

REGIONAL PATHOLOGY SERVICES Date: May 2022

## **Lab Alert: Genetics News**

Regional Pathology Services provides genetic testing and expertise in molecular genetics, cytogenetics, FISH, microarray, Human Leukocyte Antigen (HLA), Next-Generation Sequencing (NGS), and forensics and paternity testing.

#### **NEW TESTING**

**FLTTKD FLT3 TKD Mutation** 

> This test detects mutations in the FLT3 tyrosine kinase domain (TKD) in blood or bone marrow from AML patients. FLT3 TKD mutations are targetable

with FLT3 inhibitors such as gilteritinib and midostaurin.

**STPP** Solid Tumor Precision Panel

> Replaces the Colorectal Cancer Panel, Gastrointestinal Stromal Tumor (GIST) Panel, Lung Cancer Panel, and Melanoma Panel

**TMBO** Tumor Mutational Burden

> The Tumor Mutation Load (TML) Assay is a targeted next-generation sequencing assay that is designed for providing an accurate assessment of TMB (mutations/Mb). Studies have shown that tumors that have a high tumor mutation burden/load (TML or TMB) potentially have a better response to immunotherapy.

## **FEATURED TESTING**

## **ACUTE MYELOID LEUKEMIA (AML) FISH PANEL**

The Human Genetic Laboratory's AML FISH panel now includes NUP98 [11p15] as recommended by the World Health Organization (WHO). The addition of this probe increases the detection of cryptic rearrangements and helps characterize prognostic outcomes.

FISH-Acute Myeloid Leukemia (AML) Panel **FAML** 

> 8 centromere, 20q12, CBFB [16q22], D7S486 [7q31] / 7 centromere, EGR1 [5q31], KMT2A (MLL) [11q23], MECOM (EVI1) [3q26.2], NUP98 [11p15], PML :: RARA [t(15;17)], RARA [17q21], RUNX1T1::RUNX1 (ETO / AML1) [t(8;21)]

#### CHRONIC MYELOID LEUKEMIA (CML) FISH PANEL

The Human Genetic Laboratory's CML FISH panel when ordered without chromosomes, often requested for follow-up studies, has a new reflex algorithm.

- BCR::ABL1 negative: no reflex will occur
- BCR::ABL1 positive: reflex to MECOM, TP53, 8 centromere, 19p13 /19q13

### FORENSICS AND PATERNITY TESTING

The Human DNA Identification Laboratory utilizes industry standard methods compliant with CODIS (U.S. Combined DNA Index System) to determine person of origin in various biological samples. Application of our methodology can be used to resolve identity in the following circumstances:

- Paternity / parentage
- Physical evidence for law enforcement agencies and private attorneys
- Patient tissue / body fluid misidentification

For more information on forensic and paternity testing visit our website:

https://www.reglab.org/services/paternity-and-forensic-testing/



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# **NEW TEST BUILD INFORMATION:**

Test Code	Test Name	ResultCode	ResultName	CPT
<u>FLTTKD</u>	FLT3 TKD Mutation	FLTKD	FLT3 TKD Mutation	81479
<u>STPP</u>	Solid Tumor Precision Panel	STPPR	Solid Tumor Precision Panel	81445
<u>TMBO</u>	Tumor Mutational Burden	TMB	Tumor Mutational Burden	81479

### **DISCONTINUED TESTING**

The following testing has been discontinued and when available a replacement test code has been indicated.

<b>Deactivated Test</b>	Deactivated Test	Replacement Test
Code	Name	Code
50GCP / 50GCPO	50 Gene Cancer Panel	<u>STPP</u>
CCMP	Colorectal Cancer Panel	<u>STPP</u>
EGFR	EGFR Mutation	No replacement
GIST	Gastrointestinal Stromal Tumor (GIST) Panel	STPP
KRASM	KRAS Mutation Detection	No replacement
LCMP	Lung Cancer Mutation Panel	<u>STPP</u>
MMP	Melonoma Mutation Panel	<u>STPP</u>
PIK3CA	PIK3CA Mutation Detection	No replacement
TP53P	TP53 Mutation on fixed tissue	TP53 *fresh tissue only*

### **UPDATED REQUEST FORMS**

- Hematopathology Molecular Oncology/Consultation Request form
- Solid Tumor FISH/Molecular Oncology Test Request Form
- Oncology Test Request Form

Request forms are found on our website and are also available through the supply portal. If you need access to the supply portal your account manager can assist.