FAST ONSET/HIGH SATISFACTION





+ Glabellar Lines (Frown Lines)1

> 10 units (U) at each of 5 injection sites (total: 50 U¹)

2 injections per corrugator muscle¹ + 1 injection in procerus muscle¹

Lateral Canthal Lines (Crow's Feet)1

> 10 U at each of 3 injection sites per side (total: 30 U/side; 60 U for both sides¹)

3 injection sites per crow's feet side¹







Active toxin: Dysport Aesthetic delivers a high amount of 150 kDa active toxin in each injection³

LONG-LASTING RESULTS

Clinical effect of Dysport Aesthetic may last up to 4 months²

FAST ONSET

Natural-looking results as early as 24 hours¹

HIGHLY SATISFIED PATIENTS AND CLINICIANS

97% of patients want to have Dysport Aesthetic again⁴





APPROVED FOR GLABELLAR LINES AND LATERAL CANTHAL LINES (CROW'S FEET)1

Patients seek aesthetic procedures to look younger and fresher/appear more rested⁵⁻⁷

Results as soon as 24 hours with a median time to onset of 3 days.

The clinical effect of Dysport Aesthetic may last up to 4 months.² Prospective, comparative, triple-blinded, single-centre, randomized, internally controlled clinical trial in 85 patients (18-65 years) with moderate-to-severe wrinkles in the glabellar or crow's feet area. Patients were given a single injection on Day 0 with Dysport Aesthetic (20 U: glabellar lines, 30 U: crow's feet) and onabotulinumtoxinA (8 U: glabellar lines, 10 U: crow's feet) injected on opposite sides of the face.

CTIVE TOXIN/LASTING RESULTS

FAST ONSET/HIGH SATISFACTION

REAL RESULTS/PORTFOLIO

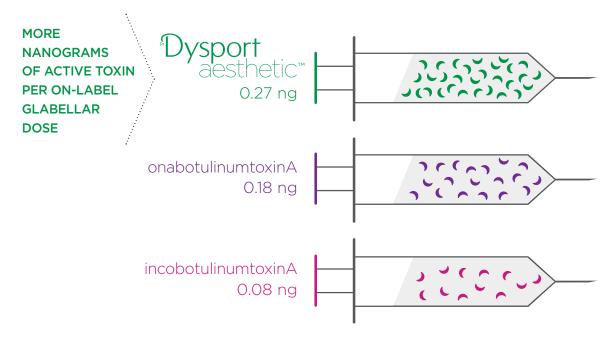
a HIGH AMOUNT OF active toxin in Each injection³



- All BoNT-As have the same mechanism of action
- Clinical activity of botulinum toxins is mediated by the
 150 kDa active toxin

HOW DO BOTULINUM TOXIN TYPE AS DIFFER?

The difference is in the amount of active toxin



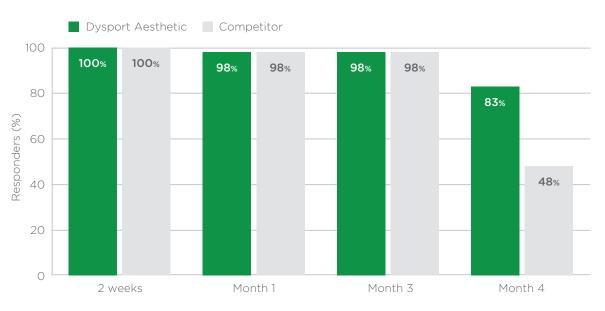




LASTING RESULTS

Efficacy MAY LAST UP TO 4 months²

INVESTIGATORS' ASSESSMENT OF TREATMENT RESPONSE
(IMPROVEMENT ≥1 GRADE) AT MAXIMUM CONTRACTION (n=59)²*



Adapted from Kassir R, et al. 2013²

83%

were still responding at 4 months²

^{*} Prospective, comparative, triple-blinded, single-centre, randomized, internally controlled clinical trial in 85 patients (18-65 years) with moderate-to-severe wrinkles in the glabellar or crow's feet area. Patients were given a single injection on Day 0 with Dysport Aesthetic (20 U: glabellar lines, 30 U: crow's feet) and onabotulinumtoxinA (8 U: glabellar lines, 10 U: crow's feet) injected on opposite sides of the face.²

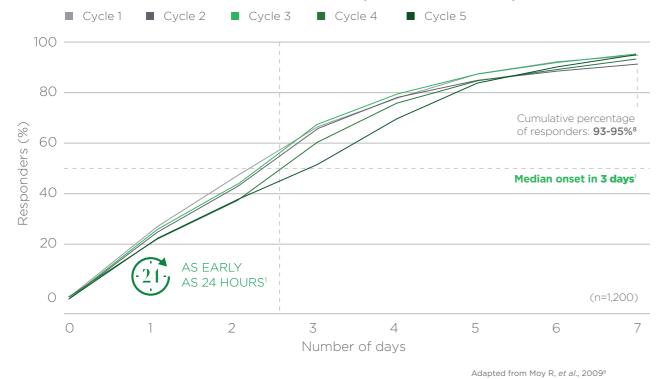
FAST ONSET/HIGH SATISFACTION

FAST ONSET

Natural-looking results as Early as 24 hours¹

MEDIAN TIME TO ONSET OF RESPONSE WAS 3 DAYS1

PATIENT ASSESSMENT OF RESPONSE (GLABELLAR LINES)⁵



At least 93% of patients saw results by day 7 post-treatment8



Multicentre, phase 3, open-label, repeat-dose study to assess the long-term safety of repeat administrations of Dysport Aesthetic 50 units in the treatment of glabellar lines.8

Satisfaction with dysport Aesthetic

LASTING PATIENT AND CLINICIAN SATISFACTION



Clinician satisfaction: **91%** mean overall practitioner satisfaction* across 5 treatment cycles⁹

Patient satisfaction: Multicentre, prospective, observational study in 533 patients (18-64 years) with moderate to severe glabellar lines at maximal frown. 48% were treatment-naïve; 51% were previously treated with a competitive botulinum toxin type A product in the same area. All patients received Dysport Aesthetic at the discretion of the investigator and subject. Satisfaction was assessed at 3 weeks and 4 months.

Clinician satisfaction: Retrospective multi-country study. Satisfaction rated on a 3-point scale (satisfactory, not satisfactory, unknown) with repeated Dysport Aesthetic use in routine clinical practice (5.9-6.5 months median treatment interval).¹⁰

^{*} Calculated as the mean percentage across the 5 treatment cycles.9

Real results ~

BEFORE AND AFTER

GALDERMA EXCLUSIVE aesthetics COLLECTION

Photos are real and unretouched.



Before Dysport Aesthetic



After (Day 10)* Area treated: glabellar lines



Before Dysport Aesthetic



After (Week 4)*
Area treated: crow's feet

* Individual results may vary.

Help give patients the relaxed, refined, refreshed and renewed look they desire^{1,11-20}

RELAX



*Dysport aesthetic**



REFINE

Restylane

REFRESH



Kestylane° SKINBOOSTERS™

Sculptra



Kestylane DEFYNE

Restylane®

Restylane REFYNI

Restylane®

Restylane

Restylane VOLYME

Restylane

SKINBOOSTERS VITAL

Restylane

Restylane SKINBOOSTERS VITALLIGH

For more information, please visit
DysportCanada.ca,
Restylane.ca
and Sculptra.ca



Safety profile

TEAES WERE GENERALLY MILD TO MODERATE AND THE MAJORITY WERE CONSIDERED **UNRELATED OR UNLIKELY TO BE RELATED TO** TREATMENT¹

	DYSPORT AESTHETIC n=2,041 (%)*	PLACEBO n=480 (%)*
Patients with at least one TEAE	852 (42)	141 (29)
EYE DISORDERS		
Eyelid ptosis	33 (2)	1 (<1)
GENERAL DISORDERS AND ADMIN	ISTRATION SITE CONDITIONS	
Injection site pain	69 (3)	7 (1)
Injection site bruising	34 (2)	7 (1)
INFECTIONS AND INFESTATIONS		
Nasopharyngitis	123 (6)	19 (4)
Upper respiratory tract infection	58 (3)	4 (<1)
Sinusitis	55 (3)	4 (<1)
Influenza	31 (2)	3 (<1)
NERVOUS SYSTEM DISORDERS	'	
Headache	110 (5)	16 (3)

^{*} Patients who received treatment with placebo and Dysport Aesthetic are counted in both treatment columns.

TEAEs (>1% INCIDENCE) FOLLOWING THE 1 ST CYCLE OF TREATMENT FROM BOTH STUDIES IN THE CROW'S FEET REGION AT 30 U ¹			
	DYSPORT AESTHETIC n=306 (%)	PLACEBO n=137 (%)	
EYE DISORDERS			
Eyelid edema	5 (1.6)	-	
GENERAL DISORDERS AND ADMINISTRATION SITE CONDITIONS			
Injection site reactions (hematoma, pruritus and swelling)	7 (2.4)	1 (0.7)	
INFECTIONS AND INFESTATIONS			
Nasopharyngitis	5 (1.7)	3 (2.2)	
NERVOUS SYSTEM DISORDERS			
Headache	13 (4.3)	3 (2.2)	



IMPORTANT safety information

CONTRAINDICATIONS:

- Patients with infection at the proposed injection sites.
- Patients with a known allergy to cow's milk protein.

MOST SERIOUS WARNINGS AND PRECAUTIONS:

Distant spread of toxin effect: The effects of all botulinum toxin products may spread from the area of injection to produce symptoms consistent with botulinum toxin effects, which can be life-threatening. The risk of symptoms is probably greatest in children treated for spasticity and in adults who have underlying conditions that would predispose them to these symptoms. Dysport Aesthetic should only be administered by physicians with the appropriate qualifications and experience in the treatment and the use of required equipment. Dysport Aesthetic units are non-interchangeable with other products.

OTHER RELEVANT WARNINGS AND PRECAUTIONS:

- Use only as directed.
- Do not use dosage recommendations and potency units applied to other botulinum toxin products.
- Caution in patients with neuromuscular junction disorders or when excessive weakness or atrophy is present in the target muscle, prolonged

bleeding times, surgical alterations to the facial anatomy, marked facial asymmetry, inflammation at the injection site(s), ptosis, excessive dermatochalasis, deep dermal scarring, or thick sebaceous skin. Muscle weakness may temporarily impair the ability to drive or operate machinery.

- Risk of muscle weakness remote to the site of injection. Extreme caution in patients with a history of underlying neurologic disorders, facial palsy, dysphagia and/or aspiration.
- Immediate medical care if swallowing, speech or respiratory disorders arise.
- Risk for transmission of Creutzfeldt-Jakob disease (CJD).
- Caution in patients with eve disorders.
- Anaphylactic reaction may occur, necessary precautions should be taken.
- Caution in pregnancy.
- · Not recommended during lactation.

FOR MORE INFORMATION:

Please consult the Product Monograph at www.galderma.com/sites/g/ files/jcdfhc196/files/inline-files/Dysport-aesthetic-PM-E.pdf for important information relating to adverse reactions, interactions and dosing information, which have not been discussed in this advertisement.

The Product Monograph is also available by calling us at 1-800-467-2081.

References

- 1. PrDysport Aesthetic™ Product Monograph. Galderma Canada Inc. October 24, 2018.
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- 7. Molina B, et al. Patient satisfaction after the treatment of glabellar lines with botulinum toxin type A (Speywood Unit): a multi-centre European observational study. JEADV 2015;29:1382-1388.

- 8. Moy R, et al. Long-term safety and efficacy of a new botulinum toxin type A in treating glabellar lines. Arch Facial Plast Surg 2009;11(2):77-83.
- 9. Rzany B, et al. Treatment of glabellar lines with botulinum toxin type A (Speywood Unit): a clinical overview. JEADV 2010;24(Suppl. 1):1-14.
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- 15. Restylane® Silk Instructions for Use.
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- 17. Restylane® Volyme Instructions for Use.
- 18. Restylane® Skinboosters™ Vital Instructions for Use.
- 19. Restylane® Skinboosters™ Vital Light Instructions for Use.
- 20. Sculptra® Instructions for Use.
- 21. Data on File. Galderma Canada Inc.



THE DYSPORT AESTHETIC



I ASTING **RESULTS**

Efficacy may last up to 4 months^2

A **HIGH** AMOUNT

of active toxin in each $injection^{3}$

DEPTH OF **experience**

- Over 28 years of use in 73 countries²¹
- 12,800+ patients in clinical trials²¹

A PROVEN, VERSATILE Solution

• Glabellar lines • Lateral canthal lines





NATURAL-LOOKING RESULTS⁴

98%
thought their results looked natural

HIGH PATIENT SATISFACTION⁴

970/0 wanted Dysport Aesthetic again

