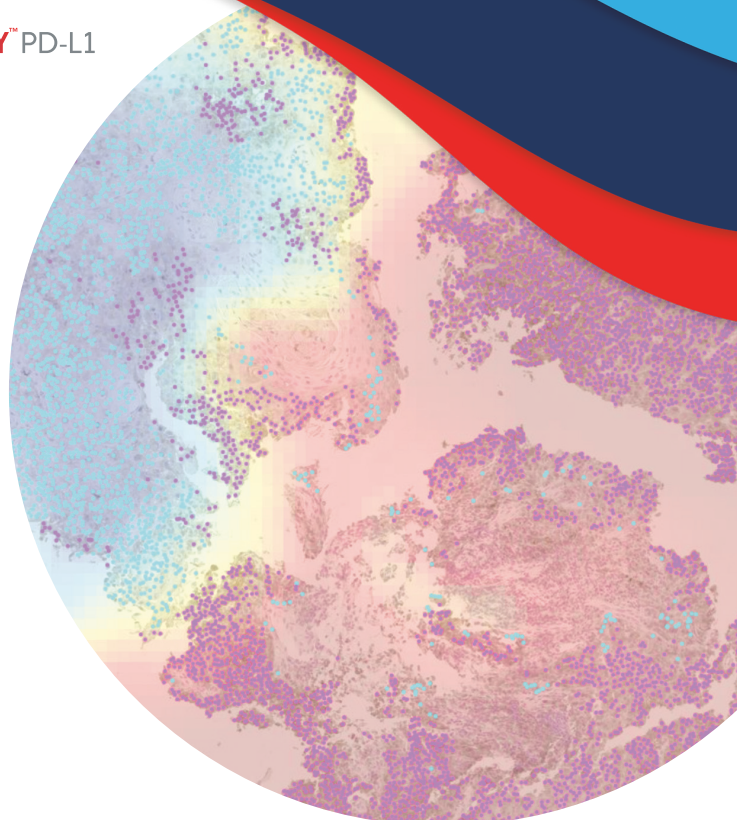




A MORE PRECISE

PD-L1 TEST FOR NSCLC¹

With an AI-backed digital pathology algorithm, part of the new Guardant Galaxy™ technology suite



>20% IMPROVED PD-L1 DETECTION*¹

AI-powered PD-L1 scoring accurately classified 22.2% of NSCLC patients initially identified as Tumor Proportion Score (TPS) <1%¹



Scan to read
full *European
Journal of
Cancer* article

ACCURATE SCORING HELPS TO PREDICT RESPONSE TO IMMUNOTHERAPY

In **KEYNOTE-189**, overall response rates ranged from 33.1-62.1% depending on TPS²



NOW VALIDATED FOR MULTIPLE CANCERS

Including breast, colorectal, gastric, hepatobiliary, pancreatic, prostate, and urothelial cancers

1. [https://www.ejancer.com/article/S0959-8049\(22\)00215-5/fulltext](https://www.ejancer.com/article/S0959-8049(22)00215-5/fulltext)
2. [https://www.annalsofoncology.org/article/S0923-7534\(21\)01146-7/fulltext](https://www.annalsofoncology.org/article/S0923-7534(21)01146-7/fulltext)

*Compared to manual pathologist interpretation in the most challenging cases in NSCLC.
AI: Artificial Intelligence; NSCLC: non-small cell lung cancer

Guardant360 TissueNext panel

Covers clinically relevant biomarkers, including TMB, MSI status, and PD-L1. Results are reported 2-3 weeks from sample receipt.

Point Mutations (SNVs) and Deletion Variants (Indels) (84 Genes)								Amplifications (20 Genes)		Fusions (12 Genes)		MSI status Qualitative result
AKT1	ALK	APC	AR	ARAF	ARID1A	ATM	BRAF	AR	BRAF	ALK	MET	
BRCA1	BRCA2	CCND1	CCND2	CCNE1	CDH1	CDK4	CDK6	CCND1	CCND2	BRAF	NTRK1	
CDK12	CDKN2A	CHEK2	CTNNB1	DDR2	EGFR	ERBB2	ESR1	CCNE1	CDK4	EGFR	NTRK2	
EZH2	FANCA	FBXW7	FGFR1	FGFR2	FGFR3	GATA3	GNA11	CDK6	EGFR	FGFR1	NTRK3	
GNAQ	GNAS	HNF1A	HRAS	IDH1	IDH2	JAK2	JAK3	ERBB2	ESR1	FGFR2	RET	
KEAP1	KIT	KRAS	MAP2K1	MAP2K2	MAPK1	MAPK3	MET	FGFR1	FGFR2	FGFR3	ROS1	
MLH1	MPL	MSH6	MSH2	MTOR	MYC	MYCN	NF1	KIT	KRAS			
NFE2L2	NOTCH1	NPM1	NRAS	NTRK1	NTRK2	NTRK3	PALB2	MET	MYC			
PDGFRA	PIK3CA	PMS2	PTEN	PTPN11	RAD51D	RAF1	RB1	MYCN	PDGFRA			
RET	RHEB	RHOA	RIT1	ROS1	SMAD4	SMO	STK11	PIK3CA	RAF1			
TERT [†]	TP53	TSC1	VHL									

NSCLC guideline-recommended genes shown in bold.

[†]Includes *TERT* promoter region

TMB
Mutations per
Megabase

PD-L1 status
AI-powered PD-L1
test now available
for multiple cancers[^]



Guardant Access Program

- Patient contacted if out-of-pocket costs for each genomic test >\$100
- Financial assistance offered if eligible
- Patient given opportunity to cancel test if financially burdened



For comprehensive support, call **855.698.8887** or email **clientservices@guardanthealth.com**

[^]Some cases may require conventional determination of PD-L1 status without AI.

Important Note: Guardant360 TissueNext and Guardant Galaxy PD-L1 tests were developed, and their performance characteristics determined by Guardant Health Clinical Laboratory in Redwood City, CA, USA, which is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) as qualified to perform high complexity testing. These tests have not been cleared or approved by the US FDA.