

Sun Pharma launches plaque psoriasis drug in Japan

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ILUMYA is Sun Pharma's first innovative drug to be launched in the Japanese market



Indian firm Sun Pharmaceutical Industries has announced that its wholly-owned Japanese subsidiary has launched ILUMYA[®] Subcutaneous Injection 100 mg Syringe (Nonproprietary name: tildrakizumab? (genetical recombination), "ILUMYA") in Japan for the treatment of plaque psoriasis in adult patients who have an inadequate response to conventional therapies.

ILUMYA is a humanized IgG1/k monoclonal antibody designed to selectively bind to the p19 subunit of IL-23 and inhibit its interaction with the IL-23 receptor, leading to inhibition of the release of pro-inflammatory cytokines and chemokines.

Junichi Nakamichi, Country Head, Sun Pharma Japan, said, "ILUMYA is Sun Pharma's first innovative drug to be launched in the Japanese market. We are pleased to introduce a new, safe and effective treatment option for plaque psoriasis to doctors and patients in our country. This is an important milestone for Sun Pharma as we expand our product portfolio in Japan."

ILUMYA is contraindicated in patients with a previous serious hypersensitivity reaction to ILUMYA or to any other excipients. Cases of angioedema and urticaria occurred in ILUMYA -treated subjects in clinical trial.

Treatment with ILUMYA should not be initiated in patients with a clinically important active infection until the infection resolves or is adequately treated. Consider the risks and benefits of treatment prior to prescribing ILUMYA in patients with a chronic infection or a history of recurrent infection.

The most common (?1%) adverse reactions associated with ILUMYA include upper respiratory infections, injection site reactions, and diarrhea. Adverse reactions that occurred at rates less than 1% but greater than 0.1% in the ILUMYA group and at a higher rate than in the placebo group included dizziness and pain in extremity.