becomes different from that measured during scratching. The gap between the two profiles could reach a maximum of about 160 nm at the end of scratching, which corresponds to the diamond elastic spring-back that represents 20% of the total strain at maximum load.

The evolution of the friction coefficient as a function of the penetration distance is shown in Fig. 6b. The friction coefficient, μ , is the ratio of the lateral versus the normal force imposed by the indenter. This coefficient increased sharply when the tip began to penetrate the Al layer, reaching a value of 0.3 at a scratching distance of 180 μm (penetration depth equal to -304 nm, normal load equal to 12.92 mN). At this normal load, a slight perturbation of the friction coefficient was observed, indicative of different mechanical behaviour due to the removal of Al chips during scratching. Thus, the friction coefficient continued to increase to reach a value of 0.5 when the indenter reached the Si/Al interface. Finally, the friction coefficient began to fall and stabilize at a value of around 0.23 at the end of the test when the tip reached the diamond substrate.

Fig. 7 shows the morphology of the two scratches formed on the Si/Al bilayer. From the beginning of the scratching process, the Al film underwent plastic deformation due to its low hardness, with “bead” formation on the edges. At the end of the scratching, initial formation of chips was observed due to the cohesive failure of Al. This damage confirms the evolution of the recorded friction coefficient and the abrupt topography variations. No delamination or chip formation was observed when the tip passed through the silicon layer, evidencing therefore, the good adhesion of this layer to diamond. This result was also confirmed by the gradual evolution of the friction coefficient when the tip passed through the Si layer and scratched the diamond.

Fig. 4. Chemical analyses using EDX profilometry of the nano-indent formed on Si/Al multilayer. (a) Morphology of the indent and the direction of analysis and (b) compositions along the chosen analysis direction.

Fig. 5. Nano-indentation data for the Si/Al multilayer using DCM-type tip. (a) Loading/unloading curves and (b) average hardness and Young's modulus against penetration into surface.

4. Study of Ti/Pt/Au metallization system

4.1. Morphology of the surface

The micrographs acquired at low magnification show defect repartition over the entire sample surface (Fig. 8). These defects are either blisters with diameters varying between 5 and 30 μm or peeling of the metallization species. From the micrographs obtained at higher magnification, the average grain size of the Au film was determined as 2 μm . These defects appear to be generated during the deposition process. Blisters often result from the presence of residual stresses in the layers and/or brittle interfacial areas.

In order to determine the location of these defects, EDX analyses were performed. The composition of the layers and origin of the defects on the surface were determined by chemical analysis of two characteristic zones of the sample surface: a defect-free zone (region 1) and an area corresponding to peeling (region 2) (Fig. 8).

Chemical analysis of region 1 shows the presence of a gold layer on the substrate surface. The other layers were not detected due to their low thickness with respect to the gold layer. Conversely, chemical analysis of region 2 revealed only the presence of carbon, corresponding to diamond. Peeling seemed to be localized between Ti and diamond or between Pt and Ti. Several EDX analyses were performed in the peeling areas using incident electrons at small acceleration voltages (~ 5 kV). The aim was to analyse thin layers while maintaining a correct statistical measurement by performing long acquisitions. For the analyses carried out in the peeling areas, only carbon was highlighted. However, it is difficult to precisely decide on the peeling area in view of the low thicknesses of the Ti and Pt layers. During the deposition process, non-negligible compressive residual stress was produced in the Ti/C interface. When the stress in the metallization system is compressive, buckling-driven interface delamination occurs, as observed for the Ti/Pt/Au metallization system.

4.2. Mechanical properties of Ti/Pt/Au system

4.2.1. Nano-indentation with MTS XP indenter

Fig. 9a shows the corresponding loading-unloading curves for tests performed on the diamond substrate with the Ti/Pt/Au metallization

system. The test results were reproducible. The general shape of the curves was similar to that of those already described for Si/Al metallization on diamond. The load increased slightly in the first stage due to the low stiffness and hardness of gold. As previously observed, no influence of the intermediate layers of Ti and Pt was observed because of the thinness of these layers. The load increased rapidly to reach a maximum depth of 1600 nm at the end due to the high stiffness of the diamond substrate. The elastic spring-back represents nearly 50% of the total penetration distance. This is higher than that observed for Al/Si metallization. The present value is mainly due to the spring-back of the diamond substrate.

Fig. 9b shows the change in the stiffness and hardness of the system as a function of the penetration depth. Due to the progressive increase of the depth from gold to diamond, the stiffness increased to reach a maximum value of about 750 GPa, which is greater than that found for the Si/Al metallized substrate. This value corresponds to the Young's modulus of diamond. From this depth (600 nm), the stiffness decreased considerably with many perturbations, the origin of which is certainly related to the appearance of cracks within the diamond substrate itself, as seen above. However, the hardness continued to increase to a maximum average value of 26 GPa, which is very close to the value determined for the Si/Al metallized substrate.

The observation of the nano-indent morphology shows the presence of three cracks (Fig. 10a). These cracks appeared on the faces that were in contact with the indenter and propagated to the edges (~ 3 μm). These cracks were separated from the nano-indent centre by 500 nm. All of the obtained nano-indents were characterized by these cracks on their inner faces. Mapping showed that the apparent surface in the centre was predominantly that of diamond. Due to its low thickness, the Pt layer (0.05 μm) was not detectable.

Pile-up occurred during plastic deformation of the ductile Au, and superposed plies were observed at the crater edges. This is expected due to the strain hardening nature of Au. The pile-up tends to increase the contact area of the indenter and to enhance the gradient of plastic deformation.

The chemical composition of the nano-indent throughout the section was determined as shown in Fig. 11. These chemical analyses confirm the location of the cracks in diamond based on consideration of the size of the zone in contact with diamond.

Fig. 6. Nano-scratching data for the deposited Si/Al bilayer. (a) Depth profiles and (b) variation of the friction coefficient during the scratch test for Si/Al bilayer.

Fig. 7. SEM Micrographs showing the morphology of the scratches formed on Si/Al bilayer.

Fig. 8. Ti/Pt/Au surface micrographs obtained by SEM observations at different magnifications. The micrographs show the presence of several blisters of constant size over the surface and peeling occurred due to compressive stress.

4.2.2. Nano-indentation with DCM indenter

Nano-indentation tests at small displacements were carried out to characterize the intrinsic properties of the surface gold layer (Fig. 12a). The behaviour of the metallization species was almost plastic given the very small spring-back measured during unloading. In fact, this spring-back is too small compared to the maximum penetration depth. In addition, creep displacement is produced at the hold but it is not as significant as the one observed for the Si/Al metallization. Displacement creep is not exceeding 15 nm.

Fig. 12b shows the changes in the average mechanical properties as a function of the penetration depth. A plateau was logged for a depth of 40 nm. This plateau corresponds to the intrinsic properties of the gold layer; the stiffness was nearly 80 GPa and the hardness was close to 1 GPa.

4.2.3. Nano-scratching tests

The scratching profiles obtained during the tests performed on the Ti/Pt/Au multilayer are shown in Fig. 13a. The profile indicates that the penetration depth increased rapidly until the end of the scratch when it reached the diamond substrate. The profile of the "post scan" shows that the spring-back started very early during scratching and corresponded to that of the entire metallization system with diamond. This spring-back was very high compared to the one obtained for the Si/Al metallization system and represents more than the half of the total penetration depth at the end of the test. However, the profile of the "post scan" became irregular and fluctuated in the middle of the scan. SEM observations are required to understand these variations.

The friction coefficient increased rapidly to 0.6 when the indenter crossed gold and platinum, then decreased and stabilized at 0.25 when the diamond substrate was reached (Fig. 13b).

The morphology of the scratches formed on the Ti/Pt/Au multilayer is shown in Fig. 14. The beads on the edges of the scratch were obtained due to the ductility of gold. Peeling of the multilayer occurred from a scratch distance of 160 μm, even before the tip reached the diamond substrate. At this distance, the indenter was at a depth corresponding to the thickness of Ti for a critical scratch load of about 25 mN. These variations appeared due to the deformation of the multilayer, and particularly due to the mechanically affected Ti/C interface. At a scratch distance of over 230 μm, the ductile fracture continued to the end of the scratch.

5. Conclusion

The Si/Al metallization system had a uniform morphology with some Al clusters evenly distributed on the surface. The Al surface comprised grains with uniform diameters. Changes in the stiffness and hardness of the bilayer against the penetration depth were determined by a nano-indentation test. The mechanical properties of aluminium ($H_{al} = 1$ GPa and $E_{al} = 70$ GPa) and diamond ($H_C = 25$ GPa and $E_C = 750$ GPa) were determined. It was not possible to determine the mechanical properties of the Si layer given its low thickness and because this layer was sandwiched between two materials with strong mechanical properties. A significant creep displacement was noticed at room

Fig. 9. Nano-indentation data for the Ti/Pt/Au multilayer using XP-type tip. (a) Loading/unloading curves and (b) average hardness and Young's modulus against penetration into surface.

Fig. 10. Nano-indent obtained from nano-indentation test performed for Ti/Pt/Au metallization system: (a) SEM micrograph of the deformation morphology and (b) EDX mapping showing the composition of the multilayer.

temperature during the hold time. This creep displacement attests of a viscoplastic behaviour of the metallization system. Moreover, due to the ductile nature of Al, a pile-up phenomenon was observed on the edges of the indent leading to significant plastic deformation. Nano-scratching tests showed that the failure in aluminium is cohesive; the interface strength can therefore be taken as being greater than the shear strength of aluminium. The silicon layer appears to adhere well to the diamond substrate due to formation of a SiC interphase at the interface during heat treatment, as previously documented in many studies.

Unlike Si/Al metallization systems, the diamond substrate with the Ti/Pt/Au metallization system has many defects distributed on its surface, formed by blistering or peeling. EDX spectrometric analysis indicated that peeling of the layers possibly occurred at the Ti/C interface, involving a buckling-driven interface delamination mechanism. High load nano-indentation tests revealed the presence of cracks on the nano-indents. These cracks initiate within the diamond substrate and

propagate to the metallization system, as shown by EDX spectrometric analysis and indent micrographs. The intrinsic mechanical properties of the gold layer and diamond were determined as $H_{Au} = 1$ GPa, $E_{Au} = 80$ GPa, $H_C = 26$ GPa, and $E_C = 750$ GPa. Unlike the Si/Al metallization system, Ti/Pt/Au metallization system exhibits small creep displacement. Nano-scratching tests reveal the brittleness of the Ti/C interface, indicated by the formation of several chips on the scratch edge when the tip reached the Ti layer.

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Fig. 11. Chemical analyses using EDX profilometry of the nano-indent formed on Ti/Pt/Au multilayer. (a) Morphology of the indent and the direction of analysis and (b) compositions along the chosen analysis direction.

Fig. 12. Nano-indentation data for the Ti/Pt/Au multilayer using DCM-type tip. (a) Loading/unloading curves and (b) variation of hardness and Young's modulus against penetration into surface.

Fig. 13. Nano-scratching data for the deposited Ti/Pt/Au multilayer. (a) Depth profiles and (b) variation of the friction coefficient during the scratch test.

Fig. 14. Micrographs of scratch formed on Ti/Pt/Au multilayer: morphology of the scratch.

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