Abstract
A new non-destructive method was applied in order to assess bone integrity. The method is based on measurement of bone resorption. It was first published in 1966 and since then, in sensitive patients, is known to cause serious problems in joint replacement surgery. Titanium niobium nitride (TiNbN) can act as a surface coat for knee arthroplasty to "hide" the cobalt chrome (CoCr) femoral component beneath, therefore affording an immunoprivileged state. The aim of this study is to determine the wear properties of titanium niobium nitride against Ultra High Molecular Weight Polyethylene (UHMWPE) compared to cobalt chrome and to examine the metallic alloy surface of knee prostheses after loading cycles using a knee simulator. Three TiNbN coated and one CoCr Vanguard total knee femoral components were articulated against standard UHMWPE grade tibial inserts in the Stanmore-Instron knee simulator. Surface roughness, UHMWPE mass, lowest point, surface profiles and volumetric change were measured every one million cycles up to five million cycles. After five million cycles the average roughness of the cobalt chrome medial and lateral femoral condyles was over three times that of the TiNbN coated femoral condyles. There was no obvious difference in weight loss, volume loss or progression of lowest points of the tibial inserts articulating with the TiNbN coated and the cobalt chrome femoral component. Despite a clear reduction in roughness progression over the course of this in vitro test, there was no demonstrable improvement in UHMWPE wear measured gravimetrically or by surface profiling. The TiNbN implant tested may still be of great benefit to patients who are metal sensitive, but the coat offers no benefit in UHMWPE wear.

Keywords: Knee arthroplasty; Titanium niobium nitride; Knee simulator; Wear

Introduction
Hypersensitivity to an orthopaedic implant was first published in 1966 [1] and since then it was suggested it can pose serious problems in joint replacement surgery. 10-20% of the general population is reported to have metal sensitivity [2-4]. These patients are susceptible to type IV hypersensitivity whereby the release of nickel, cobalt and chromium ions in their cobalt chrome (CoCr) implant can cause a cell-mediated immune reaction [5]. There is still debate within the orthopaedic community about the presence and significance of deep metal sensitivity associated with orthopaedic implants. However, many patients complain about pain with no radiographic evidence of implant malpositioning and the diagnosis of metal sensitivity is often based just on positive skin patch tests. This involves application of the potential allergen to bare skin and the positive identification of a contact dermatitis reaction within 48 hours, effectively inducing a type IV hypersensitivity reaction on the skin. Additionally patients requiring knee joint replacements often indicate that they are metal sensitive and this is associated with a positive skin patch test.

To address this issue, various strategies possible solutions have been investigated: the CoCr femoral components can be coated or alternative femoral component materials such as ceramics can be used. Titanium niobium nitride (TiNbN) can act as a surface coat to "hide" the cobalt chrome femoral component beneath, therefore affording an immunoprivileged state [6]. Titanium niobium nitride is a coating applied by physical vapour deposition (PVD). PVD is a process by which positively charged metal ions are evaporated in a vacuum chamber and react with inert gases introduced to the chamber. The surfaces that are to be coated are negatively charged to allow a strong bond to form between the substrate and the coating [7,8].

The aim of this study is to determine whether there is a difference in Ultra High Weight Polyethylene (UHMWPE) wear articulating against cobalt chrome and titanium niobium nitride and to examine the metallic alloy surface after 5 million loading cycles in a knee simulator.

Materials and Methods
Four CoCr Vanguard (Biomet, Bridgend, UK) total knee replacements were used. Three femoral components were coated with TiNbN via physical vapour deposition, the other made from CoCr. Four corresponding tibial base-plates were all made of CoCr. Six implant grade UHMWPE tibial inserts of 10 mm were used; four in the simulator wear testing, and two as soak controls.

The Stanmore-Instron four station, force driven knee simulator was used, providing six degrees of freedom of motion (Anterior/Posterior, Lateral/Medial, Superior/Inferior) in accordance with ISO standard 14243 providing proven clinical wear performance. The components were fixed using Simplex P, acrylic bone cement (Stryker Orthopaedics, Mahwah, NJ, USA) to the simulator. Before commencing the test all the components were aligned. The femoral and tibial components were aligned to 0° of flexion and 0° antero-posterior tilt, varus/valgus and axial rotation respectively. The knee simulator was run at 1 Hz using axial rotation respectively. The knee simulator was run at 1 Hz using

*Corresponding author: Kunalan Maruthainar, John Scales Centre for Biomedical Engineering, The Institute of Orthopaedics and Musculoskeletal Science, Royal National Orthopaedic Hospital Trust, Brockley Hill, Stanmore, Middlesex, UK

Received July 11, 2013; Accepted August 24, 2013; Published August 30, 2013


Copyright: © 2013 Malikian R, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.
a standard walking cycle at a temperature of 37°C. Figure 1 illustrates the load, torque, displacement, rotation, soft tissue load and anterior-posterior shear load throughout the simulated gait cycle.

Prior to commencing testing and after completion of each million cycles all components were cleaned under aseptic conditions in Decon (Decon Laboratories Limited East Sussex, UK) and washed with distilled water to remove lubricant and proteins. They were placed in an incubator at 37°C to dry for 24 hours.

The weight of UHMWPE inserts was measured using a Precisa 92SM-202A scale (Precisa Instruments AG, Dietikon, Switzerland) accurate to 5 decimal points. In order to keep fluid absorption in consideration the two soak controls were weighed and the net difference was calculated at each cycle interval. If a mass difference in the controls was measured it would be used to correct the mass of the test components to ensure any changes were only due to wear. The surface profile of the tibial inserts were measured using an NPL soft probe profiler (3D Digital Design, Southgate, England) controlled by a computer programme driven machine (Heidenhain, West Sussex, England). Prior to testing, three indentations were milled around the same area is identified over repeated tests for consistent analysis. This act as calibration start points for the surface profiler to ensure the test could be derived. The lowest points on the medial and lateral condyles were symmetrical distribution, with bimodal peaks of greatest surface roughness in the middle region of both medial and lateral condyles. All surface roughness, with greatest surface roughness being in the posterior plane on the medial and the lateral condyle. These were at 10° intervals from 0° to 30° flexion. This resulted in 24 measurements per femoral component, from which an average roughness was calculated.

As suggested in the ISO 14243 standards, a lubricant of 30% fetal calf serum and 0.8% sodium azide was used. The soak controls were held at 37°C. Apart from the motion and loading, they were subjected to the same treatment as the test samples. The serum was changed every five days and disposed of. The test was run for five million cycles, which is similar to five years total knee replacement use in vivo [9].

Average surface roughness (Ra) was measured using a Mitutoyo Surftest (Mitutoyo, Kawasaki, Japan) with a 0.08 mm cut-off. Six measurements were taken at four points perpendicular to the anterior-posterior plane on the medial and the lateral condyle. These were at 10° intervals from 0° to 30° flexion. This resulted in 24 measurements per femoral component, from which an average roughness was calculated.

**Results**

**Surface roughness**

Prior to the start of the test, the average roughness of the titanium niobium nitride implants (0.035 micrometers) was greater than that of the cobalt chrome implants (0.018 micrometers), although there were no visible scratches. At the end of the test the average roughness of the cobalt chrome medial and lateral femoral condyles was a factor of 2.23 and 4 greater than that of the average roughness of the titanium niobium nitride coated femoral condyles respectively. CoCr lateral condyle roughness increased over the five million cycles by a factor of 17.41 from 0.017 to 0.296. The corresponding change for the titanium niobium nitride coated lateral condyles was from 0.036 to 0.074, equivalent to a rise of a factor of 2.04. The cobalt chrome medial condyle roughness rose from 0.019 to 0.256 resulting in an increase by a factor of 13.69. The titanium niobium nitride coated medial femoral condyle average roughness rose over the course of the test from a start figure of 0.034 to 0.115. This equated to a rise of a factor of 3.37. The difference in average roughness was greater for the lateral condyle than the medial condyle. The graphical representations of the roughness are presented in Figure 2.

CoCr femoral condylar surface roughness had a relatively symmetrical distribution, with bimodal peaks of greatest surface roughness in the middle region of both medial and lateral condyles. All Titanium niobium nitride condyles had an asymmetrical distribution of surface roughness, with greatest surface roughness being in the medial condyles, seen as an obvious spike on the graph in Figure 3.

**Gravimetric wear**

There was no obvious difference in the weight loss of the tibial inserts articulating with the titanium niobium nitride coated and the cobalt chrome femoral components. The mean weight loss of the three titanium articulating inserts was 0.04222g over five million cycles,

![Figure 1: Graphs demonstrating the forces and movements in the simulator at 5 million cycles.](image1)

![Figure 2: Average femoral roughness.](image2)
whereas, for the one insert articulating against cobalt chrome was 0.02611g over the test period. At 2,000,000 cycles there was a negative mass change with the TiNbN 3 component, indicating mass. The gravimetric wear of the individual inserts is presented in Figure 4.

Surface profiling

The results of the surface profiling which were used to calculate volume loss of UHMWPE showed no clear difference between the two bearing surfaces (Figure 5). There was no evidence of a reduction in volume loss with a titanium niobium nitride surface.

Lowest points

The lowest points show a downward progression over the time of the simulation. There was no significant difference between the progression of the lowest points of the titanium niobium nitride coated and the cobalt chrome over the course of the test (Figure 6 medial, Figure 7 lateral).

Discussion

Surface roughness of femoral components

After five million cycles in the knee simulator the surface roughness of the three TiNbN coated components was lower than the post-test roughness of the cobalt chrome coated component. This is despite the CoCr component starting with a pre-test roughness less than the TiNbN components.

From the measurements, the TiNbN appears to exhibit a scratch resistance greater than that of cobalt chrome. This finding is in accordance with work in pin on plate, and knee simulators [10-12]. The medial roughness rose from 0.034 to 0.115 for the titanium components, compared to a rise from 0.019 to 0.256 for the cobalt chrome. The scratch resistance of titanium niobium nitride has been demonstrated before in a hip simulator [7]. It must be noted that there was a greater difference in average roughness progression for the lateral condyles when compared to the medial. The average roughness for the CoCr lateral condyle increased over the five million cycles from 0.017 to 0.296, whilst the titanium lateral condyles changed from 0.036 to 0.074. The average roughness assessed across both condyles rose from 0.018 to 0.276 for the CoCr by the end of the test. Such a rise was far higher than that of the TiNbN rising from 0.0353 to 0.095. The difference between the medial and lateral components may be due to the direction of measurement relative to the direction of sliding during testing. Measurements were taken perpendicular to the anterior-posterior sliding and may over-estimate the roughness. Taking an average of 24 recordings for each condyle aimed to reduce the significance of error.

It is important to note that the influence of surface roughness on wear is minimal between 0.15 and 0.28 micrometers. After 5 million cycles only the CoCr surface roughness was greater than this at 0.296 on the medial condyle.

The use of a single cobalt chrome component was not optimum, however the decision to use three TiNbN and only one CoCr control in a four station knee simulator was based on the ISO14243 standards requiring analysis of three samples of the test material. At the end of five million cycles the titanium niobium nitride coat was intact to the naked eye conferring that it was retained [13-16]. Further experiments would have extra cobalt chrome controls to increase the power and enable
the serum or scanning electron microscopy of UHMWPE may have provided evidence for the specific wear mechanism.

Surface of the UHMWPE tibial inserts

Analysis of the surface of the UHMWPE did not reveal any differences between the volumetric wear of UHMWPE of the titanium and cobalt chrome. When interpreting surface profiling, creep must always be considered to be a cause of possible bias.

Conclusion

Despite a clear reduction in roughness progression over the course of this in vitro test, there was no demonstrable improvement in UHMWPE wear measured gravimetrically and by surface profiling. The implant tested may still be of great benefit to patients who are metal sensitive, but the coat offers no benefit in UHMWPE wear. The implant has not been marketed as an alternative to cobalt chrome. In those that suffer severe knee osteoarthritis but as an option to patients with metal ion allergies.

Acknowledgement

We are indebted to Mr Bob Skinner, Mark Harrison and Keith Rayner for their hard work for this study.

Conflict of interest

Supported in part by a research grant funded by Biomet. Biomet had no role in the study design, analysis or manuscript content. The authors have no affiliation with the funding company.

References


