## ONCOTYPE MAP™ PAN-CANCER TISSUE TEST

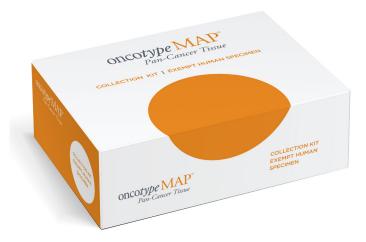
RAPID COMPREHENSIVE GENOMIC PROFILING TO AID THERAPY SELECTION

## **Every day matters. Why wait?**

Oncotype MAP Pan-Cancer Tissue test delivers rapid, comprehensive tumor profiling to aid therapy selection for patients with advanced, metastatic, refractory, or recurrent cancer. The test identifies actionable genomic alterations within 3-5 business days to guide timely treatment decisions, usually before the next follow-up visit.

Utilizing next generation sequencing (NGS) and a broad array of immunohistochemical (IHC) stains and panels, the Oncotype MAP Pan-Cancer Tissue test identifies clinically-relevant genomic alterations from patient specimens as small as 3 mm<sup>2</sup> of tissue or 2-3 slides<sup>‡</sup> allowing you to identify appropriate patient therapy based on tumor characteristics.

Results are provided in an easy-to-interpret report with NCCN® actionable indications for therapy, along with potential evidence-based therapies and clinical trials, to guide treatment options for a breadth of solid tumor types.



# Answers you need at the speed your patients deserve



#### **Fast**

- Results reported in 3-5 business days\*†
- Quantity Not Sufficient (QNS) results typically reported within 48 hours



#### Comprehensive

- 257 genes, 11 select fusions
- 22 IHC stains including PD-L1 (22C3, SP142)
  - 25+ disease-specific panels
- Copy number variants, tumor mutation burden, microsatellite instability, and protein expression.



#### Actionable

- 100+ therapies, 45+ combination therapies, 650+ clinical trials
- NCCN Compendium recommendations



#### Accurate

- FNA and FFPE samples as small as 3 mm² with ≥15% tumor content
- >99% sensitivity for SNVs and Indels, detection down to ≥7.5% mutant allele frequency (MAF) with >99% specificity\*

Oncotype MAP Pan-Cancer Tissue test helps guide therapy selection for today and captures emerging markers that may find their way into guidelines tomorrow.

Order now for your patients with solid tumors



# Oncotype MAP™ Pan-Cancer Tissue Test Markers

Immunohistochemistry										
Single IHC stains		Tumor-specific Panels								
ALK AR CAIX ER hENTI HER2 IDO MET MGMT MLHI MMR (4 IHC) MSH2 MSH6	PD1 PD-L1 (22C3) PD-L1 (SP142) PMS2 PR PTEN ROS1 TOP1 TP TRKpan TS TUBB3	Anal Carcinoma: PD-L1 (22C3), PD-1, TS, TUBB3 Appendix: HER2, PD-L1 (22C3), TOP1, PTEN Bladder: PD-L1 (22C3), PD-L1 (SP142), hENTI, TUBB3 Bone Cancer: TOP1, MGMT, CAIX, HENTI Breast: AR, PD-L1 (2P12), PD-L1 (2P12								3, hENTI, TS
257 NGS Gene Panel										
ABCB1 ABCC1 ABCC2 ABL1 ACVR1 ACVR1B ACVR2B ACVRL1 ADAMTS1 ADAMTS1 ADAMTS6 ADAMTS18 ADAMTS18 ADAMTS18 ADAMTS11 AKT1 AKT1 AKT2 AKT3 ALK AMER1 APC APLNR AR ARAF	AREG ARIDIA ARIDIB ARID2 ATM ATR ATRX AURKA AURKB AXINI AXL B2M BAPI BARDI BCOR BMP6 BMPRIA BMPRIB BNIP3 BRAF BRCA1 BRCA2 BRIP1 BTK	BUB1B CALR CBL CCND1 CCND2 CCND3 CCNE1 CD274 CDA CDC73 CDH1 CDK4 CKD6 CDK12 CDK12 CDKN2A CHEK1 CHEK2 CHFR CHKA CIC CREBBP CSFIR CTLA4 CTNNB1	CYP19A1 CYP1A1 CYP2D6 CYP3A4 CYSLTR2 DCK DDR2 DICER1 DNMT3A EGFR EMSY EP300 EPCAM EPHA5 EPHA7 ERBB2 ERBB3 ERBB4 ERCC1 ERCC2 ERCC3 ERRFII ESR1 ESR2	EWSRI EZH2 FAM175A FANCA FANCC FANCD2 FANCE FANCF FANCG FANCH FATT FBXW7 FCGR2A FGD4 FGF3 FGF4 FGFR1 FGFR2 FGFR3 FGFR4 FLT3 FLT4 FOXL2 FUBP1	GAS6 GATA3 GLI1 GNAQ GNAS GSTP1 HAMP HDAC2 HGF HNF1A HRAS HSD3B1 IDH1 IDH2 IGFIR IKZF1 ILGR JAK1 JAK2 JAK3 KDM5C KDM6A KDR	KEAPI KIT KRAS MAF MAP2KI MAP2KI MAP3KI MAPKI MAPKAPKS MDM2 MDM2 MED12 MEN1 MET MGMT MLH1 MPL MRE11A MSH2 MSH6 MTHFR MTOR MUTYH	MYC MYCN MYODI NBN NF1 NF2 NFE2L2 NOTCH1 NOTCH2 5 NOTCH3 NPMI NRAS NTRK1 NTRK2 NTRK3 PALB2 PBRM1 PDCDILG2 PDGFRA PDGFRA PDGFRA PDGFRB PIK3CA PIK3CB PIK3CCB	PIK3RI PIMI PLCB4 PLCG1 PMS2 POLD1 POLE PP2RIA PTCH1 PTEN PTPNII RAD50 RAD51C RAD51D RAF1 RB1 RBM10 RECQL RET RHEB RICTOR RITI RNF43 ROS1	RPTOR RRMI SDHB SDHC SETD2 SF3BI SMAD1 SMAD2 SMAD4 SMAD5 SMAD9 SMARCA4 SMARCBI SMO SOCSI SPOP STAG2 STAT3 STAT5A STAT5B STKII SUFU TERT-p TGFBI	TGFB2 TGFB3 TGFBR1 TGFBR2 TNFA1P3 TNK1 TOP2A TP53 TSC1 TSC2 TSHR TYMS VEGFA VHL WTI XRCC1 YES1

Genetic Structures Tested: Single nucelotide variants (SNVs) and insertions/deletions in coding regions of genes listed above; UTRs and splice junctions when actionable (e.g. MET exon 14 skipping and EGFRVIII). MSI; mutation burden (SNV's, insertions, deletions) based on -1 megabase; select fusions involving ALK, BRAF, FGFR1, FGFR2, FGFR3, MET, RET, ROS1, NTRK1, NTRK2, NTRK3 (ETV6); and copy number variants.

Turnaround time is based on when sample is received. Mutation calls may not be available from some regions due to pseudogenes or sequence context. Select IHCs may not be run if already performed within the last six months unless indicated in the notes section. HER2 equivocal by IHC will be reflexed to FISH testing in select tumor types. Reflex testing will exceed standard turnaround time for results. MMR includes the following IHCs: MLH1, MSH2, MSH6, PMS2.





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### **About Exact Sciences**

A leading provider of cancer screening and diagnostic tests, Exact Sciences helps people get the answers they need to make more informed decisions across the cancer continuum. Building on the success of the Cologuard and Oncotype DX tests, Exact Sciences is investing in its product pipeline to take on some of the deadliest cancers and improve patient care. Through an innovative, rigorous approach, and with the support of visionary collaborators, we're helping advance the fight against cancer.

#### References

\* Data on file, in-house assay optimization protected by trade-secret/patent regulations. † Turnaround time is based on sample receipt.

‡ Morris S, Subramanian J, Gel E, Runger G, Thompson E, Mallery D, et al. Performance of next-generation sequencing on small tumor specimens and/or low tumor content samples using a commercially available platform. PLoS ONE. (2018); 13(4): e0196556. https://doi.org/10.1371/journal.pone.0196556.

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