

# SERVICES PORTFOLIO



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# **MOLECULAR TESTING OF TUMOR TISSUE SAMPLES**

**NUCLEIC ACID EXTRACTION**

PP-T-001	<b>Macrodissection and DNA extraction from FFPE tissue</b>
PP-T-002	<b>Macrodissection and RNA extraction from FFPE tissue</b>

**COMBINED STUDIES**

CS-T-001	<p><b>SOLID TUMOR (<i>Recommended for LUNG, MELANOMA, BREAST and other Solid tumors</i>)</b></p> <p><b>CS-T-001</b></p> <p><b>NGS Somatic QIaseq DNA Custom Extended Panel (30 genes)</b>          Genes: <i>ALK, ARID1A, BRAF, CDK4, CDK6, EGFR, ERBB2, ERBB4, FAT1, FGFR1, FGFR2, FGFR3, IDH1, IDH2, KEAP1, KIT, KRAS, MET, MYC, NFE2L2, NRAS, PDGFRA, PIK3CA, POLD1, POLE, RET, ROS1, SETD2, STK11, TP53</i></p> <p><b>nCounter RNA Fusion and Gene Expression Tumor Panel*</b>          Fusions: <i>ALK, FGFR1,2,3, NRG1, NTRK1,2,3, RET, ROS1</i>          Splicing variant: <i>EGFRvIII, METΔ14</i>          Gene expression analysis: <i>ALK, ATM, BARD1, BID, BRCA1, BRCA2, CCNE1, CD274, CD4, CD8A, CDK2, EGFR, FGFR1,2,3, FOXP3, GZMM, IFNG, KRAS, NRG1, NTRK1,2,3, PDCD1, PTEN, RET, ROS1, STK11, TP53</i></p>
CS-T-002	<p><b>CS-T-002</b></p> <p><b>Combined study CS-T-001 and PD-L1 IHC</b></p>

*\*The nCounter RNA Pangaea Solid Tumor Panel could be replaced by the NGS RNA Fusion XP Lung Panel (ask for prices)*

CS-T-003	<p><b>UROTHELIAL carcinoma study</b></p> <p><b>CS-T-003</b></p> <p><b>NGS Somatic QIaseq DNA Custom Extended Panel (30 genes)</b></p> <p><b>nCounter RNA Fusion Urothelial Carcinoma Panel</b>          Fusions: <i>FGFR2 and FGFR3</i>  <b>Identification of the variants: <i>FGFR2-BICC1, FGFR2-CASP7, FGFR3ex17-TACC3ex10, FGFR3ex17-TACCex11, FGFR3-BAIAP2L1</i></b></p>
CS-T-004	<p><b>COLON study</b></p> <p><b>CS-T-004</b></p> <p><b>KRAS, BRAF, NRAS sequencing (NGS Custom Panel) (19 genes)</b>          Genes: <i>ALK, BRAF, CDK4, CDK6, EGFR, ERBB2, ERBB4, FGFR1, IDH1, IDH2, KIT, KRAS, MET, NRAS, PDGFRA, PIK3CA, ROS1, STK11, TP53</i></p> <p><b>Mismatch Repair (MMR) status (IHC)</b>          Genes: <i>MLH1, MSH2, MSH6, PMS2</i></p>
MS-T-001	<p><b>GIST study</b></p> <p><b>c-KIT, PDGFR sequencing (NGS Custom Panel) (19 genes)</b></p>

## — MULTIGENIC MOLECULAR PANELS

MS-T-002	<b>NGS somatic QIAseq DNA Breast-Gynecological custom Panel (11 genes)</b> Genes: <i>AKT, BRCA1, BRCA2, CTNNB1, ERBB2, ESR1, PALB2, PIK3CA, POLE, PTEN, TP53</i>
MS-T-003	<b>NGS Somatic QIAseq DNA Custom Extended Panel (30 genes)</b> Genes: <i>ALK, ARID1A, BRAF, CDK4, CDK6, EGFR, ERBB2, ERBB4, FAT1, FGFR1, FGFR2, FGFR3, IDH1, IDH2, KEAP1, KIT, KRAS, MET, MYC, NFE2L2, NRAS, PDGFRA, PIK3CA, POLD1, POLE, RET, ROS1, SETD2, STK11, TP53</i>
MS-T-004	<b>NGS RNA Fusion XP Lung Panel (13 genes)</b> Fusions: <i>ALK, BRAF, EGFR, FGFR1,2,3, MET, NRG1, NTRK1,2,3, RET, ROS1</i> Splicing variant: <i>METΔ14</i>
MS-T-005	<b>NGS Somatic QIAseq DNA Repair Genes Panel (28 genes)</b> Genes: <i>APC, ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CDKN2A, CHEK2, CTNNB1, EPCAM, FANCA, MLH1, MRE11A, MSH2, MSH6, MUTYH, NBN, PALB2, PIK3CA, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, STK11, TP53</i>
MS-T-006	<b>nCounter RNA Fusion and Gene expression Tumor Panel</b> Fusions: <i>ALK, FGFR1,2,3, NRG1, NTRK1,2,3, RET, ROS1</i> Splicing variants: <i>EGFRvIII, METΔ14</i> Gene expression analysis: <i>ALK, ATM, BARD1, BID, BRCA1, BRCA2, CCNE1, CD274, CD4, CD8A, CDK2, EGFR, FGFR1,2,3, FOXP3, GZMM, IFNG, KRAS, NRG1, NTRK1,2,3, PDCD1, PTEN, RET, ROS1, STK11, TP53</i>
MS-T-007	<b>nCounter RNA Pangaea Urothelial Carcinoma Panel</b> Fusions: <i>FGFR2 and FGFR3</i> <b>Identification of the variants: <i>FGFR2-BICC1, FGFR2-CASP7, FGFR3ex17-TACC3ex10, FGFR3ex17-TACCex11, FGFR3-BAIAP2L1</i></b>
MS-T-008	<b>PROSIGNA</b> Genetic prognostic signature for breast cancer

## — INDIVIDUAL GENE MUTATIONS (TaqMan® allelic discrimination qPCR assay)

ISM-T-001	<b>One hotspot mutation (<i>BRAF, EGFR, KRAS, PIK3CA</i>)</b>
ISM-T-002	<b><i>EGFR</i> exon 19 del and <i>EGFR</i> exon 21 (L858R, L861Q) mutations</b>
ISM-T-003	<b><i>EGFR</i> exon 19 de, <i>EGFR</i> exon 21 (L858R, L861Q) and <i>EGFR</i> exon 20 T790M mutations</b>
ISM-T-004	<b><i>EGFR</i> exon 20 resistance mutation T790M</b>
ISM-T-005	<b><i>EGFR</i> exon 20 resistance mutation C797S</b>
ISM-T-006	<b><i>KRAS</i> hotspot mutations (codons 12, 13 and 61)</b>
ISM-T-007	<b><i>BRAF</i> hotspot mutations (V600E, V600K, V600R)</b>
ISM-T-008	<b><i>PIK3CA</i> hotspot mutations (codons 542, 545 and 1047)</b>
ISM-T-009	<b><i>ESR1</i> hotspot mutations (codons 380, 536, 537, 538)</b>

## — SPlicing VARIANTS AND FUSIONS (nCounter Technology)

IST-T-001	<b>Fusion variant or <i>MET</i> splicing variant single analysis</b> Fusions: <i>ALK, FGFR1,2,3, NRG1, NTRK1,2,3, RET, ROS1</i> Splicing variants: <i>EGFRvIII, METΔ1</i>
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## ANATOMIC PATHOLOGY

## — ANATOMICAL PATHOLOGY DIAGNOSIS

PP-T-003	<b>Anatomical pathology diagnosis (H&amp;E and 4 IHC analysis)</b>
PP-T-004	<b>Confirmation of anatomical pathology diagnosis (Previous AP report required)</b>

## — MAIN FISH ASSAYS\*

ISF-T-001	<b>HER-2/neu amplification**</b>
ISF-T-002	<b><i>ALK</i> fusions**</b>
ISF-T-003	<b><i>ROS1</i> fusions</b>
ISF-T-004	<b><i>RET</i> fusions</b>
ISF-T-005	<b><i>MET</i> amplification</b>
ISF-T-006	<b><i>EGFR</i> amplification</b>
ISF-T-007	<b><i>KRAS</i> amplification</b>
ISF-T-008	<b><i>NTRK1</i> fusions</b>
ISF-T-009	<b><i>NTRK2</i> fusions</b>
ISF-T-010	<b><i>NTRK3</i> fusions</b>
ISF-T-011	<b><i>FGFR1</i> fusions</b>
ISF-T-012	<b><i>FGFR1</i> amplification</b>

ISF-T-013	<b>FGFR2 fusions</b>
ISF-T-014	<b>FGFR2 amplification</b>
ISF-T-015	<b>FGFR3 fusions</b>
ISF-T-016	<b>FGFR3 amplification</b>
ISF-T-017	<b>BCL2 fusions</b>
ISF-T-018	<b>BCL6 fusions</b>
ISF-T-019	<b>NRG1 fusions</b>
ISF-T-020	<b>MYC fusions</b>
ISF-T-021	<b>MYC amplification</b>
ISF-T-022	<b>1p/19q loss</b>

\* Ask for other IHC or FISH assays if required

\*\* FDA approved

### — MAIN PRONOSTIC AND PREDICTIVE IMMUNOHISTOCHEMISTRY (IHC) ASSAYS\*

ISI-T-001	<b>ALK (D5F3) **</b>
ISI-T-002	<b>PDL1 (22C3)</b>
ISI-T-003	<b>Mismatch Repair (MMR) status (MLH1, MSH2, MSH6, PMS2)</b>
ISI-T-004	<b>MET (SP44)</b>
ISI-T-005	<b>BRG1 (SMARCA4) (E8V5B)</b>
ISI-T-006	<b>ERBB2 (4B5)</b>

\* Ask for other IHC or FISH assays if required

\*\* FDA approved

### EXTERNALIZED SERVICES

PO-E-001	<b>ONCOPDO 3D Organoid Test (Invitrocue)</b>
PO-E-002	<b>Oncotype DX</b>
PO-E-003	<b>Foundation ONE CDx</b>

# **MOLECULAR TESTING OF TUMOR LIQUID BIOPSY SAMPLES**

## NUCLEIC ACID EXTRACTION

PP-BL-001	cfDNA extraction from liquid biopsies
PP-BL-002	cfRNA extraction from liquid biopsies
PP-LB-003	DNA extraction from white blood cells

## COMBINED STUDIES

CS-BL-001	<p><b>CS-BL-001 (Recommended for all type of tumors)</b>  <b>NGS Somatic QIAsq DNA Custom Panel (19 genes)</b>          Genes: <i>ALK, BRAF, CDK4, CDK6, EGFR, ERBB2, ERBB4, FGFR1, IDH1, IDH2, KIT, KRAS, MET, NRAS, PDGFRA, PIK3CA, ROS1, STK11, TP53</i></p> <p><b>nCounter RNA Fusion Solid Tumor Panel for liquid biopsies (6 genes)</b>          *Fusions: <i>ALK, NRG1, NTRK1, RET, ROS1</i>          Splicing variants: <i>METΔ14</i>          *More frequently found fusions</p>
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## MULTIGENIC MOLECULAR PANELS

MS-BL-001	<p><b>NGS Somatic QIAsq DNA Custom Panel (19 genes)</b>          Genes: <i>ALK, BRAF, CDK4, CDK6, EGFR, ERBB2, ERBB4, FGFR1, IDH1, IDH2, KIT, KRAS, MET, NRAS, PDGFRA, PIK3CA, ROS1, STK11, TP53</i></p>
MS-BL-002	<p><b>NGS Somatic QIAsq DNA Custom Extended Panel (30 genes)</b>          Genes: <i>ALK, ARID1A, BRAF, CDK4, CDK6, EGFR, ERBB2, ERBB4, FAT1, FGFR1, FGFR2, FGFR3, IDH1, IDH2, KEAP1, KIT, KRAS, MET, MYC, NFE2L2, NRAS, PDGFRA, PIK3CA, POLD1, POLE, RET, ROS1, SETD2, STK11, TP53</i></p>
MS-BL-003	<p><b>nCounter RNA Pangaea Solid Tumor Panel for Liquid Biopsies (6 genes)</b>          *Fusions: <i>ALK, NRG1, NTRK1, RET, ROS1</i>          Splicing variants: <i>METΔ14</i>          *More frequently fusions</p>
MS-BL-004	<p><b>NGS somatic QIAsq DNA Breast-Gynecological custom Panel (11 genes)</b>          Genes: <i>AKT, CTNNB1, ERBB2, ESR1, PALB2, PIK3CA, POLE, PTEN, TP53</i></p>

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ISM-BL-004	<i>EGFR</i> exon 20 resistance mutation T790M
ISM-BL-005	<i>EGFR</i> exon 20 resistance mutation C797S
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ISM-BL-007	<i>BRAF</i> hotspot mutations (V600E, V600K, V600R)
ISM-BL-008	<i>PIK3CA</i> hotspot mutations (codons 542, 545 and 1047)
ISM-BL-009	<i>ESR1</i> hotspot mutations (codons 380, 536, 537, 538)

## — SPLICING VARIANTS AND FUSIONS (nCounter Technology)

IST-BL-001	<b>Fusion variant or <i>MET</i> splicing variant single analysis</b> Fusions: <i>ALK</i> , <i>NRG1</i> , <i>NTRK1</i> , <i>RET</i> , <i>ROS1</i> Splicing variants: <i>MET</i> Δ14
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**GERMLINE MOLECULAR  
TESTING ASSOCIATED  
WITH INHERITED  
PREDISPOSITION  
SYNDROMES**

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**MULTIGENIC MOLECULAR PANELS**

<b>MS-G-001</b>	<p><b>NGS QIAseq Inherited Cancer Panel (28 genes, germline)</b></p> <p>Genes: <i>APC, ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CDKN2A, CHEK2, CTNNB1, EPCAM, FANCA, MLH1, MRE11A, MSH2, MSH6, MUTYH, NBN, PALB2, PIK3CA, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, STK11, TP53</i></p>
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**INMUNOHISTOCHEMISTRY (IHC) ASSAYS (Recommended for Lynch syndrome)**

<b>ISI-T-003</b>	<p><b>Mismatch Repair (MMR) status (<i>MLH1, MSH2, MSH6, PMS2</i>)*</b></p>
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*\*Sample required: FFPE Block*

# **MOLECULAR TESTING FOR OPHTHALMOLOGICAL DISEASES**

**MULTIGENIC MOLECULAR PANELS**

OF-BL-001	<b>nCounter Gene Expression Lacrima Dx (24 genes)</b> Genes: <i>B2M, CCL4, CD46, CFB, CTSS, GAPDH, HLA-A, HLA-DMB, HLA-DRB3, IFITM1, IRF7, JAK1, LGALS3, MUC1, MX1, MYD88, OAZ1, PSMB8, RARRES3, SERPING1, STAT1, STAT2, TAP1, TAPBP</i>
OF-BL-002	<b>nCounter Gene Expression Lacrima Extended Dx (96 genes)</b> Genes: <i>ABCf1, ADA, AHR, ALAS1, ARHGDI1B, ATG5, B2M, BAX, BCAP31, BCL10, C14orf166, CASP3, CCL4, CCND3, CCRL1, CD14, CD19, CD46, CD59, CD83, CD99, CEACAM6, CFB, CFH, CFI, CTNNA1, CTSC, CTSS, CXCR2, DUSP4, EEF1G, FADD, FKBP5, G6PD, GAPDH, GUSB, HLA-A, HLA-C, HLA-DMB, HLA-DPA1, HLA-DQB1, HLA-DRA, HLA-DRB1, HLA-DRB3, HPRT1, IFI16, IFI35, IFITM1, IFNAR1, IFNG, IKZF2, IKBKG, IL13RA1, IL32, IL6ST, IL8, IRF5, IRF7, JAK1, LGALS3, LTF, MAPK1, MUC1, MX1, MYD88, NFKB1, NFKBIA, OAZ1, PIGR, PLA2G2A, PLA2, PML, POLR1B, POLR2A, PPIA, PSMB10, PSMB5, PSMB7, PSMB8, PTK2, PYCARD, RARRES3, RUNX1, S100A9, SDHA, SERPING1, SLC2A1, SMAD3, STAT1, STAT2, TAP1, TAPBP, TBP, TUBB</i>

**EXTERNAL SERVICES**

PO-E-004	<b>CentoVision Panel (Inherited Diseases)</b>
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# SERVICES PORTFOLIO



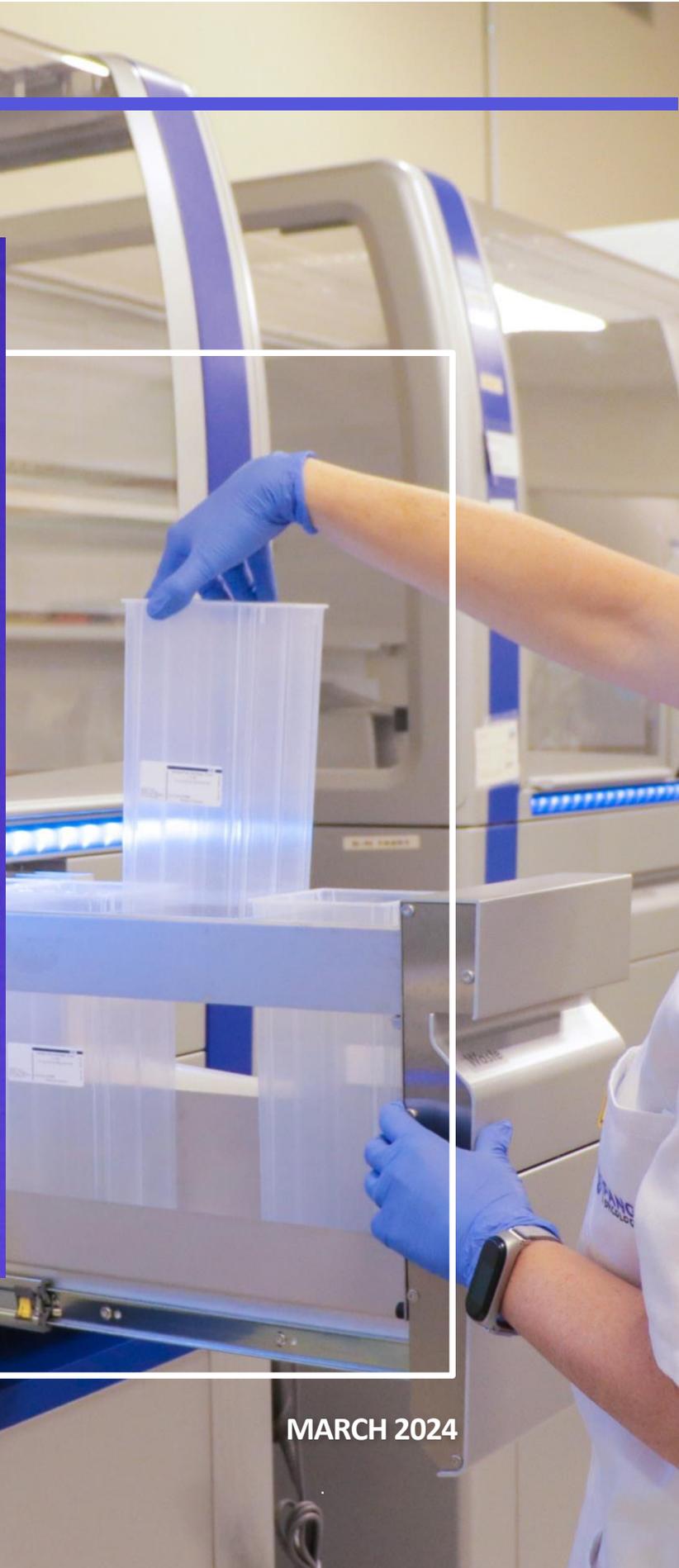
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