

OncolIndx<sup>®</sup> offers comprehensive genomic profiling across solid tumors. Its unique design identifies key actionable insights for personalising cancer treatment.



## OncolIndx<sup>®</sup> in practice

- Challenges of managing complex cancers are addressed through evidence-based treatment choice.
- NGS report enhanced with artificial intelligence and updated clinical studies help identify the most beneficial therapeutic options.
- An easy digital interface to interpret and understand complex cancer genomics.

## Where is OncolIndx<sup>®</sup> needed?

- Cancers progressing on standard-of-care treatment.
- Cancers where personalised treatment could enhance survival.
- Cancers showing resistance to conventional treatment.
- Cancers where combination of target drugs and/or immunotherapy could benefit.

### Clinical and Technical Validation of OncolIndx<sup>®</sup> Assay A Comprehensive Genome Profiling Assay for Pan-Cancer investigations

Alteration	* PPV	* NPV	Accuracy	Specificity	Sensitivity
SNVs	100	100	100	100	100
Small INDELs	100	94.03	97.40	100	95.60
CNA	100	100	100	100	100
Fusions	100	96.43	98.48	100	97.44

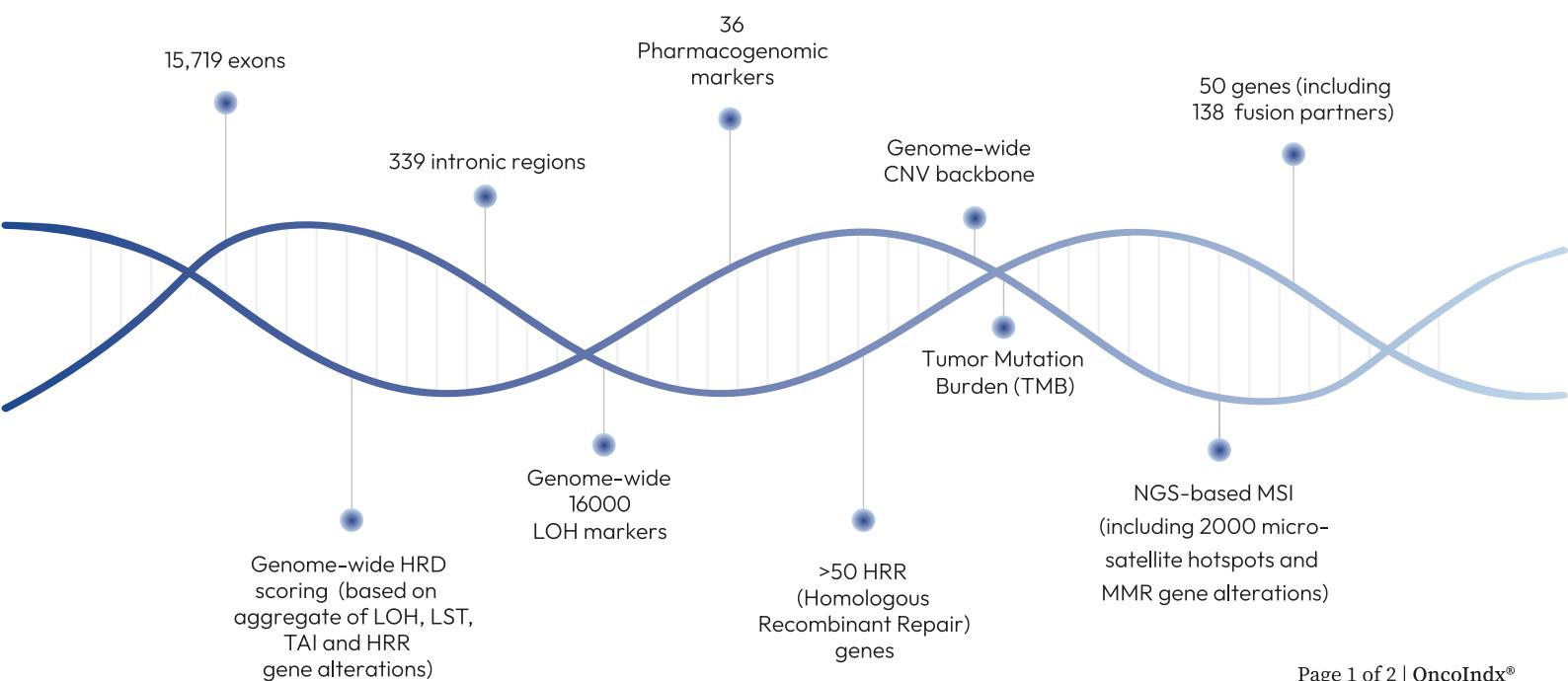
\* NPV: Negative predictive value. PPV: Positive predictive value.

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## Designed for oncologists managing complex, advanced and refractory cancer patients.

- Proprietary panel of 1,000+ genes
- Longitudinal monitoring markers
- PD-L1 through IHC (TBx) / immunofluorescence on CTCs (LBx)
- CTC enumeration
- Traceability to ongoing clinical trials worldwide



## Key Actionable Genes Include:



### Lung

EGFR, ALK, HRAS, NRAS, KRAS, BRAF, TERT, ROS1, MET, RET, NTRK1/2/3, ERBB2, PIK3CA



### Breast

BRCA1, BRCA2, HER2, PIK3CA, MSH6, PMS2, NTRK1/2/3, MLH1, BRAF



### Melanoma

BRAF, KIT, ROS1, NTRK1/2/3



### Bladder

FGFR2/3, NTRK1/2/3



### Colorectal

BRAF, KRAS, NRAS, HRAS, ERBB2, MLH1, MSH2, MSH6, PMS2, NTRK1/2/3



### Ovary

BRCA1, BRCA2, HER2, PMS2, MLH1, MSH2, MSH6, NTRK1/2/3



### Gastrointestinal

ALK, HRAS, HER2, NRAS, KRAS, BRAF, TERT, ROS1, MET, RET, NTRK1/2/3



### Thyroid

ALK, HRAS, NRAS, KRAS, BRAF, TERT, RET, NTRK1/2/3



### Prostate

BRCA1, BRCA2, MLH1, MSH2, MSH6, PMS2, AR, NTRK1/2/3

## Features of Oncolndx report

- In accordance to international guidelines.
- Clear, unambiguous and actionable genomic findings.
- Based on highly reliable and internally curated databases.

## Techincal Specification

Specimen	Liquid (CSF, Plasma, Pleural Fluid)	Tissue
Panel Size	3 MB	3 MB
Sample Type	cfDNA (TagFlexLB/Norgen/PaxGene Tube) + 1 tube of 8.5 ml Peripheral Whole Blood (~8-10 days of washout period required if patient under therapy e.g., Chemotherapy, Radiation, Immunotherapy, etc.)	1 FFPE tissue block with 20-30 % tumor content (not more than 3-4 months old)*
Sample Collection	Peripheral blood using butterfly needles/ 22 gauge syringe	FFPE tissue block
Sample volume/quantity	8.5 ml (1 tube)	1 FFPE block with 20-30% tumor content
cfDNA/DNA input amount	> 1ng	> 1ng
Sequencing platform	Illumina	Illumina
Read Length	2 x 150bp	2 x 150bp
Mean Sequencing Depth	10000x	2000x
Sequencing Chemistry	Hybrid capture	Hybrid capture
Indications	Solid Tumor	Solid Tumor
Analysis and Reporting	iCare™- AI based analysis platform with active reporting options	

Oncolndx® is a well-validated NGS test to extract therapy information from tumor DNA. It is possible on both tissue as well as blood sample.

1000+ genes panel

This NGS test includes 1000+ genes of targetable/ therapeutic significance.

>99% on target coverage

The test has a mean coverage of 98-100% across all tested genes.

2000x / 10000x mean depth

The test is performed at a mean depth of 2000x for tissue samples and 10000x for blood.



**1Cell.Ai™**

Ai-Powered Precision Oncology



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