



Medtronic

Neuromodulation Products

HCPCS II and C-Codes

2011

Medtronic provides this information for your convenience only. It is not intended as a recommendation regarding clinical practice. It is always the provider's responsibility to determine coverage and submit appropriate codes, modifiers, and charges for the services that were rendered. This document provides assistance for FDA approved or cleared indications. Where reimbursement is requested for a use of a product that may be inconsistent with, or not expressly specified in, the FDA cleared or approved labeling (e.g., instructions for use, operator's manual or package insert) consult with your billing advisors or payers for advice on handling such billing issues. Some payers may have policies that make it inappropriate to submit claims for such items or related service. Contact your Medicare contractor or other payer for interpretation of coverage, coding, and payment policies.

HCPCS II Codes¹

HCPCS II codes are a supplement to CPT® codes.² Although some HCPCS II codes are for procedures and services not classified in CPT, the majority of HCPCS II codes are for supplies, DME, drugs, and medical devices. In many situations, CPT and HCPCS II codes must be used together to completely describe a service. In particular, CPT codes indicate the procedure performed and HCPCS II codes identify the specific device, supply, DME, or drug utilized in the procedure.

Example

CPT	63650	Percutaneous implantation of neurostimulator electrode array, epidural
HCPCS II	L8680	Implantable neurostimulator electrode, each

HCPCS II codes are utilized by the entity that purchased and supplied the medical device, DME, drug, or supply to the patient. HCPCS II codes for medical devices usually begin with an E or an L.

C-Codes

C-codes are a special type of HCPCS II code. They are used only by hospitals when billing Medicare for medical devices utilized in the outpatient setting. Although other payers may accept C-codes, HCPCS II E and L codes are generally used for billing non-Medicare payers. For Medicare, billing C-codes is mandatory for medical devices utilized in the outpatient setting.

Device Edits Medicare

Medicare's Consolidated Device Edits require that when specific CPT procedure codes for device implantation are billed, associated HCPCS II codes for the devices must also be billed.³ When a hospital outpatient bill is received that contains one of the specific CPT procedure codes without one of the required HCPCS II codes, the claim is returned to the provider for correction. Some of the edits also work in reverse; when a device HCPCS II code is present on the bill, the associated CPT procedure code must also be present.

This edit is necessary to ensure that device charges are correctly submitted and are uniquely identifiable on hospital outpatient claims. This is essential because Medicare uses the claims data to set future payment rates for outpatient hospital procedures under the APC prospective payment system.

1. Healthcare Common Procedure Coding System (HCPCS) Level II codes are maintained by the Centers for Medicare and Medicaid Services. More information can be found at: http://www.cms.hhs.gov/MedHCPCSGenInfo/01_Overview.asp#TopOfPage

2. CPT copyright 2010 American Medical Association. All rights reserved. CPT is a registered trademark of the American Medical Association. No fee schedules, basic units, relative values, or related listings are included in CPT. The AMA assumes no liability for the data contained herein. Applicable FARS/DFARS restrictions apply to government use.

3. Device edits are updated once per quarter and can be found at http://www.cms.hhs.gov/HospitalOutpatientPPS/02_device_procedure.asp#TopOfPage.

Medtronic Deep Brain Stimulation Therapy

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Neurostimulators			
Kinetra® Neurostimulator	7428	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Soletta® Neurostimulator	7426	L8686 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Activa® SC	37602 37603	L8686 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Activa® PC Neurostimulator	37601	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Activa® RC Neurostimulator	37612	L8687 (includes the generator, extension, patient programmer, and recharging system)	C1820 (includes the generator, extension, patient programmer, and recharging system)
Patient Programmers			
DBS Patient Programmer (for Activa® RC and Activa® PC)	37642	L8681 ³ (programmer, replacement only)	C1787
Access® Therapy Controller	7436	L8681 ³ (programmer, replacement only)	C1787
Access® Review Therapy Controller	7438	L8681 ³ (programmer, replacement only)	C1787
Recharger			
Patient Recharger	37651	L8689 ⁴ (recharger, replacement only)	N/A (included in C1820)
Leads			
DBS Lead Kit (1.5 mm Spaces)	3387S-40	L8680	N/A ²
DBS Lead Kit (0.5 mm Spaces)	3389S-40	L8680	N/A ²
DBS Lead Kit (4.0 mm Spaces)	3391S-40	L8680	N/A ²
Lead Extensions			
Low Profile DBS Extension	7482A25 7482A40 7482A51 7482A66 7482A95	Included in L8686 and L8688	C1883
8×4 Extension	37085-40 37085-60 37085-95	Included in L8687 (rechargeable) L8688 (non-rechargeable)	C1883 (adaptor)

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Medtronic Deep Brain Stimulation Therapy *continued*

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Fiducial Markers			
Unibody® Fiducial Markers	FM-4007 FM-4010 FM-4013	A4648 ⁵	C1879
Pocket Adaptor			
Pocket Adaptor (1×4)	64001	Included in L8687 (rechargeable) L8688 (non-rechargeable)	C1883 (adaptor)
Pocket Adaptor (2×4)	64002	Included in L8687 (rechargeable) L8688 (non-rechargeable)	C1883 (adaptor)

1. Healthcare Common Procedure Coding System (HCPCS) Level II codes are maintained by the Centers for Medicare and Medicaid Services. More information can be found at: http://www.cms.hhs.gov/MedHCPCSGenInfo/01_Overview.asp#TopOfPage.

2. CPT Code for lead implantation is classified as inpatient only.

3. L8681 is to be used only for replacement of programmer and is not to be separately coded with the initial generator. L8681 defined as: Patient Programmer (external) for use with implantable programmable pulse generator, replacement only.

4. L8689 is to be used only for replacement of recharger and is not to be separately coded with the initial generator. L8689 defined as: External recharging system for battery (internal) for use with implantable neurostimulator, replacement only.

5. Code A4648 cannot be used by physicians for fiducials implanted in the office (see Transmittal 604, CR6579, November 27, 2009).

Drug Delivery Systems

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Pumps			
SynchroMed® II Infusion Pump, programmable	863720	E0783 (entire system: pump, catheter, connectors)	C1772 (pump)
	863740	E0786 (pump replacement excluding catheter)	
Personal Therapy Manager			
myPTM® Personal Therapy Manager	8835	A9900 ³	N/A
Refill Kits			
IsoMed® Pump Refill Kit	8553	A4220	N/A
SynchroMed® Pump Refill Kit	8551	A4220	N/A
Catheters			
Intrathecal Catheter (Sutureless Pump Connector)	8731SC	E0785 (replacement catheter only)	C1755 (catheter)
InDura® Free Flow Intrathecal Catheter	8711	E0785 (replacement catheter only)	C1755 (catheter)
InDura® 1P Intrathecal Catheter	8709	E0785 (replacement catheter only)	C1755 (catheter)
InDura® 1P Intrathecal Catheter (Sutureless Pump Connector)	8709SC	E0785 (replacement catheter only)	C1755 (catheter)
Percutaneous Intraspinial Trial Catheter Kit	8516	E1399 (trial catheter only)	C1755 (catheter)
AlgoLine Intraspinial Catheter	81102	E0785 (replacement catheter only)	C1755 (catheter)
	81192		

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2. This miscellaneous code can be used for replacement of the myPTM®. There is no code for initial myPTM®. Check with payer for appropriate coding of initial myPTM®.

Drug

Drug Name and Description	NDC Number	Medtronic Model #	HCPCS II Code	Total Drug Quantity	Billing Units	C-Code and Description
Lioresal™ Intrathecal (baclofen injection) Screening ampule of 0.05 mg/1 mL (50 mcg/mL)	58281-0562-01	8563S	J0476	.05 mg	1	N/A
Lioresal™ Intrathecal (baclofen injection) Refill Kit, one ampule of 10 mg/20 mL (500 mcg/mL)	58281-0560-01	8561	J0475	10 mg	1	N/A
Lioresal™ Intrathecal (baclofen injection) Refill Kit, two ampules of 10 mg/5 mL (2,000 mcg/mL)	58281-0561-02	8562	J0475	20 mg	2	N/A
Lioresal™ Intrathecal (baclofen injection) Refill Kit, one ampule of 40 mg/20 mL (2,000 mcg/mL)	58281-0563-01	8564	J0475	40 mg	4	N/A
Lioresal™ Intrathecal (baclofen injection) Refill Kit, two ampules of 10 mg/20 mL (500 mcg/mL)	58281-0560-02	8565	J0475	20 mg	2	N/A
Lioresal™ Intrathecal (baclofen injection) Refill Kit, two ampules of 40 mg/20 mL (2,000 mcg/mL)	58281-0563-02	8566	J0475	80 mg	8	N/A
Infumorph™ (preservative-free morphine sulphate sterile solution), per 10 mg			J2275	10 mg	1	N/A
Prialt® (ziconotide intrathecal infusion, 1 mcg)			J2278	1 mcg	1	N/A

Infumorph™ is a trademark of Baxter Healthcare Corporation.
 Prialt® is a registered trademark of Azur Pharma International Limited.

Enterra® Therapy

Medtronic Product Name and Description	Current Medtronic Model #	Previous Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Neurostimulators				
Enterra® Neurostimulator	3116	3116	L8688	C1767
Leads				
Enterra® Leads	435135 435150D	4351	L8680	C1778

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InterStim® Therapy for Urinary Control

Medtronic Product Name and Description	Current Medtronic Model #	Previous Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Neurostimulators				
InterStim® Neurostimulator	3023	30230134	L8688	C1767
InterStim® II Neurostimulator	3058	30580134	L8686	C1767
Patient Programmers				
InterStim® Patient Programmer	3031A	3031A	L8681 ² (programmer, replacement only)	C1787
InterStim iCon® Patient Programmer	3037	3037	L8681 ² (programmer, replacement only)	C1787
Lead Introducer				
Lead Introducer Kit	355018	355018	N/A	C1894
Lead Introducer (box of 6)	042294	042294	N/A	C1894
Leads				
InterStim® Test Stimulation Lead Kit (sacral nerve stimulation)	3065USC TL22	3065USC TL22	A4290	C1897
Test Kit Leads	3057-1SC (single lead) 3057-6SC (six leads)		L8680	C1778
InterStim® Leads	3093-28 3093-33 3093-41 3889-28 3889-33 3889-41	309328 309333 309341 388928 388933 388941	L8680	C1778
Extensions				
All Extensions	3095-10 3095-25 3095-51	309510 309525 309551	N/A	C1883

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2. L8681 is to be used only for replacement of programmer and is not to be separately coded with the initial generator. L8681 defined as: Patient Programmer (external) for use with implantable programmable pulse generator, replacement only.

Neurostimulation

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Neurostimulators			
RestoreUltra® implantable rechargeable neurostimulator (INS)	37712	L8687 (includes the generator, extension, patient programmer and external recharging system)	C1820 (includes the generator, extension, patient programmer and external recharging system)
RestoreAdvanced® system kit	37713KIT	L8687 (includes the generator, extension, patient programmer and external recharging system)	C1820 (includes the generator, extension, patient programmer and external recharging system)
RestoreAdvanced® implantable rechargeable neurostimulator (INS)	37713	L8687 (includes the generator, extension, patient programmer and external recharging system)	C1820 (includes the generator, extension, patient programmer and external recharging system)
PrimeAdvanced® system kit	37702KIT	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
PrimeAdvanced® implantable non-rechargeable neurostimulator (INS)	37702	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Restore® system kit	37711KIT	L8687 (includes the generator, extension, patient programmer and external recharging system)	C1820 (includes the generator, extension, patient programmer and external recharging system)
Restore® implantable rechargeable neurostimulator (INS)	37711	L8687 (includes the generator, extension, patient programmer and external recharging system)	C1820 (includes the generator, extension, patient programmer and external recharging system)
RestorePrime® (implantable non-rechargeable neurostimulator (INS)	37701	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Synergy® EZ system kit	7729US	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Synergy® implantable neurostimulator (INS)	7427	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Synergy Versitrel® EZ system kit	7719US	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)

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Neurostimulation *continued*

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Synergy Versitrel [®] implantable neurostimulator (INS)	7427V-NP	L8688 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Itrel [®] 3 EZ system kit	7425EZ	L8686 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Itrel [®] 3 implantable neurostimulator (INS)	7425	L8686 (includes the generator, extension, and patient programmer)	C1767 (includes the generator, extension, and patient programmer)
Patient Programmers			
MyStim [®] Patient Programmer (for use with Restore [®] family of neurostimulators)	37742 37743	L8681 (patient programmer; replacement only)	C1787 (patient programmer; replacement only)
Synergy [®] EZ patient programmer (hand-held)	7435	L8681 (patient programmer; replacement only)	C1787 (patient programmer; replacement only)
Itrel [®] EZ patient programmer (hand-held)	7434A	L8681 (patient programmer; replacement only)	C1787 (patient programmer; replacement only)
Rechargers			
External patient recharger system (for use with Restore [®] , RestoreAdvanced [®] , and RestoreUltra [®] neurostimulators)	37752	L8689 (external recharging system; replacement only)	N/A (included in C1820)

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Neurostimulation *continued*

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Leads			
1×8 Compact lead kit	3778-45 3778-60 3778-75	L8680	C1778 (lead)
1×8 Subcompact lead kit	3776-45 3776-60 3776-75	L8680	C1778 (lead)
1×8 Standard lead kit	3777-30 3777-45 3777-60 3777-75	L8680	C1778 (lead)
2×4 Hinged Specify® lead kit	3999-30 3999-45 3999-60	L8680	C1778 (lead)
On-Point® peripheral nerve stimulation lead kit	3987A 3987A-60	L8680	C1778 (lead)
Pisces-Quad® Compact lead kit	3887-33 3887-45 3887-56	L8680	C1778 (lead)
Pisces-Quad® lead kit	3487A-28 3487A-33 3487A-45 3487A-56	L8680	C1778 (lead)
Pisces-Quad® Plus lead kit	3888-33 3888-45 3888-56	L8680	C1778 (lead)
Resume II® lead kit	3587A-25 3587A-70	L8680	C1778 (lead)
Resume® TL lead kit	3986A 3986A-45 3986A-60 3986A-70	L8680	C1778 (lead)
Specify® lead kit	3998 399821	L8680	C1778 (lead)
Specify® 2x8 lead kit	39286-30 39286-65	L8680	C1778 (lead)
Specify® 5-6-5 lead kit	39565-30 39565-65	L8680	C1778 (lead)

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Neurostimulation *continued*

Medtronic Product Name and Description	Medtronic Model #	HCPCS II Code ¹ and Description	C-Code and Description
Adaptors/Extensions			
1×8 Extension	3708120 3708140 3708160	N/A (included in L8686, L8687, or L8688)	C1883 (extension)
Low Profile Quad Extension (with in-line connector) and accessories	748910 748925 748940 748951 748966	N/A (included in L8686, L8687, or L8688)	C1883 (extension)
Percutaneous Extension (in-line connector for quadripolar leads only)	3550-05	N/A (included in L8686, L8687, or L8688)	C1883 (extension)
Single Stretch-Coil™ Extension	3708320 3708340 3708360	N/A (included in L8686, L8687, or L8688)	C1883 (extension)
Bifurcated Stretch-Coil™ Extension	3708220 3708240 3708260	N/A (included in L8686, L8687, or L8688)	C1883 (extension)
1×4 Pocket Adaptor	74001	N/A (included in L8686, L8687, or L8688)	C1883 (adaptor)
2×4 Pocket Adaptor	74002	N/A (included in L8686, L8687, or L8688)	C1883 (adaptor)
Trial Leads			
1×8 Compact Trial lead kit	387445 387460	L8680	C1897 (lead)
1×8 Standard Trial lead kit	387345 387360	L8680	C1897 (lead)
1×8 Subcompact Trial lead kit	3875-45 3875-60	L8680	C1897 (lead)

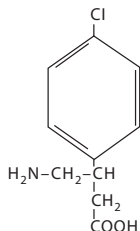
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LIORESAL® INTRATHECAL (baclofen injection)

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 postimplant clinician and patient information (see WARNINGS)

DESCRIPTION

LIORESAL INTRATHECAL (baclofen injection) is a muscle relaxant and antispastic. Its chemical name is 4- amino- 3-(4- chlorophenyl) butanoic acid, and its structural formula is:



Baclofen is a white to off-white, odorless or practically odorless crystalline powder, with a molecular weight of 213.66. It is slightly soluble in water, very slightly soluble in methanol, and insoluble in chloroform.

LIORESAL INTRATHECAL is a sterile, pyrogen-free, isotonic solution free of antioxidants, preservatives or other potentially neurotoxic additives indicated only for intrathecal administration. The drug is stable in solution at 37° C and compatible with CSF. Each milliliter of LIORESAL INTRATHECAL contains baclofen U. S. P. 50 mcg, 500 mcg or 2000 mcg and sodium chloride 9 mg in Water for Injection; pH range is 5.0 - 7.0. Each ampule is intended for SINGLE USE ONLY. Discard any unused portion. **DO NOT AUTOCLAVE.**

CLINICAL PHARMACOLOGY

The precise mechanism of action of baclofen as a muscle relaxant and antispasticity agent is not fully understood. Baclofen inhibits both monosynaptic and polysynaptic reflexes at the spinal level, possibly by decreasing excitatory neurotransmitter release from primary afferent terminals, although actions at supraspinal sites may also occur and contribute to its clinical effect. Baclofen is a structural analog of the inhibitory neurotransmitter gamma-aminobutyric acid (GABA), and may exert its effects by stimulation of the GABA_B receptor subtype.

LIORESAL INTRATHECAL when introduced directly into the intrathecal space permits effective CSF concentrations to be achieved with resultant plasma concentrations 100 times less than those occurring with oral administration.

In people, as well as in animals, baclofen has been shown to have general CNS depressant properties as indicated by the production of sedation with tolerance, somnolence, ataxia, and respiratory and cardiovascular depression.

Pharmacodynamics of LIORESAL INTRATHECAL:

Intrathecal Bolus:

Adult Patients: The onset of action is generally one-half hour to one hour after an intrathecal bolus. Peak spasmolytic effect is seen at approximately four hours after dosing and effects may last four to eight hours. Onset, peak response, and duration of action may vary with individual patients depending on the dose and severity of symptoms.

Pediatric Patients: The onset, peak response and duration of action is similar to those seen in adult patients.

Continuous Infusion:

LIORESAL INTRATHECAL'S antispastic action is first seen at 6 to 8 hours after initiation of continuous infusion. Maximum activity is observed in 24 to 48 hours.

Continuous Infusion: No additional information is available for pediatric patients.

Pharmacokinetics of LIORESAL INTRATHECAL:

The pharmacokinetics of CSF clearance of LIORESAL INTRATHECAL calculated from intrathecal bolus or continuous infusion studies approximates CSF turnover, suggesting elimination is by bulk-flow removal of CSF.

Intrathecal Bolus: After a bolus lumbar injection of 50 or 100 mcg LIORESAL INTRATHECAL in seven patients, the average CSF elimination half-life was 1.51 hours over the first four hours and the average CSF clearance was approximately 30 ml/hour.

Continuous infusion: The mean CSF clearance for LIORESAL INTRATHECAL (baclofen injection) was approximately 30 ml/hour in a study involving ten patients on continuous intrathecal infusion. Concurrent plasma concentrations of baclofen during intrathecal administration are expected to be low (0- 5 ng/ml).

Limited pharmacokinetic data suggest that a lumbar-cisternal concentration gradient of about 4: 1 is established along the neuroaxis during baclofen infusion. This is based upon simultaneous CSF sampling via cisternal and lumbar tap in 5 patients receiving continuous baclofen infusion at the lumbar level at doses associated with therapeutic efficacy; the interpatient variability was great. The gradient was not altered by position. Six pediatric patients (age 8- 18 years) receiving continuous intrathecal baclofen infusion at doses of 77- 400 mcg/ day had plasma baclofen levels near or below 10 ng/ml.

INDICATIONS

LIORESAL INTRATHECAL is indicated for use in the management of severe spasticity. Patients should first respond to a screening dose of intrathecal baclofen prior to consideration for long term infusion via an implantable pump. For spasticity of spinal cord origin, chronic infusion of LIORESAL INTRATHECAL via an implantable pump should be reserved for patients unresponsive to oral baclofen therapy, or those who experience intolerable CNS side effects at effective doses. Patients with spasticity due to traumatic brain injury should wait at least one year after the injury before consideration of long term intrathecal baclofen therapy. LIORESAL INTRATHECAL (baclofen injection) is intended for use by the intrathecal route in single bolus test doses (via spinal catheter or lumbar puncture) and, for chronic use, only in implantable pumps approved by the FDA specifically for the administration of LIORESAL INTRATHECAL into the intrathecal space.

Spasticity of Spinal Cord Origin: Evidence supporting the efficacy of LIORESAL INTRATHECAL was obtained in randomized, controlled investigations that compared the effects of either a single intrathecal dose or a three day intrathecal infusion of LIORESAL INTRATHECAL to placebo in patients with severe spasticity and spasms due to either spinal cord trauma or multiple sclerosis. LIORESAL INTRATHECAL was superior to placebo on both principal outcome measures employed: change from baseline in the Ashworth rating of spasticity and the frequency of spasms.

Spasticity of Cerebral Origin: The efficacy of LIORESAL INTRATHECAL was investigated in three controlled clinical trials; two enrolled patients with cerebral palsy and one enrolled patients with spasticity due to previous brain injury. The first study, a randomized controlled cross-over trial of 51 patients with cerebral palsy, provided strong, statistically significant results; LIORESAL INTRATHECAL was superior to placebo in reducing spasticity as measured by the Ashworth Scale. A second cross-over study was conducted in 11 patients with spasticity arising from brain injury. Despite the small sample size, the study yielded a nearly significant test statistic (p= 0.066) and provided directionally favorable results. The last study, however, did not provide data that could be reliably analyzed.

LIORESAL INTRATHECAL therapy may be considered an alternative to destructive neurosurgical procedures. Prior to implantation of a device for chronic intrathecal infusion of LIORESAL INTRATHECAL, patients must show a response to LIORESAL INTRATHECAL in a screening trial (see Dosage and Administration).

CONTRAINDICATIONS

Hypersensitivity to baclofen. LIORESAL INTRATHECAL is not recommended for intravenous, intramuscular, subcutaneous or epidural administration.

WARNINGS

LIORESAL INTRATHECAL is for use in single bolus intrathecal injections (via a catheter placed in the lumbar intrathecal space or injection by lumbar puncture) and in implantable pumps approved by the FDA specifically for the intrathecal administration of baclofen. Because of the possibility of potentially life-threatening CNS depression, cardiovascular collapse, and/ or respiratory failure, physicians must be adequately trained and educated in chronic intrathecal infusion therapy.

The pump system should not be implanted until the patient's response to bolus LIORESAL INTRATHECAL injection is adequately evaluated. Evaluation (consisting of a screening procedure: see Dosage and Administration) requires that LIORESAL INTRATHECAL be administered into the intrathecal space via a catheter or lumbar puncture. Because of the risks associated with the screening procedure and the adjustment of dosage following pump implantation, these phases must be conducted in a medically supervised and adequately equipped environment following the instructions outlined in the Dosage and Administration section.

Resuscitative equipment should be available.

Following surgical implantation of the pump, particularly during the initial phases of pump use, the patient should be monitored closely until it is certain that the patient's response to the infusion is acceptable and reasonably stable.

On each occasion that the dosing rate of the pump and/ or the concentration of LIORESAL INTRATHECAL (baclofen injection) in the reservoir is adjusted, close medical monitoring is required until it is certain that the patient's response to the infusion is acceptable and reasonably stable.

It is mandatory that the patient, all patient caregivers, and the physicians responsible for the patient receive adequate information regarding the risks of this mode of treatment. All medical personnel and caregivers should be instructed in 1) the signs and symptoms of overdose, 2) procedures to be followed in the event of overdose and 3) proper home care of the pump and insertion site.

Overdose: Signs of overdose may appear suddenly or insidiously. Acute massive overdose may present as coma. Less sudden and/ or less severe forms of overdose may present with signs of drowsiness, lightheadedness, dizziness, somnolence, respiratory depression, seizures, rostral progression of hypotonia and loss of consciousness progressing to coma. Should overdose appear likely, the patient should be taken immediately to a hospital for assessment and emptying of the pump reservoir. In cases reported to date, overdose has generally been related to pump malfunction or dosing error. (See Drug Overdose Symptoms and Treatment.)

Extreme caution must be used when filling an FDA approved implantable pump. Such pumps should only be refilled through the reservoir refill septum. However, some pumps are also equipped with a catheter access port that allows direct access to the intrathecal catheter. Direct injection into this catheter access port may cause a life-threatening overdose.

Withdrawal: Abrupt withdrawal of intrathecal baclofen, regardless of the cause, has resulted in sequelae that included high fever, altered mental status, exaggerated rebound spasticity and muscle rigidity that in rare cases progressed to rhabdomyolysis, multiple organ-system failure, and death. In the first 9 years of post-marketing experience, 27 cases of withdrawal temporally related to the cessation of baclofen therapy were reported; six patients died. In most cases, symptoms of withdrawal appeared within hours to a few days following interruption of intrathecal baclofen therapy. Common reasons for abrupt interruption of intrathecal baclofen therapy included malfunction of the

catheter (especially disconnection), low volume in the pump reservoir, and end of pump battery life; human error may have played a causal or contributing role in some cases. 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.

All patients receiving intrathecal baclofen therapy are potentially at risk for withdrawal. Early symptoms of baclofen withdrawal may include return of baseline spasticity, pruritus, hypotension, and paresthesias. Some clinical characteristics of the advanced intrathecal baclofen withdrawal syndrome may resemble autonomic dysreflexia, infection (sepsis), malignant hyperthermia, neuroleptic-malignant syndrome, or other conditions associated with a hypermetabolic state or widespread rhabdomyolysis.

Rapid, accurate diagnosis and treatment in an emergency-room or intensive-care setting are important in order to prevent the potentially life-threatening central nervous system and systemic effects of intrathecal baclofen withdrawal. The suggested treatment for intrathecal baclofen withdrawal is the restoration of intrathecal baclofen at or near the same dosage as before therapy was interrupted. However, if restoration of intrathecal delivery is delayed, treatment with GABA-ergic agonist drugs such as oral or enteral baclofen, or oral, enteral, or intravenous benzodiazepines may prevent potentially fatal sequelae. Oral or enteral baclofen alone should not be relied upon to halt the progression of intrathecal baclofen withdrawal.

Seizures have been reported during overdose and with withdrawal from LIORESAL INTRATHECAL as well as in patients maintained on therapeutic doses of LIORESAL INTRATHECAL.

Fatalities:

Spasticity of Spinal Cord Origin: There were 16 deaths reported among the 576 U.S. patients treated with LIORESAL INTRATHECAL (baclofen injection) in pre- and post-marketing studies evaluated as of December 1992. Because these patients were treated under uncontrolled clinical settings, it is impossible to determine definitively what role, if any, LIORESAL INTRATHECAL played in their deaths. As a group, the patients who died were relatively young (mean age was 47 with a range from 25 to 63), but the majority suffered from severe spasticity of many years duration, were nonambulatory, had various medical complications such as pneumonia, urinary tract infections, and decubiti, and/or had received multiple concomitant medications. A case-by-case review of the clinical course of the 16 patients who died failed to reveal any unique signs, symptoms, or laboratory results that would suggest that treatment with LIORESAL INTRATHECAL caused their deaths. Two patients, however, did suffer sudden and unexpected death within 2 weeks of pump implantation and one patient died unexpectedly after screening.

One patient, a 44-year-old male with MS, died in hospital on the second day following pump implantation. An autopsy demonstrated severe fibrosis of the coronary conduction system. A second patient, a 52-year-old woman with MS and a history of an inferior wall myocardial infarction, was found dead in bed 12 days after pump implantation, 2 hours after having had documented normal vital signs. An autopsy revealed pulmonary congestion and bilateral pleural effusions. It is impossible to determine whether LIORESAL INTRATHECAL contributed to these deaths. The third patient underwent three baclofen screening trials. His medical history included SCI, aspiration pneumonia, septic shock, disseminated intravascular coagulopathy, severe metabolic acidosis, hepatic toxicity, and status epilepticus. Twelve days after screening (he was not implanted), he again experienced status epilepticus with subsequent significant neurological deterioration. Based upon prior instruction, extraordinary resuscitative measures were not pursued and the patient died.

Spasticity of Cerebral Origin: There were three deaths occurring among the 211 patients treated with LIORESAL INTRATHECAL in pre-marketing studies as of March 1996. These deaths were not attributed to the therapy.

PRECAUTIONS

Children should be of sufficient body mass to accommodate the implantable pump for chronic infusion. Please consult pump manufacturer's manual for specific recommendations.

Safety and effectiveness in pediatric patients below the age of 4 have not been established.

Screening

Patients should be infection-free prior to the screening trial with LIORESAL INTRATHECAL (baclofen injection) because the presence of a systemic infection may interfere with an assessment of the patient's response to bolus LIORESAL INTRATHECAL.

Pump Implantation

Patients should be infection-free prior to pump implantation because the presence of infection may increase the risk of surgical complications. Moreover, a systemic infection may complicate dosing.

Pump Dose Adjustment and Titration

In most patients, it will be necessary to increase the dose gradually over time to maintain effectiveness; a sudden requirement for substantial dose escalation typically indicates a catheter complication (i.e., catheter kink or dislodgement).

Reservoir refilling must be performed by fully trained and qualified personnel following the directions provided by the pump manufacturer. Refill intervals should be carefully calculated to prevent depletion of the reservoir, as this would result in the return of severe spasticity and possibly symptoms of withdrawal.

Strict aseptic technique in filling is required to avoid bacterial contamination and serious infection. A period of observation appropriate to the clinical situation should follow each refill or manipulation of the drug reservoir.

Extreme caution must be used when filling an FDA approved implantable pump 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.

Additional considerations pertaining to dosage adjustment: It may be important to titrate the dose to maintain some degree of muscle tone and allow occasional spasms to: 1) help support circulatory function, 2) possibly prevent the formation of deep vein thrombosis, 3) optimize activities of daily living and ease of care.

Except in overdose related emergencies, the dose of LIORESAL INTRATHECAL should ordinarily be reduced slowly if the drug is discontinued for any reason.

An attempt should be made to discontinue concomitant oral antispasticity medication to avoid possible overdose or adverse drug interactions, either prior to screening or following implant and initiation of chronic LIORESAL INTRATHECAL infusion. Reduction and discontinuation of oral anti-spasmodics should be done slowly and with careful monitoring by the physician. Abrupt reduction or discontinuation of concomitant antispasmodics should be avoided.

Drowsiness: Drowsiness has been reported in patients on LIORESAL INTRATHECAL. Patients should be cautioned regarding the operation of automobiles or other dangerous machinery, and activities made hazardous by decreased alertness. Patients should also be cautioned that the central nervous system depressant effects of LIORESAL INTRATHECAL (baclofen injection) may be additive to those of alcohol and other CNS depressants.

Precautions in special patient populations: Careful dose titration of LIORESAL INTRATHECAL is needed when spasticity is necessary to sustain upright posture and balance in locomotion or whenever spasticity is used to obtain optimal function and care.

Patients suffering from psychotic disorders, schizophrenia, or confusional states should be treated cautiously with LIORESAL INTRATHECAL and kept under careful surveillance, because exacerbations of these conditions have been observed with oral administration.

LIORESAL INTRATHECAL should be used with caution in patients with a history of autonomic dysreflexia. The presence of nociceptive stimuli or abrupt withdrawal of LIORESAL INTRATHECAL (baclofen injection) may cause an autonomic dysreflexic episode.

Because LIORESAL is primarily excreted unchanged by the kidneys, it should be given with caution in patients with impaired renal function and it may be necessary to reduce the dosage.

LABORATORY TESTS

No specific laboratory tests are deemed essential for the management of patients on LIORESAL INTRATHECAL.

DRUG INTERACTIONS

There is inadequate systematic experience with the use of LIORESAL INTRATHECAL in combination with other medications to predict specific drug-drug interactions. Interactions attributed to the combined use of LIORESAL INTRATHECAL and epidural morphine include hypotension and dyspnea.

CARCINOGENESIS, MUTAGENESIS, AND IMPAIRMENT OF FERTILITY

No increase in tumors was seen in rats receiving LIORESAL (baclofen USP) orally for two years at approximately 30-60 times on a mg/kg basis, or 10-20 times on a mg/m² basis, the maximum oral dose recommended for human use. Mutagenicity assays with LIORESAL have not been performed.

PREGNANCY CATEGORY C

LIORESAL (baclofen USP) given orally has been shown to increase the incidence of omphaloceles (ventral hernias) in fetuses of rats given approximately 13 times on a mg/kg basis, or 3 times on a mg/m² basis, the maximum oral dose recommended for human use; this dose also caused reductions in food intake and weight gain in the dams.

This abnormality was not seen in mice or rabbits. There are no adequate and well-controlled studies in pregnant women. LIORESAL should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

NURSING MOTHERS

In mothers treated with oral LIORESAL (baclofen USP) in therapeutic doses, the active substance passes into the breast milk. It is not known whether detectable levels of drug are present in breast milk of nursing mothers receiving LIORESAL INTRATHECAL. As a general rule, nursing should be undertaken while a patient is receiving LIORESAL INTRATHECAL only if the potential benefit justifies the potential risks to the infant.

PEDIATRIC USE

Children should be of sufficient body mass to accommodate the implantable pump for chronic infusion. Please consult pump manufacturer's manual for specific recommendations.

Safety and effectiveness in pediatric patients below the age of 4 have not been established.

Considerations based on experience with oral LIORESAL (baclofen USP)

A dose-related increase in incidence of ovarian cysts was observed in female rats treated chronically with oral LIORESAL. Ovarian cysts have been found by palpation in about 4% of the multiple sclerosis patients who were treated with oral LIORESAL for up to one year. In most cases these cysts disappeared spontaneously while patients continued to receive the drug. Ovarian cysts are estimated to occur spontaneously in approximately 1% to 5% of the normal female population.

ADVERSE DRUG EVENTS

Spasticity of Spinal Cord Origin:

Commonly Observed in Patients with Spasticity of Spinal Origin — In pre- and post-marketing clinical trials, the most commonly observed adverse events associated with use of LIORESAL INTRATHECAL (baclofen injection) which were not seen at an equivalent incidence among placebo-treated patients were: somnolence, dizziness, nausea, hypotension, headache, convulsions and hypotonia.

Associated with Discontinuation of Treatment — 8/474 patients with spasticity of spinal cord origin receiving long term infusion of LIORESAL INTRATHECAL in pre- and post-marketing clinical studies in the U. S. discontinued treatment due to adverse events. These include: pump pocket infections (3), meningitis (2), wound dehiscence (1), gynecological fibroids (1) and pump overpressurization (1) with unknown, if any, sequela. Eleven patients who developed coma secondary to overdose had their treatment temporarily suspended, but all were subsequently re-started and were not, therefore, considered to be true discontinuations.

Fatalities — See Warnings.

Incidence in Controlled Trials — Experience with LIORESAL INTRATHECAL (baclofen injection) obtained in parallel, placebo-controlled, randomized studies provides only a limited basis for estimating the incidence of adverse events because the studies were of very brief duration (up to three days of infusion) and involved only a total of 63 patients. The following events occurred among the 31 patients receiving LIORESAL INTRATHECAL (baclofen injection) in two randomized, placebo-controlled trials: hypotension (2), dizziness (2), headache (2), dyspnea (1). No adverse events were reported among the 32 patients receiving placebo in these studies.

Events Observed during the Pre- and Post-marketing Evaluation of LIORESAL INTRATHECAL — Adverse events associated with the use of LIORESAL INTRATHECAL reflect experience gained with 576 patients followed prospectively in the United States. They received LIORESAL INTRATHECAL for periods of one day (screening) (N = 576) to over eight years (maintenance) (N = 10). The usual screening bolus dose administered prior to pump implantation in these studies was typically 50 mcg. The maintenance dose ranged from 12 mcg to 2003 mcg per day. Because of the open, uncontrolled nature of the experience, a causal linkage between events observed and the administration of LIORESAL INTRATHECAL cannot be reliably assessed in many cases and many of the adverse events reported are known to occur in association with the underlying conditions being treated. Nonetheless, many of the more commonly reported reactions—hypotonia, somnolence, dizziness, paresthesia, nausea/vomiting and headache—appear clearly drug-related.

Adverse experiences reported during all U.S. studies (both controlled and uncontrolled) are shown in the following table. Eight of 474 patients who received chronic infusion via implanted pumps had adverse experiences which led to a discontinuation of long term treatment in the pre- and post-marketing studies.

INCIDENCE OF MOST FREQUENT (≥1%) ADVERSE EVENTS IN PATIENTS WITH SPASTICITY OF SPINAL ORIGIN IN PROSPECTIVELY MONITORED CLINICAL TRIALS

Adverse Event	Percent of Patients Reporting Events		
	N = 576	N = 474	N = 430
	Screening ^a	Titration ^b	Maintenance ^c
	Percent	Percent	Percent
Hypotonia	5.4	13.5	25.3
Somnolence	5.7	5.9	20.9
Dizziness	1.7	1.9	7.9
Paresthesia	2.4	2.1	6.7
Nausea and Vomiting	1.6	2.3	5.6
Headache	1.6	2.5	5.1
Constipation	0.2	1.5	5.1
Convulsion	0.5	1.3	4.7
Urinary Retention	0.7	1.7	1.9
Dry Mouth	0.2	0.4	3.3
Accidental Injury	0.0	0.2	3.5
Asthma	0.7	1.3	1.4
Confusion	0.5	0.6	2.3
Death	0.2	0.4	3.0
Pain	0.0	0.6	3.0
Speech Disorder	0.0	0.2	3.5
Hypotension	1.0	0.2	1.9
Ambylopia	0.5	0.2	2.3
Diarrhea	0.0	0.8	2.3
Hypoventilation	0.2	0.8	2.1
Coma	0.0	1.5	0.9
Impotence	0.2	0.4	1.6
Peripheral Edema	0.0	0.0	2.3
Urinary Incontinence	0.0	0.8	1.4
Insomnia	0.0	0.4	1.6
Anxiety	0.2	0.4	0.9
Depression	0.0	0.0	1.6
Dyspnea	0.3	0.0	1.2
Fever	0.5	0.2	0.7
Pneumonia	0.2	0.2	1.2
Urinary Frequency	0.0	0.6	0.9
Urticaria	0.2	0.2	1.2
Anorexia	0.0	0.4	0.9
Diplopia	0.0	0.4	0.9
Dysautonomia	0.2	0.2	0.9
Hallucinations	0.3	0.4	0.5
Hypertension	0.2	0.6	0.5

^a Following administration of test bolus

^b Two month period following implant

^c Beyond two months following implant

N= total number of patients entering each period

%=% of patients evaluated

In addition to the more common (1% or more) adverse events reported in the prospectively followed 576 domestic patients in pre- and post- marketing studies, experience from an additional 194 patients exposed to LIORESAL INTRATHECAL (baclofen injection) from foreign studies has been reported. The following adverse events, not described in the table, and arranged in decreasing order of frequency, and classified by body system, were reported:

Nervous System: Abnormal gait, thinking abnormal, tremor, amnesia, twitching, vasodilatation, cerebrovascular accident, nystagmus, personality disorder, psychotic depression, cerebral ischemia, emotional lability, euphoria, hypertension, ileus, drug dependence, incoordination, paranoid reaction and ptosis.

Digestive System: Flatulence, dysphagia, dyspepsia and gastroenteritis.

Cardiovascular: Postural hypotension, bradycardia, palpitations, syncope, arrhythmia ventricular, deep thrombophlebitis, pallor and tachycardia.

Respiratory: Respiratory disorder, aspiration pneumonia, hyperventilation, pulmonary embolus and rhinitis.

Urogenital: Hematuria and kidney failure.

Skin and Appendages: Alopecia and sweating.

Metabolic and Nutritional Disorders: Weight loss, albuminuria, dehydration and hyperglycemia.

Special Senses: Abnormal vision, abnormality of accommodation, photophobia, taste loss and tinnitus.

Body as a Whole: Suicide, lack of drug effect, abdominal pain, hypothermia, neck rigidity, chest pain, chills, face edema, flu syndrome and overdose.

Hemic and Lymphatic System: Anemia.

Spasticity of Cerebral Origin:

Commonly Observed — In pre- marketing clinical trials, the most commonly observed adverse events associated with use of LIORESAL INTRATHECAL (baclofen injection) which were not seen at an equivalent incidence among placebo-treated patients included: agitation, constipation, somnolence, leukocytosis, chills, urinary retention and hypotonia.

Associated with Discontinuation of Treatment — Nine of 211 patients receiving LIORESAL INTRATHECAL in pre-marketing clinical studies in the U.S. discontinued long term infusion due to adverse events associated with intrathecal therapy.

The nine adverse events leading to discontinuation were: infection (3), CSF leaks (2), meningitis (2), drainage (1), and unmanageable trunk control (1).

Fatalities — Three deaths, none of which were attributed to LIORESAL INTRATHECAL, were reported in patients in clinical trials involving patients with spasticity of cerebral origin. See Warnings on other deaths reported in spinal spasticity patients.

Incidence in Controlled Trials — Experience with LIORESAL INTRATHECAL (baclofen injection) obtained in parallel, placebo-controlled, randomized studies provides only a limited basis for estimating the incidence of adverse events because the studies involved a total of 62 patients exposed to a single 50 mcg intrathecal bolus. The following events occurred among the 62 patients receiving LIORESAL INTRATHECAL in two randomized, placebo-controlled trials involving cerebral palsy and head injury patients, respectively: agitation, constipation, somnolence, leukocytosis, nausea, vomiting, nystagmus, chills, urinary retention, and hypotonia.

Events Observed during the Pre- marketing Evaluation of LIORESAL INTRATHECAL — Adverse events associated with the use of LIORESAL INTRATHECAL reflect experience gained with a total of 211 U.S. patients with spasticity of cerebral origin, of whom 112 were pediatric patients (under age 16 at enrollment). They received LIORESAL INTRATHECAL for periods of one day (screening) (N= 211) to 84 months (maintenance) (N= 1). The usual screening bolus dose administered prior to pump implantation in these studies was 50- 75 mcg. The maintenance dose ranged from 22 mcg to 1400 mcg per day. Doses used in this patient population for long term infusion are generally lower than those required for patients with spasticity of spinal cord origin.

Because of the open, uncontrolled nature of the experience, a causal linkage between events observed and the administration of LIORESAL INTRATHECAL cannot be reliably assessed in many cases. Nonetheless, many of the more commonly reported reactions— somnolence, dizziness, headache, nausea, hypotension, hypotonia and coma— appear clearly drug-related.

The most frequent (≥1%) adverse events reported during all clinical trials are shown in the following table. Nine patients discontinued long term treatment due to adverse events.

INCIDENCE OF MOST FREQUENT (≥ 1%) ADVERSE EVENTS IN PATIENTS WITH SPASTICITY OF CEREBRAL ORIGIN IN PROSPECTIVELY MONITORED CLINICAL TRIALS

Adverse Event	Percent of Patients Reporting Events		
	N = 211	N = 153	N = 150
	Screening ^a Percent	Titration ^b Percent	Maintenance ^c Percent
Hypotonia	2.4	14.4	34.7
Somnolence	7.6	10.5	18.7
Headache	6.6	7.8	10.7
Nausea and Vomiting	6.6	10.5	4.0
Vomiting	6.2	8.5	4.0
Urinary Retention	0.9	6.5	8.0
Convulsion	0.9	3.3	10.0
Dizziness	2.4	2.6	8.0
Nausea	1.4	3.3	7.3
Hypoventilation	1.4	1.3	4.0
Hypertonia	0.0	0.7	6.0
Paresthesia	1.9	0.7	3.3
Hypotension	1.9	0.7	2.0
Increased Salivation	0.0	2.6	2.7
Back Pain	0.9	0.7	2.0
Constipation	0.5	1.3	2.0
Pain	0.0	0.0	4.0
Pruritus	0.0	0.0	4.0
Diarrhea	0.5	0.7	2.0
Peripheral Edema	0.0	0.0	3.3
Thinking Abnormal	0.5	1.3	0.7
Agitation	0.5	0.0	1.3
Asthenia	0.0	0.0	2.0
Chills	0.5	0.0	1.3
Coma	0.5	0.0	1.3
Dry Mouth	0.5	0.0	1.3
Pneumonia	0.0	0.0	2.0
Speech Disorder	0.5	0.7	0.7
Tremor	0.5	0.0	1.3
Urinary Incontinence	0.0	0.0	2.0
Urination Impaired	0.0	0.0	2.0

^a Following administration of test bolus

^b Two month period following implant

^c Beyond two months following implant

N= Total number of patients entering each period. 211 patients received drug; (1 of 212) received placebo only.

The more common (1% or more) adverse events reported in the prospectively followed 211 patients exposed to LIORESAL INTRATHECAL (baclofen injection) have been reported. In the total cohort, the following adverse events, not described in the table, and arranged in decreasing order of frequency, and classified by body system, were reported:

Nervous System: Akathisia, ataxia, confusion, depression, opisthotonos, amnesia, anxiety, hallucinations, hysteria, insomnia, nystagmus, personality disorder, reflexes decreased, and vasodilatation.

Digestive System: Dysphagia, fecal incontinence, gastrointestinal hemorrhage and tongue disorder.

Cardiovascular: Bradycardia.

Respiratory: Apnea, dyspnea and hyperventilation.

Urogenital: Abnormal ejaculation, kidney calculus, oliguria and vaginitis.

Skin and Appendages: Rash, sweating, alopecia, contact dermatitis and skin ulcer.

Special Senses: Abnormality of accommodation.

Body as a Whole: Death, fever, abdominal pain, carcinoma, malaise and hypothermia.

Hemic and Lymphatic System: Leukocytosis and petechial rash.

DRUG OVERDOSE

Special attention must be given to recognizing the signs and symptoms of overdosage, especially during the initial screening and dose- titration phase of treatment, but also during reintroduction of LIORESAL INTRATHECAL after a period of interruption in therapy.

Symptoms of LIORESAL INTRATHECAL Overdose: Drowsiness, lightheadedness, dizziness, somnolence, respiratory depression, seizures, rostral progression of hypotonia and loss of consciousness progressing to coma of up to 72 hr. duration. In most cases reported, coma was reversible without sequelae after drug was discontinued. Symptoms of LIORESAL INTRATHECAL overdose were reported in a sensitive adult patient after receiving a 25 mcg intrathecal bolus.

Treatment Suggestions for Overdose:

There is no specific antidote for treating overdoses of LIORESAL INTRATHECAL (baclofen injection); however, the following steps should ordinarily be undertaken:

- 1) Residual LIORESAL INTRATHECAL solution should be removed from the pump as soon as possible.
- 2) Patients with respiratory depression should be intubated if necessary, until the drug is eliminated.

Anecdotal reports suggest that intravenous physostigmine may reverse central side effects, notably drowsiness and respiratory depression. Caution in administering physostigmine is advised, however, because its use has been associated with the induction of seizures and bradycardia.

Physostigmine Doses for Adult Patients: Administer 2 mg of physostigmine intramuscularly or intravenously at a slow controlled rate of no more than 1 mg per minute. Dosage may be repeated if life- threatening signs, such as arrhythmia, convulsions or coma occur.

Physostigmine Doses for Pediatric Patients: Administer 0.02 mg/ kg physostigmine intramuscularly or intravenously, do not give more than 0.5 mg per minute. The dosage may be repeated at 5 to 10 minute intervals until a therapeutic effect is obtained or a maximum dose of 2 mg is attained.

Physostigmine may not be effective in reversing large overdoses and patients may need to be maintained with respiratory support.

If lumbar puncture is not contraindicated, consideration should be given to withdrawing 30- 40 ml of CSF to reduce CSF baclofen concentration.

DOSAGE AND ADMINISTRATION

Refer to the manufacturer's manual for the implantable pump approved for intrathecal infusion for specific instructions and precautions for programming the pump and/ or refilling the reservoir. There are various pumps with varying reservoir volumes and there are various refill kits available. It is important to be familiar with all of these products in order to select the appropriate refill kit for the particular pump in use.

Screening Phase: Prior to pump implantation and initiation of chronic infusion of LIORESAL INTRATHECAL (baclofen injection), patients must demonstrate a positive clinical response to a LIORESAL INTRATHECAL bolus dose administered intrathecally in a screening trial. The screening trial employs LIORESAL INTRATHECAL at a concentration of 50 mcg/ ml. A 1 ml ampule (50 mcg/ ml) is available for use in the screening trial. The screening procedure is as follows. An initial bolus containing 50 micrograms in a volume of 1 milliliter is administered into the intrathecal space by barbotage over a period of not less than one minute. The patient is observed over the ensuing 4 to 8 hours. A positive response consists of a significant decrease in muscle tone and/ or frequency and/ or severity of spasms. If the initial response is less than desired, a second bolus injection may be administered 24 hours after the first. The second screening bolus dose consists of 75 micrograms in 1.5 milliliters. Again, the patient should be observed for an interval of 4 to 8 hours. If the response is still inadequate, a final bolus screening dose of 100 micrograms in 2 milliliters may be administered 24 hours later.

Pediatric Patients: The starting screening dose for pediatric patients is the same as in adult patients, i.e., 50 mcg. However, for very small patients, a screening dose of 25 mcg may be tried first. **Patients who do not respond to a 100 mcg intrathecal bolus should not be considered candidates for an implanted pump for chronic infusion.**

Post- Implant Dose Titration Period: To determine the initial total daily dose of LIORESAL INTRATHECAL following implant, the screening dose that gave a positive effect should be doubled and administered over a 24-hour period, unless the efficacy of the bolus dose was maintained for more than 8 hours, in which case the starting daily dose should be the screening dose delivered over a 24-hour period. No dose increases should be given in the first 24 hours (i.e., until the steady state is achieved).

Adult Patients with Spasticity of Spinal Cord Origin: After the first 24 hours, for adult patients, the daily dosage should be increased slowly by 10- 30% increments and only once every 24 hours, until the desired clinical effect is achieved.

Adult Patients with Spasticity of Cerebral Origin: After the first 24 hours, the daily dose should be increased slowly by 5- 15% only once every 24 hours, until the desired clinical effect is achieved.

Pediatric Patients: After the first 24 hours, the daily dose should be increased slowly by 5-15% only once every 24 hours, until the desired clinical effect is achieved. If there is not a substantive clinical response to increases in the daily dose, check for proper pump function and catheter patency. Patients must be monitored closely in a fully equipped and staffed environment during the screening phase and dose- titration period immediately following implant. Resuscitative equipment should be immediately available for use in case of life- threatening or intolerable side effects.

Maintenance Therapy:

Spasticity of Spinal Cord Origin Patients: The clinical goal is to maintain muscle tone as close to normal as possible, and to minimize the frequency and severity of spasms to the extent possible, without inducing intolerable side effects. Very often, the maintenance dose needs to be adjusted during the first few months of therapy while patients adjust to changes in life style due to the alleviation of spasticity. During periodic refills of the pump, the daily dose may be increased by 10-40%, but no more than 40%, to maintain adequate symptom control. The daily dose may be reduced by 10-20% if patients experience side effects. Most patients require gradual increases in dose over time to maintain optimal response during chronic therapy. A sudden large requirement for dose escalation suggests a catheter complication (i.e., catheter kink or dislodgement).

Neuromodulation Products

Maintenance dosage for long term continuous infusion of LIORESAL INTRATHECAL (baclofen injection) has ranged from 12 mcg/ day to 2003 mcg/ day, with most patients adequately maintained on 300 micrograms to 800 micrograms per day. There is limited experience with daily doses greater than 1000 mcg/ day. Determination of the optimal LIORESAL INTRATHECAL dose requires individual titration. The lowest dose with an optimal response should be used.

Spasticity of Cerebral Origin Patients: The clinical goal is to maintain muscle tone as close to normal as possible and to minimize the frequency and severity of spasms to the extent possible, without inducing intolerable side effects, or to titrate the dose to the desired degree of muscle tone for optimal functions. Very often the maintenance dose needs to be adjusted during the first few months of therapy while patients adjust to changes in life style due to the alleviation of spasticity. During periodic refills of the pump, the daily dose may be increased by 5 - 20%, but no more than 20%, to maintain adequate symptom control. The daily dose may be reduced by 10-20% if patients experience side effects. Many patients require gradual increases in dose over time to maintain optimal response during chronic therapy. A sudden large requirement for dose escalation suggests a catheter complication (i.e., catheter kink or dislodgement).

Maintenance dosage for long term continuous infusion of LIORESAL INTRATHECAL (baclofen injection) has ranged from 22 mcg/ day to 1400 mcg/ day, with most patients adequately maintained on 90 micrograms to 703 micrograms per day. In clinical trials, only 3 of 150 patients required daily doses greater than 1000 mcg/ day.

Pediatric Patients: Use same dosing recommendations for patients with spasticity of cerebral origin. Pediatric patients under 12 years seemed to require a lower daily dose in clinical trials. Average daily dose for patients under 12 years was 274 mcg/ day, with a range of 24 to 1199 mcg/ day. Dosage requirement for pediatric patients over 12 years does not seem to be different from that of adult patients. Determination of the optimal LIORESAL INTRATHECAL dose requires individual titration. The lowest dose with an optimal response should be used.

Potential need for dose adjustments in chronic use : During long term treatment, approximately 5% (28/627) of patients become refractory to increasing doses. There is not sufficient experience to make firm recommendations for tolerance treatment; however, this "tolerance" has been treated on occasion, in hospital, by a "drug holiday" consisting of the gradual reduction of LIORESAL INTRATHECAL over a 2 to 4 week period and switching to alternative methods of spasticity management. After the "drug holiday," LIORESAL INTRATHECAL may be restarted at the initial continuous infusion dose.

Stability

Parenteral drug products should be inspected for particulate matter and discoloration prior to administration, whenever solution and container permit.

Delivery Specifications

The specific concentration that should be used depends upon the total daily dose required as well as the delivery rate of the pump. LIORESAL INTRATHECAL may require dilution when used with certain implantable pumps. Please consult manufacturer's manual for specific recommendations.

Preparation Instruction:

Screening

Use the 1 ml screening ampule only (50 mcg/ml) for bolus injection into the subarachnoid space. For a 50mcg bolus dose, use 1 ml of the screening ampule. Use 1.5 ml of 50 mcg/ml baclofen injection for a 75 mcg bolus dose. For the maximum screening dose of 100 mcg, use 2 ml of 50 mcg/ml baclofen injection (2 screening ampules).

Maintenance

For patients who require concentrations other than 500 mcg/ml or 2000 mcg/ml, LIORESAL INTRATHECAL **must be diluted**.

LIORESAL INTRATHECAL **must be diluted** with sterile preservative free Sodium Chloride for Injection, U.S.P.

Delivery Regimen:

LIORESAL INTRATHECAL is most often administered in a continuous infusion mode immediately following implant. For those patients implanted with programmable pumps who have achieved relatively satisfactory control on continuous infusion, further benefit may be attained using more complex schedules of LIORESAL INTRATHECAL delivery. For example, patients who have increased spasms at night may require a 20% increase in their hourly infusion rate. Changes in flow rate should be programmed to start two hours before the time of desired clinical effect.

HOW SUPPLIED

LIORESAL INTRATHECAL (baclofen injection) is available in single use ampules of 10 mg/20 ml (500 mcg/ ml) or 10 mg/ 5 ml (2000 mcg/ml) or 40 mg/20 ml (2000 mcg/ml) packaged in a Refill Kit for intrathecal administration. For screening, LIORESAL INTRATHECAL is available in a single use ampule of 0.05 mg/ 1 ml.

Model 8561 LIORESAL INTRATHECAL Refill Kit contains one ampule of 10 mg/ 20 ml (500 mcg/ml) (NDC 58281-560-01).

Model 8562 LIORESAL INTRATHECAL Refill Kit contains two ampules of 10 mg/ 5 ml (2000 mcg/ml) (NDC 58281-561-02).

Model 8563s LIORESAL INTRATHECAL contains one ampule of 0.05 mg/ 1 ml (NDC 58281-562-01).

Model 8564 LIORESAL INTRATHECAL Refill Kit contains four ampules of 10 mg/ 5 ml (2000 mcg/ml) (NDC 58281-561-04) or one ampule of 40 mg/20 ml (2000 mcg/ml) (NDC 58281-563-01).

Model 8565 LIORESAL INTRATHECAL Refill Kit contains two ampules of 10 mg/ 20 ml (500 mcg/ml) (NDC 58281-560-02).

Model 8566 LIORESAL INTRATHECAL Refill Kit contains eight ampules of 10 mg/ 5 ml (2000 mcg/ml) (NDC 58281-561-08) or two ampules of 40 mg/20 ml (2000 mcg/ml) (NDC 58281-563-02).

STORAGE

- Does not require refrigeration.
- Do not store above 86° F (30° C).
- Do not freeze.
- Do not heat sterilize.

Manufactured by Novartis Pharma Stein AG, Stein, Switzerland, for Medtronic, Inc., Minneapolis, Minnesota 55432- 5604 USA.

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