

THE DISTILLERY

This week in techniques

Disease modelsHuman embryonic stem cells (hESCs) generated from somatic cell nuclear transfer (SCNT) using postnatal somatic cellsSCNT could be useful for creating patient-matched hESCs for disease modeling and therapeutic applications. Previous efforts to reliably generate hESCs with nuclear transfer protocols have been limited to using nuclei from fetal as opposed to postnatal somatic cells. Fibroblasts from a 32-year-old female with type 1 diabetes or a newborn male were fused to enucleated donor human oocytes and activated with an oocyte activation protocol. A subset of the oocytes bearing the diploid genome of the donor fibroblasts developed into blastocysts, which were used to establish stable hESC lines. Next steps include comparing induced pluripotent stem (iPS) cell lines to nuclear transfer cell lines of the same genetic makeup to understand key differences between the two types of cells.Patent application fibroblasts developed into blastocysts, which were used to establish stable hESC lines. Next steps include comparing induced pluripotent stem (iPS) cell lines to nuclear transfer cell lines of the same genetic makeup to understand key differences between the two types of cells.Patent application fibroblasts developed into blastocysts, which were used to establish stable hESC lines. Next steps include comparing induced pluripotent stem (iPS) cell lines to nuclear transfer cell lines of the same genetic makeup to understand key differences between the two types of cells.Patent application fibroblast developed into blastocysts, which were used to establish stable hESC lines.Next steps include comparing induced pluripotent stem (iPS) tell into the same genetic makeup to understand key differences between the two types of cells.Patent application fibroblasts developed into blastocysts, which were used to est	Approach	Summary	Licensing status	Publication and contact information
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