For U.S. Healthcare Professionals

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Dedicated Support

for Your Dysport® Patients and Their Families

Please see Indications and Important Safety Information on page 4 and accompanying full Prescribing Information, including BOXED WARNING.

IPSEN CARES®
Coverage, Access, Reimbursement & Education Support

Dysport®
(abobotulinumtoxinA)

Getting Started with IPSEN CARES®

IPSEN CARES team members can help your enrolled patients understand and navigate their insurance coverage so they can start treatment with Dysport® (abobotulinumtoxinA).

How to Enroll

Enrolling your patient is as easy as filling out an IPSEN CARES Enrollment Form with your patient and having the patient review and sign the patient authorization.

Choose the method that works best for your office.

Fill Out and Submit Online

Complete and submit the form with your patient online at IPSENCARES.com

Fill Out Digitally and Fax

Download the form online at IPSENCARES.com, print, sign, and fax to 888-525-2416

Fill Out on Paper and Fax

Download and print the form, available at IPSENCARES.com or from your Patient Access Manager, then sign and fax to 888-525-2416

Patients may sign the Patient Authorization Form online at IPSENCARES.com.

We'll Take It From There

- Upon receiving a completed enrollment form, a Patient Access Manager (PAM) from IPSEN CARES will reach out to your patient
- The PAM will then conduct a Benefits Verification, assist with any access challenges, and identify financial support options for which your patient may qualify

IPSEN CARES supports patients in many ways– personalized support offerings are detailed on the next page

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Personalized Support Offerings

Dysport® (abobotulinumtoxinA)

IPSEN CARES® serves as a central point of contact among patients/caregivers, healthcare providers, insurance companies, and specialty pharmacies during treatment with Dysport®.

Financial & Insurance Assistance

- Benefits investigation to help understand the patient's health insurance coverage
- Financial assistance through the Copay Assistance Program
  - Patients may pay as little as $0 for each Dysport prescription if they have commercial insurance and meet eligibility criteria*
  - Submit claims for reimbursement by uploading required documentation via the copay website
- Free drug through the Patient Assistance Program for eligible* patients
- Prior authorizations and appeals support
- Help navigating the medical claims submission process

Dedicated, Individualized Support

- Information and support to facilitate interactions between healthcare providers, patients and their families, the insurance company, and the specialty pharmacy
- Guidance based on the patient's specific situation and healthcare needs

Medication Support Nurse Program

Patients prescribed Dysport who are enrolled in IPSEN CARES can received individualized support offerings provided by an IPSEN CARES nurse to help them through their treatment journey:

- Information to help patients understand how to work with their care team to set and achieve treatment goals
- Support to help patients receive their injections in a timely manner
- Medication Support Nurses are provided by Ipsen and do not work under the direction of the patient's healthcare provider or give medical advice

* Terms and conditions apply. Visit IPSENCARES.com to learn more.

Patient Access Managers are available
Monday – Friday, 8:00 AM – 8:00 PM ET
IPSENCARES.com
Phone: 866-435-5677 Fax: 888-525-2416

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INDICATIONS & IMPORTANT SAFETY INFORMATION

INDICATIONS

DYSPORT (abobotulinumtoxinA) for injection is indicated for the treatment of:

- spasticity in patients 2 years of age and older
- cervical dystonia in adults

IMPORTANT SAFETY INFORMATION

WARNING: DISTANT SPREAD OF TOXIN EFFECT

Postmarketing reports indicate that the effects of DYSPORT and all botulinum toxin products may spread from the area of injection to produce symptoms consistent with botulinum toxin effects. These may include asthenia, generalized muscle weakness, diplopia, blurred vision, ptosis, dysphagia, dysphonia, dysarthria, urinary incontinence and breathing difficulties. These symptoms have been reported hours to weeks after injection. Swallowing and breathing difficulties can be life threatening and there have been reports of death. The risk of symptoms is probably greatest in children treated for spasticity, but symptoms can also occur in adults treated for spasticity and other conditions, particularly in those patients who have underlying conditions that would predispose them to these symptoms. In unapproved uses and in approved indications, cases of spread of effect have been reported at doses comparable to or lower than the maximum recommended total dose.

Contraindications

DYSPORT is contraindicated in patients with known hypersensitivity to any botulinum toxin products, cow's milk protein, or to any of the components in the formulation, or infection at the proposed injection site(s). Serious hypersensitivity reactions including anaphylaxis, serum sickness, urticaria, soft tissue edema, and dyspnea have been reported. If such a serious reaction occurs, discontinue DYSPORT and institute appropriate medical therapy immediately.

Warnings and Precautions

Lack of Interchangeability Between Botulinum Toxin Products

The potency Units of DYSPORT are specific to the preparation and assay method utilized. They are not interchangeable with other preparations of botulinum toxin products and, therefore, units of biological activity of DYSPORT cannot be compared to or converted into units of any other botulinum toxin products assessed with any other specific assay method.

Dysphagia and Breathing Difficulties

Treatment with DYSPORT and other botulinum toxin products can result in swallowing or breathing difficulties. Patients with pre-existing swallowing or breathing difficulties may be more susceptible to these complications. In most cases, this is a consequence of weakening of muscles in the area of injection that are involved in breathing or swallowing. When distant effects occur, additional respiratory muscles may be involved. Deaths as a complication of severe dysphagia have been reported after treatment with botulinum toxin. Dysphagia may persist for several weeks and require use of a feeding tube to maintain adequate nutrition and hydration. Aspiration may result from severe dysphagia and is a particular risk when treating patients in whom swallowing or respiratory function is already compromised. Treatment of cervical dystonia with botulinum toxins may weaken accessory muscles of ventilation, which may result in a critical loss of breathing capacity in patients with respiratory disorders who may have become dependent upon these muscles. Patients treated with botulinum toxin may require immediate medical attention should they develop problems with swallowing, speech, or respiratory disorders. These reactions can occur within hours to weeks after injection with botulinum toxin.

Pre-existing Neuromuscular Disorders

Individuals with peripheral motor neuropathic diseases, amyotrophic lateral sclerosis, or neuromuscular junction disorders (e.g., myasthenia gravis or Lambert-Eaton syndrome) should be monitored particularly closely when given botulinum toxin. Patients with neuromuscular disorders may be at increased risk of clinically significant effects including severe dysphagia and respiratory compromise from typical doses of DYSPORT.

Human Albumin and Transmission of Viral Diseases

This product contains albumin, a derivative of human blood. Based on effective donor screening and product manufacturing processes, it carries an extremely remote risk for transmission of viral diseases and variant Creutzfeldt-Jakob disease (vCJD). There is a theoretical risk for transmission of Creutzfeldt-Jakob disease (CJD), but if that risk actually exists, the risk of transmission would also be considered extremely remote. No cases of transmission of viral diseases, vCJD, or CJD have ever been identified for licensed albumin or albumin contained in other licensed products.

Intradermal Immune Reaction

The possibility of an immune reaction when injected intradermally is unknown. The safety of DYSPORT for the treatment of hyperhidrosis has not been established. DYSPORT is approved only for intramuscular injection.

Pre-existing Conditions at the Injection Site

Caution should be exercised when DYSPORT is used where the targeted muscle shows excessive weakness or atrophy.

Adverse Reactions

- The most common adverse reactions (≥4%) in adults with upper limb spasticity include muscular weakness; in adults with lower limb spasticity (≥5%) include falls, muscular weakness, and pain in extremity
- The most common adverse reactions (≥10%) in pediatric patients with upper limb spasticity include upper respiratory tract infection and pharyngitis; in pediatric patients with lower limb spasticity include nasopharyngitis, cough, and pyrexia
- The most common adverse reactions (≥5%) in adults with cervical dystonia include muscular weakness, dysphagia, dry mouth, injection site discomfort, fatigue, headache, musculoskeletal pain, dysphonia, injection site pain, and eye disorders

Drug Interactions

Co-administration of DYSPORT and aminoglycosides or other agents interfering with neuromuscular transmission (e.g., curare-like agents) should only be performed with caution because the effect of the botulinum toxin may be potentiated. Use of anticholinergic drugs after administration of DYSPORT may potentiate systemic anticholinergic effects such as blurred vision. The effect of administering different botulinum neurotoxins at the same time or within several months of each other is unknown. Excessive weakness may be exacerbated by another administration of botulinum toxin prior to the resolution of the effects of a previously administered botulinum toxin. Excessive weakness may also be exaggerated by administration of a muscle relaxant before and after administration of DYSPORT.

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