Eon Mini Rechargeable IPG

The feature-filled Eon Mini[™] IPG–the smallest IPG on the market and the longest-lasting in its class–offers a powerful combination of size and performance to meet evolving pain management needs.¹⁻⁴



	ANS Eon Mini	Medtronic RestoreULTRA	Boston Scientific Precision
Volume	18 cc	22 cc	22 cc
Weight	29 g	45 g	33 g
Battery life approval	10 years, open-ended ^a	9-year end-of-life ^b	5 years, open-ended
Battery life approval settings	High	Medium	Medium
Current delivery	Constant current	Constant voltage	Constant current
Charging system	Mobile	Mobile	Mobile
Charging coil type	Internal	Internal	Internal
Maximum recommended implant depth	2.5 cm	1.0 cm	2.0 cm
Contacts	16	16	16
Amplitude	0-25.5 mA	0-10.5 V	0-12.7 mA°
Pulse width	50-500 µs	60-1000 µs	20-1000 µs
Frequency	2-1200 Hz	2-1200 Hz	2-1200 Hz
Maximum sustainable frequency	1200 Hz for 1 stim set 600 Hz for 2 stim sets 300 Hz for 4 stim sets 200 Hz for 6 stim sets 150 Hz for 8 stim sets	1200 Hz for 1 Stim Set 600 Hz for 2 Stim Sets 300 Hz for 4 Stim Sets	1200 Hz for 1 stim set 130 Hz for 2 stim sets 80 Hz for 4 stim sets
Maximum amplitude with single cathode at 524 ohms	24.7 mA	21.8 mA	12.7 mA
Recharge interval	Days between necessary recharges		
Low (3 mA, 200 µs, 40 Hz)	121	54	61
Medium (6.5 mA, 225 µs, 70 Hz)	22	19	20
High (8.6 mA, 238 us, 160 Hz)	6	6	6

The Eon Mini IPG is best in class.

a At least 24 hours between recharges at 10 years at high settings

b Circuitry renders the device inoperable after 9 years

c $\,$ 20 mA is possible if using more than one cathode $\,$



References

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Chart sources

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NEUROMODULATION

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Indications for Use: Chronic, intractable pain of the trunk and limbs. Contraindications: Demand-type cardiac pacemakers, patients who are unable to operate the system or who fail to receive effective pain relief during trial stimulation. Warnings/Precautions: Diathermy therapy, cardioverter defibrillators, magnetic resonance imaging (MRI), explosive or flammable gases, theft detectors and metal screening devices, lead movement, operation of machinery and equipment, postural changes, pediatric use, pregnancy, and case damage. Patients who are poor surgical risks, with multiple illnesses, or with active general infections should not be implanted. Adverse Events: Painful stimulation, loss of pain relief, surgical risks (e.g., paralysis). Clinician's manual must be reviewed prior to use for detailed disclosure. Caution: U.S. federal law restricts this device to sale and use by or on the order of a physician.

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