

**INVASIVE PAPILLARY DUCTAL CARCINOMA OF THE BREAST: A CASE REPORT****Amani S Hadi<sup>1</sup>, Gamal Abdul Hamid<sup>2\*</sup>, Refaat Al-Areque<sup>3</sup> and Wafa Abdullah<sup>4</sup>**<sup>1</sup>National Oncology Center, Aden, Yemen,<sup>2</sup>Faculty of Medicine, Aden University.<sup>3</sup>Alamal Oncology Center, Taiz, Yemen.<sup>4</sup>Alamal Oncology Center Taiz, Yemen.**\*Corresponding Author: Gamal Abdul Hamid**

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**ABSTRACT**

The general rate of invasive papillary carcinoma (IPC) is uncommon, representing for less than 1-2 % of invasive breast cancers. They are most generally observed in postmenopausal females and uncommon in males. Invasive papillary carcinomas are low grade tumors originating from large or dilated ducts. They are made out of all around outlined solid nodules of monotonous neoplastic cell separated by network of fibrovascular cores, IPC is a remarkable sort of breast cancer and regarded of whether it is in-situ or invasive, it has brilliant prognosis. We presenting two cases of invasive papillary carcinoma in male and female; A case of 55years postmenopausal female who presented with history of left breast mass, which this mass noticed after trauma same site for 1year ago the mass gradually increase in size no tenderness, no signs of inflammation. Excisional biopsy was performed and specimen was histopathology diagnosed as invasive papillary carcinoma, left MRM was performed and histopathology diagnosis confirmed and without residual tumor seen in submitted slides and all submitted lymph nodes were free of tumor infiltration (0/14). IHC show ER and PR negative with HER-2 positive. The second case 70 years male presented with right breast mass and history of post-trauma since one year back with gradual increase in size, right radical mastectomy done and histopathology diagnosed as invasive papillary carcinoma, IHC was done ER and PR positive with HER2- negative.

**KEYWORDS:** invasive papillary carcinoma, Low grade tumors, excellent prognosis.**INTRODUCTION**

Breast cancer is the commonest malignancy of females in Yemen and all over the world and the second leading cause of death due to cancer among females.<sup>[1-3]</sup>

Papillary lesions of the breast have assessed in wide range going from benign intra ductal papilloma (with or without atypia) to papillary carcinoma in situ and invasive papillary carcinoma.<sup>[4]</sup> Among this group of lesions, solid papillary carcinoma (SPC) establishes an unmistakable element clinically and morphologically. Solid papillary carcinoma are Low grade tumors originating from large or dilated ducts. They are made out of all around encompassed solid nodules of monotonous neoplastic cells separated by network of fibrovascular cores.<sup>[5-8]</sup>

These lesions usually present as subareolar mass and /or nipple discharge, most frequently in elderly women and represent less than 2% of breast carcinomas in females.<sup>[9]</sup>

It is predominately seen in postmenopausal women.<sup>[10]</sup> Breast cancer in male is rare it account for 0.6 of all

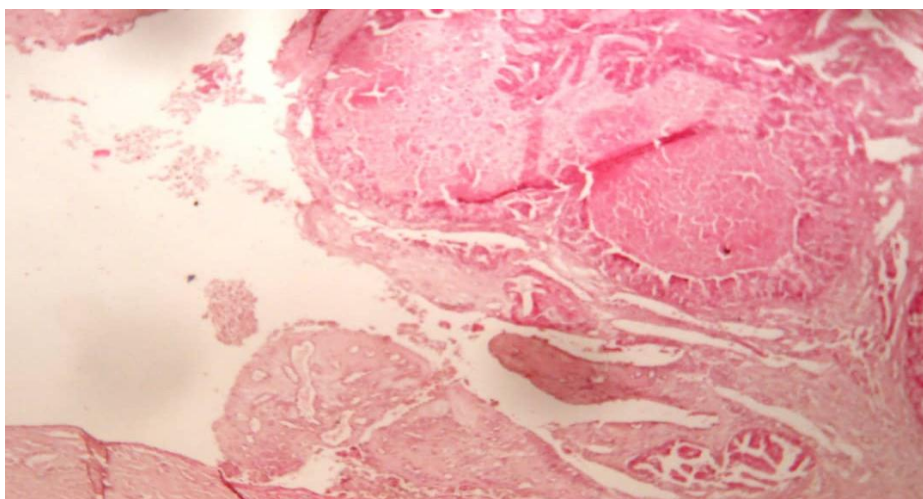
breast carcinoma and less than 1% of all malignancy in men.<sup>[11]</sup>

**CASE PRESENTATION**

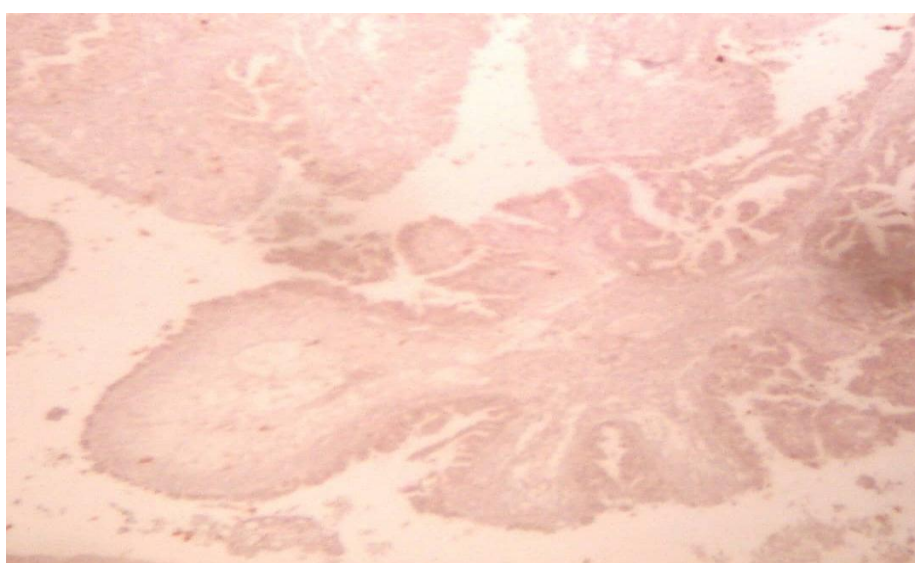
Case1: A 55years postmenopausal female who presented with history of left breast mass, which noticed after trauma in same site for 1year ago the mass gradually increase in size without tenderness or signs of inflammation.

ultrasonography revealed heterogeneously hypoechoic mass measure 4.5cmx5.1cm.

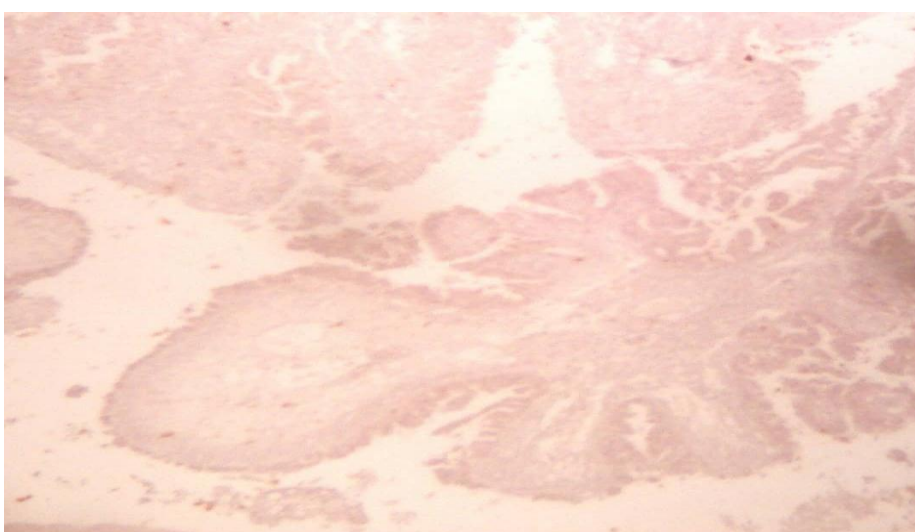
True cut biopsy was done for histopathology shows atypical hyperplasia, DCIS cannot excluded. excision biopsy was performed and specimen was histopathology diagnosed as invasive papillary carcinoma, left MRM was performed and histopathology diagnosed as no residual tumor seen in submitted slides and all submitted lymph nodes are free of tumor (0/14). IHC done ER and PR negative with HER-2 positive. (microphotograph- 1,2,3).



**Microphotograph-1: Papillary structure aligned around fibrovascular core(H&E,10X).**



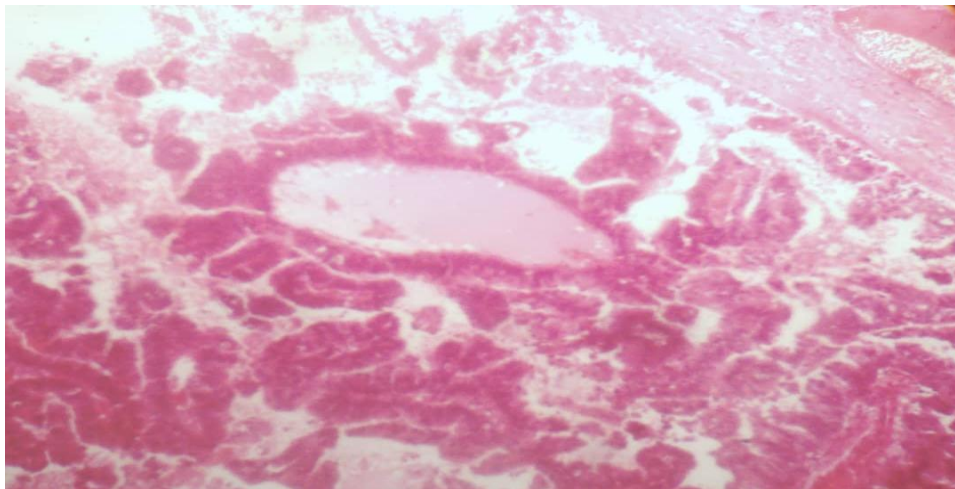
**Microphotograph-2: Immunihistochemstery show ER negativity.**



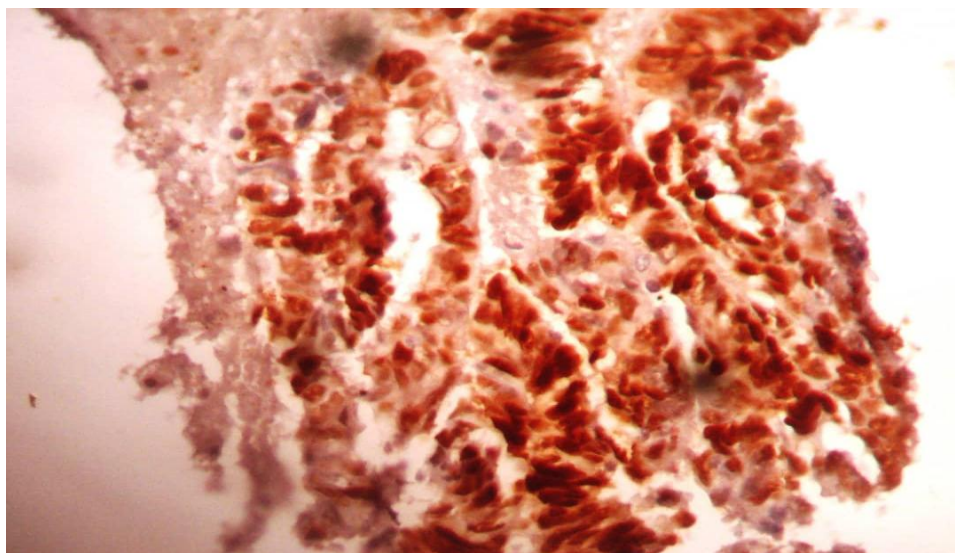
**Microphotograph 3: Immunihisochemistry show PR negativity.**



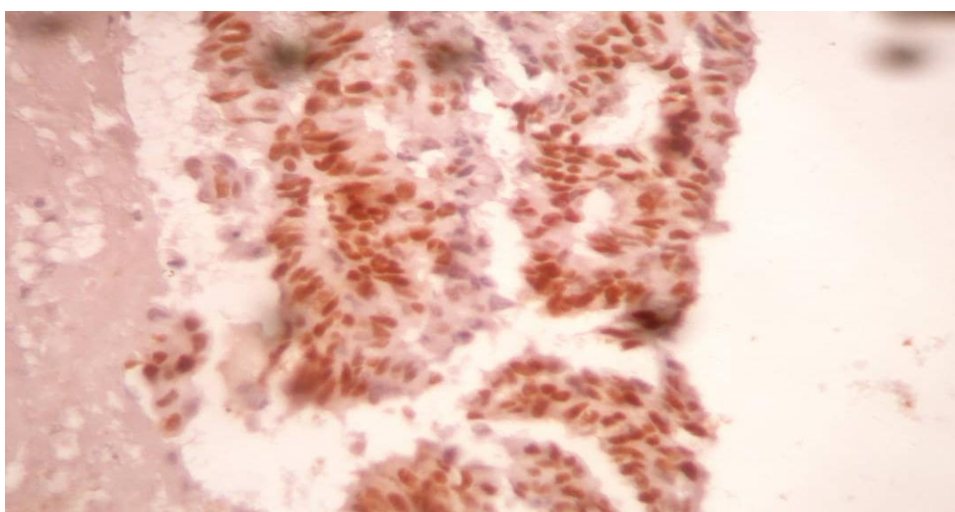
**Case 2:** A 70 years male presented with history of right breast mass post-trauma before one year with increase gradually in size, right radical mastectomy done and histopathology diagnosed as invasive papillary carcinoma, IHC was done ER and PR positive with HER-2 negative. (Microphotograph 4,5,6,7).



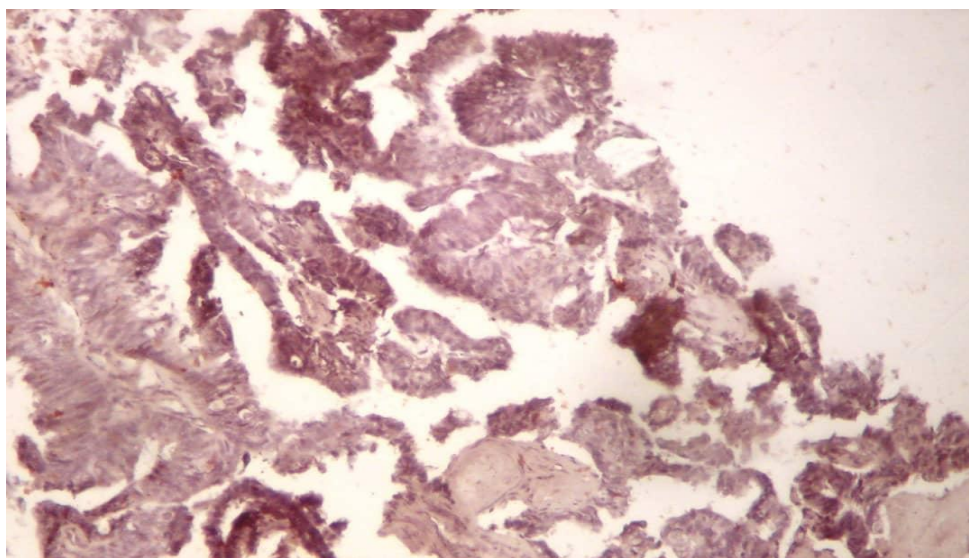
**Microphotograph 4:** Papillary arrangement of neoplastic cells within cyst like space (Hematoxyline and Eosion,40X).



**Microphotograph 5:** Immunihistochemistry showing ER Positivity.



**Microphotograph 6:** Immunihisochemistry showing PR Positivity.



**Microphotograph 7: Immunihistochemistry, HER-2 Negative**

## DISCUSSION

Invasive papillary carcinoma of breast cancer are rare, accounting for less than 1% of invasive breast cancer. They occur most frequently in postmenopausal women in the 6<sup>th</sup> to 8<sup>th</sup> decade of life and non white women. IPC in man is usually reported among those of an older age group (67 to 84 years).<sup>[12]</sup> More than 75% of the cases include a ductal carcinoma in situ (DCIS) component [10-13]. Lymph node involvement and distant metastasis are uncommon and has favorable prognosis.<sup>[13,14]</sup> All malignant papillary proliferation of breast lack an intact myoepithelial cell layer (MCL) within the papilla or at the periphery of the tumor, which is an important feature allowing distinction from benign intraductal papilloma.<sup>[15]</sup> In assessing the presence of complete myoepithelial layer, P63 is often used as an adjunct to assess the presences and distribution of myoepithelial cells in papillary neoplasm of the breast.<sup>[16]</sup> Other immunohistochemical markers, such as estrogen/progesterone receptor, C-erbB2, and Ki-67, provide prognostic information.<sup>[17]</sup> Papillary breast carcinoma is usually estrogen/progesterone receptor-positive and C-erbB2 negative as demonstrated by immunohistochemical results in male present case.<sup>[18]</sup> These molecular expression correspond with luminal A-like subtype, which is associated with a lower recurrence rate and longer disease-free interval.<sup>[17]</sup> Variable management strategies are considered when dealing with this rare form of breast cancer. Treatment options for the breast can involve breast conserving surgery in the form of wide local excision, with or without adjuvant RT, or mastectomy.<sup>[19-21]</sup>

Though there is no definite guidelines for management of IPC in male, Grabowski *et al.* suggested that surgery is the mainstay of treatment, which can be either conversion or mastectomy. Since the prognosis of IPC is excellent with low locoregional and distant recurrence rates, mastectomy is usually not necessary, unless it is technically unavoidable.<sup>[22]</sup> There is controversy

regarding axillary lymph node clearance. Sentinel node biopsy may be an excellent alternative to full axillary dissection in patients with IPC and associated invasive carcinoma.<sup>[23]</sup> There is no clear indication in adjuvant chemotherapy even in hormone responsive cases. The addition of hormonal treatment does not appear to have impacted the outcome.<sup>[22]</sup> Recently Fayanju *et al.* reviewed that patient having DCIS or micro invasion disease in association with IPC were more likely to receive radiotherapy and tamoxifen.<sup>[24]</sup>

## CONCLUSION

We highlight this rare variant of invasive breast cancer presenting with unusually large tumor size. Even though it commonly occurs in postmenopausal woman, it can rarely appear in perimenopausal age group.

Invasive papillary carcinoma is very rare entity in male but it has favorable prognosis. Clinical and radiological features are not specific and there is chance of under diagnosis in aspiration cytology. Therefore correct histopathological diagnosis and invasion status is required. The mainstay of treatment is surgical resection with adjuvant hormonal therapy. Accurate diagnosis of papillary lesion remains challenging only by *hematoxylin and eosin stain*, IHC for myoepithelial cells is a useful diagnostic tool in differentiating benign from malignant lesion with implication on management. Regardless of its invasive nature, it carries an excellent prognosis and thus, awareness of the entity is important to avoid over treatment.

Adjuvant therapy is still controversial and prognosis is excellent with 10-years survival rate for IPC is 100%, the recurrence-free survival rate is 96% and 77% at two and ten years, respectively.

## REFERENCES

1. Bawazir A.A, Abdul Hamid G., Morales G: Available data on cancer in southeastern governorates of Yemen. Eastern Mediterranean Health Journal, 1998; 1(4): 107-113. [https://www.researchgate.net/publication/328228933\\_Gynecological\\_malignancies\\_in\\_Aden\\_Yemen\\_An\\_overview\\_of\\_4\\_years](https://www.researchgate.net/publication/328228933_Gynecological_malignancies_in_Aden_Yemen_An_overview_of_4_years).
2. Abdul Hamid G., Tayeb M.S, Bawazir A.A. Breast cancer in south-east republic of Yemen. East Mediterr Health JEMHJ, 2001; 7: 5.
3. Al-Kahiry W, Omer HH, Saeed NM, Hamid GA: Late presentation of breast cancer in Aden, Yemen. Gulf J Oncolog, 2011; 9: 7–11.
4. Reid-Nicholson MD, Tong G, Cangiarella JF, Moreria AL. Cytomorphologic feature of papillary lesion in the male breast: a study of 11 cases. Cancer, 2006; 108: 1100-1107. [http://www.breastcancer.org/symptoms/type/rar\\_idc/papillary.jsp](http://www.breastcancer.org/symptoms/type/rar_idc/papillary.jsp).
5. Maluf HM, Koener FC. Solid papillary carcinoma of the breast: a form of intraductal carcinoma with endocrine differentiation frequently associated with mucinous carcinoma. Am J Surg Pathol, 1995; 19(11): 1211-1214.
6. Nasser H. Solid papillary carcinoma of the breast. Pathol Case Rev, 2009; 14(4): 157-161.
7. Tasng WY, Chan JK. Endocrine ductal carcinoma in situ (E-DCIS) of the breast: a form of low-grade DCIS with distinctive clinicopathologic and biologic characteristics. Am J Surg Pathol, 1996; 20(8): 921-931.
8. Collins LC, Carlo VP, Hwang H, Barry TS, Gown AM, Schmitt SJ. Intra cystic papillary carcinomas of the breast: A reevaluation using a panel of myoepithelial cell markers. Am J Surg Pathol, 2006; 30(8): 1002-1007.
9. MacGrogan G, Moinfar F, Raju U. Intraductal papillary neoplasms. In Tavassoli FA, Devilee P eds. Pathology and Genetics of tumors of the breast and female Genital organs Lyon: IARC Press 2003, 76-80.
10. Mulligan AM, O'Malley FP. papillary lesions of the breast: A review. Adv Anat Pathol, 2007; 14: 108-19.
11. Anderson WF, Devesa SS. In situ male breast carcinoma in surveillance, Epidemiology, and results database of National cancer Institute. Cancer, 2005; 104: 1711-1741.
12. Sinha S, Hughes RG, Ryley NG. Papillary carcinoma in male breast cyst: a diagnostic challenge. Ann R Coll of Surg Engl, 2006; 88(5): W11-W5.
13. M. Hanby and C. Walker, "invasive papillary carcinoma," in WHO Classification of tumors of the Breast, P.64, IARC Press, Lyon, France, 2012.
14. Vural, A. Alnak, and K. Altundag, "invasive papillary carcinoma of the breast: An overview of twenty-four cases," Ann Surg Oncol, 2012; 19: 144-145.
15. Pal SK, Lau SK, Kruper L, Nwoye U, Gruberogio C, Gupta RK. Papillary carcinoma of the breast; An overview. Breast cancer Res treat, 2010; 122: 611-615.
16. M. Tse, P.H, and T. Moriya. "The role of immunohistochemistry in the differential of papillary lesion of the breast," Journal of clinical pathology, 2009; 62(5): 407-411.
17. Vasconcelose, A. Hussainzanda, S. Berger et al., "The St. Gallen Surrogate classification for breast cancer subtypes successfully predicts tumor presenting features, nodal involvement, recurrence patterns and disease free survival," Breast, 2016; 29: 181-185.
18. Duprez, P. M. Wilkerson, M. Lacroix-Triki et al., "immunophenotypic and genomic characterization of papillary carcinoma of the breast," Journal of pathology, 2012; 226: 427-441.
19. Solorzano CC, Middleton LP, Hunt KK, Mirza N, Merice F, Kuerer HM, Ross MI, Ames FC, Feig BW, Pollock RE, Singletary ES, Babiera G. Treatment and outcome of patients with intracystic papillary carcinoma of the breast. Am J Surg, 2002; 184(4): 464-8.
20. Prajapati ZN, Rathod GB, Gonsai RN, TB. Matrix producing mammary carcinoma –A rare breast tumor. International journal of Current Research and Review, 2012; 4(15): 62-65.
21. Rathod GB, Rahual Goyal, Rippal Kumar Bhimani, Goswami SS. Metaplastic Carcinoma of breast in 65 years old female-A case report. Medical Science, 2014; 10(119): 77-81.
22. Grabowski J, Salzstein SL, Sader GR, Blair S. Intracystic Papillary carcinoma: a review of 917 cases. Cancer, 2008; 111: 916-920.
23. Solorzano CC, Middleton LP, Hunt KK, et al. Treatment and outcome of patients with intracystic papillary carcinoma of the breast. Am J Surg. 2002; 184(4): 464-468.
24. Fayanju OM, Ritter J, Gillanders WE, et al. Therapeutic management of intracystic papillary carcinoma of the breast: the roles of radiation and endocrine therapy. Am J Surg. 2007; 194: 497-500.