





At-A-Glance

- Provides personalized cancer risk information and access to education
- Supports tailored medical recommendations such as increased screening
- Facilitates connection to third-party telehealth genetic counseling
- Integrates seamlessly into your existing workflow

WORKFLOW SOLUTIONS SUPPORTING PATIENTS AND PROVIDERS AT EVERY STEP



DIGITAL HISTORY COLLECTION AND ASSESSMENT

Suite of digital tools collect and analyze patient medical and family history and weigh against medical guidelines



GENETIC TESTING

Ordered through Ambry for qualified individuals



POST-TEST GENETIC COUNSELING

Made available to all patients



PRE-TEST EDUCATION

How genetic testing can guide personalized, proactive healthcare



RESULTS DELIVERY

Results delivered to the provider, and in most cases, to the patient, using the CARE platform



DOCUMENTATION

Transparency at each step, improving patient and provider experience



CANCER PREVENTION AND EARLY DETECTION BEGIN WITH KNOWING YOUR RISK.

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DID YOU KNOW?

*At least 42% of newly diagnosed cancer cases in the US (~797,000) are potentially avoidable⁴

*excludes non-melanoma skin cancer

People are
diagnosed with

Million¹ cancer each year in
the United States

~100/0
OF ALL CANCER IS
HEREDITARY

\$26,000,000,000° cost savings from early detection of cancer

References

- 1. cdc.gov/chronicdisease/about/costs/index.htm
- 2. Z. Kakushadze, et al, in September 2017—Cornell University
- 3. cancer.gov/about-cancer/causes-prevention/genetics
- 4. cancer.org/content/dam/cancer-org/research/cancer-facts-and-statistics/annual-cancer-facts-and-figures/2021/cancer-facts-and-figures-2021.pdf

Shape the future of healthcare with solutions focused on proactive, personalized care.





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