

Motivation: Combat infant hypothermia

- Hypothermia contributes to 18% - 42% of 4 million annual infant deaths in the developing world [1].
- Current solutions (e.g., incubators and kangaroo care) can be expensive, high maintenance, or impractical.
- Hospitals lack sufficient staff, space, and funding.
- Mulago Hospital in Kampala, Uganda:
 - Two nurses for 60 neonates per day in the NICU
 - Only two (out of 20) incubators working

The lack of affordable and functioning incubators in low-resource settings makes it difficult to effectively combat infant hypothermia in the third world.



Objective: Provide a warm and safe environment

Our modular system is an innovative, effective, and viable solution against infant hypothermia.

Functional Requirements

- Increase infant temperature
 - Based on gestation, age, and weight of the baby
 - Controlled heating to achieve normal temperature range of 36.5-37.5 °C [2,3]
- Maintain temperature range
 - Continuous monitoring of infant temperature
 - Feedback mechanism
 - Alarm LED which signals infant overheating or fever

Constraints

- Affordable
- Biocompatible and Safe
- Durable
- Easy to Maintain and Repair
- Easy to Sterilize and Clean
- High Controllability
- High Infant Visibility
- Intuitive
- Easy to Use
- Versatile

Our Solution: Warm water circulation system

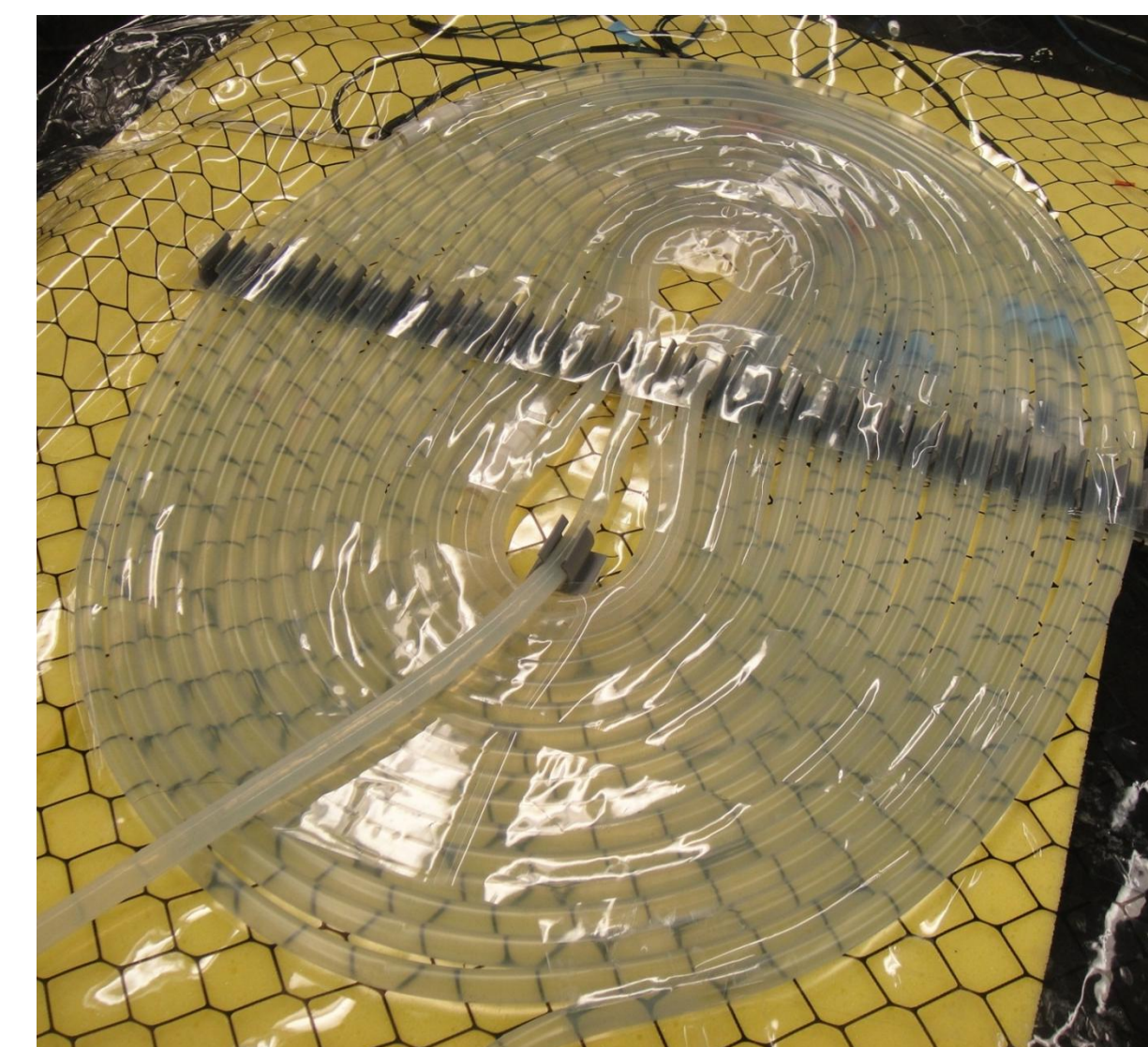


Figure 1. Mat prototype

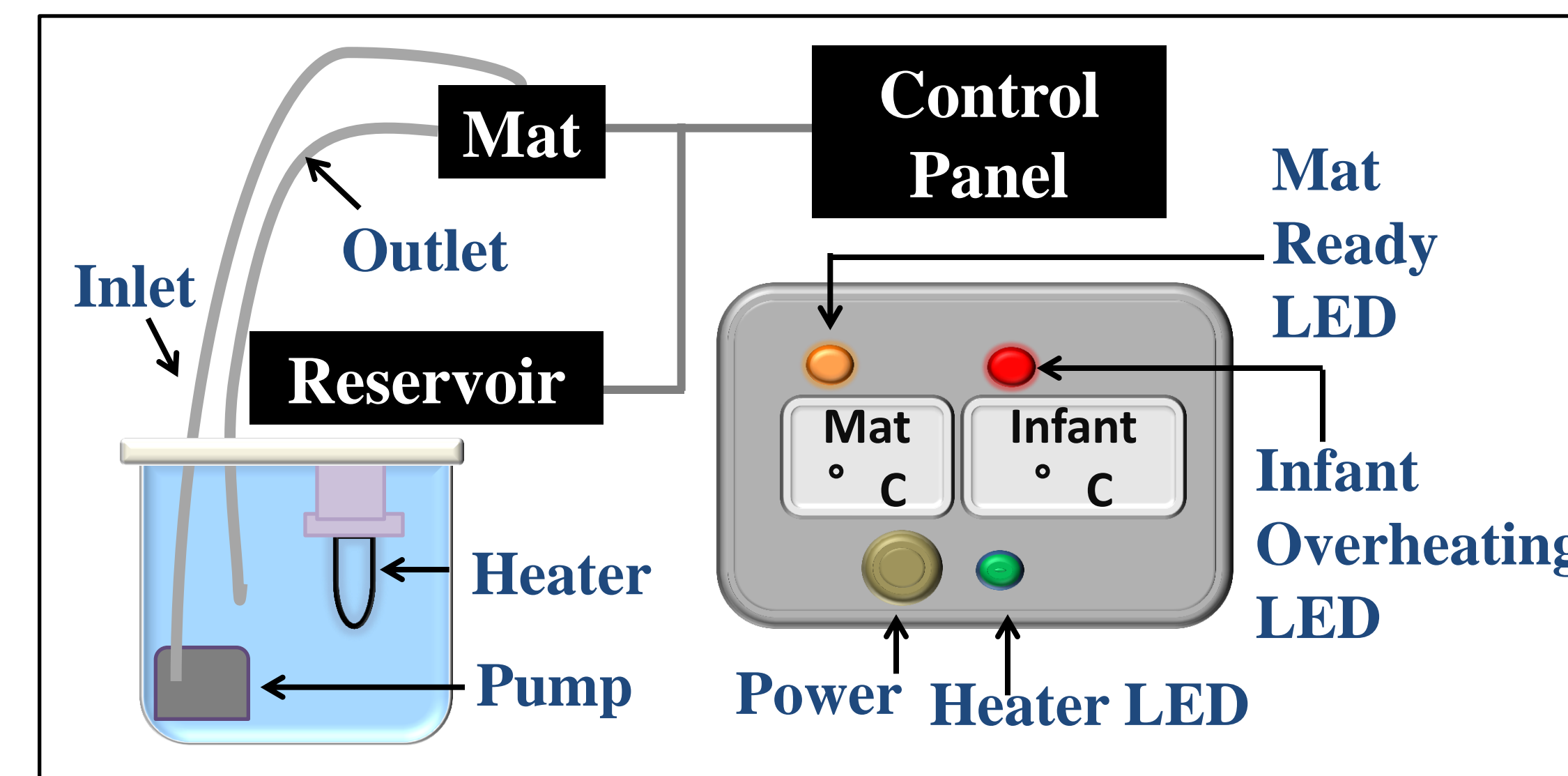


Figure 2. System diagram

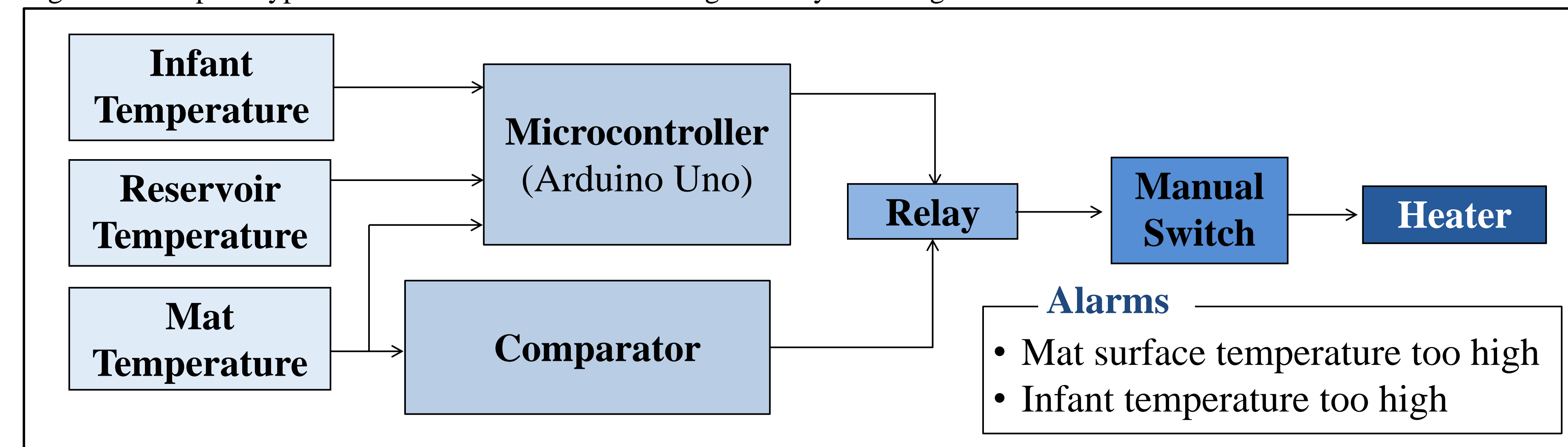


Figure 3. Schematic diagram of the feedback system

Feedback Algorithm

- Heats and maintains the reservoir at a certain temperature
- Once the mat reaches the set temperature, feedback switches to maintaining mat temperature
- Monitors the temperature of the infant and maintains it at 37 °C

Fail-safes

Circuitry

- Resistors to drive voltage down in case of thermistor disconnection
- Comparator in case the microcontroller fails

Code

- Heater turns off if:
- Thermistor disconnects
 - System overheats
 - Mat overheats
 - Infant overheats

Test: Raise and maintain biofluid temperature

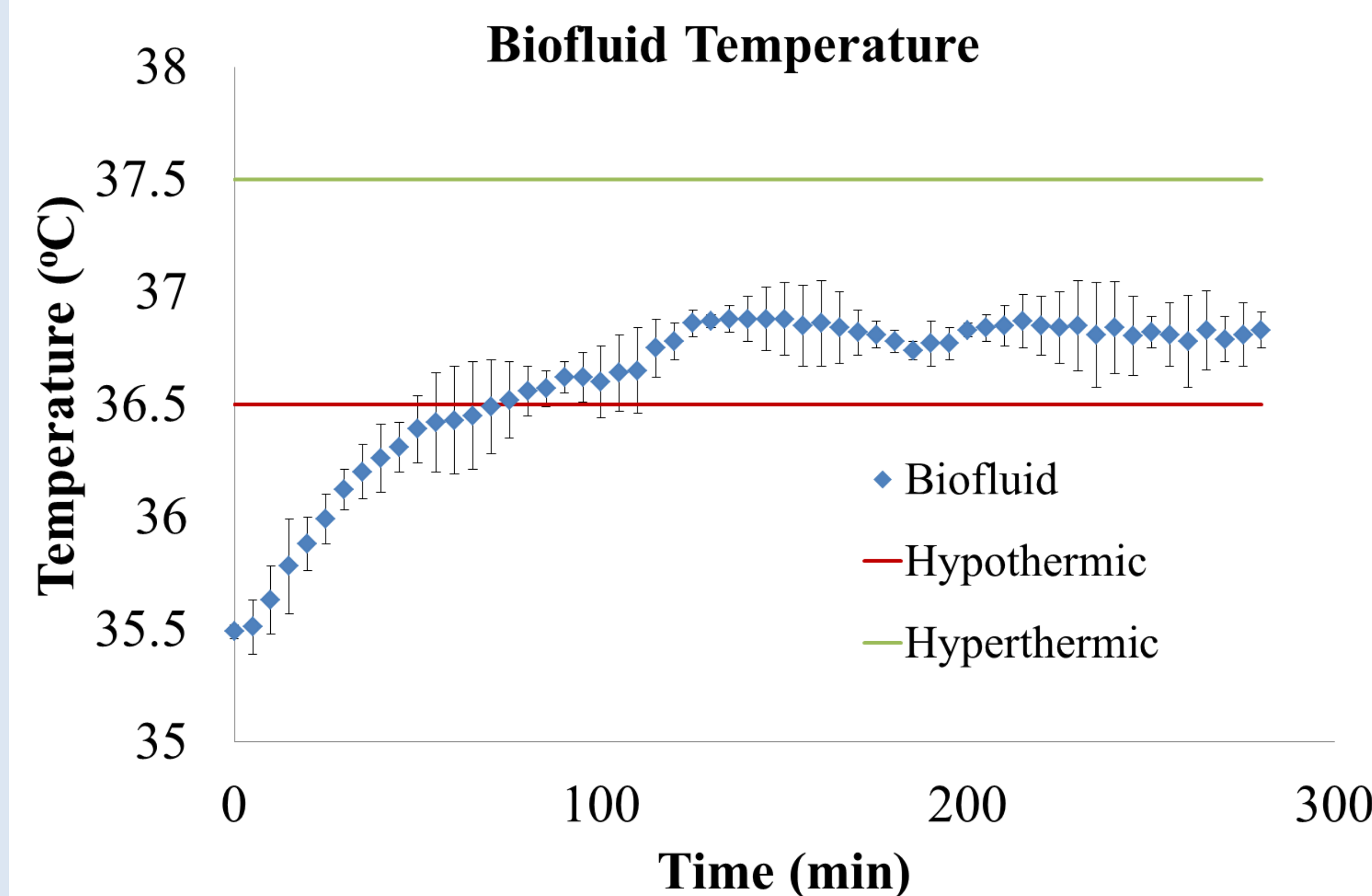


Figure 4. Average biofluid temperature vs. time (n = 3)

Procedure

- 1L biofluid model (35.5 °C) was placed on the mat. The temperature of the biofluid was monitored.

Results

- Heated and maintained the biofluid to within normal range of 36.5 - 37.5 °C
 - Control (no heating): biofluid decreased from 37 °C to 34 °C in 18 minutes
- 2.3 kWh (\$0.36) per day with continuous use
- Equivalent to using a 100W light bulb for 24hr

Cost: Use of readily available and cheap components

Component	Cost (\$)	Modularity
PVC Tubing	9.20	<ul style="list-style-type: none"> Thermostat heater Microcontroller (\$0.50) Tubing Case/Mat
PVC Sheet	12.34	
1000 W Heater	11.00	
200 gal/hr Pump	19.36	
3.5 gal Bucket and Lid	13.96	
Arduino Uno	15.88	Sterilization <ul style="list-style-type: none"> Water: bleach PVC sheet and tubing: ethanol
Electronics & Misc	25.00	
Total	106.74	

Table 1. Manufacturing cost

Conclusions and Future Work

Conclusions

- Can effectively raise the temperature of a biofluid baby model
- Can maintain the temperature of the biofluid within the normal range (36.5-37.5 °C) for up to 5 h. Expect that the heating time can be significantly extended

Future Work

- Optimize design and minimize cost and power consumption
- Fold over component to heat infant from both sides.
- Increase heating unit versatility to allow multiple mat unit warming
- Implement a back-up power source (e.g., battery) in case of power failure

References

- UNICEF. Progress for children: a world fit for children statistical review. New York, NY: United Nations Children's Fund; 2007.
- Voorhoeve, H. Aspects of adapted newborn care in rural hospitals. University of Leiden, The Netherlands. October 2005.
- Hypothermia. *The Merck Manuals: The Merck Manual for Healthcare Professionals*. <http://www.merckmanuals.com/professional/print/sec21/ch319/ch319d.html>. Accessed March 1st, 2012.

Acknowledgements

We would like to thank our instructors, advisors, and consultants: Aaron Kyle, Ph.D.¹, Elizabeth Hillman, Ph.D.¹, Keith Yeager¹, Sarah De Leo¹, David Jangraw¹, Lance Kam, Ph.D.¹, Genevieve Brown¹, Margaret Nakakeeto-Kijjambu, MD², Richard Polin, M.D.³, Rakesh Sahni, M.D.³, Helen Towers, M.D.³, Yvonne Vaucher, M.D.⁴, and David Vallancourt, Ph.D.⁵. Funding was provided by the department of biomedical engineering at Columbia University.

¹Biomedical Engineering Dept., Columbia University, ²Mulago Hospital, Uganda, ³CUMC Pediatrics, ⁴UCSD, ⁵Electrical Engineering Dept., Columbia University.