



Review Article

Immunity and Autoimmune system-a review

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ABSTRACT

Infection happened once a microorganism invaded the body cells and reproduced. Infection sometimes caused associated degree of responses. If the reaction was swift and efficient, the infection would be eradicated or controlled before the illness develops. In most cases, infections led to sickness. Diseases occurred once Immunity is minimal or non-existent, once the microorganism's pathogenicity (its proclivity for wreaking havoc on host cells) is increased, and after the number of infections in the body has reached a critical level. betting on the communicable disease, symptoms vary greatly. The immune system is weakened by major immunological disease disorders, also known as primary immune diseases or primary immunological disorder, which makes infections and other health issues more likely. Fever and fatigue are common signs that the immune system isn't working properly, despite the fact that the symptoms of immune diseases vary. Immune deficits are usually detected by blood tests that assess the number of immune components present or their usefulness.

INTRODUCTION

Immunity defined as a balanced condition in which cellular entities had sufficient defences to resist infection, illness, or other undesirable biological invasions while still having sufficient tolerance to prevent hypersensitive reactions and response disorders [1]. The 'behavioural immune system' consists of mechanisms that evolved as a method of facilitating behaviours that reduced infection risk and increased fitness. Many factors like sleep, diet, stress and hygiene had an effect on the immune system's performance, and any offsets in these behaviours would cause mayhem on immune perform [2]. Hand laundry (Personal & Environmental Hygiene), Sleep Cycles & hydrocortisone Levels; The system is influenced by the sleep-wake cycles of our unit of time rhythms. Studies counselled that whereas we were sleeping, we had got attenuated levels of the strain internal secretion hydrocortisone, which might suppress immune perform, and accumulated signals that activate the system. Nutrients from Food and Supplement Intake greatly affected the system [3]. Fever was a frequent reaction to infections; a further temperature was to amplify the response and provide microorganisms with a hostile environment. Inflammatory, or swelling caused by an increase of fluid within the infected region, might indicate that white blood cells were attacking and involved in the reaction. Vaccination worked by inducing a specific reaction that results in the production of B and T cells that are associated to a certain pathogen. These lymphocytes cells triggered a quick and efficient reaction when the body came into contact with the microbe [4].

Autoimmune Diseases

A condition in which the immune system to attack your body imprecisely is known as an autoimmune disorder (AD). In normal circumstances, the system defends against pathogens like bacteria and viruses. When it recognised these foreign invaders, it sends out an army of fighter cells to combat them. In most cases, the team was able to distinguish between invader and your own cells. However, with AD, the immune system of an individual misidentifies a biological part, including such the joints or skin, as a foreign microbe. Autoantibodies are proteins that the body produces that attack healthy cells. Some response illnesses were limited to a single organ. The exocrine gland is harmed by one polygenic



illness. Alternative illnesses, such as systemic lupus erythematosus (SLE) or cancer, affects the whole body [5]. There are quite a number of completely different reaction diseases. Here, we list a number of the foremost common ones.

Type I Diabetes

Insulin is produced by the duct gland and aids in the regulation of glucose levels. In type one diabetes, the system attacks and destroys insulin-producing cells within the duct gland. High glucose levels injure blood vessels as well as organs of the human body, kidneys, eyes, and nerves [6].

Rheumatoid arthritis

The immune system assaults the joints in AD. As a result of the invasion of foreign particles, the joints get red, swelled, hot, hurting, and stiff. Unlike arthritis, which normally affects people as they become older, RA can strike as early as your 30s or even earlier [7]. In a very explore for aspects of behaviour associated with an ovulatory drug use which could justify the favourable impact of contraceptive method on the onset of RA, coffin nail smoking and alcohol consumption square measure doable risk factors. The adjusted risk of RA in girls United Nations agency preserved a minimum of one coffin nail on a daily basis was zero.61 (95% confidence interval (CI) zero.42 to 0.89). The low incidence of alcohol consumption within the patients with RA can be because of the termination of alcohol consumption when malady onset. The low incidence of coffin nail smoking within the patients may mirror a protecting impact of coffin nail smoking on RA onset, probably induced by changes within the system [8].

Psoriasis/psoriatic arthritis

When skin cells are no longer needed, they multiply and subsequently shed. When you have a skin condition, your skin cells proliferate too quickly. The extra cells pile up and cause inflammatory red areas on the skin, which are usually accompanied by silver-white plaque scales. Swelling, stiffness, and discomfort in the joints can occur in up to 30% of people with skin disease. Rheumatoid arthritis is the name for this sort of sickness [9].

Multiple sclerosis

Multiple sclerosis (MS) affects the case in your central nervous system, which is the protective covering that covers nerve cells. The transmission speed of signals between your brain and funiculus to and from the rest of your body is slowed by damage to the case. Symptoms of this injury include weakness, balance issues, and difficulty walking. The illness manifests itself in a variety of ways and progresses at varying speeds. According to 2012 research, around half of those with MS aim to improve their walking abilities within fifteen years of the disease starting [10].

Systemic lupus erythematosus (SLE)

Though lupus was first identified as a skin condition in the 1800s due to the rash it causes, the general kind, which is the most prevalent, affects a variety of organs, including the joints, kidneys, brain, and heart. The most frequent symptoms are joint discomfort, weariness, and rashes [11].

Inflammatory bowel disease

Inflammatory bowel disease (IBD) is a term used to describe diseases in which the lining of the internal organ wall becomes inflamed. Every type of IBD affects a different section of the canal. Crohn's disease can cause inflammation anywhere throughout the canal, from the mouth to the asshole. Colitis is a condition that affects just the lining of the large intestine (colon) and a body component [12].

Addison's disease Adrenal glands are glands that create hormones corticosteroid and mineralocorticoid as well as steroid hormones, are affected by Addison's disease. If you don't get enough corticosteroids, it will affect how your body consumes and stores carbs and sugar (glucose). Mineralocorticoid deficiency can lead to atomic number 11 loss and excess metal in the blood. Weakness, weariness, weight loss, and low glucose are all symptoms [13].

Graves' disease

Graves' disease is a disorder which effects and involves the ductless gland in the neck, causing it to produce an excessive quantity of hormones. Thyroid hormones regulate metabolism, which is how the body uses energy. An overabundance of those hormones stimulates your body's activity, causing symptoms such as anxiousness, a racing heart, heat intolerance, and weight loss. Bulging eyes, often known as symptom, is one possible sign of this illness [14]. Sjogren's syndrome The glands that lubricate the eyes and lips are attacked by this illness. Sjogren's syndrome is characterised by dry eyes and xerostomia, although it will also affect the joints and skin [15].

Hashimoto's thyroiditis

Hashimoto's redness is caused by a deficiency in internal secretion production. Symptoms include weight gain, cold intolerance, exhaustion, baldness, and thyroid swelling (goitre) [16].

Myasthenia gravis

The neuromuscular junction is affected by illness, which disrupts nerve signals that help the brain govern the muscles. Signals can't drive muscles to contract if connection between neurons and muscles is disrupted. Muscle weakness is the

most prevalent symptom, which worsens with activity and improves with rest. Muscles that control eye motions, protective fold gap, swallowing, and facial movements are generally affected [17].

Autoimmune vasculitis

Inflammation occurs when the immune system attacks blood vessels. As a result of the inflammation, the arteries and veins constrict, allowing less blood to flow through them [18].

Pernicious anemia

This problem results in a lack of a super molecule called factor, which is produced by the lining cells of the abdomen and is essential for the small internal organ to absorb B-12 from meals. Without enough of this food, one might get anaemia, and the body's capacity to produce polymers correctly will be harmed. Malignant anaemia is more frequent among the elderly. According to 2012 research, it affects 0.01% of people in the general population, but over 2% of people over sixty [19].

Celiac disease

People who have stomach discomfort can't eat meals that include protein, a macromolecule present in wheat, rye, and other grains. When protein enters the gut, the body's immune system targets this area of the digestive tract, causing inflammation [20]. Cancer; Cancers that are bound, such as leukemia, malignant neoplastic illness, and malignant neoplasm, have a direct influence on the system. The tumour developed when immune cells expand out of control [21].

Sepsis

Infection is your body's immune system's wonderful response to an infection. The body's response to the infection causes widespread inflammation, which puts in motion a series of events that leads to organ damage, failure, and death [21].

Autoimmunity

The reaction in illness is similar to that in infection, with the exception of certain self-antigens are (or will be) a target for the immune system. Certain self-antigens might trigger a process within a particular organ, such as the thyroid (Grave's disease, Hashimoto's thyroiditis) or the brain (Grave's disease, Hashimoto's thyroiditis) (multiple sclerosis). Or, in response to them, a general inflammatory state may develop (e.g., general LE [SLE]) [22]. When our immune system assaults our own tissues, we get autoimmune disease. T-cell receptors and B-cell receptors target particular antigens, as do all adaptive immune responses. In contrast to infection, these cells recognise antigens from proteins inside the organ, which triggers a persistent inflammatory response that affects the tissue's normal function [15].

Immunodeficiency

Many persons with basic immunological disorders are born without some of the body's inherent defences or with a malfunctioning immune system, making them more vulnerable to bacteria that cause illnesses [17]. There are about 300 different types of primary immunological disease disorders, and researchers are continually discovering new ones. They will be divided into six teams, each supporting a different aspect of the system: Defective phagocytes, Complement deficits, B cell (antibody) deficiencies, T cell deficiencies, Combination B and lymphocyte shortages, Unknown (idiopathic) [18].

Allergies

When your body misinterprets a normally innocuous chemical for a harmful invader, a hypersensitive reaction occurs. Antibodies are then produced by the system to keep an eye out for that specific chemical. These antibodies will unharness a range of system chemicals, such as amine, that trigger hypersensitive response symptoms when you are exposed to the toxin again. Common allergens that trigger hypersensitivity reaction includes, Allergens in the air, such as spore, animal dander, dirt mites, and moulds; Peanuts, tree nuts, wheat, soy, fish, shellfish, eggs, and milk; Stings from insects such as bees and wasps; Antibiotics, particularly penicillin-based antibiotics, are among the most often prescribed medications; Latex or other things you ingested might create allergic responses on your skin [13].

Lupus

Lupus develops when your body's immune system destroys healthy tissue (autoimmune disease). Lupus is very certainly caused by a combination of your genetics and your environment. It appears that people with a genetic susceptibility to lupus may get the disease if they inherit contact with anything in the environment that causes lupus. However, in the vast majority of instances, the cause of lupus is unclear. Daylight is one of the probable causes. In those with lupus, exposure to the sun may cause skin lesions or induce an indoor reaction. Infections. In some people, having an associate degree infection might trigger lupus or induce a recurrence. Medications. Blood pressure meds, anti-seizure drugs, and antibiotics are all known to cause lupus. Symptoms of drug-induced lupus generally improve after the patient stops taking the medication. Symptoms may continue even after the medicine is stopped in rare cases [23].

Alzheimer's disease

The system, in addition to memory and the brain, deteriorates with age. Immunity could be a consider heart condition, stroke, cancer, disease, polygenic disease and Alzheimer's (AD). This level could also be a result of cellular or mitochondrial pathology. These ageing consequences raise inflammation in the body over time, which can conjointly have an effect on the progression of a human malady state. Changes within the body related to aging, like a weaker system, square measure inflicting the build-up of the supermolecule related to Alzheimer's malady. The disease of the system produced by aberrant amide folding inside the central nervous system is one of the most critical progressors of AD. Despite the fact that the systems of elderly patients are more vulnerable in terms of operation, immune regulation and therapy are now recognised as some of the most effective treatments for age-related disorders [14]. The system's function in the degenerative process of Alzheimer's disease is notable, but it is by no means limited to the brain. Numerous findings from clinical and experimental studies point to an important, but largely ignored, factor in the pathogenesis of Alzheimer's disease: general immune signals originating outside the brain. The importance of a close practical connection between the system and the central system is becoming increasingly apparent. Aging is the most significant risk factor in late-onset AD. We have a propensity to outline the distinct molecular and cellular modifications within the edge of ageing individuals and AD patients here, from an associate medicine standpoint. Furthermore, we have a propensity to discuss recent findings that link certain peripheral system arms and different types of immune responses to the modulation of AD development. Taken together, our findings highlight the dynamic importance of a symphony of brain-extrinsic, peripheral signals in ageing and chronic CNS processes. We have a propensity to assume that a scientific paper combining a large number of current findings will help guide the development of next-generation therapy and shape future AD research strategies [18].

The Mucosal Immune System and its Autophagy Regulation

The alimentary canal poses a unique challenge to the membrane system, which must constantly monitor the vast surface for the presence of pathogens while maintaining tolerance to helpful or harmless antigens. Many aspects of membrane immune responses are influenced by the method of autophagy, according to recent research. Initially thought to be a "self-eating" survival system that allows for nutrition use during famine, autophagy has since been linked to a variety of cellular responses, as well as many elements of immunology. The discovery that autophagy will degrade animate creature microorganisms led to the first linkages between autophagy and host immunity. Following research revealed that autophagy affects substance processing, thymic choice, leukocyte balance, and the control of immunological serum globulin and protein release, indicating that it plays a much larger role in immune responses. This animation depicts the inflammatory cycle that occurs in the brains of Alzheimer's sufferers [21].

Diagnosis & Treatment

Allergic diseases can also be assessed using blood tests or allergic response skin testing to determine which allergens cause symptoms. Medication that reduces the reaction, such as corticosteroids or other immune-suppressive drugs, may be extremely beneficial in active or reactive circumstances [22]. "In some immune deficiency disorders, replacing of missing or deficient components might potentially be a therapeutic," Lau added. "Antibody infusions to combat infections might possibly be involved." Antibodies from the organism may also be used in treatment. A monoclonal antibody is a type of macromolecule generated in a research facility that binds to molecules in the body. They'll be used to modulate the components of the response that cause inflammation. Biological antibodies are being used to treat cancer, according to the National Cancer Institute. They'll deliver drugs, poisons, and heated things to cancer cells [23].

REFERENCES

1. Zimmermann P, Curtis N. Factors That Influence the Immune Response to Vaccination. *Clin Microbiol Rev.* 2019;32(2):e00084-18. doi: 10.1128/CMR.00084-18.
2. Pinckard RN, Weir DM & McBride WH. Factors influencing the immune response. I. Effects of the physical state of the antigen and of lymphoreticular cell proliferation on the response to intravenous injection of bovine serum albumin in rabbits. *Clinical and experimental immunology*, 1967,2(3), 331–341.
3. Mombeini T, Hashabi F. Review of pharmacological effects of phytoestrogens on brain and behavior. *Basic & Clinical Neuroscience* 2014. 2014 Oct 29.
4. Alberts B, Johnson A, Lewis J, et al. *Molecular Biology of the Cell*. 4th edition. New York: Garland Science; 2002. Lymphocytes and the Cellular Basis of Adaptive Immunity. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK26921/>

5. Jin R, Yang G, Li G. Inflammatory mechanisms in ischemic stroke: role of inflammatory cells. *Journal of leukocyte biology*. 2010;87(5):779-89. doi: 10.1189/jlb.1109766.
6. Dregan A, Charlton J, Chowienczyk P, Gulliford MC. Chronic inflammatory disorders and risk of type 2 diabetes mellitus, coronary heart disease, and stroke: a population-based cohort study. *Circulation*. 2014;130(10):837-44. doi: 10.1161/CIRCULATIONAHA.114.009990.
7. Escames G, López LC, García JA, García-Corzo L, Ortiz F, Acuña-Castroviejo D. Mitochondrial DNA and inflammatory diseases. *Human genetics*. 2012;131(2):161-73. doi: 10.1007/s00439-011-1057-y.
8. Liu X, Tedeschi SK, Barbhuiya M, Leatherwood CL, Speyer CB, Lu B, et al. Impact and Timing of Smoking Cessation on Reducing Risk of Rheumatoid Arthritis Among Women in the Nurses' Health Studies. *Arthritis care & research*. 2019;71(7):914-24. doi: 10.1002/acr.23837.
9. Hardy J. The Amyloid Hypothesis of Alzheimer's Disease: Progress and Problems on the Road to Therapeutics. *Science*. 2002;297(5580):353-356. doi: 10.1126/science.1072994.
10. Selkoe DJ, Hardy J. The amyloid hypothesis of Alzheimer's disease at 25 years. *EMBO molecular medicine*. 2016 Jun;8(6):595-608. doi: 10.15252/emmm.201606210.
11. Kabat AM, Pott J, Maloy KJ. The mucosal immune system and its regulation by autophagy. *Frontiers in immunology*. 2016;7:240. doi: 10.3389/fimmu.2016.00240.
12. Yap YA, Mariño E. An insight into the intestinal web of mucosal immunity, microbiota, and diet in inflammation. *Frontiers in immunology*. 2018:2617. doi: 10.3389/fimmu.2018.02617.
13. Blom M, Zetterström RH, Stray-Pedersen A, Gilmour K, Gennery AR, Puck JM, et al. Recommendations for uniform definitions used in newborn screening for severe combined immunodeficiency. *Journal of Allergy and Clinical Immunology*. 2021. doi: 10.1016/j.jaci.2021.08.026.
14. Weetman AP. Graves' disease following immune reconstitution or immunomodulatory treatment: should we manage it any differently?. *Clinical Endocrinology*. 2014;80(5):629-32. doi: 10.1111/cen.12427.
15. Kramer JM. Early events in Sjögren's Syndrome pathogenesis: The importance of innate immunity in disease initiation. *Cytokine*. 2014;67(2):92-101. doi: 10.1016/j.cyto.2014.02.009.
16. Ehlers M, Schott M. Hashimoto's thyroiditis and papillary thyroid cancer: are they immunologically linked?. *Trends in Endocrinology & Metabolism*. 2014;25(12):656-64. doi: 10.1016/j.tem.2014.09.001.
17. Berrih-Aknin S, Le Panse R. Myasthenia gravis: a comprehensive review of immune dysregulation and etiological mechanisms. *Journal of autoimmunity*. 2014;52:90-100. doi: 10.1016/j.jaut.2013.12.011.
18. Hid Cadena R, Abdulahad WH, Hospers GA, Wind TT, Boots AM, Heeringa P, Brouwer E. Checks and balances in autoimmune vasculitis. *Frontiers in Immunology*. 2018;9:315. doi: 10.3389/fimmu.2018.00315.
19. Zulfiqar AA, Andres E. Association pernicious anemia and autoimmune polyendocrinopathy: a retrospective study. *Journal of medicine and life*. 2017;10(4):250. PMID: PMC5771255.
20. Mayassi T, Ladell K, Gudjonson H, McLaren JE, Shaw DG, Tran MT, et al. Chronic inflammation permanently reshapes tissue-resident immunity in celiac disease. *Cell*. 2019;176(5):967-81. doi: 10.1016/j.cell.2018.12.039.
21. Chen DS and Ira M. Oncology Meets Immunology: The Cancer-Immunity Cycle. *Immunity*. 2013, 39(1):1-10. doi.org/10.1016/j.immuni.2013.07.012.
22. Herrada AA, Escobedo N, Iruretagoyena M, Valenzuela RA, Burgos PI, Cuitino L & Llanos C. Innate Immune Cells' Contribution to Systemic Lupus Erythematosus. *Frontiers in Immunology*, 2019,10: 772. doi.org/10.3389/fimmu.2019.00772
23. Pan L, Lu MP, Wang JH, Xu M, Yang SR. Immunological pathogenesis and treatment of systemic lupus erythematosus. *World Journal of Pediatrics*. 2020;16(1):19-30. doi: 10.1007/s12519-019-00229-3.