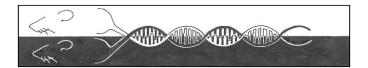


Transgenic and Stem Cells Service Unit



TRANSGENIC SERVICE REQUEST

Date						
Principal Investigator's Name Name of Person Conducting Experiment						
Institution						
Lab Contact	Lab Contact E-Ma	uil				
Lab Contact Telephone						
Billing Contact	Billing Contact E-	Mail				
Billing Telephone	Billing Fax					
Billing Address						
Gene Name IMM Abbreviated Name						
Agent characteristics (gene function):						
Expression of foreign gene/ protein produce	ed:					
Are there any toxic molecules or oncogenes	s produced: Yes No					
Method Used to Purify DNA (e.g. Gene Classical Control of the Classical	'ean)					
Type of Service: Transgenic Line Tra	nnsient Transgenics Mo	ouse Strain				
PO (or account # for UTHealth) for project	: :					
Acknowledgements: By signing this form, the principal investige Service Unit in any publication that describ were generated at the University of Texa Cells Service Unit. We wish to thank Dr. E expertise in generating the mice.).	pes the genetically-engineers s Health Science Center-	ed mice (e.g. Transgenic mice Houston, Transgenic and Stem				
Principal Investigator's signature		Date				
Dr. Zsigmond's signature		Date				

INSTRUCTIONS

Shipment of DNA and Forms

Fill form out and fax or mail it with the DNA construct to:

University of Texas Health Science Center- Houston

The Brown Foundation Institute of Molecular Medicine,

Transgenic and Stem Cells Service Unit

c/o Aleksey Domozhirov

1825 Pressler Street, Suite 611, Houston, TX 77030

Telephone: (713) 500-2452 **Fax**: (713) 500-2208 **E-Mail**: transgenic@uth.tmc.edu

DNA Preparation

The requirements for an acceptable linearized DNA preparation are:

- (1) 260/280 of 1.7 to 2.0
- (2) Concentration of 0.1 to 0.2 micrograms/microliter
- (3) Volume of 50 to 70 microliters

Mouse Strain

Specify mouse strain needed as background strain. The standard mouse strain is C57BL/6.

Type of Service

Select "Transgenic Line" or "Transient Transgenics".

Transient transgenics are made when the investigator wishes to examine the offspring prior to birth for developmental studies, or if there is embryonic lethality.

<u>Table:</u> Please fill out the first line only.

Method	Dilution	O.D. 260	O.D. 280	260/280	DNA concentration

<u>Picture:</u> Please run 1-2 microliters of the DNA construct on a gel next to a molecular weight marker.

Place Picture Here