

Stem Cell Research Oversight Committee Reporting Form

Pursuant to Health and Safety Code §125119, at least once per year a Stem Cell Research Oversight (SCRO) Committee must conduct continuing review of each human stem cell research project under its purview and report to the California Department of Public Health (CDPH) on each project. This mandate does not apply to research that is funded by the California Institute for Regenerative Medicine.

Please electronically send the following information to CDPH Human Stem Cell Research Program by *June 30, 2008* at stemcell@cdph.ca.gov. In addition to the information you provide, each facility that performs oocyte retrieval or procurement for a research project must also provide to CDPH research subject information specified in Health and Safety Code §125330 and Chapter 2 (commencing with §125330).

[CDPH Human Stem Cell Research Information](#)

SCRO Committee Information

Company or Institutional Affiliation of SCRO Committee:		
Contact Information:		
Name	Phone, Fax, Email	Address
Number of human stem cell research projects the SCRO Committee has reviewed this reporting year, January 1, 2007- June 30, 2008:		

Please provide the following information for each research project you have reviewed this reporting year:

1. Project title
2. Principle and Co-Investigators
3. Contact information
4. Protocol review status
5. Description and status of the project
6. Description of any unanticipated problems or serious continuing investigator noncompliance issues
7. Description of SCRO Committee actions taken to respond to problems/issues

Research Project Information

Research Project Title:		
SCRO Committee #:		
Principle Investigator:		
Name	Institution or Company Affiliation	
Co-Investigator(s):		
Name	Institution or Company Affiliation	
Co-Investigator(s):		
Name	Institution or Company Affiliation	
Contact Information:		
Name	Phone, Fax, Email	Address
Anticipated Duration of Project:		
Project Commencement Date:		
Expected Completion Date:		
Protocol Status:		
<input type="checkbox"/> New <input type="checkbox"/> Modification <input type="checkbox"/> Renewal		

Stem Cell Information

This Research Project Involves (check all that apply):	
<input type="checkbox"/> Human adult stem cells <input type="checkbox"/> Purely in vitro research using hSC lines <input type="checkbox"/> Derivatives of pluripotent cells or hESC <input type="checkbox"/> Somatic cell nuclear transfer (SCNT)	<input type="checkbox"/> Deriving or creating human pluripotent stem cell lines <input type="checkbox"/> Human pluripotent cells <input type="checkbox"/> Registered hESC lines from the NIH hESC Registry (http://stemcells.nih.gov/research/registry/)

Other Project Information

<p>Is this a clinical trial? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>	<p>If applicable, indicate which trial phase:</p>
<p>Funding Sources (check all that apply):</p> <p><input type="checkbox"/> Academic institution <input type="checkbox"/> California Institute for Regenerative Medicine</p> <p><input type="checkbox"/> Gifts <input type="checkbox"/> Private sponsorship</p> <p><input type="checkbox"/> Federal <input type="checkbox"/> Other:</p>	
<p>For Projects Involving the Use of Human Oocytes:</p> <p>If any oocytes were procured from outside California, please indicate the facility or institution they came from, including contact information, and how many from each site.</p> <p>1.</p> <p>2.</p>	
<p>List and Briefly Describe In-State, Out-of-State, and International Collaborations with Institutions/Organizations:</p> <p>1.</p> <p>2.</p> <p>3.</p>	
<p>Lay Summary/Abstract of Research Project (if applicable, include provenance of human stem cell lines and rationale for using human oocytes, embryos, or SCNT):</p>	
<p>Unanticipated Problems/Investigator Noncompliance Issues:</p>	
<p>Response of SCRO Committee to these Issues:</p>	