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(54) Title: TRANSORAL ROBOTIC SIMULATOR

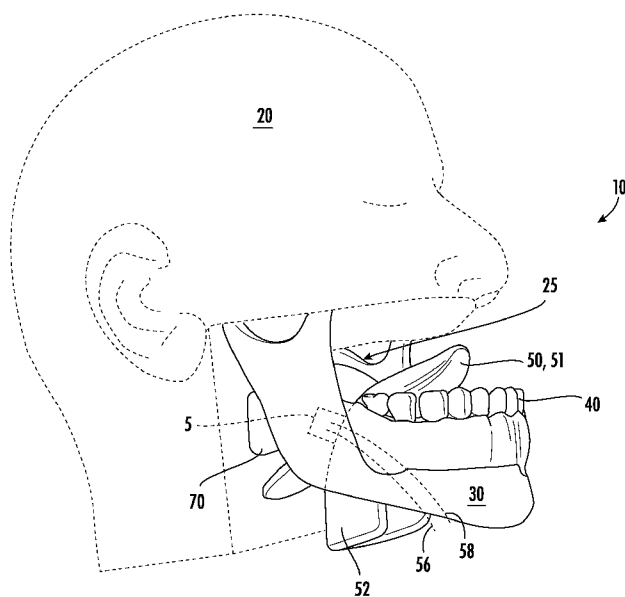


FIG. 6B

(57) Abstract: A transoral robotic surgery (TORS) simulator system includes a three-dimensional human head model having an oral cavity made of a synthetic material. The oral cavity has a mandible structure and a simulated human tongue and/or a simulated human tonsil. The simulated human tongue and/or the simulated human tonsil are made from a silicone material. The simulator system also includes artificial tissue(s) attached within the oral cavity, the artificial tissue being formed to mimic a biological tissue, whether cancerous (e.g., a tumor) or otherwise. The simulator system also includes a marker material present on and/or within the artificial tissue; the marker material allows a user of the simulator system to visually differentiate between the artificial tissue and synthetic material of the oral cavity.

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TRANSORAL ROBOTIC SIMULATOR

CROSS-REFERENCE TO RELATED APPLICATIONS

- 5 This application claims priority to U.S. Patent Application Serial No. 63/162,320, filed on March 17, 2021, the disclosure of which is incorporated by reference herein in its entirety.

GOVERNMENT SUPPORT

- 10 This invention was made with government support under grant number DK007386, awarded by the National Institutes of Health. The government has certain rights in the invention.

BACKGROUND

- 15 Within the last decade the use of robotic platforms has grown, with the da Vinci robot (Intuitive Surgical Sunnyvale, CA), being the most widely used system to date. Compared to laparoscopic techniques, current robotic systems are capable of restoring hand-eye coordination, augmenting surgeon dexterity, improving visualization, and providing ergonomic positioning.⁴
- 20 Technologic advances in robotic surgery have resulted in novel surgical applications, including transoral robotic surgery (TORS) for the treatment of oropharyngeal malignancies.

- The Da Vinci robot received FDA approval for treatment of T1 and T2 oropharyngeal malignancies in 2009; since then, TORS has been applied to
- 25 larger lesions and expanded to treat other conditions, including sleep apnea, pediatric airway lesions, and submandibular sialolithiasis. A key factor limiting advancements in TORS is the amount of training and exposure, or lack thereof, required for novice surgeons to safely operate the Da Vinci surgical robot. While TORS has proven to be an effective therapy in experienced
- 30 hands, the learning curve for the robotic procedures is slow and poorly understood, as the novice surgeon lacks the case volume to rapidly improve surgical competency; currently known training systems, devices, and methods are not capable of simulating TORS procedures.

Robotic training for TORS with the Da Vinci robot is highly variable by institution; however, successful implementation of robotic training curricula is increasing. Although there is no mandated credentialing process for robotic surgery, in 2015 the AHNS Education committee, AAO-HNS Robotic Task
5 Force, and the AAO-HNS Sleep Disorders Committee put forth training standards, which recommended a comprehensive robotic training curriculum for surgeons, including a skills acquisition phase that includes virtual reality, inanimate, and cadaveric simulation modalities.

Even at present, the need for TORS-specific simulator systems,
10 devices, and methods for trainees (e.g., novice surgeons, at least with respect to aspects of surgical techniques performed using robotics) to acquire necessary psychomotor skills remains unmet. In order to provide optimized integration into a robotic surgery curriculum, the TORS-specific simulator systems, devices, and methods disclosed herein maximize fidelity and
15 usability while minimizing operating cost. Further, the allow trainees to develop hands-on robotic experience with repetitive, task-specific exercises while maintaining a balance between fidelity and usability. Furthermore, results of a pilot study in which the capability of an example embodiment of a novel TORS-specific surgical simulator are presented herein, in which
20 operators of varying experience using the Da Vinci SI robot are differentiated.

SUMMARY

The presently disclosed subject matter now will be described more fully hereinafter, in which some, but not all embodiments of the presently disclosed
25 subject matter are described. Indeed, the presently disclosed subject matter can be embodied in many different forms and should not be construed as limited to the embodiments set forth herein; rather, these embodiments are provided so that this disclosure will satisfy applicable legal requirements.

According to an example embodiment, a transoral robotic surgery
30 (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil,

wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the TORS simulator system is configured for use with a robotic surgical system controlled by the user.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the oral cavity comprising synthetic material is made via an additive manufacturing process.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the simulated human tongue is molded using a mold formed via additive manufacturing.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the mandible structure, the simulated human tongue, and/or the simulated human tonsil is/are modular, such that each of the mandible structure, the simulated human tongue, and/or the simulated human tonsil are individually removable and/or replaceable from the head model.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible

structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the marker material is injected at a base of the at least one artificial tissue.

10 According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil
15 comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; and one or more electrically conductive wires or materials integrated into the oral cavity, optionally, wherein the simulator
20 system is configured for monopolar capability.

 According to an example embodiment, a transoral robotic surgery
25 (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil
30 comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a

user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the at least one artificial tissue is configured to comprise monopolar capability.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; and one or more electrically conductive wires or materials integrated into the oral cavity, optionally, wherein the simulator system is configured for monopolar capability; wherein the one or more electrically conductive wires are connected to an electrical ground for electrical grounding, allowing for monopolar electrocautery capabilities.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the marker material comprises a colored dye, wherein the colored dye is visible to the user of the simulator system

performing a simulated surgical procedure and is configured to provide a margin for reference during resection of the at least one artificial tissue.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system
5 comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the
10 oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the simulated human tongue and/or the
15 simulated human tonsil are configured to allow for attachment of the at least one artificial tissue thereto for the user to perform a tonsillectomy and/or base of tongue (BOT) resection.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system
20 comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the
25 oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; and a perfusion channel integrated into the
30 synthetic material and/or the at least one artificial tissue, wherein the perfusion channel is configured to contain, transport and/or be perfused with a liquid configured to mimic blood and, optionally, wherein the perfusion channel is integrated into the at least one artificial tissue.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; and a copper tape, mesh, wire, or other electrically conductive material that is integrated into, or attached within, the oral cavity and is configured to detect contact with a surgical instrument during use of the simulator system.

According to an example embodiment, a transoral robotic surgery (TORS) simulator system is disclosed herein, the TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material; at least one artificial tissue attached within the oral cavity, wherein the artificial tissue is configured to mimic a biological tissue; and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; wherein the biological tissue is a cancerous growth tissue.

According to an example embodiment, a robotic surgical training system is disclosed herein, the robotic surgical training system comprising a TORS simulator system according to any of the example embodiments described herein; and an evaluative assessment criteria configured to assess the user's use of the simulator system.

According to an example embodiment, a robotic surgical training system is disclosed herein, the robotic surgical training system comprising a TORS simulator system according to any of the example embodiments described herein; and an evaluative assessment criteria configured to assess the user's use of the simulator system; wherein the evaluative assessment criteria comprises a Global Evaluative Assessment of Robotic Skill (GEARS), or a modified version thereof.

According to an example embodiment, a robotic surgical training system is disclosed herein, the robotic surgical training system comprising a TORS simulator system according to any of the example embodiments described herein; an evaluative assessment criteria configured to assess the user's use of the simulator system; and one or more cameras configured to capture images during use of the TORS simulator system and/or video recording devices configured to capture video during use of the TORS simulator system.

According to an example embodiment, a robotic surgical training system is disclosed herein, the robotic surgical training system comprising a TORS simulator system according to any of the example embodiments described herein; an evaluative assessment criteria configured to assess the user's use of the simulator system; and a computing device communicatively connected to the TORS simulator system, the robotic surgical system, and/or the one or more cameras and/or video recording devices.

According to an example embodiment, a robotic surgical training system is disclosed herein, the robotic surgical training system comprising a TORS simulator system according to any of the example embodiments described herein; an evaluative assessment criteria configured to assess the user's use of the simulator system; and a graphical user interface to display images, videos, and/or data from the surgical training system.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and

a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the evaluative assessment is performed using an evaluative assessment criteria for assessing the user's use of the simulator system.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising

synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the evaluative assessment provided to the practitioner is based on data from a Global Evaluative Assessment of Robotic Skill (GEARS), or modified version thereof.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the user performs the simulated TORS procedure on the TORS simulator system using a robotic surgical system controlled by the user.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method

comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the
5 simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a
10 user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein providing the TORS simulator system comprises forming
15 the oral cavity via additive manufacturing.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising
20 synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological
25 tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; providing an evaluative
30 assessment to the user regarding the user's performance of the TORS procedure; and forming a mold for the simulated human tongue via additive manufacturing, wherein providing the TORS simulator system comprises molding the simulated human tongue within the mold.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the mandible structure, the simulated human tongue, and/or the simulated human tonsil is/are modular, such that each of the mandible structure, the simulated human tongue, and/or the simulated human tonsil are individually removable and/or replaceable from the head model.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative

assessment to the user regarding the user's performance of the TORS procedure; wherein providing the TORS simulator system comprises injecting the marker material at a base of the at least one artificial tissue.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein providing the TORS simulator system comprises integrating one or more electrically conductive wires or materials into the oral cavity, optionally, wherein the simulator system is configured for monopolar capability.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a

user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the at least one artificial tissue comprises monopolar capability.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the at least one artificial tissue comprises monopolar capability; and wherein providing the TORS simulator system comprises connecting the one or more electrically conductive wires to an electrical ground for electrical grounding, allowing for monopolar electrocautery capabilities.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a

silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the marker material comprises a colored dye, wherein the colored dye is visible to the user of the simulator system performing a simulated surgical procedure and is configured to provide a margin for reference during resection of the at least one artificial tissue.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein the simulated human tongue and/or the simulated human tonsil are configured to allow for attachment of the at least one artificial tissue thereto for the user to perform a tonsillectomy and/or base of tongue (BOT) resection.

According to an example embodiment, a method for training a user in transoral robotic surgery (TORS) techniques is disclosed herein, the method

comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the
5 simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a
10 user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS procedure; wherein providing the TORS simulator system comprises forming
15 a perfusion channel in the synthetic material and/or the at least one artificial tissue, wherein the perfusion channel is configured to contain, transport and/or be perfused with a liquid configured to mimic blood and, optionally, wherein the perfusion channel is integrated into the at least one artificial tissue.

According to an example embodiment, a method for training a user in
20 transoral robotic surgery (TORS) techniques is disclosed herein, the method comprising providing a TORS simulator system comprising a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the
25 simulated human tongue and/or the simulated human tonsil comprise a silicone material, at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue, and a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a
30 user of the system between the at least one artificial tissue and the synthetic material of the oral cavity; having the user perform a simulated TORS procedure on the TORS simulator system; and providing an evaluative assessment to the user regarding the user's performance of the TORS

procedure; wherein a copper tape, mesh, wire, or other electrically conductive material is integrated into, or attached within, the oral cavity and is configured to detect contact with a surgical instrument during use of the simulator system.

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BRIEF DESCRIPTION OF THE DRAWINGS

The presently disclosed subject matter can be better understood by referring to the following figures. The components in the figures are not necessarily to scale, emphasis instead being placed upon illustrating the principles of the presently disclosed subject matter (often schematically). In the figures, like reference numerals designate corresponding parts throughout the different views. A further understanding of the presently disclosed subject matter can be obtained by reference to an embodiment set forth in the illustrations of the accompanying drawings. Although the illustrated embodiment is merely exemplary of systems for carrying out the presently disclosed subject matter, both the organization and method of operation of the presently disclosed subject matter, in general, together with further objectives and advantages thereof, may be more easily understood by reference to the drawings and the following description. The drawings are not intended to limit the scope of this presently disclosed subject matter, which is set forth with particularity in the claims as appended or as subsequently amended, but merely to clarify and exemplify the presently disclosed subject matter.

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For a more complete understanding of the presently disclosed subject matter, reference is now made to the following drawings in which:

FIG. 1 is an isometric view of an example embodiment of a mold of a human mandible, including mucosa, suitable for use in the simulator systems, devices, and methods disclosed herein.

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FIGS. 2 and 3 are isometric views of respective portions of an example embodiment of a mold used for forming an analogue of a human tongue.

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FIG. 4 is an isometric view of an example embodiment an analogue of a human tongue formed using the example mold portions shown in FIG. 2 and 3, with a growth (e.g., simulated tumor) on the base of the tongue for use in the simulator systems, devices, and methods disclosed herein.

FIG. 5 is an isometric view of an example embodiment of a mold of a human tonsil with a growth (e.g., simulated tumor) on the base thereof for use in the simulator systems, devices, and methods disclosed herein.

FIG. 6A is an isometric view of an example embodiment of a portion of a transoral robotic surgery (TORS) simulator system.

FIG. 6B is a perspective view of the portion of the TORS simulator system shown in FIG. 6A.

FIG. 7 is a perspective view of an example embodiment of a TORS simulator system, including the portion thereof shown in the example embodiment shown in FIGS. 6A and 6B.

FIG. 8 is a table of results for users of the example embodiment of the TORS simulator system shown in FIG. 7.

FIG. 9 is a graphical plot of results for users of the example embodiment of the TORS simulator system shown in FIG. 7, based on prior experience of the user with surgical robots.

FIG. 10 is a graph and table showing results based on user experience of various metrics for users of the TORS simulator system shown in FIG. 7.

FIG. 11 is a graphical plot of post-simulation survey responses from users of the TORS system shown in FIG. 7.

DETAILED DESCRIPTION

The subject matter disclosed herein is directed generally towards simulator systems for use in training users in transoral robotic surgery (TORS). Example embodiments of systems, devices, and/or methods of such TORS simulator systems, devices, and/or methods are thus provided herein, for purposes of illustration and not limitation.

Robotic surgery is the most recent and powerful evolution in minimally invasive surgery. Addressing many of the limitations of laparoscopic surgery, current robotic systems have restored hand-eye coordination, augmented surgeon dexterity, improved visualization, and provided ergonomic positioning. Within the last decade the use of robotic platforms has grown, with the da Vinci robot, developed by Intuitive Surgical (Sunnyvale, CA), being the most widely used system to date. Within the field of otolaryngology, the

most common application of the da Vinci robot system has been in transoral oral robotic surgery (TORS) for oropharyngeal malignancy.

FIGS. 6A and 6B show a portion of an example embodiment of such a TORS simulator system. Thus, the illustration of FIGS. 6A and 6B can be referred to herein as a simulator device, generally designated **10**. The simulator device **10** comprises a head-and-neck structure **20**, which can be formed using any suitable fabrication technique, including, for example, molding, additive manufacturing (e.g., 3D printing), and the like. The head-and-neck structure **20** is generally in the shape of a portion of a human head and is configured to have replaceable artificial tissue components attached thereto. As shown in FIGS. 6A and 6B, the replaceable artificial tissue components are an analogue of a human mandible **30**. As noted elsewhere herein, the mandible **30** is removable from the head-and-neck structure **20**, such that the simulator device **10** is in the form of a generally modular TORS simulator device. The simulator device **10** is configured for use with a robotic surgical system, including the Da Vinci robotic surgical system.

FIG. 1 shows aspects of an example embodiment of a human mandible, generally designated **30**, which can be used in the TORS simulator system **1** shown in FIG. 7. The mandible **30** is designed to allow for placement of artificial tissue components for a simulated tonsillectomy and base of tongue (BOT) resection. The mandible **30** comprises a mucosal surface **32** that substantially conforms to a typical inner surface of a human mouth, or portion thereof. Teeth **40** are attached to the mucosal surface **32**. The mucosal surface **32** defines an oral cavity, generally designated **25**, into a portion of which the base portion **52** of the tongue **50** extends, as shown in FIGS. 6A and 6B. The mucosal surface **32** can be, in some embodiments, painted or otherwise colored to be substantially similar to that of oral mucosa, so as to provide enhanced fidelity of the resultant simulator system. The mandible **30** is configured to be connected to the head-and-neck structure **20** at the upper ends **34** of the mandible **30**.

FIGS. 2 and 3 show aspects of portions of a mold that, when assembled together, form an interior region in the shape of a human tongue. Thus, in FIG. 2, a front mold, generally designated **50A**, is shown, which has an inner

surface **51A** that is generally in the shape of a portion (e.g., a front portion) of a human tongue. Similarly, in FIG. 3, a rear mold, generally designated **50B**, is shown, which has an inner surface **51B** that is generally in the shape of a portion (e.g., a rear portion) of a human tongue. When the front mold **50A** is assembled with the rear mold **50B**, an internal volume is formed that has a volumetric shape that generally corresponds to (e.g., is substantially similar to) that of a human tongue. The inner surfaces **50A**, **50B** of the front and rear molds **50A**, **50B**, respectively, are advantageously designed to allow for placement of artificial tissue components for a simulated tonsillectomy and base of tongue (BOT) resection.

An example embodiment of an analogue of a human tongue, generally designated **50**, that is formed by assembly of the front and rear molds **50A**, **50B** together and providing silicone therein, is shown in FIG. 4. In the example embodiment shown and described herein, the tongue **50** is formed from silicone, but the material used in forming the tongue **50** is not necessarily so limited and may include any suitable material. In order to increase anatomical fidelity of the tongue **50**, dye(s) can be mixed with the silicone material before the silicone was introduced (e.g., poured) into the mold formed by the front and rear molds **50A**, **50B** and the silicone material sets, or cures, a prescribed duration of time to form the tongue **50**. The front and rear molds **50A**, **50B** are advantageously reusable to produce a suitable quantity of tongues **50** for use in the TORS simulator system **100** disclosed herein. The tongue **50** comprises a tip section **51** and a base section **52**.

Since, in the example embodiment shown herein, the TORS simulator system **100** is used in training users in surgical base of tongue resection techniques, a tumor **5** is formed on the base section **52** of the tongue **50**. The tongue **50** has a perfusion channel **56** formed internal to the tongue **50**, from one side of the tongue **50** to another (e.g., opposite) side of the tongue **50** for passage to a perfusion tube through the perfusion channel **56**. The perfusion channel **56** terminates at, or within, the area of the tongue **50** where the tumor **5** is attached to the tongue **50**. The tumor **5** is formed using artificial tissue material (e.g., such as is available from LifeLike Biotissues) having suitable electrical conductance and customizability. In forming the tumor **5**, the artificial

tissue material is cut into a prescribed shape (e.g., corresponding to that of a typical base of tongue tumor). The tumor **5** is then advantageously injected at the base thereof with a dye of a contrasting color (e.g., methylene blue). As used herein, the term “base” generally means the portion of the tumor **5** immediately adjacent to the surface of the tongue **50** onto which the tumor **5** is, or will be, attached. The injection of a dye of a contrasting color into the base of the tumor **5** is advantageous to provide a margin for reference during resection of the tumor **5** from the tongue **50**. The tumor **5** advantageously comprises one or more copper wires (e.g., “copper wiring”) embedded within the artificial tissue material. The tongue **50** has a grounding wire channel **58** formed internal to the tongue **50**, from one side of the tongue **50** to another (e.g., opposite) side of the tongue **50** for passage to an electrically conductive wire, or wires, (e.g., electrically conductive wires connected to an electrical ground) through the grounding wire channel **58**. The grounding wire channel **58** terminates at, or within, the area of the tongue **50** where the tumor **5** is attached to the tongue **50** and the copper wires within the tumor **5** are connected, via the electrically conductive wire(s) within the grounding wire channel **58**, to a Bovie pad for electrical grounding, thereby allowing for monopolar electrocautery capabilities. In some embodiments the tumor **5** can represent any lesion or abnormality for which surgical training is desired.

An example embodiment of an analogue of a human tonsil, generally designated **70**, is shown in FIG. 5. In the example embodiment shown and described herein, the tonsil **70** is formed from silicone, but the material used in forming the tonsil **70** is not necessarily so limited and may include any suitable material. In order to increase anatomical fidelity of the tonsil **70**, dye(s) can be mixed with the silicone material used to form the tonsil **70**. The silicone material sets, or cures, a prescribed duration of time to form the tonsil **70**.

Since, in the example embodiment shown herein, the TORS simulator system **100** is used in training users in surgical tonsillectomy techniques, a tumor **5** is formed on the tonsil **70**. The tonsil **70** has a perfusion channel **76** formed internal to the tonsil **70**, from one side of the tonsil **70** to another (e.g., opposite) side of the tonsil **70** for passage to a perfusion tube through the perfusion channel **76**. The perfusion channel **76** terminates at, or within, the

area of the tonsil **70** where the tumor **5** is attached to the tonsil **70**. The tumor **5** is formed using artificial tissue material (e.g., such as is available from LifeLike Biotissues) having suitable electrical conductance and customizability. In forming the tumor **5**, the artificial tissue material is cut into
5 a prescribed shape (e.g., corresponding to that of a typical palatine tumor). The tumor **5** is then advantageously injected at the base thereof with a dye of a contrasting color (e.g., methylene blue). As used herein, the term “base” generally means the portion of the tumor **5** immediately adjacent to the surface of the tonsil **70** onto which the tumor **5** is, or will be, attached. The injection of
10 a dye of a contrasting color into the base of the tumor **5** is advantageous to provide a margin for reference during resection of the tumor **5** from the tonsil **70**. The tumor **5** advantageously comprises one or more copper wires (e.g., “copper wiring”) embedded within the artificial tissue material. The tonsil **70** has a grounding wire channel **78** formed internal to the tonsil **70**, from one
15 side of the tonsil **70** to another (e.g., opposite) side of the tonsil **70** for passage to an electrically conductive wire, or wires, (e.g., electrically conductive wires connected to an electrical ground) through the grounding wire channel **78**. The grounding wire channel **78** terminates at, or within, the area of the tonsil **70** where the tumor **5** is attached to the tonsil **70** and the copper wires within the
20 tumor **5** are connected, via the electrically conductive wire(s) within the grounding wire channel **78**, to a Bovie pad for electrical grounding, thereby allowing for monopolar electrocautery capabilities.

In the TORS simulator system **100** shown in FIG. 7, the tongue **50** has a simulated tumor **5** attached (e.g., adhesively, such as by glue) to the base
25 section **52** of the tongue **50**. As shown in FIGS. 2 and 3, the front and rear molds **50A**, **50B** have formed therein generally longitudinally-extending sections that form passages (e.g., perfusion channel **56** and grounding wire channel **58**) from the underside of the tongue **50** to the location on the base section **52** where the tumor **5** is, or will be, attached. In one of these passages
30 of the tongue **50**, a ground wire is positioned therein and connected between the tumor (e.g., the one or more copper wires formed on and/or in the tumor **5**) and a Bovie pad to create an electrical conductance of the tumor and also allowing resection of the tumor **5** via monopolar cautery. In the example

embodiment described herein, a grounding wire is fed through the grounding wire channel **58**, from the underside of the tongue **50**, and is connected between the ground wire embedded within the tumor **5** and a Bovie (grounding) electrical pad to provide electrical conductance of the tumor **5** and also allowing for resection of the tumor **5** from the tongue **50** with monopolar cautery.

In some embodiments, the TORS simulation system **100** has a simulated tumor **5** attached (e.g., adhesively, such as by glue) to an outer surface of a simulated tonsil **70**. In forming the tonsil **70**, one or more passages (e.g., perfusion channel **76** and grounding wire channel **78**) are formed through all or a portion of the tonsil **70**. In one of these passages of the tonsil **70**, a ground wire is positioned therein and connected between the tumor (e.g., the one or more copper wires formed on and/or in the tumor **5**) and a Bovie pad to create an electrical conductance of the tumor and also allowing resection of the tumor **5** via monopolar cautery. In the example embodiment described herein, a grounding wire is fed through the grounding wire channel **78**, from the underside of the tonsil **70**, and is connected between the ground wire embedded within the tumor **5** and a Bovie (grounding) electrical pad to provide electrical conductance of the tumor **5** and also allowing for resection of the tumor **5** from the tonsil **70** with monopolar cautery.

In some embodiments, the TORS simulation system **100** can be used with a tongue **50** having a tumor **5** attached thereto, as described elsewhere herein, and also a tonsil **70** having a tumor **5** attached thereto, as described elsewhere herein.

In embodiments of the TORS simulator system **100** in which the tongue **50** and/or the tonsil **70** have a simulated tumor **5** attached thereto, it is advantageous to provide increased fidelity for users to provide a live perfusion simulation upon resection of the tumor **5**, whether from the tongue **50** and/or the tonsil **70**. Thus, it is advantageous to simulate bleeding from a blood vessel (e.g., artery) during the surgical resection, so that users of the TORS simulator system **100** will be trained in treating such bleeding during surgery on a living patient. In FIG. 7, an example of live perfusion from a tumor **5** on the tongue **50** is illustrated, but persons having ordinary skill will understand that the tube

providing the fluid flow to the tumor **5** of the tongue **50** can, either instead of or in addition to the tube shown, be connected through a passage formed in the tonsil **70** to the tumor **5** attached to the tonsil **70**. Thus, in order to provide live perfusion functionality, the TORS simulator system **100** comprises a fluid source **92** containing a fluid **96**, a pump **90** connected to the fluid source **90** to withdrawn the fluid **96** from the fluid source **92**, and an injection tubing **94** that is inserted through on the of the passages (e.g., perfusion channel **56** or **76**) through the tongue **50** or the tonsil **70**, as the case may be, to provide a flow of fluid to the underside of the base of the tumor **5**.

10 The fluid **96** is a simulated blood and the pump **90** is configured to pump the fluid **96** from the fluid source **92**, through the injection tubing **94**. In some embodiments, the injection tubing **94** is inserted through the passage entirely, such that the distal end of the injection tubing **94** is internal to (e.g., at least partially inside of) the tumor **5**. The flow rate of the fluid **96** may be controlled
15 via the pump **90** to provide a realistic flow rate that is generally the same as would occur during an incision being made in a tumor in a human. Various fluids can be used in the live perfusion model in order to simulate sanguination during resection of the tumor **5** from either the tongue **50** or the tonsil **70**. An example of a fluid that can be used as artificial blood for live perfusion in the
20 example system **100** disclosed herein is a mixture of water and corn starch that is dyed with red food coloring to resemble human blood in both consistency and color. In the example embodiment disclosed herein, the pump **90** provides a pulsatile fluid flow at variable flow rates. An example of a pump **90** that can provide such pulsatile fluid flow at variable flow rates is a
25 INTLLAB 12V DC DIY Peristaltic Liquid Pump Dosing Pump (2mm ID x 4mm OD).

 In order to validate the example embodiment of the TORS simulator system **100**, 16 users (surgeons) having differing robotic skill levels (e.g., novice, experienced, and robotically trained) participated in a surgical
30 simulation of a partial tonsillectomy and tongue base tumor resection, using the tongue mold **50** shown in FIG. 4 and the tonsil mold **70** shown in FIG. 5. Using Maryland forceps and the unipolar Bovie arm, the users interacted with the TORS simulator system **1** shown in FIG. 7, which had a 3D-printed oral

cavity and oropharyngeal architecture and artificial tissue lesions. Video recordings of each participant were graded by a blinded robotically trained surgeon using a GEARS criterion adapted for the TORS simulator system.

5 Prior to simulator testing participants completed informed consent documentation. Participants were offered brief instruction on robotic controls and EndoWrist® instruments including the use of Maryland forceps, monopolar Bovie, and the camera arm. Brief oral instruction was provided on the simulated tonsillectomy and BOT resection including the interpretation of blue dye at the base of each lesion as the “normal tissue” margin for depth. 10 Participants were instructed to attempt a complete resection along this margin. Prior to beginning the recorded simulation, participants were given 5-10 minutes to practice movements with the robotic arms.

Participants began by resecting the simulated palatine tonsil followed by resection of the simulated BOT lesion. Participants were able to ask clarify 15 questions during the simulation. After completion of the simulation, participants were given a post-simulation survey to complete, detailing their experience with the robot and simulator.

Simulations were timed and recorded using the da Vinci recording software and other camera recording devices when appropriate. Simulation 20 recordings were de-identified and assigned a coded file number before uploading to an encrypted, shared folder to which only the research team had access. Following each participant simulation, pictures were taken of tumor margins to assess resection completion; pictures were uploaded to each participant’s profile which included the recorded simulations.

25 To test the prototype’s ability to differentiate operator skill level, the Global Evaluative Assessment of Robotic Skill (GEARS), a validated 5-point Likert tool for the evaluation of robotic surgical skill, was selected to evaluate operator use of the prototype. To increase the simulators capability in differentiating between expert and novice operators performing transoral 30 robotic surgery, the GEARS criteria was modified to appropriately grade the simulated BOT resection and tonsillectomy. Autonomy and third arm use categories were removed from the GEARS criteria because the simulation did not include third arm manipulation or assistance from a second operator. The

simulator was designed to assess skill at performing oncologically-sound transoral robotic surgery, therefore, a resection completion category was added to the assessment criteria.

One of the challenges of transoral robotic surgery is space. The upper
5 aerodigestive tract is a relatively narrow conduit within which to fit rigid robotic arms and camera. The confined rigid operative dimension created by the bony framework of the mandible, spine and maxilla creates a challenging environment within which to move instruments. To gain access to the operative field in TORS, the robot arms are placed through a shared narrow
10 window of the oral cavity, this increases the risk for instrument collision, and incidental injury to structures in the oral cavity, such as the lip, teeth, mandible, mucosa and palate. The operator at the console has a limited view of the operative field which requires a comprehensive knowledge and appreciation for the spatial relationships of the arms to each other and the surrounding oral
15 tissues. In comparison, port access in abdominal robotic surgery is done under insufflation, providing a safe space for the movement of proximal portions of the robotic arms not visible under the console, an advantage not afforded to robotic arm manipulation during transoral robotic surgery

Therefore, instrument collision was also added to the GEARS
20 assessment tool. Each simulation recording was evaluated by a blinded, robotically trained surgeon using a modified GEARS assessment tool, shown in FIG. 12. With the modified GEARS assessment tool, participants were graded on the following skill sets: depth perception, bimanual dexterity, efficiency, force sensitivity, robotic control, collisions, resection completion,
25 and time. The tonsillectomy and BOT resection were performed in sequence; however, each task was individually timed and graded.

Average GEARS scores were differed in a 3-way head-to-head comparison of participant groups: Experience v. No Experience ($P = .004$), Experience v. Robotically trained ($P < .050$), and Robotically Trained v. No
30 Experience ($P = .018$). Robotic surgical skills were tested and found to be statistically significant between novice and experienced operators in depth perception, bimanual dexterity, efficiency, force sensitivity, robotic control, and resection completion.

Using a modified GEARS criterion, the prototype simulator successfully differentiated operators of differing skill levels. These preliminary findings support the construct validity in our TORS simulator.

5 The 16 users each used the TORS simulator system **100** twice, yielding a total of 32 simulations during validation. The users self-identified as either novice users of surgical robots (8), users with some experience using surgical robots (5), or users trained in using surgical robots (3) prior to using the TORS simulation system **100**.

10 As shown in FIG. 8, the total GEARS scores correlated with reported experience level, with novice operators awarded 54% of total points awarded at 19, operators with some experience awarded 82.3% of total points at 28.8, and robotically trained operators awarded 97.1% of total points at 34. In a head-to-head comparison of all three operator groups, a statistically significant difference was detected in the mean total GEARS scores between each individual operator group ($p < 0.0001$); no robotic experience versus some robotic experience ($p < 0.001$), no robotic experience versus robotically trained ($p = 0.004$), and some robotic experience versus robotically trained ($p = 0.024$) (Figure 3).

20 When evaluating the individual metrics of the GEARS assessment, the simulation successfully differentiated novice from experienced and robotically trained operators in the following robot skills: depth perception ($p = 0.0001$), bimanual dexterity ($p = 0.0006$), efficiency ($p = 0.003$), force sensitivity ($p = 0.0155$), robotic control ($p = 0.022$), and resection completion ($p = 0.0097$) (Table 1) (Figure 5). However, regarding frequency of instrument collisions during the simulated tonsillectomy and BOT resection, there was no difference when comparing novice operators to robotically trained and operators reporting some robot experience, ($p = 0.2012$) (Table 1).

25 After the completion of the simulation, each participant completed a survey to evaluate the simulator and their overall experience on a 5-point Likert scale. In assessing the ability of the investigative team to coordinate the simulation, participants rated Level of Instruction (4.4) and Smoothness of Transition (4.7). With regards to the simulated experience participants rated Ease of Use (4.4), Realistic Application of Operative Skill (4.1), and Realism

of Overall Experience (4.0). Lastly, participants rated their Comfort with the Robot After the Simulation at 4.5 (Figure 6).

FIG. 9 shows a 3-way head-to-head comparison of mean total GEARS scores between operator groups: A statistically significant difference in average total GEARS scores was observed between operator groups of different experience levels ($p < 0.0001$). Self-reported experience with the robot was associated with higher mean total GEARS scores compared to no reported robotic experience (7.5pt difference; $p < 0.001$). Formal robotic training was associated with higher mean total GEARS compared to self-report experience with the robot (6.5pt difference; $p = .0238$) and no reported experience with the robot (14pt difference; $p = 0.004$).

FIG. 10 shows a comparison of mean GEARS scores in all metrics (5-point Likert Scale) between operator groups. Higher scores in depth perception were associated with both robotic training ($p=0.015$) and experience with the robot ($p<0.008$) compared to no robot experience. Higher scores in bimanual dexterity were associated with both robotic training ($p=0.014$) and experience with the robot ($p=0.005$) compared to no robot experience. Robotic training was associated with higher efficiency scores compared to both some robot experience ($p<0.028$) and no robotic experience ($p=0.014$). Robotic training was associated with higher scores in force sensitivity compared to no robot experience ($p=0.032$). Robotic training was associated with higher scores in robotic control compared to no robot experience ($p=0.043$).

FIG. 11 shows a mean post-simulation survey grading by question as completed by the users themselves.

Thus, in some embodiments, disclosed herein are surgical training simulators for transoral robotic surgery (TORS). Such simulators can in some embodiments comprise a three-dimensional human head model comprising oral cavity and oropharyngeal synthetic material, wherein the oral cavity and oropharyngeal structure comprises one or more of a mandible structure, a tongue structure and/or a tonsil structure, wherein the tongue structure comprises a silicone material; an artificial tissue implanted in a region of the oropharynx, wherein the artificial tissue is configured to mimic a tumor tissue;

and a marker material configured to differentiate between the artificial tissue and synthetic material of the oropharynx. In some aspects, the training simulator is configured to be used with a robotic surgical system. In some aspects, the oral cavity and oropharynx comprising synthetic material is made
5 via an additive manufacturing process. In some aspects, the tongue structure comprising a silicone material is molded from an additive manufactured mold. In some embodiments, the mandible structure, the tongue structure and/or the tonsil structure is modular whereby the structure can be removed and/or replaced within the head model. In some aspects, the marker material is
10 injected at a base of the artificial tissue mimicking a tumor tissue. In some aspects, the marker material comprises a colored dye, wherein the colored dye is visible to a practitioner performing a simulated surgery and provides a margin for reference during resection.

In some embodiments, the simulators can further comprise a series of
15 electrically conductive wires or materials integrated into the oral cavity and oropharynx, optionally wherein the simulator is configured for monopolar capability. In some aspects, the artificial tissue is configured to comprise monopolar capability. In some aspects, the artificial tissue embedded with copper wiring which was subsequently connected to a Bovie pad for electrical
20 grounding, allowing for monopolar electrocautery capabilities. In some aspects, the marker material comprises a colored dye, wherein the colored dye is visible to a practitioner performing a simulated surgery and provides a margin for reference during resection. In some aspects, the mandible component and/or tongue component is configured to allow for placement of
25 the artificial tissue mimicking a tumor for a simulated tonsillectomy and base of tongue (BOT) resection. In some embodiments, such simulators can further comprise conduit integrated into the synthetic material and/or artificial tissue, wherein the conduit is configured to contain, transport and/or be perfused with a liquid configured to mimic blood, optionally wherein the conduit is integrated
30 into the artificial tissue mimicking the tumor. In some embodiments, such simulators can further comprise a copper tape, mesh or wire, or other conductive material, integrated into the oral cavity and oropharynx and configured to detect instrument collision during use of the simulator.

Also disclosed herein are robotic surgical training systems, the systems comprising a surgical training simulator of any of the above claims; and an evaluative assessment criteria configured to assess an operator's use of the simulator. In some embodiments, the evaluative assessment criteria
5 comprises a Global Evaluative Assessment of Robotic Skill (GEARS), or modified version thereof. In some embodiments, such training systems can further comprise one or more cameras and/or video recording devices configured to capture images and/or video during use of the simulator. In some embodiments, such training systems can further comprise a computing
10 device communicatively connected to the simulator, robotic surgical system and/or the one or more cameras and/or video recording devices. In some embodiments, such training systems can further comprise a graphical user interface to display images, videos and/or data from the surgical training system.

15 Provided herein are also methods for training a practitioner for transoral robotic surgery (TORS), that methods comprising providing a surgical training simulator and/or a robotic surgical training system of any of the above claims; having the practitioner perform a simulated TORS procedure on the simulator or training system; and providing an evaluative assessment to the practitioner.
20 The evaluative assessment provided to the practitioner can be based on data from a Global Evaluative Assessment of Robotic Skill (GEARS), or modified version thereof.

The terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting of the presently disclosed
25 subject matter.

While the following terms are believed to be well understood by one of ordinary skill in the art, the following definitions are set forth to facilitate explanation of the presently disclosed subject matter.

All technical and scientific terms used herein, unless otherwise defined
30 below, are intended to have the same meaning as commonly understood by one of ordinary skill in the art. References to techniques employed herein are intended to refer to the techniques as commonly understood in the art, including variations on those techniques or substitutions of equivalent

techniques that would be apparent to one skilled in the art. While the following terms are believed to be well understood by one of ordinary skill in the art, the following definitions are set forth to facilitate explanation of the presently disclosed subject matter.

5 In describing the presently disclosed subject matter, it will be understood that a number of techniques and steps are disclosed. Each of these has individual benefit and each can also be used in conjunction with one or more, or in some cases all, of the other disclosed techniques.

 Accordingly, for the sake of clarity, this description will refrain from
10 repeating every possible combination of the individual steps in an unnecessary fashion. Nevertheless, the specification and claims should be read with the understanding that such combinations are entirely within the scope of the invention and the claims.

 Following long-standing patent law convention, the terms “a”, “an”, and
15 “the” refer to “one or more” when used in this application, including the claims. Thus, for example, reference to “a cell” includes a plurality of such cells, and so forth.

 Unless otherwise indicated, all numbers expressing quantities of ingredients, reaction conditions, and so forth used in the specification and
20 claims are to be understood as being modified in all instances by the term “about”. Accordingly, unless indicated to the contrary, the numerical parameters set forth in this specification and attached claims are approximations that can vary depending upon the desired properties sought to be obtained by the presently disclosed subject matter.

25 As used herein, the term “about,” when referring to a value or to an amount of a composition, mass, weight, temperature, time, volume, concentration, percentage, *etc.*, is meant to encompass variations of in some embodiments $\pm 20\%$, in some embodiments $\pm 10\%$, in some embodiments $\pm 5\%$, in some embodiments $\pm 1\%$, in some embodiments $\pm 0.5\%$, and in some
30 embodiments $\pm 0.1\%$ from the specified amount, as such variations are appropriate to perform the disclosed methods or employ the disclosed compositions.

 The term “comprising”, which is synonymous with “including”

“containing” or “characterized by” is inclusive or open-ended and does not exclude additional, unrecited elements or method steps. “Comprising” is a term of art used in claim language which means that the named elements are essential, but other elements can be added and still form a construct within the scope of the claim.

As used herein, the phrase “consisting of” excludes any element, step, or ingredient not specified in the claim. When the phrase “consists of” appears in a clause of the body of a claim, rather than immediately following the preamble, it limits only the element set forth in that clause; other elements are not excluded from the claim as a whole.

As used herein, the phrase “consisting essentially of” limits the scope of a claim to the specified materials or steps, plus those that do not materially affect the basic and novel characteristic(s) of the claimed subject matter.

With respect to the terms “comprising”, “consisting of”, and “consisting essentially of”, where one of these three terms is used herein, the presently disclosed and claimed subject matter can include the use of either of the other two terms.

As used herein, the term “and/or” when used in the context of a listing of entities, refers to the entities being present singly or in combination. Thus, for example, the phrase “A, B, C, and/or D” includes A, B, C, and D individually, but also includes any and all combinations and subcombinations of A, B, C, and D.

The subject matter disclosed herein can be implemented in software in combination with hardware and/or firmware. For example, the subject matter described herein can be implemented in software executed by a processor. In one exemplary implementation, the subject matter described herein can be implemented using a computer readable medium having stored thereon computer executable instructions that when executed by a processor of a computer control the computer to perform steps. Exemplary computer readable mediums suitable for implementing the subject matter described herein include non-transitory devices, such as disk memory devices, chip memory devices, programmable logic devices, and application specific integrated circuits. In addition, a computer readable medium that implements

the subject matter described herein can be located on a single device or computing platform or can be distributed across multiple devices or computing platforms.

- 5 It will be understood that various details of the presently disclosed subject matter may be changed without departing from the scope of the presently disclosed subject matter. Furthermore, the foregoing description is for the purpose of illustration only, and not for the purpose of limitation.

CLAIMS

1. A transoral robotic surgery (TORS) simulator system comprising:
a three-dimensional human head model comprising an oral cavity
5 comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material;
at least one artificial tissue attached within the oral cavity, wherein the
10 artificial tissue is configured to mimic a biological tissue; and
a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity.
15
2. The TORS simulator system of claim 1, wherein the TORS simulator system is configured for use with a robotic surgical system controlled by the user.
- 20 3. The TORS simulator system of any of claims 1 or 2, wherein the oral cavity comprising synthetic material is made via an additive manufacturing process.
4. The TORS simulator system of any of claims 1-3, wherein the
25 simulated human tongue is molded using a mold formed via additive manufacturing.
5. The TORS simulator system of any of claims 1-4, wherein the mandible structure, the simulated human tongue, and/or the simulated human tonsil
30 is/are modular, such that each of the mandible structure, the simulated human tongue, and/or the simulated human tonsil are individually removable and/or replaceable from the head model.

6. The TORS simulator system of any of claims 1-5, wherein the marker material is injected at a base of the at least one artificial tissue.

7. The TORS simulator system of any of claims 1-6, comprising one or more electrically conductive wires or materials integrated into the oral cavity, optionally, wherein the simulator system is configured for monopolar capability.

8. The TORS simulator system of any of claims 1-7, wherein the at least one artificial tissue is configured to comprise monopolar capability.

9. The TORS simulator system of claim 7, wherein the one or more electrically conductive wires are connected to an electrical ground for electrical grounding, allowing for monopolar electrocautery capabilities.

15

10. The TORS simulator system of any of claims 1-9, wherein the marker material comprises a colored dye, wherein the colored dye is visible to the user of the simulator system performing a simulated surgical procedure and is configured to provide a margin for reference during resection of the at least one artificial tissue.

20

11. The TORS simulator system of any of claims 1-10, wherein the simulated human tongue and/or the simulated human tonsil are configured to allow for attachment of the at least one artificial tissue thereto for the user to perform a tonsillectomy and/or base of tongue (BOT) resection.

25

12. The TORS simulator system of any of claims 1-11, comprising a perfusion channel integrated into the synthetic material and/or the at least one artificial tissue, wherein the perfusion channel is configured to contain, transport and/or be perfused with a liquid configured to mimic blood and, optionally, wherein the perfusion channel is integrated into the at least one artificial tissue.

30

13. The TORS simulator system of any of claims 1-12, comprising a copper tape, mesh, wire, or other electrically conductive material that is integrated into, or attached within, the oral cavity and is configured to detect contact with a surgical instrument during use of the simulator system.

5

14. The TORS simulator system of any of claims 1-13, wherein the biological tissue is a cancerous growth tissue.

15. A robotic surgical training system, the system comprising:
10 a TORS simulator system of any of claims 1-14; and
an evaluative assessment criteria configured to assess the user's use of the simulator system.

16. The robotic surgical training system of claim 15, wherein the evaluative
15 assessment criteria comprises a Global Evaluative Assessment of Robotic Skill (GEARS), or a modified version thereof.

17. The robotic surgical training system of any claim 15 or 16, comprising
20 one or more cameras configured to capture images during use of the TORS simulator system and/or video recording devices configured to capture video during use of the TORS simulator system.

18. The robotic surgical training system of claim 17, comprising a
25 computing device communicatively connected to the TORS simulator system, the robotic surgical system, and/or the one or more cameras and/or video recording devices.

19. The robotic surgical training system of any of claims 17 or 18,
30 comprising a graphical user interface to display images, videos, and/or data from the surgical training system.

20. A method for training a user in transoral robotic surgery (TORS) techniques, the method comprising:

providing a TORS simulator system comprising:

a three-dimensional human head model comprising an oral cavity comprising synthetic material, wherein the oral cavity comprises a mandible structure and a simulated human tongue and/or a simulated human tonsil, wherein the simulated human tongue and/or the simulated human tonsil comprise a silicone material;

at least one artificial tissue attached within the oral cavity, wherein the at least one artificial tissue is configured to mimic a biological tissue; and

a marker material present on and/or within the at least one artificial tissue, wherein the marker material is configured to visually differentiate for a user of the system between the at least one artificial tissue and the synthetic material of the oral cavity;

having the user perform a simulated TORS procedure on the TORS simulator system; and

providing an evaluative assessment to the user regarding the user's performance of the TORS procedure.

21. The method of claim 20, wherein the evaluative assessment is performed using an evaluative assessment criteria for assessing the user's use of the simulator system.

22. The method of claim 21, wherein the evaluative assessment provided to the practitioner is based on data from a Global Evaluative Assessment of Robotic Skill (GEARS), or modified version thereof.

23. The method of claim 20, wherein the user performs the simulated TORS procedure on the TORS simulator system using a robotic surgical system controlled by the user.

24. The method of any of claims 20-23, wherein providing the TORS simulator system comprises forming the oral cavity via additive manufacturing.

25. The method of any of claims 20-24, comprising forming a mold for the simulated human tongue via additive manufacturing, wherein providing the TORS simulator system comprises molding the simulated human tongue within the mold.

5

26. The method of any of claims 20-25, wherein the mandible structure, the simulated human tongue, and/or the simulated human tonsil is/are modular, such that each of the mandible structure, the simulated human tongue, and/or the simulated human tonsil are individually removable and/or replaceable from the head model.

10

27. The method of any of claims 20-26, wherein providing the TORS simulator system comprises injecting the marker material at a base of the at least one artificial tissue.

15

28. The method of any of claims 20-27, wherein providing the TORS simulator system comprises integrating one or more electrically conductive wires or materials into the oral cavity, optionally, wherein the simulator system is configured for monopolar capability.

20

29. The method of any of claims 20-28, wherein the at least one artificial tissue comprises monopolar capability.

30. The method of claim 29, wherein providing the TORS simulator system comprises connecting the one or more electrically conductive wires to an electrical ground for electrical grounding, allowing for monopolar electrocautery capabilities.

25

31. The method of any of claims 20-30, wherein the marker material comprises a colored dye, wherein the colored dye is visible to the user of the simulator system performing a simulated surgical procedure and is configured to provide a margin for reference during resection of the at least one artificial tissue.

30

32. The method of any of claims 20-31, wherein the simulated human tongue and/or the simulated human tonsil are configured to allow for attachment of the at least one artificial tissue thereto for the user to perform a
5 tonsillectomy and/or base of tongue (BOT) resection.

33. The method of any of claims 20-32, wherein providing the TORS simulator system comprises forming a perfusion channel in the synthetic material and/or the at least one artificial tissue, wherein the perfusion channel
10 is configured to contain, transport and/or be perfused with a liquid configured to mimic blood and, optionally, wherein the perfusion channel is integrated into the at least one artificial tissue.

34. The method of any of claims 20-33, wherein a copper tape, mesh, wire,
15 or other electrically conductive material is integrated into, or attached within, the oral cavity and is configured to detect contact with a surgical instrument during use of the simulator system.

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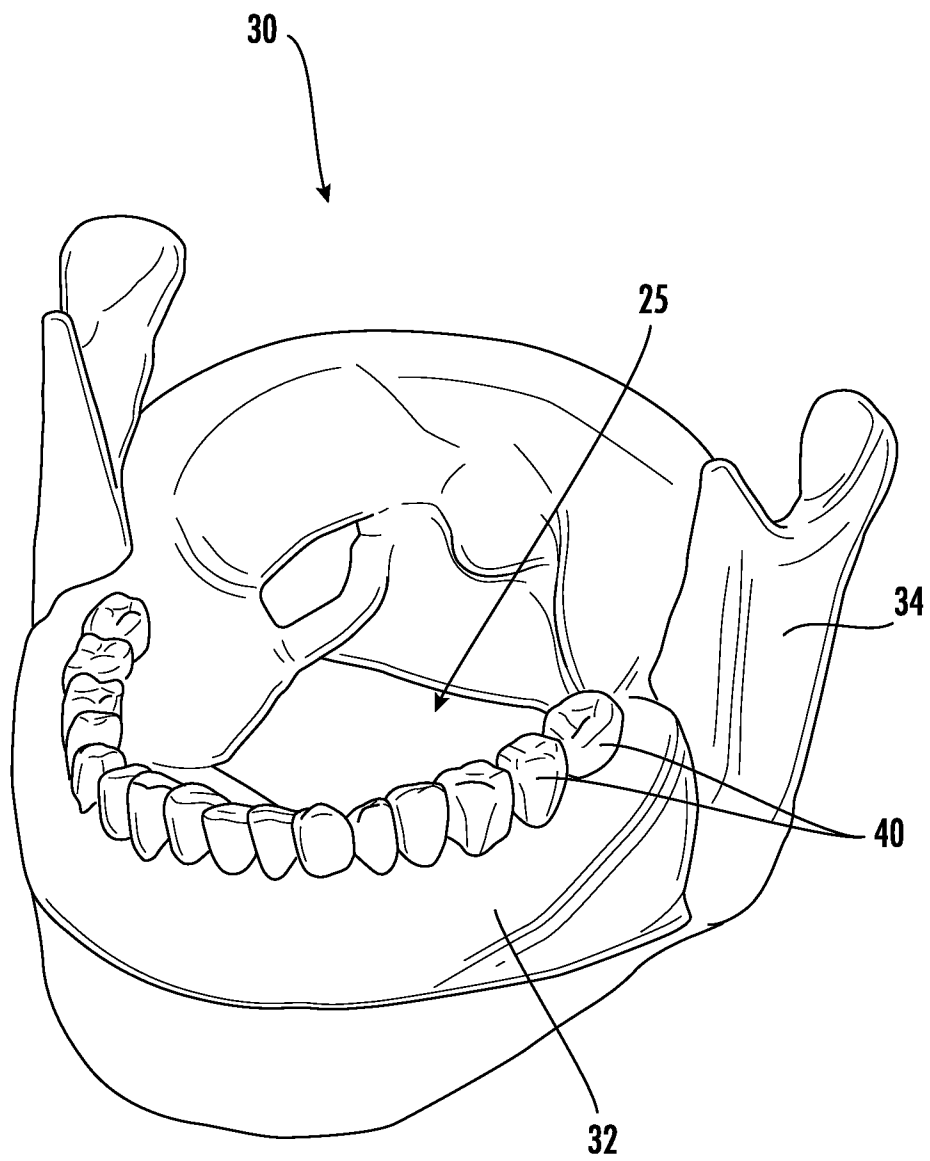


FIG. 1

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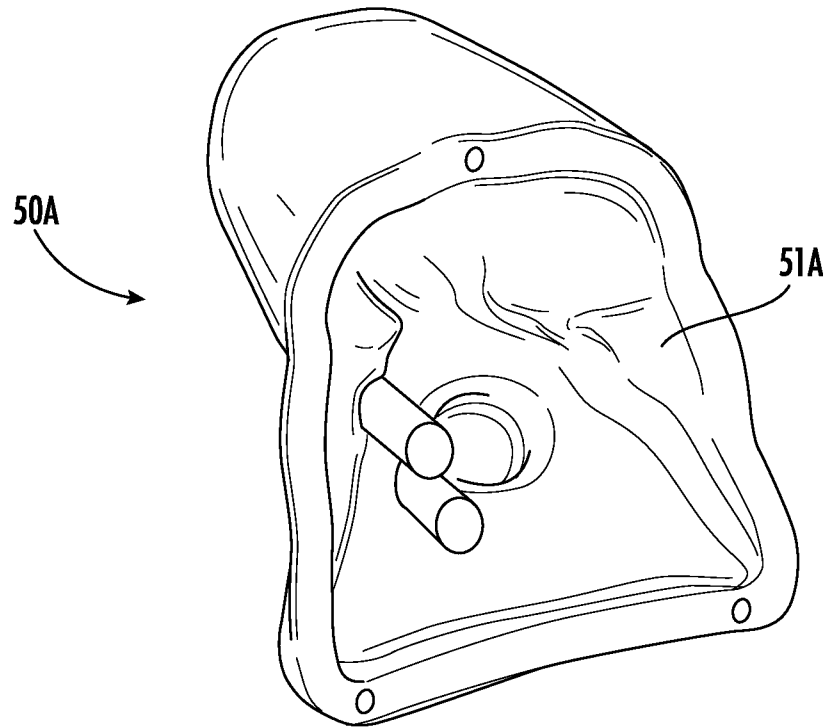


FIG. 2

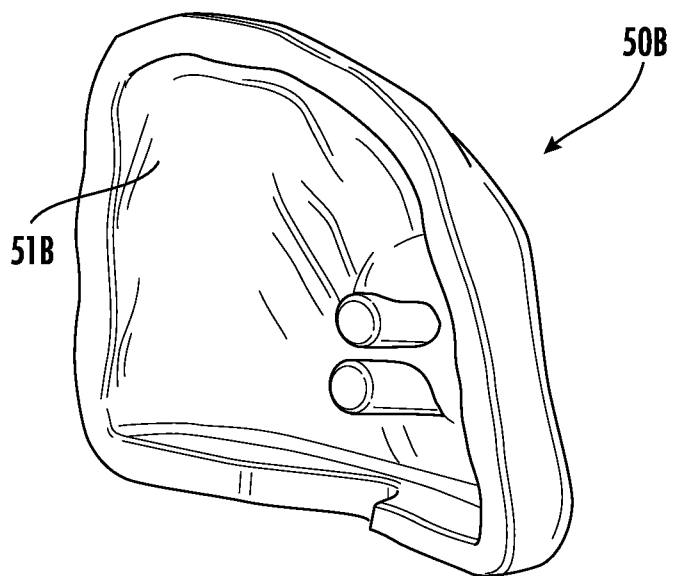


FIG. 3

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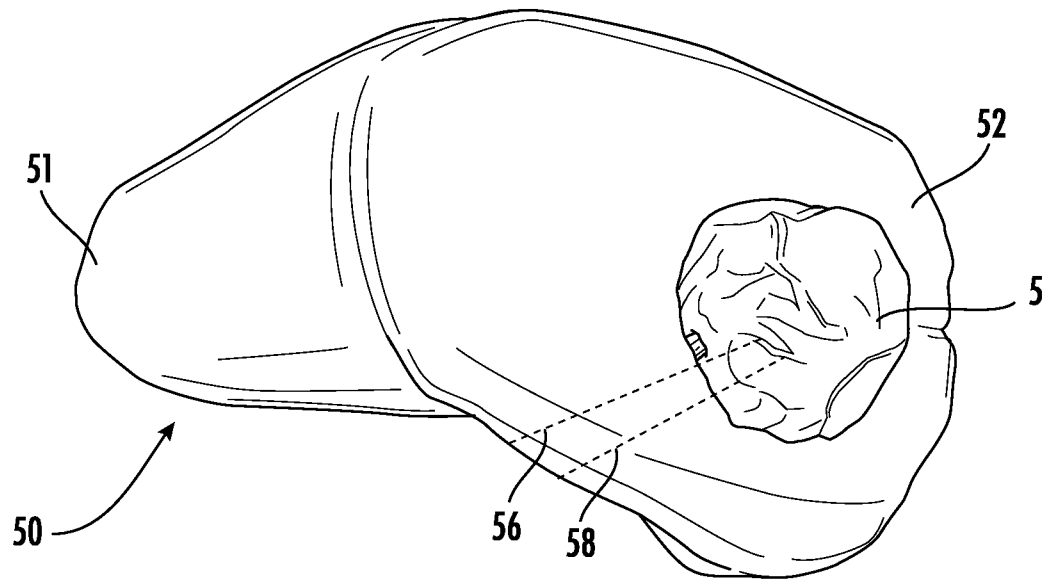


FIG. 4

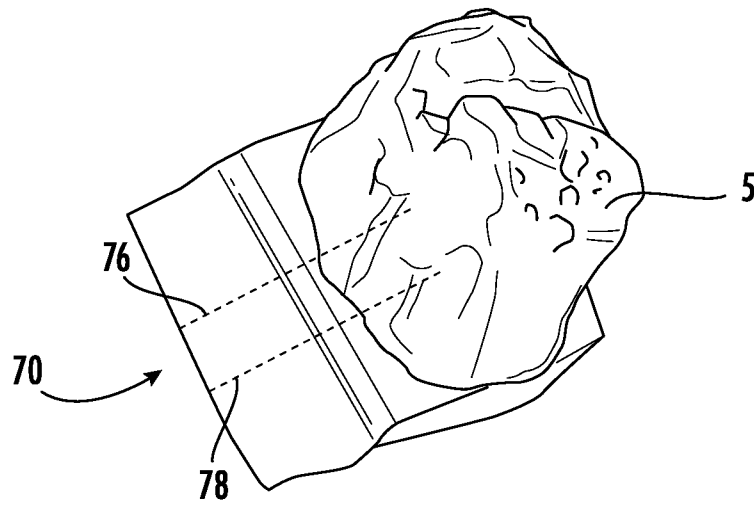


FIG. 5

4/11

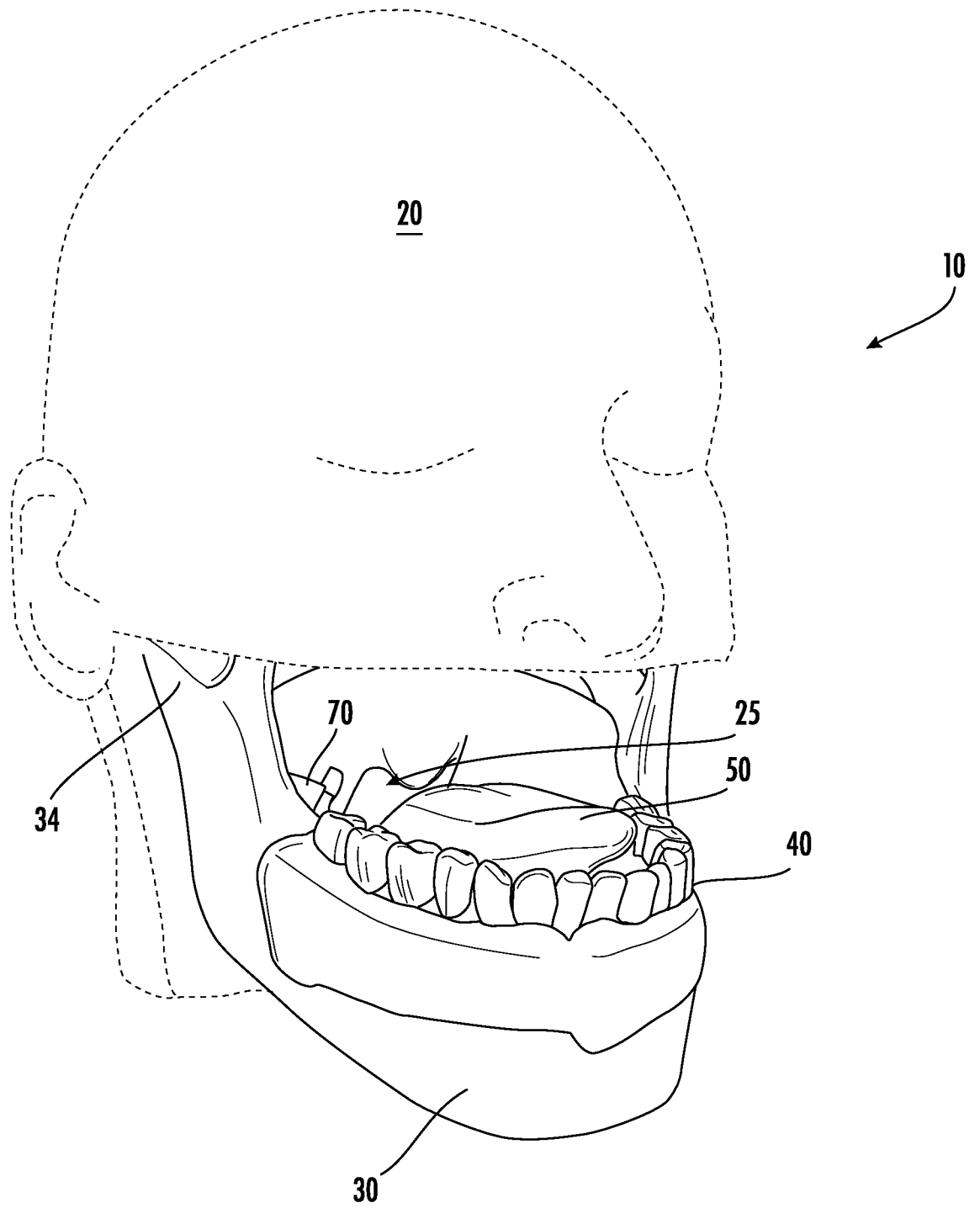


FIG. 6A

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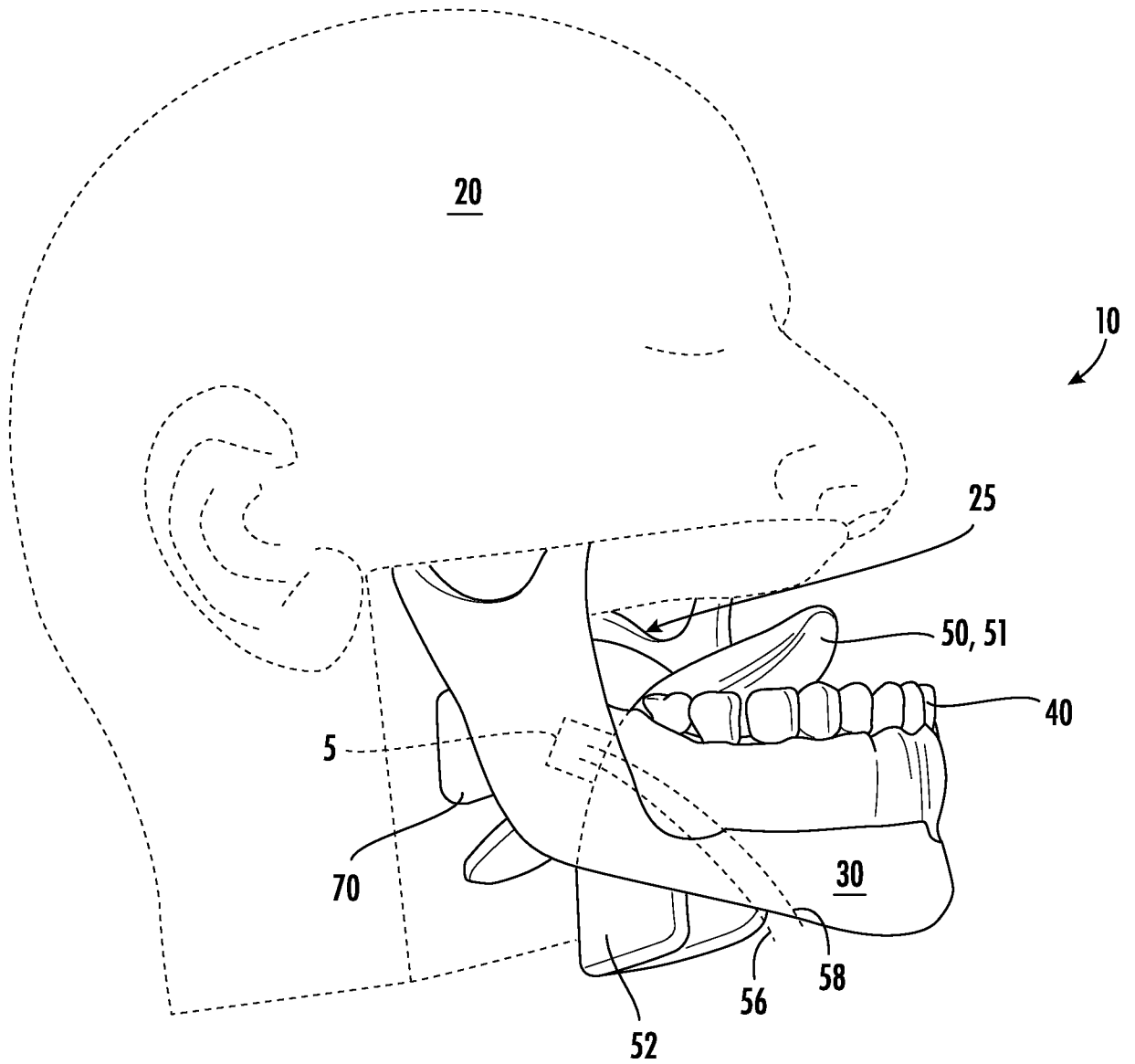


FIG. 6B

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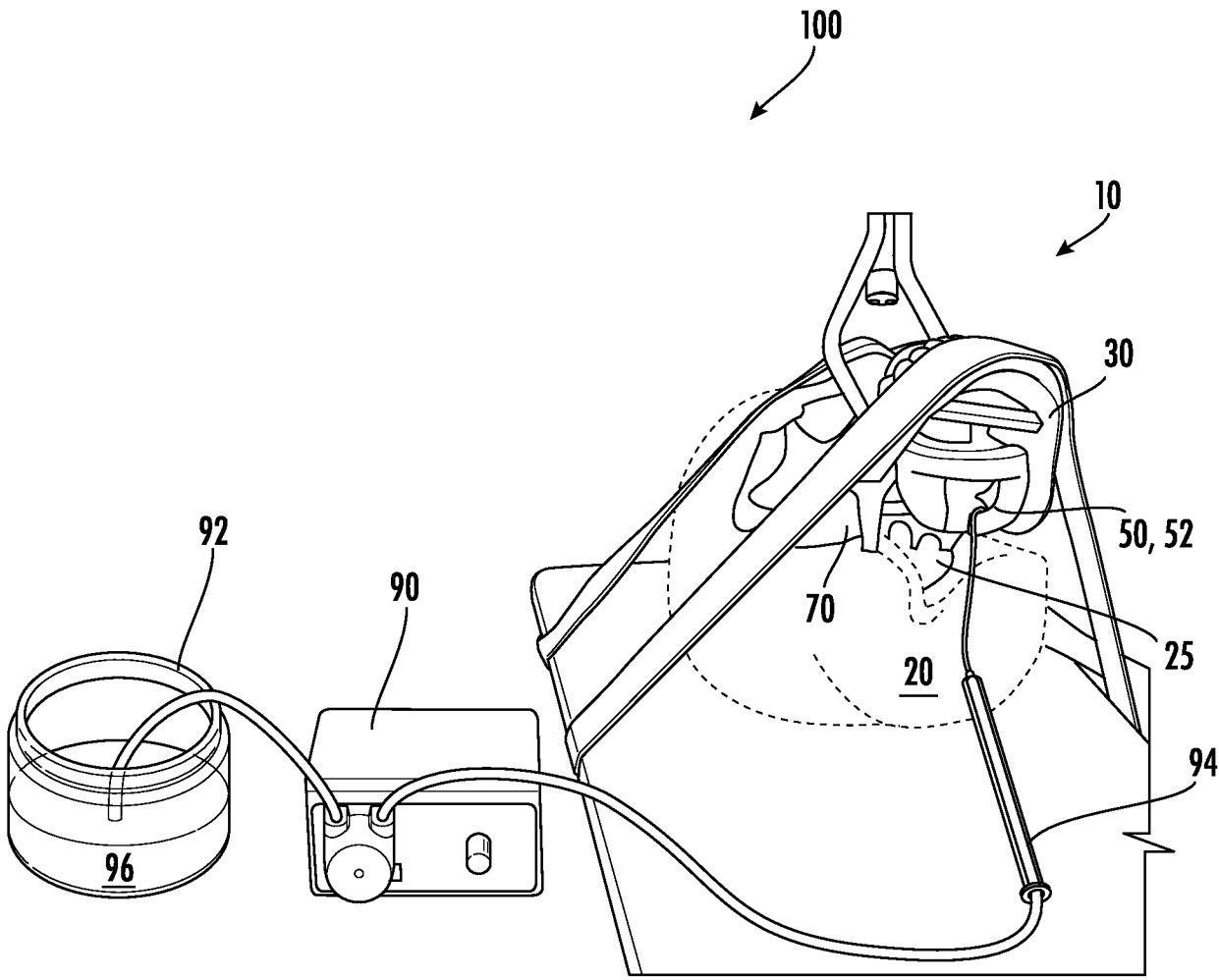


FIG. 7

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	NO EXPERIENCE (N=8)		SOME EXPERIENCE (N=5)		ROBOTICALLY TRAINED (N=3)		*P-VALUE
	MEAN	SD	MEAN	SD	MEAN	SD	
DEPTH PERCEPTION (1-5)	2.5	1.1	4.2	0.4	5.0		0.0001
BIMANUAL DEXTERITY (1-5)	3.0	0.5	4.4	0.5	4.7	0.6	0.0006
EFFICIENCY (1-5)	2.4	1.2	3.4	0.5	5.0		0.0030
FORCE SENSITIVITY (1-5)	2.9	1.4	4.0	0.7	5.0		0.0155
ROBOTIC CONTROL (1-5)	2.5	1.4	3.4	0.5	4.7	0.6	0.0220
COLLISIONS (1-5)	3.6	0.9	4.4	0.9	4.7	0.6	0.2012
RESECTION COMPLETION (1-5)	2.9	1.7	5.0	0.0	5.0		0.0097
GEARS SCORE (1-35)	19.0	4.5	28.8	2.6	34.0	1.7	0.0001

FIG. 8

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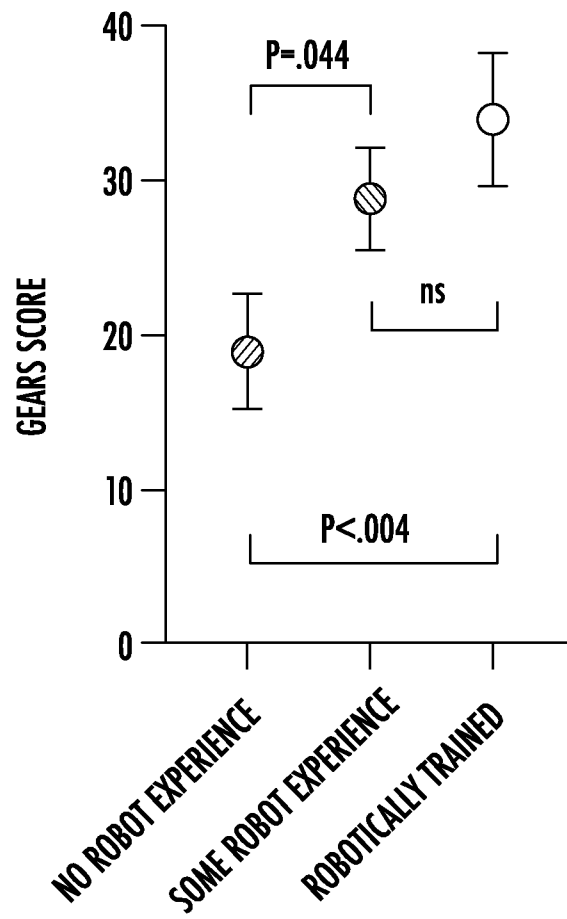


FIG. 9

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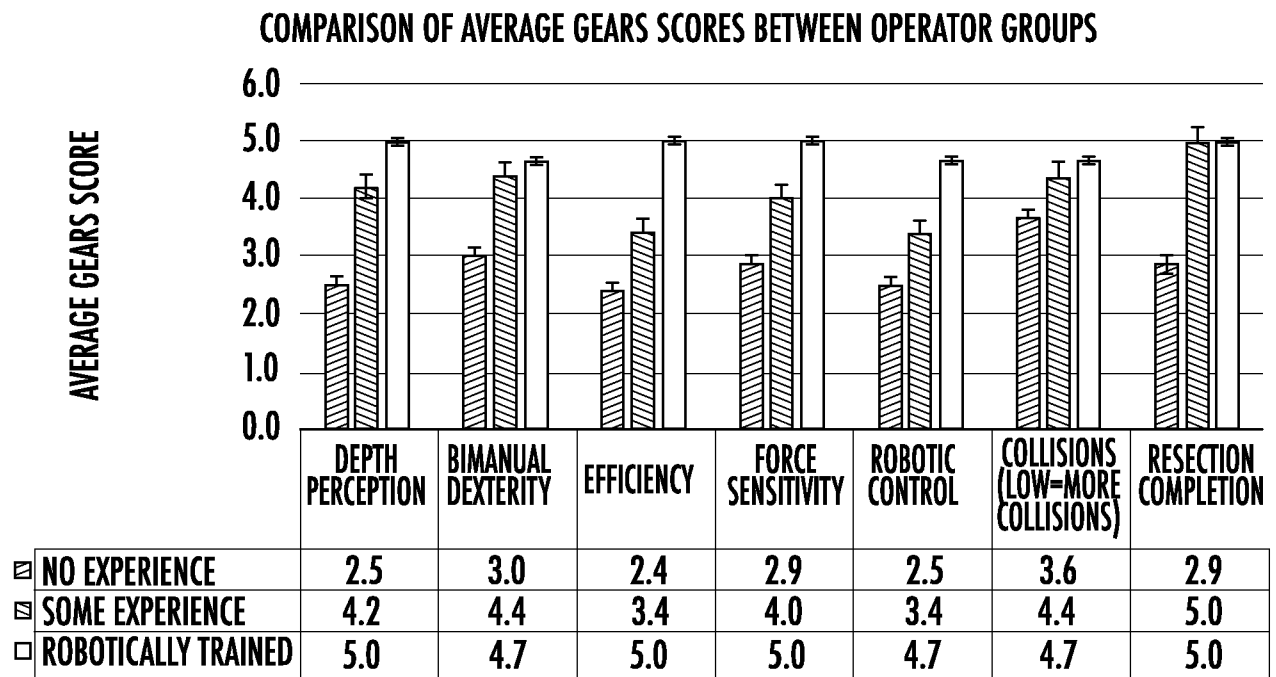


FIG. 10

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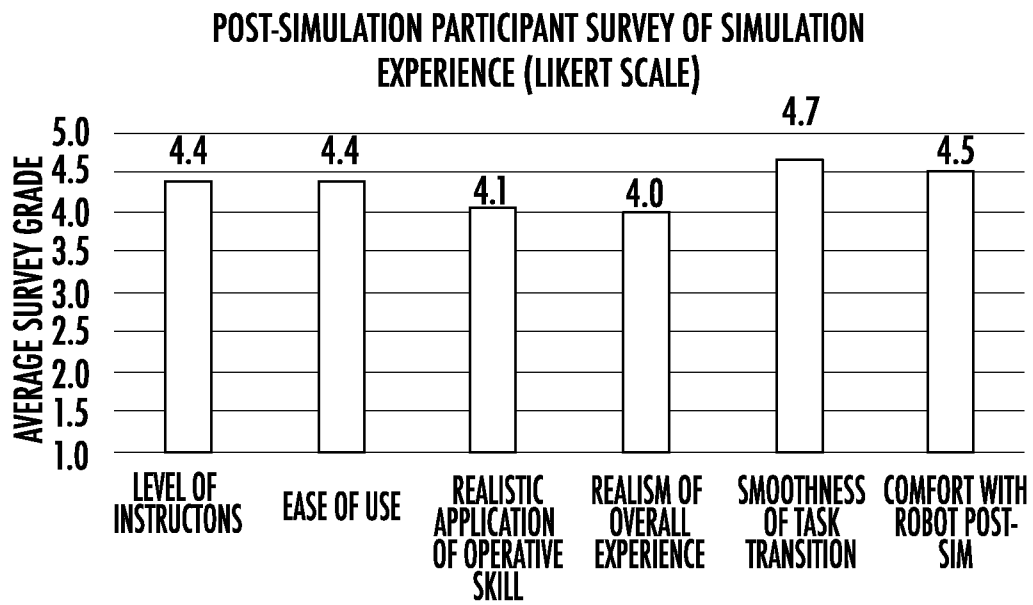


FIG. 11

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MODIFIED GLOBAL EVALUATIVE ASSESSMENT OF ROBOTIC SKILLS (GEARS) 2018

PARTICIPANT ID: _____

DATE SIMULATION: _____

EVALUATOR: _____

PLEASE CIRCLE THE NUMBER CORRESPONDING TO THE CANDIDATE'S PERFORMANCE IN EACH CATEGORY, IRRESPECTIVE OF TRAINING LEVEL.

DEPTH PERCEPTION:	1	2	3	4	5
	CONSTANTLY OVERSHOOTS TARGET, WIDE SWINGS, SLOW TO CONTACT		SOME OVERSHOOTING OR MISSING OF TARGET, BUT QUICK TO CORRECT		ACCURATELY DIRECTS INSTRUMENTS IN CORRECT PLANE TO TARGET
BIMANUAL DEXTERITY:	1	2	3	4	5
	USES ONLY ONE HAND, IGNORES NON-DOMINANT HAND, POOR COORDINATION		USES BOTH HANDS, BUT DOES NOT OPTIMIZE INTERACTIONS BETWEEN HANDS		EXPERTLY USES BOTH HANDS IN A COMPLEMENTARY WAY TO PROVIDE BEST EXPOSURE
EFFICIENCY:	1	2	3	4	5
	INEFFICIENT EFFORTS; MANY UNCERTAIN MOVEMENTS, CONSTANTLY CHANGING FOCUS OR PERSISTING WITHOUT PROGRESS		SLOW, BUT PLANNED MOVEMENTS ARE REASONABLY ORGANIZED		CONFIDENT, EFFICIENT AND SAFE CONDUCT, MAINTAINS FOCUS ON TASK, FLUID PROGRESSION
FORCE SENSITIVITY:	1	2	3	4	5
	ROUGH MOVES, TEARS TISSUE, INJURES NEARBY STRUCTURES, POOR CONTROL, FREQUENT SUTURE BREAKAGE		HANDLES TISSUES REASONABLY WELL, MINOR TRAUMA TO ADJACENT TISSUE, RARE SUTURE BREAKAGE		APPLIES APPROPRIATE TENSION, NEGLIGIBLE INJURY TO ADJACENT STRUCTURES, NO SUTURE BREAKAGE
COLLISIONS:	1	2	3	4	5
	MULTIPLE MINOR AND MAJOR COLLISIONS		SOME MINOR COLLISIONS/ FEW MAJOR COLLISIONS		FEW MINOR COLLISIONS/ NO MAJOR COLLISIONS
ROBOTIC CONTROL:	1	2	3	4	5
	CONSISTENTLY DOES NOT OPTIMIZE VIEW, HAND POSITION, OR REPEATED COLLISIONS EVEN WITH GUIDANCE		VIEW IS SOMETIMES NOT OPTIMAL. OCCASIONALLY NEEDS TO RELOCATE ARMS. OCCASIONAL COLLISIONS AND OBSTRUCTION OF ASSISTANT.		CONTROLS CAMERA AND HAND POSITION OPTIMALLY AND INDEPENDENTLY. MINIMAL COLLISIONS OR OBSTRUCTION OF ASSISTANT
RESECTION COMPLETION:	N/A	1	2	3	4
	<80% OF TUMOR RESECTED			80-90% OF TUMOR RESECTED	>90% OF TUMOR RESECTED

TIME TO COMPLETION: _____

TOTAL SCORE _____/45

FIG. 12

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2022/020721

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - G09B 23/30; A61C 11/00; G09B 23/00; G09B 23/28; G09B 23/32; G09B 23/34 (2022.01)

CPC - G09B 23/30; A61C 11/00; G09B 23/00; G09B 23/28; G09B 23/32; G09B 23/34 (2022.05)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

see Search History document

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

see Search History document

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

see Search History document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2019/246525 A1 (THE REGENTS OF THE UNIVERSITY OF CALIFORNIA) 26 December 2019 (26.12.2019) entire document	1-3, 20-24
A	US 3,520,071 A (ABRAHAMSON et al) 14 July 1970 (14.07.1970) entire document	1-3, 20-24
A	US 2007/0178429 A1 (BELL) 02 August 2007 (02.08.2007) entire document	1-3, 20-24
A	US 2016/0140879 A1 (HANANEL et al) 19 May 2016 (19.05.2016) entire document	1-3, 20-24

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"D" document cited by the applicant in the international application

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

28 May 2022

Date of mailing of the international search report

JUN 21 2022

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
P.O. Box 1450, Alexandria, VA 22313-1450

Facsimile No. 571-273-8300

Authorized officer

Taina Matos

Telephone No. PCT Helpdesk: 571-272-4300

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2022/020721

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claims Nos.: 4-19, 25-34
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.