<table>
<thead>
<tr>
<th>Proposed Material</th>
<th>Information, Examples, etc.</th>
<th>What You Need to Provide (Provenance)</th>
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</thead>
</table>
| Registered Pluripotent Cell Lines | • NIH Registered  
• CIRM Registered  
• UK Stem Cell Bank  
• UK Human Fertilization & Embryology Authority  
• Canadian National Stem Cell Oversight Committee  
• Japanese Guidelines for Derivation & Utilization of Human Embryonic Stem Cells | Appropriate registry name and registry number for each line. Some lines may require approval documentation from the registry. |
| Non-Registered Pluripotent Cell Lines | Cell lines provided by a company, institution, or collaborator that do not appear on one of the above registration lists | 1. A sample Consent Document  
2. IRB (or equivalent) Approval Letter |
| Fresh Somatic Cells for the purpose of generating re-programmed cells | Usually in the form of tissue (e.g., punch biopsy, foreskin, residual tissue from surgical procedure, etc.) | 1. A sample Consent Document  
2. IRB (or equivalent) Approval Letter  
3. If provenance is not available, and the material is de-identified please complete the Petition to Designate a Stem Cell Line as Acceptably Derived form. |
| Archived Somatic Cells for the purpose of generating re-programmed cells | Usually obtained from a tissue bank | 1. If provenance is available:  
   • Sample Consent Document  
   • IRB (or equivalent) Approval Letter  
2. If provenance is not available, and the material is de-identified, please complete the Petition to Designate a Stem Cell Line as Acceptably Derived form. |
| Induced Pluripotent Cells (iPS) | Source Material:  
• Fresh Somatic Cells: see guidance above  
• Archived Somatic Cells: see guidance above | Please provide the appropriate documentation based on the source material. |
| Fetal Tissue derived stem cells | | Please follow the guidance above for Fresh Somatic Cells |
| Fetal Tissue | Aborted pregnancy materials, pathology samples | Provide the following statements as outlined in Code of California Regulations title 17 § 100085: Use of Fetal Tissue.  
• Statement signed by woman donating the material  
• Statement signed by attending physician  
• Statement signed by Principal Investigator |
| Embryos/Oocytes | | 1. Consent Document (either sample IRB-approved Consent or Consent document proposed for use)  
2. In both instances (Embryos and Oocytes) IRB Approval is required |