

Screen for dozens of cancers with a simple blood draw



Elevate member care with the Galleri® test

The Galleri test incorporates a first-of-its-kind technology for multi-cancer early detection blood testing. With a simple blood draw, the Galleri test screens for many of the deadliest cancers that don't have recommended screenings today, like pancreatic, ovarian, liver, and more. 1,2,3* The test helps health plans support its members to be more proactive with their health.

The Galleri test does not detect a signal for all cancers, and not all cancers can be detected in the blood. False positive and false negative results do occur. The Galleri test identifies DNA in the bloodstream shed by cancer cells and does not predict future genetic risk for cancer. The Galleri test should be used in addition to screening tests recommended by a healthcare provider.

Cancer's effect on health plans and their members

Cancer has historically been one of the most challenging areas for health plans to address. The Galleri test can help do more to screen for cancer.

Innovation



STATUS QUO

Nearly 70% of deaths are caused by cancers that don't have recommended screenings.1,2**



Up to 99% of cancers impacting members ages 50+ can be screened for by the Galleri test.4

Cost of treatment



STATUS QUO

Of commercial cancer claims above \$100,000, 81% are caused by cancers without recommended screening.4



Modeled data shows that adding Galleri led to a 50% shift of late-stage cancers to earlier stages, potentially resulting in a 24% reduction in direct medical spending on members with metastatic diagnoses.4,5***

Proactive intervention



STATUS QUO

Later-stage cancer is treated quickly, aggressively, and in a costly manner,⁶ leaving little time to engage members in lowercost, best-in-class treatment and care management options.



WITH GALLERI

By screening for more cancers earlier, modeled data shows that adding the Galleri test to usual care could potentially result in a 50% reduction in the proportion of cases diagnosed in Stage III and Stage IV.5***

^{*}Sensitivity in study participants with - Pancreatic cancer: 83.7% overall (61.9% stage I, 60.0% stage II, 85.7% stage III, 95.9% stage IV). Ovarian cancer: 83.1% overall (50.0% stage I, 80.0% stage II, 87.1% stage III, 94.7% stage IV). Liver/bile duct cancer: 93.5% overall (100% stage I, 70.0% stage II, 100% stage III, 100% stage IV).

^{**}Assumes screening is available for all prostate, breast, cervical, and colorectal cancer cases and 43% of lung cancer cases (based on the estimated proportion of lung cancers that occur in screen-eligible individuals older than 40 years).

^{***}Results based on modeled data, with an earlier version of Galleri added to usual care, and costs analyzed by GRAIL.





All cells, healthy and cancerous, release DNA into the bloodstream. But DNA from cancer cells is different from DNA from healthy cells and act like a signal of cancer. Using next-generation DNA sequencing, machine learning, and AI, the Galleri test can screen for a cancer signal in the blood and predict the tissue or organ associated with the cancer signal to help guide next steps.^{3,7}



The Galleri test is supported by robust clinical data.

Dependable

0.5% false positive rate³

Found in study participants without cancer.

Impactful

76.3% sensitivity

In cancers responsible for every 2 of 3 US cancer deaths.^{3,9}* Overall test sensitivity was 51.5%.

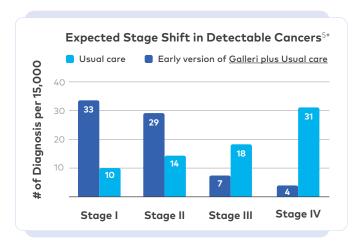
Actionable

93.4% accuracy

In predicting Cancer Signal Origin, found in study participants with cancer diagnosis after a Cancer Signal Detected test result.

^{*}The group of cancers responsible for two-thirds of annual US cancer deaths included anus, bladder, colon/rectum, esophagus, head and neck, liver/bile duct, lung, lymphoma, ovary, pancreas, plasma cell neoplasm, and stomach.





Cancer won't wait. Screen members for more.

Adding a multi-cancer early detection test to recommended cancer screenings allows your health plan to go further, screening for dozens of deadly cancers, including many that don't have recommended screening.²

*Late stage defined as stages III + IV. MCED results based upon fast tumor growth model. Usual care represents real-world cancer diagnostic processes (e.g., screening, incidental detection, symptomatic workup) as captured by SEER. Results based on performance data from Liu, et al. Ann Oncol. 2020 modeled with stage-specific incidence and survival of all invasive cancers diagnosed in persons aged 50–79 between 2006 and 2015 from the US Surveillance, Epidemiology, and End Results (SEER) program. Performance of MCED test in a state-transition model (interception model) when added to usual care. Assumes long-term screening results with optimized screening interval.

Galleri is a highly-ranked health benefit.10

#1

Employees rank the Galleri test as the #1 health benefit¹⁰*

#3

Medicare beneficiaries rank the Galleri test as the most important supplemental benefit after vision and dental^{11**}

*Market research conducted by Ipsos Group. Study demographics included 1,000 respondents who were full-time employees working at companies with at least 3,000 employees: 15% of sample aged 65+, 70% of sample aged 50-64; 15% aged under 50 with cancer risk factors. Respondents were a representative mix across gender, region, race, ethnicity.

A comprehensive and member-oriented experience

GRAIL offers several implementation options that do not impact medical policy:

- Health Plan Administrator (HPA) collaboration
- Medicare Advantage supplemental benefit
- · Employee benefits offering pilot
- · Fully insured commercial pilot



From helping eligible members schedule their blood draw to facilitating follow-up consultations to explain their results, the Galleri test experience is designed to support your members at each step:



Awareness and education with customized materials



Online test
request, followed by
in-person blood draw
(Test requires physician
approval. On-site blood
draw events are also
an option.)



Return of results to provider within 10-14 business days of receipt at GRAIL lab



Support services for Cancer Signal Detected test results

in the event a cancer signal is detected

^{**}Market research Study demographics included 1,000 respondents who were Medicare Advantage eligible age 63-80. Respondents were a representative mix of gender, region, race and ethnicity, and Income.

"Promoting early screenings, building awareness about preventative techniques, and developing strategic partnerships remain the most effective measures for cancer prevention."

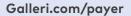
- Health Payer Intelligence, February 22, 2022¹²

Be the innovative partner your clients and members demand by making available the Galleri multi-cancer early detection test.



Contact us to find out more about the Galleri test for payers, including comprehensive marketing campaigns and the support for integrating this benefit into your organization.







payer@grailbio.com

Important Safety Information

The Galleri test is recommended for use in adults with an elevated risk for cancer, such as those age 50 or older. The test does not detect all cancers and should be used in addition to routine cancer screening tests recommended by a healthcare provider. The Galleri test is intended to detect cancer signals and predict where in the body the cancer signal is located. Use of the test is not recommended in individuals who are pregnant, 21 years old or younger, or undergoing active cancer treatment. Results should be interpreted by a healthcare provider in the context of medical history, clinical signs, and symptoms. A test result of No Cancer Signal Detected does not rule out cancer. A test result of Cancer Signal Detected requires confirmatory diagnostic evaluation by medically established procedures (e.g., imaging) to confirm cancer. If cancer is not confirmed with further testing, it could mean that cancer is not present or testing was insufficient to detect cancer, including due to the cancer being located in a different part of the body. False positive (a cancer signal detected when cancer is not present) and false negative (a cancer signal not detected when cancer is present) test results do occur. **Rx only.**

Laboratory / Test Information

The GRAIL clinical laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) and accredited by the College of American Pathologists. The Galleri test was developed — and its performance characteristics were determined — by GRAIL. The Galleri test has not been cleared or approved by the Food and Drug Administration. The GRAIL clinical laboratory is regulated under CLIA to perform high-complexity testing. The Galleri test is intended for clinical purposes.

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