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Second Quarter 2017 Highlights

- Presented new results from a large, community-based, multi-center clinical validation study conducted at Kaiser Permanente at the American Urological Association (AUA) 2017 Annual Meeting. The results confirmed that the Oncotype DX GPS test is a strong independent predictor of prostate cancer-specific death and disease progression (metastases) at 10 years in men with localized prostate cancer.
- Data from two Oncotype DX GPS test analyses based on a prospective, multi-center, 1,200-patient study were also presented at AUA, including results that represent the first-ever prospective validation of a tissue-based molecular marker in prostate cancer. The second analysis, presented at AUA and published online in *Urology*, reinforced that the GPS test significantly increases use of and persistence on active surveillance.
- Initiated an early access Clinical Utility Program for select centers around the country in preparation for the commercial launch of the Oncotype DX AR-V7 Nucleus Detect™ test in collaboration with Epic Sciences.
- The 15th St. Gallen International Breast Cancer Conference Expert Panel published updated guidelines endorsing the Oncotype DX Breast Recurrence Score test for guiding treatment decisions on adjuvant chemotherapy in both node-negative and node-positive patients (up to three nodes).
- Breast Cancer Research and Treatment published positive five-year clinical outcomes results from the PlanB study led by the West German Study Group (WSG) in 93 centers across Germany. The study showed that women with Recurrence Score® results of 11 or less who were treated with hormonal therapy alone had excellent outcomes with 94 percent disease-free survival rates at five years despite having high-risk disease by traditional parameters.
- Presented results from eight studies across breast, prostate and kidney cancers that provide additional evidence of the value of Oncotype DX tests in predicting clinically meaningful endpoints and outcomes across cancer types at the 2017 American Society of Clinical Oncology (ASCO) Annual Meeting in Chicago.

This fact sheet contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Forward-looking statements are subject to risks and uncertainties that could cause actual results to differ materially, and reported results should not be considered as an indication of future performance. These risks and uncertainties include, but are not limited to: our business model; the regulation of our tests; the applicability of clinical study results to actual outcomes; our ability to independently develop and commercialize and collaborate with companies to commercialize new tests and expand into new markets domestically and internationally; the risk that we may not obtain or maintain sufficient levels of reimbursement, domestically or abroad; competition; unanticipated costs or delays in research and development efforts; our ability to obtain capital when needed; and the other risks and uncertainties set forth in our filings with the Securities and Exchange Commission, including the risks set forth in our most recent Annual Report filed on Form 10-K and our subsequently filed Quarterly report(s) filed on Form 10-Q. These forward-looking statements speak only as of the date hereof. Genomic Health disclaims any obligation to update these forward-looking statements.

NOTE: The Genomic Health logo, Oncotype, Oncotype DX, Recurrence Score, DCIS Score, Oncotype SEQ, Liquid Select, Genomic Prostate Score, Oncotype DX AR-V7 Nucleus Detect and Oncotype IQ are trademarks or registered trademarks of Genomic Health, Inc. All other trademarks and service marks are the property of their respective owners.