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Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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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 WM3060-01-0010**WM3060 Viable Cells****Overview**

Description:	WM3060 Viable Cells - WM3060-01-0010
Item No.:	WM3060-01-0010
Size:	10 x 1 million cells
Origin:	Human

Product Details

Background:	WM3060 is a metastatic human melanoma cell line established from a metastatic site (left axillary nodes) in a 55-year-old male with level III malignant melanoma. The subject displayed VGP, and RGP with a tumor thickness of 3.0 mm. This cell line was derived from the same patient as the cell lines WM2013, and WM1960. This cell line contains a Q61K mutation at position 61 in the N-RAS gene which causes increased signaling via the extracellular signal-regulated MAPK/ERK kinase pathways to enhance proliferation. This cell line is wild type for BRAF, PTEN, c-KIT, and CDK4. WM3060 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:	<ul style="list-style-type: none">• 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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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.
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Cell Line Data

Cell Line:	Human Melanoma
Product Type:	Viable Cells
Cell Viability:	Yes
Stage:	Metastasis
BRAF:	WT
CDK4:	WT
C-Kit:	WT
N-RAS:	Q61K
PTEN:	WT
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 1:3 every 5 days using 0.25% trypsin/EDTA.
Incubation:	36°C with 5% CO ₂

Formulation

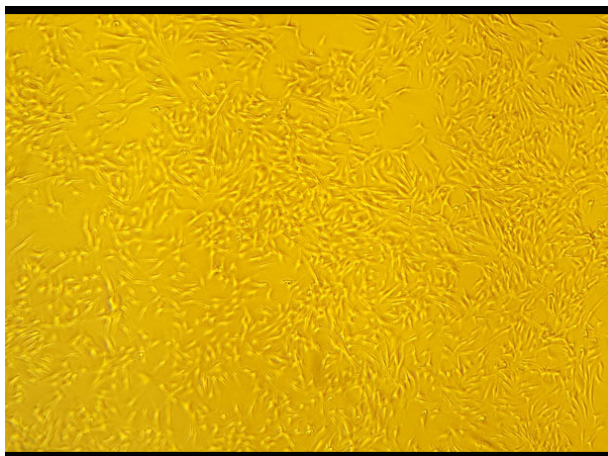
Physical State:	Frozen Cell Suspension
Concentration:	1.0 million cells/mL Count By Hemocytometer
Buffer:	None
Preservative:	None
Stabilizer:	None

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.

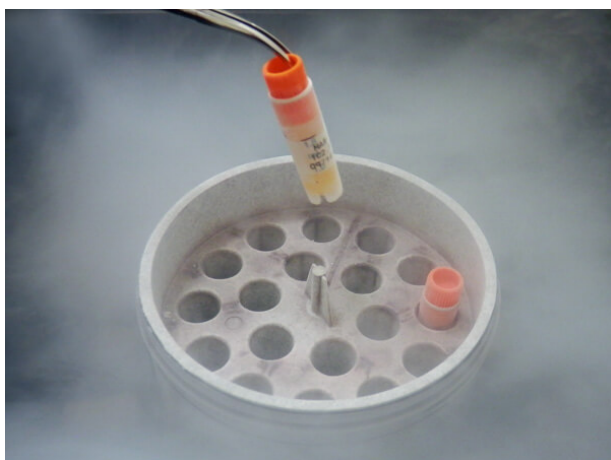
Expiration: Expiration date is two (2) years from date of receipt.

Images



Viable cell growth

Established WM3060 viable cell growth in culture using appropriate Tumor Specialized Media with 2%FBS.



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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