

Adjuvant Trastuzumab with Docetaxel or Vinorelbine for HER-2-Positive Breast Cancer

Gianluigi Ferretti, Paola Papaldo, Alessandra Fabi, Paolo Carlini, Alessandra Felici, Francesco Cognetti

Division of Medical Oncology A, Regina Elena Cancer Institute, Rome, Italy

Colozza et al. [1] recently presented an overview of four adjuvant trials with trastuzumab in patients with HER-2-overexpressing and/or amplified breast cancers (the North Central Cancer Treatment Group Trial N9831, the National Surgical Adjuvant Breast and Bowel Project Trial B31, the Breast Cancer International Research Group Trial 006, and the HERceptin[®] Adjuvant study). A very recent study showed that a short course of trastuzumab administered concomitantly with docetaxel or vinorelbine is effective in women with breast cancer who have an amplified *HER-2/neu* gene [2].

In vitro studies indicated that the combination of trastuzumab and vinorelbine exerts synergistic activity [3]. In fact, the results of recent clinical studies of this combination inuntreated or heavily pretreated patients with HER-2-positive metastatic breast cancer [4, 5] have shown high objective response rates. In vitro, the combination of trastuzumab with docetaxel exerts synergistic activity, as well [2]. This combination in untreated patients with HER-2-positive metastatic breast tumors showed remarkable efficacy [6].

Concerning the adjuvant setting, Joensuu et al. [2] randomly assigned 1,010 women with axillary node-positive or high-risk node-negative cancer to receive three cycles of docetaxel or vinorelbine, followed by three cycles of fluorouracil, epirubicin, and cyclophosphamide.

The 232 women whose tumors had an amplified HER-2/ neu gene were assigned to receive or not to receive nine weekly trastuzumab infusions. Within the subgroup of patients with HER-2/neu-positive breast cancer, those who received trastuzumab had better 3-year recurrencefree survival than those who did not (hazard ratio [HR], 0.42; p = .01). Interestingly, the hazard ratio remained similar (0.41) when adjustment was made according to the type of chemotherapy given (trastuzumab combined with docetaxel or vinorelbine). Thus, even though, globally, recurrence-free survival at 3 years was better with docetaxel than with vinorelbine (HR, 0.58; p = .005), the concurrent administration of trastuzumab with one of these drugs was able to overcome this difference. Moreover, in the adjuvant setting, delayed administration of trastuzumab may be less effective than concurrent administration with paclitaxel [7].

Thus synergy for trastuzumab plus cytotoxic drug combinations is specific for HER-2-overexpressing tumor cells [3] and is not observed in HER-2-negative cells, which show different sensitivity to docetaxel or vinorelbine.

DISCLOSURE OF POTENTIAL CONFLICTS OF INTEREST

The authors indicate no potential conflicts of interest.

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Correspondence: Gianluigi Ferretti M.D., Ph.D., Division of Medical Oncology A, Regina Elena Cancer Institute, Via Elio Chianesi 53, 00144, Rome, Italy. Telephone: 00390652665354; Fax: 00390652665637; e-mail: gia.fer@flashnet.it Received April 13, 2006; accepted for publication May 26, 2006. ©AlphaMed Press 1083-7159/2006/\$20.00/0

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