

TITLE: KROMATID CHROMOSOME ANALYSIS REPORT

I. ASSAY INFORMATION

Project Quote #	Q200401
Specimen Type	iPSC
Body Site	N/A
Sample ID	100 Lead Het (AB) (S005292)
	Male
Cell Line	
Gender	
Passage number	N/A
(or N/A)	
Study Objective	The purpose of this study is to characterize iPS cells grown <i>in vitro</i> , designated for cytogenetic analysis.

II. CELL MAINTENANCE

Culture vessel	N/A
Media	N/A
Density (estimated)	N/A
Culture atmosphere	N/A
Culture maintenance	N/A

DocuSign Env	/elope ID: C3	1530E5-73	4A-4C8C	-8BD1-090	C7D483FA87
kr	nm	2			Docu

gn Envelope ID: C31530E5-734A-4C8	7D483FA87			
romali	Document Code:	FORM-0068A	Document Type:	FORM
	Revision:	3.0	Date Effective:	08-27-21

Culture Maintenance Process Description	N/A
	Analyst Initial/Date: N/A

III. **CULTURE HARVEST**

Culture Harvest Process Description	N/A
	Analyst Initial/Date: N/A

Material	Usage information	
Harvest materials (trypsin, EDTA, etc.)	Type: N/a LN/ Exp. Date: N/a	
Colcemid	LN/ Exp. Date: N/a Concentration: 0.1 µg/ml (10 µl/ml) Incubation time: N/a	
Hypotonic	LN/ Exp. Date: N/a Solution: N/a Incubation time:	
Fixative	Prepared Fresh, day-of-use	



IV. STAINING

Solution Type	Lot#	Exp. Date	Solution Type	Lot#	Exp. Date
Isoton II Diluent	4710610	07/12/22	Wright Stain	210317	03/17/22
Pancreatin	SLCD9444	01/11/23	Gurr Buffer	210818	09/18/21
FBS	19J079	11/4/22	Permount	210201-01	02/1/23

Description	A sample of fixed cells labeled 100 Lead Het (AB) (KromaTiD Sample ID S005292) was received at KromaTiD on 08-26-21. The fixed cells were washed twice with fixative (prepared fresh day-of-use) and the O.D. was adjusted to 0.0250. Drops of the final cell suspension were placed on clean slides and aged for 60 minutes at 90°C. Slides were digested in a pancreatin solution with Isoton II diluent. The enzymatic reaction was then stopped by rinsing with FBS, followed by application of a stain solution (3:1 Wright/Gurr buffer) which was poured on the slides so that it covered the entire surface. After staining for up to 1 minute, slides were washed with de-ionized water for 1-5 seconds and air dried. The mounting medium Permount was applied to the slides, a coverslip was placed on the slide and the slides were scanned on the microscope.
	Analyst Initial/Date: MV 08-30-21 through 09-15-21

TEST DESCRIPTION:

G-banding with trypsin treatment and Giemsa stain (GTG-banding) is used in cytogenetics to produce a visible karyotype by staining metaphase chromosomes. This technique allows each chromosome to be distinguished by its characteristic banding pattern. G-banding is useful in assessing structural abnormalities in individual chromosomes, as well as extra or missing chromosomes within a cell. Industry-standard protocols for scoring and describing results were used (ISCN 2016: An International System for Human Cytogenomic Nomenclature).

KromaliD

V. **RESULTS**

Cells Counted	20	Total Karyograms	2
Cells Analyzed	20	Average Band Resolution	350
Image File Location	1	Jax Gbanding_S005292	

5.1 Chromosome Count per 20 Metaphases

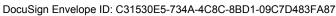
Of the 20 cells counted, 16 contained 46 chromosomes (80%). *Cells containing greater than 57 chromosomes are recorded as polyploid.* The polyploid frequency was 0%, based on the metaphases counted.

5.2 CHROMOSOME ABERRATION DATA

The chromosome aberration data via G-band for the 20 metaphases examined is summarized in attached case report cell list. 0 chromosome aberrations were found in the 20 cells analyzed with 0% of the cells aberrant.

*Note: Cells with an euploidy gain/loss were found to be non-clonal, and therefore not included in the aberration data below.

	Tech Summary	Additional Comments
Karyotype	46,XY[20]	Normal Male Karyotype
Cells Analyzed	20	
Normal Cells	20	Random loss/gain cells normalized
Abnormal Cells	0	
Aberration	N/A	
Туре		
Aberration %	0%	





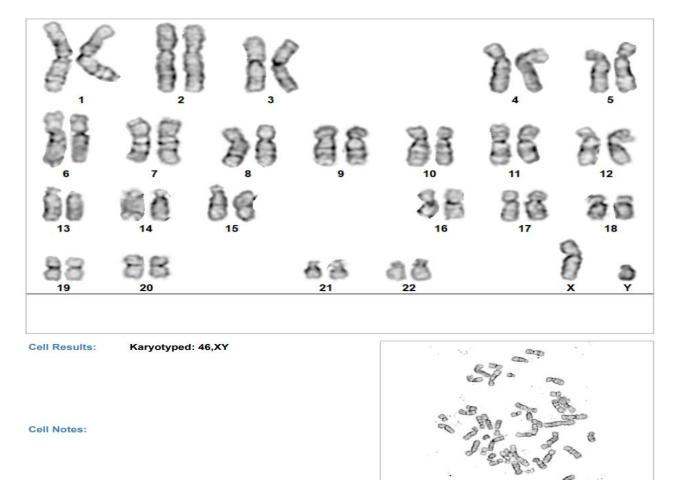
-09071	J483FA87			
	Document Code:	FORM-0068A	Document Type:	FORM
	Revision:	3.0	Date Effective:	08-27-21

5.3. INTERPRETATION/ SIGNIFICANCE:

G-banded chromosome analysis of metaphase cells designated as 100 Lead Het (BB) (KromaTiD Sample ID S005292) show a normal male karyotype 46,XY[20].

The other abnormalities/aberrations detected were non-clonal, and were designated as low level mosaicism or random gain/loss.

5.4 REPRESENTATIVE IMAGES:



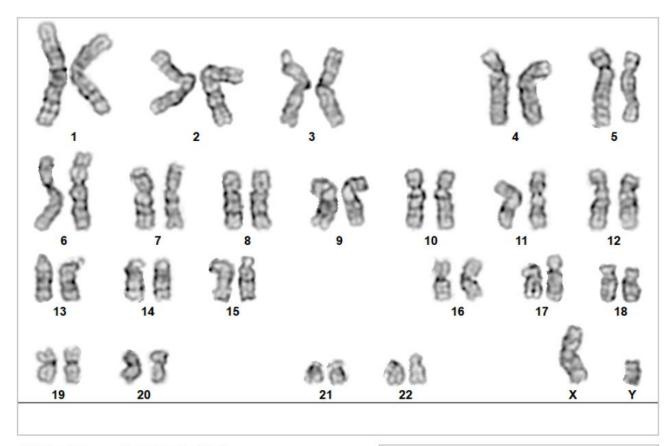
Label - Slide/Cell: S005292 - 1/13

X,Y: 10.5 , 20.0

Report Date: Tuesday, September 21, 2021



Document Code:	FORM-0068A	Document Type:	FORM
ъ	2.0		09 07 01
Revision:	3.0	Date Effective:	08-27-21



Cell Results:

Karyotyped: 46,XY

Cell Notes:



Label - Slide/Cell: S005292 - 1/40

X,Y: 10.0 , 34.9

Report Date: Tuesday, September 21, 2021

DocuSign Envelope ID: C31530E5-734A-4C8C-8BD1-09C7D483FA87



J483FA87			
Document Code:	FORM-0068A	Document Type:	FORM
Revision:	3.0	Date Effective:	08-27-21

Limitations: This assay allows for microscopic visualization of numerical and structural chromosome abnormalities. The size of structural abnormality that can be detected is >3-10Mb, dependent upon the G-band resolution obtained from this specimen. For the purposes of this report, band level is defined as the number of G-bands per haploid genome. Detection of heterogeneity of clonal cell populations in this specimen is limited by the number of metaphase cells analyzed, documented above as "number of cells counted". Results are for Research Use Only, and should not be used for clinical purposes.

Completed By/Date:

e: Michael Vernich Michael Vernich Cytogenetics Associate

Reviewed By/ Date:

DocuSigned by 9/24/2021 Ivan furs 78884964FB9848A... Ivan Perez Cytogenetics Technologist III

Approved By/Date:

DocuSianed by: Marssa Jalugues/24/2021

5FAF272CE1894D0... Marissa Rodrigues QA Manager

DocuSign Envelope ID: C31530E5-734A-4C8C-8BD1-09C7D483FA87 Case: Jax G-banding_S005292

LIMS ID:

S005292

LIMS Sample Name: 100 Lead Het (AB)

# Slide Cell Coordinates Results Analysis State By							
"	Cinat	0011	Coordinatoo		State	Cluic Dy	
Slide	e: Name: 1	Label: S	005292	'			
1	1	1	13.23 X 8.03	Karyotyped: 45,XY, −16	Karyotyped	mvernich	
2	1	4	15.68 X 12.49	Karyotyped: 46,XY	Karyotyped	mvernich	
3	1	5	9.08 X 16.28	Karyotyped: 46,XY	Karyotyped	mvernich	
4	1	7	9.42 X 17.63	Karyotyped: 46,XY	Karyotyped	iperez	
5	1	8	14.05 X 17.60	Karyotyped: 46,XY	Karyotyped	mvernich	
6	1	9	10.48 X 18.73	Karyotyped: 46,XY	Karyotyped	mvernich	
7	1	13	10.52 X 20.00	Karyotyped: 46,XY	Karyotyped	mvernich	
8	1	17	11.19 X 20.70	Karyotyped: 46,XY	Karyotyped	mvernich	
9	1	21	9.02 X 22.12	Karyotyped: 46,XY	Karyotyped	mvernich	
10	1	35	11.15 X 31.01	Karyotyped: 44,XY, -4, -20	Karyotyped	mvernich	
11	1	40	10.00 X 34.94	Karyotyped: 46,XY	Karyotyped	mvernich	
12	1	41	7.75 X 35.18	Karyotyped: 46,XY	Karyotyped	mvernich	
13	1	45	16.22 X 41.02	Karyotyped: 46,XY	Karyotyped	mvernich	
Slide	: Name: 2	2 Label: S	005292				
14	2	10	16.42 X 13.94	Karyotyped: 46,XY	Karyotyped	mvernich	
15	2	26	11.19 X 31.97	Karyotyped: 46,XY	Karyotyped	mvernich	
Slide	: Name: 3	B Label: S	005292				
16	3	4	10.77 X 18.40	Karyotyped: 45,XY, -12	Karyotyped	mvernich	
17	3	19	9.81 X 29.39	Karyotyped: 46,XY	Karyotyped	mvernich	
18	3	38	10.87 X 33.78	Karyotyped: 45,XY, -3	Karyotyped	mvernich	
19	3	47	4.09 X 36.09	Karyotyped: 46,XY	Karyotyped	mvernich	
20	3	49	17.40 X 37.26	Karyotyped: 46,XY	Karyotyped	mvernich	