Carcinoma of stomach and breast with lymphoid stroma: localisation of Epstein-Barr virus

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Abstract

Aims—To determine the presence of Epstein-Barr virus (EBV) genome in six patients (three with gastric and three with breast carcinoma) with severe small lymphoid cell infiltration.

Methods—The polymerase chain reaction and in situ hybridisation were used to detect EBV genome. The number and distribution of T and B lymphocytes were evaluated by immunohistochemistry.

Results—Histologically all of the patients had poorly differentiated tumours. Immunohistochemistry showed that T cells predominated in three cases, B cell in two, and almost equal numbers in one case. PCR showed that the EBV genome was present in two cases each of gastric and breast carcinoma. In situ hybridisation for EBV genome provided positive signals only in the small lymphoid cells in one gastric and two breast carcinomas giving a positive reaction for EBV genome by PCR. The gastric and breast cancer cells did not give positive signals.

Conclusion—Severe lymphoid cell infiltration in gastric and breast carcinoma does not necessarily indicate that these tumours are associated with EBV. Larger numbers of cases will need to be studied to confirm this.

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An oncogenic role for Epstein-Barr virus (EBV) in B cell lymphoma and nasopharyngeal carcinoma has been suggested.1 Recent studies have shown the presence of EBV sequences in carcinoma cells of the thymus,2 stomach,1 and salivary gland.3 The EBV related epithelial tumours have generally shared common morphological features (poorly differentiated tumour cells accompanied by severe lymphoid cell infiltration—so-called lympho-epithelioma). Gastric lympho-epithelioma-like carcinoma is thought to be similar to the gastric carcinoma with lymphoid stroma reported by Watanabe et al in 1976.4

Gastric carcinoma with lymphoid stroma is relatively rare and constitutes about 4% of all gastric carcinomas.4 Because these patients survive longer than those without lymphoid stroma,1,5 the infiltration by small lymphoid cells in the tumours has been interpreted as a host reaction to the tumour. Therefore, there are two theories on the cause of carcinomas with lymphoid stroma: one is a central role for EBV, resulting in the lympho-epithelioma-like pattern; the other is a host reaction to tumour growth. In the present study, the presence of EBV genome in three cases each of gastric and breast cancer showing severe lymphoid cell infiltration and around the tumours was studied by using the polymerase chain reaction (PCR) and in situ hybridisation, originally established by us as a highly sensitive method.6

Methods

Three patients with gastric carcinoma and three with breast carcinoma showing massive infiltration of lymphoid cells in and around the primary tumour were selected for study. All patients were treated by surgical resection of tumour. Histological specimens were fixed in 10% formalin and routinely processed for paraffin wax embedding.

Sections, cut at 4 μm, were stained with haematoxylin and eosin. The tumours were classified according to the "General Rules for Gastric Cancer Study"7 or "General Rules for Clinical and Pathological Recording of Mammary Cancer" (Japan).8 The avidin-biotin-peroxidase complex staining procedure was used. Sections were incubated with normal horse serum for 30 minutes and incubated overnight at 4°C with mouse antihuman antibodies Mx-PanB(CD20) (Kyowa Medex, Tokyo; diluted at 1 in 50); MB-1, MT-1(CD43) (Bioscience, Emmenbrucke; 1 in 50); and UCHL-1 (Dakopatts, Copenhagen; 1 in 100). Subsequent reactions were accomplished using a VectaStain avidin biotin-peroxidase complex kit (Vector Laboratories, Burlingame, California, USA). The peroxidase reaction was blocked in phosphate buffered saline (pH 7·4) containing 0·005% H2O2 and 0·03% 3', 3-diaminobenzidine tetrahydrochloride.

PCR

DNA was extracted from formalin fixed, paraffin wax embedded tissues. PCR was performed using heat-stable Thermus thermophilus (Tth) DNA polymerase (Toyocho, Osaka, Japan). Primers and probes were synthesised using a DNA synthesizer (391 DNA synthesiser) and purified in an Oligonucleotide Purification Cartridge (Applied Biosystems, Foster, California). EBV primers were located in the long internal repeat 1, used previously by Ubara et al.9 The PCR products were electrophorased on 2% agarose gel and visualised with ethidium bromide staining under ultraviolet light. The gel was transferred on to a nylon membrane and hybridised with the 32P-end labelled specific oligonucleotide probe.
As positive controls, pellets of Raji cells and an EBV positive Burkitt’s lymphoma cell line were examined. A pellet of Ramos cell, derived from EBV negative Burkitt’s lymphoma cells, was used as negative control for EBV. Raji and Ramos cells were provided by the Japanese Cancer Research Resources Bank (JCRB) (Tokyo).

IN SITU HYBRIDISATION
Recently we established a highly sensitive in situ hybridisation procedure using digoxigenin-11-dUTP labelled probe using PCR. With this method, we were able to examine the localisation of EBV genomes in tissue sections. Using 12 sets of primers, the BamH1-W fragment of EBV was amplified with labelled substrate in individual PCRs. The 12 probes, with an average size of 120 base pairs, were mixed together and hybridised with the histological sections. Cytospin specimens of the EBV positive cell line, Raji (gift of Dr H Mizusawa, National Institute of Hygiene) and the EBV negative cell line MT-1 (gift of the Japanese Cancer Research Resources Bank, Tokyo) were hybridised with the probes. Numerous dots were observed over the nucleus of the Raji cells. No signals could be found in MT-1 cells. The specificity of the probes and the staining were evaluated by the following control studies: (1) the slides were incubated with the hybridisation buffer containing unlabelled DNA, subsequently incubated with antigen to digoxigenin antibody-alkaline phosphatase conjugate and its substrate; (2) signals were observed using digoxigenin labelled probe for JD, repeat sequence of mouse gene. This probe was prepared by random primed labelling, as described above.

Results
The clinical and pathological findings of the six patients are summarised in the table. All of the gastric carcinomas were poorly differentiated adenocarcinoma. Depth of invasion had extended beyond the muscularis propria in two cases and within the submucosa in one. All of the mammary carcinomas were classified as invasive solid tubular carcinoma. The lymphoid stroma consisted of small lymphoid and plasma cells in every tumour, with lymphoid follicle formation in one gastric carcinoma (case 2). Eosinophils were common in three gastric carcinomas and neutrophils in one (case 1). Small lymphoid cells had invaded the cancer cell nests in one gastric (case 2) and one breast carcinoma (case 4). Immunohistochemical staining showed that B lymphocytes outnumbered T lymphocytes in two cases (cases 2 and 4); T lymphocytes predominated in three (cases 1, 3, and 5), with an almost equal number of both subsets in one case.

The PCR method showed that the EBV genome was present in two (cases 2 and 4) out of three gastric and two breast carcinomas (cases 4 and 6) (figure). In situ hybridisation for the EBV genome provided positive signals in the small lymphoid cells in one case of gastric (case 3) and two of breast carcinoma (cases 4 and 6) with a positive reaction for EBV genome by the PCR. The signals were observed as a few dots in the nuclei. No signals were detected in the gastric and breast carcinoma cells.

Discussion
The PCR showed that about 70% of the gastric and breast carcinomas in this series contained EBV genome; this high positivity rate of EBV agreed with previously published findings on undifferentiated carcinomas with severe lymphoid infiltration of the salivary gland, thymus, and stomach. Using in situ hybridisation, a previous study had shown that the EBV genome was localised in the carcinoma cells of salivary gland and stomach. However, Min et al reported that in situ hybridisation failed to define the presence of the EBV genome in three cases of gastric carcinoma with EBV genome, confirmed by PCR, in the tumour tissues. In our study positive EBV genome signals were detected by in situ hybridisation only in the small lymphoid cells. The gastric and breast carcinoma cells never gave positive signals.

The present cases of carcinoma with severe lymphoid cell infiltration superficially resemble the previously reported cases showing similar morphological features. These tumours had undifferentiated or poorly differentiated morphology. All of our cases were poorly differentiated carcinomas. Shibata et al reported that their cases of undifferentiated carcinoma with intense lymphoid infiltration were similar to the gastric carcinoma with lymphoid stroma described by Watanabe et al.
Meanwhile Watanabe et al. reported the presence of limited areas of differentiated adenocarcinoma in 12 of 42 cases.

The immunological characteristics of the infiltrating lymphoid cells in carcinoma with severe lymphoid cell infiltration have seldom been mentioned before. Min et al. reported that most of the lymphocytes surrounding cancer cell nests in their cases of gastric cancer were T cells. It is well known that patients with undifferentiated carcinoma showing severe lymphoid cell infiltration survive longer than those without. In our cases, the T:B cell ratio was not constant; T cells predominated in three cases and B cells in two cases. Although the follow up was short, our patients did not show a favourable prognosis: two out of six died within two years.

In conclusion, this study has shown that severe lymphoid cell infiltration in gastric carcinoma, and possibly breast carcinoma, does not necessarily indicate that these tumours are neoplasms related to EBV. Because of the small numbers studied, more cases need to be investigated.

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