Axonics SNM System® Remote Control

Remote Control User Manual



Axonics SNM System® Remote Control





Axonics®, Axonics Modulation®, Axonics Modulation Technologies®, Axonics Sacral Neuromodulation System® and r-SNM® are trademarks of Axonics Modulation Technologies, Inc., registered or pending registration in the U.S. and other countries.

Caution: The Axonics Sacral Neuromodulation System should be used only on the order of a doctor

Contents

Introduction	5
Manual overview	6
Refer to your Patient Therapy Guide	Error! Bookmark not defined.
Remote Control precautions	Error! Bookmark not defined.
Recommended use and care	Error! Bookmark not defined.
How to use your Remote Control	Error! Bookmark not defined.
Summary of buttons and lights	Error! Bookmark not defined.
How your Remote Control communicates with your Stimulator	Error! Bookmark not defined.
1) Connect to your Stimulator	Error! Bookmark not defined.
2) View Stimulator Status	Error! Bookmark not defined.
3) Turn Down your stimulator level	Error! Bookmark not defined.
4) Turn up your stimulator level	Error! Bookmark not defined.
5) Turn your stimulation off	
6) Turn your stimulation on	Error! Bookmark not defined.
7) Monitor charge status – Permanent Stimulator only	Error! Bookmark not defined.
8) Switch the active program – <i>Trial Stimulator only</i>	Error! Bookmark not defined.
9) See a system error	Error! Bookmark not defined.
10) Turn off your Remote Control	
Troubleshooting	
Additional Information	
Replacement and disposal	Error! Bookmark not defined.
Specifications	
Axonics contact information	9
Label Symbols	10

Introduction

The Axonics Sacral Neuromodulation (SNM) therapy is provided to relieve the symptoms of urinary fecal dysfunction. Patients are given a Remote Control to monitor the battery status and settings of their Axonics SNM System.

This manual describes how to use the Remote Control.

Axonics recommends you read this manual before using your Remote Control.



Manual overview

This manual describes the steps for safe use of your Remote Control. The manual includes:

- Description of Remote Control buttons, indicators and functions
- Steps for using your Remote Control
- Troubleshooting Remote Control issues
- Additional information about the Remote Control

Note: The Remote Control may behave differently when communicating with the Trial Stimulator (TS) and the Permanent Stimulator (PS). This manual will highlight when certain behavior is specific to the TS or PS. Otherwise both stimulators are referred to commonly as the Stimulator.

Specifications

Item Specification

F
ons
5°C - 40°C (41°F – 104°F)
70kPa – 106kPa
15% - 95%
ent Conditions – Short Term (3 days or less)
-25°C - 70°C (-13°F – 158°F)
57kPa – 106kPa
15% - 95%
ent Conditions – Long Term (over 3 days)
20°C - 30°C (68°F – 86°F)
70kPa – 106kPa
30% - 85%
Lithium-ion battery (non-rechargeable)
7.1 x 3.8 x 1.6 cm
60g
5 years
Radiofrequency telemetry
Model: 2301 FCC ID: 2AEEGP IC: 20225-P Quality of Wireless Service: This device operates in the 401-406 MHz frequency and the maximum effective radiated power is below the limit of 25 μW ERP/EIRP as specified in EU: EN ETSI 301 839 and USA: FCC 47 CFR Part 95; Subpart I. The Remote Control has to be within 1 meter from the Stimulator for successful communication. Wireless Security: The Remote Control is uniquely paired to a specific Stimulator. Additional mechanisms exist to ensure the integrity of radio data.

^{*} Expected battery life based on typical use scenario

Specifications

FCC Compliance

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

This transmitter is authorized by rule under the Medical Device Radio Communication Service (in part 95 of the FCC Rules) and must not cause harmful interference to stations operating in the 400.150–406.000 MHz band in the Meteorological Aids (i.e., transmitters and receivers used to communicate weather data), the Meteorological Satellite, or the Earth Exploration Satellite Services and must accept interference that may be caused by such stations, including interference that may cause undesired operation. This transmitter shall be used only in accordance with the FCC Rules governing the Medical Device Radio communication Service. Analog and digital voice communications are prohibited. Although this transmitter has been approved by the Federal Communications Commission, there is no guarantee that it will not receive interference or that any particular transmission from this transmitter will be free from interference.

IC Compliance

This device complies with Industry Canada license-exempt RSS standard(s). Operation is subject to the following two conditions: (1) this device may not cause interference, and (2) this device must accept any interference, including interference that may cause undesired operation of this device.

FCC and IC Compliance

This device may not interfere with stations operating in the 400.150–406.000 MHz band in the Meteorological Aids, Meteorological Satellite, and Earth Exploration Satellite Services and must accept any interference received, including interference that may cause undesired operation.

Note: Changes and modifications to the Remote Control not authorized by Axonics could void FCC and IC certification and negate your authority to use the product.

Axonics contact information

Please contact your doctor if you have questions about your health or the Axonics SNM therapy.

Axonics Patient Support

Axonics, located in Irvine, CA (USA) can help answer questions or concerns about your Axonics SNM therapy. Please note that Axonics cannot discuss or comment on your medical condition.

■ Call: +1-949-396-6320

■ Hours: M-F, 8:00 am - 5:00 pm (Pacific Time)

Label Symbols

	Remote Control Remote Control User Manual (this document)
∩ R	,
₩	
SN	Product Serial Number
<u>~</u>	Manufacturer
REF	Product Model Number
M	Manufacturing Date
 ∱	EC 60601-1/EN60601-1, Type BF Equipment
(°i)) No	on ionizing electromagnetic radiation
TI	Conformité Européenne (European Conformity): 2018. This symbol means that the device fully complies with AIMD Directive 90/385/EEC Notified Body reviewed) and RED 2014/53/EU (self-certified)
IC In	ndustry Canada certification number
° 61.	Classified by CSA with respect to safety
UPA DY ONLY C	or USA audiences only Caution: US Federal law restricts this device for sale by or on the order of a Physician
∏i R€	efer to instructions for use (Consult accompanying documents)
1 To	emperature limitation
Ø H	Humidity limitation
€ Pi	Pressure limitation
(A) D	Oo not use if package is damaged
EC REP	Authorized representative in the European community
	rotection from the amount of dust that would interfere with the operation of the evice.
Pr	rotection against temporary immersion in water.
l 🙈 🗛	his device complies with all applicable Australian Communications and Media authority (ACMA) regulatory arrangements and applicable electrical equipment afety requirements.





Axonics Modulation Technologies, Inc. 26 Technology Drive Irvine, CA 92618 (USA) www.axonicsmodulation.com Tel. +1-949-396-6320 Fax +1-949-396-6321



HealthLink Europe Services BV De Tweeling 20-22 5215 MC's - Hertogenbosch The Netherlands



