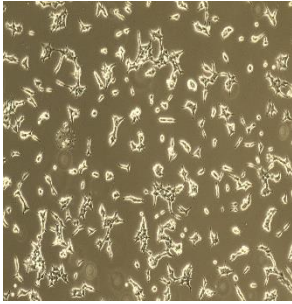
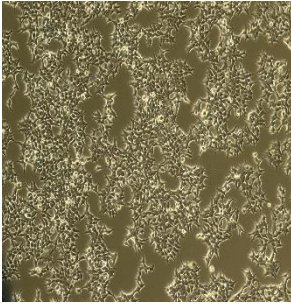


# HEK293T CRISPRn



Institut et hôpital neurologiques de Montréal  
Montreal Neurological Institute and Hospital

Product Information		
Cell Line	HEK293T PDI CRISPRn BA5	
Parental	HEK293T	
Product ID	HEK293T PDI CRISPRn BA5	
Product Batch	HEKn-190430	
Genotype	WT Doxycycline inducible Cas9	
Passage	P5	
Date of Production	2019-04-30	
Properties		
Volume	1 ml/vial	
Storage Conditions	Liquid Nitrogen	
Cell Number/ Vial	2.7 x 10 <sup>6</sup> cells/ml	
Viability	97%	
Quality Control		
Test	Test Method	Pass/Fail
Viability	Post thawing culture	Pass
Mycoplasma	MycoAlert™ Mycoplasma Detection Kit (Lonza)	Pass
Cell Line Characterization	Sanger Sequencing (DNA)	Pass
Morphology Images	10x objective      24h Post-Thaw	48h Post-Thaw
		
Growth Conditions		
Culture Media	Dulbecco's Modified Eagle's Medium (DMEM) supplemented with FBS 10%, L-glutamine 2mM, Penicillin-Streptomycin 100U/ml	
Passage Method	Trypsin	
Freezing Media	FBS with 10% DMSO	
Recommended Subculture	Cells are cultured as a monolayer at 37°C in a humidified atmosphere with 5% CO <sub>2</sub> . Cells should be passaged every 5-7 days. Split at 80-85% confluency, approximately 1:10-1:20.	
Cell Line Revival	Rapidly thaw cells in a 37°C water bath. Transfer contents into a tube containing 5 ml pre-warmed media. Centrifuge cells, remove supernatant wash cells with 10 ml PBS, centrifuge cells, remove PBS and seed into a 10 cm flask containing pre-warmed media.	