

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Datasheet for WM983B-01-0001

WM983B Viable Cells

Overview

Description:	WM983B Viable Cells - WM983B-01-0001
Item No.:	WM983B-01-0001
Size:	1 million cells
Applications:	Cellular Assay, IF, Other, WB
Origin:	Human

Product Details

Background: WM983B is a human metastatic melanoma cell line that was established from an inguinal node

metastatic site in a 54-year-old male patient. The subject displayed malignant melanoma of polypoid type, level IV with a tumor thickness of 2.5 cm. MW983B line contains the BRAF V600E mutation that causes increased signaling via the extracellular signal-regulated MAPK/ERK kinase pathways to enhance proliferation. WM983B is wild type for PTEN, N-ras, c-KIT, and CDK4. This cell line was derived from the same patient as the cell lines WM983A, WM983B BR, and

WM983C. WM983B cells produce xenograft tumors when injected into immunocompromised

Melanoma, patient derived tumor, tumor models, skin cancer, xenograft

Species of Origin: Human

Target Details

Synonyms:

Purity/Specificity: Cells are sterile, validated by short tandem repeat profiling, and are tested as negative for

mycoplasma. It is recommended that cell lines are tested for mycoplasma contamination and short tandem repeat (STR) profiling every 10 passages or each time a frozen seed stock is made.

See cell culture protocol for additional details.

Relevant Links: • Cell Line EULA

Melanoma Cell Culture Protocol

Application Details

Suggested Applications: Cellular Assay, IF, Other, WB (Based on references)

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Application Note:	The key applications of these cell lines include genetic studies, xenograft production, drug testing, and drug target discovery. These cell line models can be used in various biological assays, and for identifying critical target genes, and cell signaling pathways.
Assay Dilutions:	All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.

Cell Line Data

Product Type:Viable CellsMorphology:fibroblasticCell Viability:YesStage:MetastasisBRAF:V600ECDK4:WTC-Kit:WTN-RAS:WTPTEN:WTPaired:YesMedium:Tumor Specialized Media with 2% HI-FBSSub-culture:Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.Incubation:36°C with 5% CO2	Cell Line:	Human Melanoma
Cell Viability: Yes Stage: Metastasis BRAF: V600E CDK4: WT C-Kit: WT N-RAS: WT PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	Product Type:	Viable Cells
Stage: Metastasis BRAF: V600E CDK4: WT C-Kit: WT N-RAS: WT PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	Morphology:	fibroblastic
BRAF: V600E CDK4: WT C-Kit: WT N-RAS: WT PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	Cell Viability:	Yes
CDK4: WT C-Kit: WT N-RAS: WT PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	Stage:	Metastasis
C-Kit: WT N-RAS: WT PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	BRAF:	V600E
N-RAS: WT PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	CDK4:	WT
PTEN: WT Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	C-Kit:	WT
Paired: Yes Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	N-RAS:	WT
Medium: Tumor Specialized Media with 2% HI-FBS Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	PTEN:	WT
Sub-culture: Cells should be maintained between 30 – 95% confluence in tumor specialized medium with 2% FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	Paired:	Yes
FBS; split cultures 1:10 every 5-6 days using 0.25% trypsin/EDTA.	Medium:	Tumor Specialized Media with 2% HI-FBS
Incubation: 36°C with 5% CO2	Sub-culture:	•
	Incubation:	36°C with 5% CO2

Formulation

Physical State:	Frozen Cell Suspension
Concentration:	1.0 million cells/mL Count By Hemocytometer
Buffer:	None
Preservative:	None
Stabilizer:	None

Shipping & Handling

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Shipping Condition:	Dry Ice
Storage Condition:	Cells are frozen with 90% FBS/10% DMSO solution at about 1×10^6 cells/ml. Store vial in liquid nitrogen upon arrival.
Expiration:	Expiration date is two (2) years from date of receipt.

Images



Flask

Human melanoma tumor cells with known gene mutations, disease stage, STR, and RPPA profiling

References

- Rajasekaran S, Cheng S, Gajendran N, Shekoohi S, Chesnokova L, Yu X, and Witt SN. Transcriptomic analysis of melanoma cells reveals an association of α -synuclein with regulation of the inflammatory response. *Sci Rep.* (2024)
- Mousson A et al. ---Inhibiting FAK-Paxillin Interaction Reduces Migration and Invadopodia-Mediated Matrix Degradation in Metastatic Melanoma Cells. *Cancers (Basel)*. (2021)
- Juraleviciute M et al. MX2 mediates establishment of interferon response profile, regulates XAF1, and can sensitize melanoma cells to targeted therapy. *Cancer Med.* (2021)
- Hanniford D et al. Epigenetic silencing of CDR1as drives IGF2BP3-mediated melanoma invasion and metastasis. *Cancer Cell.* (2021)

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No test method can provide total assurance that the hepatitis B virus, hepatitis C virus, human immunodeficiency virus, or any other infectious agents are absent. Thus, all blood products, including purified proteins derived from human blood sources, should be handled at Biosafety Level 2 as recommended by the CDC\NIH manual entitled Biosafety in Microbiological and Biomedical Laboratories for potentially infectious human serum, blood specimens or proteins derived from same. Source material for the human blood product supplied to your facility has been tested for the detection of HIV antibody, Hepatitis B surface antigen, antibody to Hepatitis C, HIV 1 antigen(s), antibody to HTLV - I/II, and syphilis by FDA guidelines. All units were found to be non-reactive/negative for these tests. All human blood source material is collected in FDA licensed centers and is tested with FDA approved test kits.

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