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Introduction

Prostate cancer is the second most common form of cancer found in men. A prostate biopsy is done to collect small tissue samples when the prostate-specific antigen level in a person's blood is elevated, or when an abnormality is found during a person's digital rectal exam. Trials¹⁻² suggest that magnetic resonance imaging (MRI) is preeminent in prostate biopsies due to its ability to differentiate between diseased and normal tissue accurately. However, recent studies suggest that MRI-guided prostate biopsies suffer from inaccuracy due to stiff anatomical structures, such as the pelvic diaphragm, which causes major deflections³⁻⁴. <u>In-vivo force measurement is</u> crucial for differentiating stiff structures and potentially improving the needle placement <u>accuracy. We developed and validated an</u> <u>attachable needle force sensor that is</u> <u>MRI-compatible (ANFS-M) to achieve this goal.</u>



Results

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Methods



Fig. 1. Design and force distribution of the ANFS-M <u>Validation:</u> Our working hypothesis is that the



Fig. 3. We assume the following simple mathematical model to predict the thrust force on the needle: F = aD + bv + c

where is the force (N), is the displacement of the needle tip from the entry point (mm), and is the velocity of the needle tip (mm/s). and are the weighting factors for the displacement and the velocity respectively, whereas c is the offset.

Discussion

During the phantom punctures the displacement of the needle tip was found to be proportional to the force, as the deeper the needle entered in the phantom the more friction it faced. Additionally, the force also depends on the velocity of the needle tip, though the relationship varies based on the stiffness of the phantom. In Fig. 3. A. & B. the pliability of the phantoms along with human instinct to reduce speed when meet with a force caused velocity to be low when the force was high. In Fig. 3. C. the greater stiffness allowed for more uniform velocities. Still, the ANFS-M differentiated the stiffness of the phantoms as evident by the steeper slopes for the phantoms made with higher concentrations of agar present in Fig. 2 & 3 (15g \rightarrow 2.65 * 10⁻³ < 20g \rightarrow 3.82*10⁻³ < 25g \rightarrow 0.0434).

ANFS-M can differentiate tissues with different stiffnesses. The ANFS-M was used in needle insertions into three homogeneous phantoms made from different concentrations of agar (0.0291/15g, 0.0385/20g, and 0.0483/25g) labeled accordingly as 15g, 20g, and 25g. Prior to experimentation the load cell was calibrated through the use of known masses. The displacement of the needle tip was recorded using an optical 3D tracking system (Polaris Vicra, Northern Digital Inc.). We used the force data collected by the ANFS-M, along with the displacement and velocity to model the total axial force and friction force exerted on the tip of the needle.



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