

# Access to Cancer Testing and Treatment

The Access to Cancer Testing and Treatment program, also known as **Project ACTT**, provides high quality cancer genomic testing to patients with metastatic or recurrent lung, breast, or colorectal cancer. Formerly known as **Canadian Early Access Program**, **Canexia Health** initially launched the program in 2019 to provide oncologists and patients with access to a **blood-based, multiplex hotspot mutation panel** known as **Follow It**, using next-generation sequencing (NGS).



## Follow It<sup>®</sup>

Follow It is a clinically actionable, blood-based circulating tumour DNA (ctDNA) NGS multiplex panel that tests for 146 hotspots and 23 exons in 30 cancer genes that are known to be important in the prognosis and treatment of multiple solid adult tumours. Follow It provides a detailed clinical report, which includes interpretations of results, available clinical trials, and current treatment options based on the molecular profile of the tumour.



### Benefits To Your Patients


- Minimally invasive sample collection.
- Rapid, actionable results.



### Benefits To Your Practice

- Rapid turnaround time for patient treatment information.
- Identification of targeted therapies for your patient, and potentially avoiding treatments that might not be clinically beneficial.

## Follow It<sup>®</sup> Clinical Report:


Follow It<sup>®</sup> Report

**Patient Name:** Jane Doe  
**Report ID:** 544

**Date of Birth:** 11/05/1957  
**Surgical ID:**

**Patient ID:** pt-DNA-10599-CG001QV40Run26-22  
**Panel:** Full

**PATIENT INFORMATION**

Patient Name: Jane Doe  
Date of Birth: 11/05/1957 Sex: Female  
Care Card #: 010101 Province of Issue: BC  
Diagnosis: None  
Reason for Referral: Diagnostic Evaluation  
Previous Molecular Tests: N/A  
Test Requested: FOLLOW IT  
Date of Receipt: None  
Date of Report: 2019-08-02 11:41:22

**HEALTHCARE PROVIDER INFORMATION**

Referring Physician: Nathanael Down  
Institution: ABC General Hospital  
Address:  
Phone: (999) 000-999 Fax:  
Pathologist: Nigel Nigel  
Institution: ABC General Hospital  
Address:  
Phone: (999) 000-999 Fax:

**SPECIMEN INFORMATION**

Specimen Collection Date: None  
Specimen Source: Blood  
Specimen Type: Plasma  
Primary Site of Tumour: Endometrium  
Histologic Type: Endometrioid carcinoma  
Sample: DNA-10599-CG001QV40Run26-22

This test is an amplicon based hotspot next-generation sequencing assay (NGS) that interrogates clinically actionable gene alteration in circulating tumour DNA extracted from plasma. The test results, interpretations and clinical trials included in this report are provided in the context of a primary cancer type as reported by the referring physician.

**CONSIDERATIONS**

\* POLE mutations are associated with an ultra-mutated tumour phenotype and an excellent prognosis in endometrial endometrial carcinoma.  
\*\* May require immunotherapy in the non-metastatic setting.  
\*\*\* PMS2 and PTEN mutations. In the context of POLE-mutated endometrial endometrial carcinoma, PMS2 and PTEN mutations are considered secondary events and are not currently associated with prognostic or therapeutic significance.

**SUMMARY OF TEST RESULTS**

Gene	cDNA change	Amino Acid Change	Exon	Allelic Ratio (%)	Therapeutic Implication	Level of Evidence	Clinical Trials Available
POLE	C.1251G>T (NM_026271.5)	V411L	15	19.4	Associated with an excellent prognosis in endometrial endometrial carcinoma. May respond to immunotherapy in the non-metastatic setting.	Tier II D • Metastatic Tier III D • Metastatic	0
PMS2A	C.268G>A (NM_026271.5)	R88C	2	28.1	Alterations in PMS2A are frequent (70%) in POLE-mutated endometrial carcinoma and are considered secondary events and are not currently associated with prognostic or therapeutic significance.	Tier II D • Metastatic	0
PTEN	C.399G>A (NM_003144.4)	R130D	5	24.9	Alterations in PTEN are frequent (94%) in POLE-mutated endometrial carcinoma and are considered secondary events and are not currently associated with prognostic or therapeutic significance.	Tier II D • Metastatic	0
TP53	C.638G>A (NM_005045.5)	R213D	6	3.1	NA (see other biology section)	NA	NA

TABLE 1: Mutations Present

## Important Information about ACTT

<b>Tumor Types Tested</b>	<ul style="list-style-type: none"> <li>Breast cancer</li> <li>Lung cancer</li> <li>Colorectal cancer</li> </ul>
<b>Eligible Patient Population</b>	<ul style="list-style-type: none"> <li>Patients with suspected or known relapsed disease where ctDNA testing can detect resistance mutations.</li> <li>Patients with metastatic disease where ctDNA testing can detect resistance mutations.</li> <li>Patients with tumour not easily amenable to biopsy.</li> <li>Very ill patients with metastatic disease.</li> </ul>
<b>Turnaround Time</b>	10 working days
<b>Specimen Type</b>	Blood
<b>Specimen Collection</b>	<ul style="list-style-type: none"> <li>BC, Ontario, and Saskatchewan patients will take their TRF provided by the referring oncologist to a LifeLabs location,</li> <li>Quebec Patients will contact Genolife at 1-844-440-5454 to provide their TRF and schedule an appointment at a Genolife sample collection clinic.</li> </ul>
<b>Pricing</b>	Testing is available at NO COST for 2000 patients till March 31st, 2021
<b>Further Information</b>	If you would like further information, please send us an email <a href="mailto:test@canexiahealth.com">test@canexiahealth.com</a>

## About Canexia Health

Canexia Health (formerly Contextual Genomics) makes high-quality cancer genomic information accessible with our clinically-validated assays, informatics, and support. Our suite of genomics-based cancer tests is clinical actionable and cost-effective, designed to improve cancer treatment and monitoring. With our extensive scientific experience, specialized genomics-based tests, and support from pharmaceutical and diagnostics partners, we are leading the shift towards precision oncology.

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[www.canexiahealth.com](http://www.canexiahealth.com)



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## Genes And Associated Cancer Types

Gene	Therapeutic Options
AKT1	●
ALK	● ● ● ● ●
AR	● ●
BRAF	● ● ● ● ●
CTNNB1	●
DDR2	● ● ● ● ●
EGFR	● ● ● ● ●
ERBB2	● ● ● ● ●
ESR1	● ● ●
GNA11	●
GNAQ	●
GNAS	●
HRAS	● ●
IDH1	●
IDH2	●

Gene	Therapeutic Options
KIT	● ● ● ● ●
KRAS	● ● ● ● ●
MAP2K1(MEK1)	● ● ● ● ●
MAP2K2(MEK2)	● ● ● ● ●
MET	● ● ● ● ●
NRAS	● ● ● ● ●
PDGFRA	● ● ● ● ●
PIK3CA	● ● ● ● ●
POLE	●
PTCH1	● ● ● ● ●
PTEN	● ● ● ● ●
RET	● ● ● ● ●
ROS1	● ● ● ● ●
SMO	● ● ● ● ●
TP53	● ● ● ● ●

Key	Health Canada Approved Drugs	Off-Label Drugs	Clinical Trial	Resistance
	●	●	●	●

Cancer	Associated Genes
Breast	AKT1, ERBB2, ESR1, PIK3CA
Colorectal	BRAF, KRAS, NRAS, PIK3CA
Endometrial	CTNNB1, PIK3CA, POLE
Melanoma	BRAF, KIT, NRAS
GIST	BRAF, PDGFRA, KIT
Glioma	BRAF, IDH1, IDH2
NSCLC	BRAF, EGFR, ERBB2, KRAS, MET
Sarcoma	GNAS, IDH1, IDH2

