

Cosentyx Rheum - Mechanism of action - HCP

Prescribing information

Image



Image



Cosentyx® (secukinumab): Mechanism of action

Cosentyx is indicated for the treatment of moderate to severe plaque psoriasis (PsO) in adults, children and adolescents from the age of 6 years who are candidates for systemic therapy; active psoriatic arthritis (PsA) in adult patients (alone or in combination with methotrexate [MTX]) when the response to previous disease-modifying anti-rheumatic drug therapy has been inadequate; active ankylosing spondylitis (AS) in adults who have responded inadequately to conventional therapy; active nonradiographic axial spondyloarthritis (nr-axSpA) with objective signs of inflammation as indicated by elevated C-reactive protein and/or magnetic resonance imaging evidence in adults who have responded inadequately to non-steroidal anti-inflammatory drugs; active enthesitis-related

arthritis (ERA) in patients 6 years and older (alone or in combination with MTX) whose disease has responded inadequately to, or who cannot tolerate, conventional therapy; active juvenile psoriatic arthritis (JPsA) in patients 6 years and older (alone or in combination with MTX) whose disease has responded inadequately to, or who cannot tolerate, conventional therapy.¹

Cosentyx Summary of Product Characteristics (SmPC) can be found here

Cosentyx blocks IL-17A, a key cytokine associated with inflammation in PsA, axSpA and JIA^{1-11}

Image



Increased levels of IL-17A are found in the tissues of:

- PsA patients with psoriatic plaques¹
- blood of patients with PsA and AS^{7,14}

Image

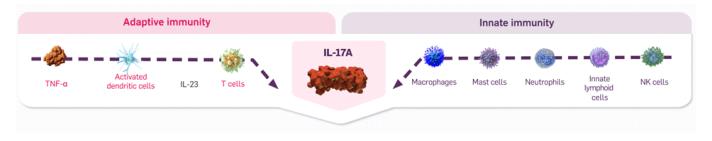


IL-17A is a pro-inflammatory cytokine. Its overproduction may contribute to signs and symptoms seen in patients with PsA, AS, and nr-axSpA^{1,11,15}

Cosentyx blocks 1L-17A, irrespective of its source^{1,12,13}

Watch the video below to discover how Cosentyx works.

Image



How does Cosentyx work?



This video has been produced and funded by Novartis Pharmaceuticals Ltd. Intended for UK healthcare professionals only.

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Image

Efficacy in PsA

Efficacy in axSpA

Efficacy in JIA

Dosing

Safety profile

HCP resources

Cosentyx is intended for use under the guidance and supervision of a physician experienced in the diagnosis and treatment of conditions for which Cosentyx is indicated. Please refer to the Cosentyx SmPC for full product information before prescribing. ¹
Therapeutic Indications ¹ Cosentyx is indicated for the treatment of moderate to severe plaque psoriasis (PsO) in adults, children and adolescents from the age of 6 years who are candidates for systemic

therapy; active psoriatic arthritis (PsA) in adult patients (alone or in combination with

therapy has been inadequate; active ankylosing spondylitis (AS) in adults who have

C-reactive protein and/or magnetic resonance imaging evidence in adults who have

responded inadequately to conventional therapy; active nonradiographic axial

methotrexate [MTX]) when the response to previous disease-modifying anti-rheumatic drug

spondyloarthritis (nr-axSpA) with objective signs of inflammation as indicated by elevated

responded inadequately to non-steroidal anti-inflammatory drugs; active moderate to severe hidradenitis suppurativa (HS; acne inversa) in adults with an inadequate response to conventional systemic HS therapy; active enthesitis-related arthritis (ERA) in patients 6 years and older (alone or in combination with MTX) whose disease has responded inadequately to, or who cannot tolerate, conventional therapy; active juvenile psoriatic arthritis (JPsA) in patients 6 years and older (alone or in combination with MTX) whose disease has responded inadequately to, or who cannot tolerate, conventional therapy.¹

AS, ankylosing spondylitis; axSpA, axial spondyloarthritis; CRP, C-reactive protein; DMARD, disease modifying anti-rheumatic drug; ERA; enthesitis-related arthritis; HS, hidradenitis suppurativa; IL-17A, interleukin 17A; IL-23, interleukin 23; JIA, juvenile idiopathic arthritis; MRI, magnetic resonance imaging; MTX, methotrexate; NK, natural killer; nr-axSpA, non-radiographic axial spondyloarthritis; NSAID, non-steroidal anti-inflammatory drug; PsA, psoriatic arthritis; PsO, plaque psoriasis; SmPC, Summary of Product Characteristics; TNF α , tumour necrosis factor alpha.

References

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Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to Novartis online through the pharmacovigilance intake (PVI) tool at www.novartis.com/report, or alternatively email medinfo.uk@novartis.com or call 01276 698370.

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