

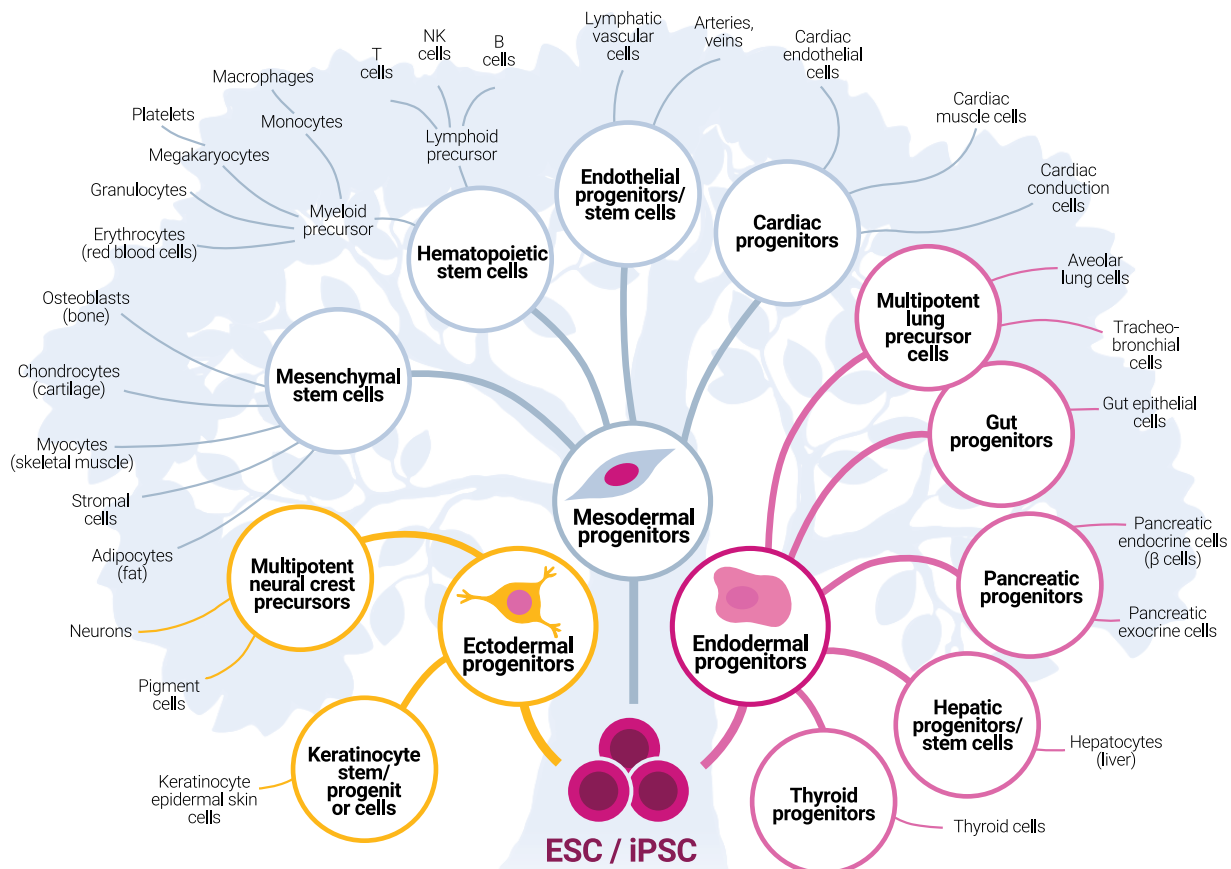
# Differentiation of embryonic and induced pluripotent stem cells


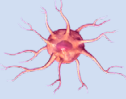
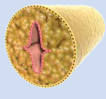

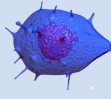
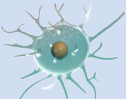
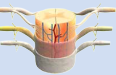

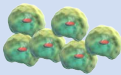





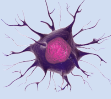




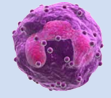






Pluripotent stem cells possess the capacity for indefinite self-renewal and the potential to differentiate into all adult cell types of the three primary germ layers. These features make pluripotent stem cells important candidates in the field of regenerative medicine due to their exceptional potential for replacement of lost, damaged, or diseased cells.

While embryonic stem cells (ESCs) are the most well-known type of pluripotent stem cell, the generation of ESCs from human embryos has resulted in a great deal of controversy. The introduction of induced pluripotent stem cells (iPSCs) has been a major breakthrough in the field of regenerative medicine, as it allows for the generation of pluripotent stem cells directly from

adult cells. Since iPSCs generated from adult cells can offer an unlimited supply of autologous cells, this circumvents the controversial use of human embryos while helping to minimize the risk of immune rejection.

The use of ESCs and iPSCs in cell therapies and research requires not only their procurement but also their *in vitro* differentiation into fully functioning, specialized cell types. Differentiation can be influenced and controlled through exposure to specific chemical and physical signals. Common chemical signals include cytokines, growth factors, and small molecules, to either activate or inhibit specific cellular pathways to achieve a desired cell fate.



Pathway	PeproTech cytokines and growth factors	Cell type									
<b>Self-renewal</b>	FGF-basic TGF-β1	 Embryonic stem cells (ESCs), induced pluripotent stem cells (iPSCs)									
<b>Neurogenesis</b>	BDNF CNTF EGF FGF-8a, b FGF-basic GDNF IGF-I β-NGF Noggin NT-3, -4 PDGF-AA, -AB, -BB, -CC Sonic hedgehog (Shh)	 Astrocytes	 Peripheral neurons	 Oligodendrocytes	 GABAergic neurons	 Dopaminergic neurons	 Motor neurons	 Glutamatergic neurons	 Neural crest		
<b>Adipogenesis</b>	BMP-2, -4, -7 FGF-basic TGF-β1	 White adipocytes	 Brown adipocytes								
<b>Hematopoiesis</b>	BMP-4 EPO IL-2, -3, -4, -6, -7, -11, -15 Flt3-ligand G-CSF GM-CSF SCF TPO VEGF <sub>165</sub> , VEGF <sub>121</sub>	 T cells	 NK cells	 B cells	 Dendritic cells	 Platelets	 Erythrocytes	 Eosinophils	 Basophils	 Neutrophils	 Macrophages
<b>Gastrointestinal</b>	Activin A EGF FGF-4, -10 Noggin R-spondin-1 Wnt-3a	 Intestinal tissue	 Stomach tissue								
<b>Cardiomyogenesis</b>	Activin A BMP-4 DKK-1 FGF-4 FGF-8a, b FGF-basic VEGF <sub>165</sub> , VEGF <sub>121</sub>	 Cardiac muscle	 Cardiomyocytes								
<b>Osteogenesis</b>	BMP-2, -4, -6 FGF-basic IGF-I IL-1α, -1β, -6, -7, -11, -15 LIF M-CSF PTHrP sRANK ligand SDF-1α (CXCL12) SDF-1β (CXCL12) TGF-β1	 Mature osteoblasts, osteocytes (bone)									

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