

TITLE: KROMATID CHROMOSOME ANALYSIS REPORT

I. ASSAY INFORMATION

Project Quote #	Q200401
Specimen Type	iPSCs
Body Site	N/A
Sample ID	KOLF2.1J-P11 (S008110)
Cell Line Gender	Male
Passage number (or N/A)	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

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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			Solution Type	Lot#	Exp. Date
Isoton II Diluent	4710610	07/12/22	Wright Stain	210817-Wright	08/17/22
Pancreatin	SLCD9444	01/11/23	Gurr Buffer	220329-Gurr	04/29/22
FBS	20J481	01/06/24	Permount	210201-01	02/1/23

A sample of fixed cells KOLF2.1J-P11 (KromaTiD Sample ID S008110) was received at KromaTiD on 4-13-22. The fixed cells were washed twice with fixative (prepared fresh day-of-use) and the O.D. was adjusted. 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 4/26/22

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).

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V. RESULTS

Cells Counted	60	Total Karyograms	2
Cells Analyzed	60	Average Band Resolution	425
Image File Location	l	Jax Gbanding_S008110	

5.1 CHROMOSOME COUNT PER 20 METAPHASES

Of the 60 cells counted, 47 contained 46 chromosomes (78.33%). 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 60 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 aneuploidy gain/loss were found to be non-clonal, and therefore not included in the aberration data below.

	Tech Summary	Additional Comments			
		Normal Male Karyotype			
Cells Analyzed	60				
Normal Cells	60	Random loss/gain cells normalized			
Abnormal Cells	0				
Aberration	N/A				
Туре					
Aberration %	0%				





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5.3. INTERPRETATION/ SIGNIFICANCE:

G-banded chromosome analysis of metaphase cells designated KOLF2.1J-P11 (KromaTiD Sample ID S008110) shows a normal male karyotype 46,XY[60].

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

X,Y: 15.8, 42.9

5.4 REPRESENTATIVE IMAGES: 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 X Y Cell Results: Karyotyped: 46,XY Cell Notes:

Label - Slide/Cell: S008110 - 1/96

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	13	14 14	15		3 6	2	18		
	19	88 20		66	22	×	1 Y		
	Cell Results: Cell Notes:	Karyotyped:	46,XY				AD AD		
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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:

Billiousigned by: Michael Vernich Michael Vernich Cytogenetics Supervisor

4/30/2022

Approved By/Date:

Greg Husar Gregory Husar Operations Manager 4/30/2022

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Table 1: Chromosome Analysis for sample KOLF2.1J-P11 (S008110). 60 cells were analyzed.

# Slide: I	Slide Name: 1 La	Cell	Coordinates	Results	Analysis State	State By			
1	1	10	5.93 X 15.17	Karyotyped: 46,XY	Karyotyped	mvernich			
2	1	28	8.24 X 23.17	Karyotyped: 45,XY, -15	Karyotyped	mvernich			
3	1	70	6.23 X 37.46	Karyotyped: 46,XY	Karyotyped	mvernich			
4	1	94	12.03 X 42.22	Karyotyped: 46,XY	Karyotyped	mvernich			
5	1	96	15.77 X 42.88	Karyotyped: 46,XY	Karyotyped	mvernich			
6	1	99	12.03 X 43.33	Karyotyped: 46,XY	Karyotyped	mvernich			
7	1	103	7.56 X 43.52	Karyotyped: 46,XY	Karyotyped	skeables			
Slide: Name: 2 Label: S008110									
8	2	28	4.36 X 24.42	Karyotyped: 43,XY, −2, −7, −9	Karyotyped	mvernich			
9	2	59	11.59 X 30.95	Karyotyped: 46,XY	Karyotyped	mvernich			
10	2	66	12.83 X 32.97	Karyotyped: 46,XY	Karyotyped	skeables			
11	2	76	14.51 X 35.72	Karyotyped: 45,XY, −22	Karyotyped	mvernich			
Slide: I	Name: 3 La	abel: S00	8110						
12	3	12	13.62 X 22.44	Karyotyped: 46,XY	Karyotyped	mvernich			
13	3	18	14.27 X 26.86	Karyotyped: 46,XY	Karyotyped	mvernich			
14	3	30	13.03 X 32.86	Karyotyped: 46,XY	Karyotyped	mvernich			
15	3	35	4.63 X 35.54	Karyotyped: 44,XY, −18, −19	Karyotyped	mvernich			
16	3	41	13.55 X 39.95	Karyotyped: 46,XY	Karyotyped	mvernich			
Slide: I	Slide: Name: 4 Label: S008110								
17	4	2	6.79 X 13.10	Karyotyped: 46,XY	Karyotyped	mvernich			
18	4	7	14.26 X 20.26	Karyotyped: 42,XY, −8, −9, −13, −19	Karyotyped	mvernich			
19	4	13	11.75 X 24.41	Karyotyped: 46,XY	Karyotyped	mvernich			
20	4	22	9.67 X 34.05	Karyotyped: 47,XY, +17	Karyotyped	mvernich			

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	21	4	23	11.24 X	34.37	Karyotype −21	ed: 44,XY, −8,	Karyotyped	mvernich	
	22	4	25	12.38 X	34.11	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	23	4	30	16.00 X	41.68	Karyotype	ed: 46,XY	Karyotyped	mvernich	
Slide: Name: 5 Label: S008110										
	24	5	10	6.77 X 1	7.30	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	25	5	27	7.24 X 2	24.59	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	26	5	34	6.49 X 3	80.06	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	27	5	52	6.00 X 3	39.08	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	28	5	57	6.24 X 4	13.03	Karyotype −22	ed: 44,XY, −12,	Karyotyped	mvernich	
	29	5	60	17.52 X	42.64	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	Slide: N	lame: 6 Lat	oel: S008	110						
	30	6	5	13.85 X	14.11	Karyotype	ed: 45,XY, −12	Karyotyped	mvernich	
	31	6	19	9.78 X 2	23.05	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	32	6	34	7.40 X 2	27.87	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	33	6	46	8.74 X 3	37.29	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	34	6	52	10.16 X	42.79	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	Slide: N	lame: 7 Lat	oel: S008	110						
	35	7	1	14.54 X	9.83	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	36	7	4	16.78 X	10.75	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	37	7	24	15.70 X	24.30	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	38	7	32	16.28 X	28.12	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	39	7	35	14.66 X	31.31	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	40	7	40	11.23 X	33.82	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	41	7	42	13.93 X	34.55	Karyotype	ed: 46,XY	Karyotyped	mvernich	
	42	7	55	5.84 X 4	12.18	Karyotype −16	ed: 44,XY, −2,	Karyotyped	mvernich	

Slide: Name: 8 Label: S008110

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43	8	3	14.09 X	10.20	Karyotyped: 46,XY		Karyotyped mvernich		vernich
44	8	6	16.19 X	13.21	Karyotype −1, −10, -	ed: 42,X, -Y, -13	Karyotyped	m	vernich
45	8	13	9.62 X 1	5.65	Karyotype	ed: 46,XY	Karyotyped	m	vernich
46	8	22	17.05 X	24.47	Karyotype	ed: 46,XY	Karyotyped	m	vernich
47	8	23	16.39 X	23.61	Karyotype −18, −21	ed: 43,XY, −17,	Karyotyped	m	vernich
48	8	28	4.18 X 2	25.82	Karyotype	ed: 46,XY	Karyotyped	m	vernich
49	8	34	15.75 X	30.09	Karyotype	ed: 46,XY	Karyotyped	m	vernich
50	8	39	5.89 X 3	32.48	Karyotype	ed: 46,XY	Karyotyped	m	vernich
51	8	42	7.62 X 3	34.34	Karyotype	ed: 46,XY	Karyotyped	m	vernich
52	8	45	9.93 X 3	33.97	Karyotype	ed: 45,X, -Y	Karyotyped	m	vernich
53	8	67	8.05 X 4	3.65	Karyotype	ed: 46,XY	Karyotyped	m	vernich
Slide: N	Name: 9 Lal	bel: S008	3110						
54	9	41	15.76 X	35.76	Karyotype	ed: 46,XY	Karyotyped	m	vernich
Slide: I	Name: 10 La	abel: S00	8110						
55	10	21	11.92 X	34.51	Karyotype	ed: 46,XY	Karyotyped	m	vernich
Slide: Name: 11 Label: S008110									
56	11	12	15.24 X	18.64	Karyotype	ed: 46,XY	Karyotyped	m	vernich
57	11	33	17.22 X	29.02	Karyotype	ed: 46,XY	Karyotyped	m	vernich
58	11	43	12.23 X	34.39	Karyotype	ed: 46,XY	Karyotyped	m	vernich
59	11	44	12.32 X	34.50	Karyotype	ed: 46,XY	Karyotyped	m	vernich
60	11	50	8.09 X 3	88.47	Karyotype	ed: 46,XY	Karyotyped	m	vernich