

Datasheet for WM1341D-01-0005**WM1341D Viable Cells****Overview**

Description:	WM1341D Viable Cells - WM1341D-01-0005
Item No.:	WM1341D-01-0005
Size:	5 x 1 million cells
Applications:	Cellular Assay, Functional Assay, IF, IHC, Multiplex, WB
Origin:	Human

Product Details

Background:	WM1341D is a tumorigenic (VGP) primary melanoma cell line with competence for metastasis. These cells display epithelial square and triangular morphology in culture. This cell line features the specific V600R (Val600Arg) mutation at codon 600 in the BRAF gene. The V600R mutation results in an amino acid substitution at position 600 in BRAF, from a valine (V) to an arginine (R). This mutation occurs within the activation segment of the kinase domain. Mutations at V600 result in increased kinase activity and are transforming in vitro. WM1341D 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

Suggested Applications:	Cellular Assay, Functional Assay, IF, IHC, Multiplex, WB (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:	epithelial square and triangle
Cell Viability:	Yes
Stage:	VGP
BRAF:	V600R
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 10 days using 0.25% trypsin/EDTA.
Incubation:	36°C with 5% CO ₂

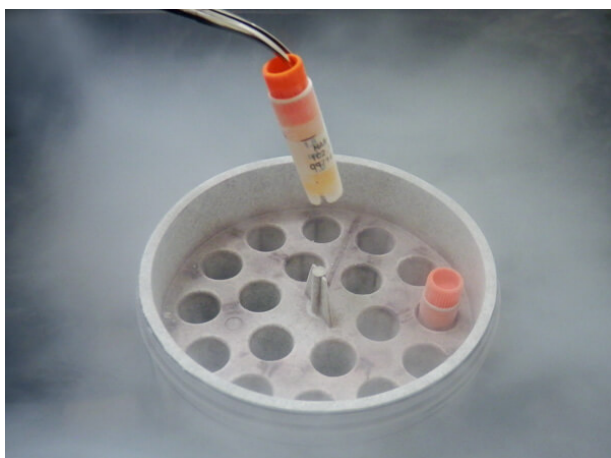
Formulation

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 5×10^6 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 WM1341D viable cell growth in culture using appropriate Tumor Specialized Media with 2%FBS.

References

- Wadzynska J et al. The impact of cellular elements of TME on melanoma biology and its sensitivity to EGFR and MET targeted therapy. *Biochim Biophys Acta Mol Cell Res.* (2023)
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- Makowiecka A et al. Changes in Biomechanical Properties of A375 Cells Due to the Silencing of TMSB4X Expression Are Not Directly Correlated with Alterations in Their Stemness Features. *Cells.* (2021)
- Kupczyk P et al. PARP1 as a Marker of an Aggressive Clinical Phenotype in Cutaneous Melanoma—A Clinical and an In Vitro Study. *Cells.* (2021)
- Malek N et al. The origin of the expressed retrotransposed gene ACTBL2 and its influence on human melanoma cells' motility and focal adhesion formation. *Sci Rep.* (2021)
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- Makowiecka A et al. Thymosin β 4 regulates focal adhesion formation in human melanoma cells and affects their migration and invasion. *Front Cell Dev Biol.* (2019)
- Makowiecka A et al. Varying effects of EGF, HGF and TGF β on formation of invadopodia and invasiveness of melanoma cell lines of different origin. *Eur J Histochem.* (2016)

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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