Interleukin-17 in human inflammatory diseases

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Abstract
Human Th17 pro-inflammatory cells are currently defined as cells that produce IL-17A and F, tumor necrosis factor (TNF)-α, IL-6, IL-21, IL-22 and IL-23. Recently discovered related molecules are forming a family of cytokines, the IL-17 family, IL-17A, IL-17B, IL-17C, IL-17D, IL-17E and IL-17F. The associated receptors for the IL-17 family identified are IL-17R, IL-17RH1, IL-17RL (receptor like), IL-17RD and IL-17RE. This review introduces the roles of IL-17 and Th17 cells in human autoimmune diseases. Studies have shown that T cells with inflammatory effects on epithelial, endothelial and fibroblast cells express IL-17. Th17 cells are supposed to be involved in various autoimmune diseases, such as rheumatoid arthritis, psoriasis, multiple sclerosis, and inflammatory bowel diseases. Based on the biologic functions and regulation, IL-17 has regulatory roles in host defense and chronic inflammation which result in tissue damage and autoimmunity. So the IL-17 links links innate and adaptive immunity and has both beneficial and pathological effects on the immune system. This paper will focus on the possible roles of IL-17 in autoimmune diseases, a fundamental player in immune regulation.

Key words: interleukin-17, T cells, inflammation, cytokines, cytokine receptors, autoimmunity.

Identification of interleukin-17, T cells, inflammation, cytokines, cytokine receptors, autoimmunity.

Two decades ago, Mossman and Coffman suggested that CD4⁺ T cells differentiate into two subsets with common functions and patterns of cytokine secretion, characterized by CD4⁺ T helper 1 (Th1) or T helper 2 (Th2) cells. Th1 cells are designated by production of interleukin-2 (IL-2) and interferon-γ (IFN-γ) and induction of cell-mediated immunity against intracellular pathogens and generation of delayed type hypersensitivity responses, while Th2 cells secrete interleukin-4 (IL-4), IL-5, and IL-13 through signal transducer and activator of transcription 4 (STAT4), and stimulate humoral immunity against parasitic helminthes [1].

Although CD4⁺ T-cell subpopulations such as T helper 3 (Th3) cells, T regulatory type 1 (Tr1) cells, transforming growth factor (TGF)-induced regulatory T cells (iTregs), regulatory T cells developed in the thymus (natural or nTregs) and follicular helper T (TFH) cells have been described, they are not defined as a separate lineage. IL-17 producing T cells represented a subset distinct from other CD4⁺ T-cell subsets, known as Th17. Th17 cells provide protection against some infection and are associated to the development of autoimmune diseases. Th17 cells as a pleiotropic cytokine can trigger an acute inflammatory response that is dominated by recruitment of granulocyte lineage, especially neutrophils against extracellular bacteria and some fungi [2]. IL-17 was defined as a proinflammatory cytokine produced by activated T-cells in response to stimulation through the T cell receptor. Th17 cells secrete not only IL-17A, but also IL-17F, IL-21, IL-22 and IL-23; these cytokines most likely cooperate to induce multiple inflammatory and hematopoietic effects on epithelial, endothelial, and fibroblastic cells [3].

Pathogens can induce Th17 response, including gram-positive Propionibacterium acnes, gram-negative Citrobacter rodentium, Porphyromonas gingivalis, Klebsiella pneumoniae, bacteroides species and Borrelia species,
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*Mycobacterium tuberculosis*, parasitic infections like *Toxoplasma gondii*, and fungi such as *Candida albicans* [4, 5]. Now, several autoimmune diseases are supposed to be Th17-mediated diseases, because the biologic functions of IL-17 are consistent with the chronic and destructive nature of inflammation. This review introduces the roles of IL-17 and Th17 cells in human autoimmune diseases.

The IL-17 family and its receptors

IL-17 is a prototype member of the IL-17 family of cytokines, which contains six structurally related isoforms: IL-17A, IL-17B, IL-17C, IL-17D, IL-17E (IL-25), and IL-17F that share 20–50% homology (conserved cysteines) to IL-17. IL-17E (or IL-25) is not produced by Th17 cells, but it is produced by Th2 cells. IL-25 induces the expression of Th2-type cytokines and chemokines such as CCL5 (RANTES) and CCL11 (Eotaxin) [6]. The human IL-17 gene was mapped on human chromosome 6p12. The gene product is a protein of 150 amino acids with a molecular weight of 15 kDa and is secreted as a disulfide linked dimer of 30–35 kDa glycoproteins. IL-17 polypeptide weight of 15 kDa and is secreted as a disulfide linked product is a protein of 150 amino acids with a molecular weight of 33–60 kDa. IL-17RD reveals a remarkable relationship between the IL-17R family and conserved molecules in immunity. IL-17RC (IL-17RL) is 22% identical and 34% similar to IL-17RA sequences. IL-17RB signaling promotes Th2-type immunity. IL-17RC (IL-17RL) is 22% identical and 34% similar to the human IL-17RA with molecular weights of 33–60 kDa. IL-17RD reveals a remarkable relationship between the IL-17R family and conserved molecules involved in embryonic development. Whereas IL-17RA and IL-17RC are the receptors for IL-17A and IL-17F, IL-17RB is not only the receptor for IL-17E (IL-25) but also binds IL-17B with low affinity [14].

IL-17 family members have one transmembrane glycoprotein receptor) seems to comprise two IL-17RA subunits and one IL-17RC subunit. IL-17RC exists in a large number of splice isoforms that differ in the extracellular domain. IL-17RA is expressed in hematopoietic tissue highly in response of IL-17A in epithelial, endothelial, fibroblast cells, macrophages, and DCs [13]. IL-17B binds to IL-17 with a relatively high affinity. The IL-17RB protein has 19.2% protein sequence identity to the human IL-17RA sequences. IL-17RB signaling promotes Th2-type immunity. IL-17RC (IL-17RL) is 22% identical and 34% similar to the human IL-17RA with molecular weights of 33–60 kDa. IL-17RD reveals a remarkable relationship between the IL-17R family and conserved molecules involved in embryonic development. Whereas IL-17RA and IL-17RC are the receptors for IL-17A and IL-17F, IL-17RB is not only the receptor for IL-17E (IL-25) but also binds IL-17B with low affinity [14].

The receptor for IL-17A (type I trans-membrane glycoprotein receptor) seems to comprise two IL-17RA subunits and one IL-17RC subunit. IL-17RC exists in a large number of splice isoforms that differ in the extracellular domain. IL-17RA is expressed in hematopoietic tissue highly in response of IL-17A in epithelial, endothelial, fibroblast cells, macrophages, and DCs [13]. IL-17B binds to IL-17 with a relatively high affinity. The IL-17RB protein has 19.2% protein sequence identity to the human IL-17RA sequences. IL-17RB signaling promotes Th2-type immunity. IL-17RC (IL-17RL) is 22% identical and 34% similar to the human IL-17RA with molecular weights of 33–60 kDa. IL-17RD reveals a remarkable relationship between the IL-17R family and conserved molecules involved in embryonic development. Whereas IL-17RA and IL-17RC are the receptors for IL-17A and IL-17F, IL-17RB is not only the receptor for IL-17E (IL-25) but also binds IL-17B with low affinity [14].

IL-17RA and IL-17F are also key cytokines for the recruitment, activation, and migration of neutrophils. IL-17, through modulation of chemokine activity, was described as a factor, which leads to the formation of germinal centers (GC) of lymph follicles containing B cells within GCs and increasing somatic hyper mutation. Follicular helper cells express CXCR5, respond to the lymph follicle, associated with chemokine CXCL13 and home, and help create the light zone of GCs, where further help can be given from cognate B cells that have undergone immunoglobulin isotype switching and somatic hyper mutation in the GC dark zone. Thus, T follicular helper cells in the GC light zone induce further differentiation and selection of B cells. Both Th17 and T follicular helper cells are the main source of IL-21, and thus both play a significant role in setting up productive GC reactions [10, 11].
**Differentiation of Th17 cells**

Th17 cells as a distinct subpopulation of CD4+ T helper cells are specialized by the unique identification of differentiation factors and transcription factors. Tumor necrosis factor-α (TNF-α), IL-1β, transforming growth factor-β (TGF-β) and IL-6 can enhance Th17 differentiation [15]. IL-17 does not inhibit Th1 or Th2 differentiation so Th1 and Th2 cells typically dominate over Th17 cells. One of the ways in which TGF-β can promote Th17 differentiation is suppressing the production of the inhibitory cytokines IFN-γ and IL-4. TGF-β synergizes with IL-6 to induce expression of the transcription factor RORγt, a key regulator of Th17 differentiation [16, 17]. In humans, retinoic acid receptor-related orphan receptor-γt (RORγt), IFN-regulatory factor 4 (IRF-4), aryl-hydrocarbon receptor (AHR), and the transcription factor signal transducer and activator of transcription 3 (STAT3) expression are induced by IL-1β, IL-6, and IL-23 [18–20].

**IL-17 signaling pathways**

Studies have demonstrated that IL-17 is capable of activating nuclear factor κB (NF-κB) transcription factors in many cell types, including fibroblasts, macrophages, chondrocytes, intestinal epithelial cells, and colonic and pancreatic myofibroblasts [21]. The NF-κB has been activated in response to IL-17D, IL-17E and IL-17F. IL-17 receptor (IL-17R) activates extracellular signal-regulated protein kinase (ERK1 and ERK2), stress-induced c-Jun N-terminal kinases (JNK-1 and JNK-2), and mitogen-activated protein kinases (p38 MAPKs) pathways. These signaling pathways result in up-regulation of IL-6, IL-1, and NF-κB [22].

**The cellular source of IL-17**

IL-17 cytokines are now known to be secreted by other cell types apart from CD4+ T cells, γδ T cells, natural killer T (NKT) cells, natural killer (NK) cells, monocytes, macrophages, dendritic cells (DC), microglia, neutrophils, eosinophil, astrocytes, and oligodendrocytes. Thus, cells of both the innate and the adaptive immune systems as well as non-immune cells [23] produce IL-17A and IL-17F.

**Cytokines expressed by Th17 cells**

Th17 lineage expresses IL-21 after activation and autocrine IL-21 plays a significant role in RORγt and IL-17 expression [24]. After IL-6 or IL-21 induces IL-23 receptor expression, IL-23 in combination with TGF-β can induce RORγt and IL-17 expression. IL-6 upregulates IL-21, then both IL-6 and IL-21 deregulate IL-23 receptor, and eventually IL-23 acts to deregulate the effectors function and pathogenicity in Th17 cells [25]. Human IL-21 deregulated IL-17F and IL-17A production and T-cell proliferation and to counteract suppression by Tregs. Other pro-inflammatory cytokines secreted from human Th17 cells include TNF-α, IL-22, and IL-26 [26]. A subset of Th17 cells co-expresses with IFN-γ and IL-10, however IL-10, an anti-inflammatory cytokine is generally supposed to include Th2 cells and various types of Tregs [27]. IL-10 produced by Th17 cells may serve as an important protective role in limiting inflammation and tissue damage which is normally caused by IL-17. Both IL-17A and IL-17F induce the production of antimicrobial peptides (β defensin-2, S100A7, S100A8, and S100A9), mucins (MUC5B and MUC5AC) [28], cytokines like IL-6, granulocyte colony-stimulating factor (G-CSF), and granulocyte-macrophage colony-stimulating factor (GM-CSF), chemokines (CXCL1, CXCL5, IL-8, CCL2, and CCL7), and matrix metalloproteinase (MMP1, MMP3, MMP9, MMP12, and MMP13) from fibroblasts, endothelial cells, and epithelial cells [29].

IL-17A induces intercellular cell adhesion molecule 1 (ICAM-1) for granulocyte recruitment and inflammation in keratinocytes, IL-1 and tumor necrosis factor (TNF) in macrophages, inducible nitric oxide synthase (iNOS) and cyclooxygenase-2 (COX-2) in chondrocytes, and COX-2-dependent prostaglandin E2-mediated receptor activator of nuclear factor κB (NF-κB) ligand (RANKL) expression in osteoblasts. Furthermore, IL-17A promotes stem cell factor (SCF) and G-CSF-mediated granulopoiesis. IL-17A is also involved in neutrophil recruitment with CXCL8 (IL-8), CXCL1 (Gro-α), CXCL2 (MIP2), and CXCL5 (LIX). CCL5 (RANTES) is involved in recruiting T cells, monocytes, basophils, and eosinophils [30].

** Trafficking of Th17 cells**

Pro-inflammatory cytokines, chemokines and chemokine receptors produced by Th17 cells may result in the inflammation and pathogenesis that is mediated by activated T cells trafficking into involved tissues. Human Th17 cells express the chemokine receptor CCR6 from healthy peripheral blood and inflamed tissue. In addition, the majority of the RORC expression was restricted to the CCR6+/CCR4+ population, with a small amount in the CCR6+ CXC3+ population [31]. CCR6 which mediates homing to skin and mucosal tissues, plays an important role in recruitment of pathogenic T cells in IL-17-associated inflammatory diseases including psoriasis, inflammatory bowel diseases (IBD), and rheumatoid arthritis (RA). The CCR6 ligand CCL20 is overexpressed in stromal cells by IL-17; Th17 cells draw further Th17 and Th1 cells to inflamed tissues [32].

**IL-17 and human inflammatory diseases**

IL-17A in humans are associated with pathology in numerous autoimmune and inflammatory conditions, such as rheumatoid arthritis (RA), multiple sclerosis (MS), psoriasis, Crohn’s disease, systemic lupus erythematosus (SLE), asthma, Behçet’s disease, and hyper IgE syndrome [33].
Rheumatoid arthritis

The RA is considered as a chronic inflammation of the synovitis of the joints. The major characteristic of inflamed articular joints is proliferating synovial fibroblasts, joint and cartilage progressive destruction, infiltrating CD4+ T cells and autoantibody-producing plasma cells (rheumatoid factor and anti-cyclic citrullinated peptide (CCP) antibodies), increased numbers of innate immune cells as well as DC, granulocytes and macrophages and ectopic germinal centers (GC) in joints [34]. High levels of IL-17 were found in the rheumatoid synovium of patients with RA, promote both inflammation, and bone degradation [35]. Th17 cells induce the production of pro-inflammatory cytokines such as TNF-α, IL-1β and IL-6 from cartilage, synoviocytes, macrophages and bone cells. Th17 cells up-regulate RANK ligand in osteoblasts, stimulate the activity of matrix metalloproteinase (MMP) 1, 2, 3, 9 and 13, extracellular matrix degradation of the joint, and bone resorption. The IL-17 also stimulates the expression of multiple chemokines such as IL-8/CXCL8, CXCL1 (KC/GRO-α), CXCL2 (MIP2α/GRO-β), CCL20 (MIP-3α), CCL2 (MCP1) and CCL7 (MCP3). These chemokines can recruit neutrophils, macrophages and lymphocytes to the synovium, thereby enhancing inflammation with more severe joint damage [36, 37].

Psoriasis

Psoriasis is a chronic inflammatory skin disorder with dermal hyperplasia and skin infiltration by immune cells, which have migrated from blood. The main histological features of psoriatic skin are epidermal keratinocyte hyper-proliferation, dermal angiogenesis, and infiltration of dendritic cells (DCs), T lymphocytes, neutrophils, monocytes and macrophages [38]. Both Th1 and Th17 cytokines have been found in psoriatic skin, as well as IL-17A, IL-17F, IL-19, IL-20, IL-22, IL-26 and TNF-α in serum and lesion skin. IL-17 and IL-22 synergistically up-regulated expression of skin antimicrobial peptides such as β-defensin-2 (BD-2), S100A7–9 (psoriasin), cathelicidin (LL37) and S100A8/9 (calprotectin) which may cause more resistant to skin infections than people without psoriasis [39].

Systemic lupus erythematosus

Systemic lupus erythematosus (SLE) is a chronic multi-organ systemic autoimmune disease caused by hyper gammaglobulinaemia and autoantibody to nuclear antigens result in inflammation and damage to multiple organs especially kidneys. Increased plasma levels of IL-17 have been identified in the serum of SLE patients as a consequence of too much IL-17 synthesis by CD4+ T cells and CD3+CD4-CD8- double negative (DN) T cells [40]. IL-17 and IL-23 can be detected in the kidneys of patients with lupus nephritis, which may contribute to renal failure [41]. IL-17 promotes B cell survival both alone and synergistically with B cell-activating factor (BAFF), to protect B cells from apoptosis, thereby increasing the number of autoantibody producing cells. Increased BAFF expression is found in 22–25% of SLE serum samples. Hence, the impact of IL-17 on B cells may explain its role in contributing to SLE pathogenesis. Th17 differentiation by IL-6, IL-21, and IL-23 induce transcription of the RORγt and increase phosphorylation of STAT3 in SLE patients [42].

Inflammatory bowel disease and Crohn’s disease

The roles of Th17 cells in inflammatory bowel disease (IBD) and Crohn’s disease (CD) are more complicated than in other autoimmune diseases [43]. In both diseases, chronic relapsing inflammatory disorder in the mucosa leads to destruction of the lamina, with complications such as perforations, and internal or external fistulas. The presence of lymphocytic infiltrates in the inflamed colonic mucosa has shown the involvement of IL-17 in the pathogenesis of inflammatory bowel disease (IBD) [44]. Furthermore, elevated serum IL-17 concentration was found in patients with active IBD in comparison to healthy individuals. The key cytokines IL-6, IL-8, IL-12, IL-17, IL-23, and MCP-1 were found overexpressed in lesions of Crohn’s disease [45].

Multiple sclerosis

Multiple sclerosis (MS) is a CD4+ T cell-mediated autoimmune disease involving the central nervous system. Migration and accumulation of inflammatory immune cells from blood occurred through the brain blood barrier. Chronic inflammation of the brain leads to the destruction of myelin sheaths leading to a decrease in influx transmission and function loss. The expression of IL-6, IL-1β, IFN-γ and, TNF-α in monocytes, microglial cells, and astrocytes was associated with active MS [46]. IL-17 over-expressed in brain biopsies of MS patients was found in the blood and cerebrospinal fluid (CSF) of MS patients. Studies on pathogenic T-cell response in MS indicated recruitment of myeloid cells like monocytes/macrophages and neutrophils to the site of the inflammation and brain lesions by development of human Th17 cells dependent on IL-1β and IL-6, IFN-γ and, TNF-α [47]. Memory T cells producing IL-17 and IL-22 disrupt the tight junctions of the blood-brain barrier and enable Th17 cells to migrate into the central nervous system (CNS) and cause neuronal loss [48].

Systemic sclerosis

Systemic sclerosis (SSc) is a progressive sclerosis of the skin, lungs, gastrointestinal tract and by microvascular abnormalities of the skin and visceral organs dysfunction which leads to progressive fibrosis, defined as scleroderma [49]. A higher expression of IL-17A and IL-17F
has been demonstrated in peripheral blood and fibrotic lesions of the skin and lungs of patients with scleroderma. Furthermore, IL-17A and IL-17F can enhance the secretion of the proinflammatory and pro-fibrotic cytokines such as IL-1α, IL-6, IL-23, IL-8, chemokines (CXC12 and its receptor CXCR4, CCL20), and adhesion molecules (L-selectin and ICAM-1) from fibroblasts and endothelial cells [50–53].

So, IL-17 as a key event in SSc pathogenesis induces synthesis of several cytokines and proteins engaged in tissue remodeling, proliferation and differentiation of B lymphocytes leading to inflammation, auto-antibody production, auto-agression, angiogenesis, and disturbed turnover of extracellular matrix, microvascular damage, and fibrosis [54, 55].

Behçet’s disease

Patients with active Behçet’s disease (BD) had significantly higher CD4 (+) CD25 (+) T cells compared with healthy controls. Noticeable increase in Th17 cells were identified in the peripheral blood of patients with active BD. Th17 cells regulate inflammation via production of distinct cytokines such as the IL-17 family. Previous studies confirmed that Th17 cells are pathological in several human autoimmune and inflammatory diseases [56]. Th17 cells predominantly produce IL-17A-F, IL-21, IL-22 and TGF-β, IL-6 and TGF-β induce the differentiation of Th17 cells from naïve T cells. A high level of TBX 21 (Th1), RORC (Th17) and Foxp3 (Treg) were confirmed in neuro-BD [57]. The presence of the IL-21 and IL17-A producing T cells was demonstrated in the cerebrospinal fluid, brain parenchyma inflammatory infiltrates, and intracerebral blood vessels of patients with active BD and central nervous system involvement. IL-21 represents a promising objective for novel therapy in BD [58, 59].

Hyper IgE syndrome

The hyper IgE syndrome, a primary immunodeficiency, is associated with very high levels of IgE, skin and lung manifestations, bone disorders and infections. IgE is the prototypic immunoglobulin produced under the action of the Th2 cytokines [60]. Recently, mutations in the STAT3 gene, one of a family of transcription activators, have been identified, which nullify the ability to increase Th17 response. Indeed defects in Th17 cells have been observed in these hyper IgE patients. These findings result from the balance between the Th17 and Th2 pathways, where the Th17 pathway inhibits the Th2 pathway and IgE production. Patients have relapsing *Candida albicans* and *Staphylococcus aureus* infection in the skin and lungs [61].

Conclusions

IL-17 and Th17 subset is a new revival of the contribution of some T cells to chronic inflammation and extra-cellular matrix destruction. A developing list of diseases has been associated with IL-17 but the final demonstration of its contribution to disease pathogenesis is still unknown. Tools are now getting ready to test these concepts in the clinic. Recent studies support the notion that deregulated production of IL-17 and IL-21, cytokines produced by Th17 cells, may participate in the pathogenesis of autoimmune diseases. Knowledge of the molecular networks responsible for the regulation of this T helper cell subset is accumulating at a rapid pace. This information will undoubtedly be critical for the development of innovative therapeutic strategies aimed at targeting autoimmune diseases.

Conflict of interest

There is no conflict of interest.

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