

it is thus the first organ in all animal species to become predominantly lymphoid. It continues to grow till about the 12th year after puberty. It undergoes spontaneous progressive involution. Indicating that it functions best in early life

The thymus is located behind the upper part of the sternum. Aberrant thymic tissues are often found in neighboring sites. It has two lobes surrounded by a fibrous capsule. Septa arising from the capsule divide the gland into lobules which are differentiated into an outer cortex and an inner medulla. The cortex is crowded with actively proliferating, small lymphocytes. The medulla consists mainly of epithelial cells and mature lymphocytes amidst which are Hassall's corpuscles, which are whorl-like aggregations of epithelial cells.

Till recently, the thymus was an organ without any recognised function. The fortuitous observations by GOOD (1954) of thymoma and impaired immunity in a patient, and by MILLER (1961) of immunodeficiency in neonatally thymectomised mice, paved the way for the understanding of the pivotal role of the organ in the development of the cell-mediated immunity. The primary function of the thymus is the production of the thymic lymphocytes. It is the major site for lymphocyte production, only about one percent leave the thymus. The rest are destroyed locally. The reason for this apparently wasteful process is not known. In the thymus, the lymphocytes acquire new surface antigens (thymic antigens). Lymphocytes produced in the thymus are called thymus (T) dependent lymphocytes or T cells. Unlike lymphocyte proliferation in the peripheral organs, in the thymus it is not dependent on antigen stimulation. In fact, peripheral antigenic stimuli do not lead to any immune response in the thymus. Antigen

introduced directly into the thymus may lead to a local immune response.

(Fig...) development of T and B-cell systems.

the thymus confers immunological competence on the lymphocytes during their stay in the organ, prethymic lymphocytes are not immunocompetent, in the thymus they are educated so that they become capable of mounting cell mediated immune response against appropriate antigens, this at least in part is effected by hormone like factor produced by the thymus such as thymulin thymosin and thymoprotein, the importance of thymus in lymphocytes proliferation and development of CMI is evident from the lymphopenia deficient graft rejection and the so called runt disease seen in neonatally thymectomised mice. Deficient CMI is also seen in congenital aplasia of the thymus in human being (DiGeorge syndrome) and in mice (nude mice).

T- lymphoid are selectively seeded into certain sites in the peripheral lymphatic tissues, being found in the white pulp of the spleen, around the central arterioles and in the paracortical areas of lymph nodes, these regions have been termed thymus dependent as they found grossly depleted after neonatal thymectomy, while thymectomy affects CMI primarily, it also diminishes antibody response to certain types of antigens (thymus dependent antigens) such as sheep erythrocytes and bovine serum albumin, Humoral response to other antigens is unaffected.

Bursa of Fabricius:-

This is a lymphoepithelial organ arising as a pouch from the dorsal part of the cloaca in birds. Its development, structure and function parallel those of the thymus. It originates from an epithelial anlage, becomes a lymphoid organ by about the 15th day of embryonation, develops full functional ability near hatching and starts involuting by 7-13 weeks of age. Corresponding to the age of puberty, this bursa is also a site of lymphocytic proliferation and differentiation, stem cells from the yolk sac, fetal liver and bone marrow enter the bursa, proliferate and develop into immunocompetent 'bursal lymphocytes' or B-cells (B for Bursa or Bone marrow). These migrate and seed selective areas in the peripheral lymphoid organ the mantle, the germinal follicles and perifollicular regions of the spleen, and the outer cortical areas and medullary cords of lymph nodes, these are known as bursa dependent or thymus independent areas following appropriate antigenic stimulation, B-lymphocytes transform into plasma cells and secrete antibodies.

The vital role of the bursa in humoral immunity was discovered accidentally by Glick and Chang (1956) when they found no antibody response to a bacterial antigen in chickens which had been bursectomized at hatching. Immunocompetence is conferred on the lymphocytes by the bursa in stages, competence for IgM production is acquired early (about the 14th day of embryonation) and for IgG late (about the 21st day synthesis IgM, but not IgG).

