UMMS TRANSGENIC ANIMAL MODELING CORE (TAMC)

iPSC INDUCTION (Reprogramming) INDUCTION OF PLURIPOTENCY IN HUMAN PRIMARY CELLS

The Facility will perform:

- 1. Transduction of cells in 6-well plate format with non-integrating Sendai virus to transduce the four Yamanaka factors (Klf4, Oct4, Sox2, and cMyc).
- 2. Splitting transduced cells onto 10 cm dishes with inactivated feeder cell layers or feeder-free extra-cellular matrices. Once iPS colonies are formed, up to 20 colonies displaying the most ES cell- like morphology will be picked individually and expanded.
- 3. Based on morphology, 5 to 6 clones will be selected and further propagated as individual clones.
- 4. After 6 passages loss of the virus will be tested by immunostaining against viral antigens. Retention of pluripotency in the viral-negative lines will be confirmed by immunostaining.
- 5. Up to 5 individual iPS lines derived from a given primary sample will be provided to the PI as frozen cells and/or in culture. Sister-vial of frozen cells will be retained by the TAMC if the Core is to provide further services.
- 6. If required, the Core will also provide frozen cell pellets to facilitate subsequent nucleic acid analysis.

The time required to generate iPS lines from a given primary sample will vary, but is estimated to require 8-12 weeks. Although the pluripotency of generated lines (as determined by *in vivo* assays) cannot be guaranteed for any sample, the TAMC will do all it can to ensure successful induction, and repeat certain steps if the initial performance was judged to be sub-optimal.

Charges: \$4,000.00 per cell line

Please also see the description of extended "iPS induction - Plus" service.

PI Name		
Department	Notes:	
Account Number		
Number of Lines	-	
Date Received		
Total Charges \$		
x	_ X	
UMMS Investigator Date	UMMS TAMC	Date