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A Nanoindentation Study of the Fatigue Properties of

Al/a-Si Core-Shell Nanostructures

A Nanoindentation Study of the Fatigue Properties of Al/a-Si Core-Shell Nanostructures

An Undergraduate Honors College Thesis

in the

Department of Mechanical Engineering College of Engineering University of Arkansas Fayetteville, AR

by

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ABSTRACT

Nanostructure-textured surfaces can reduce friction and increase the reliability of micro- and nanoelectromechanical systems (NEMS/MEMS). For MEMS incorporating moving parts, the fatigue properties of nanostructures pose a challenge to their reliability in long-term applications. In this study, the fatigue behavior of hemispherical Al/a-Si core-shell nanostructures (CSNs), bare hemispherical Al nanodots, and a flat Al/a-Si layered thin film have been studied using nanoindentation and nano-scale dynamic mechanical analysis (nano-DMA) techniques. Fatigue testing with nano-DMA shows that the deformation resistance of CSNs persists through 5.0×10^4 loading cycles at estimated contact pressures greater than 15 GPa. When the a-Si shell is removed, as in the Al nanodots, significant nanostructure deformation results due to repeated cyclic loading. In addition, for the Al/a-Si layered thin film, which lacks the geometry and core-confinement properties of CSNs, cyclic loading results in fatigue failure of the a-Si layer. CSNs demonstrate none of the failure mechanisms exhibited by these control structures. The unique properties displayed by CSNs when subjected to fatigue testing establish their prolonged durability when implemented in micro- and nano-scale applications.

KEY WORDS: core-shell nanostructure; nanoindentation; fatigue; dynamic mechanical analysis; continuous stiffness measurement

NOTE: This thesis, for the fulfillment of the requirements for the Mechanical Engineering Honors Program, is submitted in a journal format.

1. Introduction

Over the past decade, advancements in nanotechnology have allowed for the development of micro-/nano-electromechanical systems (MEMS/NEMS), which show great promise in nearly every product category. The applications of MEMSs range from aerospace to health care, and are commonly implemented as accelerometers [1], sensors [2], and magnetic storage devices [3]. However, when systems are designed at the micro- and nano-scale, the van der Waals force, capillary force, and electrostatic force are accentuated by the large surface-to-volume ratios compared to macroscale systems [4,5]. This results in a vulnerability to several failure mechanisms, including adhesion, stiction, and wear [4-7].

It is well understood that surfaces patterned with nanostructures, known as nanotextured surfaces (NTSs), significantly reduce adhesion and friction between the contact surfaces of many materials, and thus provide a solution to the current challenges of MEMS [8-14]. However, the individual nanotextures composing NTSs are susceptible to permanent deformation and fatigue failure under the contact stresses typically experienced in micro-scale devices [15-17]. For example, while microscale friction testing of Ni nanodot patterned surfaces showed a reduction in adhesion and friction, significant deformation was seen after testing [15,16]. This behavior has also been seen in Al nanotextured surfaces [14] and amorphous carbon surfaces [17]. Deformation and fracture in turn diminish the tribological properties initially provided by the textured surface.

Fatigue is of particular interest to the reliability of MEMS incorporating moving parts, since repetitive loading may lead to failure as a result of crack growth and the accumulation of plastic deformation [6,7,18-22]. For example, single crystal silicon is the most predominantly used material in MEMS/NEMS due to its desirable electronic properties and mechanical resilience; however, research shows that at the micron-scale it has inferior wear and adhesion characteristics, as well as poor fatigue properties [22-25]. If NTSs are to be successfully implemented in micro-scale devices, the individual nanotextures must also not be susceptible to fatigue damage or wear.

Core-shell nanostructures (CSNs) composed of a soft Al core within a hard a-Si shell have been discovered to be highly deformation resistant, in addition to possessing superior tribological properties when implemented as deformation-resistant nanotextures [14,26]. Several studies have been performed to gain fundamental understanding of the mechanisms that contribute to their deformation resistance [27-29]. The sample surface consists of patterned hemi-spherical Al nanodots with 100 nm – 300 nm diameters on a silicon substrate. A 300 nm thick conformal a-Si shell is then deposited on the Al nanodot-patterned surface, forming core-shell nanostructures of a soft Al inner core encompassed by a hard a-Si shell. This nanotextured surface showed no permanent deformation when subjected to nanoscratch testing, with a maximum applied normal load of 8,000 μ N with a 100 μ m diamond tip. In addition, these structures have displayed superior resilience in nanoindentation testing, being subjected to high contact pressures with no residual deformation. However, their nanomechanical properties and deformation-resistance have not been studied beyond 4 loading/unloading cycles per structure and their structural integrity under prolonged repetitive loading has not been determined.

Nano-scale dynamic mechanical analysis (DMA) is a recently developed method of nanoindentation for characterizing visco-elasticity and fatigue life at the nano-scale [30,31]. By

superimposing a sinusoidal load-oscillation upon a nominally increasing indenter load, this technique allows for continuous measurement of the material properties of a sample. It has been shown previously that contact stiffness is sensitive to damage formation, and that continuous measurement of contact stiffness allows for precise characterization of a material's fatigue behavior [20]. This method has been used to determine the fatigue properties of Si nanobeams, Cu thin-films at the nano-scale, and amorphous carbon coatings [20,21,31]. In each study, an abrupt change in the contact stiffness indicated that fatigue damage had occurred.

In this paper, the method of nano-scale DMA is developed for studying the fatigue behavior of individual core-shell nanostructures. The fatigue properties of deformation-resistant Al/a-Si CSNs are then compared to the response of Al nanodots and flat Al/a-Si layered thin films subjected to repetitive loading. Scanning electron microscope images of the structures and surfaces are presented to better understand the mechanisms of failure, as well as to establish the deformation response of core-shell nanostructures under extended cyclic loading.

2. Experimental

2.1. Nanostructure Fabrication

Arrays of Al nanodots were fabricated using electron beam lithography (EBL) and a metal lift-off procedure. First, a positive tone electron resist, 495k MW PMMA at 4% dilution, was spin coated onto a (100) crystalline Si wafer at 3,000 rpm. An electron beam writer (JBX-9300FS, JEOL Ltd.) was used to selectively expose the electron resist, using a 50 kV accelerating voltage, 1 nA of current, and a beam

dose of 1,000 μ C/cm². The patterned arrays were then developed in a 1:3 mixture of methyl isobutyl ketone (MIBK) and isopropyl alcohol (IPA) for 45 s, and rinsed with pure IPA for 15 seconds. This resulted in patterned arrays of holes in the PMMA film, and was followed by a low-pressure oxygen plasma etch to smooth the edges of the holes. Next, 100 nm of Al was deposited onto the patterned PMMA film using thermal evaporation (Auto 306D, Edwards Vacuum) at a rate of 0.4 nm/s. Finally, the remaining PMMA was removed by immersion in a Remover PG (MicroChem Corp.) bath heated to 75° C for 48 hours. Using this fabrication method, ordered arrays of Al nanodots with base diameters of 100 nm and 300 nm were fabricated, each 100 nm in height.

The patterned arrays of 100 nm diameter Al nanodots were then coated with a-Si using plasma enhanced chemical vapor deposition (PECVD; Plasma-Therm SLR730). The rf power, substrate temperature, and silane flow rate during a-Si deposition were 20 W, 250° C, and 85 sccm, respectively. Using this method, a-Si was uniformly deposited on the sample, and Al/a-Si CSNs with 100 nm base diameter Al cores and an a-Si shell thickness of 300 nm were produced.

The flat Al/a-Si thin-films were fabricated using thermal evaporation followed by PECVD. First, 100 nm of Al deposited onto a (100) single crystal Si wafer using thermal evaporation (Auto 306D, Edwards Vacuum) at a rate of 0.4 nm/s. Then, 300 nm of a-Si was deposited onto the Al film by using PECVD with an rf power, substrate temperature, and silane flow rate of 20 W, 250° C, and 85 sccm, respectively. This resulted in flat Al/a-Si thin-films with an Al layer 100 nm thick and an a-Si layer 300 nm thick. Fig. 1 shows schematics of the three fabricated nanostructure geometries.

2.2. Fatigue and Nanoindentation Tests

Nanoindentation and fatigue experiments were carried out using a TriboIndenter (Hysitron Inc.) equipped with a NanoDMA I module. The Triboindenter operates by electrostatic force actuation and measures displacements using a capacitive sensing scheme, with a force resolution of 3 nN and a displacement resolution of 0.02 nm. Within the NanoDMA I module, a lock-in amplifier is used to apply a sinusoidal dynamic load to the indenter tip concurrently with a given quasi-static force, at frequencies between 0.1 Hz and 200 Hz. The lock-in amplifier continuously measures the displacement amplitude of the indenter tip and the phase shift between the indenter and the applied signal. From this data the dynamic material properties of the sample are determined. In this study, a spherical diamond indenter tip of 1 µm radius of curvature was used, and integrated scanning probe microscopy (SPM) within the TriboIndenter was used to accurately locate and indent the individual nanostructures with the same 1 µm tip. A large indenter tip radius, compared to the size of the nanostructures, was chosen to provide compression loading to the nanostructures, rather than penetration into the samples.

Fig. 2a shows a schematic of the tests performed on the Al nanodots and Al/a-Si thin film to determine the approximate load of failure due to dynamic loading. These tests were conducted in a load-controlled mode, incorporating a constant dynamic load amplitude and a mean quasi-static load increasing linearly throughout the experiment. To maintain uniform testing parameters across all geometries, Hertzian contact theory for sphere-on-sphere contact was used to estimate the contact pressure applied to the Al nanodots at the determined critical load [32]. The indentation force which applies an equivalent contact pressure to the CSNs was then calculated and used for fatigue testing of CSNs.

Fig. 2b shows a schematic of nanoindentation fatigue tests on hemi-spherical nanostructures and flat thin films. Fatigue testing was performed by maintaining a constant quasi-static load on the sample while applying a dynamic force at a given frequency. The maximum load amplitude was set to 70% of the determined critical load, and the excitation frequency was set to 60 Hz for all experiments. Contact stiffness change was used as an indicator of damage formation, and the number of cycles was determined by the time elapsed during each test [20]. To encourage deformation of the CSNs during fatigue loading, indents with larger quasi-static and dynamic load levels were also performed.

The nanoscale surface topography and morphology of the nanostructures were characterized with a combination of scanning electron microscopy (SEM; Nova NanoLab, FEI) and integrated SPM on the nanoindenter.

3. Results and Discussion

3.1. Nanostructure Characterization and Morphology

By using EBL to fabricate the patterned Al nanodots, very uniform arrays of Al nanodots and CSNs were produced. SEM micrographs of surfaces patterned with 300 nm diameter Al nanodots and CSNs with 100 nm core diameter and 300 nm shell thickness are shown in Fig. 3. Through X-ray diffraction measurements on a similarly prepared Al film, it was determined that the Al nanodots were polycrystalline, and composed of a mixture of (111) and (200) crystallites. Fig. 4 shows SEM images of an individual Al nanodot and CSN. The PECVD process followed for depositing a-Si resulted in small-scale surface roughness in the shell and film morphologies, but due to the large size of the indenter tip

used during nanoindentation, it is expected that this will not interfere with mechanical characterization of the structures.

3.2. Ramping Load Nanoindentation

3.2.1. Al Nanodots

Dynamic indentation experiments with a linearly increasing quasi-static load were performed to characterize the response of Al nanodots to dynamic compression loading, as well as to determine the load at which structural failure occurs. Fig. 5a shows a nanoindentation loading profile as a function of time from 50 μ N to 300 μ N increasing at 0.75 μ N/s. This profile was used in conjunction with a 30 μ N peakto-peak dynamic load at an oscillation frequency of 60 Hz, such as shown in Fig. 2a, to indent a 300 nm diameter Al nanodot, while continuously measuring the contact stiffness of the nanostructure. Fig. 5b shows the contact stiffness and indenter displacement as functions of time for a 300 nm diameter Al nanodot indented with the previously defined loading profile. It is observed that a transition in contact stiffness occurs from linearly-increasing with time to scattered and increasing more gradually. In addition, it is seen that this transition coincides with a jump in displacement and a peak in contact stiffness, each occurring at a total applied load of approximately 120 µN. Fig. 5d and Fig. 5e show SPM topography and gradient images after indentation which indicate that cracking in the nanostructure has occurred. It is understood that the propagation of cracks within the nanostructure would manifest as discontinuities in the contact stiffness or displacement response, and it is strongly suggested that these correspond to the propagation of the observed cracks. Therefore, the critical load at which crack propagation occurs in a 300 nm Al nanodot due to dynamic loading was determined to be ~120 μ N.

It should be noted that two additional smaller peaks in contact stiffness with corresponding displacement discontinuities were observed at mean applied loads of approximately 70 and 95 μ N. The presence of two additional cracks in the nanostructure suggest that these signatures correspond to propagation of the second and third cracks. However, in order to investigate the most severe loading condition when applied to CSNs, 120 μ N was chosen as the critical applied load due to dynamic loading.

3.2.2. Al/a-Si Layered Thin Film

Fig. 6a shows applied load versus displacement for a quasi-static indent peaking at 8000 µN on a flat Al/a-Si thin film with 100 nm of Al and 300 nm of a-Si deposited on Al. Here it is observed that two jumps in displacement occur at ~2000 and ~6000 μ N, which are potential indications of mechanism activity within the structure. Fig. 6b shows contact stiffness versus mean applied load for a ramping load dynamic indentation test on the film. It is observed that as the applied load increases, a jump discontinuity in contact stiffness occurred at a maximum load of 1850 µN, with a magnitude of 1.8 N/mm. Looking at an SEM micrograph of the thin film surface after indentation at 8000 µN, shown in Fig. 7, it is clear that radial cracking, circumferential cracking, and delamination of the a-Si layer occur. Circumferential variations in contrast of the surface are potentially due to charging effects during SEM, which would result from a discontinuity between the two layers of the film. In addition, delamination of the a-Si film would occur prior to catastrophic failure, due to the distribution of tensile stress being maximized at the interface between the two materials. Therefore, it is suggested that the contact stiffness signature at 1850 µN seen in dynamic indentation corresponds to delamination of the a-Si layer. This is assumed to represent the same event which occurred at 2000 µN in quasi-static loading, since dynamic indentation applies a more severe loading condition and would result in a more rapid onset of failure. This conclusion is consistent with data from traditional nanoindentation of this structure at a variety of load levels, which show that catastrophic failure of Al/a-Si thin films, including the propagation of cracks, occurs at approximately 6000 μ N, as seen in Fig. 6a. Therefore, the mean critical load at which a flat Al/a-Si thin film fails due to quasi-static loading was determined to be ~2000 μ N.

3.3. Fatigue Testing

3.3.1. Al Nanodot Fatigue Behavior

Nanoindentation fatigue testing was performed on 300 nm Al nanodots to characterize their behavior when subjected to cyclic loading. Fig. 8 shows contact stiffness as a function of cycles for a 300 nm diameter Al nanodot indented for 5.0×10^4 cycles at a mean load of 70 µN and a load oscillation amplitude of 30 μ N. It was observed that the contact stiffness increased linearly during the first ~1.0 \times 10⁴ cycles of loading, after which the stiffness plateaued and remained more or less constant for the remainder of the test. This initial trend of increasing contact stiffness may be attributed to both increasing contact area between the indenter tip and the nanostructure [33], as well as strain hardening due to dislocation nucleation and propagation in metals [34,35]. Fig. 9c and d show SPM images taken before and after fatigue testing, where it is seen that significant permanent deformation resulted in the nanostructure. Since the residual deformation is very large in comparison to the total height of the nanostructure, there is a large corresponding increase in contact area between the interfacing bodies. Therefore, increasing contact area is credited as the predominant source of the change in contact stiffness. Also, since all measurements are made after the first load cycle is applied, it is known that any delayed phenomena must be an effect of the applied load oscillation. Because of this, a trend of increasing contact stiffness strongly suggests that plastic deformation is occurring with each subsequent loading cycle, up until the contact stiffness plateaus at ~ 1.0×10^4 cycles.

To further investigate this hypothesis, fatigue tests at the same load level were performed at up to 1.0×10^4 cycles, and contact stiffness versus cycles for this experiment is reported in Fig. 8. As before, the contact stiffness also transitioned from linear to constant. However, in contrast to the 5.0×10^4 cycle experiments, the time at which the structure remains at constant contact stiffness is greatly reduced because of the shorter testing time. Fig. 9a and b shows SPM micrographs of the Al nanodot after fatigue testing for 1.0×10^4 cycles, and Fig. 9c and d show SPM images of another Al nanodot tested for 5.0×10^4 cycles. Here it is observed that both nanostructures exhibit a permanent ~30 nm reduction in height. Thus, the cycles at which deformation ceases is defined as the point where contact stiffness remains constant, meaning that no further deformation is occurring past the first ~ 1.0×10^4 cycles, there remains significant permanent deformation in the nanostructure. This amount of residual deformation would render the structure ineffective for tribological and surface texturing applications, and should be classified as ductile failure of the nanostructure.

3.3.2. Fatigue Behavior of Flat Al/a-Si Thin Film

Fatigue testing was performed on flat Al/a-Si thin films to characterize the response of a layered material which lacks the geometric and core-confinement properties of standard CSNs. Fig. 10 shows contact stiffness versus cycles for fatigue tests on the Al/a-Si thin film at a 1300, 1400, 1500, and 1600 μ N mean load and a unanimous 500 μ N oscillating load amplitude. Due to the probabilistic nature of the failure signature appearing, fatigue tests at a variety of loads were conducted to establish the applied load which best captures fatigue failure, as well as to demonstrate the relationship between fatigue life and

applied mean stress. Fig. 10 shows contact stiffness for a 1300 μ N mean applied load, where no change in stiffness is observed. This indicates that a 1300 μ N mean load is not large enough to damage the film. At 1400 μ N, a jump in contact stiffness of 1 N/mm was observed at 4.5 × 10⁴ cycles. At subsequently higher applied mean loads, the number of cycles before which the failure signature appears decreases until 1600 μ N, where the jump appears at only 0.4 × 10⁴ cycles into testing. Following the argument presented for quasi-static nanoindentation experiments on this structure, a jump in contact stiffness at approximately 2000 μ N corresponded to delamination between the a-Si and Al layers. Since the same failure signature presents itself in fatigue tests at slightly depressed load levels and with delayed occurrence, it is suggested that this discontinuity also corresponds to delamination or subsurface fracture of the film. Thus, the critical fatigue load is identified as 1400 ± 250 μ N (estimated contact pressure of ~19.4 GPa), below which no fatigue damage is induced within the structure for the duration tested.

3.3.3. Deformation Resistant CSNs

Nanoindentation fatigue testing was performed on CSNs to characterize their mechanical response to cyclic nanoindentation loading. From the experiments on 300 nm Al nanodots, the maximum contact pressure induced by an 85 μ N applied load was estimated to be ~17.5 GPa. Hertzian Contact Theory was then used to transpose this pressure into the indenter force which applies an equivalent contact pressure to a CSN with a 100 nm core diameter and 300 nm thick shell, where it was assumed the CSN has an equivalent Young's modulus and Poisson's ratio based on the volumetric ratio of 20% Al and 80% a-Si. In this calculation, the values used for the Young's moduli are 170 GPa and 179 GPa for Al and a-Si, respectively, and the Poisson's ratios are 0.35 and 0.25 for Al and a-Si, respectively. This analysis yielded a mean indenter load of 55 μ N and a peak-to-peak load amplitude of 30 μ N.

Fig. 11a shows contact stiffness as a function of cycles for fatigue testing CSNs at a variety of load levels, including 55, 100, 150, and 200 µN mean loads superimposed with 30, 50, 75, and 100 µN oscillating load amplitudes, respectively. Three repetitions of each test show overlap in the stiffness response of individual nanostructures, which illustrates uniformity between nanostructures as well as repeatability of the testing procedure. Fig. 11b isolates one contact stiffness curve for each load level tested. The 55 \pm 15 μ N indentation displays a contact stiffness response which increases gradually throughout the entirety of the test, resulting in a total stiffness increase of ~1.5 N/mm across 5.0×10^4 cycles. Unlike the fatigue response of bare Al nanodots, an increasing trend in contact stiffness of CSNs may correspond to mechanisms involving both core and shell materials. These include the accumulation of dislocations and strain hardening within the Al core, increasing contact area beneath the indenter tip, pressure induced phase transformations in a-Si [36], as well as a-Si densification occurring within the shell [25,37]. In Fig. 12a and b, SPM images of a CSN following $55 \pm 15 \mu$ N indentation show that no residual deformation is present in the nanostructure. In addition, nanoindentation testing on multiple structures resulted in a mean height change of 2 nm at $55 \pm 15 \mu$ N. This eliminates increasing contact area as the primary source of this response, since progressive indenter displacement necessarily requires permanent deformation of the sample (i.e., if the elastic limit had not been surpassed by the initial loading cycle, there would be no further deformation due to subsequent cycles at the same load). In addition, nanoindentation experiments on a flat a-Si thin film show no evidence of a pressure-induced phase transformation up to an estimated contact pressure of ~22 GPa [29]. Since a-Si phase transformations and increasing contact area have been eliminated as potential sources of the increase in contact stiffness, it is suggested that a-Si densification and the accumulation of dislocations within the Al core are responsible.

This conclusion is consistent with experiments of repeated indentation on individual CSNs, which show hardening behavior with each subsequent indent [29].

Since no residual deformation was observed at the equivalent contact pressure of ~17.5 GPa, fatigue tests at incrementally higher applied loads were conducted to further investigate the resistance of CSNs to cyclic loading. It was observed that indents at $200 \pm 50 \mu$ N showed a rapid increase in contact stiffness, followed by a well-defined plateau beginning at 1.0×10^4 cycles. Data from indents at intermediate load levels illustrate a relationship between increasing applied load and the rate of increasing contact stiffness, while the magnitude of total increase unanimously remained ~1.5 N/mm. Fig. 12c shows an SEM image of a CSN after indentation at $200 \pm 50 \mu$ N for 5.0×10^4 cycles, where it is seen that even at the estimated contact pressure of ~25.1 GPa, there is no cracking or significant deformation present in the nanostructure. This is in contrast to the experiments conducted on the flat Al/a-Si layered structure, which exhibited visible cracking at an estimated ~19.4 GPa contact pressure.

To further investigate the response of CSNs subjected to high-pressure fatigue, fatigue testing was conducted at multiple test durations. Fig. 13 shows contact stiffness versus cycles for two separate CSNs undergoing fatigue testing at 200 μ N mean load superimposed with 100 μ N peak-to-peak oscillating load amplitude at 60 Hz frequency for 1.0×10^4 and 5.0×10^4 cycles. The two indentation curves closely overlap, displaying the same transition signature occurring at 1.0×10^4 cycles. Fig. 14 shows SPM images of CSNs fatigue tested at 200 μ N mean load for 1, 1.0×10^4 , and 5.0×10^4 cycles, which display a residual change in height of 3, 11, and 9 nm, respectively. Through indentation of 3 structures for each test duration, the mean change in height was determined to be 4.3, 8.6, and 9.6 nm for each testing time,

respectively, with standard deviations of 1.3, 2.7, and 3.4 nm, respectively. These results are consistent with the observed contact stiffness trends, and show that deformation of the nanostructure, while minor, occurs mostly within the first 1.0×10^4 cycles of loading. In addition, the absence of further height reduction in the longest test duration entails that there is no permanent deformation occurring past the transition to constant contact stiffness, and suggests that the CSN exhibits superior fatigue life past this point.

It should be noted that although similar contact stiffness trends were witnessed in both Al nanodots and CSNs, the response provided by the CSNs is more consistent between independent nanostructures than the bare Al cores, in addition to lacking significant residual deformation. This suggests that a nanotexture composed of patterned CSNs would possess greater uniformity, and thus higher reliability and consistency in application. As a result, it is clear that even at high contact pressures, CSNs are very resistant to deformation when subjected to prolonged cyclic loading and CSNs do not exhibit the fatigue failure mechanisms present in either bare Al nanodots or flat Al/a-Si thin films.

4. Conclusions

The mechanical fatigue response of hemi-spherical Al nanodots, flat Al/a-Si layered thin films, and novel Al/a-Si core-shell nanostructures was characterized using nanoindentation and nano-scale DMA. The fatigue behavior of each nanostructure was analyzed through the change in contact stiffness throughout the applied loading cycles. The CSNs demonstrate superior deformation resistant properties when subjected to cyclic compression loading for 5.0×10^4 cycles, even at contact forces up to 250 µN (estimated contact pressure of 25.1 GPa). When the a-Si shell is removed, bare Al cores demonstrate significant residual deformation due to repeated cyclic loading. The Al/a-Si thin films demonstrate delamination due to fatigue loading at contact pressures less than those applied to the CSNs. An analysis of the contact stiffness response of the CSNs show that dislocation nucleation and a-Si densification are occurring within the structure, resulting in hardening of the CSN with repeated loading. This study explicates that the novel deformation resistance of Al/a-Si CSNs persists through repeated loading cycles, and establishes their prolonged durability when implemented in nanomechanical applications.

5. Future Work

In this study, the experiments conducted to determine the fatigue behavior of a flat Al/a-Si layered thin film were not necessarily conclusive. An issue arose that although the nanoindentation results strongly suggest delamination between the a-Si and Al layers, this was not able to be captured through SEM imaging. When imaging was attempted, the fatigue testing location was not clearly defined, and it was not conclusive that failure of the surface occurred. This is likely due to one of two causes: recovery of the film surface over time, such that the indentation is not visually distinct, or displacement of the layers causing delamination, but the material recovers such that the a-Si and Al surfaces are separated but coincident.

To investigate the possibility of time-dependent recovery of the layered thin-film, SPM imaging before and after nanoindentation may be used. By imaging the indent location immediately afterwards, a measurable divot is present. Then, imaging the same location sometime later will allow direct analysis of whether the depth of the residual impression changes after indentation. This analysis will be sufficient in determining if time-dependent recovery of the nanostructure is occurring. In order to further investigate the mechanisms responsible for these failure signatures, focused ion beam (FIB) microscopy may be used to cut a fatigue indentation impression along its cross-section. Then, SEM imaging may be used to directly analyze the interface between the Al and a-Si layers, where it will be clear whether delamination occurred.

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Figure 3. SEM micrographs of NTSs composed of (a) 300 nm diameter Al nanodots and (b) CSNs with 100 nm diameter cores and 300 nm shell thicknesses.

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