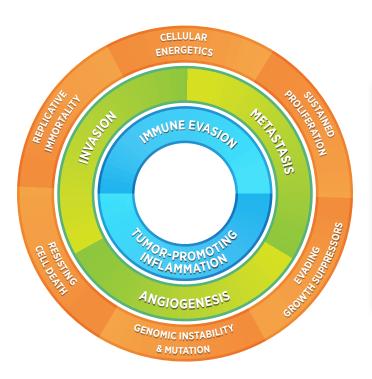
# nCounter® Tumor Signaling 360 Panel Gene Expression Panel

## Targeted Therapeutics • Tumor Signaling • Immune Response

Covering hundreds of genes involved in tumorigenesis, metastasis, and inflammation, the nCounter Tumor Signaling 360 Panel offers a holistic view of the biology of the tumor, microenvironment, and immune response with an emphasis on dysfunctional cell signaling in cancer. Identify targets for novel therapies and understand the mechanism of action for current ones with a comprehensive look at 40+ pathways involved in tumor biology, immune evasion, and remodeling of the microenvironment.



## **Product Highlights**

- Profile 780 human and mouse across 40+ annotated pathways involved in tumor signaling
- Assess altered signaling pathways
- · Identify targets for novel therapeutics
- Understand the mechanism of action of targeted therapies
- Measure the anti-tumor immune response
- · Quantify the relative abundance of immune cell types

Feature	Specifications
Number of Targets	780 (Human), 780 (Mouse), including internal reference genes
Sample Input - Standard (No amplification required)	25-300 ng
Sample Input - Low Input	As little as 1 ng with nCounter Low Input Kit (sold separately)
Sample Type(s)	Cultured cells/cell lysates, sorted cells, FFPE-derived RNA, total RNA, fragmented RNA, PBMCs, and whole blood/plasma
Customizable	Add up to 55 unique genes with Panel Plus
Time to Results	Approximately 24 hours
Data Analysis	nSolver™ Analysis Software (RUO) and the ROSALIND® Platform

#### **Core Themes and Annotations**

The nCounter 360 series of cancer gene expression panels have been developed to comprehensively profile the tumor, immune response, and microenvironment. As therapeutics that target tumor signaling evolve, a greater understanding of tumor signaling is required as well as a better understanding of the interactions of tumor cells with the tissue milieu. The Tumor Signaling 360 Panel is intended to play this exact role, enabling deeper profiling of the tumor that is complete, yet focused on signaling pathways of interest for targeted therapeutic development.

Tumor Signaling								
Cellular Energetics (Hs/Mm)	Sustained P (Hs/Mm)	roliferation	Evading Growth Suppressors (Hs/Mm)	Enabling Replicative Immortality (Hs/Mm)	Genomic In: Mutation (Hs/Mm)	stability &	Resisting Cell Death (Hs/Mm)	
105/104 Genes	223/221 Ger	nes	79/79 Genes	48/48 Genes	111/111 Genes		23/23 Genes	
Autophagy	Androgen Signaling		Cell Cycle	Immortality & Stemness	DNA Damage & Repair		Apoptosis	
Glucose Metabolism	EGFR Signaling		Senescence	Epigenetic & Transcriptional Regulation	TNF Superfamily			
Glutamine Metabolism	ERBB2 Signaling			p53 Signaling				
Lipid Metabolism	Estrogen Sig	gnaling						
mTOR Signaling	FGFR Signaling							
Nrf2 & Oxidative Stress	Hedgehog							
	MAPK Signa	lling						
	MET Signalii	ng						
	Мус							
	Notch Signa	ling						
	PI3K-Akt Signaling TGF-beta Signaling							
	Wnt Signalir	ng						
	Immune Response		Tumor Microenvironme			nt		
Immune Evasion Tumor-P (Hs/Mm)		romoting Inflammation (Hs/Mm)	Activating Invasion and Metastasis (Hs/Mm)			Angiogenesis (Hs/Mm)		
122/124 Genes		178/171 Genes 146/146 Genes				67/67 Genes		
Antigen Presentation		Chemokine	Signaling	Signaling Cell Adhesion & Motility		HIF1 Signaling		
B cell Function	Inflammatio		n	ECM Remodeling & Metastasis		PDGF Signaling		
Cytotoxicity	Interferon F		sponse EMT			VEGF Signaling		
Myeloid Immune Evasion JAK-STAT		JAK-STAT S	ignaling	Hippo Signaling				
T cell Co-stimulation		NF-kB Sign	aling					
T cell Exhaustion								
TCR Signaling								
Tumor Antigen								

To view the annotated gene lists for the Tumor Signaling 360 Panel, visit nanostring.com/TS360

### nSolver™ Analysis Software

NanoString offers advanced software tools that address the continuous demands of data analysis and the need to get simple answers to specific biological questions easily. Genes included in the Tumor Signaling 360 Panel are organized and linked to various advanced analysis modules.

#### ROSALIND® Platform

ROSALIND is a cloud-based platform for life science research that enables scientists to analyze and interpret differential gene expression data without need for bioinformatics or programming skills. ROSELIND makes analysis for nCounter data easy, with guded modules for:

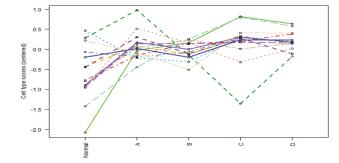
Normalization / Quality Control / Individual Pathway Analysis Differential Expression / Gene Set Analysis

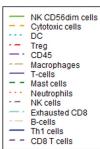
nCounter customers can access ROSALIND free of charge at  $\underline{\textbf{rosalind.bio/nanostring}}.$ 

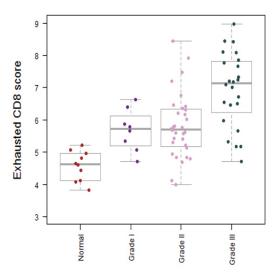
#### **Immune Cell Profiling Feature**

Genes included in the Tumor Signaling 360 Panel provide unique cell profiling data to measure the relative abundance of 14 different human immune cell types. The table below summarizes each cell type represented by gene content in the panel, as qualified through biostatistical approaches and selected literature in the field of immunology.

Cell Type	Associated Human Genes
B Cells	BLK, CD19, FAM30A, FCRL2, MS4A1, PNOC, SPIB, TCL1A, TNFRSF17
CD45	PTPRC
CD8	CD8A, CD8B
Cytotoxic Cells	CTSW, GNLY, GZMA, GZMB, GZMH, KLRB1, KLRD1, KLRK1, NKG7, PRF1
Dendritic Cells	CCL13, CD209, HSD11B1
Exhausted CD8	CD244, EOMES, LAG3, PTGER4
Macrophages	CD163, CD68, CD84, MS4A4A
Mast Cells	CPA3, HDC, MS4A2, TPSAB1/B2
Neutrophils	CEACAM3, CSF3R, FCAR, FCGR3A/B, FPR1, S100A12, SIGLEC5
NK Cells	NCR1, XCL1/2
NK CD56dim Cells	IL21R, KIR2DL3, KIR3DL1, KIR3DL2
T Cells	CD3D, CD3E, CD3G, CD6, SH2D1A, TRAT1
Th1 Cells	TBX21
Treg	FOXP3







# **Tumor Inflammation Signature**

Included within the Tumor Signaling 360 panel is the Tumor Inflammation Signature (TIS). This 18-gene signature measures activity known to be associated with response to PD-1/PD-L1 inhibitors.

- Measures 4 axes of biology to characterize a peripherally suppressed immune response
  - Antigen Presenting Cells
  - T Cell/NK Presence
  - IFNγ Biology
  - T Cell Exhaustion

- Tissue-of-origin agnostic (Pan-cancer)
- Potential surrogate for PD-L1 and mutational load in a research setting

### **Ordering Information**

Gene Expression Panels arrive ready-to-use and generally ship within 24 hours following purchase.

Product	Product Description	Quantity	Catalog Number
nCounter Human Tumor Signaling 360 Panel	Gene Expression CodeSet profiling (780 genes) 760 tumor signaling related genes + 20 internal reference controls. No Master Kit	12 Reactions	XT-CSO-H-TS360-12
nCounter Human Tumor Signaling 360 Panel Standard	Standard containing a pool of synthetic DNA oligonucleotides that correspond to the target sequence of each of the 780 unique probe targets in the panel.	12 Reactions	PSTD-H-T360-12
nCounter Mouse Tumor Signaling 360 Panel	Gene Expression CodeSet profiling (780 genes) 760 tumor signaling related genes + 20 internal reference controls. No Master Kit	12 Reactions	XT-CSO-M-TS360-12
nCounter Mouse Tumor Signaling 360 Panel Standard	Standard containing a pool of synthetic DNA oligonucleotides that correspond to the target sequence of each of the 780 unique probe targets in the panel.	12 Reactions	PSTD-M-T360-12
nCounter Analysis System Master Kit Reagents and Cartridges	Reagents, cartridges, and consumables necessary for sample processing on the nCounter Analysis System	12 Reactions	NAA-AKIT-012
nCounter SPRINT Cartridge (1 Cartridge, 12 lanes)	Sample Cartridge for nCounter SPRINT System	12 Reactions	SPRINT-CAR-1.0
nCounter SPRINT Reagent Pack	nCounter SPRINT Reagent Pack containing Reagents A, B, C, and Hybridization Buffer	192 Reactions	SPRINT-REAG-KIT
Low Input RNA Reagent Kit	48 rxn kit for profiling from low sample input amounts	48 Reactions	LOW-RNA-48

#### **Selected Publications**

- 1. Bieging KT and Attardi LD. Deconstructing p53 Transcriptional Networks in Tumor Suppression. Trends Cell Biol. 2012; 22 (2), 97-106.
- 2. Bild, AH et al. Oncogenic Pathway Signatures in Human Cancers as a Guide to Targeted Therapies. Nature. 2006;439 (7074), 353-7.
- 3. Chiaradonna, F. Ras-dependent Carbon Metabolism and Transformation in Mouse Fibroblasts. Oncogene. 2006;25 (39), 5391-404.
- 4. Cordenonsi, M et al. The Hippo Transducer TAZ Confers Cancer Stem Cell-Related Traits on Breast Cancer Cells. Cell. 2011;147(4):759-72.
- 5. Hanahan D and Weinberg, RA. Hallmarks of cancer: the next generation. Cell. 2011;144(5):646-74.
- 6. Danaher, P et al. Gene Expression Markers of Tumor Infiltrating Leukocytes. J Immunother Cancer. 2017;21(5):18.
- 7. Jia, P and Zhao Z. Characterization of Tumor-Suppressor Gene Inactivation Events in 33 Cancer Types. Cell Reports. 2019;26, 496-506.
- 8. Malta, TM et al. Machine Learning Identifies Stemness Features Associated With Oncogenic Dedifferentiation. Cell. 2018;173(2): 338-354.e15.
- 9. Sanchez-Vega F et al. Oncogenic Signaling Pathways in The Cancer Genome Atlas. Cell. 2018; 173 (2), 321-337.e10.
- 10. Semenza, GL Hypoxia-inducible Factor 1: Oxygen Homeostasis and Disease Pathophysiology. Trends Mol Med. 2001;7(8):345-50.
- Singh, A et al. RNAi-mediated Silencing of Nuclear Factor erythroid-2-related Factor 2 Gene Expression in Non-Small Cell Lung Cancer Inhibits Tumor Growth and Increases Efficacy of Chemotherapy. Cancer Res. 2008; 68 (19), 7975-84.
- 12. Sun, J et al. A systematic analysis of FDA-approved anticancer drugs. BMC Systems Biology. 2017, 11(Suppl 5):87.
- 13. Sweet-Cordero A et al. An Oncogenic KRAS2 Expression Signature Identified by Cross-Species Gene-Expression Analysis. Nat Genet. 2005; 37 (1), 48-55.
- 14. Way, GP et al. Machine Learning Detects Pan-cancer Ras Pathway Activation in The Cancer Genome Atlas. Cell Rep. 2018;23(1): 172-180.e3.

## For more information, please visit nanostring.com

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