

# Certificate of Analysis

CELL LINE NAME	<b>BIHi268-A-18</b>	hPSCreg Link: <a href="https://hpscereg.eu/user/cellline/edit/BIHi268-A-18">https://hpscereg.eu/user/cellline/edit/BIHi268-A-18</a>	
DONOR GENDER/AGE:	<input type="checkbox"/> Male <input checked="" type="checkbox"/> Female <input type="checkbox"/> unknown Age:		
DISEASE PHENOTYPE / GENETIC VARIANT			
BANK	Master Bank, ID . , Passage 40 ...., Freezing Date: 14.04.2021		
FREEZING METHOD	Bambamker		
CULTURE PLATFORM	Wählen Sie ein Element aus.		
	Medium: E8	Coating: Geltrex	
REPROGRAMMING	Reprogramming Method Vector details (e.g. Kit, Pub, AddgeneNr):		
GENETIC MODIFICATION	<input checked="" type="checkbox"/> yes <input type="checkbox"/> no Targeting Vector: <input type="checkbox"/> TALEN, <input checked="" type="checkbox"/> CRISPR, <input type="checkbox"/> ZNF, Addgene: <input type="checkbox"/> Isogenic control/SNP <input checked="" type="checkbox"/> Gene knock-out <input type="checkbox"/> Transgene knock-in		
	Parental/isogenic cell line	BIHi043-A	
	Target gene/Transgene/Locus (het/hom?)	SORL1	
	Validation (e.g. PCR, sequencing)	Sequencing	

TEST DESCRIPTION	Test Method	Test Specification	Result
STERILITY (viral pathogens)	<input type="checkbox"/> Blood screening Donor <input type="checkbox"/> PCR (primary cells) <input checked="" type="checkbox"/> PCR (iPS clone/subclone)	HBV, HCV, HIV negative	not done
STERILITY (mycoplasma)	Test Method	No contamination detected	Pass
STERILITY (bacteria/ yeast/ fungi)	Culture for 7 days in antibiotic free medium	No contamination detected	Pass
REPROGRAMMING VECTORE CLEARANCE	<input type="checkbox"/> PCR <input type="checkbox"/> AB staining <input type="checkbox"/> Confirmed in parental line	Vector not present	not done
VIABILITY / MORPHOLOGY	Phase contrast microscopy of cells at 24, 48, and 72 hrs	Growth rate and confluency typical of hPSCs	Pass
UNDIFFERENTIATED PHENOTYPE	Markers for undifferentiated hPSCs <input type="checkbox"/> IF-Staining <input checked="" type="checkbox"/> FACS <input type="checkbox"/> other	Expression of at least three pluripotency markers detected	Pass
	<input type="checkbox"/> Pluritest	Pluripotency and Novelty Scores above threshold	not done
PLURIPOTENT DIFFERENTIATION POTENTIAL	3-germ layer differentiation: <input type="checkbox"/> spontaneous (e.g. EB formation)	Detection of markers for cells from the three germ layers	not done
	<input type="checkbox"/> directed differentiation	Successful differentiation to cells of all three germ layers	Result
	<input type="checkbox"/> Teratoma formation	Observation of tissues derived from the three germ layers	Result
KARYOTYPE	PerkinElmer KaryoLite BoBs™	Karyotype matches Donor	Result

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	Virtual karyotyping using Illumina OMNI-EXPRESS-8v1.6 Chip	No significant changes compared to the primary cells detected	Pass
	G-Banding	Karyotype matches Donor	Result
IDENTITY (STR ANALYSIS)	Promega GenePrint® 10 System	Identical to profile of primary cells	Result

date / signature: 16-02-2022

/Narasimha Telugu