Key features

- Developed by iPSC expert manufacturers
- High cell survival rate and genomic stability
- Compatible for low density iPSC culture
- Suitable for 2D/3D, feeder-free/feeder cultures
- GMP-compatible composition for smooth transition from research to clinical manufacturing
- GMP-grade version and FDA DMF coming soon

Feature 3:

2D feeder free, 2D feeder culture, 3D floating culture

Feature 1: High cell survival and growth rate



High growth and survival rates are maintained even in feeder-free culture. Low density cell culture (2x10^3/ 6well) is also possible, suitable for cell thawing and cell cloning after gene editing.

Feature 2: Maintain high genomic stability and pluripotency

OCT3,						h.	10
	dente -	- Angela				No. Control of the second s	URACIA URACIA
	and and a	-	2002 2010	5115 (735)	10000	the LOSS Res J LOSS	a ato
NANC	12	11	10	9	1	7	6
	88	88	88		66	88	發發
	10		16		n		13
	ą	43	0 Ŕ	68		88	0.0
	Y	Х	22	21		20	19



Normal karyotype was maintained after 50 passages (n = 6 lines). Pluripotent marker expression was also comparably high.

enfree culture Feeder culture 3D-expa

Cells efficiently proliferate in 2D feeder-free and 2D feeder culture. Also suitable for large scale expansion in 3D suspension culture.



Reprogramming rine-derived cells

Fibroblasts, peripheral mononuclear cells, urine-derived cells are efficiently reprogrammed into iPSCs when cultured in Puel. Other iPSC culture media have low reprogramming efficiency.

Feature 5: Maintain high differentiation potential



Puel can effectively maintain pluripotency. After long-term culture in Puel, iPSCs can differentiate into neurons, myo-precursor cells, cardiomyocytes, hematopoietic cells, NK cells, T cells and epidermal cells.

Feature 4: High efficiency iPSC induction



