

Datasheet for WM3282-01-0001**WM3282 Viable Cells****Overview**

Description:	WM3282 Viable Cells - WM3282-01-0001
Item No.:	WM3282-01-0001
Size:	1 million cells
Applications:	Other
Origin:	Human

Product Details

Background:	WM3282 is a tumorigenic (VGP) primary melanoma cell line with competence for metastasis. These cells display pigmented, mesenchymal morphology in culture. This cell line contains V600K (Val600Lys) mutation at codon 600 in the BRAF gene. This mutation occurs within the activation segment of the kinase domain and causes constitutively active kinase activity and activation of MEK and ERK signaling pathway. WM3282 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

Suggested Applications:	Other (Based on references)
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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:	pigmented, mesenchymal
Cell Viability:	Yes
Stage:	Primary/VGP
BRAF:	V600K
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; split cultures 1:3 every 1 week using 0.25% trypsin/EDTA.
Incubation:	36°C with 5% CO ₂

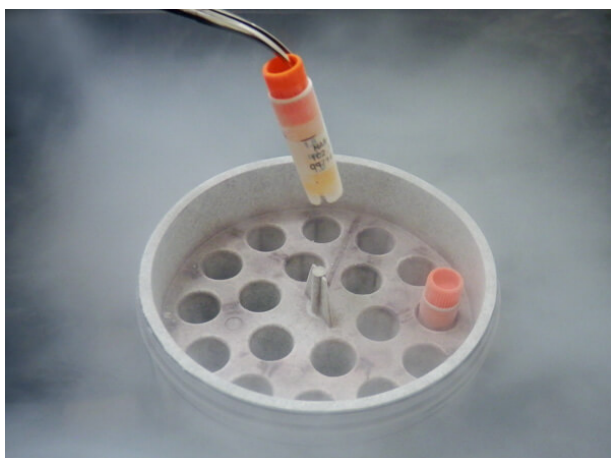
Formulation

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

Shipping & Handling

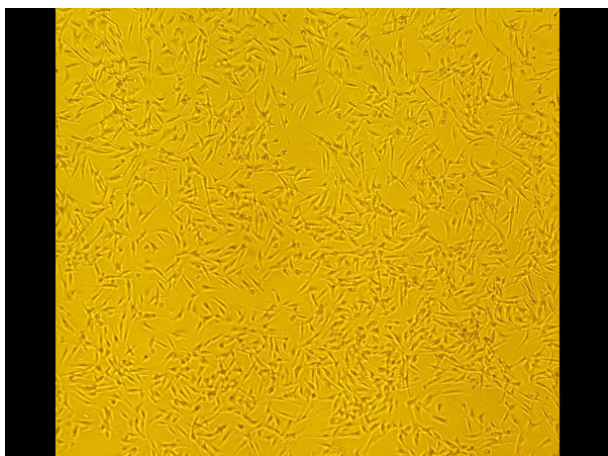
Shipping Condition:	Dry Ice
Storage Condition:	Cells are frozen with 90% FBS/10% DMSO solution at about 1×10^6 cells/ml. Store vial in liquid nitrogen upon arrival.
Expiration:	Expiration date is two (2) years from date of receipt.

Images



Flask

Human melanoma tumor cells with known gene mutations, disease stage, STR, and RPPA profiling



Viable cell growth

Established WM3282 viable cell growth in culture using appropriate Tumor Specialized Media with 2%FBS.

References

- Ashida A et al. Circulating tumour DNA for monitoring treatment response to anti-PD-1 immunotherapy in melanoma patients. *Acta Derm Venereol.* (2017)

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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