

ReDefyne Your Expressions

The newest dermal fillers in the *Restylane*[®] family of products include *Restylane*[®] Refyne and *Restylane*[®] Defyne. These are two next-generation hyaluronic acid (HA) dermal fillers created with **XpresHAN Technology**[™], to help smooth moderate lines and provide the right amount of support and flexibility through a range of facial expressions.^{1,2}

Quick Facts:

- Maintain **expression in motion**³
- **Natural-looking results** and long-lasting effects
- Over **30 million** *Restylane*[®] treatments (including both NASHA and XpresHAN technologies) worldwide*⁴

Restylane[®] Refyne and *Restylane*[®] Defyne are poised to **ReDefyne** how aesthetic professionals think about individual assessment, experience and satisfaction. These *Restylane*[®] products are approved for patients over the age of 21.

Restylane
REFYNE

Restylane[®] Refyne is FDA-approved, for patients over the age of 21, for the treatment of moderate to severe facial wrinkles and folds, such as nasolabial folds.

Restylane
DEFYNE

Restylane[®] Defyne is FDA-approved, for patients over the age of 21, for the treatment of moderate to severe, deep facial wrinkles and folds, such as nasolabial folds.

Here is what you can expect from these dermal fillers:

- **Integrates with your skin** and acts like your body's own HA⁵
- **Gives the support you need** while allowing a range of facial expressions^{1,2}
- **Offers the flexibility you want** with consistent texture for natural expression in motion^{2,3}
- **Provides minimal downtime** with a comfortableness in returning to social engagement⁶



*Actual patient. Individual results may vary. Images have not been retouched.
Before and after 1.0 mL of Restylane[®] Defyne in nasolabial folds, 1.0 mL Restylane[®] Defyne to marionette lines. Post-injection timing: 2 weeks.*

For more information about *Restylane*[®] Refyne and *Restylane*[®] Defyne, visit www.RestylaneUSA.com or speak with a licensed healthcare professional to see if these products are right for you.

* *Restylane*[®] Refyne and *Restylane*[®] Defyne were first approved in Europe in 2010 under the brand name *Emerve*[®] and have proven safety profile demonstrated by more than one million treatments worldwide.

Important Safety Information

The *Restylane* family of products includes *Restylane*[®], *Restylane-L*[®], *Restylane*[®] *Lyft with Lidocaine*, *Restylane*[®] *Silk*, *Restylane*[®] *Refyne*, and *Restylane*[®] *Defyne*.

APPROVED USES

Restylane[®] and *Restylane-L*[®] are for mid-to-deep injection into the facial tissue for the correction of moderate to severe facial wrinkles and folds, such as nasolabial folds. *Restylane*[®] and *Restylane-L*[®] are also indicated for injection into the lips in patients over the age of 21.

Restylane[®] *Lyft with Lidocaine* is for deep implantation into the facial tissue for the correction of moderate to severe facial wrinkles and folds, such as nasolabial folds and for cheek augmentation and for the correction of age-related midface contour deficiencies in patients over the age of 21.

Restylane[®] *Silk* is for lip augmentation and for correction of perioral wrinkles in patients over the age of 21.

Restylane[®] *Refyne* is for mid-to-deep injection into the facial tissue for the correction of moderate to severe facial wrinkles and folds, such as nasolabial folds, in patients over the age of 21.

Restylane[®] *Defyne* is for mid-to-deep injection into the facial tissue for the correction of moderate to severe deep facial wrinkles and folds, such as nasolabial folds, in patients over the age of 21.

Are there any reasons why I should not use products within the *Restylane*[®] family? (Contraindications)

To ensure a safe procedure, your doctor will talk to you about your medical history to determine if you are an appropriate candidate for treatment. You should not use products within the *Restylane* family if:

- You have severe allergies with a history of severe reactions (anaphylaxis)
- You are allergic to lidocaine or to any of the gram-positive bacterial proteins used to make hyaluronic acid
- You are prone to bleeding or have been diagnosed with a bleeding disorder

Are there other precautions that I should discuss with my doctor?

- Tell your doctor if you are breastfeeding, pregnant, or trying to become pregnant. The safety of these products for use during pregnancy, or in women who are breastfeeding, has not been studied
- *Restylane*, *Restylane-L*, *Restylane*[®] *Lyft with Lidocaine*, *Restylane Refyne* and *Restylane Defyne* are intended to treat facial wrinkles and folds, such as nasolabial folds. *Restylane* and *Restylane-L* are also intended for lip enhancement. Treatments in other areas of the face have not been evaluated in clinical studies.
- The safety and effectiveness of *Restylane*[®] *Silk* for areas other than the lips and perioral area have not been evaluated in clinical studies
- Tell your doctor if you have any history of scarring, particularly thick and stiff scars, or any pigmentation (skin color) disorders. These side effects can occur with hyaluronic acid fillers in general.
- Tell your doctor if you are planning other laser treatments or a chemical peel, as there is a possible risk of inflammation at the treatment site if these procedures are performed after treatment
- Patients who experience skin injury near the site of injection with these products may be at a higher risk for side effects
- Tell your doctor if you are on any medications to decrease your body's immune response (immunosuppressive therapy). Using these medications may increase your risk of bruising or bleeding at the gel injection site.
- Tell your doctor if you are using any "blood thinners" such as aspirin, warfarin, or any other

medications that affect bleeding. Using these medications may increase your risk of bruising or bleeding at the gel injection site.

- The use of these products on gel injection sites with skin sores, pimples, rashes, hives, cysts, or infections should be postponed until healing is complete. Use of product in these areas could delay healing or make your skin problems worse.

What are the possible side effects?

The most commonly observed side effects are swelling, redness, pain, bruising, headache, tenderness, lump formation, and itching at the injection site. These are typically mild in severity and typically resolve in less than 7 days in nasolabial folds and less than 14 days in lips. Serious but rare side effects include delayed onset infections, recurrence of herpetic eruptions, and superficial necrosis at the injection site.

One of the risks with using this product is unintentional injection into a blood vessel. The chances of this happening are very small, but if it does happen, the complications can be serious, and may be permanent. These complications, which have been reported for facial injections, can include vision abnormalities, blindness, stroke, temporary scabs, or permanent scarring of the skin.

As with all skin injection procedures, there is a risk of infection.

To report a side effect with any of the Restylane products, please call Galderma Laboratories, L.P at 1-855-425-8722.

The *Restylane* family of products is available only through a licensed practitioner. Complete Instructions for Use are available at www.RestylaneUSA.com.

¹ Segura, S., Anthonioz, L., Fuchez, F., et al. A Complete Range of Hyaluronic Acid Filler With Distinctive physical properties Specifically Designed for Optimal Tissue Adaptations. *J. Drugs Dermatol.* 2012; 11(1 suppl.):s5-s8. **Restylane*[®] Refyne is known as *Emerve*[®] Classic outside the U.S. *Restylane*[®] Defyne is known as *Emerve*[®] Deep, respectively, outside the U.S. XpresHAn Technology[™] is also known as Optimal Balance Technology (OBT)[™] outside the U.S.

² Data on file. MA-32418 Clinical Study Report. Fort Worth, TX: Galderma Laboratories, L.P., 2016.

³ Data on file. MA-30083 Clinical Study Report. Fort Worth, TX: Galderma Laboratories, L.P., 2016.

⁴ Data on file. Global Launch Learnings. Fort Worth, TX: Galderma Laboratories, L.P., 2016.

⁵ Data on file. Fort Worth, TX: Galderma Laboratories, L.P., 2016.

⁶ Data on file. GLI.04.SPR.US10346 Clinical Study Report. Fort Worth, TX: Galderma Laboratories, L.P., 2016.