The Effects of Acid Passivation, Tricresyl Phosphate Presoak, and UV/Ozone Treatment on the Tribology of Perfluoropolyether-Lubricated 440C Stainless Steel Couples

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THE EFFECTS OF ACID PASSIVATION, TRICRESYL PHOSPHATE PRE-SOAK, AND UV/OZONE TREATMENT ON THE TRIBOLOGY OF PERFLUOROPOLYETHER-LUBRICATED 440C STAINLESS STEEL COUPLES

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ABSTRACT

The boundary-lubrication performance of two perfluoropolyether (PFPE) thin films in the presence of passivated 440C stainless steel is presented. The study used a standard ball on disk (BoD) tribometer in dry nitrogen and a vacuum spiral orbit tribometer (SOT). Stainless steel surfaces were passivated with one of four techniques: high and low temperature chromic acid bath, a tricresyl phosphate (TCP) soak, or UV/Ozone treated for 15 minutes. After passivation, each BoD disk had a 400Å film of Krytox 16256 (PFPE) applied to it. The lifetimes of these films were quantified by measuring the number of sliding cycles before an increase in friction occurred. The lubricated lifetime of the 440C couple was not altered as a result of the various passivation techniques. The resulting surface chemistry of each passivation technique was examined using X-ray photoelectron spectroscopy (XPS). The SOT was used to examine the effects of the TCP treatment on the lubricated lifetime of another PFPE, Brayco 815Z, under rolling conditions. None of the passivation techniques were found to dramatically increase the oxide film thickness or lubricated lifetimes.

INTRODUCTION

Pretreatment of instrument-bearing components with either a tricresyl phosphate (TCP) presoak [1-10] or a chromic acid passivation bath [2,4], has been performed for over 30 years. It has been repeatedly shown that these pretreatment procedures improve the bearing's performance in low-speed, boundary lubricated contacts [1,2,9,10]. Over the years, these pretreatments have been incorporated in standard bearing component preconditioning procedures in an attempt to provide additional protection while in boundary lubrication conditions [2,8]. However, all known reports of improved boundary lubrication performance have used either mineral oils or synthetic esters. This is not surprising since the vast majority of TCP and chromic acid passivation research was performed in the 1960's, when these were the dominant classes of lubricants. These lubricant classes are vastly different than the perfluoropolyether class commonly used today. There has only been one study [11] that focused on the longevity of a TCP passivated surface lubricated with Krytox, a PFPE, and none on chromic acid passivated surfaces. Wedeven et al. [11] concluded that TCP passivation had no effect on the longevity of the PFPE lubricated system.

Perfluoropolyethers (PFPEs) have been the preferred liquid lubricants for space applications for over three decades because of their proven
tribological performance and other attractive properties [12] including low vapor pressure, low chemical reactivity, and wide liquid temperature range. PFPEs are used in many space mechanisms including actuators, antenna pointing mechanisms, filter wheels, gyroscopes, and scanning mirrors [13,14]. However, there have been many incidents during which PFPE lubricated components have shown anomalous behavior [15,16]. These anomalies are thought to be the result of PFPE degradation. In order to understand how various passivation techniques might affect the boundary lubricating performance of PFPEs, it is first necessary to understand how PFPEs degrade.

Several studies have focused on modeling the degradation of PFPE lubricants. Research has repeatedly shown that PFPEs degrade while in boundary lubricated, sliding/rolling contact [17-22] and at elevated temperatures [23-26], losing their desirable properties, including lubricity. Although few models have been proposed to explain the degradation, it has been shown that the rate of degradation is dependent upon the substrate surface chemistry and the presence and relative strength of Lewis acids.

Kasai [23] has shown that the thermal catalytic degradation of a PFPE is dependent upon the strength of the Lewis acid present and has proposed a chemical degradation mechanism for PFPEs. He postulated a mechanism in which Lewis acids act as a catalyst that promotes an intramolecular disproportionation reaction, which results in polymer-chain degradation.

Other static degradation studies, conducted to look at the effects of certain metals or metal alloys in the presence of particular PFPEs, have shown that PFPE degradation is highly dependent upon the chemistry of the substrate. For example, an unbranched PFPE lubricant showed more degradation at 288°C in the presence of M-50 steel and Ti-4Al-4Mn than in contact with either pure titanium or pure aluminium [25].

Some studies have applied the concept of modifying one or both of the surface chemistries by coating or ion implantation and observing the associated tribological characteristics. Carré [20] demonstrated that the PFPE, boundary lubricated, wear life was increased by a factor of 5 to 10 by using TiN-coated 440C test specimens. TiC-coated 440C and SiN$_2$ ceramic ball bearings have also been shown to have greater longevity compared to their 440C counterparts when PFPE lubricated [19-21]. In Carré's studies [20,21] it was concluded that the hard coatings and ceramic material prevented the formation of Lewis acids, thus halting degradation of the PFPE. Shogrin et al. [27] demonstrated that implanting 440C substrates with Ti either with or without a carbon (CH$_4$) back pressure enhanced the lubricating lifetimes of a PFPE. It was concluded that the resulting microstructures inhibited the formation of reactive sites.

Another way to inhibit tribologically induced degradation may be to passivate the contacting surfaces, which might inhibit the formation of Lewis acids in the presence of fluorinated oils, or decrease the availability of reactive sites on the surface, thus, extending the lubricated lifetime.

The objective of this work was to study the effect of various passivation techniques on the lifetime of PFPE lubricated (Krytox 16256 and Brayco 815Z), 440C stainless steel couples in sliding and rolling contact. Chromic acid passivation (similar to 2,4) and TCP (similar to 1-10), as well as a modified chromic acid passivation technique (which uses a higher temperature and a longer reaction time), and UV/Ozone treatment [28] were studied in sliding contact. TCP and UV/Ozone treatments were also examined in rolling contact. UV/Ozone treatment has been used to remove carbonaceous contamination from surfaces. A demonstration of their effectiveness in prolonging lubricity is sought rather than a detailed description of the associated chemical degradation mechanisms.

**EXPERIMENTAL**

**TRIBOMETERS**

**Ball-on-disk (BoD)**

Tribological testing of each disk/ball couple was performed on the fixed ball-on-disk tribometer shown in Figure 1. All tests were performed using a relative sliding speed of 0.05 m/s and a 3N normal load, which generated an initial Hertzian contact pressure of -0.69 GPa. Tests were performed in dry N$_2$ (< 2% RH) at ambient temperature (~25°C). Friction force as a function of disk revolutions (cycles) was recorded for each test.

After a test, the ball was rotated to expose an unworn surface and the fixture was extended so the ball would contact the disk at a different radius.
By doing this, three wear tracks, 1.59 mm (1/16") apart, could be worn on each disk.

Figure 1 – Ball-on-disk (BoD) tribometer

Figure 2 – The spiral orbit tribometer (SOT)

**Vacuum Spiral-Orbit-Tribometer (SOT)**

Lubricant lifetimes in rolling contact under vacuum were examined using the spiral orbit tribometer (SOT) shown in Figure 2. The SOT is essentially a thrust bearing with a single ball and no retainer. The lower disk is fixed and the top disk rotated. The ball rotates in a spiral orbit and the guide plate is used to return the ball to its original orbit once per revolution. The force the ball exerts on the guide plate is measured and friction force can be calculated. Accelerated testing is achieved by limiting the amount of lubricant available to micrograms, which is completely consumed during the test. Lifetime is defined as the number of orbits before the friction coefficient exceeds a predetermined value. The tribometer is further described in References 29-32 and its previous usages appear in References 33-35.

The effects of TCP passivation and UV/ozone treatment were examined using the SOT. All tests were performed using a mean Hertzian stress of 1.5 GPa, room temperature (~23°C), a vacuum <1.3x10^6 Pa, 200-RPM rotational speed, and 50 μg lubricant.

**MATERIALS**

**Ball on Disk Tribometer**

Disks and bearing balls made from 440C stainless steel (fully hardened) were used. The 440C disks were 17.5 mm (11/16") in diameter and 4.76 mm (3/16") thick. Each disk was polished to a mirror finish having an arithmetic roughness average, \(R_a\), of ~ 0.01 μm. The bearing balls were Grade 10 and had a diameter of 9.53 mm (3/8").

**Spiral Orbit Tribometer**

Disks and bearing balls made from 440C stainless steel (fully hardened) were used. The 440C disks were 50.8 mm (2") in diameter. The disks and guide plate were polished to a mirror finish having an arithmetic roughness average, \(R_a\), of ~ 0.01 μm. The bearing balls were Grade 25 and had a diameter of 12.7 mm (1/2").

**Lubricant**

The surface preparations tested were: unlubricated and untreated (unpassivated), unlubricated and treated (passivated), and passivated and unpassivated disks that were lubricated. Throughout this report the term "unpassivated" refers to pairs that have not been passivated or
UV/Ozone treated. The lubricant used for the BoD was Krytox 16256. Typical properties of this PFPE are given in Table 1(a). The lubricant used for the SOT tests was Brayco 815Z and its properties are listed in Table 1(b).

<table>
<thead>
<tr>
<th>Table 1(a) – Selected properties of Krytox 16256</th>
</tr>
</thead>
<tbody>
<tr>
<td>20°C</td>
</tr>
<tr>
<td>---</td>
</tr>
<tr>
<td>Vapor Pressure (Torr)</td>
</tr>
<tr>
<td>Kinematic Viscosity (cS)</td>
</tr>
<tr>
<td>Density (g/cm³)</td>
</tr>
<tr>
<td>Surface Tension at 25°C</td>
</tr>
<tr>
<td>Average Molecular Weight</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Table 1(b) – Selected properties of Brayco 815Z</th>
</tr>
</thead>
<tbody>
<tr>
<td>20°C</td>
</tr>
<tr>
<td>---</td>
</tr>
<tr>
<td>Vapor Pressure (Torr)</td>
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<td>Kinematic Viscosity (cS)</td>
</tr>
<tr>
<td>Density (g/cm³)</td>
</tr>
<tr>
<td>Average Molecular Weight</td>
</tr>
</tbody>
</table>

**SPECIMEN PREPARATION**

**Ball on Disk Tribometer**

All disk/ball pairs were ultrasonically cleaned in hexane, acetone, and methanol for approximately ten minutes each. The pairs were then blown dry with nitrogen and covered to minimize dust contamination. Disk/ball pairs that were not passivated or lubricated were placed in a N₂ atmosphere (R.H. < 2%) at ambient temperature where they remained until tested. Unpassivated disks that had PFPE applied, were lubricated and then placed in a N₂ atmosphere, with their accompanying unpassivated ball. Disk/ball pairs were treated using one of the four following procedures.

**Spiral Orbit Tribometer**

The disks and ball were ultrasonically cleaned in hexane, methanol, and distilled water for ten minutes each. The pairs were then rinsed in methanol and blown dry with nitrogen. The disks and ball were treated. The ball was lubricated and then the parts were placed into the vacuum chamber at ambient temperature where they remained until tested. Disks and the ball were treated using one of the four following procedures.

**Chromic-Acid Passivation**

Clean components were passivated in one of two ways using a solution consisting of 2.0 gm sodium dichromate (Na₂Cr₂O₇-2H₂O), 20 ml concentrated nitric acid (HNO₃), and 80 ml deionized H₂O. Specimens were immersed in the solution either for 30 minutes while heated to 46°C, or for 60 minutes while heated to 56°C. The samples were then removed and thoroughly rinsed using two successive deionized H₂O baths. Samples were then blown dry with nitrogen and covered.

**Tricresyl Phosphate (TCP) Passivation**

Components were immersed for 48 hours in neat tricresyl phosphate (TCP) heated to 107°C. (The TCP was purchased in accordance with the requirements of TT-T-656b). The samples were then ultrasonically cleaned in hexane, acetone, and methanol for approximately 10 minutes each. Samples were then blown dry with nitrogen and covered.

**UV/Ozone Treatment**

Disk/ball pairs were UV/ozone treated for 15 minutes to remove carbonaceous contamination [28], which begins to reform immediately after this treatment.

Treated disks that did not have PFPE applied to them, and all treated bearing balls were placed in a dry N₂ atmosphere where they remained until tested. Treated disks that had PFPE applied to them were lubricated (UV/ozone passivated disks were lubricated within 5 minutes of passivation) and then placed in the same dry N₂ environment.

**XPS Analysis of Passivated Surfaces**

A disk passivated by each of the four techniques and a baseline disk (untreated) were analyzed using X-ray photoelectron spectroscopy (XPS). XPS analyzes the top several atomic layers of the sample providing both elemental and chemical state information. An area of 2 mm x 5 mm was analyzed on each surface. Survey spectra, from 0 to 1200 eV binding energy, were taken to determine which elements were present followed by high-resolution energy windows for each of the major elements detected on the surface to determine their chemical state. There is typically a 5 to 10% error associated with the resulting calculated atomic concentrations.
APPLICATION OF LUBRICANT FILMS

Ball on Disk Tribometer

Thin film deposition was accomplished by using the apparatus shown in Figure 3. This is a derivative of the deposition process known as dipping. In this process, a specimen is lubricated by slowly withdrawing it at a constant speed from a solution of the fluid in a volatile solvent. As the solvent evaporates, a thin film of the non-volatile fluid remains on the surface. In this procedure, both solution concentration and withdrawal speed govern the applied film thickness [36]. For the apparatus used in this study, gravity driven solution flowed past the stationary specimen. The surface speed that the solution attained was approximately 0.8 mm/sec. Each disk was lubricated using a solution with a concentration of 5.0 g of PFPE lubricant in 100 ml of solvent (a perfluorinated cyclobutane).

LUBRICANT FILM THICKNESS MEASUREMENTS (BALL ON DISK TRIBOMETER)

The PFPE thickness on each disk was measured using an infrared microscope (μ-FTIR) in the reflectance mode using a Grazing Angle Objective (GAO) with a Mercury Cadmium Telluride (MCT) detector. The instrument's spectral resolution was 4 cm⁻¹. A gold-coated glass slide was used as the background. Absorbance as a function of wave number was determined for each film at three different radii on the disk's surface, approximately 120° apart using an analysis spot diameter of 100 μm.

Theoretical thin film optical calculations have been performed to model the PFPE film's behavior in the infrared spectral region [37]. These results are plotted as absorbance of the 1313 cm⁻¹ band as a function of film thickness. After experimentally obtaining the absorbance of the 1313 cm⁻¹ band, the corresponding film thickness was calculated.

The FDD apparatus produced PFPE film thicknesses of ~400Å. Lubricant uniformity across any one surface was ± 15%. The error associated with surface-to-surface film thickness repeatability was ± 20%.

RESULTS AND DISCUSSION

BALL ON DISK TRIBOMETER

Establishment of the Lubricant-Failure Criterion

Lubricant failure occurred when the friction coefficient increased to a value typical of an unlubricated, unpassivated disk/ball couple. When failure occurred, it was argued that the protective layers (applied lubricant, surface oxide, and/or additional passivation layer) either wore off, ruptured, or broke down. Figure 4 shows the mean friction coefficient as a function of cycle number for three wear tracks on two separate unlubricated, unpassivated couples. Mean friction coefficient was the mean of the friction coefficients of the complete revolution over the indicated cycle. Typically, friction coefficient fluctuations over a
cycle were ±0.06 (maximum: ±0.15) about the mean. The mean friction coefficient data all lied between 0.63 and 0.8. Thus, an increase in mean friction coefficient above 0.63 was indicative of failure and was defined as the failure limit. The number of rotational cycles that can be applied to the lubricant film on a wear track before the friction rose above the failure limit was the film's lubricating lifetime.

It was also important to know if the passivation process itself changed the failure limit. Therefore, one disk passivated at each condition was tested unlubricated. Initially, the chromic-acid and TCP passivation samples exhibited lower mean friction coefficients (0.26-0.3), but then raised near linearly above the failure limit at different rates. The value of the TCP passivated surface continued to rise to -0.8, where it then dropped to a value of -0.53 in the course of one cycle, after which it again rose to failure and stayed. The UV/Ozone treated surfaces showed no improvements and behaved the same as the untreated samples depicted in Figure 5. Table 2 summarizes this data. The UV/Ozone treated results are not shown because no improvements were observed with this technique. In this table, the average values appear outside the parentheses, whereas the ranges are contained within the parentheses. These data suggest that both chromic-acid passivation techniques improved the friction coefficient for a brief time, whereas TCP passivation extended the low friction coefficient for an extended time.

Table 2 – Unlubricated wear test results

<table>
<thead>
<tr>
<th>Surface Passivation Treatment</th>
<th>Initial Mean Friction Coefficient</th>
<th>No Of Cycles Until Failure</th>
<th>Tracks Tested</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chromic-Acid, Low Temp</td>
<td>0.27 (0.24-0.27)</td>
<td>5 (4-6)</td>
<td>5</td>
</tr>
<tr>
<td>Chromic-Acid, High Temp</td>
<td>0.26 (0.24-0.27)</td>
<td>9 (7-10)</td>
<td>4</td>
</tr>
<tr>
<td>Tricresyl Phosphate</td>
<td>0.29 (0.27-0.34)</td>
<td>68 (6-85)</td>
<td>5</td>
</tr>
</tbody>
</table>

Figure 4 – Typical friction coefficient data for unpassivated, unlubricated disk/ball couples [27]

LUBRICATED RESULTS

Lubricated, unpassivated disks/ball couples were tested. A typical plot of mean friction coefficient vs. cycle number is shown in Figure 5. This data was taken from a disk with a ~70 Å PFPE film. The mean friction coefficient begins to rise from <0.3 near 200 cycles and crosses the failure line of 0.63 at ~370 cycles (lubricant lifetime = 370 cycles).

This general trend was typical for the samples used in this study. In these tests the maximum variation of the friction coefficient before it began to rise (here at ~200 cycles) was ±0.02. The relatively small variation indicated that the entire wear track was uniformly experiencing the benefits of full boundary lubrication from the PFPE. After the initial rise, the variation of friction coefficient grew as the number of cycles increased until, at the failure limit, it typically had a variation of ±0.1 (maximum: ±0.20, minimum: ±0.03).

As a test progressed, different locations on the wear track lost their lubricity at varying rates, which accounted for the spread in friction coefficient values seen over one cycle (indicated by the variation noted above). Although the rate of lubricity loss over the entire wear track varied, and the variation increased as the number of cycles increased, the mean friction coefficient continued to rise, signifying an overall loss of lubricity.
Beyond failure, variations of the friction coefficient over a rotational cycle, as well as the mean friction coefficient were sporadic. On some wear tracks, the mean friction coefficient dipped below the failure line after it had reached failure (rarely dipping below ~0.4). This decrease was caused by the rapid growth of the wear scar's width coming into contact with fresh, unused PFPE lubricant. This fresh lubricant decreased the friction coefficient before it too degraded, causing the mean friction coefficient to again rise above the failure limit. These sporadic variations were also considered an indicator of failure.

After performing a number of the tests, it was apparent that the lifetimes had a significant spread in their values for each of the unpassivated and passivated conditions. These spreads in lifetime appeared random, as they were not dependent upon either a particular disk or radius of wear track. Weibull statistics [38] were used to characterize this data.

The Weibull plot for the unpassivated disk/ball couples lubricated with Krytox 16256 is shown in Figure 7. Each data point represents the lubricated lifetime of one wear track. The Median Rank Line represents the most probable failure history for the samples, that is, the fraction of a group of samples that can be expected to fail (i.e. exceed 0.63 friction coefficient) as a function of the number of sliding cycles. This line suggests 60% of a given number of tests can be expected to undergo failure at or before ~3700 cycles. Given an infinite number of tests, the 90% confidence bands were in the region in which the lifetimes of 90% of those tests would reside.

The usefulness of a Weibull plot can be demonstrated by relating it to the reliability of a component. For a typical space bearing, which cannot generally be retrieved for maintenance, reliability is very important. If greater than 99% reliability is desired in a lubricated couple, then the number of cycles that it is allowed to experience should be less than the value that will generally produce failure 1% of the time. For the unpassivated samples in Figure 6, 1% of them are predicted to fail by ~120 cycles. By comparing each passivation technique's Weibull plot to the plot of the unpassivated specimens (Figure 6), and to each other, produces their relative performance. Their relative performance is summarized after the next section.
PASSIVATED - LUBRICATED RESULTS

The Weibull lifetime distributions of the four passivation techniques and the unpassivated technique (from Figure 6) are shown in Figure 7. This figure indicates that the predicted lifetimes are all very similar. The 90% confidence bands of the five plots (not shown) all overlap over most of the range of each Median Rank Line. Therefore, within 90% assurance, it is predicted that no one passivation technique out performs another. These plots indicate that none of the passivation techniques exhibited superior or grossly sub-standard performances relative to the unpassivated surfaces.

Given the sliding nature of these tests and the thinness of both the lubricant and the passivation layers, one might have predicted this result. However, the unlubricated tests did show measurable differences between the various techniques. In addition, this same procedure was used to evaluate differences in various ion-implanted surfaces [27]. The thicknesses of the implanted layers were of the same order as the thicknesses of the passivated layers. Finally, it has been known for decades that the TCP presoak and the low temperature chromic acid passivation techniques provide additional protection in boundary lubricated contact while lubricated with either mineral oils or esters [1-10]. Thus, the authors conclude that none of the passivation techniques tested increased the longevity of the perfluoropolyether (16256) lubricated wear couples.

Table 3 – Summary of data from Weibull plots

<table>
<thead>
<tr>
<th>Surface Passivation/Treatment</th>
<th>1% Failure</th>
<th>60% Failure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Unpassivated</td>
<td>120</td>
<td>3700</td>
</tr>
<tr>
<td>Chromic-Acid, Low Temp</td>
<td>440</td>
<td>3600</td>
</tr>
<tr>
<td>Chromic-Acid, High Temp</td>
<td>440</td>
<td>7000</td>
</tr>
<tr>
<td>Tricresyl Phosphate</td>
<td>1000</td>
<td>2300</td>
</tr>
<tr>
<td>UV/Ozone Treatment</td>
<td>240</td>
<td>5000</td>
</tr>
</tbody>
</table>

XPS ANALYSIS OF THE PASSIVATED SURFACES

By using XPS oxide-to-metal peak area ratios, it was estimated that the oxide film thickness for all the surfaces (CrO\x26y, FeO\x26y, or FePO\x26y) was less than ~50 Å. A summary of the atomic percent of selective elements obtained from the XPS analysis of a surface passivated using each of the four techniques and the unpassivated surface is shown in Table 4. The unpassivated surface had Cr to Fe ratio of 0.34, and a distinct carbonaceous contaminating layer as indicated by the C atomic percent.

The low temperature chromic-acid passivated sample had a high Cr/Fe ratio (2.3). This indicates that the passivation solution selectively attacked and dissolved the near-surface iron oxide, while leaving the more inert chromium oxide intact. This would account for the change in Cr to Fe ratio in the first few atomic layers of the surface. The higher nitrogen atomic percent on this surface was caused by the nitric acid.
The surface of the sample passivated with the high temperature chromic-acid solution has a lower Cr to Fe ratio than the low temperature chromic-acid passivated surface, but a higher ratio than the unpassivated surface. The difference in the Cr to Fe ratios for the two passivation techniques is considered to be within the random error associated with the passivation procedure. It is noted that the two passivation techniques resulted in very similar surface chemistries, which indicates that increasing the passivation time and temperature was relatively ineffective. The fact that the Cr to Fe ratio is higher than the unpassivated surface indicates that the Fe atoms are being selectively removed at a higher rate than the Cr atoms.

Table 4 – Elemental surface concentration for each passivating treatment obtained by XPS

<table>
<thead>
<tr>
<th>Atomic % of</th>
<th>C</th>
<th>O</th>
<th>Fe</th>
<th>Cr</th>
<th>Cr/Fe Ratio</th>
<th>Misc.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Unpassivated</td>
<td>28</td>
<td>54</td>
<td>13</td>
<td>4.4</td>
<td>0.34</td>
<td>0.5N</td>
</tr>
<tr>
<td>Chronic-Acid, High Temp</td>
<td>23</td>
<td>61</td>
<td>4.3</td>
<td>10</td>
<td>2.30</td>
<td>0.9N</td>
</tr>
<tr>
<td>Chronic-Acid, Low Temp</td>
<td>19</td>
<td>61</td>
<td>10</td>
<td>9.3</td>
<td>0.93</td>
<td>0.9N</td>
</tr>
<tr>
<td>Tricresyl Phosphate</td>
<td>24</td>
<td>58</td>
<td>4.6</td>
<td>1.1</td>
<td>0.24</td>
<td>0.8Na  \ 12P</td>
</tr>
<tr>
<td>UV/Ozone Treatment</td>
<td>18</td>
<td>58</td>
<td>18</td>
<td>6</td>
<td>0.33</td>
<td>&lt;0.1N</td>
</tr>
</tbody>
</table>

Evidence for iron phosphate formation from TCP passivation is shown in Figure 9. The figure compares the XPS spectra of the Cr region of the high-temperature chromic-acid passivated surface to that of the TCP passivated surface. The chromic-acid passivated surface has a large Cr\(^{3+}\):Cr\(^{6+}\) metal peak, whereas, the TCP passivated spectra contains a much smaller Cr\(^{3+}\):Cr\(^{6+}\) metal peak. This, along with the presence of the 12 atomic percent P, and the decreased Cr to Fe ratio are consistent with the formation of an iron phosphate surface layer.

As evidenced by a decrease in carbon on the UV/ozone treated surfaces, UV/ozone treatment preferentially removes the carbonaceous contamination layer. UV/ozone treatment was not found to increase the oxide film thickness of these samples. This fact has been previously documented [28].

Figure 9 – XPS spectra showing the chromium oxide region of both the high temperature chromic-acid and TCP passivated surfaces

SPIRAL ORBIT TRIBOMETER

The SOT results also showed no difference in lubricated lifetime of Brayco 815Z when comparing the TCP passivated surface to the UV/Ozone treated surface. Lifetime for the UV/Ozone test was 57 ± 28 orbits per microgram vs. 64 ± 42 orbits per microgram for the TCP passivated surfaces. The SOT mimics the operating parameters of a real bearing operating in the boundary lubrication regime. These results support the conclusion that none of the passivation techniques tested increased the lubricated lifetimes using perfluoropolyethers.

SUMMARY OF RESULTS

1. None of the passivation techniques significantly increased the oxide film thickness.
2. Using X-ray photoelectron spectroscopy, it was found that chromic acid passivation increased the Cr to Fe ratio of the surface. Couples passivated with chromic acid, tested unlubricated, experienced an improvement in friction coefficient for a short time. However, chromic acid passivation did not affect the PFPE lubricating lifetime.
3. TCP passivation resulted in a few-monolayer-thick FePO\(_4\) layer on the surface. TCP passivated couples tested unlubricated, experienced improved, low friction for a short duration. However, an improvement in PFPE-lubricating lifetime was not observed after TCP passivation.
4. UV/Ozone treatment removed the carbonaceous contamination layer on the surface but did not alter either the unlubricated or the PFPE-lubricated lifetimes.
CONCLUSIONS

1. The PFPE lubricated lifetime (Krytox 16256) of sliding 440C couples did not statistically change as a result of the various passivation techniques.
2. The PFPE lubricated lifetime (Brayco 815Z) of 440C couples in boundary lubricated rolling contact did not statistically change as a result of the TCP passivation technique.

REFERENCES

The Effects of Acid Passivation, Tricresyl Phosphate Presoak, and UV/Ozone Treatment on the Tribology of Perfluoropolyether-Lubricated 440C Stainless Steel Couples

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The boundary-lubrication performance of two perfluoropolyether (PFPE) thin films in the presence of passivated 440C stainless steel is presented. The study used a standard ball on disk (BoD) tribometer in dry nitrogen and a vacuum spiral orbit tribometer (SOT). Stainless steel surfaces were passivated with one of four techniques: high and low temperature chromic acid bath, a tricresyl phosphate (TCP) soak, or UV/Ozone treated for 15 min. After passivation, each BoD disk had a 400 Å film of Krytox 16256 (PFPE) applied to it. The lifetimes of these films were quantified by measuring the number of sliding cycles before an increase in friction occurred. The lubricated lifetime of the 440C couple was not altered as a result of the various passivation techniques. The resulting surface chemistry of each passivation technique was examined using X-ray photoelectron spectroscopy (XPS). The SOT was used to examine the effects of the TCP treatment on the lubricated lifetime of another PFPE, Brandy 815Z, under rolling conditions. None of the passivation techniques were found to dramatically increase the oxide film thickness or lubricated lifetimes.