

ACTDrug[®] +

Actionable Insights for Timely Treatment Options



101 Genes

Assesses clinically relevant gene alterations aligned with FDA-approved and NCCN guideline-recommended targeted therapies, supporting informed first-line treatment decisions



Whole Coding-Region Design

Comprehensive variant detection and identification of clinically actionable alterations



Resistance Mutation Insight

Provides genomic insights into variants associated with treatment resistance



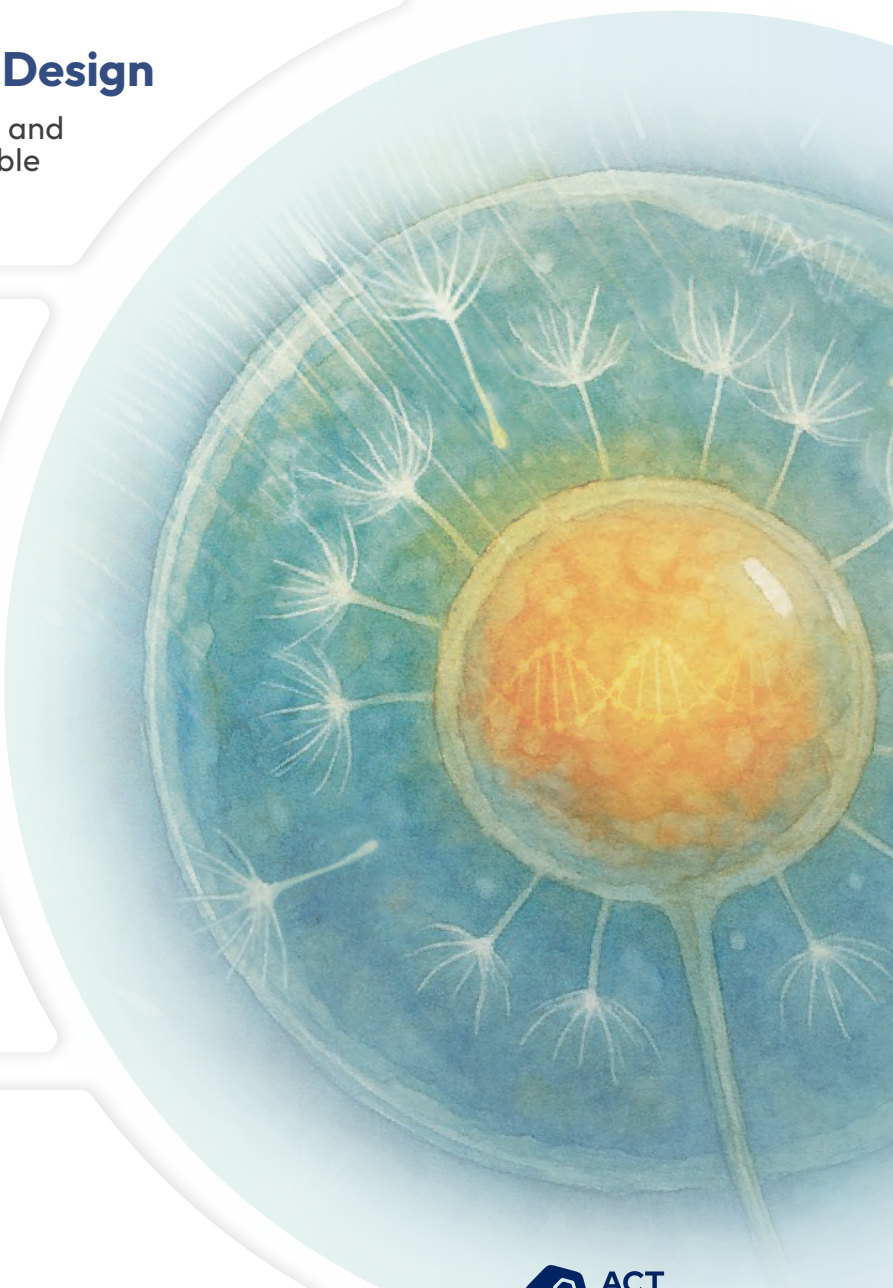
RNA-based NGS Fusion Detection

Enables assessment of known and novel gene fusion events, supporting identification of patients who may benefit from targeted therapy



7 Working Days

Rapid turnaround time



The Specification of The New ACTDrug® +

ACTDrug® +	Number of Genes Tested	101
	Types of Gene Mutations Analyzed	• Single nucleotide variants (SNVs); Small insertions and deletions (InDels) • Copy number alterations (CNAs) and Fusion genes • Large genomic rearrangement (LGR) • Microsatellite Instability (MSI)
	Sensitivity*	SNV and Indels: 100%, CNAs: 95%, LGR (BRCA1/2): 100%, Fusion genes 100%, MSI: 100%
	Specificity*	SNV and Indels: 100%, CNAs: 100%, LGR (BRCA1/2): 94%, Fusion genes 100%, MSI: 94%
	Specimen Requirements	FFPE Tumor Tissue 5-20 unstained sections (5 µm/slide, surface area ≥ 125 mm ²) 1 H&E-stained slide (5 µm)
	Turnaround Time	7 Working Days (from receipt of qualified samples at our CAP-accredited laboratory)

*Analytical performance was established under defined validation conditions

Gene List

◆ SNV / InDel ■ CNA ● Fusion

AKT1 ◆■	AKT3 ●	ALK ◆■●	APC ◆■	AR ◆■●	ATM ◆■
ATR ◆■	ATRX ◆■	BARD1 ◆■	BRAF ◆■●	*BRCA1 ◆■	*BRCA2 ◆■
BRIP1 ◆■	CCND1 ◆■	CCNE1 ◆■	CDK12 ◆■	CDK4 ◆■	CDK6 ◆■
CDKN2A ◆■	CDKN2B ◆■	CHEK1 ◆■	CHEK2 ◆■	CTNNB1 ◆■	EGFR ◆■●
ERBB2 ◆■●	ERBB3 ◆■	ERBB4 ◆■●	ERG ●	ESR1 ◆■●	EZH2 ◆■
FANCA ◆■	FANCL ◆■	FBXW7 ◆■	FGFR1 ◆■●	FGFR2 ◆■●	FGFR3 ◆■●
H3F3A (H3-3A) ◆■	HDAC2 ◆■	HIST1H3B (H3C2) ◆■	HRAS ◆■	IDH1 ◆■	IDH2 ◆■
JAK1 ◆■	JAK2 ◆■	KEAP1 ◆■	KIT ◆■●	KRAS ◆■	MAP2K1 ◆■
MAP2K2 ◆■	MDM2 ◆■	MDM4 ◆■	MET ◆■●	MLH1 ◆■	MRE11 ◆■
MSH2 ◆■	MSH6 ◆■	MTAP ◆■	MTOR ◆■	MUTYH ◆■	MYC ◆■
MYCN ◆■	NBN ◆■	NF1 ◆■	NF2 ◆■	NRAS ◆■	NRG1 ●
NRG2 ●	NTRK1 ◆■●	NTRK2 ◆■●	NTRK3 ◆■●	NUTM1 ●	PALB2 ◆■
PAX8 ●	PDGFRA ◆■	PIK3CA ◆■●	PIK3R1 ◆■	PMS2 ◆■	POLD1 ◆■
POLE ◆■	PPARG ●	PPP2R1A ◆■	PTEN ◆■	RAD51B ◆■	RAD51C ◆■
RAD51D ◆■	RAD54L ◆■	RAF1 ●	RB1 ◆■	RET ◆■●	ROS1 ◆■●
RSPO2 ●	SMAD4 ◆■	SMARCB1 ◆■	STK11 ◆■	*TERT ◆■	TFE3 ●
TPRSS2 ●	TP53 ◆■	TSC1 ◆■	TSC2 ◆■	VHL ◆■	

*genes provide large genomic rearrangement (LGR) information
#TERT promoter is also covered in the ACTDrug+ panel

Actionable and Emerging Biomarkers

Cancer Type	Clinically actionable biomarkers with FDA-approved therapies and NCCN guideline support	Potential biomarkers
Pan-Solid Tumor	BRAF V600E, NTRK1/2/3 fusions, RET fusions, MSI	
Lung Cancer	ALK, BRAF, EGFR, ERBB2, FGFR1/2/3, KRAS, MET, NRG1, ROS1	KEAP1, MTAP, PIK3CA, RB1, STK11, TP53
Breast Cancer	AKT1, BRCA1/2, ERBB2, ESR1, FGFR1/2/3, PALB2, PIK3CA, PTEN	ATM, BRIP1, BRAD1, CCND1, CCNE1, CHECK2, HRAS, KRAS, MYC, NRAS, RB1, TP53
Colorectal	BRAF, ERBB2, HRAS, KRAS, NRAS, PIK3CA, POLE, POLD1	APC, ALK, HER2, MET, MLH1, MSH2, MSH6, MUTYH, PMS2, PTEN, ROS1, SMAD4, STK11, TP53
Gastric	ERBB2	ATM, ATR, EGFR, FGFR2, MET, MLH1, MSH2, MSH6, PIK3CA, PMS2, POLE, TP53
Biliary Tract	ERBB2, FGFR2, IDH1, KRAS	FGFR1, FGFR3, IDH2, MDM2, MET, MTAP, PIK3CA, PTEN, TP53
GIST	KIT, FGFR1/2/3, NF1, PDGFRA	KRAS, PIK3CA
Thyroid	ALK, HRAS, KRAS, NRAS, PAX8, TERT	ATM, FGFR3, PIK3CA, PTEN, TP53
Brain	ATRX, CDKN2A/B, IDH1/2, EGFR, H3-3A, H3C2, TERT	ALK, CDK4/6, FGFR1/2/3, MET, TSC1/2, MDM2, MTAP, NF1, PIK3CA, POLE, PTEN, PDGFRA, ROS1

NCCN Clinical Practice Guidelines in Oncology, Non-Small Cell Lung Cancer Version 3.2026; Breast Cancer Version 5. 2025; Colon Cancer Version 5. 2025; Gastric Cancer Version1. 2026, Biliary Tract Cancer Version 2. 2025; Thyroid Carcinoma Version 1. 2025; Central Nervous System Cancers Version3. 2025; Oncologist. 2025 Mar 10;30(3):oyae357; Hum Pathol. 2025 Aug;162:105881; Cancer Discov. 2020 August ; 10(8): 1174–1193.; Genes (Basel). 2023 Jun 28;14(7):1364.; Cancers (Basel). 2024 Nov 19;16(22):3870; Clin Oncol (R Coll Radiol). 2025 Jan;37:103661; J Cancer Res Clin Oncol. 2023 Jan;149(1):467–481; Trends Mol Med. 2025 Dec;31(12):1089–1102.; Am J Pathol. 2025 Mar;195(3):437–452.; ESMO Open. 2022 Jun 10;7(3):100505.; ESMO Open. 2018 Apr 6;3(3):e000335.; Eur Thyroid J. 2025 Jul 7;14(4):e250024.; Neuro Oncol. 2024 Oct 10;27(2):331–337. Nat Med. 2024 Mar;30(3):730–739

ACTDrug® Breast

Number of Genes Tested

16

Turnaround Time

6 working days

Types of gene Mutations Analyzed

SNV/ InDels, CNAs (including BRCA1/2 LGR), Fusion genes, MSI

Gene List

◆ SNV / InDel

■ CNA

● Fusion

AKT1 ◆■

#BRAF ◆■●

*BRCA1 ◆■

*BRCA2 ◆■

ERBB2 ◆■●

ESR1 ◆■●

FGFR1 ◆■●

FGFR2 ◆■●

FGFR3 ◆■●

NTRK1 ◆■●

NTRK2 ◆■●

NTRK3 ◆■●

PALB2 ◆■

PIK3CA ◆■●

PTEN ◆■

RET ◆■●

*Indicates genes with large genomic rearrangement (LGR) coverage.

#genes also provide exon-skipping alteration information

ACTDrug® GI

Number of Genes Tested

20

Turnaround Time

6 working days

Types of gene Mutations Analyzed

SNV/ InDels, CNAs, Fusion genes, MSI

Gene List

◆ SNV / InDel

■ CNA

● Fusion

#BRAF ◆■●

ERBB2 ◆■●

FGFR1 ◆■●

FGFR2 ◆■●

FGFR3 ◆■●

HRAS ◆■

IDH1 ◆■

IDH2 ◆■

KIT ◆■●

KRAS ◆■

NF1 ◆■

NRAS ◆■

NTRK1 ◆■●

NTRK2 ◆■●

NTRK3 ◆■●

PDGFRA ◆■

PIK3CA ◆■●

POLD1 ◆■

POLE ◆■

RET ◆■●

#genes also provide exon-skipping alteration information



**Newly Diagnosed
Advanced/ Meta-
Static Patients**



**Biomarker
Associated
Drug
Information**



**6~7 Working
Days
Turnaround
Time**



**Bioinformatic
Analysis
Based on Curated
International
Databases**

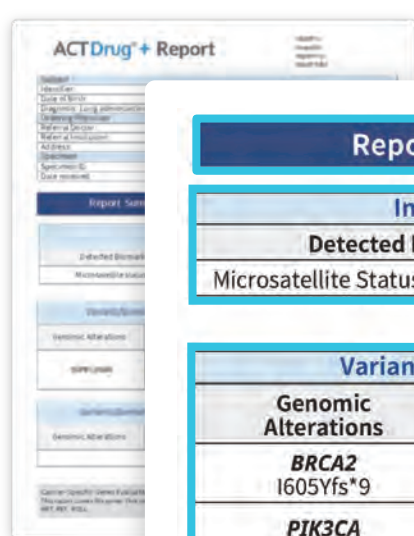


**Easy-to-
Interpret
Medical
Report**

CAP

**Quality
Assured
Testing**

ACTDrug® + Report



1

Actionable summary of the detected variant and associated therapies

2

MSI status reported to support clinical decision-making related to immunotherapy

Report Summary for Actionable Variants/Biomarkers

Immune Checkpoint Inhibitor (ICI) Related Biomarkers

Detected Biomarker Status	Corresponding Therapies
Microsatellite Status (MSI): Microsatellite stable	-

S Sensitive R Resistant

Variants/Biomarkers with Clinical Significance (Target Therapy)

Genomic Alterations	Evidence Level 1, 2 (FDA-approved, NCCN guideline)	Evidence Level 3A, 3B, 4 (Others)
BRCA2 I605Yfs*9	S Olaparib, Talazoparib	S Niraparib, Rucaparib
PIK3CA R88Q	S Alpelisib, Capivasertib, Inavolisib	S Everolimus
PIK3CA Amplification	-	S Everolimus

Variants/Biomarkers with Clinical Significance (Hormone Therapy)

Genomic Alterations	Evidence Level 1, 2 (FDA-approved, NCCN guideline)	Evidence Level 3A, 3B, 4 (Others)
Not detected		

Cancer-specific genes evaluated

FDA-Approved Biomarkers Assessed by this Assay: *ALK, BRAF, EGFR, ERBB2, KRAS, MET, RET, ROS1*.

3

Variant reporting and classification are based on levels of evidence, prioritizing treatment-related evidence supported by international publications and guidelines. List out the cancer-specific genes according to the patient's cancer type and clinical guidelines.

5

Highlight FDA-approved biomarkers assessed by this assay to enable clear and rapid interpretation for clinical decision-making.

4

Icons are used to indicate drug sensitivity or resistance for faster interpretation.

ACT Genomics Co., Ltd.

Member of Delta Group



www.actgenomics.com



service@actgenomics.com



+886-2-2795-3660



©2026 ACT Genomics Co., LTD. All Rights Reserved. It is possible that the test may return with no abnormal mutation identified for certain genes or part of the test results may not be available due to the technical limitations of the test itself and/or an individual's genetic differences and/or intra- or inter-tumoral heterogeneity even the test has been conducted under standard process. This material (including the test and the results) therefrom are intended for educational purposes only for the use of healthcare professionals and do not replace independent professional judgement. This material (including the test and the results) shall at no time be deemed a diagnostic or medical treatment recommendation to any individual. No representation, warranty, express or implied, is made as to, and no reliance should be placed on the fairness, accuracy, completeness or correctness of the information or opinions which may be contained herein. The test takers shall always consult their healthcare professionals for any enquiries of clinical interpretation of the test results.

ACT GENOMICS® and ACTDrug® are registered trademarks of ACT Genomics Co., LTD.

WW_DM_20260306_ACTDrug_Brochure_V01