

# 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

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# Datasheet for WM51-01-0001

# **WM51 Viable Cells**

#### **Overview**

Description:	WM51 Viable Cells - WM51-01-0001
Item No.:	WM51-01-0001
Size:	1 million cells
Applications:	ELISA
Origin:	Human

#### **Product Details**

Background:	WM51 is a metastatic human melanoma cell line that was established from a metastatic tissue site of a male patient. These cells display a mixed morphology and require sub-confluent culture conditions. 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 expresses wild type N-RAS, c-KIT, and CDK4 genes. WM51 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**

Tested Applications: ELISA

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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:	mixed morphology
Cell Viability:	Yes
Stage:	Metastasis
BRAF:	V600E
CDK4:	WT
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, keep cultures sub-confluent; split cultures 1:3 every 1 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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<b>Shipping Condition:</b>	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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