

MANUFACTURER SPECIFIC TRANSLATION OF ICD PROGRAMMING RECOMMENDATIONS[‡]

‡ The manufacturer specific programming settings/choices set forth below are based on a compilation of clinical expertise and clinical trial data as reported in the 2015 Consensus Statement On Optimal ICD Programming and Testing, of which this Appendix B is a part. These recommended settings/choices represent a diligent and good faith effort on the part of the Writing Committee to translate the Consensus Statement recommendations to device settings/choices for the four specified clinical issues/ ICD therapies where the writing committee considered that there was sufficient consensus and supporting data to make recommendations intended to improve the safety, morbidity and mortality profile of patients with these clinical issues/ICD therapies. They are the recommendations of the Writing Committee only. They do not represent the position or recommendations of HRS, EHRA, SOLAECE, or APhRS, nor are they the manufacturer's nominal settings or the precise programming tested during clinical trials of these devices, nor are they necessarily the settings/choices that would be recommended by the manufacturer. These recommended settings/choices are not applicable in all circumstances. As stated in the Introduction to the Consensus Statement – "Care of individual patients must be provided in the context of their specific clinical condition and data available on that patient." Each treating physician must carefully consider the circumstances of their individual patient and determine whether these recommended settings/choices are appropriate to their patient's circumstances.

BIOTRONIK

All "I-family" devices *Iforia, Ilestio, Idova, Iperia, Itrevia and Inventra and Lumax 740*; Single, Dual chamber devices, CRT-D devices and VR-DX devices.

Brady

Single Chamber
VVI 40 bpm

Dual Chamber
DDD-CLS with IRS Plus / I OPT or
DDD with Vp Suppression +/- Rate Response

CRT
DDD-CLS or Rate Response

Detection

In patients with no VT history¹
VF: 231 bpm, 24/30 intervals (30/40 in *Iperia, Itrevia and Inventra*)
VT2: 188 bpm², 30 intervals
VT1: Monitor zone at user discretion

In patients where VT CL is known
VF 231 bpm (minus Safety Margin³), 24/30 intervals
VT2: 188 bpm² (minus Safety Margin³), 30 intervals
VT1: Therapy at 10-20bpm < VT CL or Monitor zone at user discretion

¹ SVT discriminators are linked to Detection Zones. An alternative configuration would be VF 250 bpm, VT2 230 bpm and VT1 188 bpm with therapy (i.e. no Monitor zone) if >1 ATP attempt desired up to 250 bpm.

² 194 or 200 bpm are also within guidelines for the lower limit of this zone

³ Safety Margin: If clinical VT CL is known, a detection zone should be set at 10-20 bpm below documented rate

Therapy

VF: ATP One-Shot, 1 burst of 8 pulses at 85% CL, then full output shocks
VT2: ATP ≥1 bursts of 8 pulses at 85% CL, 10ms scan decrement, then shocks
VT1: Therapy as for VT2 (favoring more ATP) or Monitor zone

SVT Discriminators

Single Chamber
MorphMatch ON in *Iperia, Itrevia and Inventra*⁴
Onset ON 20% (default)
Stability ON 40ms (unless likely polymorphic VT then Stability OFF)
Sustained VT Timer OFF

Dual Chamber/CRT-D
SMART ON at default settings (or adapted to known VT)

⁴ In order to have MorphMatch on in the VT2 zone, device must be programmed to three zones with VT1 zone as monitor or therapy zone as required. This does not apply to dual chamber SVT discriminators.

Sensing Rejection

Lead Integrity check ON (if available)
BIOTRONIK HomeMonitoring ON (if available)

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Boston Scientific

Most settings are nominal in newest devices. Those that are not nominal are marked with an asterisk.

Brady	<u>Single Chamber</u> VVI, 40bpm*
	<u>Dual Chamber</u> DDD with RYTHMIQ™* or AV Search +* (± rate response)
	<u>CRT</u> DDD ± rate response Smart Delay™ Optimization of AV delays
Detection	<u>In patients with no VT history</u> VF: 8 of 10 intervals plus 5 second delay*, 250 bpm* VT: 8 of 10 intervals plus 12 second delay*, 185 ¹ bpm* VT-1: Monitor zone at user discretion, ≥12 second duration*
	<u>In patients where VT cycle length is known</u> VF: 5 second duration*, 250 bpm* VT: 12 second duration*, 185 ¹ bpm* or 10-20bpm < VT rate VT-1: Therapy at 10-20bpm < VT CL, ≥12 second duration* or Monitor zone

¹ 190, 195 or 200bpm are also within guidelines for the lower limit of this zone

Therapy	VF: Quick Convert™ ON to 300bpm* (if available) All shocks: Maximum output
	VT: ATP-1: Scan, ≥1 bursts, 8 pulses* at 84%* coupling interval and cycle length (Minimum 200ms*), 10ms decrement* ATP-2: OFF* All shocks: Maximum output
	VT-1: As for VT, favoring more ATP

SVT Discriminators	<u>ICD</u> Rhythm ID®: ON
	<u>CRT-D</u> Onset/Stability: ON or Rhythm ID®: ON*
	Sustained Rate Duration (SRD): OFF* SVT Discriminators apply only up to 230 bpm

Oversensing Rejection	Non-physiological Signal Detected: ON (LATITUDE™)
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Others	Turn on "Beep When Out-of-Range" Daily Lead Measurements*
	RV Pacing Impedance Abrupt Change alert ON (LATITUDE™) Single Chamber: Consider %RV pacing alert ON (LATITUDE™)
	Dual Chamber: Consider %RV pacing alert in non AVB patients ON (LATITUDE™) CRT-D: Consider CRT % pacing alert ON (LATITUDE™)

SUBCUTANEOUS ICD (EMBLEM™ S-ICD)	
Settings	Shock Zone: From 230bpm
	Conditional Zone: From 200bpm
	Consider post-shock pacing ON

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Medtronic

Most settings are nominal in newest devices. Those that are not nominal are marked with an asterisk.

Brady	<u>Single Chamber</u> VVI 40bpm
	<u>Dual Chamber</u> Managed Ventricular Pacing (MVP; AAI ↔ DDD) ± rate response
	<u>CRT</u> DDD
	Patient with Intact AV conduction – Adaptive BiV & LV* (if available) Patient with prolonged PR interval – Adaptive BiV (if available)

Detection	<u>In patients with no VT history</u> VF:30/40 intervals, 188 bpm ¹ FVT: OFF ² VT:OFF VT Monitor: User discretion
	<u>In patients where VT CL is known</u> VF:30/40 intervals, 188 bpm ¹ FVT: OFF ² VT:24* intervals ³ , 10-20bpm < VT rate VT Monitor: User discretion

¹ 194 or 200 bpm are also within guidelines for the lower limit of this zone

² Use of ATP Before/During Charging in the VF zone achieves the same functionality as use of the FVT zone

³ Consecutive count in VT zone; hence, lower NID as per PainFree SST data

Therapy	VF:	ATP Before* Charging; ChargeSaver ON All shocks: Full output
	VT (if ON):	Rx1: ATP, ≥1 bursts of 8 pulses at 88% VT CL, 10ms Decrement Rx2-6: shocks

SVT Discriminators	<u>Single Chamber</u> Wavelet: ON Limit: 260ms (230 bpm) Stability: OFF Onset: OFF
	<u>Dual Chamber/CRT-D</u> PR Logic: ON (Other 1:1 not turned ON until after lead stabilized at ~3 months) Wavelet: ON (if available) SVT Limit: 260ms (230 bpm) Stability: OFF Onset: OFF

Oversensing Rejection	Lead Integrity Alert: ON T wave Over-Sensing: ON (if available) RV Lead Noise: ON* without timeout (if available)
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LIVANOVA (formerly Sorin Group)

(OVATIO, PARADYM, PARADYM RF, INTENSIA and PLATINIUM high voltage devices)

Brady

Single Chamber

VVI 40bpm

Dual Chamber

SafeR (± rate response) or DDD (± rate response)

CRT

DDD (± rate response) – with weekly AV + VV SonR optimisation ON*

*requires SonRtip atrial lead with integrated haemodynamic sensor

Detection

In patients with no VT history

VF:	>255 bpm	20 cycle + 6/8 majority
FVT:	230 bpm	20 cycle + 6/8 majority
VT:	185 bpm ¹	20 or 30 cycle + 6/8 majority
Slow VT:	Monitor zone at user discretion	

In patients where VT CL is known

VF:	>255 bpm	20 cycle + 6/8 majority
FVT:	230 bpm	20 cycle + 6/8 majority
VT:	Clinical rate less 10-20 bpm	≥20 cycle + 6/8 majority
Slow VT:	Monitor zone at user discretion	

¹ 190, 195 or 200 bpm are also within guidelines for the lower limit of this zone

Therapy

VF:	6 x 42 J
FVT:	If stable*: 1 x ATP (Burst @ 85% x 8 beats) then 6 x 42 J
	If unstable: 6 x 42J
VT:	≥1 x ATP (Burst + Scan @ 85% x 8 beats; Scan 8 ms) then 6 x 42 J

*Satisfaction of stability (nominal value = 30 ms) in the Fast VT zone will not prevent therapy but rather activate programmable burst pacing prior to shock therapy.

SVT Discriminators

Single Chamber

Single button programming; Stability+/Acc

Rate, Stability, Degree of Onset, VT long cycle search

Nominal settings: Onset 19%, Stability 65ms (Slow VT, VT); Long cycle extension 10 cycles; Long cycle gap 170 ms.

Dual Chamber/CRT-D

Single button programming; PARAD+

Rate, Stability, AV association analysis, Degree and Chamber of Onset, VT long cycle search

Nominal settings: Onset 25%, Stability 65ms (Slow VT, VT); Long cycle extension 10 cycles; Long cycle gap 170 ms.

Oversensing Rejection

Daily check Lead impedance ON
Daily check Lead coil continuity ON
Daily check V over-sensing alerts ON
T-wave filtering and noise detection are hardcoded in firmware

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St. Jude Medical

Brady

Single Chamber

VVI 40bpm

Dual Chamber

DDD with Ventricular Intrinsic Preference (VIP) ± rate response

CRT

DDD ± rate response

QuickOpt at implant and follow-up

Detection

In patients with no VT history

VF:30 intervals, 250 bpm

VT2: 30 intervals, 187 bpm¹

VT:Monitor, at user discretion

In patients where VT CL is known

VF:30 intervals, 250 bpm

VT2: 30 intervals, 187bpm or 10-20bpm < VT rate

VT:Therapy at 10-20bpm < VT rate or Monitor zone

¹ 190, 193, 196 or 200 bpm are also within guidelines for the lower limit of this zone

Therapy

VF: ATP While Charging, 8 pulses at 85% VT CL

All shocks: Maximum output (Note: 1st shock 4-6J lower than full output)

VT2: ATP, ≥1 bursts of 8 pulses at 85% VT CL

Scan step 10ms, Re-adaptive ON, Minimum CL 200ms

All shocks

VT: As for VT2, favoring more ATP

SVT Discriminators

Single Chamber

Morphology: ON, 90%, 3 of 10

All others: "Passive"

Dual Chamber/CRT-D

Morphology: ON, 90%, 3 of 10

Arrhythmia onset: ON (default settings)

Interval Stability: ON (default settings)

If ALL

For CRT: Template Auto Update 30 days and Template Pacing Hysteresis ON
or Template Auto Update OFF

SVT Upper Limit: 230 bpm

SVT Discrimination Timeout: OFF

VT Therapy Timeout: OFF

Oversensing Rejection

Low Frequency Attenuation: ON

SecureSense RV Lead Noise Discrimination: ON