

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Datasheet for WM3211-01-0010 WM3211 Viable Cells

Overview

Description:	WM3211 Viable Cells - WM3211-01-0010
Item No.:	WM3211-01-0010
Size:	10 x 1 million cells
Applications:	Biochemical Assay, ELISA, IF, WB
Origin:	Human

Product Details

Background: WM3211 is a tumorigenic (VGP) primary melanoma cell line with competence for metastasis.

These cells display fibroblastic morphology in culture. This cell line was established from a metastatic site (lymph node) in a 34-year-old female. This cell line contains a L576P mutation at position 576 in the c-KIT gene. The L576P mutation results in an amino acid substitution at position 576 in KIT, from a Leucine (L) to a proline (P). This mutation occurs within the juxtamembrane domain. Mutant KIT proteins have increased kinase activity and transforming activity in vitro. This cell line is wild type for BRAF, PTEN, N-RAS, and CDK4. WM3211 cells

produce xenograft tumors when injected into immunocompromised mice.

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

Species of Origin: Human

Target Details

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: Biochemical Assay, ELISA, IF, 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

Cell Line:	Human Melanoma
Product Type:	Viable Cells
Morphology:	fibroblastic
Cell Viability:	Yes
Stage:	VGP
BRAF:	WT
CDK4:	WT
C-Kit:	L576P
N-RAS:	WT
PTEN:	WT
Paired:	No
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 1:3 every 5-6 days using 0.25% trypsin/EDTA.
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 $10x10^6$ 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



Viable cell growth

Established WM3211 viable cell growth in culture using appropriate Tumor Specialized Media with 2%FBS.

References

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- Tong S et al. A Small Peptide Increases Drug Delivery in Human Melanoma Cells. Pharmaceutics. (2022)
- Prouteau A et al. Canine Oral Melanoma Genomic and Transcriptomic Study Defines Two Molecular Subgroups with Different Therapeutical Targets. *Cancers (Basel)*. (2022)
- Ohira T et al. PITX1 inhibits the growth and proliferation of melanoma cells through regulation of SOX family genes. Sci Rep. (2021)

Disclaimer

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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