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LIFESPAN

## ENGRAFTMENT STABILITY

• A functional immune system.

• T cell-dependent inflammatory responses.

• No donor cell immune reactivity towards host.

Long-term > 12 months Stable through the lifespan of the mouse.

• Enables short-term studies requiring

• Strong effector and memory T cell function.

• T cell-driven GvHD.

Short-term < 3 months

Stable through the lifespan of the mouse.

• Faster immune cell repopulation\*.

• Higher myeloid cell engraftment\*.

• Faster lymphoid cell engraftment\*.

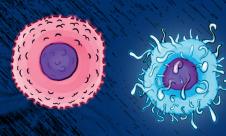
• Higher T cell and dendritic cell populations\*.

• Better AML engraftment\*.

Long-term > 12 months Stable through the lifespan of the mouse.

\* When compared to hu-CD34-NSG™

## **IMMUNE DEVELOPMENT**



B and T cells (MHCrestricted CD4 and CD8)





Macrophages



Dendritic cells



T cells (HLA-restricted CD4 and CD8)



Natural Killer cells



Monocytes

T cells (MHC-restricted CD4 and CD8)

Macrophages

Dendritic cells

## **EFFICACY STUDIES ON DEMAND START YOUR** SELECT A **EFFICACY STUDY 8 CHOOSE A STUDY** PDX LIVE<sup>™</sup> MODEL **MONITOR DATA LIVE**

## **FAST-TRACK YOUR EFFICACY TESTING PROJECTS**

PDX-engrafted mice are a fast and cost-effective platform to simulate trials, evaluate multiple drugs alone or in combination and produce predictive data.

NSG™ strains engrafted with patient-derived tumors (PDX) offer improved retention of tumor heterogeneity and architecture, providing a platform for compound efficacy testing that is more reflective of patient response.

Our searchable PDX database: tumor.informatics.iax.org/ mtbwi/pdxSearch.do

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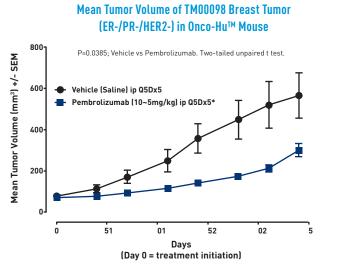
LADDER	TM00015	PIK3CA H1047R
	TM00020	TP53 E336*; KDR 0472H; PTEN T321fs
REAST	TM00089	TNBC ER-/PR-/HER2-; BRCA1 V757fs
	TM00090	TNBC ER-/PR-/HER2-
	TM00096	TNBC ER-/PR-/HER2-
	TM00098	TNBC ER-/PR-/HER2-
	TM00284	ER+/PR+/HER2-
	TM00386	ER+/PR+/HER2-
DLON	TM00179	BRAF V600E/PIK3CA G1049R
ING	TM00302	KRAS G12D/KDR Q472H/TP53 R158L
	TM00784	EGFR L858R
VARY	TM00335	CA125 & MUC16 mRNA elevated
	TM00916	ER+/PR+
ROSTATE	J000079754	Castration-resistant
	TM00298	TP53 R273C/PTEN R233*/PTEN L265fs, AR+
KIN	TM00702	BRAF V600V
	TM01149	BRAF V600V
REAST	TM00095**	ER-,PR-, HER2+
JNG	TM00199**	EGFR L858R
	TM00206**	ELM4-ALK fusion

\*\* Cohorts from these PDX models have slower growth rates and vill take longer to expand and run on study.

# ONCO-HU<sup>™</sup> MODELS THE NEXT GENERATION OF CANCER MODELING

Onco-Hu<sup>™</sup> models are a robust immuno-oncology platform for efficacy testing of novel immunotherapies targeting T cells and myeloid cells to help destroy cancers *in vivo*.

The Onco-Hu<sup>™</sup> platform is based on NSG<sup>™</sup> and NSG<sup>™</sup>-SGM3 mice, dually engrafted with human CD34+ hematopoietic stem cells (HSC) and clinically relevant PDX Live™ low passage tumors.



Onco-Hu™ models engrafted with PDX Live™ clinically relevant breast or lung tumors allow the evaluation of the efficacy of immunomodulators -alone or in combination therapies- to treat cancer.

jax.org/onco-hu

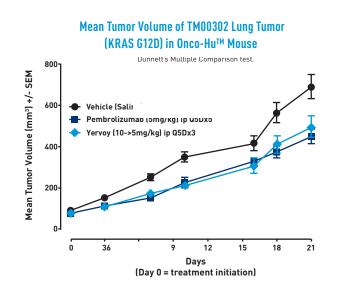
## FOR MORE INFORMATION **CONTACT TECHNICAL INFORMATION SERVICES** REGIONAL REPRESENTATIVE Find your regional rep. Visit:

#### JAX<sup>®</sup> Mice. Clinical & Research Services

The Jackson Laboratory Bar Harbor, Maine | Farmington, Conn. | Sacramento, Calif.

Technical Information Services

micetech@jax.org 1-800-422-6423 (US, Canad<u>a & Puerto Rico)</u> 1-207-288-5845 (from any location)





eading the search

for tomorrow's cures

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# THE NEXT ERONTIER DISEASE MODELING

JAX<sup>®</sup> offers humanized NSG<sup>™</sup> and NSG<sup>™</sup>-based strains with additional manipulations of the host (including the NSG<sup>™</sup>-SGM3 strain that expresses the human cytokines IL-3, GM-CSF, and SCF) for an enhanced ability to recapitulate specific functions of human disease in vivo. These humanized NSG™ strains have superior T cell-dependent immune responses and myeloid engraftment.

These highly immunodeficient mouse models, only available from The Jackson Laboratory, provide the unmatched ability to model normal and malignant tissues and cells notoriously difficult to engraft and study in other mouse strains, including:

- Hematopoietic stem cells
- Patient-derived xenografts (PDX)

PDX-engrafted NSG™ mice from our *In Vivo* Pharmacology Services are early passage to maintain tumor structure fidelity and cellular heterogeneity that is seen in the patient. These models serve as a clinically relevant platform for validation of compounds and investigations into cancer pathology, including immuno-oncology.

## Which model is best for your research?

JAX<sup>®</sup> humanized NSG<sup>TM</sup> and NSG<sup>TM</sup>-SGM3 mice generate functional human immune systems with different capabilities.

# Empowering Clinically Relevant Modeling Using NSG<sup>™</sup> Strains

Whether you are investigating the interactions between host immune cells and pathogens, tumors and immune cells, o require an oncological platform to test your therapeutic compounds, NSG™ strains have revolutionized how infectious disease and cancer research is conducted. By supporting the engraftment of human immune cells and tumors, NSG™ strains have transformed the approach taken by the biomedical community to understand human diseases.

## **RESEARCH FIELDS**

