

Accessing Genetic Testing in Australia

Genetic testing can give you important health information. Obtaining a genetic test in Australia can be complicated. Depending on your personal circumstances, genetic testing can be accessed through different pathways. Here we try to simplify the process for you.

In Australia, clinicians use your family history and other factors to determine the eligibility for a publicly-funded genetic test. There's a very high threshold to qualify for a publicly-funded genetic test, so many people are not eligible and therefore can't find out their genetic risk.

A new analysis from Invitae, a leading medical genetics company, reveals that

up to two-thirds of Australian breast cancer patients with hereditary gene variants

(such as BRCA, PALB2, Lynch Syndrome and more) would not qualify for genetic testing due to the nation's eligibility criteria and potential wait times and would be unable to access what can be life-changing knowledge.¹

Pink Hope's mission is to empower people to be able to access a genetic test at the earliest possible point whether you are considering taking a test proactively to better understand your risk or if you have cancer yourself and are looking for answers or more personalised treatments.

We are here to help you.



INVITAE

**pink
hope**



pinkhope.org.au

Pathway for Patients



Cancer Services

Public Genetic Testing

www.hgsa.org.au/resources/new-south-wales

www.genetics.edu.au/genetic-services/cancer-genetics-clinics

Private Genetic Testing

www.hgsa.org.au/asgc/find-a-genetic-counsellor

www.invitae.com/australia

www.eugenelabs.com

References: 1. Invitae, data on file 2. Haverfield EV et al. Physician-directed genetic screening to evaluate personal risk for medically actionable disorders: a large multicenter cohort study. BMC Med. 2021 Aug 18;19(1):199. *Invitae's study was a retrospective analysis of Australian guidelines applied to a US-based study population. Prospective studies are needed in the Australian population to evaluate the real-world efficacy of the guidelines.