



Certificate of Analysis - Amended Distribution Lot

Product Description	H7 (WA07) WiCell Distribution Lot
Cell Line Provider	WiCell Research Institute (Madison, WI, USA)
Distribution Lot Number	H7-WCDL-4 (lot 4)
Date Vialled	20 July 2007
Passage Number	27
Culture Method	SOP-CC-030B, SOP-CC-020B
Cryopreservation Method	SOP-CC-034B

The following testing specifications have been met for the specified product lot:

Test Description	Test Method	Test Specification	Result
Post-Thaw Viable Cell Recovery	SOP-CH-305A	Viable cells recovered	Pass
Identity by STR	SOP-CH-302B	Positive identity	Pass
Sterility - Direct transfer method	SOP-CH-304A	No contamination detected	Pass
Mycoplasma	SOP-SS-002A	No contamination detected	Pass
Karyotype by G-banding	SOP-CH-003B	Normal karyotype	Pass

Electronic versions of this certificate of analysis (CoA) complete with electronic copies of individual reports, results, and procedures are available on our website, www.wicell.org. There are also archived CoAs for past cell lots.

Please visit the technical service portion of the website for assistance with your human ES Cells. The knowledgeable technical support staff can assist with embryonic stem cell culture concerns, training, and any other customer service concerns you may encounter.

Amendment(s):

Reason for Amendment	Date
CoA updated to include copyright information, electronic signature, and WiCell logo. Links updated.	See signature
Original CoA	07-Oct-2007

Date of Lot Release	Quality Assurance Approval
07-October-2007	<p style="text-align: right;">1/3/2014</p> <p>X AMC</p> <p>AMC Quality Assurance Signed by [REDACTED]</p>

Short Tandem Repeat Analysis*

Sample Report: H7p33

UW HLA#: 56968

Sample Date: 08/26/07

Lab Received 08/27/07

Requestor: WiCell Research Institute

Test Date: 08/29/07

File Name: 070831

Report Date: 09/04/07

Sample Name: (label on tube) H7p33**Description:** WI Cell Cytogenetics provided
genomic DNA of H7p33 (DNA 044)

Locus	Repeat #	STR Genotype
D16S539	5, 8-15	12,13
D7S820	6-14	10,11
D13S317	7-15	11,12
D5S818	7-15	11,13
CSF1PO	6-15	12,12
TPOX	6-13	8,11
Amelogenin	NA	X,X
TH01	5-11	6,6
vWA	11, 13-21	14,15

92ug/mL 260/280=1.80

Comments: Based on the H7p33 DNA submitted by WI Cell dated 08/26/07 and received on 08/27/07, this sample (UW HLA# 56968) matches exactly the STR profile of the human stem cell line H7 comprising 14 allelic polymorphisms across the 8 STR loci analyzed. No STR polymorphisms other than those corresponding to the human H7 stem cell line were detected and the concentration of DNA required to achieve an acceptable STR genotype (signal/ noise) was equivalent to that required for the standard procedure (~1 ng/amplification reaction) from human genomic DNA. This result suggest that the DNA sample submitted corresponds to the H7 stem cell line and was not contaminated with any other human stem cells or a significant amount of mouse feeder layer cells. Sensitivity limits for detection of STR polymorphisms unique to either this or other human stem cell lines is ~5%. A preliminary copy of this report was issued via electronic mail to JJ of WI Cell Research Institute on Thursday, September 6, 2007.

* Testing to assess engraftment following bone marrow transplantation was accomplished by analysis of human genetic polymorphisms at STR loci. This methodology has not yet been approved by the FDA and is for investigational use only.



University of Wisconsin
Hospital and Clinics
HLA/Molecular Diagnostics Laboratory

Histocompatibility/Molecular Diagnostics Laboratory
Madison, WI 53792-2472

HLA/Molecular Diagnostics Laboratory

* Testing to assess engraftment following bone marrow transplantation was accomplished by analysis of human genetic polymorphisms at STR loci. This methodology has not yet been approved by the FDA and is for investigational use only.

File: Final STR Report

Report Number
753370
Page 1 of 1

WiCell Research Institute
Madison, WI 53719
Attn: - -

August 24, 2007
P.O. #: RP1370

STERILITY TEST REPORT

Sample Information: Human embryonic stem cell line H7 on mouse feeder layer, H7

Date Received: July 25, 2007
Date in Test: August 09, 2007
Date Completed: August 23, 2007

Test Information: Test Code: 30744
Immersion, USP / 21 CFR 610.12
Procedure #: BS210WCR.02

TEST PARAMETERS	PRODUCT	
Approximate Volume Tested	0.45 mL	0.45 mL
Number Tested	1	1
Type of Media	SCD	FTM
Media Volume	200 mL	200 mL
Incubation Period	14 Days	14 Days
Incubation Temperature	20 °C to 25 °C	30 °C to 35 °C
RESULTS	1 NEGATIVE	1 NEGATIVE

QA Reviewed: S. Salyer 08-27-07

Reviewed: George P. Brown 08/24/07

Testing conducted in accordance with current Good Manufacturing Practices.



APPENDIX I

Document #: DCF3008A
Edition #: 06
Effective date: 9/17/2003
Title: DNA FLUOROCHROME ASSAY RESULTS

DNA-FLUROCHROME ASSAY RESULTS
Procedures 3008, 3009, 3011

Sample ID # 49713 M-250 Date Rec'd: 08/29/2007 P.O. # RP1452

Indicator Cells Inoculated: Date/Initials: 8/30/07 / KG

Fixation: Date/Initials: 9/3/07 / KG

Staining: Date/Initials: 9/9/07 / JA

TEST/CONTROL ARTICLE:

H7 p33+ SOP-CC-023A.H7 p33

LOT# NA

Distribution Lab
WiCell Research Institute

Phone:

Fax #:

DNA FLUROCHROME ASSAY RESULTS:

NEGATIVE: A reaction with staining limited to the nuclear region, which indicates no mycoplasmal contamination.

POSITIVE: A significant amount of extranuclear staining which strongly suggests mycoplasmal contamination.

INCONCLUSIVE:
 A significant amount of extranuclear staining consistent with low - level mycoplasmal contamination or nuclear degeneration.

 A significant amount of extranuclear staining consistent with bacterial, fungal or other microbial contaminant or viral CPE. Morphology not consistent for mycoplasmal contamination.

COMMENTS: _____

Date: 9/4/07 Results Read by: JA Date of Review: 9/4/07 Reviewed by: U



APPENDIX IV

Document#: DCF3013D
Edition#: 10
Effective Date: 07/15/2003
Title: **M-250 FINAL REPORT SHEET**

M-250 FINAL REPORT

Direct Specimen Culture
Procedure 3008, 3011, 3013

TO: **Distribution Lab
WiCell Research Institute**

Madison, WI 53719
PHONE#:

FAX#:

BTL SAMPLE ID#: **49713** P.O.#: **RP1452** DATE REC'D: **08/29/2007**

TEST/CONTROL ARTICLE:

H7 p33 + SOP-CC-023A.H7 p33

LOT#: **NA**

DIRECT CULTURE SET-UP (DAY 0)

DATE: **08/29/2007**

INDICATOR CELL LINE (VERO)

SEE DNA FLUOROCHROME RECORD SHEET

				DATE
THIOGLYCOLLATE BROTH	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>
BROTH-FORTIFIED COMMERCIAL <u>0.5</u> mL SAMPLE	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>
<u>6.0</u> mL BROTH	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>
BROTH-MODIFIED HAYFLICK <u>0.5</u> mL SAMPLE	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>
<u>6.0</u> mL BROTH	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>
BROTH-HEART INFUSION <u>0.5</u> mL SAMPLE	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>
<u>6.0</u> mL BROTH	DAY 7	+	⊖	<u>09/05/2007</u>
	DAY 28	+	⊖	<u>09/26/2007</u>

(See Reverse)

Document#: DCF3013D
 Edition#: 10
 Effective Date: 07/15/2003
 Title: M-250 FINAL REPORT SHEET


SAMPLE ID#:		AEROBIC	MICROAEROPHILIC	DATE
49713 AGAR PLATES-FORTIFIED COMMERCIAL	DAY 7	+ ⊖	+ ⊖	<u>09/05/2007</u>
	DAY 14	+ ⊖	+ ⊖	<u>09/12/2007</u>
	DAY 21	+ ⊖	+ ⊖	<u>09/19/2007</u>
AGAR PLATES-MODIFIED HAYFLICK	DAY 7	+ ⊖	+ ⊖	<u>09/05/2007</u>
	DAY 14	+ ⊖	+ ⊖	<u>09/12/2007</u>
	DAY 21	+ ⊖	+ ⊖	<u>09/19/2007</u>
AGAR PLATES-HEART INFUSION	DAY 7	+ ⊖	+ ⊖	<u>09/05/2007</u>
	DAY 14	+ ⊖	+ ⊖	<u>09/12/2007</u>
	DAY 21	+ ⊖	+ ⊖	<u>09/19/2007</u>

BROTH SUBCULTURES (DAY 7) DATE: 09/05/2007

AGAR PLATES-FORTIFIED COMMERCIAL	DAY 7	+ ⊖	+ ⊖	<u>09/12/2007</u>
	DAY 14	+ ⊖	+ ⊖	<u>09/19/2007</u>
	DAY 21	+ ⊖	+ ⊖	<u>09/26/2007</u>
AGAR PLATES-MODIFIED HAYFLICK	DAY 7	+ ⊖	+ ⊖	<u>09/12/2007</u>
	DAY 14	+ ⊖	+ ⊖	<u>09/19/2007</u>
	DAY 21	+ ⊖	+ ⊖	<u>09/26/2007</u>
AGAR PLATES-HEART INFUSION	DAY 7	+ ⊖	+ ⊖	<u>09/12/2007</u>
	DAY 14	+ ⊖	+ ⊖	<u>09/19/2007</u>
	DAY 21	+ ⊖	+ ⊖	<u>09/26/2007</u>

RESULTS: No detectable mycoplasmal contamination

9/26/07
Date


 Director, Technical Services
 Ph.D.

M-250 Procedural Summary: The objective of this test is to ascertain whether or not detectable mycoplasmas are present in an in vitro cell culture sample, be it a primary culture, hybridoma, master seed stock or cell line. This procedure combines an indirect DNA staining approach to detect non-cultivable mycoplasmas with a direct culture methodology utilizing three different mycoplasma media formulations. The indirect approach involves the inoculation of the sample into a mycoplasma-free VERO (ATCC) indicator cell line and performing a DNA fluorochrome assay after 72-120 hours of incubation. The direct culture aspect of the test utilizes three different mycoplasma media including both broth and agar formulations. The sample is inoculated into each of the 3 broth formulations and also onto duplicate plates (0.1 mL/plate) for each of the 3 agar formulations. Subculture from broth to fresh agar plates is carried out after 7 days incubation. Agar plates are incubated aerobically and microaerophilically in order to detect any colony forming units morphologically indicative of mycoplasma contamination. Issuance of the final report with signature of the Scientific Director/Study Director signifies that the required controls were performed concurrently with the test sample(s) as detailed in the referenced SOPs and that all test conditions have been found to meet the required acceptance criteria for a valid test, including the appropriate results for the positive and negative controls.

Report Date: October 09, 2007

Case Details:

Cell Line: H7

Passage #: 33

Date Completed: 8/30/2007

Cell Line Gender: female

Investigator: JL

Specimen: hESC on MEF feeder

Date of Sample: 8/26/2007

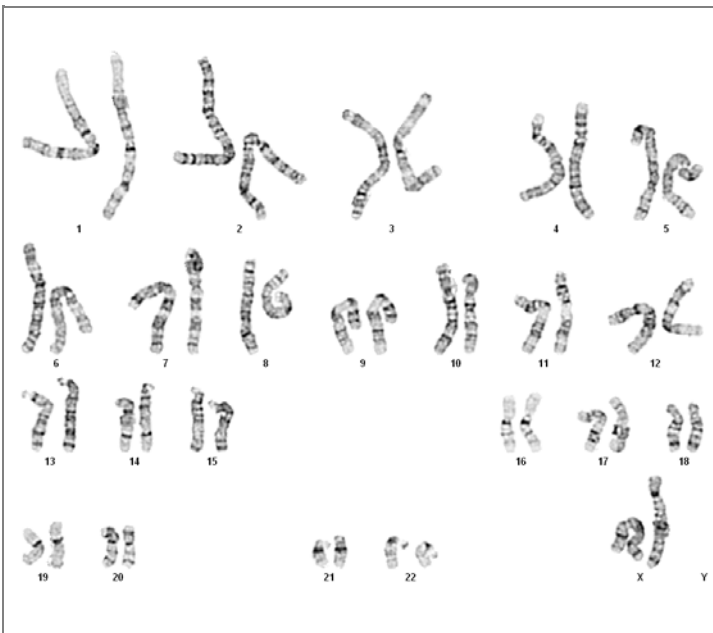
Tests, Reason for: Distribution lot testing

Results: 46,XX

Completed by CS, CLSp(CG), on 8/28/2007

Reviewed and interpreted by KDM, PhD, FACMG, on 8/30/2007

Interpretation: No abnormalities were detected at the stated level of resolution.



Cell: S01-05

Slide: C

Slide Type: Karyotyping

Cell Results: Karyotype: 46,XX

of Cells Counted: 20

of Cells Karyotyped: 5

of Cells Analyzed: 8

Band Level: 450-525

Results Transmitted by Fax / Email / Post
Sent By: _____

Date: _____
Sent To: _____