AKT signaling pathway in invasive ductal carcinoma of the breast: correlation with ER α , ER β and HER-2 expression

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ABSTRACT

Aims and background. Estradiol exerts most of its effects by direct binding to the estrogen receptor in breast carcinoma, ER β expression is a useful biomarker for breast cancer in a manner that is independent of ER α expression. However, studies evaluating ER β expression with certain tumor variables, such as tumor grade and disease-free survival, had produced conflicting results. The Akt signaling pathway currently attracts considerable attention as a new target for effective therapeutic strategies. The current study attempted to compare the relative associations of variables including ER α , ER β , HER-2/neu and AKT staining with the presence of metastases or survival.

Methods and study design. Immunohistochemical staining was employed to determine the expression of $ER\alpha$, $ER\beta$, pAkt and HER-2/neu in 110 cases of primary breast carcinoma.

Results. Positive ER α , ER β , pAkt and HER-2/neu expressions were respectively observed in 46.4% (51/110), 59.1% (65/110), 40.9% (45/110) and 31.8% (35/110) of the tumors. pAkt was significantly associated with HER-2/neu overexpression (*P* <0.005) and axillary lymph node metastasis (*P* <0.05). However, there was no significant relationship between pAkt and ER α , ER β , p53 (*P* >0.05) expressions. Survival analysis showed that pAkt positivity was associated with poor disease-free survival of the patients.

Conclusions. The current study suggested that activity of the Akt signaling pathway may indicate a poor prognosis in patients with breast carcinoma. The results implied that estrogen can activate the PI3K-Akt pathway through $ER\alpha$ and $ER\beta$ -independent mechanisms in breast cancer.

Key words: Akt, breast neoplasm, estrogen receptor- α , estrogen receptor- β , HER-2.

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