MINI-REVIEW

Estrogen Receptor α Roles in Breast Cancer Chemoresistance

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Abstract

Resistence to chemotherapy treatment, which may lead to limited efficacy of systemic therapy in breast cancer patients, is multifactorial. Among the mechanisms of resistance to chemotherapy treatment, there are those closely related to estrogen receptor α, P-glycoprotein, multidrug resistance-related protein, glutathione S-transferase pi and topoisomerase-II. ERα is a ligand-activated transcription factor that regulates gene expression and plays a critical role in endocrine signaling. In previous preclinical and clinical studies, positive ERα expression in breast cancer cells was correlated with decreased sensitivity to chemotherapy. This article reviews current knowledge on the predictive value of ERα with regard to response to chemotherapy. Better understanding of its role may facilitate patient selection of therapeutic regimens and lead to optimal clinical outcomes.

Keywords: Estrogen receptor α - chemoresistance - breast cancer - Fulvestrant

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Introduction

The estrogen receptor α (ERα) plays an important role in the progression of breast cancer. About 65% of the human breast cancers were estrogen dependent and express ER α (Puhalla et al., 2012; Xu et al., 2012). Chemotherapy is a critically important treatment choice for most breast cancer patients, which has been shown to substantially improve disease-free survival and overall survival, but chemotherapy often fails to cure the disease due to unexpected drug resistance (Karroum et al., 2012; Natarajan et al., 2012). In recent years, accumulative data from clinical trials and laboratory trials suggested that ERα status might also affect the efficacy of chemotherapy. Specifically, it has been observed that some chemotherapeutic agents may be less effective in patients with ERα+ tumors than those with ERα- tumors (Precht et al., 2010; Bailey et al., 2012; Lips et al., 2012). These findings indicate that ERα status may play an important role in determining the sensitivity of breast tumors to chemotherapy.

Drug resistance is one of the major obstacles limiting the success of breast cancer chemotherapy. In this field, inherent or acquired chemotherapy resistance, which can include development of resistance to multiple drugs, is a frequent phenomenon in breast cancer cells. Forty percent of breast cancer patients with surgery and 80% of breast cancer patients with unresectable disease have poor response to chemotherapy (Petit et al., 2001; Osako et al., 2012). Several mechanisms of drug resistance have been examined. Overexpression of a membrane efflux transporter, P-glycoprotein (P-gp) (Shi et al., 2011), overexpression of multidrug resistance (MDR)-associated protein (Romero et al., 2011), changes in topoisomerase II activity (Romero et al., 2011), modifications in glutathione S-transferase (Romero et al., 2012), and altered expression of apoptosis-associated protein Bcl-2 (Bjerre et al., 2012; Larsen et al., 2012) and tumor suppressor protein p53 (Bailey et al., 2012). ERα status, although lesser known, have also been studied in connection with MDR.

Estrogen receptor and its isoforms

Estrogen receptors are ligand-regulated transcription factors; its biological action is mediated by binding to estrogen. Ligand-binding induces conformation changes in the receptor leading to dimerization, protein-DNA interaction, recruitment of co-regulator proteins and other transcription factors and ultimately the formation of the pre-initiation complex. The Estrogen receptors including isoforms ERα and ERβ, both receptors are expressed in breast cancer. ERα and ERβ have a common structural architecture composed of three functional domains including the N-terminal A/B domain, the C or DNA-binding domain and the D/E/F or ligand-binding domain. ERα isoform has attracted a great deal of attention as a key molecule in the progression of breast cancer.

The ERα consists of at least three different variants. ERα66, which was a full-length counterpart divided into six distinct regions (A-F), was the first identified estrogen receptor (Figure 1). The N-terminal domain A/B is the ligand-independent transactivation function (AF-1). Domain C is the receptor dimerization and the binding to specific DNA sequences. The C-terminal domain E/F is
the ligand-dependent transactivation (AF-2). In contrast to its full-length counterpart, the truncated ERα46 lacks the transactivation domain AF-1 and encodes a protein with a predicted molecular weight of 46kDa. This ERα36 variant lacks both transactivation domains AF-1 and AF-2 compared to ERα66 and encodes a 36kDa protein.

**Estrogen receptor α function**

Estrogen receptor, ERα, for 17β-oestradiol, was first identified from rat uteri in 1966. Human ERα is located on human chromosome 6q25, first cloned in 1986. Estrogens are steroid hormones, which is associated with the human female reproductive cycle, interaction with estrogen receptors affects the growth, differentiation, and function of target tissues. Binding of ERs causes a conformational change allowing disassociation from co-repressors and recruitment of co-activator molecules. Estrogen-bound ERα translocation to the nucleus, where they increased transcription of target genes by two different mechanisms, classical pathway where ERα binds to estrogen-response element (ERE) and modulates target genes (Figure 2) and non-classical pathway where ERα binds to co-activator molecules, such as specificity protein 1 (Sp1), activating protein 1 (AP-1), or nuclear factor kappa b (NF-kB), associated with their recognition sites in enhancer elements and modifies their function.

**Significance of ERα in breast cancer diseases**

The attempt to define the prognostic and treatment value of ERα expression concerning mainly breast cancer patients treated has already been undertaken by many researchers.

The breast cancer patients with high ERα expression were benefit from endocrine therapy (selective estrogenreceptor modulators or aromatase inhibitors) (Kemp et al., 2011; Geisler et al., 2012; Goncalves et al., 2012; Walker et al., 2012; Walker et al., 2012). Tamoxifen (selective estrogenreceptor modulators) is currently used in both premenopausal and postmenopausal breast cancer patients with ERα positive expression (Fisher et al., 1996; Fleeman et al., 2011). The Early Breast Cancer Trialists Collaborative Group meta-analysis of 194 randomized trials employing adjuvant chemotherapy and endocrine therapy demonstrated the 15-year breast cancer recurrence rate was reduced from 45% to 33% with use of tamoxifen, and reduction of breast cancer mortality by 35% (EBCTCG et al., 2008). This meta-analysis determined that the risk of the breast cancer patients with ERα positive expression was reduction with tamoxifen both in premenopausal and postmenopausal women.

**Resistance to chemotherapy in ERα positive breast cancer**

The potential association between expression of ERα in breast tumors and resistance to chemotherapy treatment was first reported about two decades age (Tokuda et al., 2012). In a retrospective study, thirty-four of 45 patients with low or absent ERα expression level had objective responses to chemotherapy, whereas only three of 25 patients with higher ERα expression responses to chemotherapy. Since then, many papers have indicated that expression level of ERα may interfere with the therapeutic efficacy of chemotherapy in breast tumors (Dziadyk et al., 2004; Mutoh et al., 2006; Tabuchi et al., 2009). To identification the role of ERα in breast cancer threatment, the 97 breast cancer patients were studied, including the expression level of Bcl-2, ERα, PS3 HER2 and Ki-67 associated with neoadjuvant chemotherapy response. The results showed that ERα negativity is associated with better chemotherapy response (Wang et al., 2009). The similar results were showed in others (Lv et al., 2011; Jeong et al., 2012).

The neoadjuvant chemotherapy permits the assessment of pathologic response of the primary cancer tissues
ICI 182,780 caused a gradual but dramatic decrease in ERα expression in the human tissues, but no change in ERβ expression (Oliveira et al., 2003). ICI 182,780, a steroidal pure antiestrogen agent, dramatically down-regulates ERα protein levels.

In this regard, several studies have shed light on the role of ERα in breast cancer chemotherapy. For instance, Sui et al. demonstrated that ICI 182,780 could significantly sensitize ERα breast cancer cells to chemotherapy drug-induced G2/M arrest and changes in IκBa and bcl-2 (Sui et al., 2007). Moreover, ICI 182,780 was also reported to enhance drug toxicity or more favorable tumor response with ICI 182,780 treatment in breast cancer cells with ERα expression, other studies did not substantiate these findings. Further work in this area is clearly needed.

Given this, in the foreseeable future, drug treatment may be guided by individualized genotype databases that can enable customized drug dosing to minimize toxicity and to enhance therapeutic effect. The past several years have provided mounting evidence for expression of ERα in breast cancer and ERα expression frequently correlates with chemotherapy-resistant disease.

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References


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