

SECTION 2: Cell Line Details

Cell Line Name:					
Cell Type: <i>(if iPSC or iPSC derived cell line, complete the following:)</i>					
• Reprogramming Method: <i>(please specify genes used)</i>					
• Somatic Starting Material:					
• Is the original cell line available?	Yes				No
• Are gene insertions present in the reprogrammed cell? <i>(List methods used to confirm absence or presence of gene insertion events)</i>	Yes			No	Unknown
Normal: <i>(if no, provide disease details)</i>	Yes				No
Donor Age:					
Donor Gender:	Male				Female
Donor Ethnicity:					
Donor Blood Type:	A	B	AB	O	Unknown
Passage Number:					
Population Doubling Time: <i>(In Recommended Culture Conditions – from Section 4)</i>					
Date Derived: <i>(if iPSC derived cell line, include iPSC derivation date and passage number at differentiation.)</i>					
Genetic Modifications : <i>(if yes, provide details including any elements protected as intellectual property.)</i>	Yes				No
Reporters: <i>(if yes, provide details including any elements protected as intellectual property.)</i>	Yes				No



SECTION 3 : Characterization Details

Test Name	Required Result	Testing Performed	
Mycoplasma detection (method)	Negative	Yes	No
Sterility assessment	Sterile	Yes	No
Karyotype		Yes	No
Identity (STR)		Yes	No
Human Virus Testing		Yes	No
MAP		Yes	No
Bovine pathogens		Yes	No
Porcine pathogens		Yes	No
In Vivo (in apparent Viruses)		Yes	No
28 Day In Vitro		Yes	No
Co-cultivation		Yes	No
ABO/Rh		Yes	No
HLA		Yes	No
FACS		Yes	No
Embryoid Body		Yes	No
Teratoma		Yes	No
Whole genome sequencing		Yes	No
Epigenic Analysis		Yes	No
Other tests		Yes	No
Please attach any relevant documentation of the characterization details.			

SECTION 4: Culture Conditions

Were the cells co-cultured? <i>(if yes, identify supporting cells.)</i>		Yes	No
Attachment Substrate/Matrix			
Culture Medium			
Passage Reagent			
Freezing Medium and Method			
Details of critical culture conditions <i>(please attach main culture protocols):</i>			



SECTION 5: Consent Information*If redacted consent form is available, please attach.*

Check all that apply:

For Research Purposes

For Commercial Purposes

For Therapeutic Purposes

Are there any restrictions on the use of the cell lines?

(if yes, provide details)

Yes

No

Is there any available medical information on the donor(s), including infection disease screening?

(if yes, provide details.)

Yes

No

Is there any available clinical, observational, or diagnostic information about the donor(s)?

Yes

No

SECTION 6: Related Publications

Are there any publications related to this line? If yes, please list:

SECTION 7: Declaration

By submitting this deposit for the NIH Center for Regenerative Medicine, I certify that the statements and Assurance herein are true, complete, and accurate to the best of my knowledge.

Signed on behalf of Host Institution

(Person responsible e.g., Scientific Director/Department Head)

Signed by Cell Line Provider:

(Person listed in Section 1)

x Date:

x Date:

Name and title of Signatory for Host Institution

Address of Host Institution:

(if different than address in Section 1)

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Tel: 301-402-6956 • Fax: 301-480-6367 • E-mail: nihrm@mail.nih.gov<http://commonfund.nih.gov/stemcells>

Identity and growth

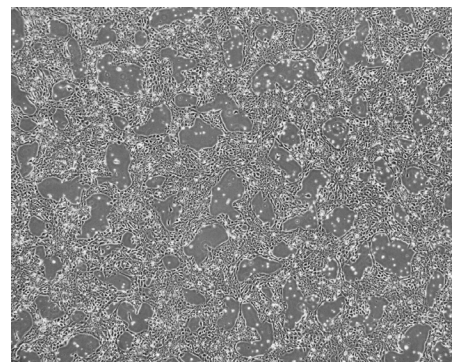
Date - 05/15/2013 - modified 01/28/2015

Cell line ID -NCRM-5
 Subline - N/A
 Cell line submitter - NIH Center for Regenerative Medicine
 RUID SNP panel - available upon request
 Source cell type - CD34⁺ cord blood
 Reprogramming method - episomal plasmid
 Growth conditions - mTeSR and Matrigel
 Passage method - Dispase (SOP SC14)
 Pre-freeze mycoplasma testing - pass

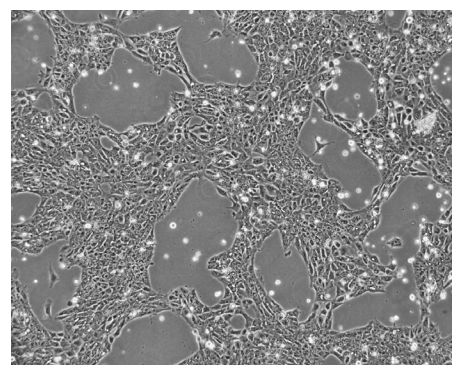
Cryopreservation

Cryopreservation date - 09/06/2012
 Cryopreservation method - Accutase and mFreSR (SOP SC13)
 Passage number - 12
 Post thaw mycoplasma testing - negative

Post thaw morphology - pass



4x magnification 3 days after thaw



10x magnification 3 days after thaw

Freeze ID	R127303685	R127452457
Recommended thaw density per 9.6cm ² (1 well of a 6 well plate)	1e6	2e6
At least 10 colonies after 3 days (+/-)	+	+
At lease 50% pluripotent based on morphology (+/-)	+	+

Post thaw FACS analysis - 98.8% Oct4/Tra-1-60 double positive

