BOTOX® Cosmetic (onabotulinumtoxinA) IMPORTANT SAFETY INFORMATION (continued)

Serious and/or immediate allergic reactions have been reported. They include: itching, rash, red itchy welts, wheezing, asthma symptoms, or dizziness or feeling faint. Get medical help right away if you are wheezing or have asthma symptoms, or if you become dizzy or faint.

Do not receive BOTOX® Cosmetic if you: are allergic to any of the ingredients in BOTOX® Cosmetic (see Medication Guide for ingredients); had an allergic reaction to any other botulinum toxin product such as *Myobloc®* (rimabotulinumtoxinB), *Dysport®* (abobotulinumtoxinA), or *Xeomin®* (incobotulinumtoxinA); have a skin infection at the planned injection site.

Tell your doctor about all your muscle or nerve conditions, such as ALS or Lou Gehrig's disease, myasthenia gravis, or Lambert-Eaton syndrome, as you may be at increased risk of serious side effects including difficulty swallowing and difficulty breathing from typical doses of BOTOX® Cosmetic.

Tell your doctor about all your medical conditions, including: plans to have surgery; had surgery on your face; have trouble raising your eyebrows; drooping eyelids; any other abnormal facial change; are pregnant or plan to become pregnant (it is not known if BOTOX® Cosmetic can harm your unborn baby); are breast-feeding or plan to (it is not known if BOTOX® Cosmetic passes into breast milk).

Please see additional Important Safety Information on following page.

Moderate to severe crow's feet lines

Before

After (Day 30)

Actual patient. **Results may vary.**

Photos taken at maximum smile before treatment with BOTOX® Cosmetic (onabotulinumtoxinA) and taken at maximum smile after treatment with BOTOX® Cosmetic at day 30.

In clinical trials at day 30, 26.1% (58/222) and 20.3% (62/306) of patients demonstrated a ≥ 2-grade improvement from baseline in crow's feet line severity at maximum smile as compared to 1.3% (3/223) and 0% (0/306) in placebo, as assessed by both investigators and subjects.[1]

Allergan

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