

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

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet for WM3702-01-0005

WM3702 Viable Cells

Overview

Description:	WM3702 Viable Cells - WM3702-01-0005
Item No.:	WM3702-01-0005
Size:	5 x 1 million cells
Origin:	Human

Product Details

Background: WM3702 is a metastatic human melanoma cell line that displays pigmented, thin elongated

morphology in culture. This cell line was established from a right upper flank of back in a 58-year-old male with necrotic fungating melanoma. The subject displayed metastatic melanoma forming multiple subcutaneous nodules with overlying skin ulcerations. This cell line contains a Q61L mutation at position 61 in the N-RAS gene. The Q61L mutation results in an amino acid substitution at position 61 in NRAS, from a glutamine (Q) to a leucine (L). WM3702 cells match with WM3758 and WM3682. These 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

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assavs, a	nd for	identifvir	g critica	Itarget	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:	pigmented, thin elongated
Cell Viability:	Yes
Stage:	Metastasis
BRAF:	WT
CDK4:	WT
C-Kit:	WT
N-RAS:	Q61L
PTEN:	ND
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 1:4 every week 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

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Storage Condition: Cells are frozen with 90% FBS/10% DMSO solution at about 5x10^6 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

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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