The cellular infiltrate in Hashimoto’s disease and focal lymphocytic thyroiditis

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SYNOPSIS  The lymphoid cells infiltrating the thyroid in three examples of Hashimoto’s disease and three examples of focal lymphocytic thyroiditis have been studied by light and electron microscopy. The cell types found were small lymphocytes, plasma cells and plasmablasts, immunoblasts, and cells morphologically intermediate between immunoblasts and small lymphocytes. The infiltrate was similar in the two conditions studied and resembled the cell response found in other conditions thought to be due to delayed hypersensitivity. It is considered that these similarities support the views that focal lymphocytic thyroiditis is a focal form of Hashimoto’s disease and that delayed hypersensitivity is important in the pathogenesis of these conditions.

Infiltration of the thyroid by lymphoid cells in Hashimoto’s disease is thought to reflect a delayed hypersensitivity reaction to the thyroid tissue (Lindsay, 1964).

The purpose of this paper is to describe these lymphoid cells in detail and to compare them with the cell mediators found in other examples of delayed hypersensitivity reactions. A comparison is also made with focal lymphocytic thyroiditis since this is regarded by some authorities as a focal form of Hashimoto’s disease (Doniach, 1960).

MATERIALS AND METHODS

Thyroid tissue from three cases of histologically proven Hashimoto’s disease and three cases of hyperthyroidism with focal lymphocytic thyroiditis was examined.

Tissue for electron microscopy was obtained from the fresh surgical specimens immediately after excision, cut into 1 mm³ blocks and fixed in ice-cold glutaraldehyde in phosphate buffer at pH 7·2 for two hours, postfixed in buffered osmium tetroxide, and embedded in Epon (Luft, 1961). Sections were cut on a Reichert ultratome, stained with uranyl acetate and lead citrate and examined in a Hitachi HS7S electron microscope.

In all cases representative blocks were embedded in paraffin wax, and sections stained by haematoxylin and eosin and methyl green-pyronin were examined.

For the purposes of this study focal lymphocytic thyroiditis was defined as multiple collections of lymphoid cells in the interstitial tissue throughout the gland. Askanazy cell metaplasia, lymphoid follicle formation, and fibrosis were not considered to be essential features. In selecting areas for electron microscopy germinal centres, if present, were avoided.

RESULTS

The types of infiltrating cells were similar in all the specimens examined whether examples of Hashimoto’s disease or of focal lymphocytic thyroiditis.

LIGHT MICROSCOPY  Three cell types were identified (Figs. 1 and 2), namely, small lymphocytes, plasma cells, and large lymphocytes with basophilic, pyroninophilic cytoplasm, and vesicular, sometimes infolded nuclei.

ELECTRON MICROSCOPY  The cell types identified were small lymphocytes, large lymphocytes, and plasma cells.

Small lymphocytes  These cells (Fig. 3) were 4 to 6·5 μ in diameter. Their cytoplasm varied in amount but was always relatively scanty and contained abundant single, free ribosomes in a rather electron-opaque hyaloplasm. Endoplasmic reticulum was very sparse, the Golgi apparatus was small and only seen occasionally, and there were few mitochondria. Nuclear chromatin was coarsely condensed and a nucleolus was rarely seen.

Large lymphocytes  The cell diameters varied from 7·8 to 12·6 μ. Cytoplasm was greater in amount than in the small lymphocytes and was sometimes abundant; free ribosomes were plentiful and arranged mainly as polyribosomes in an electronlucent hyaloplasm. Although better developed than in the small lymphocytes, endoplasmic reticulum was still scanty and consisted of narrow

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double strands irregularly studded with ribosomes. Mitochondria were present in moderate numbers and the Golgi apparatus was often seen to be well developed. Nuclear chromatin was less coarsely condensed than in the small lymphocytes and a nucleolus was often present. These cells (Fig. 4) were considered to be identical with the cells termed immunoblasts, large pyroninophilic cells, haemocytoblasts, etc, by various authors.

A number of cells resembled those just described except that they tended to be smaller (6 to 7.6 μm diameter) and their ribosomes were variable in number and were either arranged mainly as single free particles or as a mixture of varying proportions of polyribosomes and single ribosomes (Figure 5). In many respects they resembled cells described by de Petris, Karlsbad, Pernis, and Turk (1966) in lymph nodes draining areas of contact sensitivity and considered by them to be intermediate between immunoblasts and small lymphocytes.

**Plasma cells** Numerous morphologically typical plasma cells were seen and will not be described further (Figure 6). In addition there were a number of cells which appeared to be morphologically intermediate between immunoblasts and mature plasma cells. These cells contained more rough endoplasmic reticulum than immunoblasts and as the amount increased it became progressively more orientated into parallel arrays (Figures 6, 7, and 8). They appear to correspond to the cells termed plasma blasts and described in lymph nodes draining homografts (André-Schwartz, 1964) and in nodes draining an area of contact sensitivity (de Petris et al, 1966).

**DISCUSSION**

The mechanism of tissue damage in human and experimental lymphocytic thyroiditis is uncertain. The demonstration of circulating autoantibodies in Hashimoto’s disease (Roitt, Doniach, Campbell,
and Hudson, 1956; Witebsky, Rose, Terplan, Paine, and Egan, 1957) appeared at first to provide a satisfactory explanation and experiments in vitro with trypsinized monolayer cultures of thyroid cells indicated that Hashimoto serum was cytotoxic (Pulvertaft, Doniach, Roitt, and Hudson, 1959). The cytotoxic component was subsequently shown to be identical with complement-fixing thyroid autoantibody (Forbes, Roitt, Doniach, and Solomon, 1962). However, doubts about the significance of circulating antibodies in the pathogenesis of Hashimoto’s disease arose when it became apparent that the antibodies had no adverse effect on monolayer tissue cultures which had not been pretreated with trypsin nor on thyroid fragments in organ culture (Roitt, Jones, and Doniach, 1962; Irvine, 1962).

In experimental autoimmune thyroiditis further difficulties arose when it was shown that the degree of thyroid damage correlated better with delayed hypersensitivity skin tests than with levels of circulating autoantibody (McMaster, Lerner, and Exum, 1961; Miescher, Gorstein, Benacerraf, and Gell, 1961), and it has been demonstrated that the production of delayed hypersensitivity without antibody formation by immunizing guinea pigs with picrylated homologous thyroglobulin causes thyroiditis (Meischer et al, 1961). Furthermore, attempts to transfer thyroiditis passively by injecting serum from sensitized animals into normal animals have failed (Rose, Kite, and Doebblers, 1961) although success has been claimed by transferring lymph node cells from sensitized guinea pigs (Felix-Davies and Waksman, 1961; McMaster and Lerner, 1967).

It seems clear that circulating autoantibodies cannot be the sole factor in the pathogenesis of...
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naturally occurring lymphocytic thyroiditis or its experimental counterpart. Roitt et al (1962) have suggested that the antibodies can act only on cells whose surface membranes have first been damaged and that this damage may be caused by sensitized lymphoid cells infiltrating the thyroid.

In view of their apparent importance it is surprising that the nature of the infiltrating cells in Hashimoto's disease has not received more attention. In most histological accounts they are described as lymphocytes and plasma cells with no further qualification and Hazard (1955) referred to them as being of normal 'adult' type. Ultrastructural studies of Hashimoto's disease are few. Irvine and Muir (1963) examined four examples by electron microscopy but they were mainly concerned with epithelial cell and basement membrane changes, and the lymphoid cells were not studied in detail although it was observed that lymphocytes were frequently found within the follicular basement membranes in intimate relationship with the epithelial cells. Binet, Gennes, and Decourt (1963), in a study of two cases of Hashimoto's disease, were exclusively concerned with the nature of the infiltrating lymphoid cells.

FIG. 3. Small lymphocyte. The nuclear chromatin is coarsely condensed at the periphery of the nucleus (N). The cytoplasm contains abundant single, free ribosomes, a single strand of endoplasmic reticulum, and a few mitochondria (m). A Golgi apparatus is not seen. Uranyl acetate and lead citrate × 19,800.
FIG. 4. Large lymphocyte (immunoblast). The nucleus (N) is infolded and the nuclear chromatin is finely dispersed; a nucleolus is present. The cytoplasm contains numerous polyribosomes, scattered short strands of endoplasmic reticulum, several mitochondria (m), and a large Golgi apparatus (g). Uranyl acetate and lead citrate × 10,000.

FIG. 5. Two intermediate lymphocytes. The nuclei are similar to the nucleus of a small lymphocyte but the cells are larger, cytoplasm is more plentiful, and there are some polyribosomes as well as numerous single ribosomes. The Golgi apparatus (g) is well developed in the larger cell. Uranyl acetate and lead citrate × 10,000.
and described three cell types, namely, lymphocytes, plasma cells, and basophilic histioflyes like those found in 'retarded reactions to grafts'. The latter had a large Golgi apparatus, numerous cytoplasmic ribosomes, and no ergastoplasm.

No electron microscopic studies of focal lymphocytic thyroiditis have been reported.

In the present study the infiltrating lymphoid cells were similar in Hashimoto’s disease and focal lymphocytic thyroiditis and comprised small lymphocytes, plasma cells and plasmablasts, immunoblasts, and cells morphologically intermediate between immunoblasts and small lymphocytes, findings which are in general agreement with those of Binet et al (1963) in Hashimoto’s disease.

The similarity of the cellular infiltrate in Hashimoto’s disease and focal lymphocytic thyroiditis appears to support the view that the two conditions are essentially similar in pathogenesis, a similarity which has previously been commented on with regard to the epithelial changes in the two conditions and the correlation of focal lymphocytic thyroiditis with the occurrence of circulating antibodies of similar type to those found in Hashimoto’s disease (Senhauser, 1964).

The infiltrating cells described here are of the same types as those found infiltrating allografts undergoing unmodified rejection (Porter, Joseph, Rendall, Stolinski, Hoehn, and Calne, 1964), and in lymph nodes draining allografts (André-Schwartz, 1964) and areas of contact sensitivity (de Petris et al, 1966). This resemblance, particularly that between grafts undergoing rejection and thyroids that are affected by lymphocytic thyroiditis, is suggestive of a basic similarity between the various conditions and is considered to support the concept of the fundamental importance of cell-mediated immunity in the pathogenesis of Hashimoto’s disease and focal lymphocytic thyroiditis.

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FIG. 7. Early plasmablast. The cell resembles an immunoblast in nuclear and cytoplasmic characteristics but shows an increased amount of rough endoplasmic reticulum. Uranyl acetate and lead citrate × 8,000.

FIG. 8. Plasmablast. The nuclear characteristics are intermediate between those of a mature plasma cell and an immunoblast. The cytoplasm contains a moderate amount of rough endoplasmic reticulum showing some organization into parallel arrays. Part of a mature plasma cell is seen in the top left corner. Uranyl acetate and lead citrate × 10,000.
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REFERENCES