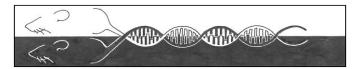


Transgenic and Stem Cells Service Unit



TRANSGENIC SERVICE REQUEST

Date	
Principal Investigator's Name	
Name of Person Conducting Experiment	
Department	
Institution	
Lab Contact	Lab Contact E-Mail
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 produced:	
Are there any toxic molecules or oncogenes produc	ed: Yes No
Method Used to Purify DNA (e.g. Gene Clean)	
Type of Service: Transgenic Line Transient T	ransgenics Mouse Strain
PO (or account # for UTHealth) for project:	
<u>Acknowledgements</u> : By signing this form, the principal investigator agree Service Unit in any publication that describes the ge- were generated at the University of Texas Health Cells Service Unit. We wish to thank Dr. Eva Zsign expertise in generating the mice.).	enetically-engineered mice (e.g. Transgenic mice a Science Center- 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.

Table:	Please fill out the first line only.	
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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