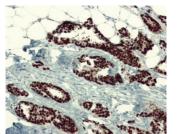


Patient Name : DOE-TEST, JANE-TEST

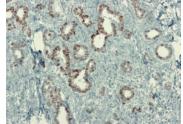
Patient Rec. #: Date of Birth: 01/01/1980 Age: 46 Sex: Female Date Collected: 09/13/22 Date Received: 09/13/22 Date Reported: 09/15/22

Specimen Number: S16-005555 UP ID: Ordering Client: THE BEST DOCTOR Ordering Physician: THE BEST DOCTOR Client Address: 7200 W. Camino Real Boca Raton, FI 33433 Telephone: (561) 453-1234 Fax:(561) 453-1238 CC Physician:

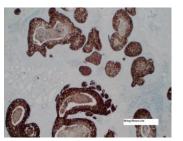
BREAST PATHOLOGY REPORT



Estrogen Receptor 73%



Progesterone Receptor 27%



HER2 = 3+

Specimen - Right Breast **DIAGNOSIS: RIGHT BREAST CORES of Infiltrating Ductal Carcinoma, not** otherwise specified/atypical medullary type and in-SITU Ductal Carcinoma, Comedo Type. Modified Scarff-Bloom-Richardson Score: 8-9 Tubule: 3 **Nuclear Pleomorphism: 3**

Mitotic activity: 2-3 Total score: 8-9

Microscopic Description: Therapeutic Biomarkers Antibody/Tests Results Interpretation: - POSITIVE - 73% ER by IHC PR by IHC - POSITIVE - 27% HER2 by IHC - POSITIVE - 3+ *HER2 IHC Results: (0/ 1+/ 2+/ 3+) Complete membrane staining: 100% **Uniform staining: (Present)** Homogeneous, dark circumferential pattern: (Present) Fixative: Formalin Duration of Fixative: 4 to 12 hours Sample adequate for analysis: Yes Control Results: External (high, low level, negative protein expression) and internal control preformed as expected.

ER (clone 6F11) / PR (clone 1A6): Breast carcinoma receptor markers. Performed on 10% formalin fixed tissue with UltraView detection system and CC1 heat induced antigen retrieval. Semi quantitative assessment of the percentage of positive tumor cells with nuclear staining.

HER2 (clone 4B5) Performed on 10% formalin fixed tissue with UltraView detection system and CC1 heat induced antigen retrieval. Semi quantitative assessment of the tumor cells' membranous staining pattern and intensity, using a scoring method of 0-3+.

GROSS DESCRIPTION:

Specimen is received in formalin and consists of several core(s) of tan gray tissue, measuring 10 cm in aggregated length and 0.2 in average thickness. Entirely submitted in one cassette.

Electronically Signed by: UP Pathologist M.D. Date: 09/13/2016

The tests utilizing analyte-specific reagents (ASR) were developed and their performance characteristics determined by Alliance Labs as required by CLIA '88 regulations. They have not been cleared or approved for specific uses by the ed for clinical purposes. * Phone Number: (561)453-1234 U.S. Food and Drug Administration (FDA). The FDA has determined that such clearance or approval is not necessary. These tests are used Alliance Laboratories * 7200 W. Camino Real, ste. 330 Boca Raton, FI 33433 Fax Number: (561)453-1238