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Design of an Affordable, Modular Implant Device for Soft Tissue Tension Assessment and Range of Motion Tracking During Total Hip Arthroplasty

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ABSTRACT **Background:** In hip arthroplasties, surgeons rely on their experience to assess the stability and balance of hip tissues when fitting the implant to their patients. During the operation, surgeons use a modular, temporary set of implants to feel the tension in the surrounding soft tissues and adjust the implant configuration. This process is naturally subjective and therefore depends on the operator. Inexperienced surgeons undertaking hip arthroplasties are twice as likely to experience errors than their experienced colleagues, leading to dislocations, pain and discomfort for the patients. **Methods:** To address this issue, a new, 3DOF force measurement system was developed and integrated into the modular, trial implants that can quantify forces and movements intraoperatively in 3D. The prototypes were evaluated in three post-mortem human specimens (PMHSs), to provide surgeons with objective data to help determine the optimal implant fit and configuration. The devices comprise a deformable polymer material providing strain-based displacements measured with electromagnetic-based sensors and an inertial measurement unit (IMU) for motion data. **Results:** Device results show a relative accuracy of approx. 2% and a sensitivity of approx. 1%. PMHS results indicated that soft tissue forces on the hip joint peak in the order of ~ 100 N and trend with positions of the leg during range of motion (ROM) tests, although force patterns differ between each PMHS. **Conclusion:** By monitoring forces and force patterns of hip soft tissues, in combination with standardised ROM tests, the force patterns could shed a light on potential anomalies that can be addressed during surgery. **Clinical and Translational Impact Statement:** The development of an instrumented hip implant device for use during surgery knowledge will eventually allow us to develop a predictive model for soft tissue balancing, that can be used for pre- and intra-operative planning for each patient on a tailored and personalised basis. Ultimately, we hope that with this device, patients will benefit from a faster recovery, from a more-precisely fitted hip, and an improved quality of life.

INDEX TERMS Medical device, surgery, hip arthroplasty, soft tissue balance, hip stability.

I. INTRODUCTION

Hip replacement is a surgical procedure undertaken to relieve osteoarthritis pain ($\sim 63\%$) [1] and to treat fractures by replacing the hip joint with an implant, commonly consisting of a shell, liner, head and broach (stem), as shown in Figure 1 (a-b). In the case of a total hip replacement (THA),

both the femoral head and acetabulum are replaced with an artificial ball and socket joint. It is in fact the most common arthroplasty procedure in The Netherlands, with more than 40.000 operations carried out in 2019 and steadily increasing [2]. Also, an increasingly younger population will be receiving this treatment [3], [4] while demanding more



FIGURE 1. The stock implant (Corail Hip Systems, DePuy Synthes, Johnson&Johnson, Warsaw IN, USA) was used as the starting basis for the design of the instrumented prototypes. (a) The permanent implant and (b) the temporary (trial reduction) implant with various modular neck and head geometries. (c) The instrumented prototype with instrumented necks and a modified head. The blue arrow indicates the 3DOF prototype and the red arrow indicates the 1DOF prototype. (d) The coordinate system used in this report; tension (or compression from the negative side of the axis) is positive while compression is reported as negative. (e) The inside of the 3DOF implant with the head removed shows the radial and axial O-rings, sensor placement and wiring.

from the treatment to suit their more active lifestyles at the same time. THA prevalence between gender is approximately 55%-45% split between women and men [4], and thus a high degree of personalised treatment is necessary due to body geometry differences. Therefore, improvements to implant fit accuracy and soft tissue balancing around the hip joint are necessary. One method to achieve this is through force measurements during the fitting of the implant to sense the tensioning of the surrounding soft tissues, which can be adjusted by changing the implant geometries during the surgery. A more well-balanced hip can lead to a quicker recovery to normal daily activities.

Soft tissue balance of THA is less investigated than that of total knee arthroplasties (TKAs) [5], [6], even though the outcomes of THAs also rely on the surgeon's experience to determine properly balanced soft tissues. Instability is the biggest cause of early implant failure, typically within five years of surgery [7]. Conventionally, a conservative approach would be to provide sufficient tension through slightly higher offsets to ensure stability, although this does come with drawbacks such as pain, fatigue and longer recovery time

until the muscles adapt over time. Larger femoral heads provide more stability and higher range of motion (ROM), but at the expense of higher impingement risks, liner fracture (due to thinner material) and higher wear rate because of the larger moment arm [8]. Too little tension, on the other hand, risks limping and even dislocation if the offset is not sufficiently restored [9]. Arguably, the optimum range of a "balanced" soft tissue tension within the hip is not simple to determine and is also subjective amongst surgical professionals.

The implant configuration is first predetermined using x-ray imaging and then confirmed intraoperatively with trial components before selecting a permanent implant by manipulating the leg in a series of movements with a ROM test, while the surgeon feels the muscles' resistance during the movements. The problem with the current ROM tests is that they are subjective, and there is no standard test protocol. Tang *et al.* [10], designed an inertial measurement unit (IMU) sensor-based implant and tested the accuracy/drift *in vitro*. Several studies indicate that soft tissue tension is related to pain (for instance in the groin and knee), dislocation, range of motion of the hip and limping [11]–[13]. Furthermore, the most common errors in THAs are made by inexperienced surgeons [7], [14], with up to twice as many dislocation incidences than experienced surgeons, decreasing by ~50% for every ten procedures performed annually, and plateauing after approximately 30 operations [15].

Researchers of the Orthoload implants developed three generations of instrumented "Hip I-III" of one-off versions of permanent hip implants that were used in clinical validation studies [16]–[18]. These latter implants were able to measure hip joint forces in everyday movement activities using embedded sensors that are powered through electromagnetic induction. While their work may initially appear to be similar to ours, our ultimate goal and design approach are fundamentally different. *The goal of this study is to measure and fine-tune the implant selection for hip replacements during the procedure.* It is also therefore not about quantifying the forces experienced in the hip after the procedure. Although we expect forces experienced in everyday activities to be different from our scope – lying down, under anaesthesia, the work from Orthoload provides a useful point of comparative reference.

We propose a measuring system for surgeons to obtain an objective force reference in combination with the standardised ROM tests. The system was developed based on the design of a commercially available trial implant (DePuy Synthes Corail) (Figure 1 (b)) to remain simple, modular, and affordable, while keeping the technical requirements including biocompatibility, water resistance, sterilisation compatibility, appropriate sizing, sensing magnitude, accuracy, safety, reliability and user-friendliness in mind. The bespoke neck and head components work with existing trial implants and the testing procedure regarding ROM trials. We first conceived a 1DOF device in the axial direction of the neck of the hip implant, as a proof of concept, since we expect

most of the forces are transmitted in that direction. Then, the sensing capability was increased to 3DOF by the addition of an IMU for motion sensing. The prototypes were tested and calibrated with a benchtop universal testing machine, and two pilot post-mortem human specimen (PMHS) studies ($n=3$ in total) were carried out to evaluate the prototypes.

II. MATERIALS AND METHODS

A. DESIGN RATIONALE

A temporary trial reduction implant based on a commercially available design has been made. The design can be swapped out easily during the intraoperative fitting process. The prototype uses linear Hall-effect sensors in combination with a compliant material to measure small deformations (~ 1 mm at 30 kg load) within the temporary implant head. Sensors ranging from strain-gauges to force-sensitive resistors (FSR) and optical sensors were evaluated. Hall-effect sensors were chosen because of their simplicity in electrical circuit design, space constraints, cost and reliability, and because they were effectively used in a knee balancer unit before [5]. These Hall sensors use a magnetic flux to determine their relative position to the source. Compliance with the structure was provided by a rubber O-ring between a neodymium magnet and the Hall-sensor to secure the head in its position and prevent fluid ingress. All components were selected to meet autoclave sterilisation (temperature, pressure, humidity) requirements; low-cost with the option to be reusable or single-use if desired. Prototypes are wired and connected to a microcontroller (Arduino UNO, Arduino AG, Turin, Italy) and a computer running Matlab (R2019a-R2020a Mathworks, Natick MA, USA).

1) IMPLANT DESIGN

In the first phase of development, we conceived a 1DOF device in the axial direction, as a working principle, since we expect most of the forces are transmitted in that direction. The instrumented implant is designed based on the trial reduction broach of the Corail Total Hip System (DePuy Synthes, Johnson & Johnson, Warsaw IN, USA) (**Figure 1 (b)**). Only the neck and head segments of the trial reduction components were utilised while retaining their exterior geometry (**Figure 1 (c)**). The head size used was $\phi 32$ mm with +1 mm offset (as the standard) and neck geometry was the 125° High Offset Coxa Vara Collared (KLA) variant. For the 3DOF prototype, additional necks were designed having variable offset (with spacers): 135° Standard offset (STD-135) and 135° High Offset (KHO) variants. Only the standard offset was used in the PMHS study, as this is the most commonly used option.

2) STERILISATION COMPATIBILITY

Electronic components were tested in an autoclave (MELAtronic 23 EN, MELAG, Berlin, Germany) through a standard cycle typical of hospital sterilisation equipment

(steam at 134 °C, 2 bar, 20 minutes) as per van der Pol, 2017 [19] to verify the survival of the components.

3) FIRST ITERATION: 1DOF PROTOTYPE

Device design: a 1DOF device was developed with milled and 3D printed stainless steel components. A hall sensor (A1302/A1308, Allegro MicroSystems, Manchester NH, USA) was implemented at the axial (z-axis) direction of the neck (see **Figure 1 (d)**). A magnet (5×1 mm circular N38) is attached to the opposite end and can slide freely (assuming small friction). Sandwiched in between is an O-ring ($10 \times 6 \times 2$ mm circular cross-section VMQ silicone rubber) acting as the spring element.

Force-voltage calibration of 1DOF: a compression test on the head was carried out to determine the relationship between applied force and voltage output of the sensors with a universal testing machine from 0-0.85 mm at 0.05 mm step size with 5 s of rest. From this, we obtain the approximate Hooke's law correlation between the amount of exerted force, compression of the O-ring, the distance between the magnet and the sensor and thus the sensor voltage reading. We also assume small hysteresis is experienced during the (de)compression of the O-ring. In other words, the energy lost between compressing and decompressing the O-ring is small enough to assume the same force-voltage relationship (indeed, this was tested and is shown in **Supplementary Figure 7**).

4) SECOND ITERATION: 3DOF PROTOTYPE

Device design: a 3DOF iteration was developed based on the 1DOF prototype with sensing in 3 axes and added an inertial measurement unit (IMU) gyroscopic motion sensor. The axial Hall sensor and O-ring setup was retained but three more Hall sensors were added on the sides of the neck to measure in the x-y plane (see **Figure 1 (d)**). The inner neck component was 3D printed using glass-filled nylon (PA 6 GF30, Hubs, Amsterdam, The Netherlands) while the neck base was made of CNC machined aluminium (7075-T651 grade aluminium, Weerg, Gardigiano, Italy). Magnets were N50 instead of N38 as used in the 1DOF version. Two larger O-rings ($15 \times 10 \times 2.5$ mm circular cross-section NBR nitrile rubber) on the neck, spaced equidistance apart from the hemisphere of the head provide displacement in the radial direction (**Figure 1(e)**). We assume small displacements of the O-rings and that any deviation/pivot about its centre causing the magnetic field and the sensor to misalign is small. Some crosstalk effects due to the proximity of different sensor-magnet pairs were isolated during the processing of the data by measuring how much the forces change in all principal directions (when compressing in one principal direction) and subtracting that deviation during data processing (see **Supplementary Figure 1**). The x-y plane Hall sensors were arranged 120° apart as shown in **Figure 2**. To convert the readings to principal forces by considering the voltage change, the location, angle and the plane of the sensor when a force is applied (thereby changing the distance between the magnets

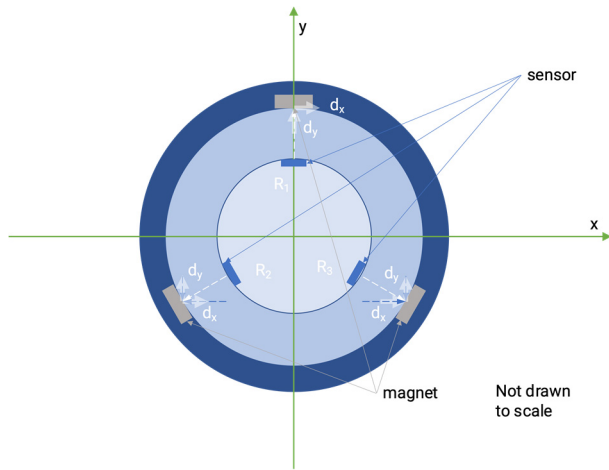


FIGURE 2. Schematic showing the x-y plane cross-section of the head with the three hall effect sensors and their resolved force vectors.

and the sensors), the following equations were used:

$$V_x = \frac{\sqrt{3}}{4} (V_3 - V_2) \tag{1}$$

$$V_y = \frac{V_1}{2} - \frac{V_2 + V_3}{8} \tag{2}$$

$$V_z = V_4 \tag{3}$$

Full derivation can be found in **Supplementary Information section 1**.

Force-voltage calibration of 3DOF: a compression test with the 3DOF prototype was carried out on the universal testing machine (LR 5K, Lloyd Instruments, Bognor Regis, United Kingdom) under a linear load-unload profile at 1 mm min⁻¹ to 1 mm displacement on 3 principal axes. An example of this is in **Supplementary Figure 2**. The peak voltage reading is correlated with the peak force reading from the indenter. The force-displacement curves resemble a linear fit, therefore, a linear scaling factor was used. The right-hand rule convention for positive direction was applied such that compression is negative. The forces reported in the 3DOF data were zeroed in the neutral position, i.e. the relative forces are presented. To test the accuracy and sensitivity of the 3DOF sensor, it was additionally validated by hand with weight pieces of 2 kg + 1 kg, at three random angles.

Isolation of forces in the principal directions: interference and crosstalk from surrounding magnets may occur when the magnet positions change due to the displacement of the O-rings. The amount of interference is proportional to the amount of compression. In the method to minimise the interference in one principal axis, the ratio between that and the cross talk from the two perpendicular axes is subtracted (for instance, the x-axis formula is shown in **Equation 4**). The overall result for this isolation method is shown in **Supplementary Figure 1** and it appears that unwanted interference was minimised. Furthermore, the raw data is shown additionally with Matlab smoothing filters of Matlab functions `medfilt1` and `sgolayfilt`. This was not used during the

PMHS experiments.

$$x_{new-x} = \frac{y^2}{x} - \frac{z^2}{x} \tag{4}$$

Hysteresis and stress relaxation: the degrees of hysteresis and stress relaxation of the compliant material – the polymer O-ring material were examined. Both the top and side O-rings underwent a linear load-unload, load-controlled compression using the universal testing machine at 0.67mm/s and 1 mm/s to 95 N (load cell limit of 100 N). For stress relaxation, a hold section of 30 s is added to the abovementioned test method between the loading and the unloading profiles. The relaxation curves were normalised for the two-term Prony series function in **Supplementary Figure 8**. The Prony coefficient for the instantaneous modulus was obtained in the same method as Crichton *et al.* [20] (see for derivation). The equation used for the two-term Prony series is as follows:

$$G(t) = 1 - g_1 \left(1 - e^{-\frac{t}{\tau_1}}\right) - g_2 \left(1 - e^{-\frac{t}{\tau_2}}\right) \tag{5}$$

Inertial measurement unit (IMU): a 9DOF IMU stick with the LSM9DS1 chip (SEN-13944, Sparkfun, Niwot CO, USA) was seated on the neck of the 3DOF unit to provide spatial coordinates to the user. The IMU connected to the same Arduino is used to process the Hall-effect sensors’ information using Matlab functions ‘`ism9ds1`’, ‘`readAcceleration`’, ‘`readAngularVelocity`’ and ‘`readMagneticField`’, complementaryFilter’ and converted to rotation in the principal axes [21]. Because of the default output of the IMU and the angle it is mounted on the neck, the principal axes were calculated, such that the x-axis points to the implant’s shoulder, and the z-axis points to the head (see **Figure 1 (d)**). The recorded ROMs were reviewed in Matlab for verification using the accelerometer-gyroscope-magnetometer fusion function ‘`ahrsfilter`’.

B. POST-MORTEM HUMAN SPECIMEN (PMHS) STUDY

1) APPROVAL STATEMENT

The PMHS studies were carried out per the Tissue Bank guidelines of Erasmus Medical Centre, Rotterdam, The Netherlands. All total body donors were part of the national donor programme and have given written consent for tissue donation for educational and scientific purposes before passing away. Under Dutch law, and these conditions no approval from the medical research ethics committee was required. Due to European privacy regulations, medical history was only available to the general practitioner of the donor and not to the receiving academic medical centre.

2) TEST PROTOCOL

The right legs of fresh (non-frozen-and-thawed/non-preserved) PMHSs (n=3, elderly age, exact age not provided) were used, having an already-exposed hip region from an earlier hip arthroplasty skills lab session using the anterior approach and bored femur and reamed acetabulum on the day. The *rigor mortis* phase past the initial time of death was

avoided. One PMHS was used for the 1DOF trial while two others were used for the 3DOF trial separately. Originally a fourth PMHS was prepared. However, the tissue conditions were unsuitable for trials and thus the fourth PMHS was excluded from the study. To prevent external electromechanical interference, we ensured that measurements were conducted without having other instruments near the surgical incision.

3) RANGE OF MOTION (ROM) PROTOCOL

The PMHS was in a supine position with the hip exposed through the anterior approach. The following tests were conducted for the 1DOF implant trial (0° means neutral position). Each movement was repeated 2-3 times.

- 1) External rotation from $\sim 0^\circ$ to 45° then to 90°
- 2) Internal rotation from $\sim 0^\circ$ to 45° then to 90°
- 3) Abduction from $\sim 0^\circ$ to 25°
- 4) Adduction from $\sim 0^\circ$ to 25°
- 5) Flexion of hip and knee from 0° to 45° then to 90°
- 6) Adduction-abduction to $\sim 25^\circ$ combined with a 45° rotation

For the 3DOF implant trial, the following tests were conducted:

- 1) Push, pull (proximal-distal)
- 2) External rotation
- 3) Flexion, extension
- 4) Flexion, external (+internal*) rotation, extension
- 5) External rotation (with dislocation)*

*For the second PMHS trial only

We allowed a three-second waiting time and zeroed the readings before the commencement of each test.

III. RESULTS

To design a soft tissue tension measuring device for hip arthroplasty, we first developed a 1DOF prototype using simple, off-the-shelf Hall-sensor based electronics and tested it in a PMHS study. This was followed up by an improved 3DOF prototype which was tested in two PMHSs.

A. DEVICE PERFORMANCE CHARACTERISTICS

1) PRINCIPAL AXES CALIBRATION

The indentation example shown in **Supplementary Figure 1** shows the raw readings (top row) converted to principal axes (middle row), and then with crosstalk effects minimised (bottom row). Both the crosstalk effects and the noise appeared to be filtered out well.

2) RUBBER O-RING CHARACTERISTICS: HYSTERESIS, STRESS RELAXATION AND STERILISATION COMPATIBILITY

In linear load-unload indentations of two speeds (0.67 mm/s and 1 mm/s), most curves (**Supplementary Figure 8**) appeared to have minimal hysteresis effects for the O-ring material used in the device (the return curve follows closely with the initial curve; the area between the outbound and inbound curves are small).

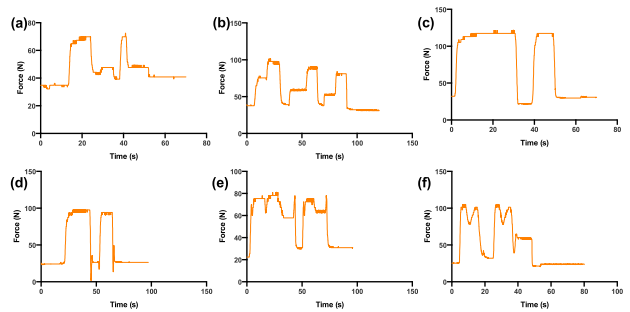


FIGURE 3. 1DOF in the axial direction showing various ranges of motion performed on one PMHS: (a) external rotation of the hip, (b) internal rotation, (c) abduction, (d) adduction, (e) Flexion and (f) abduction and adduction, plus external rotation.

In terms of stress relaxation, the load-hold-unload indentations of 30 s showed that the material's resistance to a constant force decreased sharply within 2-3 seconds after applying a constant force and that after a fitting to a two-term Prony series, we arrived at an average Prony series coefficient for each indentation velocity/O-ring-type combination of 0.92-9.96 (**Supplementary Table 1**). In other words, the forces measured may be up to approximately 8% lower than the true force in the load-hold-unload experiment after about 10 s. Consequently, we take the stress relaxation effects are small as a reasonable approximation.

B. PMHS STUDY

1) INSERTION OF THE IMPLANT ONTO THE STEM IN THE PMHS

The instrumented device was placed directly atop the stem in the same way as placing a regular neck and stem without hindrance to usability and user-friendliness. The data cable connected to the implant leads out the opening of the hip area to the Arduino and computer for recording and processing.

2) 1DOF PROTOTYPE

We observed distinct changes in force patterns of soft tissue forces acting on the implant head as the leg was moved during the range of motion trials. **Figure 3 (a-f)** shows a force range of approximately 20-120 N, depending on the ROM. Distinct patterns were observed for specific positions, for instance, (a-b) at the peak (90°) of each internal/external rotation. Notably, at the halfway point of the motion (45°), the force reading was not exactly halfway between the force values corresponding to 0° and 90° and was closer to the force value of the neutral (0°) position than the 90° position. For internal rotation (b), the midway force varied depending on the repetition, and an overall trend with decreasing in absolute force was observed. Both abduction and adduction (c-d) movements from the neutral position to $\sim 25^\circ$ produced a peak force in principal axes of approximately 120 N and 100 N, respectively. For the flexion of the hip in the combination of the flexion of the knee (e) from 0° to 45° , a forces were higher than at 90° . In the final combined ROM (f), the forces peaked at the end positions (25°) of abduction and adduction

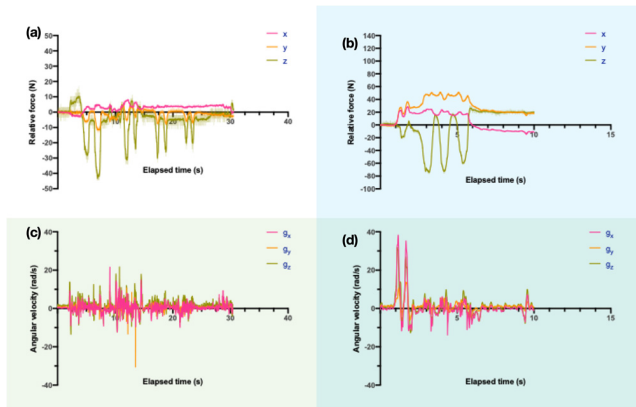


FIGURE 4. 3DOF of the push-pull (proximal-distal) range of motion performed on two PMHSs (left and right columns respectively), with the top row (a, b) showing the net resultant force in each principal direction and the bottom row showing the angular velocities (c, d) from the IMU showing the rotational movements of the hip.

of approximately 100 N, and a small dip to approximately 80 N was observed when the leg was moving from abduction to adduction. The additional exorotation movement appeared to differ between the first and second cycle – the first force reading of the first exorotation was closer to the starting force than the second exorotation. During certain movements such as an internal rotation(b), the peak forces decreased after each successive motion, although the pattern generally remained the same. In most cases, the start and finish force readings at the neutral position differ (a-b at ~40 N c-f at ~25 N), except for the (d) adduction ROM.

3) 3DOF PROTOTYPE

The results of the PMHS study carried out with the 3DOF prototype (2x replicates) are shown in the left and right columns of **Figure 4 – Figure 7**. The forces (top row) are in a range of approximately –40 N to 100 N for PMHS C1 and within –70 N to 50 N for PMHS C2. Additional angular velocity readings (bottom row) are presented, illustrating the ‘consistency’ of the surgeon’s input and speed in moving the leg through the various ROMs, which peaked at approximately 60 rad/s for most movements. The additional validation done by hand with weight pieces, shown in **Supplementary Figure 9**, shows an estimated accuracy of approx. 2 N and a sensitivity/resolution (taken as the coarsest fluctuation before Matlab filter smoothing) of approx. 0.85-1 N. Raw readings directly from the sensors are shown in **Supplementary Figure 5**.

During **push-pull**: the first movement (**Figure 4**), most of the compression force in the head is in the z-axis direction. For both replicates, the peak force for each compression is at approximately 40 N and 70 N. Note that the final position does not necessarily return to zero. The initial “moving the leg into position” in the second replicate showed a much higher (almost 40 rad/s) rotation rate than the push-pull motion, and as expected, the angular velocity for the entire

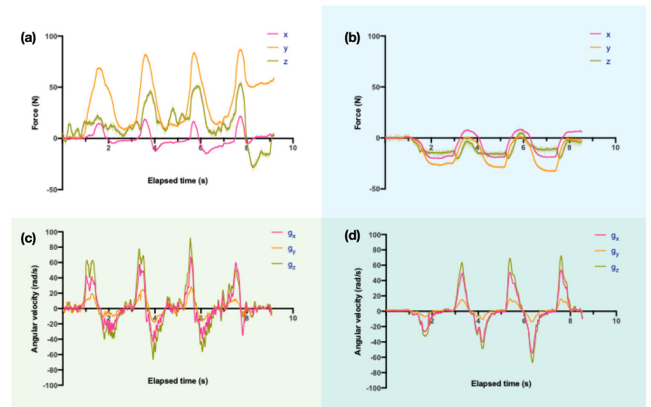


FIGURE 5. 3DOF measurements of the hip external rotation ROM performed on specimen 1 (left) and specimen 2 (right), with the top row (a, b) the net resultant force in each principal direction and the bottom row the angular velocities (c, d).

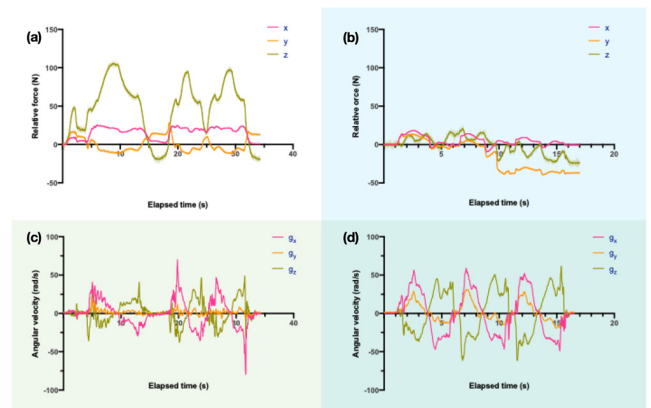


FIGURE 6. 3DOF measurements of flexion-extension ROM performed on Specimen 1 (left) and Specimen 2 (right), with the top row in (a, b) the net resultant force in each principal direction and the bottom row the angular velocities in (c, d).

ROM remains small (>10 rad/s) as the leg does not rotate (see **Supplementary Figure 3 specimen 1-2 push-pull**).

During **exorotation** (**Figure 5**), the two specimens exhibited distinct patterns during the ROM albeit different magnitudes between the two replicates. The forces all peaked at the end (limit) of the movement corresponding to where the angular velocity curves cross the x-axis. The principal y-axis direction reading of the end section of the ROM for the first replicate appeared to have plateaued at approximately 50 N instead of close to zero.

The forces observed for the **flexion** movement in **Figure 6** show, again, two different magnitudes acting on the joint, although the patterns were still discernible, especially in specimen 1. A tightening of soft tissues was observed at the peaks of the flexion movement, shown primarily with a positive z-axis reading. The y axis reading of specimen 2 shows a sudden decrease during the second cycle, but this is not reflected in the IMU data. Forces for Replicate 2 also appeared to decrease first and then increase at the peak of the movement, likely coinciding with the subtle movement

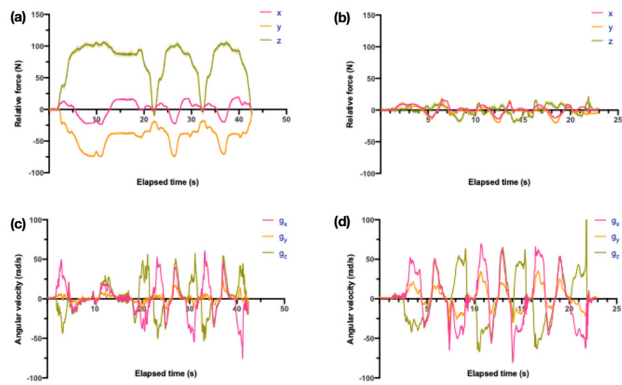


FIGURE 7. 3DOF measurement of the hip flexion, external rotation and extension ROM performed on specimen 1 (left), and an additional, internal rotation on specimen 2(right) with the top row (a, b) the net resultant force in each principal direction and the bottom row the angular velocities (c, d).

endorotation between 1-2 second timeframe. The angular velocity patterns show that the movements of the ROMs between the two were comparable, although, for specimen 1, the rotation rate was smaller than specimen 2, likely due to large resistance coming from soft tissues.

In the combined ROM of **flexion, external rotation and extension**, the data is presented in **Figure 7**. Consistently, specimen 1 exhibited higher forces (peaks at approximately -70 N to 100 N) whereas specimen 2 showed much smaller magnitudes at less than -20 N to 20 N. specimen 1's force readings and patterns were comparable to the flexion ROM as presented in **Figure 6**, but with the additional dip accounting for the external rotation. specimen 2, however, also displayed distinct patterns, although while the forces were much smaller, the external and internal movements were captured in the forces as a dip in the x and y axes and then an increase in the same axes. Instead of returning to the neutral position after an external rotation of specimen 2, the motion was continued to an internal rotation.

In the final ROM carried out additionally at the end of the experiment session, a combined **external rotation with deliberate dislocation** motion was carried out as an exploratory test following the final ROM mentioned above of specimen 2, and the result is presented in **Supplementary Figure 6**. The force curves began similar to the exorotation ROM of **Figure 5 (c)**. However, a deliberate manoeuvre of the hip joint to cause it to dislocate was identified as a jolt (just before 10 s elapsed time) in the figure.

Acceleration and magnetic field data from the IMU: the acceleration magnitudes are shown in **Supplementary Figure 3** and reported in standard gravity g (9.81 m/s²). From here we observed that the y axis is at approximately 1 when at the neutral position from gravity. The readings of all three axes illustrate the position of the leg during the range of motion with the relation between the implant reference frame (**Figure 1 (d)**) and the earth reference frame. The magnetometer data presented in **Supplementary Figure 4**

with units in *gauss*, also showing some similarities, when compared with the acceleration data in terms of rising and falling of patterns in the principal axes, is used to provide orientation reference when the motion is reanimated later in Matlab.

IV. DISCUSSION

In this paper, we presented two design iterations of a novel, low-cost medical device prototype, capable of measuring forces and orientation data for use during hip arthroplasties and trialed them on three PMHSs. The ability to obtain soft tissue tension measurements of the hip easily will especially assist inexperienced surgeons, who cannot 'feel' the right level of tension, as well as experienced surgeons to provide patients with adequate hip stability. To our knowledge, this is the first device designed to measure soft tissue tension of the hip intraoperatively, tested in an *in situ* pilot study.

A. GENERAL OBSERVATIONS

Overall, the force profiles of ROMs carried out on 3 PMHSs (1x 1DOF, 2x 3DOF with motion data) exhibited different magnitudes, with peaks up to approximately 120 N, 100 N and 50 N for 1 (1DOF), 1 and 2 (3DOF), respectively, with motion profiles showing the position at which these peaks happen – typically at the extreme ends of the motion. The rotational rates of the hip manipulated by the surgeon during the test also appeared more consistent (peaking at approximately 50 rad/s) in the 3DOF study than the forces measured. The device was inserted into the PMHSs' hips without difficulty or change to standard surgical protocol apart from the additional cable that connects the device to the Arduino. In our case, the surgeon continues with the operation and the ROM tests, while another OR team member monitors the data readout.

The forces are relative to the neutral position at the beginning of the ROM. In many cases, the forces at the end of the ROM recording do not return to zero. We suspect this could be due to some bodily fluids having shifted to other regions, and/or the fact that soft tissues are highly viscoelastic and possess hysteresis properties, so residual forces remain acting on the joint and sufficient time (~ 1 minute) and 'loosening' (i.e. a few 'wiggles') of the leg back at its neutral position should be carried out. Furthermore, the overall decreasing trend in absolute forces is probably due to the stretching and relaxing of the PMHS muscles. Additionally, the soft tissue behaviour of a PMHS may be different than that in a living patient (discussed further below in section IV-B). Regardless, in future, device accuracy could be calibrated or checked before and after an experiment to identify drifts.

Interestingly, even though the force readings appear to differ between the two 3DOF PMHSs, the angular rotational rates, as well as the acceleration vectors suggest that the ROMs were carried out consistently between the two. This discrepancy and the residual forces, most likely down to individual differences between the two PMHSs (which further stresses the need for a higher level of personalised treatments

in orthopaedic surgery with tissue stability and balance due to the variability surgeons need to recognise), could eventually be used as an indicator for surgeons to monitor for unexpected forces or force patterns at each position of the ROM tests, taking into account the body geometries and soft tissue volumes. We also expect different approaches (e.g. posterior, lateral and anterior) to have a significant influence on the overall force measurements of the soft tissues.

B. IN SITU VS IN VIVO CONDITIONS

Data collected from *in situ* conditions in our study may be different compared to *in vivo* conditions – we expect somewhat higher force readings during *in vivo* readings, at least partly due to circulation of bodily fluids [22], but it is still useful as insight and a steppingstone before the prototype is tested in clinical settings. Perhaps, we hypothesise that there will be more damping, hyperelastic and time-dependent effects and higher forces *in vivo* and would form a basis for an *in situ* vs. *in vivo* comparative study. An earlier pre- and post-mortem soft tissue (skin) mechanical behaviour study [23] may foreshadow some of the effects already. Nevertheless, we were able to establish the order of magnitude of typical forces (~100 N) experienced for *in situ* conditions and can optimise subsequent device designs to provide a higher resolution for this expected range of forces. Additionally, the values we obtained compared to published data from e.g. Orthoload were in the same order of magnitude – granted that their values were taken from living patients without anaesthesia. They reported a peak load of approximately +150-250 N for 45° flexion and ± 150 N for hip abduction (Orthoload online database) [18].

C. DEVICE DESIGN CONSIDERATIONS

Regarding the (in)accuracy of using polymer materials (rubber) as the deformable material within the implant, we believe after showing the materials have small hysteresis (e.g. <10%) and viscoelastic effects (at least for the tested compression rate and the linear load-unload tests anyway), it is acceptable for our purpose when measuring muscle tissues that have greater hyperelastic and hysteresis effects due to tissue fluids [24]. A proportional-integral-derivative (PID) loop built into the post-processing of the data may also mitigate this effect [25], although this may slow down the refresh/sampling rates of the overall system as well as introduce unnecessary complications to the device. Not every surgeon will perform the movements the same way and would be difficult to implement with human factors involved.

Rubber O-rings used herein were also found unable to withstand an autoclave sterilisation environment, so new, sterile O-rings must be used for future *in vivo* studies or a different method of sterilisation. Alternative materials such as metallic-based deformable structures could eventually substitute the O-ring (which would also eliminate any undesirable polymer characteristics such as hysteresis and stress relaxation), however, another material is required to seal the internal structure as the O-ring also doubles as a fluid

barrier and may require custom manufacturing. Furthermore, it is plausible that increasing the accuracy marginally may substantially increase the cost and complexity of this device. Regardless, to minimise capturing hyperelastic and hysteresis effects from both the rubber O-ring and biological tissues during the experiments in the current design, we propose that waiting three seconds to allow the rubber and soft tissues to enter a steady relaxation state [26], we minimise the variability from soft tissues. This should be an important step to include in all future *in situ* and *in vivo* studies.

D. FUTURE OUTLOOK AND POTENTIAL

We propose that this device eventually can be used to ensure ROMs are carried out consistently (through the use of IMU movement data) – e.g. have the surgeon check the flexion or rotation angle and the rate of motion is carried out to a predetermined position and velocity, and check if the resistance of the soft tissue tension is as expected with the force readings. With sufficient repetitions, experienced surgeons can begin devising “gold standard” guidelines on ROM protocols – at the moment, this differs per surgeon. The following step would be to use the collected (*in vivo* and *in situ* and identify differences) patient body build data (bone geometry and soft tissue volume), measured forces and movements (and through the input from multiple surgeons to identify ‘good’ and ‘bad’ cases) to build a predictive model. We propose to normalise the forces and use the force ratios at different ROM positions as a point of reference for modelling. We foresee this technology will eventually supplement current surgeries by suggesting to the surgeon intraoperatively, for a given patient and a given ROM, the ideal range of force response. If the measured force is outside the “optimum” range, then the device could prompt the surgeon to adjust the implant or to change the geometry, such as neck angle and/or offset, implant and/or head/cup sizes. This will ultimately lead to a more personalised medical approach, tailoring each implant operation more specifically to suit individual patients. At the same time, we also minimise the subjectivity of different surgeons.

E. FUTURE WORK AND LIMITATIONS

The phenomenon of the forces not zeroing at the end of the ROM trial needs future investigation. In a follow-up study, we would like to emphasise the discrepancy, variation and patterns amongst patients, and investigate how these factors influence the interpretation of the results concerning hip stability and tissue balancing. The effects of anaesthetics on muscle forces (between anaesthetised and non-anaesthetised states) also need to be further investigated. Apart from longer-term data-collection experiments already mentioned, an improvement to the current device design can further lock any potential rotation of the head to the neck, as any rotation/misalignment can potentially influence the x- and y-axis readings when the distance between the Hall sensors and the magnets changes. Therefore, we suggest changing the fitting/base of at least one of the O-rings to a triangular

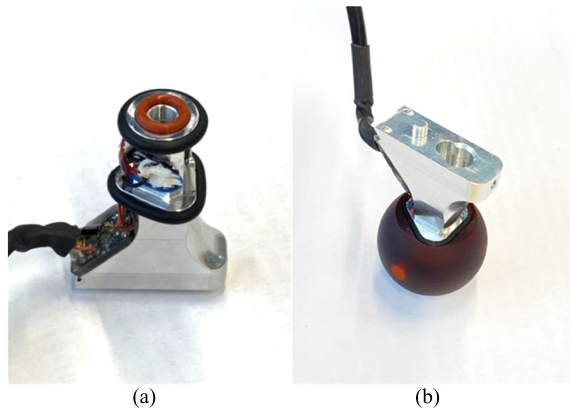


FIGURE 8. Improved triangle base design with head rotation lock allowing more precise sensor-magnet alignment for future use. (a) is without the head, viewed from the top/side, and (b) viewed from below, with the head fitted.

shape to improve the alignment of the magnet-sensor pairs. An example of this is shown in **Figure 8**.

A further study may be interesting to compare differences between *in vivo* datasets with *in situ* PMHS experiments, as well as *in vivo* anaesthetised patient experiments. We hypothesise in the latter case, the muscles will mostly be relaxed and may give lower force readings. On top of this, we were also able to distinguish that force readings correlate to positions of ROM movements. The ROM protocols we used in this study are also useful – at a minimum, surgeons should carry out a flexion-extension ROM, external and internal rotations, and a combination of those movements to capture as much of a range of the hip movement as possible.

Finally, magnet sizing and sensor-magnet pair placement should also be optimised to minimise interference. A combined simultaneous capture of force and movement *in vitro* experiment should also be carried out as further validation.

V. CONCLUSION

In this work, we developed two force-measuring hip implant prototype devices, first in 1DOF as a proof-of-principle, then in 3DOF with integrated IMU capabilities, and used them in three PMHS ROM trials. The goal is to provide surgeons with real-time, objective measurements to facilitate them during surgery in selecting the optimal implant position and geometry for the patient to improve patient outcomes, such as minimising pain and the likelihood of dislocations from an unbalanced hip. Stress relaxation and hysteresis of the device structure were deemed to be small (<10%), especially when compared to that of biological tissues. Thus, we proposed that the use of off-the-shelf materials was suitable to keep the design simple and low-cost. In the experiments, we found force values ranging in the order of ~ 100 N, also comparable to literature data, however, both force magnitudes and patterns differ between PMHSs. We suggest that using both force and position data could help identify undetected anomalies in implant fit when ROM tests are carried out by the surgeon.

Finally, we suggest that future work should focus on device refinement and optimisation for clinical use, investigate a baseline of “optimally balanced hip”, devise standardised guidelines for ROM tests and collect data in (pre)clinical trials, to build up a database of patient body geometry and force measurements. These investigations will eventually lead to a higher level of personalised treatment for individual patients, increased success rate and quality of THAs, while reducing the cost on the healthcare system (e.g. reduced recovery time) and added value to patients, healthcare providers and implant makers.

DATA AVAILABILITY

The datasets generated and/or analysed during the current study are available from the corresponding author upon reasonable request.

COMPETING INTERESTS

The authors declare no competing interests. DePuy Synthes did not perform any of the experiments nor played a role that could have influenced the outcome of the results. DePuy Synthes provided demo trial implants and CAD drawings as a collaborative partner listed in this grant.

AUTHOR CONTRIBUTIONS

Jonathan C. J. Wei wrote the manuscript and was the main contributor to the 3DOF prototype development, data analysis, and associated experiments. Bryan Blaauw performed the ROMs and provided data analysis and critical oversight of the manuscript. Dieter G. M. van der Pol was the main contributor to the 1DOF prototype development, data analysis, and associated experiments. Mauricio Cruz Saldívar assisted with the design and mechanical testing of the materials. Chung-Feng Lai assisted with validating the derivation of the 3DOF equations. Jenny Dankelman and Tim Horeman provided critical oversight to the prototype designs, data analysis, and manuscript revisions.

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