

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

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Datasheet for WM3912-01-0010 WM3912 Viable Cells

Overview

Description:	WM3912 Viable Cells - WM3912-01-0010
Item No.:	WM3912-01-0010
Size:	10 x 1 million cells
Origin:	Human

Product Details

Froduct Details	
Background:	WM3912 is a metastatic human melanoma cell line established from a neck metastatic site of a patient. This cell line thaws poorly and contains lots of spontaneous senescent cells. This cell line features the N581Y mutation at codon 581 in the BRAF gene. The N581Y mutation results in an amino acid substitution at position 581 in the BRAF, from an asparagine (N) to a tyrosine (Y). This cell line also expresses L213F mutation in CDK4 gene which results in an amino acid substitution at codon 213 in the CDK4 from leucine (L) to phenylalanine (F). WM3912 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

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.

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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
Cell Viability:	Yes
Stage:	Metastasis
BRAF:	N581Y
CDK4:	L213F
C-Kit:	WT
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 $1:3$ 1 week after established 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

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.

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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 WM3912 viable cell growth in culture using appropriate Tumor Specialized Media with 2%FBS.

Disclaimer

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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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