Regression of idiopathic thrombocytopenic purpura after endoscopic mucosal resection of gastric mucosa associated lymphoid tissue lymphoma

M Noda, N Mori, K Nomura, K Kojima, S Mitsufuji, I Yamane, S Misawa, T Okanoue

CASE REPORT

Recent reports have suggested an association between *Helicobacter pylori* infection and both gastric mucosa associated lymphoid tissue (MALT) lymphoma and thrombocytopenic purpura. Although treatments eradicating *H pylori* lead to regression of these diseases in some cases, the exact mechanisms are still controversial. This case report describes a patient with thrombocytopenic purpura accompanied by an early stage gastric MALT lymphoma. Endoscopic mucosal resection of the lesion in this patient led to dramatic regression of thrombocytopenic purpura, and t(11;18)(q21;q21), which means resistance more likely to *H pylori* eradication therapy, was confirmed by fluorescence in situ hybridisation. There is no evidence of recurrence and his platelet count is within normal limits after 24 months of follow up. This is the first case report describing regression of thrombocytopenic purpura after mucosal resection of a gastric MALT lymphoma. We suggest that while some cases of thrombocytopenic purpura may be induced by *H pylori*, others may be due to an autoreactive antibody produced by MALT lymphoma B cells.

Malignant lymphomas or mucosa associated lymphoid tissue (MALT) lymphomas are occasionally accompanied by thrombocytopenic purpura, suggesting an immunological origin for thrombocytopenic purpura. Treatments eradicating *H pylori* lead to regression of these diseases in some cases, and evidence points to an important role for *H pylori* infection in the development of idiopathic thrombocytopenic purpura (ITP). We present here a very rare but intriguing case of gastric MALT lymphoma that showed regression of thrombocytopenic purpura after endoscopic mucosal resection (EMR), and in which t(11;18)(q21;q21) was confirmed by fluorescence in situ hybridisation.

CASE REPORT

A 65 year old man was referred to our department in August 2001 with a one month history of decreased appetite and epigastric pain. He had a past history of non-insulin dependent diabetes mellitus, hyperlipidaemia, and an old myocardial infarction. Upper gastrointestinal endoscopy showed an 8 mm irregular erosion on the greater curvature of the fornix. Differential diagnosis of early gastric cancer IIc (superficial depressed) type was difficult but the histological diagnosis was gastritis with erosive change. The patient was admitted to our department for further examination and therapy because an endoscopic observation after two months demonstrated a 12 mm submucosal tumour-like lesion with a shallow ulcer on the surface (fig 1).

On admission, physical examination was unremarkable except for mild epigastric tenderness; the liver, spleen, and superficial lymph nodes were not palpable. His blood cell count however showed a platelet level of 27 000/mm$^3$. Other routine blood chemistry, liver function tests, chest radiography, and urinalysis were normal. Although we stopped administering diltiazem, famotidine, and atrovastatin to eliminate drug induced thrombocytopenia, his platelet level did not recover. A bone marrow smear showed normal cellularity and an increased number of megakaryocytes. Antiplatelet antibody titres (IgG), as tested by indirect immunofluorescence, were increased. No abnormalities in blood coagulation or in levels of autoimmune antibodies were observed.

When his platelet count was 40 000–50 000/mm$^3$, we obtained informed consent and performed EMR using a two channel scope (GIF-2T200; Olympus, Tokyo, Japan) to make an accurate histological diagnosis. Five days after EMR, his platelet count increased dramatically to 173 000/mm$^3$ (fig 2). Specimens were fixed in 10% formalin, embedded in paraffin, and routinely stained with haematoxylin and eosin (H&E). They were also sectioned for immunohistochemical study and for tissue fluorescence in situ hybridisation (T-FISH). The lesion was composed of diffuse small atypical lymphoid cells with centrocyte-like appearance (cleaved nucleus and relatively wide clear cytoplasm) that had invaded the lamina propria. The atypical lymphoid cells produced characteristic lymphoepithelial lesions, and immunohistochemically they were diffusely positive for CD20, CD79a, and CD10. By T-FISH analysis it was positive for t(11;18)(q21;q21) but negative for trisomy 3, 7, 12, and 18 (fig 3).

Ultrasonography and computed tomography showed no hepatosplenomegaly or swelling of the para-aortic lymph nodes. The tumour was diagnosed as a stage 1 extranodal marginal zone MALT B cell lymphoma. Because the rapid urease test and biopsy specimens from the gastric body and antrum were positive for *H pylori*, the patient was treated with a seven day course of lansoprazol 60 mg twice a day, amoxicillin 1500 mg twice a day, and clarithromycin 400 mg twice a day, one month after EMR. He did not receive antibiotics at any time except for this period. Three months after treatment, gastric biopsy specimens were negative for lymphoma infiltration and *H pylori* by histology and the rapid urease test, and the patient tested negative after a urea breath test. No additional therapy, such as chemotherapy or radiation, was performed. There is no evidence of recurrence.

**Abbreviations:** MALT, mucosa associated lymphoid tissue; ITP, idiopathic thrombocytopenic purpura; EMR, endoscopic mucosal resection; H&E, haematoxylin and eosin; T-FISH, tissue fluorescence in situ hybridisation
and his platelet count is within normal limits after 24 months of follow up after endoscopic removal.

MATERIALS AND METHODS

Lymphoma specimens were fixed in 10% formalin, embedded in paraffin, and stored at room temperature. Sections (4 μm thick) were cut from paraffin embedded tissue and used for histopathological observation and T-FISH. The former was performed according to routine procedures for H&E and Giemsa stainings and the latter was performed as described previously. For detection of t(11;18) (q21;q21), we used artificial chromosome clones y943b8 (specific for 18q21) and y966c4 (specific for 11q21) (obtained from MIT Center for Genome Research, Cambridge, Massachusetts, USA). Polymerase chain reaction products were labelled with either SpectrumGreen (Vysis Inc., Illinois, USA) or SpectrumOrange (Vysis Inc.) with a nick translation kit (Vysis Inc.). For detection of trisomy 3, 7, 12, and 18, we used CEP Probes CEP3 (D3Z1), CEP7 (D7Z1), CEP12 (D12Z3), and CEP18 (D18Z1) SpectrumOrange (Vysis Inc.), respectively.

Each nucleus was identified on the basis of 6-diamidino-2-phenylindole dihydrochloride staining. Images were captured with a CCD camera. Signals from at least 200 non-overlapping nuclei were evaluated per slide. For T-FISH analysis, we assessed only colocalised signals. To evaluate colocalised signals, cut off values were calculated from the mean (±2 SD), which were based on evaluation of 100 nuclei in each specimen from three patients with chronic gastritis.

DISCUSSION

MALT lymphomas are considerably mature B cell lymphomas of low malignancy that were first described by Isaacson and Wright in 1983. They were classified as marginal zone lymphomas with a shallow ulcer on the surface. A 12 mm submucosal tumour-like lesion was noted on the greater curvature of the fornix. (B) Endoscopic observation after two months revealed a slightly elevated irregular erosion at the greater curvature of the fornix. (B) Endoscopic observation after two months revealed a slightly elevated irregular erosion at the greater curvature of the fornix.

Generally, most gastric MALT lymphomas are closely linked to chronic H pylori infection, and 60–80% of low grade

Figure 2

Dramatic increase in platelet count after endoscopic mucosal resection (EMR). Suspension or re-administration of diltiazem, famotidine, and atrovastatin did not affect the platelet level.

Figure 1

(A) Endoscopic observation of the stomach showed a slightly elevated irregular erosion at the greater curvature of the fornix. (B) Endoscopic observation after two months revealed a 12 mm submucosal tumour-like lesion with a shallow ulcer on the surface.
MALT lymphomas regress in response to *H pylori* eradication therapy. In this case, however, molecular biology indicated t(11;18)(q21;q21) of the lesion, as detected by the T-FISH method, indicating a more likely resistance to *H pylori* eradication therapy. To date, radiation therapy, surgical resection, or chemotherapy have been recommended for such cases but we were able to treat the MALT lymphoma with ERM and there was no evidence of recurrence after 24 months of follow up. Furthermore, ITP also regressed after endoscopic removal of the MALT lymphoma.

In conclusion, this is the first case report on regression of thrombocytopenic purpura after mucosal resection of a gastric MALT lymphoma. The mechanisms of regression of ITP remain to be clarified but it appears that ITP may not be directly induced by *H pylori* infection in some cases but by an antibody produced by MALT lymphoma B cells or proliferated B cells in response to *H pylori* infection via T cell help.

ACKNOWLEDGEMENTS

This work was supported in part by: (1) Grant-in-Aid for Young Scientists (B) from the Ministry of Education, Culture, Sports, Science and Technology of Japan (13770268) and (2) supporting research funds from the Japanese Foundation for Research and Promotion of Endoscopy (JFE-2002).

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Accepted for publication 13 April 2004

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Figure 3 Tissue fluorescence in situ hybridisation findings on paraffin embedded tissue section. Red signals are representative of 18q21 whereas green ones are of 11q21. Arrows indicate co-localised signal (×1000).
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Gut 2004 53: 1698-1700
doi: 10.1136/gut.2003.033555

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