The Immunology of Life

What happens to your immune system as you age?

The immune system is composed of a *multitude of different cells* that interact with each other, either directly or through signaling proteins, to protect you from foreign organisms and maintain the necessary balance (homeostasis) of your body. As you grow and undergo experiences in life, *many changes occur in your immune system* that influence how you may respond to environmental stimuli. We share some of the changes that occur during life below.



Innate System

 First line of defense
Non-specific response
No immune memory generated

Adaptive System

Slower response time
Highly specific

Infancy

- Higher chance of viral and bacterial infections due to lower innate system function
- Reduced innate cell function includes: Neutrophils – poor bacteria killing abilities
 Monocytes and Macrophages – weak tissue repair and ingestion of invading pathogens
- T cells: Initial T-cell stock built in early life
- B cells: Limited neonatal antibody production

Adolescence

- During puberty, there is a maturation of innate immune cells in the central nervous system (e.g. microglia) that support puberty associated brain development
- Maturation and functional activation of innate cells continues until adulthood
- T cells: T cell production, selection and maturation continues throughout childhood but slowly ceases after puberty
- **B cells:** Approaching adulthood,

Adulthood and Old Age

- Chronic inflammation due to the presence of pro-inflammatory proteins (cytokines) known as inflammaging
- Reduced function of innate immune cells (e.g. macrophages, neutrophils) that are a necessary first line of defense
- Enhanced repertoire of memory cells for ongoing immunity against harmful organisms (*pathogens*)
- **T cells:** More dysfunctional. Impact immune function and **increase risk**

response
Generates
immune memory

Environmental Factors

• External impacts • on the immune • system

Disease Susceptibility

 Rate of disease (autoimmune, infections, cancer) Maternal antibodies (transferred to the fetus through the placenta before birth) serve as protection after birth for several months, until the adaptive immune system develops more

Initial exposures to diverse microbes (e.g. via birth mode, food, and environment) that colonize mucosal surfaces help shape the immune system and reduce risk of certain diseases

 An underdeveloped immune system leads to increased susceptibility to infections, decreased vaccine effectiveness, and increased risk of developing allergies and/or asthma in early infancy due to improper protein recognition by the immune system antibody reponses are **quicker**, **stronger**, **more specific and more durable** than those elicited in infants

Challenges to the immune system (social stress, drug use, injury) affect cell development, such as microglia in the central nervous system which can alter behaviour in adulthood

Ο

From now into adulthood, females have higher incidences of developing inflammatory diseases and of autoimmune diseases (e.g. Type 1 Diabetes, Multiple Sclerosis, Psoriasis), while males have higher cancer incidence and severe infection outcomes

of infections

 B cells: Inflammatory B cells and antibody producing cells can expand during aging and contribute to inflammaging

 Viral infections during life can negatively impact immune cell function with age

 Changes to microbes in the gut are linked to inflammation in older adults

 Reduced immune function can lead to an increased susceptibility to infections

• Reduced protective immune responses to vaccination

 Adults experience an increase in chronic diseases (e.g. diabetes) that may be impacted by immunity



Sex-related Differences

Immune
changes based
on sex

Physiological Changes



• Males: stronger innate immune responses

 Females: greater T helper cell populations (cells that assist other cells to generate immune reponses) but lower cytotoxic T cell populations (cell that directly kill infected cells)

 The thymus is largest at birth, T cells develop in the thymus which is important for the development of the adaptive immune system

- Sex hormones regulate immune function and cortisol levels that can lead to immune suppression
- Males: stronger cell-based immune responses
- Females: more effective antibody responses
- There is an achievement of fertility and the development of secondary sex characteristics
 - In later adolescence, the thymus, a primary immune tissue, begins to shrink and is replaced with fat

- Males: worsened decline in T cell function and higher proinflammatory cytokine levels
- Females: have stronger antibody (humoral) responses which can be beneficial (prevent infection) or detrimental (autoimmunity)

 During aging, there is increased tissue scarring and reduced division of cells, disruptions in the metabolism of several tissues, and alterations to cognitive and behavioural patterns which all impact health



Created by: Baweleta Isho, Christina Ditlof, Jawairia Atif, and Saad Khan Contact: baweleta.isho@mail.utoronto.ca for any questions. Some pamphlet images prepared on Biorender.com