# Q9 Describe how the body defends against infection (Sept 2010)

#### Non immune defences ->

- Physical barriers to infection →
  - Skin (prevents bacterial entry and growth)
  - Normal flora (prevent transient pathogenic organisms from colonizing the skin surface by competing for nutrients or secreting protective enzymes)
  - Mucous (traps bacteria and foreign particles)
  - Cilia (removes mucous and trapped particles)
  - Hydrochloric acid secreted in stomach (bactericidal)

## Immune defences →

#### INNATE IMMUNITY

- Phagocytes → digest and destroy microorganisms via IFN; also contribute to cell mediated immunity by presenting antigens to lymphocytes
- Natural killer cells → lymphocytes which are not B or T cells; recognize virus-infected cells and destroy them by releasing proteins that induce apoptosis
- Complement system → a collection or circulating and membrane associated proteins that, when triggered (by the alterntaivem classical or lectin pathways), results in the formation of a Membrane Attak Complex and cell apoptosis
- Acute phase proteins → A collection of proteins which play various roles (eg, CRP activates the classical complement pathway, fibronectin binds bacteria and macrophages)
- Lysosomes → present in saliva, mucous and tears → destroy bonds in bacteria wall, causing lysis

## CELL MEDIATED (ACQUIRED) IMMUNITY

- T cells → activated by the presentation of a microorganisms via the Major Histocompatibility complex MHC of an Antigen Presenting Cell. Several subtypes:
  - CD4 T cell
    - CD4 Th1 → activated by the presentation of an antigen on an MHC II molecule. Th1 cells are proinflammatory, activate macrophages and stimulate antibody production by B cells.
      Major cytokines involved are IFNy and IL12
    - CD4 Th2 → also activated by MHC II molecules but produce IL4 and IL5 to stimulate eosinophils and IgE, and maturation of B cells into antibody producing and memory B
  - CD8 T cell → activated by the presentation of an antigen on an MHC I molecule. Utilises IFNγ and IL15 to induce apoptosis of infected cell and the proliferation of memory T cells.
- B cells in the presence of costimulatory molecules from T cells and other immune cells, differentiate into either plasma cells (capable of producing specific antibodies, which coat or opsonize microorganisms for phagocytosis) or memory cells