

Eur J Cancer, Vol. 27, No. 12, p. 1710, 1991.
 Printed in Great Britain
 0277-5379/91 \$3.00 + 0.00
 Pergamon Press plc

Letters

Phase II Study of Combination Vincristine, Epirubicin and Cyclophosphamide in Advanced Breast Cancer in Chinese Patients

W. Shiu, M. Tao and T. Leung

BREAST CANCER is the commonest female cancer in Hong Kong [1]. Chemotherapy is highly effective although little is known about its effects on breast cancer in Chinese people.

From 1985 to 1989, 83 patients with advanced breast cancer were entered into a phase II study of combination chemotherapy. Their treatment was vincristine 2 mg, epirubicin 70 mg/m², and cyclophosphamide 500 mg/m² given every 3 weeks for a maximum of 8 cycles, all intravenously on day 1. Before entry, all patients had full blood counts, liver and renal function tests and an electrocardiogram. Bone scan, ultrasound of liver, and chest X-ray were also done. Inclusion criteria were histologically proven advanced breast cancer; age under 75; disease evaluable clinically or radiologically; Karnofsky performance score of more than 70; white blood counts of more than $3 \times 10^9/l$ and platelets of more than $100 \times 10^9/l$. The treatment was delayed weekly until the white blood counts were more than $3 \times 10^9/l$ and platelets were more than $100 \times 10^9/l$.

Tumour response was assessed after two cycles. If the response was static or progressive, treatment was discontinued. Patients with responsive disease were given a maximum of eight cycles. Response and toxicity to treatment were scored with WHO criteria [2]. Complete response was defined as disappearance of all evidence of disease (two observations at least 4 weeks apart). Partial response was a decrease of at least 50% of the products of the largest perpendicular diameters of evaluable lesions. It is not necessary for all evaluable lesions to have regressed to count as a partial response; however there should not have been development of new lesions. Stable disease was a decrease of less than 50% in the products of the diameters of evaluable lesions or a less than 25% increase in the same measurements. Progressive disease was a 25% or more increase in the size of at least one evaluable lesion or the appearance of a new lesion. The new development of a pleural effusion or ascites was also considered progressive disease, if this was confirmed by cytology.

3 patients were not evaluable for response due to protocol violation (no evaluable disease either clinically or radiologically). Histologically all except 1 case had infiltrating ductal carcinoma (Table 1). 21 out of 80 (26%) patients achieved a complete response; 24 (30%) achieved a partial response. Therefore the

Table 1. Clinical characteristics of patients

Median age	46 (27-70)
Previous locoregional RT	43
Previous chemotherapy	31
Median KPS	100 (70-100)
Median no. of cycles received	6 (2-8)
Site of metastases	
Liver	12
Lung	15
Chest Wall	32
Lymph node	22
Bone	1
Kidney	1
Menstrual status	
Premenopausal	48
Perimenopausal	3
Postmenopausal	29
Unknown	3

RT = radiotherapy, KPS = Karnofsky performance score.

overall response rate was 56%. 20 (25%) showed static response and 15 (19%) had progressive disease on treatment.

At the time of analysis, 64 patients had already died and 2 had defaulted follow-up; only 17 patients remained alive. The median survival of the dead patients was 40.1 weeks (range 3.4-151.4). For those who are alive, median survival is 104.4 weeks (30.3-160.6).

4 out of 83 (5%) patients had grade 4 and 15 (18%) grade 2-3 haematological toxicity. 56 (67%) had grade 2-3 nausea and vomiting. 33 (40%) had grade 1 peripheral neuropathy. Only 2 (2%) had grade I cardiac toxicity.

It is not clear whether a disease such as breast cancer in one ethnic group behaves the same as in another and it is easy to extrapolate data from the West for reference. We have done a phase II study of combination chemotherapy in a selected Chinese group with advanced breast cancer. The response rate was similar to the overall response rate of 40-80% reported in the West [3]. The overall median survival was also similar to that in the West with a medium survival of 9 months in patients with visceral metastases [4-6]. Vincristine, epirubicin and cyclophosphamide is as active in Chinese patients with breast cancer as cases in the West.

1. Hong Kong Cancer Registry 1989. Department of Health. Hong Kong Government.
2. Miller A, Hoogstraten B, Staquet M, Winkler A. Reporting results of cancer treatment. *Cancer* 1981, **47**, 207-214.
3. Muss HB, White DR, Richards F. Adriamycin versus methotrexate in five drug combination chemotherapy for advanced breast cancer: A randomised trial. *Cancer* 1978, **42**, 2141-2148.
4. Tjandra J, McLaughlin P, Russell S, Collins J, McKenzie F. Comparison of MSA with B₂-microglobulin and CEA in patients with breast cancer. *Eur J Cancer Clin Oncol* 1988, **24**, 1633-1640.
5. De Jong-Bakker, Hart A, Persijn J, Cleton F. Prognostic significance of CEA in breast cancer: a statistical study. *Eur J Cancer Clin Oncol* 1981, **17**, 1307-1313.
6. Barts S, Longo D. Tumour markers: value and limitations in the management of cancer patients. *Cancer Treat Rev* 1985, **12**, 163-207.

Acknowledgements—We thank the Hong Kong Cancer Fund (EORTC) for the research grant and Miss Annie Chan for typing the manuscript.

Correspondence to W. Shiu.

The authors are at the Department of Clinical Oncology, The Chinese University of Hong Kong, Prince of Wales Hospital, Shatin, N.T., Hong Kong.

Revised 10 July 1991; accepted 26 July 1991.