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## Pathophysiology and inhibition of IL-23 signaling in psoriatic arthritis: A molecular insight

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### Abstract

Psoriatic arthritis (PsA) is a chronic inflammatory arthritis of unknown etiology, and currently the cellular and molecular interactions that dictate its pathogenesis remain elusive. A role of the interleukin-23 (IL-23)/IL-23R (IL-23 receptor) interaction in the development of psoriasis and PsA is well established. As IL-23 regulates the differentiation and activation of innate and adaptive immunity, it pertains to a very complex pathophysiology involving a plethora of effectors and transducers. In this review, we will discuss recent advances on the cellular and molecular pathophysiological mechanisms that regulate the initiation and progression of PsA as well as new therapeutic approaches for IL-23/IL-23R targeted therapeutics.

### Keywords

Psoriatic arthritis; skin and joint inflammation; cytokines; IL-23/IL-23R pathways; human monoclonal IL-23 antibodies; therapeutics

## 1. Introduction

In the United States, an estimated 294,000 children under the age of 18 are affected by arthritis, including psoriatic arthritis [1] and up to 15% of patients who have psoriasis also

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Conflict of interest

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develop inflammatory arthritis [2]. Psoriatic arthritis (PsA) is a systemic inflammatory disease, which is characterized by inflammation of the skin and joints [3]. Epidermal hyperplasia, synovial inflammation, and bone loss are hallmarks of PsA [3]. Although there is no specific genetic marker for diagnosis of PsA, the disease could be diagnosed through clinical and imaging features, which include psoriasis, peripheral joint disease, axial disease, enthesitis, and dactylitis [2, 4]. Neutralization of pro-inflammatory cytokines of the IL-23/IL-17 axis has shown great promise in combating the disease [5–7]. Ustekinumab (IL-23p40) and secukinumab (IL-17A) are human monoclonal antibodies that specifically bind to and neutralize IL-23p40 and IL-17A, respectively [6, 7]. These antibodies have been in clinical trials for the treatment of PsA and improved signs and symptoms of PsA, suggesting that IL-23/IL-17 has an important role in the pathogenesis of PsA. We will describe the current understanding of pathogenesis of PsA and ongoing therapeutic developments for treatment of PsA.

## 2. Clinical and pathophysiological features of PsA

### 2.1. Clinical features

In human, skin is composed of three layers including the epidermis, dermis, and hypodermis. The layer of the epidermis includes the stratum corneum (the outermost layer), stratum granulosum, stratum spinosum, and stratum basale. The dermis layer contains connective tissue, hair follicles, and sweat glands whereas hypodermis is made of fat and connective tissue. During pathogenesis of psoriasis and PsA, each of these layers becomes altered which is one of major features for the diseases include thinned suprapapillary plates, collections of neutrophils in the stratum corneum (Munro's microabscesses), psoriasiform epithelial hyperplasia; elongation and edema of dermal papillae; parakeratosis, retained nuclei in the stratum corneum [2, 4]. In addition, bone and cartilage destruction with pathologic new bone formation is one of the most distinctive features of PsA [2]. Several clinical features of PsA has been characterized by Moll and Wright such as distal interphalangeal joints, arthritis mutilans, symmetric polyarthritis, ankylosing spondylitis, asymmetric distal interphalangeal joints [4].

### 2.2. Cellular Pathophysiology

Although the precise cellular mechanisms of PsA are not completely understood, it is likely that genetic, immunologic, and environmental factors are involved in the activation of multiple cell types including T cells, neutrophils, keratinocytes, and osteoclasts (Figure 1). In this part, we will focus on the immunopathogenesis of PsA.

**2.2.1. T cells**—IL-23 has been implicated in the regulation of conventional ( $\alpha\beta$ ) and non-conventional ( $\gamma\delta$ ) T cells and the production of IL-17. A number of reports suggest that IL-23 is critical for differentiation, survival, and expansion of Th17 and  $\gamma\delta$ T cells [8, 9].  $\gamma\delta$ T cells are significantly increased, in the dermis of psoriatic skin lesions, and in peripheral blood and synovium of PsA patients, compared with healthy control [10–12]. A number of different Th17 cell “flavors” have been described by the literature and shown mainly by intracellular staining to produce a number of cytokines as previously reviewed [13]. Th17 cells secreted their signature cytokines such as IL-17, TNF, IL-22 (Figure 1) [14,

15] and also produced RANKL [16] which have been shown to contribute to the pathology observed in PsA [17]. In rheumatoid arthritis, IL-22 regulates osteoclastogenesis by up-regulation of RANKL in human synovial fibroblasts [18] and recombinant IL-22 induced a significant increase in expression of antimicrobial peptides (S100A8, S100A9, defensin  $\beta$ 1) and inflammatory cytokines (IL-1 $\alpha$ , IL-6, and TNF- $\alpha$ ) and keratinocyte hyperplasia in a mouse model of psoriasis-like skin inflammation [19]. RANKL secreted by Th17 cells also directly promotes osteoclastogenesis, while the effect of RANKL on the skin is been linked with regulatory functions of peripheral CD4<sup>+</sup>CD25<sup>+</sup> regulatory T cells [16, 20]. Undoubtedly the most investigated cytokine of the Th17 cells is IL-17 itself, which is produced by both CD4<sup>+</sup> T cells and  $\gamma\delta$  T cells. IL-17 producing CD4<sup>+</sup> T cells are elevated in the synovial fluid of both psoriatic and PsA patients [17] and in a related animal model of IL-17 gene transfer in vivo it was shown that IL-17 induces bone loss and epidermal hyperplasia associated with PsA [21]. In keeping with these observations it was also demonstrated that  $\gamma\delta$  T cells were increased in human psoriatic skin lesions [10]. A recent study highlighted that TRAV15N-1 (V $\delta$ 6.3)  $\gamma\delta$ T cells were clonally expanded in mice and expressed specific cytokines, M-CSF, CCL5, CCL3 (Figure 1), which are known to act on myeloid cells, indicating that this  $\gamma\delta$  T cell subset might have distinct functions in myelopoiesis [22]. We also observed that IL-17 gene transfer induced epidermal hyperplasia was persistent in the absence of conventional ( $\alpha\beta$  Th17 cells) and non-conventional ( $\gamma\delta$ ) T cells but dependent on the activation of neutrophils (CD11b<sup>+</sup>Gr1<sup>high</sup>Ly6G<sup>+</sup>), suggesting that IL-17 pathology may occur independently of T cells [23] as discussed next.

**2.2.2. Neutrophils**—Although IL-23 promotes the secretion of IL-17 by CD4<sup>+</sup> T cells [24], neutrophils, not T cells, are the predominant cells producing IL-17 in human skin [25]. Interestingly, other groups have identified that neutrophils are the main source of IL-23 in the colon of patients with inflammatory bowel disease [26]. Additionally, IL-23 induces the expression of IL-23R, IL-17 and IL-22 on neutrophils (Figure 1) [27]. Furthermore, neutrophil direct role in epidermal hyperplasia is supported by the distinct presence of neutrophil exudates (Munro's microabscesses) in the stratum corneum of the epidermis in psoriatic patients [28, 29], suggesting that myeloid cells (neutrophil and monocytes) contribute directly to development of pathogenesis of psoriasis and PsA. We and others have shown that neutrophil depletion reduces thickness of epidermis and Munro's microabscess formation in mice [23, 30]. Neutrophil extracellular traps (NETs) was increased in human psoriatic skin, correlated with psoriasis disease severity, and induced the expression of  $\beta$ -defensin-2 [31].  $\beta$ -defensin-2, a serum biomarker for psoriasis, is strongly expressed in psoriatic epidermis [32]. Another study also showed that neutrophil-derived proteases such as cathepsin G, elastase, and proteinase-3 processed and activated IL-36 [33]. IL-36 and its receptor (IL-36R or IL-1RL2) are able to stimulate the expression of inflammatory cytokines including IL-17, IL-23, TNF, IL-8, and IL-6 in human psoriatic lesional skin [34, 35]. However, many questions remain regarding the signaling pathway(s) involved and the interplay of the IL-23/IL-17 axis with neutrophils.

**2.2.3. Keratinocytes**—In the skin, uncontrolled hyperproliferation and differentiation of keratinocytes results in epidermal hyperplasia associated with parakeratosis, which is one of the most significant features in human PsA [2, 4]. Therefore, markers of keratinocyte

differentiation and hyperproliferation such as keratin 16 (K16) and Ki67 have effectively been used as biomarkers in psoriasis [36]. In psoriasis, the transcription factor Nrf2 promotes proliferation of keratinocyte through controlling expression of K6, K16, and K17 (Figure 1) [37]. Inflammation of keratinocyte induces synthesis of chemokines (CXCL9, CXCL10, and CXCL11) that can recruit Th1 cells into the skin whereas CCL20, CXCL1, CXCL2, and CXCL8/IL-8 recruit Th17 and neutrophils [12]. Moreover, S100A7-S100A9, calcium-binding proteins, are specifically expressed in psoriasis and loss of S100A7-S100A9 improves symptoms of psoriasis-like disease [38]. We and others have previously shown that IL-23 can induce epidermal hyperplasia and stimulate the synthesis of leukotriene B4 (LTB4), a biologically active lipid inflammatory mediator, important in skin inflammation [39]. LTB4R1 is a high-affinity receptor for LTB4 predominantly expressed on neutrophils and macrophages [40] and it is also involved in joint pathology [41]. The interplay between IL-23 and the release of pro-inflammatory mediators in innate immunity is relatively understudied, and remains an area of increased interest.

**2.2.4. Osteoclasts**—In addition to T cells and neutrophils, osteoclasts also play a crucial role in the development of bone destruction [42]. Osteoclasts are multinucleated cells that perform the unique function of bone resorption [42]. During physiological bone remodeling, macrophage colony-stimulating factor (M-CSF) and receptor activator of nuclear factor kappa-B ligand (RANKL) regulate the differentiation of osteoclasts from their precursors [43]. RANKL selectively induces expression of nuclear factor of activated T cells cytoplasmic 1 (NFATc1) in the presence of M-CSF; importantly, *NFATc1*-deficient embryonic stem cells fail to differentiate into osteoclasts, and ectopic expression of NFATc1 causes precursor cells to undergo efficient differentiation without RANKL signaling indicating that NFATc1 is required for osteoclast differentiation [44]. Osteoclasts also highly express cathepsin K, matrix metalloproteinases (MMPs) which contribute differently to activities of osteoclast [45]. Bone resorption results in the release of tartrate-resistant acid phosphatase (TRAP) from osteoclasts, thus TRAP is considered as a marker of osteoclast function [46]. These factors (cathepsin K, TRAP, MMPs) mediate the activity of osteoclast and their increased expression is associated with several bone diseases serve as effective biomarkers of bone destruction. In psoriatic arthritis, expression of CD16 (FcR $\gamma$ III) was considered as a potential marker of osteoclast precursors [47]. In addition, CD16<sup>high</sup> monocytes showed a higher activity of bone erosion than CD16<sup>int</sup> and CD16<sup>low</sup> monocytes, supporting that the level of CD16 expression on monocytes/osteoclast correlated with the bone resorption activity of osteoclast in psoriatic arthritis [47]. Interestingly, we recently showed that IL-23 induces NFATc1 in the absence of exogenous RANKL and stimulates the differentiation of MDL-1<sup>+</sup>/DAP12<sup>+</sup> CD16<sup>+</sup> cells corroborating the previous findings discussed herein [48, 49].

## 2.3. Molecular Pathophysiology

**2.3.1. IL-23/IL-23R**—In 2000, Oppmann and colleagues discovered that IL-23 (p19/p40) is a novel heterodimeric cytokine of the IL-12 family and is predominantly produced by dendritic cells and macrophages [50]. IL-23 and IL-12 have a mutual p40 subunit whereas p19 is a unique for IL-23 and p35 for IL-12. IL-23 binds specifically to and signals through

its heterodimeric receptor assembly which is composed of IL-12R $\beta$ 1 and IL-23R subunits [51].

IL-23 signals via its cognate receptor IL-23R and the shared receptor IL-12R $\beta$ 1 and is thought to mediate a stoichiometric ternary complex with the ectodomains of the two receptors. While IL-12R $\beta$ 1 orthologues consistently display five extracellular domains, mammalian IL-23R consist of just three extracellular domains compared to evolutionarily more distant vertebrates, which have two to three additional membrane-proximal fibronectin type III (FnIII) domains. Current models for the assembly of cell-surface complexes mediated by IL-12 family cytokines have been largely based on the structural principles derived from the interleukin-6 (IL-6) complex with interleukin-6 receptor subunit alpha (IL-6R $\alpha$ ) and glycoprotein 130 (gp130) and other gp130 complexes [52, 53].

The structure of IL-23 in complex with its cognate receptor IL23R has been recently determined [54] providing unprecedented insights into the assembly principles underlying cytokine-receptor complexes of IL-12 family. IL-23 deploys only its IL-23p19 subunit to contact IL-23R at its N-terminal Ig-like domain, which in turn reaches out via its terminal peptide to grab onto the IL-23 scaffold (Figure 2A). The structural and functional hotspot of this interaction partially restructures the helical IL-23p19 subunit of IL-23 and leads to restraining of its IL-12p40 subunit to enable cooperative binding of the shared receptor IL-12R $\beta$ 1 with high affinity. The IL-23p19 is likely not involved in binding IL-12R $\beta$ 1 at all, so the low-affinity interaction between IL-23 and IL-12R $\beta$ 1 is exclusively mediated by the p40 subunit (Figure 2B). Moreover, the experimental evidence supports a sequential model of assembly mechanism, in which a stable binary IL-23:IL-23R complex is the mechanistic prerequisite for the recruitment of the shared receptor IL-12R $\beta$ 1 to complete the ternary cytokine-receptor complex [54]. Furthermore, the dimeric disulfide-linked form of the IL-12p40 subunit, IL-12p80, which was shown to have both agonistic signaling properties as well as antagonistic behavior over a range of activities such as macrophage chemotaxis, inflammatory responses, and protective activity in a mycobacterial model, binds IL-12R $\beta$ 1 while lacking an  $\alpha$ -helical subunit [55].

Using IL-23R<sup>GFP<sup>+/+</sup></sup> reporter mice, other studies showed that IL-23R is expressed in Th17, natural killer T (NKT) cells,  $\gamma\delta$ T cells, group 3 innate lymphoid cells (ILC), dendritic cells, macrophages, and recently neutrophils [56, 57].

**2.3.2. IL-23 signaling**—Although the cytoplasmic domain of IL-23R does not have kinase activity, it has seven tyrosine residues [51]. Three of these tyrosines are potential candidates for phosphorylation and binding-sites for Src homology 2 domain [51]. Upon engagement with IL-23, the IL-23 receptor assembly will recruit and associate with Jak2/Tyk2 leading to activation of Jak2/Tyk2 and STAT3 [58]. Moreover, NF- $\kappa$ B and PI3K/Akt pathways are activated by IL-23 in T cells, leading to phosphorylation of STAT3 by PI3K/Akt which have an important role in regulation of IL-17 gene expression [58]. In Th17 cells, retinoid-related orphan receptor (ROR)  $\gamma$ T, ROR $\alpha$ , and signal transducer and activator of transcription (STAT3) are crucial transcription factors which bind the IL-17 promoter and activate gene expression cascades [59]. In addition, interaction of IL-23 with IL-23R also induces activation of nuclear factor of kappa light chain enhancer of activated B cells (NF-

$\kappa$ B) and translocation of NF- $\kappa$ B into nucleus through regulation of inhibitory subunit of nuclear factor kappa B alpha (I $\kappa$ B $\alpha$ ) [60]. Activation of NF- $\kappa$ B will induce expression of several inflammatory genes such as TNF, IL-1 $\beta$ , IL-6, and CXCLs [61].

**2.3.3. IL-23 in the skin**—In murine model of IL-23-mediated psoriasis-like epidermal hyperplasia, IL-23 treatment induced a dermal infiltration of T cells, dendritic cells (CD11c<sup>+</sup>), macrophages (F4/80<sup>+</sup>), and neutrophils (Gr1<sup>+</sup>LY6G<sup>+</sup>) [62]. Moreover, IL-23 regulates and stabilizes the Th17 phenotype and secretion of IL-17, IL-21 and IL-22 by Th17 cells (Figure 1), which mediate the epidermal hyperplasia, keratinocyte differentiation in psoriasis [63]. Injection of IL-23 induced the expression of CCR6 (a CCL20 receptor, expressed by Th17 cells and activated neutrophils), in the epidermis layer and *Ccr6*<sup>-/-</sup> mice did not develop IL-23-induced psoriasis-like inflammation, indicating that IL-23 induced psoriasis-like inflammation is dependent on CCR6 [64]. Moreover, CCL20-derived antagonists block Th17 cell homing and prevents the signs of psoriasis in a mouse model [65].

As previously mentioned IL-23 expansion of  $\gamma\delta$ T cells in the dermis layer is a key component in the pathogenesis of psoriasis [10]. CD69 is an activation marker of skin  $\gamma\delta$ T cells, and controls the aryl hydrocarbon receptor (AhR)-dependent secretion of interleukin 22 (IL-22) by  $\gamma\delta$  T cells, which contributes to the development of psoriasis induced by IL-23 [66]. Indeed, CD69-deficient mice had lower expression of epidermal IL-22 and STAT3 and attenuated skin inflammation induced by IL-23, compared with wild-type mice, suggesting that CD69 is a key mediator of psoriasis pathogenesis [66].

IL-23 regulates not only T cells but also NK cell populations in murine psoriasis-like disease [63]. Depletion of macrophages significantly reduced the skin inflammation and inflammatory cytokines in IL-23 mediated psoriasiform skin inflammation [67]. Moreover, IL-23 regulates granulopoiesis and homeostasis of neutrophil in mice [68]. In mouse model, injection of IL-23 is able to stimulate keratinocytes proliferation and induce epidermal hyperplasia [69].

It has been showed that IL-23 induced accumulation and differentiation of Langerhans cells which may be an important mechanism in regulation of cutaneous immune responses [70]. LIM-domain only protein 4 (LMO4), a transcription factor, regulates differentiation and proliferation of keratinocyte during embryogenesis. IL-23 increased the expression of LMO4 through Jak2/Akt/ STAT3 pathway and knockdown of LMO4 using shRNA had negative effects on differentiation and proliferation of keratinocytes in the ears of IL-23-injected mice [71]. Biomarkers of keratinocytes proliferation and differentiation such as Ki67 and K16, respectively as well as skin inflammatory markers (S100A7-S100A9, TNF- $\alpha$ , IL-19) was induced by IL-23 in mouse skin [72].

**2.3.4. IL-23 in the joint**—In joint autoimmune inflammation, IL-23 induced and prolonged expression of inflammatory cytokines including IL-1 $\beta$ , IL-6, IL-17, and TNF and IL-23 deficient mice (*p19*<sup>-/-</sup>) did not exhibit any clinical symptoms of disease and were resistant to the development of joint inflammation [24]. Our group has shown that overexpression of IL-23 using *in vivo* gene delivery induced chronic arthritis, severe bone loss, and myelopoiesis in the bone marrow and spleen, which resulted in increased osteoclast

differentiation and systemic bone loss [73]. IL-23 activates the synthesis and production of leukotriene B4 (LTB4) that exacerbates synovial inflammation *in vivo* and bone resorption *in vitro*, suggesting IL-23 might induce synovial inflammation through leukotriene B4 [39, 41]. In human PBMCs, IL-23 induces osteoclast differentiation through activation of DNAX activating protein of 12 kDa and its ITAMs [48] and up-regulates activation of osteoclast-associated genes (TRAP, CalCR, MMP9) through osteoclast transcription factor NFATc1 [48] (Figure 1). Although our group could not replicate the results others have shown that IL-23 increased expression of receptor activator of NF-kappa B (RANK) in primary murine bone marrow macrophages and promoted RANKL mediated osteoclast differentiation [74].

### 3. IL-23/IL-23R blockade in PsA

Currently, several antibodies against IL-23 are being developed for the treatment of psoriasis and PsA such as ustekinumab, briakinumab, tildrakizumab, guselkumab, and risankizumab (Table 1). Although clinical trials of IL-23p40 antibodies has been done and approved earlier than IL-23p19, IL-23p40 antibodies are not only inhibiting IL-23 signaling but also the IL-12 pathway, which is not required to achieve efficacy in these patients. In fact, inhibition of IL-12 is not necessary in the treatment of psoriasis and may even have negative effects and potential risks in tumor immune surveillance and in host defense against infectious diseases [75]. On the other hand, inhibition of IL-23p19 does not increase the risks of cancer development as well as bacterial/parasite infection [76]. Therefore, targeting IL-23p19 alone may be a promising treatment approach in PsA patients, by achieving a selective down-regulation of Th17 and Th22 cell responses. In this part, we will discuss about human monoclonal antibodies, which have been approved and/or are under clinical trial for PsA treatment. In contrast to antibody-based therapeutics, non-IgG based scaffolds [77] are an attractive alternative to IgG-based antibodies and at least five non-IgG based scaffolds have been developed to target IL-23: Adnectins [78], Albumin-binding domains [79, 80], Alphabodies [81], Atrimers [82] and Nanobodies [83](Figure 3).

#### Ustekinumab

(Stelara; Janssen Biotech, Inc.) is a human IgG1 $\kappa$  monoclonal antibody that binds to the common p40 subunit of IL-12 and IL-23. Ustekinumab was approved in 2009 by the US Food and Drug Administration (FDA) and the European Medicines Agency (EMA) for the treatment of moderate-to-severe psoriasis in adult patients. In fact, the initial studies with ustekinumab were designed for the treatment of psoriasis and later studies evaluated efficacy and safety of ustekinumab for PsA treatment. Ustekinumab is subcutaneously injected into PsA patients (45 or 90 mg to 100 kg or >100 kg, respectively) at week 0 and week 4, and thereafter every 12 weeks [84]. Phase II clinical trial studies showed that ustekinumab significantly reduced signs and symptoms of PsA and diminished skin lesions compared with placebo [7]. In a phase III PSUMMIT 2 trial, ustekinumab significantly improved active PsA in a diverse population of patients with active PsA, including anti-TNF-experienced PsA patients [84, 85]. In 2013, FDA approved ustekinumab in the treatment of active PsA. Thus, anti-p40 antibody ustekinumab is considered effective for the treatment of psoriasis and PsA.

### **Briakinumab**

(ABT-874; Abbott Laboratories) is another human IgG1 $\lambda$  monoclonal antibody that targets the shared p40 subunit of IL-12 and IL-23. Briakinumab is designed for the treatment of autoimmune diseases including psoriasis [75]. However, no study has been done to evaluate efficacy and safety of briakinumab for PsA treatment. Thus, we will summary the results on efficacy and safety of briakinumab for psoriasis because psoriasis and PsA have several mutual clinical and pathological features. The clinical trial study compared efficacy and safety of briakinumab with methotrexate, a traditional systemic agent, in patients with psoriasis [75]. The results showed that briakinumab has a higher efficacy than methotrexate in reducing the signs and symptoms of moderate-to-severe psoriasis. However, patients in the briakinumab group have developed serious infections and cancers [75]. Thus, this could be one of reasons leading to withdraw from application with the FDA due to safety concerns [86].

### **Tildrakizumab**

(MK-3222, Sun Pharma and Merck & Co. Inc.) is a high-affinity, humanised, IgG1 $\kappa$  monoclonal antibody targeting IL-23p19 and designed to selectively block the IL-23 signaling pathway and it has no affinity for IL-12 [87, 88] (Figure 3). Among IL-23p19 inhibitors, tildrakizumab is the first one to show positive results in phase-3 clinical trials for the treatment of moderate-to-severe plaque psoriasis. Phase 3 trials showed that tildrakizumab (100mg and 200mg) has a higher efficiency in treatment of moderate-to-severe plaque psoriasis, compared with placebo and etanercept (a TNF inhibitor) [87]. Tildrakizumab was generally safe and well tolerated, suggesting that IL-23p19 is a crucial target for inhibition of psoriasis [88]. In 2017, tildrakizumab was approved to treat patients with moderate-to-severe plaque psoriasis.

### **Guselkumab**

(CNTO 1959, Janssen) is a fully human IgG1 $\lambda$  monoclonal antibody that targets the unique IL-23p19 subunit, thus inhibiting IL-23-specific intracellular and downstream signaling [89]. In 2017, Guselkumab was approved by FDA for treatment of moderate to severe plaque psoriasis in adults [90]. Guselkumab (100 mg) is administrated through a subcutaneous injection into the patients every eight weeks (except for the second dose, which is given four weeks after the first dose) [91]. However, efficacy of guselkumab for the treatment of PsA is currently being determined in clinical trials.

### **Risankizumab**

(BI-655066 Boehringer Ingelheim, currently ABBV-066 AbbVie) is a fully human IgG1 mAb specific for the IL-23 p19 subunit [76]. Single doses of risankizumab reduced hyperkeratosis with parakeratosis, epidermal acanthosis, and inflammation in the dermis and epidermis [76]. Moreover, treatment with risankizumab decreased expression of biomarkers associated with skin inflammation and IL-23/IL-17 signaling pathways such as K16, Ki67,  $\beta$ -defensin 2, and S100A7 as well as reduction of dermal infiltration by neutrophils, T cells, and dendritic cells [76]. Phase II trials showed that specific blockade of IL-23 with risankizumab had superior clinical responses in patients with moderate-to-severe plaque



psoriasis, compared with ustekinumab (an IL-12 and IL-23p40 antibody) [92]. Moreover, there were no serious side effects in the 180-mg risankizumab group whereas two basal-cell carcinomas and one major cardiovascular adverse event were observed in ustekinumab group [92]. The results of phase III trials on risankizumab have not yet been reported.

#### 4. Insights and Conclusions

Psoriatic arthritis is a type of chronic inflammatory arthritis that occurs in patients with psoriasis. Neutrophils and T cells play necessary roles in the development as well as the resolution of inflammation and hence can evoke and/or subside the pathological features that are observed in PsA patients. IL-23 activates neutrophils and T cells which in turn interplay with other immune cells to evoke PsA pathologies in skin and bone. IL-23 is also involved in regulation of IL-17 producing cells, which have effects directly/indirectly on keratinocyte proliferation, skin inflammation, and bone loss during development of PsA. We are very hopeful that future studies will shed more light on the pathogenic mechanisms relevant to human inflammatory PsA disease and further understanding of T cells and neutrophils participation in PsA will open new avenues of treatment for PsA patients.

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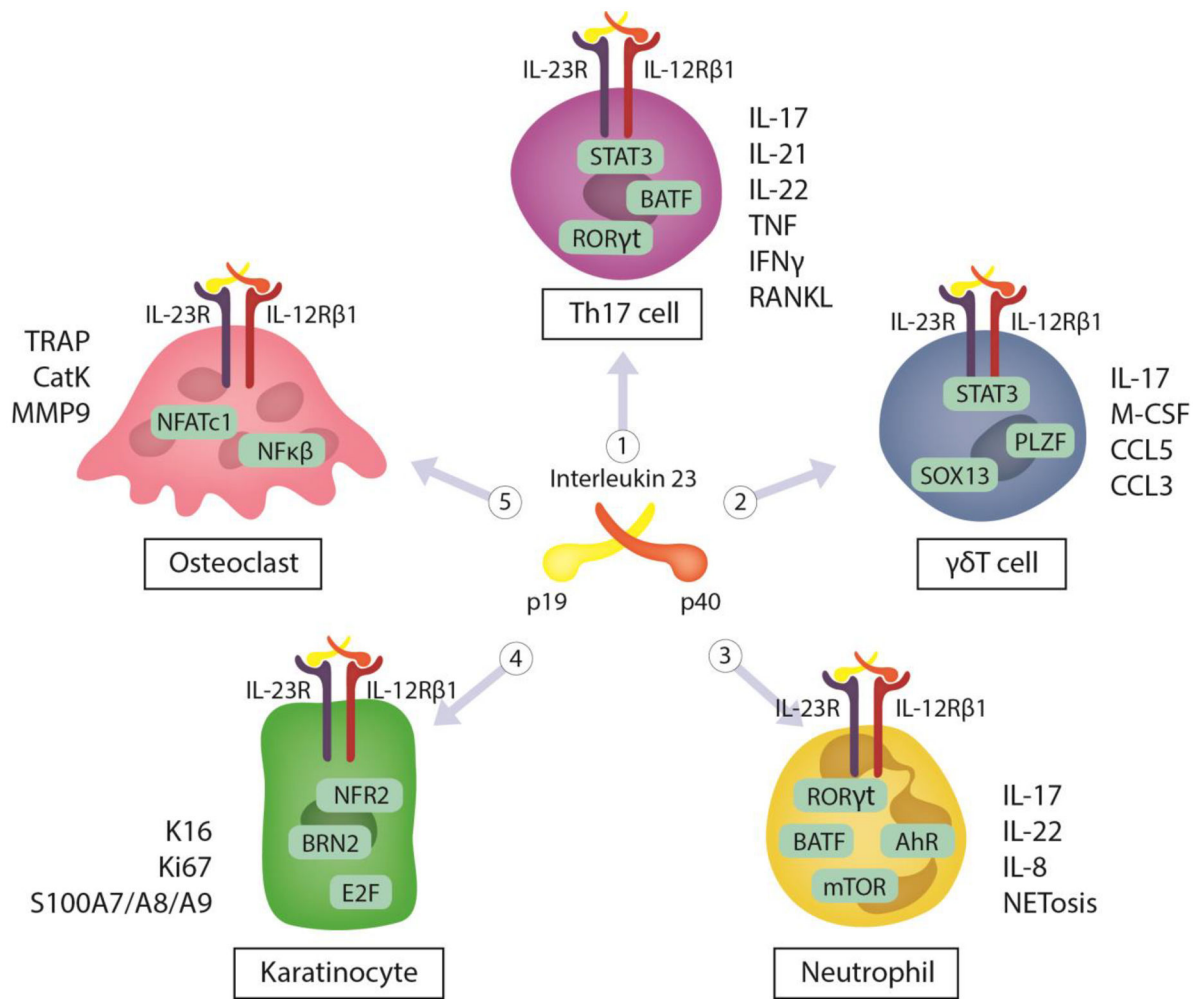
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**Highlights:**

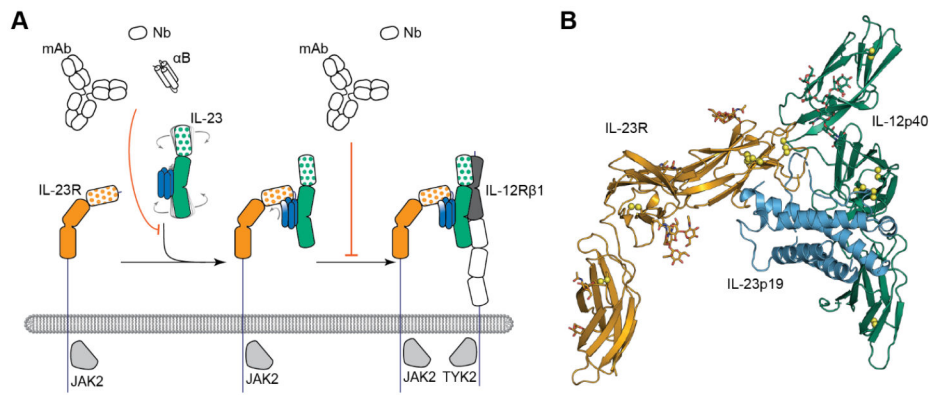
- IL-23 is critical in the pathogenesis of PsA.
- IL-23R signaling influences adaptive and innate immunity.
- Understanding the pathogenesis of PsA will lead to novel therapeutic strategies





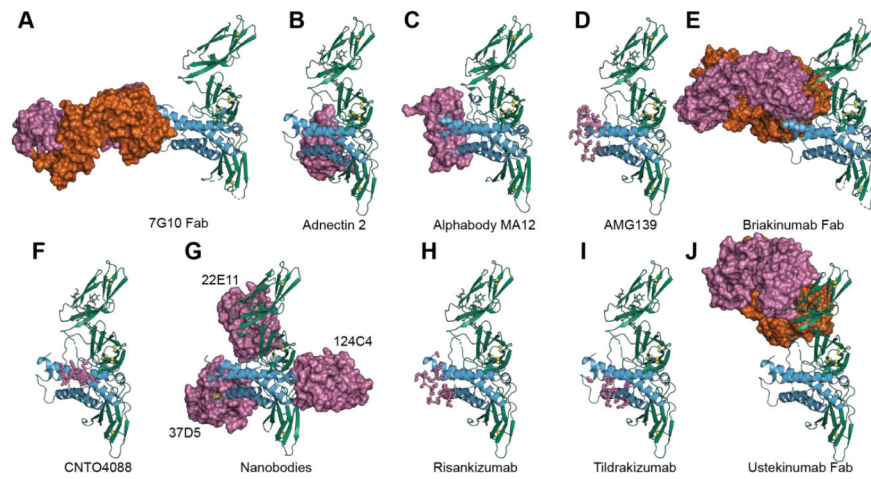
**Figure 1. Cellular pathophysiology of PsA.**

In PsA, Th17,  $\gamma\delta$ T cells, neutrophils, keratinocytes, and osteoclast are involved in the initiation of the disease. 1) Th17 cells activated by STAT3 lead to the release of IL-17, IL-21, IL-22, TNF, IFN $\gamma$  and RANKL. 2)  $\gamma\delta$ T cells are also activated by STAT3 to release of IL-17, M-CSF, CCL5, and CCL3. 3) In neutrophils, IL-23 induces the expression of IL-23R, and a number of transcriptional activators including STAT3-dependent ROR $\gamma$ t, NF- $\kappa$ B, aryl-hydrocarbon receptor (AhR), mammalian target of rapamycin (mTOR) and Basic Leucine Zipper ATF-Like Transcription Factor (BATF) pathways that regulate IL-17A, IL-17F and IL-22. NETs, IL-1 $\beta$  and IL-8 also contribute to neutrophil activity in psoriasis. 4) In keratinocytes, the transcription factor Nrf2 promotes proliferation of keratinocyte through controlling expression of K6, K16, and K17. IL-23 induces directly and/or indirectly (IL-17/IL-22) the expression of K16/Ki67 or S100A7–9 leads to induce proliferation of keratinocytes, hyperplasia, and skin inflammation. 5) In osteoclasts, IL-23 induces the expansion of osteoclast precursors and activates NFATc1 leading to the expression of TRAP, CatK, MMP9 that facilitate bone resorption.



**Figure 2. Schematic representation of the assembly mechanism of the IL-23-mediated receptor complex.**

(A) The signaling complex mediated by IL-23 proceeds via the sequential recruitment of the two cognate receptors and involves conformational selection and restructuring of IL-23 (blue and green) by IL-23R (orange) to recruit IL-12R $\beta$ 1 (black and white) with high affinity (Bloch et. al., 2018, modified). (B) Cartoon representation of the crystal structure of the IL-23:IL-23R complex (PDB 5mzv).



**Figure 3. Various antagonists developed to neutralize IL-23.**

A) Crystal structure of the IL-23 in complex with neutralizing 7G10 Fab (PDB 3D85), B) Structure of Adnectin 2, an Adnectin recognizing IL-23 (PDB 3qwr), C) Structure of MA12, an Alphabody recognizing IL-23p19 (PDB 5mj4), D) binding epitope of human antibody AMG139 mapped onto the IL-23 structure, E) Structure of briakinumab Fab, an IgG1 antibody recognizing IL-12p40 (PDB 5n2k), F) binding epitope of a recombinant mouse anti-IL23 antibody CNTO 4088 mapped onto the IL-23 structure, G) human IL-23 in complex with three nanobodies (PDB: 4grw) H) binding epitope of a humanized murine mAb Risankizumab targeting IL-23p19, mapped onto the IL-23 structure I) binding epitope of a humanized anti-IL23p19 mAb Tildrakizumab, mapped onto the IL-23 structure J) Structure of Ustekinumab-Fab, human monoclonal antibody recognizing IL-12p40 (PDB: 3hmx). In green: IL-12p40, in blue: IL-23p19, IL-23 antagonists are depicted in pink and orange surface representations.

**Table 1.**

IL-23/IL-23R targeted therapeutics for psoriasis and PsA

Drug name (Reference)	Sponsor	Target	Type	Dose (mg)	Side effects	Clinical Development Status
Ustekinumab [84]	Janssen Biotech	IL-23p40	Fully human IgG1κ monoclonal antibody	45 90	High risks infection and cancer	FDA approval (2009)
Briakinumab [75]	Abbott Laboratories	IL-23p40	Fully human IgG1 monoclonal antibody	100 200	Serious infection and cancer	FDA withdraw (2017)
Tildrakizumab [87]	Sun Pharma and Merck & Co. Inc	IL-23p19	Humanised, IgG1κ mono antibody	100 200	Non reported	FDA approval (2017)
Guselkumab [91]	Janssen	IL-23p19	Fully human IgG1λ monoclonal antibody	100	Non reported	FDA approval (2017)
Risankizumab [92]	Abb Vie Inc	IL-23p19	Fully human IgG1 monoclonal antibody	90 180	Non reported	Phase III Clinical trial

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