

| Clinical rationale for BOTOX® injection: | | |
|--|--|--|
| Clinical rationale for EMG (if performed): | | |
| Official rationale for Livia (if performed). | | |
| Comments: | | |

| Dilution Table | | | |
|--|----------------------------------|--|--|
| Diluent added (0.9% Sodium Chloride Injection) | Resulting Dose (Units/0.1 mL) | | |
| 1.0mL | 10.0 U | | |
| 2.0 mL | 5.0 U | | |
| 4.0 mL | 2.5 U | | |
| 8.0 mL | 1.25 U | | |

| Treatment date | |
|-------------------------|--|
| Dilution (Units/mL) | |
| | |
| Lot number(s) | |
| Vial expiration date(s) | |

Note: These dilutions are calculated for an injection volume of 0.1 mL. A decrease or increase in the BOTOX® dose is also possible by administering a smaller or larger injection volume from 0.05 mL (50% decrease in dose) to 0.15 mL (50% increase in dose).

BOTOX® (botulinum toxin type A) is indicated for the treatment of cervical dystonia in adults to decrease the severity of abnormal head position and neck pain associated with cervical dystonia.

IMPORTANT SAFETY INFORMATION

Contraindications: BOTOX® treatment is contraindicated in the presence of infection at the proposed injection site(s) and in individuals with known hypersensitivity to any ingredient in the formulation.

Warnings: Serious and/or immediate hypersensitivity reactions have been rarely reported. These reactions include anaphylaxis, urticaria, soft-tissue edema, and dyspnea. If such a reaction occurs, further BOTOX® injection should be discontinued and appropriate medical therapy immediately instituted. Patients with peripheral motor neuropathic diseases (eg, amyotrophic lateral sclerosis or motor neuropathy) or neuromuscular junctional disorders (eg, myasthenia gravis or Lambert-Eaton syndrome) should only receive BOTOX® treatment with caution. Patients with neuromuscular disorders may be at increased risk of clinically significant systemic effects including severe dysphagia and respiratory compromise from typical BOTOX® doses.

Adverse events: There have been rare spontaneous reports of death, sometimes associated with dysphagia, pneumonia, and/or other significant debility or anaphylaxis, after treatment with botulinum toxin. There have been rare reports of adverse events involving the cardiovascular system. The most frequently reported adverse reactions in patients with cervical dystonia following BOTOX® injection are dysphagia (19%), upper respiratory infection (12%), neck pain (11%), and headache (11%).

Focused On Innovation

PATIENT INJECTION RECORD

| Semispinalis capitis Right: Left: Sternocleidomastoid 1,2 Typical BOTOX* Dosage: 50–150 U | Typical BOTOX® Dosage: 50 –150 U Typical BOTOX® Dosage: 50 –150 U | Splenius capitis ³ Right: Left: Longissimus Right: |
|---|--|---|
| Right: Left: | Typical BOTOX® Dosage: 20 –60 U | Left: Splenius cervicis 3 Right: Left: |
| Trapezius Typical BOTOX* Dosage: 50–100 U Right: Left: | Typical BOTOX® Dosage: 25–100 U | |

FILL IN NUMBER OF UNITS INJECTED

| Splenius capitis Typical BOTOX® Dosage: 50–150 U | Typical BOTOX® Dosage: 15–75 U | Sternocleidomastoid ^{1,2} Right: |
|---|---|---|
| Right: | Typical BOTOX® Dosage: 15–50 U | Scalene |
| | | Right: |
| Levator scapulae ⁴ Typical BOTOX® Dosage: 25–100 U | | Left: |
| Right: | | % of patients/total muscle groups (Units). the sternocleidomastoid muscle to ≤ 100 U |
| _eft: | may decrease the occurrence | of dysphagia. If contributing, dose is always nocleidomastoids for anterocollis. Patients |
| Trapezius Typical BOTOX® Dosage: 50–100 U | with smaller neck muscle mas | s, or patients who require bilateral injections nuscle, have been reported to be at greater |
| Right: | risk of dysphagia. 3 When injecting in back of neck in order to avoid tilting of the h | k, it is critical that injections be symmetrical |
| | 4 Usually injected if shoulder is in | |

Some patients may require injections into additional muscles. Fill in additional locations/units injected below.

| Sites Injected | No. Injections/Muscle | Units/Injection | Total Units/Muscle |
|--------------------|-----------------------|----------------------|--------------------|
| | | | |
| | | | |
| | | | |
| | | | |
| | | | |
| Total Units Billed | | Total Units Injected | |

Note to representative: Please provide full prescribing information when presenting this material.



