

Ms. PARMINDER KAUR

PID NO: P33723514299560

Age: 42 Year(s) Sex: Female Reference: SELF

Sample Collected At:

Dr Vinay Kumar Chopra Kos Diagnostic Lab, 6349/i, Nicholson Road, Ambala Cantt, Hry 133001. Processing Location: - Metropolis Healthcare Ltd, Unit No409-416, 4th Floor, Commercial Building-1, Kohinoor Mall.Mumbai-70

VID: 230111501886119

Registered On: 22/11/2023 12:45 PM Collected On: 20/11/2023 12:44PM Reported On: 26/11/2023 08:57 PM



INTERNATIONAL & NATIONAL SUBSPECIALITY PATHOLOGY

Breast Pathology Dermato pathology Gastrointestinal Pathology Genitourinary Pathology Gynecologic Pathology Head & Neck Pathology Hematolymphoid Pathology Hepatobiliary Pathology Neuropathology

Paediatric & Perinatal Pathology **Pulmonary Pathology** Renal Pathology Soft tissue Pathology Transplant Pathology (Renal & Hepatic)

Chief Scientific Officer, Senior Consultant Oncopathologist

Dr Kirti Chadha

In - House Faculty

Senior Consultants

Dr Anuradha Murthy

Dr Amita Joshi

Dr Meenal Hastak

Dr Leena Naik Dr Vikas Kavishwar

Consultants

Dr Barodawala S.M.

Dr Kunjal Lila

Dr Shital Munde

Dr Shraddha More

IMMUNOHISTOCHEMISTRY PANEL (PREDICTIVE & PROGNOSTIC MARKER ASSESSMENT)

Case Summary

CASE NO. :23MLI17315

Clinical Notes Ca Breast.

Original H and E Right MRM - Histomorphology is consistent with Invasive

Ductal Carcinoma - Breast, Grade III.

Gross Examination Received 11 paraffin blocks labelled as HP-4279/23 A to J

RESULT:

Estrogen Receptor (ER)

Interpretation	Negative
Proportion	-
Intensity	-
Quick Score	-

Progesterone Receptor (PgR)

Interpretation	Negative
Proportion	-
Intensity	-
Quick Score	-

PROPORTION & INTENSITY SCORING - ALLRED SCORE

Proportion	Intensity	
0 = No nuclear staining	0 = No staining	
1 = <1% nuclei staining	1 = weak staining	
2 = 1-10 % nuclei staining	2 = moderate staining	
3 = 11-33 % nuclei staining	3 = strong staining	
4 = 34-66% nuclei staining		
5 = 67-100 % nuclei staining		



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METROPOLIS
HIST@XPERT
GLOBAL EXPERTISE IN SUB SPECIALTY SOLUTIONS

C-erbB-2/ Her 2 neu

Score value	0
Score result	Negative

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RESULT INTERPRETATION OF Her 2/neu. (ASCO/CAP 2018).

Score	C-erbB-2 (HER-2 protein) overexpression assessment	Staining pattern	
0	Negative	No observed staining or incomplete, faint / barely perceptible membrane staining in <= 10% of invasive tumour cells.	
1+	Negative	Incomplete, faint / barely perceptible membrane staining in > 10% of invasive tumour cells.	
2+	Equivocal	Weak to moderate complete membrane staining observed in >10% tumor cells.	
3+	Positive	Complete / intense circumferential membrane staining in > 10% of invasive tumour cells.	

Processing : Manual / Automated (Roche Ventana Benchmark XT 715030, Roche Ventana Benchmark XT 716016, Intellipath)

CAP Recommendation:

- Testing ER, PgR and Her2/neu status on all newly diagnosed invasive breast cancers (primary site and/or metastatic site), and whenever appropriate, repeat testing in patients with a known breast cancer diagnosis who now present with a local or distant recurrence.
- Fixation time: Ideally should be 6-72 hours for ER/PgR and Her2/neu.
- Cold Ischemic time: It is the time from tissue removal to initiation of fixation and should be less than or equal to one hour.



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Note:

ER / PR

- Considering an ER and PgR test performed by an IHC assay as positive if at least one percent
 of the tumor in the sample tests positive, which helps predict whether a patient is likely to
 benefit with endocrine treatment.
- Patients whose breast cancers have very low levels of staining(Quick score 2) may still benefit
 from adjuvant endocrine therapy. It is reasonable for oncologists to discuss the pros and cons of
 endocrine therapy with patients whose tumors contain low levels of ER by IHC (one percent to
 ten percent weakly positive cells) and to make an informed decision based on available
 information.
- Patients with tumours scoring less than 2 are regarded as ER / PgR negative and have a negligible chance of response to endocrine therapy.

C-erbB-2

- For equivocal cases (2+) Fluorescent in-situ hybridization (FISH) analysis for Her-2 gene amplification can be used as a secondary test.
- Prolonged cold ischemic and fixation times may lead to a false negative Her2/neu result.
- If the cold ischemic and fixation times are not known, negative Her2/neu results should be interpreted with caution.
- If the initial HER2 test result in a core needle biopsy specimen of a primary breast cancer is negative, a new HER2 test may be ordered on the excision specimen.

References:

- Fitzgibbons PL, Murphy DA, Hammond ME, et al. Recommendations for validating estrogen and progesterone receptor immunohistochemistry assays. Arch Pathol Lab Med. 2010;134;930-935
- Wolff AC, Hammond EH, Allison KH, et al. Human Epidermal Growth Factor Receptor 2 Testing in Breast Cancer: American Society of Clinical Oncology/College of American Pathologists Clinical Practice Guideline Focused Update. Arch Pathol Lab Med. 2018;142(11):1364-1382.

Quality Assurance:

- The external (positive and negative) and internal control (wherever applicable) show appropriate reactivity.
- Detection system: MACH-2 Universal HRP Polymer Detection / Ultraview Dab detection Kit.
- Clone: ER SP1; Company: Thermoscientific/Lab Vision.
- Clone: PgR PgR636; Company: Dako.
- Clone: C-erbB-2 SP3; Company: Thermoscientific/Lab Vision.

Dispatch Summary:

- 1. Blocks that are submitted are enclosed with the report.
- 2. Stained slide is archived.
- 3. Case images are available on request.

-- End of Report --



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