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## Transcript

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ON SCREEN:

Please watch the Important Safety Information following this video.

ON SCREEN:

Jordan, an actual patient with psoriatic arthritis (PsA) on COSENTYX® (secukinumab), was compensated for her time. Individual results may vary.

SUPER:

JORDAN, ACTUAL PATIENT

JORDAN:

I first started feeling joint pain mostly in my knees when I was younger.

SUPER:

COSENTYX® (secukinumab) logo

COSENTYX is indicated for adults with active PsA.

Individual results may vary.

JORDAN:

It was pretty much always just in my knees, kind of ankles, legs area.

But once I got older, I'd started getting it in different parts of my body, like kind of back and elbows. So I started feeling it more all over my full body.

SUPER:

COSENTYX® (secukinumab) logo

JORDAN:

Some days are better than others. Some days you wake up and you don't even want to get out of bed, you don't want to stand in the shower.

SUPER:

COSENTYX® (secukinumab) logo

Up to 60% of patients in a clinical trial saw at least a 20% improvement in PsA symptoms at 16 weeks (versus 18% taking placebo). Individual results may vary.

JORDAN:

COSENTYX is the first thing that really worked for me, and I hadn't heard of it before my doctor brought it to my attention.

SUPER:

COSENTYX® (secukinumab) logo

Individual results may vary.

JORDAN:

The further I got into it, I noticed the joint pain was also dissipating. Now that I feel better, I realize that it's a lot easier to pick up a case of product and put it in my car, grab my computer bag, and do all the things that I need to do.

SUPER:

COSENTYX® (secukinumab) logo

Please see full Prescribing Information, including Medication Guide, on this site or at COSENTYXHCP.com. 9/23 298791

## INDICATIONS

COSENTYX® (secukinumab) is indicated for the treatment of moderate to severe plaque psoriasis in patients 6 years and older who are candidates for systemic therapy or phototherapy.

COSENTYX is indicated for the treatment of active psoriatic arthritis (PsA) in patients 2 years of age and older.

COSENTYX is indicated for the treatment of adult patients with active ankylosing spondylitis (AS).

COSENTYX is indicated for the treatment of adult patients with active non-radiographic axial spondyloarthritis (nr-axSpA) with objective signs of inflammation.

COSENTYX is indicated for the treatment of active enthesitis-related arthritis (ERA) in patients 4 years of age and older.

## **IMPORTANT SAFETY INFORMATION**

### **CONTRAINDICATIONS**

COSENTYX is contraindicated in patients with a previous serious hypersensitivity reaction to secukinumab or to any of the excipients in COSENTYX. Cases of anaphylaxis have been reported during treatment with COSENTYX.

### **WARNINGS AND PRECAUTIONS**

#### **Infections**

COSENTYX may increase the risk of infections. In clinical trials, a higher rate of infections was observed in COSENTYX treated subjects compared to placebo-treated subjects. In placebo-controlled clinical trials in subjects with moderate to severe plaque psoriasis, higher rates of common infections, such as nasopharyngitis (11.4% versus 8.6%), upper respiratory tract infection (2.5% versus 0.7%) and mucocutaneous infections with candida (1.2% versus 0.3%) were observed with COSENTYX compared with placebo. A similar increase in risk of infection was seen in placebo-controlled trials in subjects with psoriatic arthritis, ankylosing spondylitis and non-radiographic axial spondyloarthritis. The incidence of some types of infections appeared to be dose-dependent in clinical studies. In the postmarketing setting, serious and some fatal infections have been reported in patients receiving COSENTYX.

Exercise caution when considering the use of COSENTYX in patients with a chronic infection or a history of recurrent infection.

Instruct patients to seek medical advice if signs or symptoms suggestive of an infection occur. If a patient develops a serious infection, monitor the patient closely and discontinue COSENTYX until the infection resolves.

#### **Pre-treatment Evaluation for Tuberculosis**

Evaluate patients for tuberculosis (TB) infection prior to initiating treatment with COSENTYX. Avoid administration of COSENTYX to patients with active TB infection. Initiate treatment of latent TB prior to administering COSENTYX. Consider anti-TB therapy prior to initiation of COSENTYX in patients with a past history of latent or active TB in whom an adequate course of treatment cannot

be confirmed. Monitor patients closely for signs and symptoms of active TB during and after treatment.

### **Inflammatory Bowel Disease**

Caution should be used when prescribing COSENTYX to patients with inflammatory bowel disease. Exacerbations, in some cases serious, occurred in COSENTYX treated subjects during clinical trials in plaque psoriasis, psoriatic arthritis, ankylosing spondylitis and non-radiographic axial spondyloarthritis. In addition, new onset inflammatory bowel disease cases occurred in clinical trials with COSENTYX. In an exploratory trial in 59 subjects with active Crohn's disease, there were trends toward greater disease activity and increased adverse events in the secukinumab group as compared to the placebo group. Patients who are treated with COSENTYX should be monitored for signs and symptoms of inflammatory bowel disease.

### **Eczematous Eruptions**

In postmarketing reports, cases of severe eczematous eruptions, including atopic dermatitis-like eruptions, dyshidrotic eczema, and erythroderma, were reported in patients receiving COSENTYX; some cases resulted in hospitalization. The onset of eczematous eruptions was variable, ranging from days to months after the first dose of COSENTYX.

Treatment may need to be discontinued to resolve the eczematous eruption. Some patients were successfully treated for eczematous eruptions while continuing COSENTYX.

### **Hypersensitivity Reactions**

Anaphylaxis and cases of urticaria occurred in COSENTYX treated subjects in clinical trials. If an anaphylactic or other serious allergic reaction occurs, administration of COSENTYX should be discontinued immediately and appropriate therapy initiated.

The removable caps of the COSENTYX Sensoready® pen and the COSENTYX 1 mL and 0.5 mL prefilled syringes contain natural rubber latex, which may cause an allergic reaction in latex-sensitive individuals. The safe use of the COSENTYX Sensoready pen or prefilled syringe in latex-sensitive individuals has not been studied.

**Immunizations**

Prior to initiating therapy with COSENTYX, consider completion of all age appropriate immunizations according to current immunization guidelines. COSENTYX may alter a patient's immune response to live vaccines. Avoid use of live vaccines in patients treated with COSENTYX.

**MOST COMMON ADVERSE REACTIONS**

Most common adverse reactions (>1%) are nasopharyngitis, diarrhea, and upper respiratory tract infection.

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