# VR simulation with haptic feedback for medical procedure

# **Product Design Specifications | February 2, 2018**

Client: Dr. Ryan McAdams and Dr. Brandon Tomlin

**Advisor:** Professor Beth Meyerand

**Team:** Carter Griest griest@wisc.edu (*Team Leader*)

Isaac Hale <u>ihale2@wisc.edu</u> (Communicator & BSAC)
Joseph Campagna <u>jcampagna@wisc.edu</u> (BWIG)
Roberto Romero <u>rromero4@wisc.edu</u> (BPAG)

#### **Function**

The client wishes to develop a virtual simulation environment that accurately models a neonatal intubation procedure. Tentatively, the client desires that a virtual environment be created which mimics the upper respiratory tract, throat, and mouth of a neonate. The virtual system should precisely emulate a clinical environment in order to function as a novel neonatal intubation training method for physicians. Furthermore, the virtual components should be integrated with a haptic feedback motor arm which pairs physical traits to the virtual objects, thus mimicking a clinical procedure with only the use of virtual reality and a portable haptic feedback device. Ultimately, this device should serve as a virtual, but effective, surgery training method.

## **Client Requirements:**

- The virtual environment must accurately emulate the physical characteristics of an infant's tongue, larynx, trachea, and vocal cords to a degree of precision within the scope of a neonatal intubation procedure.
- A haptic interface must allow the user to provide manual inputs, to be visually represented precisely in a virtual environment. The interface must also register interactions in virtual space between the user's input and a preprogrammed objects, and relay these interactions back to the user via a haptic motor arm.
- The system must include a user-friendly software allowing changes in procedural specifications.

## **Design requirements:**

# 1. Physical and Operational Characteristics

#### a. Performance requirements:

- The system must be constructed for use in both clinical and rural settings, demanding portability and durability.
- The system must be capable of running up to 25 full simulations per day.
- A virtual environment must be capable of simulating neonates in the range of 1-10 lbs.

- **b. Safety:** Any electronic components must be enclosed within appropriate housing to minimize the risk of injury due to electric shock.
- **c. Accuracy and Reliability:** The system must be accurate to .02 mm to compete with current haptic feedback systems and provide a realistic surgical environment.
- d. Life in Service: The system must last at least 5 years with minimal maintenance.
- **e. Shelf Life:** The device will be stored inside and will not be exposed to extreme weather conditions. It should not need maintenance while not in use.
- **f. Operating Environment:** The system should be capable of operating in a variety of environments, including clinical and outdoor settings. The virtual simulation will be perfected by using feedback from expert neonatologists to accurately emulate a neonatal intubation procedure.
- **g. Ergonomics:** The device should be intuitive to use and feel very similar to tools used during neonatal intubation such as the laryngoscope and endotracheal tube.
- h. Size: The device must be small enough be carried around in a backpack or other case.
- **i. Weight:** The device must weigh less than 40 lbs, light enough to be easily transported in a backpack or other case.
- **j. Materials:** The system will be comprised of a pair of virtual reality goggles, a haptic feedback motor arm, and any computer hardware required to render the environment and power the system.
- **k. Aesthetics, Appearance, and Finish:** The virtual reality should not be blurry. The user should be able to interact with the environment without noticeable buffering.

#### 2. Production Characteristics

- **a. Quantity:** One functional prototype will suffice for BME 301. Ultimately, however, the aim is to provide worldwide accessibility.
- **b. Target Product Cost:** The device should cost under \$3000.

## 3. Miscellaneous

- **a. Standards and Specifications:** If successful, the device would require IRB and FDA approval to serve as a credible source of medical training.
- **b. Customer:** The system will be used by training physicians who are practicing neonatal intubation procedures. Consequently, they will demand a realistic virtual environment with physical characteristics which accurately model a neonates anatomy and physiology.
- **c. Patient-related concerns:** No concerns should arise from the use of this device as it will serve as an additional form of medical training, not an alternative to current training.
- **d. Competition:** Competition exists among virtual reality platforms, but to our knowledge, there only exists a single haptic feedback system on the market currently, and there are no integrative VR and haptic feedback systems which are used for medical simulation.