

Ready for a Full-Body MRI?

Axonics™ therapy is safe for an MRI within defined conditions. This document contains some of the key information you and your healthcare professional will need when you need a full-body MRI.

What you need to do to prepare for an MRI:

- Make sure your neurostimulator is charged until the battery status light at the bottom of your remote control is green
- Bring this sheet, your patient ID card, and your remote control with you to your MRI appointment



Information for the Healthcare Professional

Patient Name: _____

Doctor Name: _____

Office Address: _____

Phone: _____

Date form completed: _____

Is the Axonics Lead located in the sacral foramen?

☐ No ☐ Yes

Is the Axonics neurostimulator located in the posterior hip/upper buttock area?

☐ No ☐ Yes

Any implanted leads NOT connected to a functioning Axonics stimulator (abandoned leads)

☐ No ☐ Yes (please list) _____

Any implants in addition to the Axonics™ SNM system?

☐ No ☐ Yes (please list) _____

* See MRI guidelines for MRI eligibility process for remote controls manufactured prior to May 2020.

Using the Patient Remote Control to Check MRI Readiness*



- 1 **Press the centre button** to connect to the stimulator. After connecting, **hold the down arrow** for 5 or more seconds to turn off stimulation (the 7 stimulation level lights at the top of the remote control are off).



- 2 When stimulation turns off, check the battery status light on the very bottom is green (if the bottom light is not green, charge your neurostimulator). **Hold the down arrow** for 5 or more seconds (release button, when programme lights 1 and 2, flash) to check MRI readiness.



- 3 When the middle 3 level lights at the top of the remote turn on, **you are ready** for an MRI. These lights will be seen for 15 seconds after which the remote control will automatically turn off.



NOTE: If a solid red light is seen, you are not eligible to have a full-body MRI, and you should contact your doctor.

Important Safety Information

Axonics SNM Therapy for urinary control is indicated for the treatment of urinary retention and the symptoms of overactive bladder, including urinary urge incontinence and significant symptoms of urgency-frequency alone or in combination, in patients who have failed or could not tolerate more conservative treatments. Axonics SNM Therapy for bowel control is indicated for the treatment of chronic faecal incontinence in patients who have failed or are not candidates for more conservative treatments.

Axonics SNM Therapy is contraindicated for patients who have not demonstrated an appropriate response to test stimulation or patients who are unable to operate the Axonics SNM Systems. With all medical procedures, there are risks associated with the procedure and the use of the device, including longterm implantation. The risks include but are not limited to: adverse change in voiding function (bowel and/or bladder), infection, pain or irritation at the implant site, lead or device migration, electrical shock, change in sensation or magnitude of stimulation which has been described as uncomfortable (jolting or shocking) by some patients, and heating or burns at the device site.

If one or more of these complications occur, you may need medical treatment or surgery. Be sure to talk with your doctor so that you thoroughly understand all of the risks and benefits associated with the use of the device.

CAUTION: The law restricts these devices to sale by or on the order of a physician. Indications, contraindications, warnings, and instructions for use can be found in the product labelling supplied with each device. Products shown for INFORMATION purposes only and may not be approved or for sale in certain countries. This material not intended for use in France.

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