

Biomet® EBI® Bone Healing System SFLX Coilette Therapeutic Treatment Coil



When you see these symbols, immediately refer to the full prescribing information described within the User Manual and Packaging Insert supplied with your non-invasive stimulator for more information. Please read the following helpful information listed below. This information will enable you to utilize this medical device safely and effectively. Retain this information for later use. This medical device is **ONLY** intended to be used with the non-invasive bone growth stimulator which was prescribed by your physician for your treatment.



INDICATIONS FOR USE

The Biomet® EBI® Bone Healing System is indicated for the treatment of fracture nonunions, failed fusions, and congenital pseudarthrosis in the appendicular system. A nonunion is considered to be established when there are no visibly progressive signs of healing.

USAGE

Follow the treatment schedule prescribed by your physician, normally ten (10) hours per day. Your compliance with the recommended ten (10) hours per day treatment is very important. A review of the original premarket clinical data demonstrated that less than the recommended use of this device possibly results in an increase in the time to heal your fracture nonunion. If you are unable to treat for ten continuous hours, it is recommended that you break up the total treatment time into more than one session. Please consult the general treatment instructions section contained within the User Manual and Packaging Insert supplied with your non-invasive stimulator for more information before beginning treatment.



ATTENTION

This medical device is a durable therapeutic electrical device intended for single patient use only under a prescription. Federal Law (U.S.A.) restricts this device to sale by or on the order of a physician. Treatment at home or in another appropriate or similar setting is acceptable. This medical device cannot be reprocessed, refurbished, disinfected, reused, sterilized, etc. with the intent to be used by another patient or for treatment other than prescribed.

CONTRAINDICATIONS

A. Nonunion fractures in which a synovial pseudarthrosis (fluid filled gap) exists.



B. Under certain conditions, electromagnetic stimulation could inhibit or impair the functioning of certain external, non-invasive and/or implanted, invasive active medical devices inclusive of "all active electrical and non-active conductive/metallic implants" as well as "worn medical devices" due to adverse events that may occur with other active electrical implants (e.g., Spinal Cord stimulators, Implantable Cardioverter-defibrillators, etc.) The impact or effect of pulsed electromagnetic fields generated by a non invasive bone growth stimulator on the function of other anatomical stimulators, pain pumps, insulin pumps, implanted spinal nerve stimulators and similar active devices has not been evaluated.



C. Use of the Bone Healing System on pregnant patients has not been evaluated; therefore, it is not recommended in these cases.



D. The Bone Healing System has not been tested for safety or been evaluated for heating in the MR environment. The effects of MRI procedures and scans using MR systems has not been determined or established; therefore, MRI scans and procedures should not be performed on patients until the device system has been completely removed. MR Unsafe-Not for MRI Use.

WARNINGS

- A. The long term effects of exposure to low level magnetic fields are not known. Routine use of these bone healing systems for over 30 years has indicated no known risks.
- B. During the treatment of patients with open epiphyses, when the epiphysis is in the pulsing field, physicians are advised that the epiphyseal growth plates should be monitored for possible effects.
- C. Use of the Bone Healing System for the spine and skull have not been evaluated.
- D. To reduce the risk of potential injury:
 - 1. AVOID touching the AC Adapter contacts when the AC Adapter is plugged into an AC wall outlet.
 - 2. DO NOT charge the battery in bed if treating while sleeping.
- E. The control unit is electrically live when connected with the AC Wall Adapter and plugged into an AC Wall outlet. To reduce the risk of serious injury by electric shock, patients are advised:
 - 1. DO NOT permit the AC Adapter to be connected when wet.
 - 2. DO NOT immerse the control unit, treatment coil, or the AC Wall Adapter in water or any liquid.
- F. No unauthorized modification of this device is allowed for any reason whatsoever.

PRECAUTIONS

The following conditions may compromise a successful treatment outcome.

- A. Nonunion fractures with gaps in excess of 1.0cm.
- B. Presence of fixation devices or instrumentation made from magnetic materials.

PLEASE NOTE: Most presently used internal or external fixation devices are constructed of 316L S.S., titanium alloys, and cobalt-chromium alloys which are non-magnetic and, therefore, compatible with the Bone Healing System.

Adverse Effects

The Bone Healing System was FDA approved in 1979. Since then, more than 450,000 systems have been commercially distributed and prescribed to patients. Based on the results of an exhaustive historical search of the MAUDE and MDR Databases, the probability of an adverse event was extremely unlikely (.0062%).

The identified hazards associated with the use of non-invasive bone growth stimulation devices are comprehensive, well-known, understood and continue to clearly establish the benefits significantly outweighing the risks.

CONTENTS

- SFLX Therapeutic Treatment Coil
- Flexion Gauge
- User, Safety & Application Instructions



Do not dispose of this device with household waste

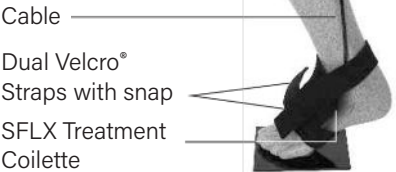
Rx Only - Prescription Only - Single Use Only
- Not for Re-Sale or Re-Distribution
- Do Not Reuse.

Flexion Gauge Instructions For The SFLX Treatment Coilette
Application Instructions For: SFLX Treatment Coilette Applies To:

Description	Treatment Coil	Suggested Placement
SFLX Coilette	1068238	Metatarsals, Scaphoid, Distal Radius, Cuboid, M-L Malleous

Application Instructions For The SFLX Treatment Coilette

Foot Application



1. Attach the SFLX Treatment Coilette to the strap by connecting it to the Velcro® hook provided on the strap.
2. Center the SFLX Treatment Coilette over the fracture site.
3. Wrap lower strap around bottom of foot and attach to Velcro® hook.
4. Wrap top strap around ankle and cable, and attach to Velcro® hook.
5. Adjust strap for comfort. Excess strap length may then be cut away. When treatment is completed for the day, remove the SFLX Treatment Coilette by loosening the straps. Leave the straps attached to the SFLX Treatment Coilette. It is ready for your next daily treatment.



Hand Application



1. Shape and place the SFLX Treatment Coilette over the treatment site as instructed by your physician or sales representative.
2. Place the SFLX Treatment Coilette onto the Velcro® hook on the strap.
3. Place the SFLX Treatment Coilette over treatment site and bring straps around each side of thumb.
4. Secure Velcro® straps onto the Velcro® hook. Adjust for comfort.
5. Excess strap length may then be cut away.
6. When treatment is completed for the day, remove the SFLX Treatment Coilette by loosening the straps. Leave the straps attached. It is ready for your next daily treatment.

NOTE: During treatment, the position of the treatment coil may shift due to normal patient activity. Often this movement is associated with patient activity, mobility or the underlying surface the treatment coil rests on (skin, shirt, cast, etc). The treatment coil may be loosened and may need to be repositioned back to a comfortable location before tightening the straps, completing the adjustment.

SFLX Treatment Coilette Specifications and Gauge Directions

SFLX Coilette Shape	Depth of Penetration	Vertical Fracture Length
Flat	2.75 cm	4.0 cm
Elliptical	3.5 cm	4.0 cm
Saddle	3.5 cm	2.0 cm

To insure proper fit and efficacious treatment with the SFLX Treatment Coilette:

1. Place the SFLX Treatment Coilette over the treatment site and shape for best fit. The treatment coilette must only be bent in one of two directions (long axis or short axis). Do not kink or twist the treatment coilette.
2. The shape of the treatment coilette will determine the specifications (see above).
3. Remove the treatment coilette and utilize the Flexion Gauge to verify the shape listed below.



Flat Shape (see coil specifications above)
The Flexion Gauge is not needed.



Elliptical Shape (see coil specifications above) Place the treatment coilette over the green portion of the gauge marked "elliptical." The Coilette bend should not exceed the elliptical curve indicated on the Flexion Gauge.

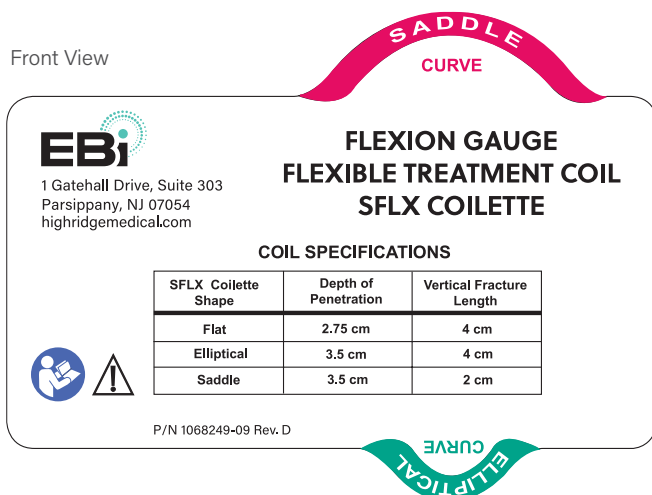


Saddle Shape (see specifications above) Place the treatment coilette over the red portion of the gauge marked "saddle." The treatment coilette bend should not exceed the saddle curve on the Flexion Gauge.

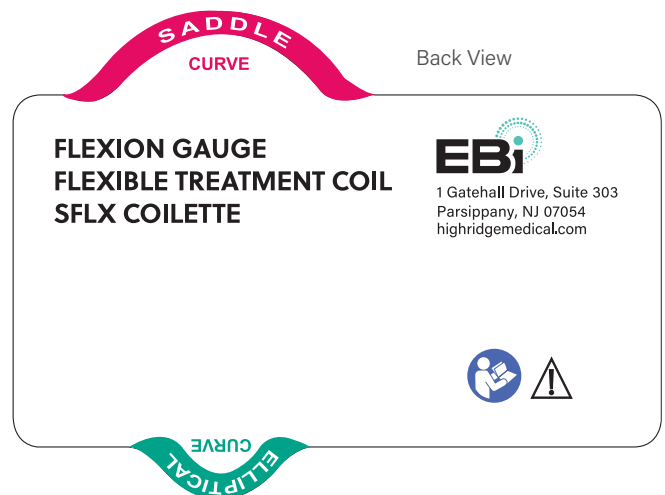
In order to ensure proper fit and efficacious treatment, the Flexion Gauge should be employed to check for the proper shape.

1. Place the treatment coilette at the treatment site and shape for best fit. Treatment coilette must only be bent in one of two directions: Long or short axis. Do not kink or twist the treatment coilette. The shape of the treatment coilette will determine the specifications.
2. Remove the shaped treatment coilette and utilize the Flexion Gauge to verify the acceptable bend according to the shape listed below.
3. If the treatment coilette shape does not fit the curve on the Flexion Gauge, contact an EBI® representative for a suitable replacement and assistance.

Front View



Back View



Not to Scale

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**Legal Manufacturer**

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Complete prescribing information including full indications, contraindications, warnings and precautions associated with the use of these devices may be found online at highridgemedical.com or by calling 800-526-2579. The Biomet® EBI® Bone Healing System is indicated for the treatment of fracture nonunions, failed fusions, and congenital pseudarthrosis in the appendicular system. Contraindicated for nonunion fractures in which a synovial pseudarthrosis exists. Electromagnetic PEMF stimulation is contraindicated for use by patients with implantable pacemakers or defibrillators. The Bone Healing System is not MR safe. Federal Law (U.S.A.) restricts this device to sale by or on the order of a physician. Rx Only. Single Patient Use Only. Do Not Reuse. All content herein is protected by copyright, trademarks and other intellectual property rights, as applicable, owned by or licensed to Highridge Medical or its affiliates unless otherwise indicated, and must not be redistributed, duplicated or disclosed, in whole or in part, without the express written consent of Highridge Medical. P/N 1068318-08 REV D. HM0696 REV A 04/25. ©2025 EBI, LLC. All rights reserved.