

Please complete and sign this form and submit to customer service via email at **gmb-sps-piramal@cordlogistics.com** or fax it to 614-652-0536.



Date	Date Needed	Customer Number
License Number		License Expiration Date

SHIPPING INFORMATION		
Name		
Address 1		
Address 2		
City	State	ZIP
Contact Name		
Purchase Order Number		
Email Address*		

BILLING INFORMATION		
Name		
Address 1		
Address 2		
City	State	ZIP
Telephone Number		
Fax Number		
Email Address*		

*To receive an order confirmation, please provide an email address.

PRODUCT	TYPE	NDC #	QUANTITY
Gablofen® (baclofen injection) Prefilled Syringes			
500 mcg/mL (10,000mcg/20mL)	Prefilled Syringe	66794-155-01	
1000 mcg/mL (20,000mcg/20mL)	Prefilled Syringe	66794-156-01	
2000 mcg/mL (40,000mcg/20mL)	Prefilled Syringe	66794-157-01	
Gablofen® (baclofen injection) Vials			
500 mcg/mL (10,000mcg/20mL)	Ready-to-use Vial	66794-155-02	
1000 mcg/mL (20,000mcg/20mL)	Ready-to-use Vial	66794-156-02	
2000 mcg/mL (40,000mcg/20mL)	Ready-to-use Vial	66794-157-02	
Gablofen® (baclofen injection) Screening Dose			
50 mcg/mL (50mcg/20mL)	Screening Dose	66794-151-01	

Special Instructions:

- I do NOT want to receive refill kits with this order
- Rush delivery; an extra \$50 charge will apply (Standard delivery is 2 business days)
- State License on File
- Faxing Order
- Emailing Order

Mailing Address
 Piramal Critical Care, Inc.
 501 Mason Road, LaVergne, TN 37086

Contact Information
 Customer Service Email:
gmb-sps-piramal@cordlogistics.com
 Customer Service Phone:
 844-529-8995
 Customer Service Fax:
 614-652-0536

Reimbursement
 Customers seeking reimbursement from Medicare, Medicaid, or any other federal or state program may be required to disclose on cost report forms any discount from Piramal Critical Care Inc.

I certify that I am a licensed practitioner eligible to receive the selected medications, as required under the Prescription Drug Marketing Act. I further certify that I am eligible to prescribe, request, and dispense the above listed medications in the state where I am licensed and that my collaborative agreement and/or formulary, if applicable, permit(s) me to do the same. I am also the responsible person for purchases made at the above mentioned address under my state license number. I will notify Piramal immediately if my responsibility status and/or relationship with this facility is changed or terminated.

Print and sign or type your name in field after verifying professional licensure above.

SEE IMPORTANT RISK INFORMATION ON BACK PAGE

INDICATIONS AND USAGE

- Gablofen® (baclofen injection) is a gamma-aminobutyric acid (GABA) ergic agonist indicated for use in the management of severe spasticity of cerebral or spinal origin in adult and pediatric patients age 4 years and above.
- Gablofen should be reserved for patients unresponsive to oral baclofen therapy, or those who experience intolerable central nervous system side effects at effective doses.
- Patients should first respond to a screening dose of intrathecal baclofen prior to consideration for long term infusion via an implantable pump.
- Spasticity due to traumatic brain injury: wait at least one year after injury before considering Gablofen therapy.

IMPORTANT RISK INFORMATION

WARNING: DO NOT DISCONTINUE ABRUPTLY

See full prescribing information for complete boxed warning

Abrupt discontinuation of intrathecal baclofen, regardless of the cause, has resulted in sequelae that include high fever, altered mental status, exaggerated rebound spasticity, and muscle rigidity, that in rare cases has advanced to rhabdomyolysis, multiple organ-system failure and death.

Prevention of abrupt discontinuation of intrathecal baclofen requires careful attention to programming and monitoring of the infusion system, refill scheduling and procedures, and pump alarms. Patients and caregivers should be advised of the importance of keeping scheduled refill visits and should be educated on the early symptoms of baclofen withdrawal. Special attention should be given to patients at apparent risk (e.g. spinal cord injuries at T-6 or above, communication difficulties, history of withdrawal symptoms from oral or intrathecal baclofen). Consult the technical manual of the implantable infusion system for additional post-implant clinician and patient information.

CONTRAINDICATIONS

- Hypersensitivity to baclofen.
- Do not use Gablofen for intravenous, intramuscular, subcutaneous or epidural administration.

WARNINGS AND PRECAUTIONS

- Risk of life-threatening overdose during pump refills. Use extreme caution when filling the Medtronic SynchroMed® II Programmable Pump which is equipped with an injection port that allows direct access to the intrathecal catheter. Direct injection into the catheter through the catheter access port may cause a life-threatening overdose.
- Use only with Medtronic SynchroMed® II Programmable Pump (or other pumps labeled for intrathecal administration of Gablofen (baclofen injection)).
- Potential for contamination due to non-sterile external surface of prefilled syringe. Although the drug solution and pathway in the Gablofen prefilled syringes are sterile, the external surface of the prefilled syringes (all strengths, including the 50 mcg/mL strength) are non-sterile and have the potential to lead to contamination and consequent adverse reactions. The use of Gablofen prefilled syringe in an aseptic setting (e.g., operating room) to fill sterile intrathecal pumps prior to implantation in patients is not recommended, unless the external surface of the prefilled syringe is treated to ensure sterility. Gablofen supplied in vials may be used with conventional aseptic technique to fill intrathecal pumps prior to implantation.

- Resuscitative equipment and trained staff must be available during screening dose, dose titration, and refills due to the potential life-threatening CNS depression, cardiovascular collapse, and/or respiratory failure.
- Overdose may cause drowsiness, lightheadedness, dizziness, somnolence, respiratory depression, seizures, rostral progression of hypotonia and loss of consciousness progressing to coma.
- Use with caution in patients with psychotic disorders, schizophrenia or confusional states as it may exacerbate condition(s).
- Fatalities have been reported with intrathecal baclofen use.
- Caution should be used in patients with a history of autonomic dysreflexia.
- Presence of infection may increase the risk of surgical complication and complicate dosing of Gablofen.
- May cause drowsiness: use caution in operation of automobiles, dangerous machinery and activity that may be hazardous by decreased alertness. Other CNS depressants and alcohol may add to this effect.
- Potential development of intrathecal mass formation. Clinicians should monitor for signs and symptoms of new neurologic symptoms including the use of imaging diagnostic modalities.
- Oral baclofen use has been associated with a dose-related increase in incidence of ovarian cysts.

ADVERSE REACTIONS

SERIOUS ADVERSE REACTIONS

- Sudden withdrawal of Gablofen can result in serious complications that include high fever, confusion, muscle stiffness, multiple organ-system failure, and death. Inform patients that early symptoms of Gablofen withdrawal may include increased spasticity, itching, and tingling of extremities. If Gablofen withdrawal or a pump malfunction is suspected, patients should be brought immediately to a hospital for assessment and treatment.
- Gablofen overdose may occur suddenly or insidiously, and symptoms may include confusion, drowsiness, lightheadedness, dizziness, slow or shallow breathing, seizures, loss of muscle tone, loss of consciousness, and coma.
- Other serious adverse events may include: potential development of intrathecal mass formation, drainage, infection, meningitis, unmanageable trunk control, CSF leakage, coma and death.

COMMON ADVERSE REACTIONS

- The most common adverse reactions in patients with spasticity of spinal origin were hypotonia (25.3%), somnolence (20.9%), dizziness, nausea/vomiting, hypotension, headache, and convulsions.
- The most common adverse reactions in patients with spasticity of cerebral origin were hypotonia (34.7%), somnolence (18.7%), headache (10.7%), agitation, constipation, leukocytosis, chills, and urinary retention.
- Other common adverse events may include hypoventilation, hypertonia, paresthesia, increased salivation, back pain, pruritus, diarrhea, peripheral edema, asthenia, pain, confusion, speech disorder, amblyopia, accidental injury, and dry mouth.

USE IN SPECIFIC POPULATIONS

- Pregnancy Category C. The effect of baclofen in labor and delivery is unknown.
- Breastfeeding: Baclofen is excreted into breast milk at oral therapeutic doses.
- Pediatric use: Safety and effectiveness in pediatric patients below the age of 4 years have not been established.

To report suspected adverse reactions, call 888.525.8114, contact the FDA at 800.FDA.1088 or <http://www.fda.gov/medwatch>.