



The EVLP System used to assess the lungs in this case study is currently undergoing clinical investigation in the United States as part of a clinical trial PXUS 14-001 (NCT02234128).

The study is sponsored by Lung Bioengineering, a wholly owned subsidiary of United Therapeutics Corporation.

The system is for investigational use only.

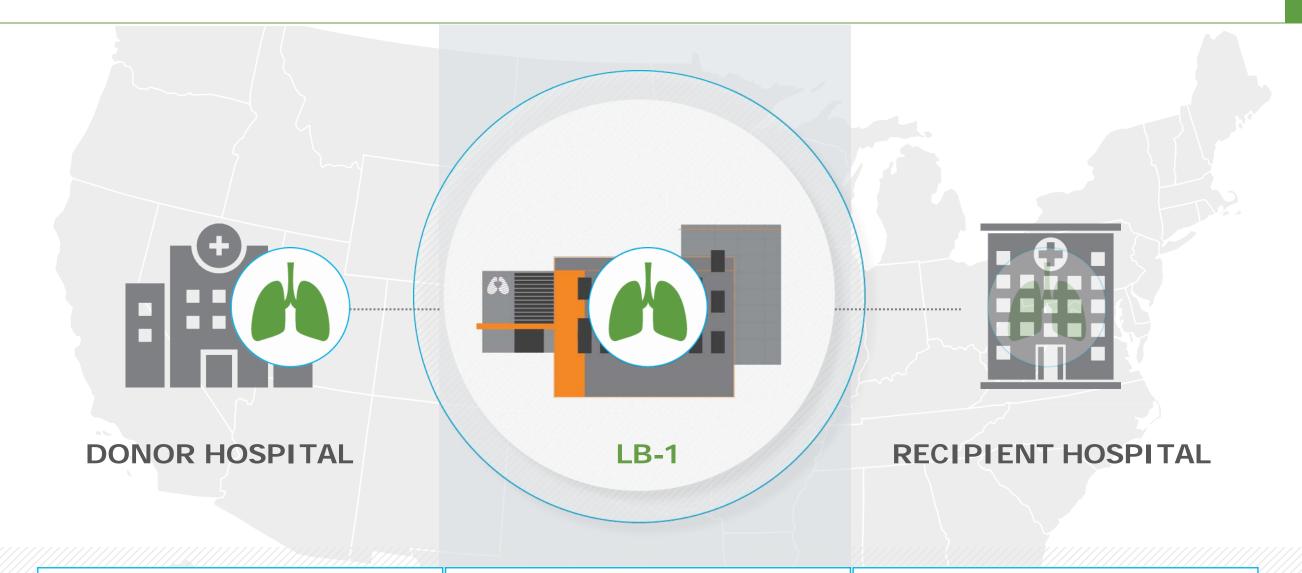
BACKGROUND



- According to Organ Procurement and Transplantation Network (OPTN) data, in 2017, the national lung utilization rate was 23.7%
- Lungs declined in the donor operating room often have little to no opportunity for reallocation due to time constraints on the recipient transplant centers and unacceptable cold ischemic times.
- Ex Vivo Lung Perfusion (EVLP) may allow an Organ Procurement Organization (OPO) extended time to allocate lung(s) to a center beyond the time of the deceased donor cross clamp time.

HOW DOES OUR STUDY WORK?

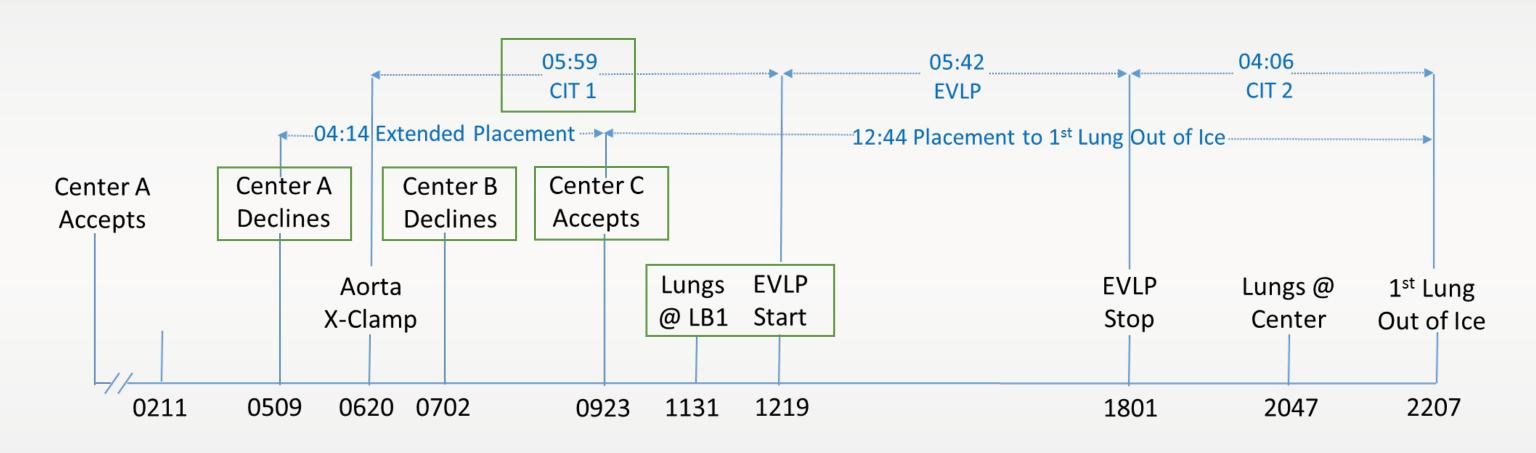




Time allowed by trial protocol

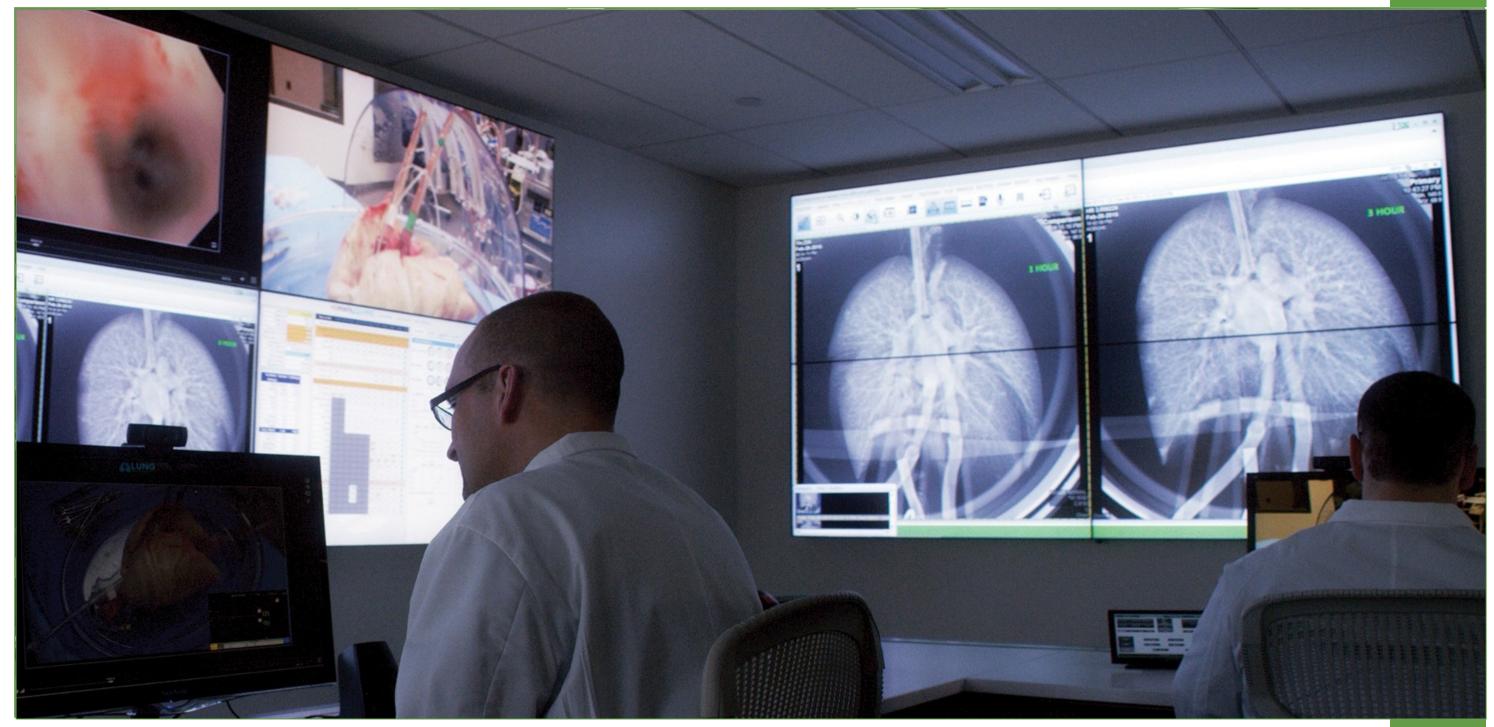
CIT-1	EVLP	CIT-2
🕚 10 h	① 3-6 h	① 6 h
Donor cross clamp to EVLP start	Normothermic Perfusion	EVLP end to start of transplant procedure





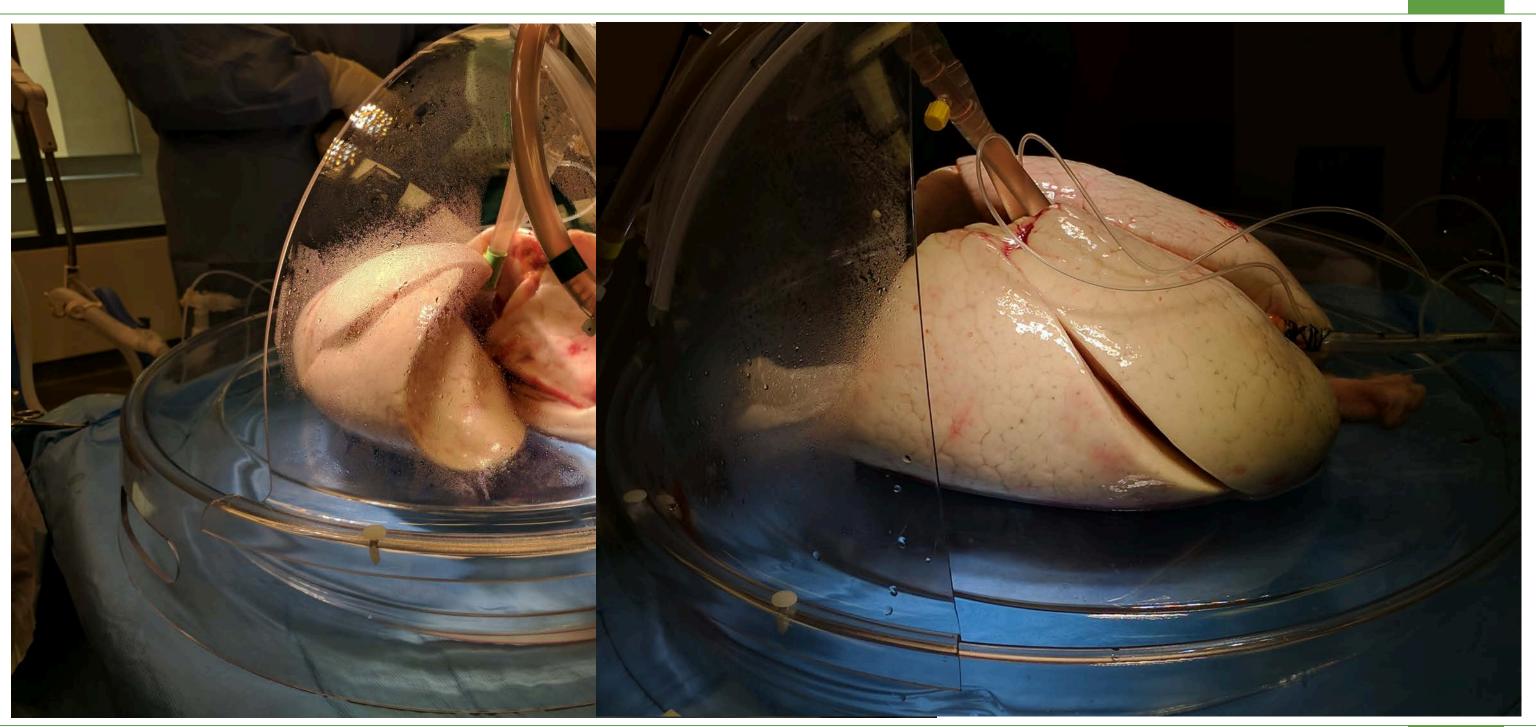
PROCEDURE ROOM





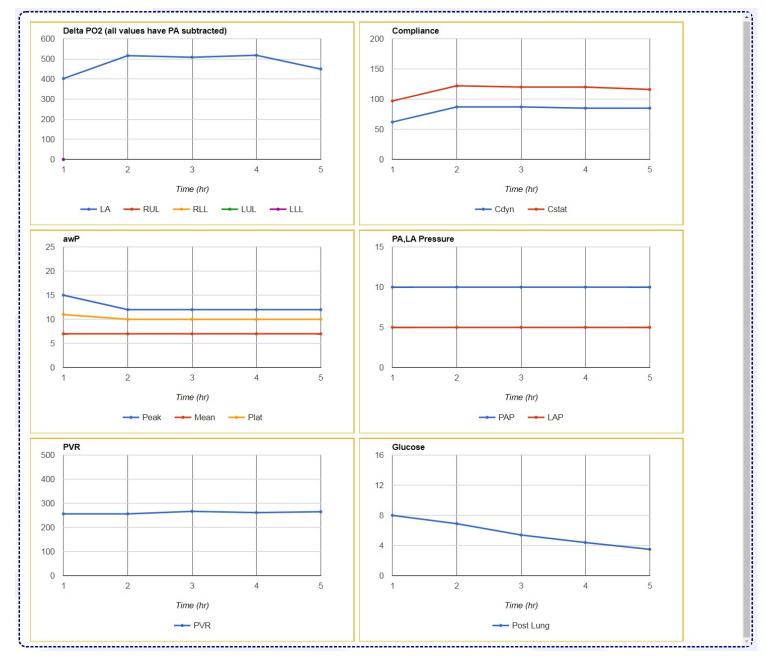






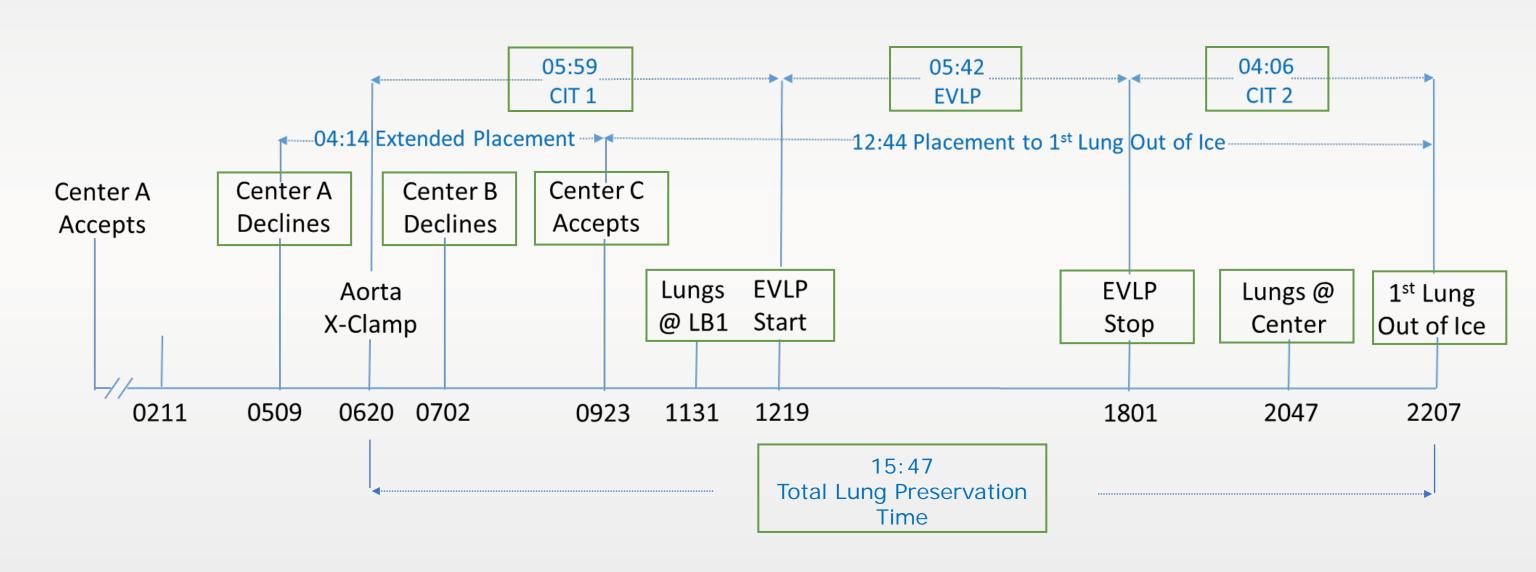
EVLP





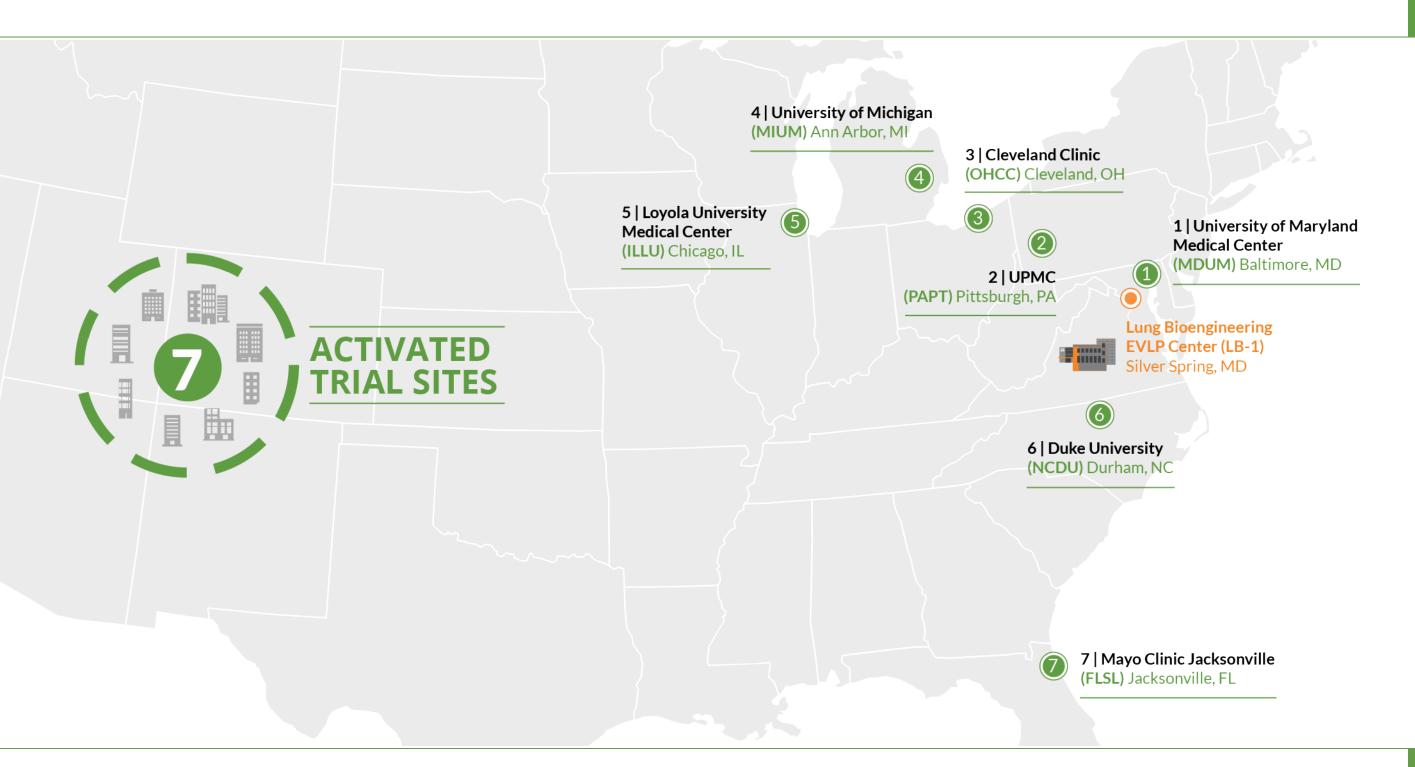






ACTIVATED CLINICAL TRIAL SITES







THANK YOU