

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

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- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in





www.rockland.com tech@rockland.com +1 484.791.3823

Datasheet for WM3704-01-0001 WM3704 Viable Cells

Overview

Description:	WM3704 Viable Cells - WM3704-01-0001		
Item No.:	WM3704-01-0001		
Size:	1 million cells		
Origin:	Human		

Product Details

Background:	WM	3704 is a	metastatic h	numan	melanoma	cell lin	e that o	displ	ays a	mesenc	hymal	morpho	ology.

This cell line was established from a lymph node metastasis site of a patient. This cell line features the specific V600E (Val600Glu) mutation at codon 600 in the BRAF gene. This mutation causes constitutively active kinase activity and activation of MEK and ERK signaling pathway. This cell line also expresses mutated PTEN gene and R24H mutation at position 24 in the CDK4 gene. The R24H mutation results in an amino acid substitution at codon 24 in CDK4, from an arginine (R) to histidine (H). This mutation is an activating mutation, as it results in growth advantages by preventing p16 binding, while maintaining the ability of CDK4 to form a functional kinase with cyclin D. WM3704 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

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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:	mesenchymal
Cell Viability:	Yes
Stage:	Metastasis
BRAF:	V600E
CDK4:	R24H
C-Kit:	WT
N-RAS:	WT
PTEN:	Mutated
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 1:6 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

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Shipping Condition:	Dry Ice
Storage Condition:	Cells are frozen with 90% FBS/10% DMSO solution at about 1x10^6 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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