Splenicpleurae

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Splenosisis the autotransplantation of splenic tissue, usually after traumatic rupture of spleen. Reports of splenosis pleurae are rare. Of the six cases reported so far, there is only one (Shaw and Shafi, 1937) in the British literature. On the other hand, splenosis peritonei has been well documented. The transplants have always followed traumatic splenic rupture, diaphragmatic rupture, and splenectomy. It is important to recognise this condition to avoid unnecessary surgical procedures.

Case report

A 17-year-old labourer presented with a few weeks' history of a painful lump in the right groin, lethargy, and weakness. A road traffic accident in 1964 had resulted in fractures of the left inferior pubic ramus and left seventh rib and ruptured spleen and diaphragm. Splenectomy and repair of the diaphragm were undertaken. He remained well until his present illness. He was afebrile, and the only abnormality was painless, generalised lymphadenopathy.

INVESTIGATIONS

Results of investigations showed haemoglobin 14·3 g/dl, and white cell count 7·3×10⁹ (normal differential count). The following tests were normal: Paul Bunnel, ESR (Westergren); viral agglutinations; serum calcium; liver function, immunoglobulins and protein electrophoresis; urea and electrolytes; electrocardiography; intravenous pyelogram; and hydatid complement fixation. Chest radiograph (fig 1) showed rounded lesions in the left mid-zone. A chest radiograph in 1964 had been normal.

At exploratory thoracotomy, multiple purple, sessile nodules, some lobulated, were removed from the parietal, visceral, and mediastinal pleura. They ranged from one to three centimetres in diameter and had the appearance of small spleens. They were soft and surrounded by thin fibrous capsules, but no definite intrasplenic trabeculae could be seen. Microscopic examination (fig 2) showed that red pulp predominated over the white. The white pulp consisted of clusters of lymphocytes with no germinal centres. In most of these clusters were arterioles either lying beside the lymphoid follicles or occasionally centrally, indistinguishable from malpighian corpuscles. The red pulp consisted of endothelial-lined sinusoids, histiocytes, and scattered lymphocytes. The connective tissue capsule, in some areas, was well vascularised and contained rather large arteries and veins. The fibrous trabeculae normally present in the spleen were not shown in any of the nodules. The adjacent lung showed patchy areas of fibrosis and old haemorrhage compatible with previous trauma.

DISCUSSION

Splenosis rarely follows traumatic splenic rupture and is an important differential diagnosis of multiple pulmonary opacities when there is a history of trauma to the spleen. Other causes of similar radiological changes are accessory spleens, endometriosis, intrathoracic and metastatic neoplasms, and haemangiomatis. Splenotic nodules presumably result from autotransplantation on the pleura after scattering of splenic red pulp fragments from the ruptured spleen. Blood-borne spread is unlikely because the splenic tissue is not seen in other organs.

Splenotic nodules were noted (Schilling, 1907) at a laparotomy, but the phenomenon of auto-transplantation and splenosis was first suggested by Buchbinder and Lipkoff (1939). Finger clubbing, dyspnoea, and cervicofacial venous congestion have been described in association with thoracic transposition of the spleen. These splenotic nodules may be responsible for a relapse of a haematological disorder after removal of the spleen for the condition (Fleming et al, 1976. The absence

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of Howell-Jolly bodies, siderocytes, and other features found in the peripheral blood film after splenectomy may suggest the presence of residual splenic tissue. Various radioisotope scanning techniques (Jacobson and De Nardo, 1971) have been used to diagnose functioning splenic tissue.

Although splenosis pleurae is a very rare condition, this being only the second case report in British literature, it should be considered in the differential diagnosis of intrapulmonary opacities in the presence of history of splenic rupture.

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References


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Splenosis pleurae.

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