Disclosure

No financial relationships to disclose.
Center for Fetal Monkey Gene Transfer for Heart, Lung, and Blood Diseases
(Established in 2001)
Program Goals

Evaluate the safety and efficiency of gene transfer strategies as they emerge

Use established monkey models to explore fetal approaches for heart, lung, and blood diseases

Provide NHLBI-funded investigators with essential expertise, services, and resources
Center for Fetal Monkey Gene Transfer for Heart, Lung, and Blood Diseases

Services Offered to Grantees

- Study development and design
- Gene transfer (all age groups)
- Transplant models (autologous, allogeneic, xenogeneic)
- In vivo imaging (ultrasound, microPET, optical)
- Physiologic assessments
- Tissue harvests (collect, process, analyze)
- Laboratory (e.g., molecular, cellular including qRT-PCR, immunophenotyping, sorting, immunoselection, morphology, morphometry, laser capture)
Access to Services

• NHLBI-funded investigators

• Submit Letter of Intent (LOI) by June 30, 2010

• If LOI is approved, a full proposal template will be provided

Look for annual calls for LOI submission in June issue of *Molecular Therapy* or check the website
Funded investigators may choose to:

• Provide their vectors for study and all work will be completed by Center personnel
• Participate in analysis of samples collected
• Participate in the study as a visiting scientist
Contact Information

Visit the website:
www.CFMGT.ucdavis.edu

E-mail Dr. Alice Tarantal at:
aftarantal@primate.ucdavis.edu
Gene Therapy Resource Program
(Initiated in March 2007)
History of Program

Major Research Challenges Identified by NHLBI Gene Therapy Working Group at June 2005 Meeting:

• Producing large scale, well-characterized viral vectors under current Good Manufacturing Practices for use in clinical trials
• Conducting pharmacology and toxicology studies in small and large animal models
• Meeting the regulatory requirements of the FDA, IBCs, IRBs, NIH RAC, and DSMB

The NHLBI responded to the recommendations of the Working Group by establishing the Gene Therapy Resource Program in March 2007
Goals

Facilitate the translation of gene therapy research into clinical interventions

Provide resources for gene therapy research primarily in heart, lung, and blood diseases

Provide resources to investigators at other NIH institutes through transfer of funds
## Program Infrastructure

**Clinical Coordinating Center**  
Social & Scientific Systems, Inc.  
PI: Susan B. Sepelak, MS, MSM

<table>
<thead>
<tr>
<th>Preclinical Vector Core</th>
<th>Clinical Lentivirus Vector Core</th>
<th>Clinical AAV Vector Core</th>
</tr>
</thead>
<tbody>
<tr>
<td>University of Pennsylvania</td>
<td>Indiana University</td>
<td>The Children’s Hospital of Philadelphia</td>
</tr>
<tr>
<td>PI: James Wilson, MD, PhD</td>
<td>PI: Kenneth Cornetta, MD</td>
<td>PI: Fraser Wright, PhD</td>
</tr>
</tbody>
</table>

| Pharmacology/Toxicology Core | | |
|-----------------------------| | |
| Lovelace Biomedical & Environmental Institute | | |
| PI: Janet Benson, PhD | | |

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Gene Therapy Resource Program
Preclinical Vector Core
University of Pennsylvania
Preclinical Vector Core

Produces large- and small-scale viral and non-viral vectors for studies in basic research directed toward clinical applications

Produces AAV (over 30 serotypes including 1, 2, 5, 6, 7, 8, 9, rh10), adenoviral, lentiviral (various pseudotypes) as well as plasmid-based vectors

Provides a variety of expression cassettes for tissue-specific expression (e.g. heart, liver and lung)

Provides immunology testing services including neutralizing antibody assays, interferon gamma ELISpot, and intracellular cytokine staining to adenoviral vectors and AAV capsid and transgene products
Clinical-Grade AAV Vector Core
Children’s Hospital of Philadelphia
Produces scalable clinical-grade adeno-associated virus (AAV) vectors for use in clinical studies

Produces GMP process-comparable AAV vectors for use in pharm/tox or other studies

Provides Chemistry, Manufacturing, and Controls (CMC) and Certificate of Analysis (COA)

Assists in vector (cis) plasmid design to achieve optimal safety and productivity

Prepares certified components (plasmid DNA, HEK293 Master Cell Bank) and reagents required for clinical vector manufacture
Clinical-Grade Lentiviral Vector Core
Indiana University
Clinical-Grade Lentivirus Vector Core

- Produces scalable clinical-grade lentiviral vectors for use in clinical studies
- Produces GMP process-comparable lentiviral vectors for use in pharm/tox or other studies
- Evaluates and optimizes lentiviral vector constructs intended for clinical use
- Provides pilot runs for pre-clinical evaluation prior to large-scale production
- Assists in release testing to certify vectors for clinical use
- Provides Chemistry, Manufacturing, and Controls (CMC) and Certificate of Analysis (COA)
Pharmacology / Toxicology Core
Lovelace Biomedical and Environmental Research Institute
Performs toxicology testing and biodistribution studies of vectors in large and small animal models as a prerequisite for use in clinical studies.

Conducts studies according to Good Laboratory Practice Guidelines and prepare final study reports.

Assists with pharmacology/toxicology study design.

- Animal species available include rodents, rabbits, ferrets, beagle dogs, and non-human primates.
- Facilities for necropsy, clinical pathology, histopathology, and assessment of vector biodistribution and transgene expression.
Manages the RSA receipt, review, disposition, and tracking processes

Provides management support to the GTRP Steering Committee

Assists NHLBI with development and maintenance of program SOPs, policies, and bylaws

Maintains the GTRP website and electronic forms

Provides regulatory assistance to investigators via the RSA process

Manages disbursement of clinical trial funds to investigators

Provides logistical and operational support for meetings and teleconferences
Clinical Coordinating Center
Social & Scientific Systems, Inc.
Access to Program Resources

Step 1: Investigator Registration and Approval

Step 2: Submit RSA

WHAT IS THE NHLBI GENE THERAPY RESOURCE PROGRAM?

The NHLBI Gene Therapy Resource Program (GTRP) facilitates the translation of gene therapy research into clinical interventions. The GTRP provides resources for gene therapy research primarily in heart, lung, and blood diseases as reflected in the NHLBI Mission (http://www.nhlbi.nih.gov/about/crg/mission.htm). Requests for resources for gene therapy research that are consistent with the missions of other NIH Institutes may also be considered by the Program.

Resources are provided in the form of preclinical and clinical-grade vector production, pharmacology/toxicology testing, immunology testing, clinical trials funding assistance, and regulatory support at no cost to the investigator. Investigators must first receive approval of their Registration with the Program in order to request resources.

The GTRP, directed by the NHLBI Gene Therapy Group, consists of three vector production cores, a pharmacology/toxicology testing core, and a clinical coordinating center. A Scientific Review Board and Steering Committee review Request for Service Applications and make recommendations to the NHLBI Gene Therapy Group regarding the applications’ scientific merit, feasibility, and compatibility with the Program’s mission.
National Eye Institute (NEI)

To support an intramural investigator requesting GMP process-comparable AAV vector production for use in pharmacology/toxicology studies related to retinoschisis.

The NIH-RAID Program and the National Institute of Neurologic Disorders and Stroke (NINDS)

To support an extramural investigator requesting GMP process-comparable AAV vector production for use in pharmacology/toxicology studies and clinical grade AAV vector production for use in a phase I clinical trial. Both studies are related to Parkinson’s Disease.
For More Information

Visit our website: www.gtrp.org
Visit our exhibit booth 311

E-mail the **Clinical Coordinating Center** at:

* gtrpccc@s-3.com

or

E-mail **Sonia I. Skarlatos, Ph.D.** at:

* skarlats@mail.nih.gov