

BETA CELL BIOLOGY CONSORTIUM

<http://www.betacell.org>

Description of project

- The mission of the BCBC is to facilitate interdisciplinary approaches that will advance our understanding of pancreatic islet development and function. The long-term scientific goal is to develop a cell-based therapy to restore normal insulin production and action to diabetic patients.

Accomplishments

- Collaborative research efforts focused on understanding fully how endogenous beta cells are made by studying pancreatic development, with the hope of producing pancreatic islets in tissue culture.
- Infrastructure to collaboratively generate necessary reagents, mouse strains, antibodies, assays, protocols and technologies that are beyond the scope of any single research effort. The BCBC Coordinating Center at Vanderbilt University oversees the BCBC website (www.betacell.org), and coordinates all activities of the BCBC, including scientific cores, reagent databases, steering committee meetings, investigator retreats, the “pilot and feasibility studies” program, and the special “seeding collaborative studies in beta cell biology” program.
 - Antibody Core/Subcommittee that is developing antibodies to mouse and human cell surface antigens and important markers of progenitor cells
 - Mouse ES Core/Subcommittee that is creating a set of genetically modified ES cells that can be used to generate mice for the purpose of studying pancreatic development and diabetes
 - EPConDB, a searchable database that contains information about genes expressed in the mouse and human pancreas during development
<http://www.cbil.upenn.edu/EPConDB/>;
Kaestner/Stoeckert, University of Pennsylvania
 - Microarray Core that is producing and distributing mouse and human cDNA microarrays <http://www.betacell.org/php/microarrays.php>;
Kaestner, University of Pennsylvania
- Ongoing stem cell-related research efforts include the development of key reagents to permit the prospective isolation of stem/progenitor cells in both neonatal and adult animals, and development of protocols to efficiently differentiate embryonic and adult stem cells into pancreatic islet tissue appropriate for transplantation, and the development of assays to test the efficacy of pancreatic stem/progenitor cells as a cure for type 1 diabetes. The Human Cell

Culture Committee is working collaboratively to conduct gene expression profiling of human pancreatic progenitor cells.

Future Directions

- The Beta Cell Biology Consortium will continue to work collaboratively to generate and distribute necessary reagents, mouse strains, antibodies, assays, protocols, and technologies that will facilitate research in pancreatic islet development and function.
- Research will be conducted in three areas of scientific focus:
 - Beta Cell Development--understanding developmental pathways required to produce a fully functioning pancreatic islet;
 - Beta Cell Regeneration--understanding the mechanisms of beta cell regeneration in the adult animal and human;
 - Stem Cell Biology--understanding the nature of stem/pancreatic progenitor cells during normal development and in adults.

Materials to be made available to researchers

The PancChip 5.0 is now available for order. For more information, please visit <http://www.cbil.upenn.edu/EPConDB/>.

Please visit <http://www.betacell.org> for more information on obtaining the following materials:

- Antibodies
- BAC Clones
- Microarrays of Mouse and Human Islet Genes
- Targeting Vectors
- Transgenic mice

Participants

Sponsor: National Institute of Diabetes and Digestive and Kidney Diseases

Participating Institutions

Hagedorn Research Institute
Harvard University

INSERM, Strasbourg
NIDDK, NIH
University of California, Irvine
University of California, San Francisco
University of California, San Diego
University of Chicago
University of Colorado, Boulder
University of Colorado, Denver
University of Minnesota
University of Pennsylvania
University of Wisconsin
Vanderbilt University
Vrije Universiteit Brussels

Steering Committee

Chair:

Mark Magnuson

Vanderbilt University

Committee Members:

Graeme Bell

University of Chicago

Mike German

University of California, SF

Joel Habener

Harvard University

John Hutton

UCHSC

Klaus Kaestner

University of Pennsylvania

Ole Madsen

Hagedorn Research Institute

Sheryl Sato

NIDDK, NIH

Lori Sussel

UCHSC

Catherine Verfaillie

University of Minnesota

Gordon Weir

Harvard University

Chris Wright

Vanderbilt University

Ex Officio:

Carol Haft

NIDDK, NIH

David Harlan

NIDDK, NIH

Marvin Gershengorn

NIDDK, NIH

Doug Melton

Harvard University

Phil Smith

NIDDK, NIH

Chris Stoeckert

University of Pennsylvania

Program Coordinator:

Lisa Rouse

Vanderbilt University