exogen®
ultrasound bone healing system

Recycling Programme

Your used EXOGEN devices can be collected for recycling, an important step forward in our commitment to environmental responsibility.

The EXOGEN recycling programme means that, as a device reaches the end of its useful life, it can be processed for material recovery. We are proud to offer a programme that supports both your clinical needs and environmental goals.

Processed with care in our partner UK facility

European Recycling Platform
A Landbell Group Company

The EXOGEN recycling programme is administered through our partner, European Recycling Platform UK Ltd, who transfers the product to an Approved Authorised Treatment Facility (AATF).

The strict licensing standards for becoming an AATF help ensure that our products are recycled safely and responsibly. By keeping the programme within the UK, we also reduce transportation emissions and support local industry.

exogen® Recycling Programme

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Instructions for Patients:

1. Take your EXOGEN device to each of your fracture care follow-up appointments. This will help your care team monitor your progress and ensure the device is functioning correctly.
2. Aim for a 100% treatment adherence rate. Your EXOGEN device will record this, providing valuable information for your care team.
3. Once you have completed your prescribed EXOGEN treatments, return the device to the clinic you received it from. This will ensure that your device is collected for recycling.

If you have purchased the device yourself, contact your local Bioventus representative or Bioventus Customer Care at CustomerCare-International@Bioventus.com to arrange for device collection.

Instructions for Clinics:

1. To participate in the programme, inform your local Bioventus representative. A device collection barrel will be provided to you free of charge.
2. When a patient returns an EXOGEN device, note the adherence rate and treatment outcome for audit purposes. Deposit the used device in your collection barrel. If the device is eligible for the EXOGEN Performance Programme, keep the device separately and contact your local Bioventus representative.
3. When your collection barrel is full, contact your local Bioventus representative to arrange a collection. A replacement barrel will be provided to you at time of collection.

By following these steps, you are supporting excellence of care and our commitment to environmental responsibility. Thank you for your participation!

Summary of Indications for Use: EXOGEN is indicated for the non-invasive treatment of osseous defects (excluding vertebra and skull) that includes the treatment of delayed unions, nonunions,* stress fractures and joint fusion. EXOGEN is also indicated for the acceleration of fresh fracture heal time, repair following osteotomy, repair in bone transport procedures and repair in distraction osteogenesis procedures.

There are no known contraindications for the EXOGEN device. Safety and effectiveness have not been established for individuals lacking skeletal maturity, pregnant or nursing women, patients with cardiac pacemakers, on fractures due to bone cancer, or on patients with poor blood circulation or clotting problems. Some patients may be sensitive to the ultrasound gel.

Full prescribing information can be found in product labeling, at EXOGEN.com/wp-content/uploads/2024/07/81087028_Sonic-2020-IFU_OUS_RevF.pdf, or by calling Bioventus Customer Service.

*A nonunion is considered to be established when the fracture site shows no visibly progressive signs of healing.

bioventus®
Innovations For Active Healing