Prospective Registry Of MultiPlex Testing

What is the purpose?

PROMPT provides patients, physicians, and researchers with an opportunity to share information about multiplex genetic testing so that we can all better understand the implications of these genetic mutations.

Updates from Researchers

Using a platform housed at **PatientCrossroads**, individuals can enroll in PROMPT and provide data by completing questionnaires about their personal and family health histories. PROMPT can also be used as a way to receive updates from physicians and researchers.

PROMPT is an online registry for patients and families who:

- Have had multiplex genetic testing
- Have been found to have a genetic variation which may be linked to an increased risk of cancer

How is personal information shared?

Patients can choose whether to share identifiable data with researchers, or whether to share only de-identified data. **PatientCrossroads** uses sophisticated security features to ensure that physicians and researchers get the data they need without breaches in confidentiality.

What is MultPlex Testing?

Multi-gene Panel Testing is a newer form of genetic testing. These panels look for mutations in several different genes all at once. All of the genes on the panels have been tied to an increased risk of cancer, but the risks associated with some of the genes are better known than the risks associated with others.

Register for PROMPT in 3 easy steps!

① Log on to your computer and go to www.promptstudy.org

② Click on the ***Register Now!*** Button

③ Fill in the required information and Start PROMPT Survey

For questions, please contact PROMPT@uphs.upenn.edu

Have a smartphone?

You can quickly access the PROMPT Registration website by scanning the QR Code below.

Supporting Organizations

MEMORIAL SLOAN-KETTERING CANCER CENTER · ESTABLISHED 1884 ·

BASSER RESEARCH CENTER *for* BRCA
Abramson Cancer Center
Penn Medicine

MAYO CLINIC

PROMPT

Prospective Registry Of MultiPlex Testing