Product Performance Report

July 2017

BIOTRONIK
excellence for life

Product Performance Report

July 2017

Cardiac Rhythm Management

Pacemakers

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1 Quality Excellence

BIOTRONIK has a long history of high quality in product design and performance. For more than 50 years, the name BIOTRONIK has been synonymous with excellent workmanship and reliable patient safety. Our quality concept follows an integrated approach and extends from preventative risk measures during a product's development phase through all the steps of the manufacturing and design process.

BIOTRONIK's quality assurance system guarantees strict adherence to internal quality standards as well as compliance with international standards and guidelines. Regular reviews of our product performance and manufacturing evaluations contribute significantly to the achievement of extraordinary quality. Our customers, patients, and physicians can rely on the highest degree of safety built into our products. We always welcome suggestions from users about how we can improve the quality of our products.

This Product Performance Report is an integral component of BIOTRONIK's commitment to provide detailed, accurate information regarding long term reliability. The Product Performance Report exemplifies BIOTRONIK's policy of transparent and timely communication with our customers.

As a means to obtain continuous improvement of the designs, BIOTRONIK carefully analyzes returned products and incorporates all findings into our quality assurance system. This Product Performance Report was prepared in accordance with International Standard ISO 5841-2: 2014 (E) [1] and is in compliance with the recommendations from the U.S. Heart Rhythm Society Task Force on Device Performance Policies and Guidelines. As an active member of AdvaMed and their Pacemaker/ICD Working Group, BIOTRONIK has worked extensively with the CRM industry to ensure comparable product performance data is reported by all manufacturers.
The data provided in BIOTRONIK's Product Performance Report incorporates the requirements and definitions as defined in AdvaMed's Requirements for Uniform Reporting of Clinical Performance of Pulse Generators, except as noted herein.

In BIOTRONIK's continuous efforts to provide accurate and transparent information and to ensure that a conservative estimate for device performance is reported, the Survival Probability calculations presented herein also consider reported pacemaker and ICD battery depletions as well as lead complications without the device having been returned for analysis.

1 The ISO 5841-2:2014(E) is replacing the previous version ISO 5841-2:2000. As part of the update, AdvaMed's Requirements for Uniform Reporting of Clinical Performance of Pulse Generators were incorporated in the new ISO 5841-2:2014(E).

Because a significant portion of this report is based on analyses of returned products, BIOTRONIK urges all physicians to return explanted devices and to notify us when a product is explanted or no longer in use for any reason.

BIOTRONIK aims to continually improve and enhance the scope of this report while integrating the latest information and data concerning the performance of our products. Please contact our U.S. Compliance Department (888) 345-0374 or Worldwide CRM Technical Service Department at ppr@biotronik.com with any comments, suggestions or questions regarding this report. Your feedback is highly appreciated and will be used to further develop this report.

BIOTRONIK, July 2017

Dr. Volker Lang
Vice President Global Quality Management
BIOTRONIK SE & Co. KG

2 Terms and Definitions

The following terms and definitions are used for Pacemakers and Implantable Cardioverter Defibrillators (ICDs) as well as pacing and ICD leads throughout this Product Performance Report. These definitions form the basis for this Product Performance Report by clearly articulating the status of each device return and product analysis classification.

Elective Replacement Indicator

All active implantable devices that are powered by an internal battery need to be replaced when their battery is depleted. BIOTRONIK pacemakers and ICDs have an Elective Replacement Indicator (ERI) feature aka Recommended Replacement Time (RRT) that notifies the health care provider when the device's battery is nearing the end of its useful life. Display of ERI is BIOTRONIK's recommendation to the user that the battery's present state will require device replacement in the near future. For further details please refer to the corresponding manual.

Battery Depletion

Battery depletions are classified as either normal (expected) or premature. Premature battery depletions are defined as device malfunctions, while normal battery depletions are not device malfunctions. Batteries of returned devices are considered to have depleted normally when (a) a device is returned with no associated complaint and the device has reached its elective replacement indicator(s) with implant time that meets or exceeds the nominal (50 percentile) predicted longevity at default (labeled) settings, or (b) a device is returned and the device has reached its elective replacement indicator(s) with implant time exceeding 75% of the expected longevity using the longevity calculation tool available at time of product introduction, calculated using the device's actual use conditions and settings. For consistency with previous product performance reports, for ICDs released prior to Lumax and pacemakers released prior to Philos II, batteries of returned devices are considered to have depleted normally if they have reached their elective replacement indicator and testing indicates that the battery and associated circuitry are within specifications.

Out of Specification

Any component or software related event that causes the device's characteristics to not meet pre-defined performance specifications and requirements while implanted and in service. Returned product analysis that determines a device to be out of specification is considered a device malfunction. Normal battery depletions are not considered device malfunctions. BIOTRONIK defines the requirements and performance specifications for each product.

Device Malfunctions

Any component or software related event that causes the device's characteristics to be out of specification while implanted and in service are considered as device malfunctions. Because it is impossible to verify that a device has malfunctioned without analyzing it, only returned devices can be classified as malfunctions for this report. Each returned lead, ICD and pacemaker is analyzed to determine if it has malfunctioned. If the analysis determines that a pacemaker or ICD failed to meet its specifications while implanted and in service, it is further classified as either a malfunction with compromised therapy or as a malfunction without compromised therapy. Devices damaged during implant, revision or after explant, damaged due to external causes (i.e. electrocautery) or due to failure to follow instructions, warnings or contraindications in its associated technical manual are not considered malfunctions. Devices damaged due to interaction with other implanted devices (i.e., leads) are also not considered as malfunctions for the purposes of this Product Performance Report.

Malfunctions with Compromised Therapy

The condition when a pacemaker or ICD is found to have malfunctioned in a manner that compromised pacing or defibrillation therapy (including complete loss or partial degradation) while implanted and in service. Therapy is considered to have been compromised if critical patient-protective pacing or defibrillation therapy is not available.
Examples include: sudden loss of battery voltage; accelerated current drain such that a depleted

battery was not detected before loss of therapy; sudden malfunction during a tachycardia or fibrillation event resulting in aborted delivery of therapy; intermittent malfunction where therapy is sporadically unavailable.

Malfunctions without Compromised Therapy

The condition when a pacemaker or ICD is found to have malfunctioned in a manner that did not compromise pacing or defibrillation therapy while implanted and in service. Therapy is not compromised as long as critical patient-protective pacing and defibrillation therapies are available as determined through device analysis.

Lead Complications

A lead performance issue where a complaint, associated with at least one of the clinical manifestations listed below, is reported and where the non-returned lead is:

- Verified by medical records to have been implanted and in-service, and
- Reported to have been removed from service,
- Modified to remedy the malfunction, or
- Left in service based on medical judgment.

Complications for leads implanted greater than 30 days are reported as Qualifying lead complications, whereas complications occurring during the first 30 days are reported as Acute lead observations.

- Failure to Capture
- Failure to Sense
- Oversensing
- Abnormal Pacing Impedance
- Abnormal Defibrillation Impedance
- Insulation Breach
- Conductor Fracture
- Lead Dislodgement
- Extracardiac Stimulation
- Cardiac Perforation
- Other

Survival Probability Estimates

The probability that a device remains operational during a discrete time interval is defined as survival probability. Survival probability, as presented in this report, is related to device survival only and not survival of the patient. The survival probability estimates in this report are based on BIOTRONIK's analysis of returned products as well as events that are reported to BIOTRONIK (e.g., battery depletions or lead complications).

Cumulative Survival Probability Estimates

The survival probability over a device's service time is the cumulative survival probability. It is calculated from all discrete survival probabilities of previous time intervals. This characteristic is calculated separately for malfunction-free survival and all-cause survival (including normal battery depletions). Specific populations that are subject to a safety advisory notification are excluded and shown separately.

Implanted Devices

Only devices remaining implanted for at least one calendar day after the implantation date are considered as implanted. Devices that are removed from the patient on the same calendar day as the implant procedure do not contribute to the survival statistics.

Active Implants

The number of devices that remain operational within a discrete observation interval are active implants. Units are removed from this cohort due to patient death or explant for any reason.

Underreporting

A device status may change without being accounted for in the Product Performance database due to a lack of information being provided to BIOTRONIK. Underreporting adjustments deemed to be necessary are detailed in this report.

Safety Advisory Notifications

Any action taken by the manufacturer to inform clinicians concerning a device performance issue that may cause the device to not meet its predefined specifications is referred to as a Safety Advisory Notification.

3 Methodology for Pacemaker and ICD Survival Estimates

3.1 Cumulative Survival Probability

This report has been prepared in accordance with ISO 5841-2:2014(E) applying actuarial analysis for the calculation of survival probabilities. Survival estimates given in this report are considered to be generally representative for worldwide performance of BIOTRONIK devices.

The Cumulative Survival Probability is an estimate based on the percentage of pacemakers and ICDs that remain implanted and operational at various points of the product's service time in absence of concurrent events such as morbidity and voluntary explants for various reasons (e.g., device upgrade). The Device Survival Estimate over time is displayed in cumulative curves (Kaplan-Meier). The product's performance is evaluated in discrete one-month intervals. The survival probability for each month is given by the number of devices that remain implanted and operational through this month divided by the number of devices that entered the interval. The cumulative survival probability for any period is given by multiplying all survival probabilities of previous months.

At the time of implantation, the cumulative survival probability is 100 %. Even though they are analyzed as part of our quality system monitoring, devices that are found to be out of specification prior to or during the implantation procedure are removed from the statistics as they do not contribute to a patient's risk of being subject to a device malfunction or replacement during the device's service time. Because this report is provided to describe product performance based on returned product analyses, the pacemaker and ICD data does not include information regarding medical complications such as erosion, infection or diaphragmatic stimulation.

In general, during the initial phase of the service time, devices which are out of specification are the primary contribution to reduction of survival probability. As the product lifecycle lengthens, normal battery depletion assumes a greater impact on the survival curve and becomes the dominating factor.

In order to make these two effects distinguishable, the cumulative survival probability curves are shown separately for devices that are confirmed to have malfunctioned only, and for total (all-cause) cumulative survival. In case of a device being subject to a safety advisory notification that significantly impacts the survival probability, this factor is displayed separately.

3.2 Data Acquisition

This report is based on the observation of BIOTRONIK's U.S. products through review of our device registration and tracking systems and analyses of returned products from all sources. Because the ability to perform decedent searches of patients with BIOTRONIK devices via the U.S. Social Security Administration, the use of U.S. data more accurately represents the active patient population for reporting purposes. In addition, device tracking regulations and vigilance reporting regulations vary throughout the world; therefore use of the U.S. data is most appropriate for accurate and consistent reporting of product performance.

In order to be included in the population under observation, a device must be registered and implanted for at least one calendar day. The cutoff date for the data included in this report is December 31, 2016. The number of U.S. devices that are implanted and remain active as well as the total number of products distributed worldwide are provided for each product family in this report. Information is provided for separate product families, in that devices with nearly identical hardware and therapy functions are combined. For example, Lexos VR and VR-T (with Home Monitoring) ICDs are combined into a single family, Lexos Single Chamber ICDs.

Survival estimates are calculated for product families having accumulated at least 10,000 cumulative implant months. Because 10,000 implant months may take some time to accumulate, there may be a gap between U.S. market release and the start of graphical representation of survival probability. Products no longer being distributed with less than 500 active implants may be excluded from this report.

ISO 5841-2 describes a method for adjusting the device survival probability to compensate for underreported malfunctions and unrelated patient deaths. The factor for underreporting of malfunctions is unknown as currently no systematic data is available that reveals this factor. Consequently, this factor remains unaccounted for this report. Patient mortality is artificially elevated if the reported rate from our registration and tracking systems is below the annual mortality in clinical studies. Normal battery depletion rate is assumed if the reported rate of depletion decreases over time.

3.3 Returned Product Analysis

Information on malfunctioning for the pacemaker and ICD portions of this report is taken exclusively from the analysis of returned products. The outcome of this analysis is the basis for the final classification of the device's cause of explantation. Only analyzed products with confirmed device malfunctions are utilized in the calculation of malfunction-free survival probability.

Every pacemaker and ICD returned to BIOTRONIK is analyzed per internal procedures and classified as functioning normally, normal battery depletion, or malfunctioning (including premature battery depletion) while implanted and in service. These device classifications are the basis for BIOTRONIK's cumulative survival estimates on pacemakers and ICDs.

As a significant portion of pacemakers and ICDs with normal battery depletion are not returned for analysis, BIOTRONIK also considers unconfirmed pacemaker and ICD battery depletions (reported, but device not returned) in the total survival estimates to ensure that a conservative estimate for device performance is reported.

3.4 Product Performance Graphs and Data

The product performance information is shown in each section in alphabetical order and by product type.

For each product, the report provides:

- Product versions that contribute to the evaluation
- U.S. and CE market release dates
- Worldwide quantity of products that have been distributed
- U.S. registered implants (number of products included in this report)
- Estimated active U.S. implants
- Number of U.S. normal battery depletions
- Number of U.S. confirmed malfunctions

The survival plots provide:

1. Total Survival

The combined cumulative survival probability for all causes that result in device removal or a system out of operation, excluding removals for clinical reasons unrelated to the device's performance (i.e., infections).

2. Malfunction-Free Survival

The cumulative survival probability free of component or software malfunctions excluding normal battery depletions, but including premature battery depletions. Normal battery depletions only have an impact on the total cumulative survival.

Products or subgroups of products may become subject to safety advisory notifications that can significantly impact the overall product performance. However, as these subgroups are clearly defined they are separated from the non-advisory devices. The impact of the advisory notification is then shown in a separate graph for total cumulative survival and for malfunction-free survival of the device population affected by the advisory notification. Current advisories are listed in chapter 10 of this report.

The cumulative survival data and the 95% confidence intervals according to the Greenwood's Formula[1] are shown in numerical form for the observed population.

1 Greenwood, M. The natural duration of cancer. Reports on Public Health and Medical Subjects 33, London: Her Majesty's Stationery Office, 1–26, 1926.

4 Performance of BIOTRONIK Pacemakers

4.1 Single-Chamber Pacemakers

4.2 Dual-Chamber Pacemakers

4.3 CRT Pacemakers

4.1 Single-Chamber Pacemakers

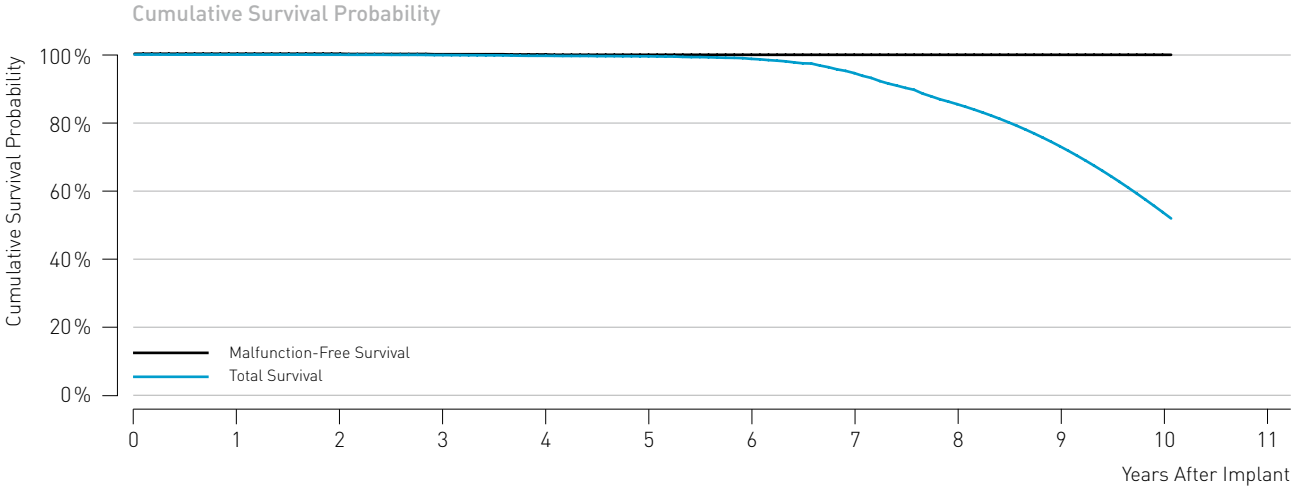
Cylos and Cylos 990

Product Details

Product Versions*	VR, 990 VR
NBG Code(s)	VVIR
US Market Release	Jan 2006
CE Market Release	Nov 2005 / Mar 2008
Worldwide Distributed Devices	25 900
Registered U.S. Implants	6 150
Estimated Active U.S. Implants	3 090
U.S. Normal Battery Depletions	453

	Quantity	Rate
U.S. Confirmed Malfunctions	4	0.07%
▪ Therapy Compromised	1	0.02%
▪ Therapy Available	3	0.05%

* While Cylos 990 VR is not distributed in the U.S., the performance is expected to be similar to the U.S. distributed products



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.
Total Survival [%]	100.0	100.0	100.0	99.9	99.6	99.5	98.8	94.6	85.6	73.4	54.2
(95% Confidence Interval)			±0.1	±0.1	±0.2	±0.2	±0.3	±0.8	±1.4	±1.6	±1.4
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	99.9	99.9	99.9	99.9	99.9	99.9	99.9
(95% Confidence Interval)				±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

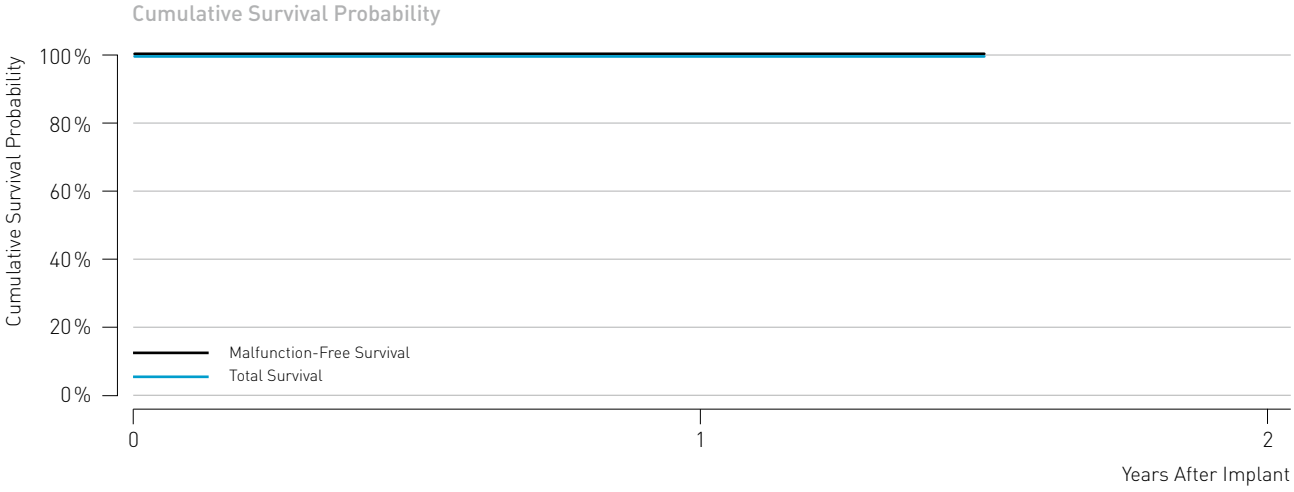
4.1 Single-Chamber Pacemakers

Eluna 8

Product Details

Product Versions	SR, SR-T
NBG Code(s)	AAIR, VVIR
U.S. Market Release	Dec 2014
CE Market Release	Aug 2014
Worldwide Distributed Devices	13 600
Registered U.S. Implants	3 280
Estimated Active U.S. Implants	3 090
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

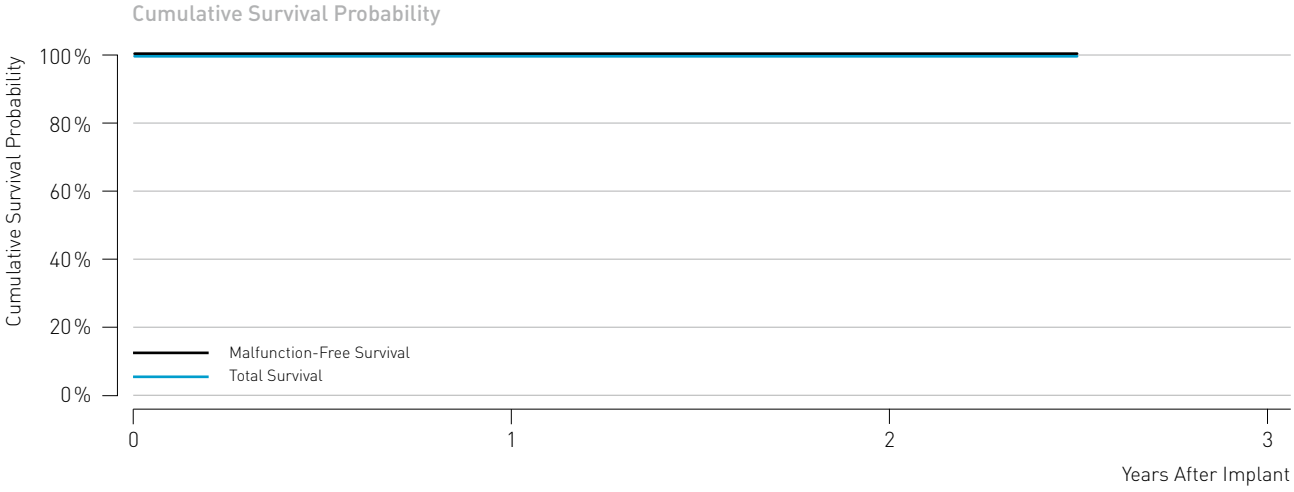
4.1 Single-Chamber Pacemakers

Entovis

Product Details

Product Versions	SR, SR-T
NBG Code(s)	AAIR, VVIR
U.S. Market Release	Jun 2010
CE Market Release	Nov 2009
Worldwide Distributed Devices	28 100
Registered U.S. Implants	2 400
Estimated Active U.S. Implants	2 090
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

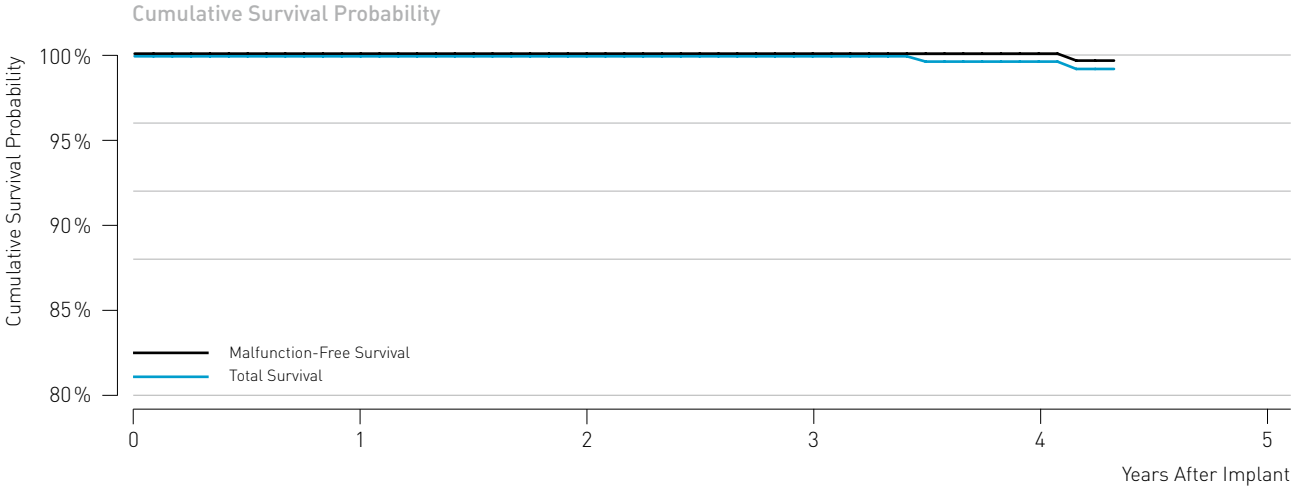
4.1 Single-Chamber Pacemakers

Estella

Product Details

Product Versions	SR, SR-T
NBG Code(s)	AAIR, VVIR
U.S. Market Release	Feb 2011
CE Market Release	Feb 2011
Worldwide Distributed Devices	21 000
Registered U.S. Implants	609
Estimated Active U.S. Implants	429
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.16%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	1	0.16%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	99.7
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1

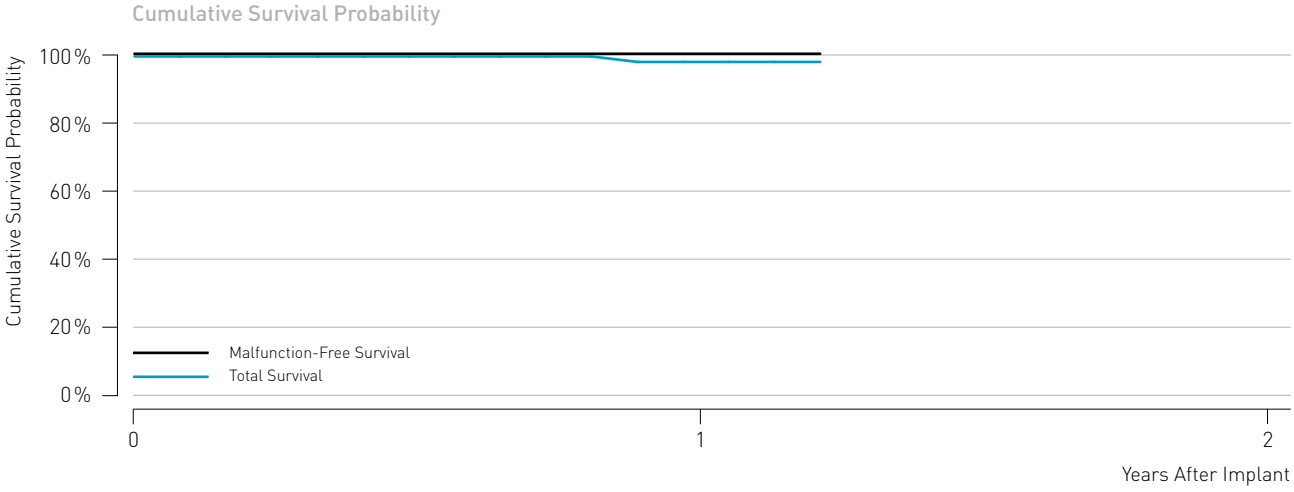
4.1 Single-Chamber Pacemakers

Etrinsa

Product Details

Product Versions	SR-T
NBG Code(s)	AAIR, VVIR
U.S. Market Release	Dec 2014
CE Market Release	Aug 2014
Worldwide Distributed Devices	36 300
Registered U.S. Implants	1 050
Estimated Active U.S. Implants	976
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	99.8
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

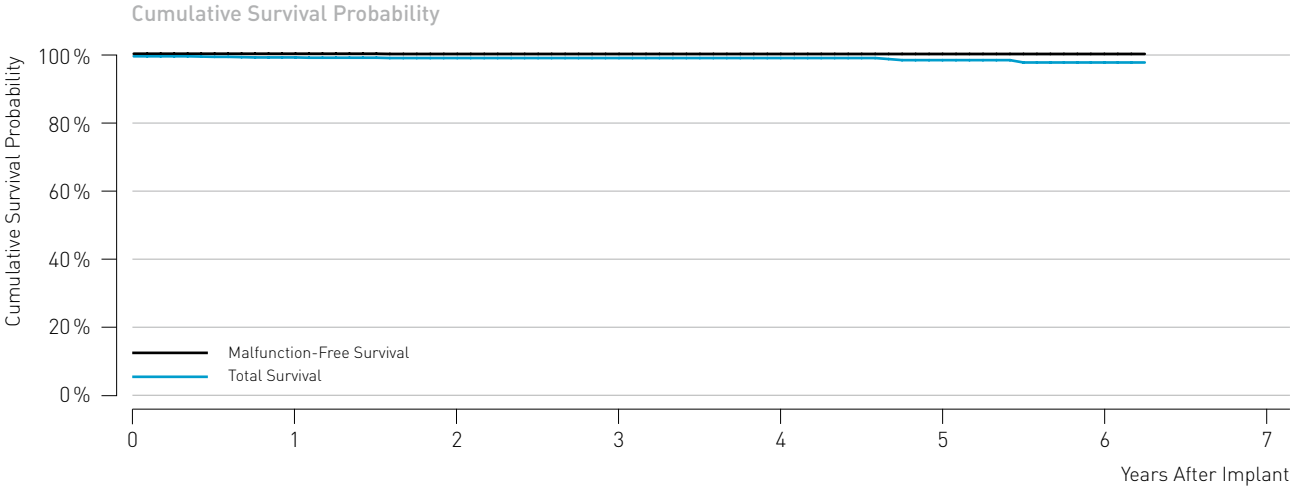
4.1 Single-Chamber Pacemakers

Evia

Product Details

Product Versions	SR, SR-T
NBG Code(s)	AAIR, VVIR
U.S. Market Release	May 2010
CE Market Release	Oct 2009
Worldwide Distributed Devices	56 500
Registered U.S. Implants	12 000
Estimated Active U.S. Implants	9 330
U.S. Normal Battery Depletions	8

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.01%
▪ Therapy Compromised	1	0.01%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.
Total Survival [%]	100.0	100.0	99.9	99.9	99.9	99.9	99.8
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

4.1 Single-Chamber Pacemakers

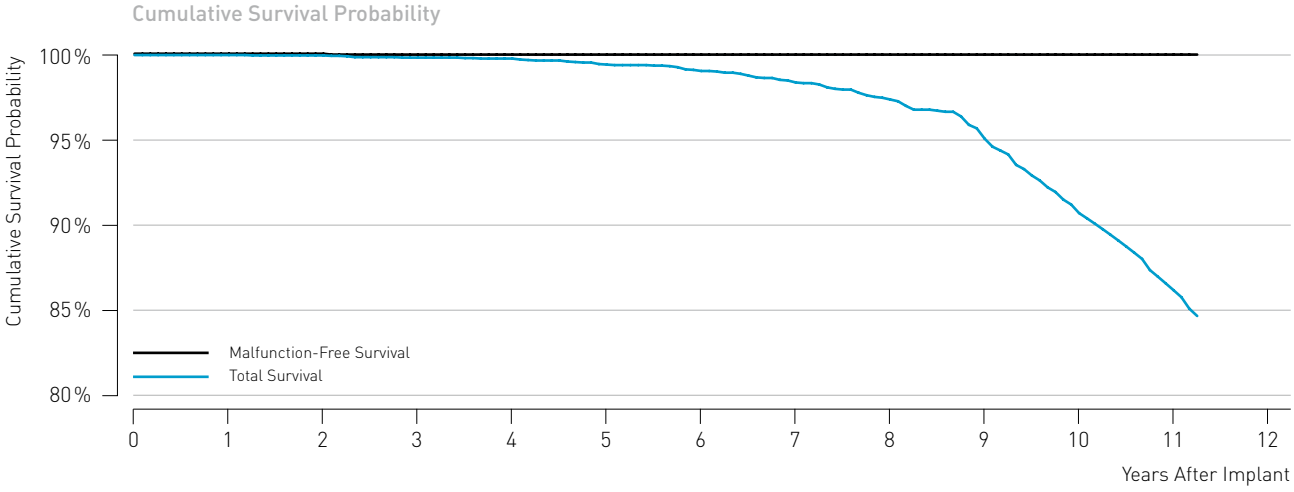
Philos II and Talos

Product Details

Product Versions*	S, SR
NBG Code(s)	SSI, SSIR
U.S. Market Release	Sep 2004
CE Market Release	Feb 2004 / May 2006
Worldwide Distributed Devices	213 000
Registered U.S. Implants	5 240
Estimated Active U.S. Implants	2 830
U.S. Normal Battery Depletions	159

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.02%
▪ Therapy Compromised	1	0.02%
▪ Therapy Available	0	0.00%

* While Talos SR and Talos S are not distributed in the U.S., their performance is expected to be similar to the U.S. distributed products



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.
Total Survival [%]	100.0	100.0	100.0	99.8	99.8	99.4	99.1	98.4	97.4	95.1	90.7	86.1
(95% Confidence Interval)				±0.1	±0.1	±0.2	±0.3	±0.4	±0.6	±0.9	±1.5	±1.9
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
(95% Confidence Interval)												

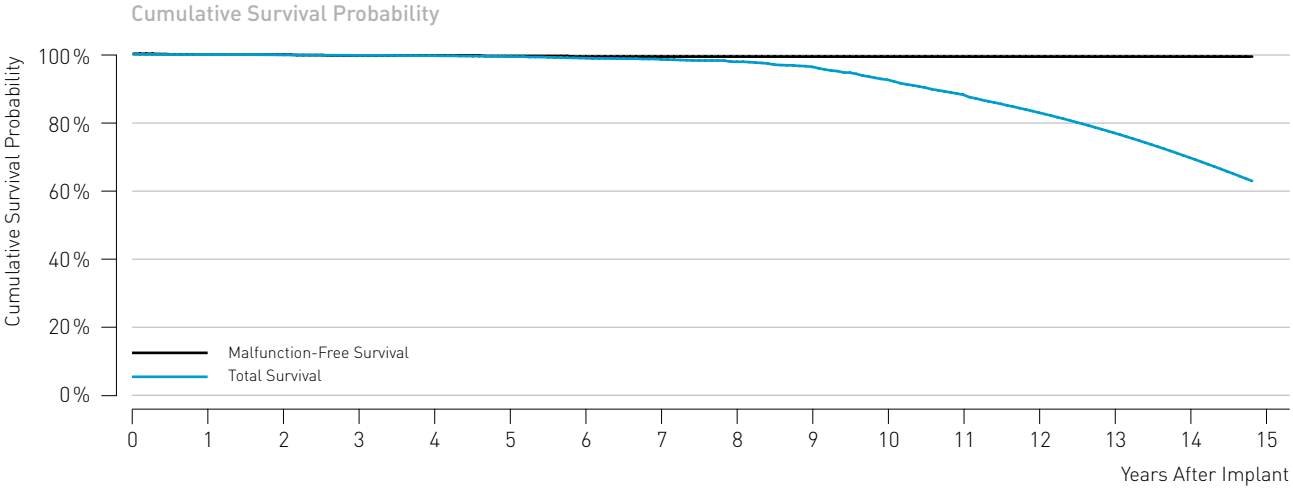
4.1 Single-Chamber Pacemakers

Philos

Product Details

Product Versions	S, SR
NBG Code(s)	SSI, SSIR
U.S. Market Release	Sep 2000
CE Market Release	Aug 2000
Worldwide Distributed Devices	109 000
Registered U.S. Implants	5 770
Estimated Active U.S. Implants	1 580
U.S. Normal Battery Depletions	244

	Quantity	Rate
U.S. Confirmed Malfunctions	7	0.12%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	7	0.12%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.
Total Survival [%]	100.0	99.9	99.8	99.7	99.6	99.4	98.9	98.5	97.8	96.3	92.5	88.1	82.9	76.9	69.7
(95% Confidence Interval)		±0.1	±0.1	±0.2	±0.2	±0.3	±0.4	±0.4	±0.6	±0.8	±1.2	±1.5	±1.7	±1.7	±1.5
Malfunction-Free Survival [%]	100.0	99.9	99.9	99.9	99.9	99.9	99.8	99.8	99.8	99.8	99.8	99.8	99.8	99.8	99.8
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

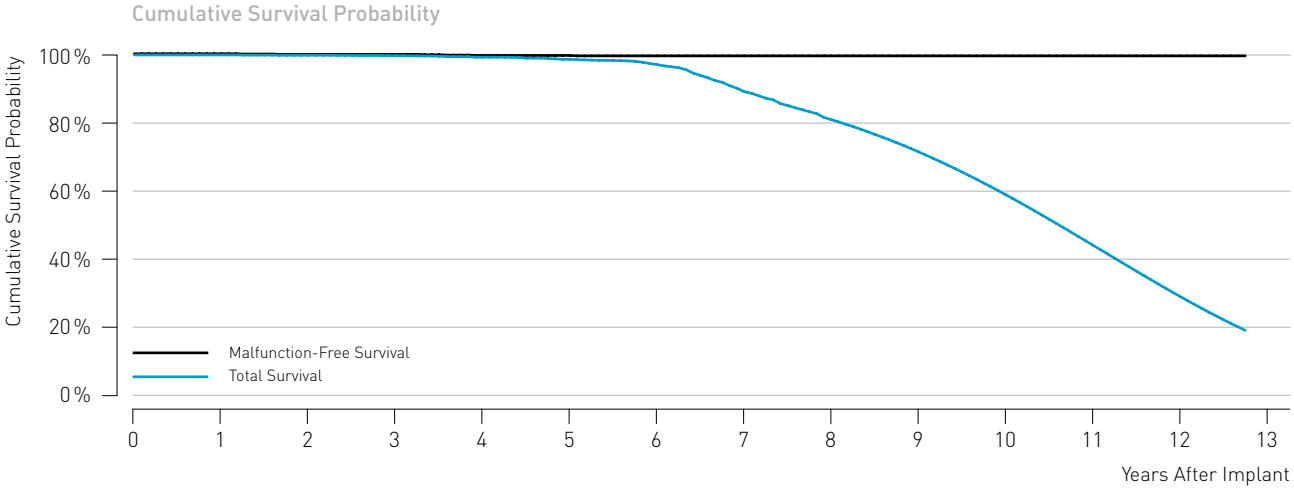
4.1 Single-Chamber Pacemakers

Protos

Product Details

Product Versions	VR/CLS
NBG Code(s)	VVIR
U.S. Market Release	Jan 2003
CE Market Release	Jul 2003
Worldwide Distributed Devices	9 820
Registered U.S. Implants	3 260
Estimated Active U.S. Implants	804
U.S. Normal Battery Depletions	306

	Quantity	Rate
U.S. Confirmed Malfunctions	6	0.18%
▪ Therapy Compromised	2	0.06%
▪ Therapy Available	4	0.12%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.
Total Survival [%]	100.0	100.0	99.9	99.8	99.3	98.7	97.1	89.2	81	71.5	58.9	44.1	29.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.2	±0.4	±0.5	±0.8	±1.6	±2.0	±2.0	±1.7	±1.3	±0.9
Malfunction-Free Survival [%]	100.0	100.0	99.9	99.9	99.8	99.8	99.7	99.7	99.7	99.7	99.7	99.7	99.7
(95% Confidence Interval)			±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2

4.2 Dual-Chamber Pacemakers

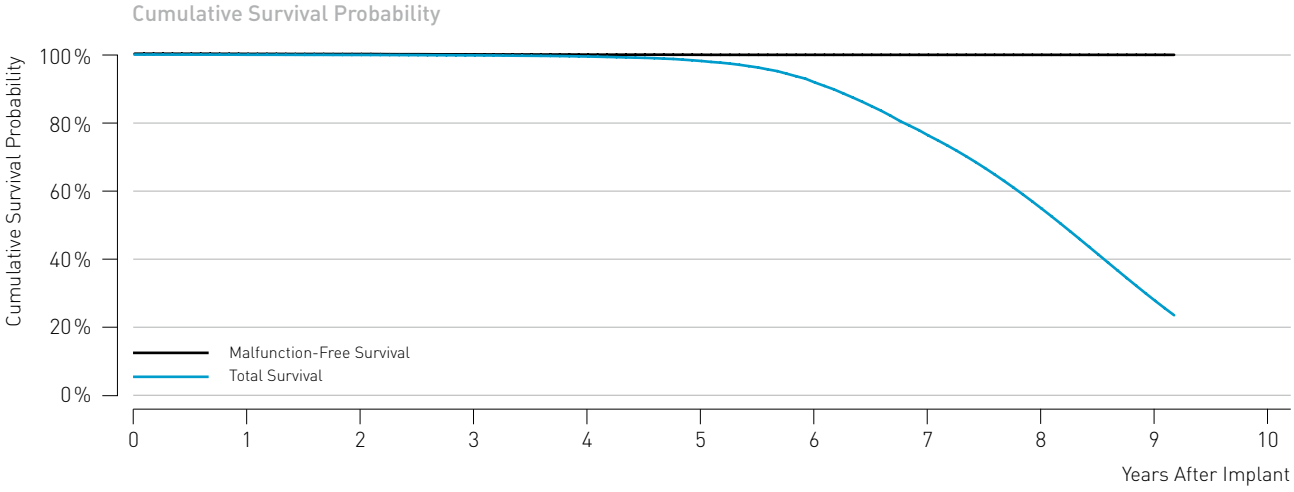
Cylos and Cylos 990

Product Details

Product Versions*	DR, DR-T
NBG Code(s)	DDDR
U.S. Market Release	Jan 2006
CE Market Release	Nov 2005 / Mar 2008
Worldwide Distributed Devices	81 300
Registered U.S. Implants	30 400
Estimated Active U.S. Implants	11 800
U.S. Normal Battery Depletions	5 654

	Quantity	Rate
U.S. Confirmed Malfunctions	27	0.09%
▪ Therapy Compromised	7	0.02%
▪ Therapy Available	20	0.07%

* While Cylos 990 DR and Cylos 990 DR-T are not distributed in the U.S., the performance is expected to be similar to the U.S. distributed products



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.
Total Survival [%]	100.0	99.9	99.9	99.8	99.4	98.1	91.8	76.3	54.8	27.8
(95% Confidence Interval)				±0.1	±0.1	±0.2	±0.4	±0.6	±0.7	±0.6
Malfunction-Free Survival [%]	100.0	100.0	100.0	99.9	99.9	99.9	99.9	99.9	99.9	99.9
(95% Confidence Interval)										

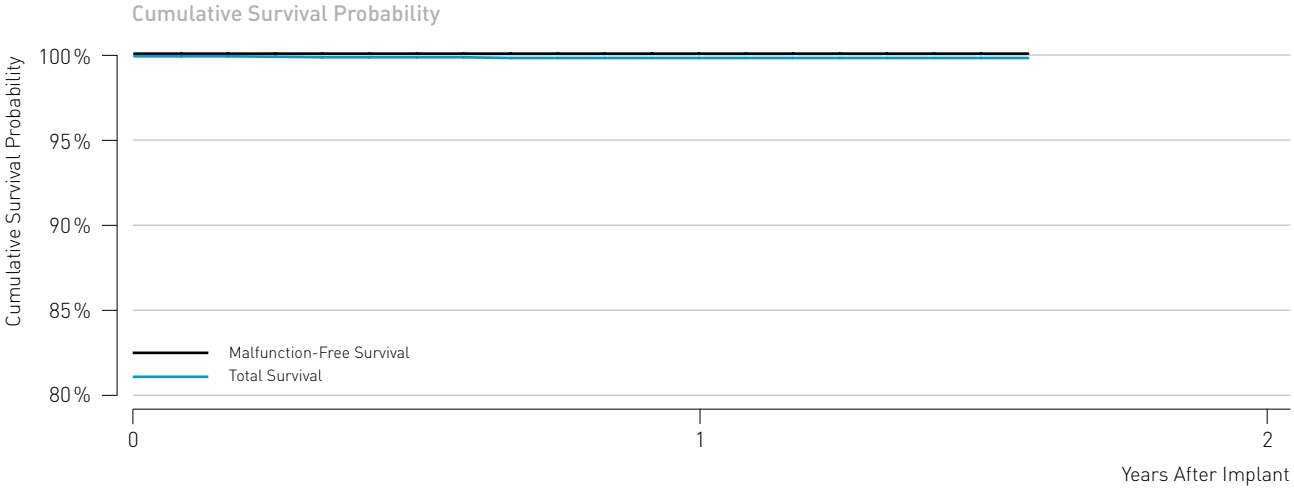
4.2 Dual-Chamber Pacemakers

Eluna 8

Product Details

Product Versions	DR, DR-T
NBG Code(s)	DDDR
U.S. Market Release	Dec 2014
CE Market Release	Aug 2014
Worldwide Distributed Devices	61 700
Registered U.S. Implants	23 100
Estimated Active U.S. Implants	22 300
U.S. Normal Battery Depletions	3

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

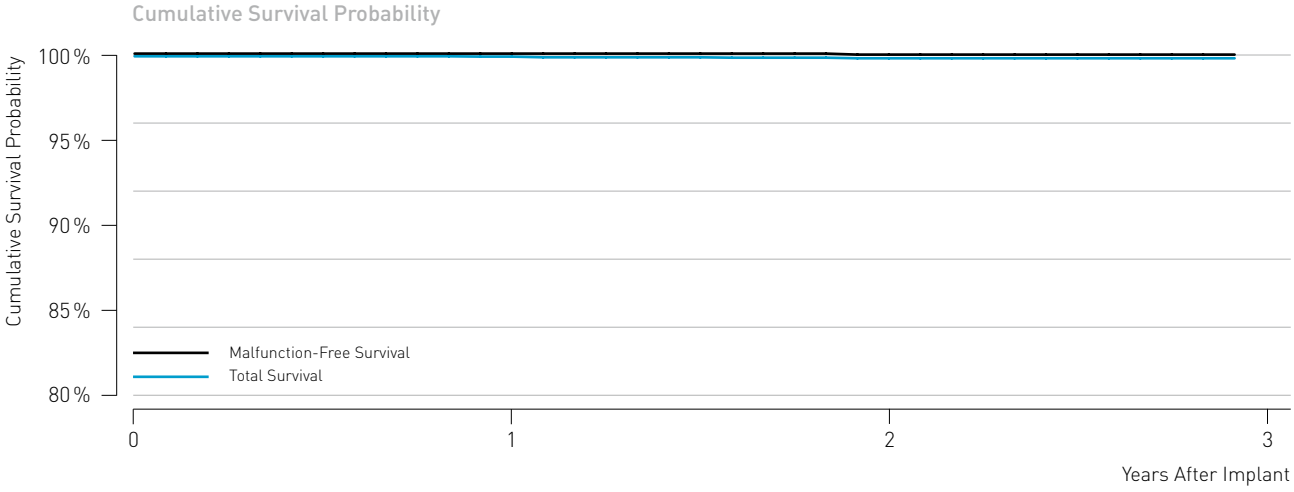
4.2 Dual-Chamber Pacemakers

Entovis

Product Details

Product Versions	DR, DR-T
NBG Code(s)	DDDR
U.S. Market Release	Feb 2010
CE Market Release	Nov 2009
Worldwide Distributed Devices	106 000
Registered U.S. Implants	12 200
Estimated Active U.S. Implants	10 700
U.S. Normal Battery Depletions	4

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.02%
▪ Therapy Compromised	1	0.01%
▪ Therapy Available	1	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	100.0	99.9
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

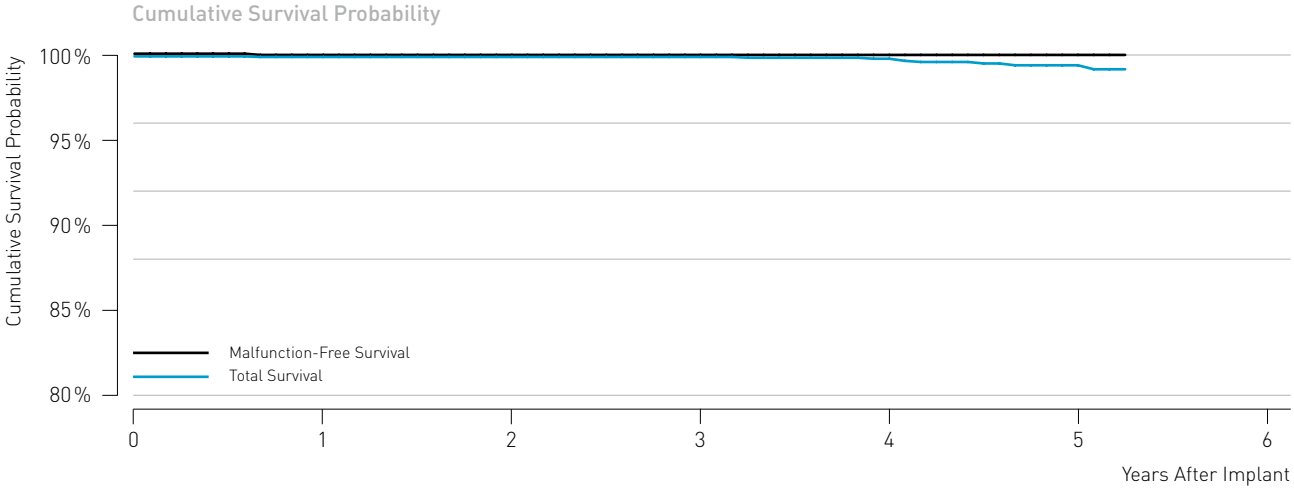
4.2 Dual-Chamber Pacemakers

Estella

Product Details

Product Versions	DR, DR-T
NBG Code(s)	DDDR
U.S. Market Release	Feb 2011
CE Market Release	Feb 2011
Worldwide Distributed Devices	28 100
Registered U.S. Implants	2 950
Estimated Active U.S. Implants	2 320
U.S. Normal Battery Depletions	8

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.07%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	2	0.07%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	99.9	99.5
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

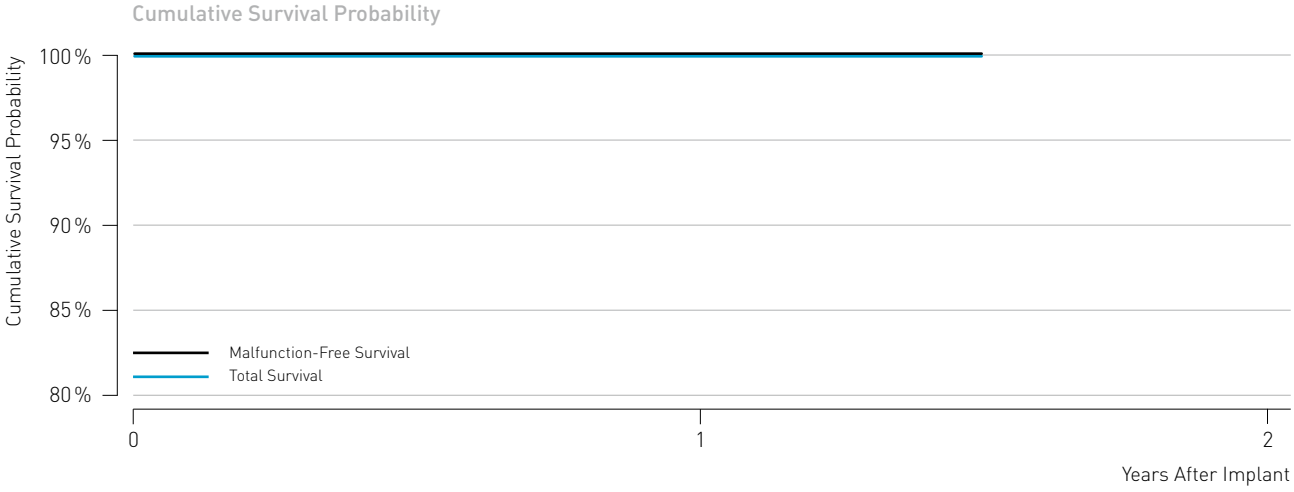
4.2 Dual-Chamber Pacemakers

Etrinsa 8

Product Details

Product Versions	DR-T
NBG Code(s)	DDDR
U.S. Market Release	Dec 2014
CE Market Release	Aug 2014
Worldwide Distributed Devices	50 700
Registered U.S. Implants	7 110
Estimated Active U.S. Implants	6 740
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

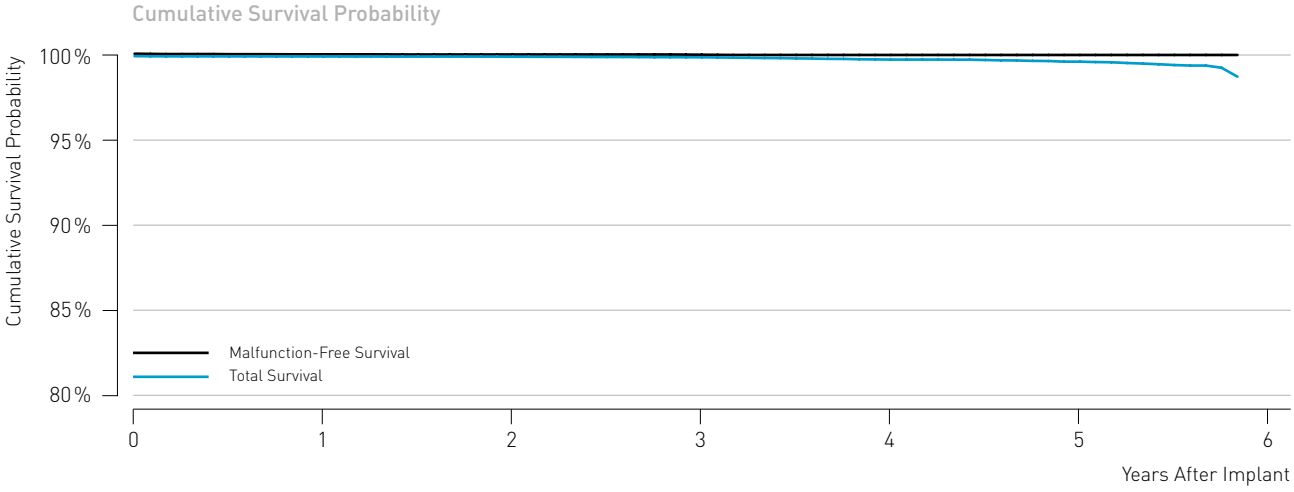
4.2 Dual-Chamber Pacemakers

Evia

Product Details

Product Versions	DR, DR-T
NBG Code(s)	DDDR
U.S. Market Release	May 2010
CE Market Release	Oct 2009
Worldwide Distributed Devices	195 000
Registered U.S. Implants	61 900
Estimated Active U.S. Implants	48 700
U.S. Normal Battery Depletions	100

	Quantity	Rate
U.S. Confirmed Malfunctions	18	0.03%
▪ Therapy Compromised	10	0.02%
▪ Therapy Available	8	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	100.0	100.0	99.9	99.8	99.7
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

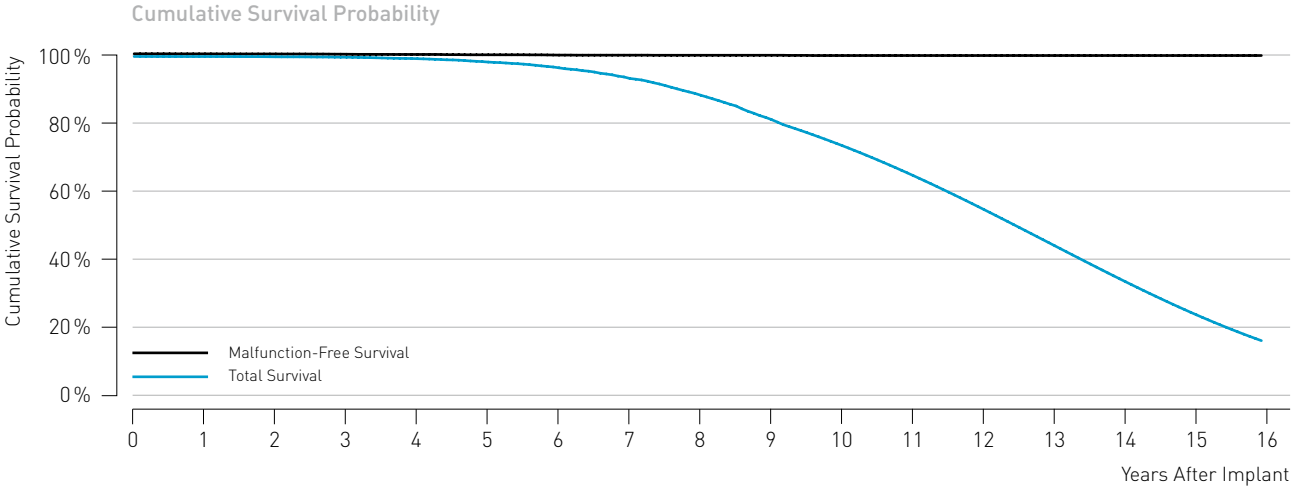
4.2 Dual-Chamber Pacemakers

Philos

Product Details

Product Versions	D, DR, DR-T, SLR
NBG Code(s)	DDD, DDDR, VDDR
U.S. Market Release	Sep 2000
CE Market Release	Aug 2000
Worldwide Distributed Devices	172 000
Registered U.S. Implants	20 700
Estimated Active U.S. Implants	5 410
U.S. Normal Battery Depletions	2 433

	Quantity	Rate
U.S. Confirmed Malfunctions	28	0.14%
▪ Therapy Compromised	5	0.02%
▪ Therapy Available	23	0.11%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.
Total Survival [%]	100.0	100.0	99.9	99.8	99.3	98.4	96.7	93.5	88.6	81.5	73.8	65.0	55.0	44.4	33.8	24.1
(95% Confidence Interval)				±0.1	±0.1	±0.2	±0.3	±0.4	±0.6	±0.8	±0.9	±0.8	±0.7	±0.6	±0.4	±0.3
Malfunction-Free Survival [%]	100.0	100.0	100.0	99.9	99.9	99.9	99.8	99.8	99.8	99.8	99.8	99.8	99.8	99.8	99.8	99.8
(95% Confidence Interval)						±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

4.2 Dual-Chamber Pacemakers

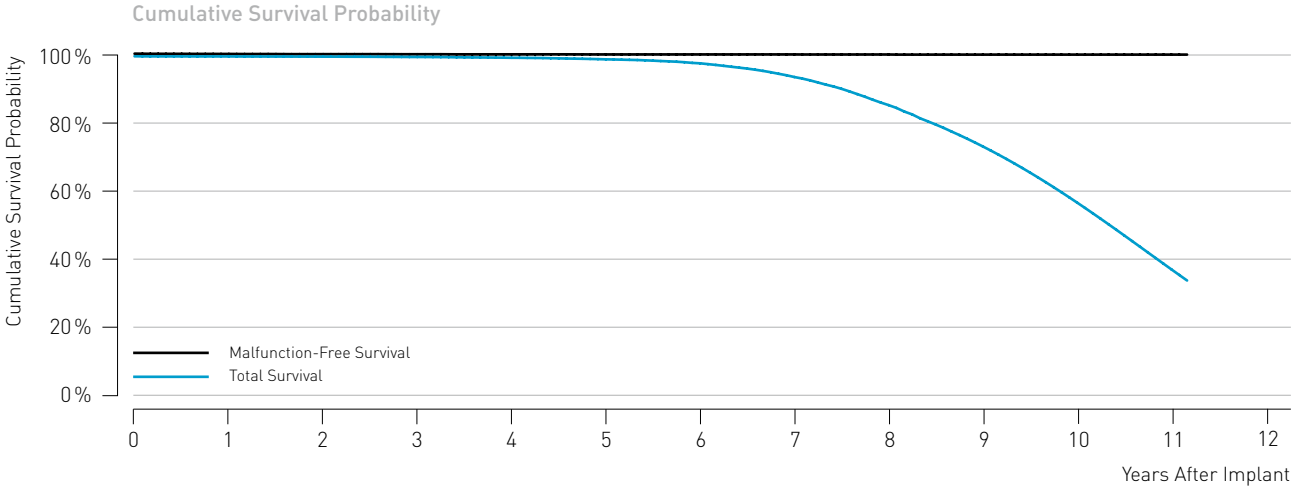
Philos II and Talos

Product Details

Product Versions*	D, DR, DR-T (Philos II only), SLR
NBG Code(s)	DDD, DDDR, VDDR
U.S. Market Release	Sep 2004
CE Market Release	Feb 2004 / May 2006
Worldwide Distributed Devices	366 000
Registered U.S. Implants	23 200
Estimated Active U.S. Implants	10 400
U.S. Normal Battery Depletions	2 830

	Quantity	Rate
U.S. Confirmed Malfunctions	21	0.09%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	21	0.09%

* While Philos II SLR, Talos D, Talos DR and Talos SLR are not distributed in the U.S., their performance is expected to be similar to the U.S. distributed products



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.
Total Survival [%]	100.0	100.0	99.9	99.8	99.6	99.1	97.9	93.9	85.6	73.4	56.9	37.2
(95% Confidence Interval)				±0.1	±0.1	±0.1	±0.2	±0.4	±0.6	±0.9	±1.0	±0.7
Malfunction-Free Survival [%]	100.0	100.0	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9
(95% Confidence Interval)										±0.1	±0.1	±0.1

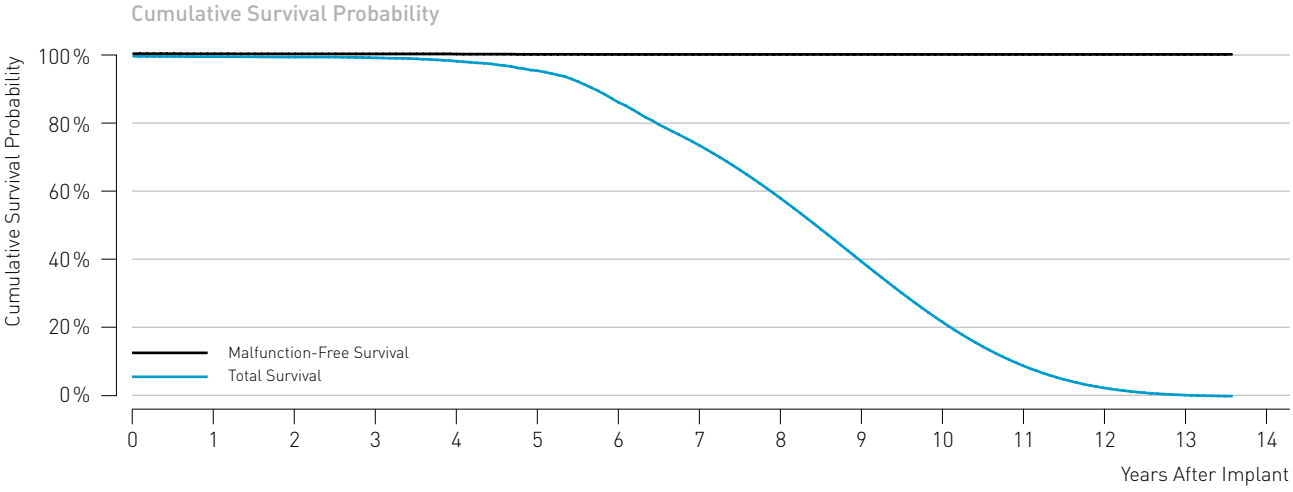
4.2 Dual-Chamber Pacemakers

Protos

Product Details

Product Versions	DR/CLS
NBG Code(s)	DDDR
U.S. Market Release	Jan 2003
CE Market Release	Jul 2003
Worldwide Distributed Devices	27 800
Registered U.S. Implants	10 800
Estimated Active U.S. Implants	2 540
U.S. Normal Battery Depletions	1 906

	Quantity	Rate
U.S. Confirmed Malfunctions	10	0.09%
▪ Therapy Compromised	2	0.02%
▪ Therapy Available	8	0.07%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.
Total Survival [%]	100.0	99.9	99.8	99.5	98.5	95.7	86.4	73.8	58.3	39.8	22	9.1	2.5	0.4
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.3	±0.5	±0.8	±1.1	±1.0	±0.7	±0.4	±0.2		
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9
(95% Confidence Interval)					±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

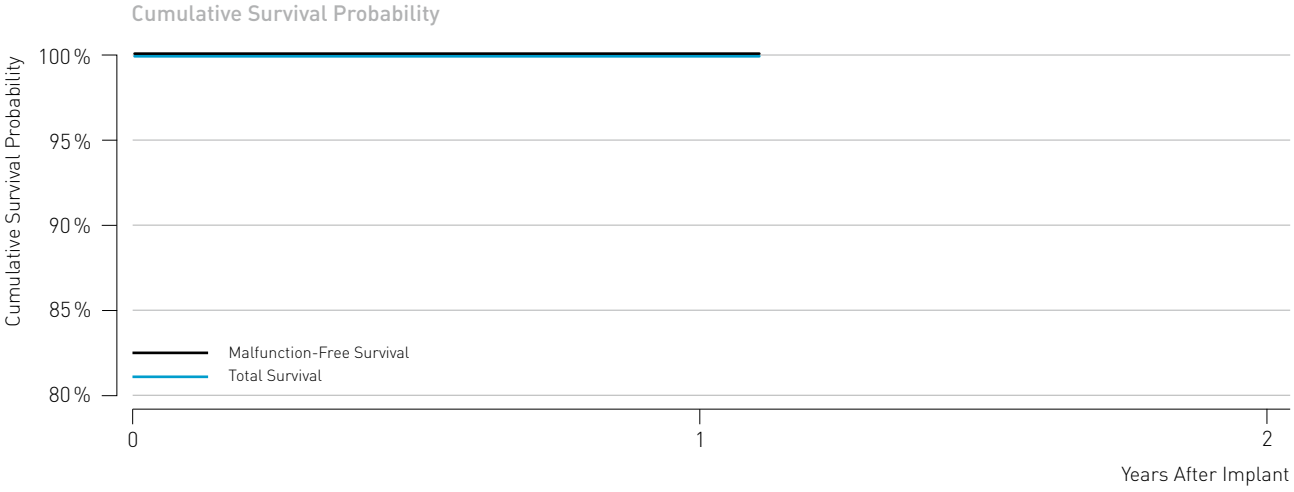
4.3 CRT Pacemakers

Etrinsa 8

Product Details

Product Versions	HF-T
NBG Code(s)	DDDRV
U.S. Market Release	Dec 2014
CE Market Release	Aug 2014
Worldwide Distributed Devices	36 300
Registered U.S. Implants	1 020
Estimated Active U.S. Implants	919
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

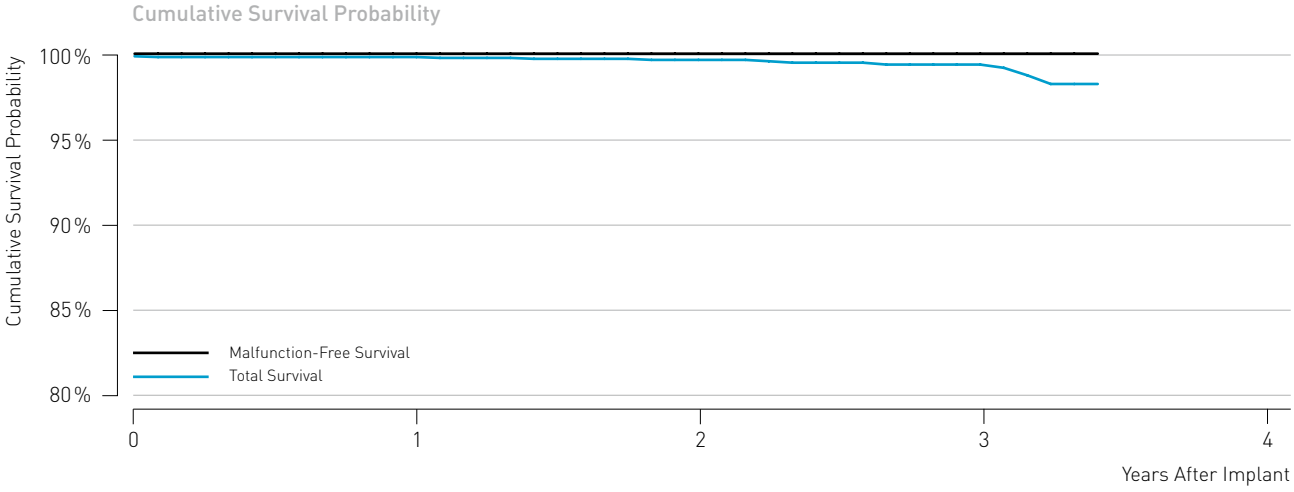
4.3 CRT Pacemakers

Evia

Product Details

Product Versions	HF, HF-T
NBG Code(s)	DDDRV
U.S. Market Release	May 2010
CE Market Release	Oct 2009
Worldwide Distributed Devices	8 450
Registered U.S. Implants	2 250
Estimated Active U.S. Implants	1 670
U.S. Normal Battery Depletions	13

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.
Total Survival [%]	100.0	100.0	99.8	99.5
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1

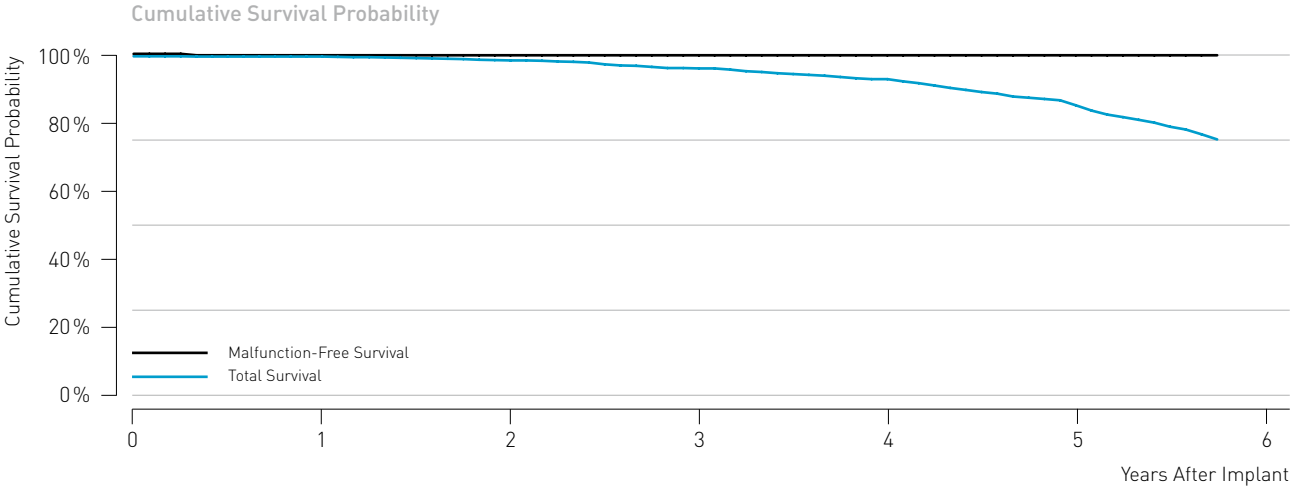
4.3 CRT Pacemakers

Stratos

Product Details

Product Versions	LV, LV-T
NBG Code(s)	DDDRV
U.S. Market Release	May 2008
CE Market Release	Nov 2002
Worldwide Distributed Devices	21 400
Registered U.S. Implants	1 310
Estimated Active U.S. Implants	484
U.S. Normal Battery Depletions	167

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.08%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	1	0.08%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	99.9	98.8	96.5	93.4	85.8
(95% Confidence Interval)		±0.2	±0.7	±1.2	±1.6	±2.6
Malfunction-Free Survival [%]	100.0	99.9	99.9	99.9	99.9	99.9
(95% Confidence Interval)		±0.2	±0.2	±0.2	±0.2	±0.2

5 Performance of BIOTRONIK ICDs

5.1 Single-Chamber ICDs

5.2 Dual-Chamber ICDs

5.3 CRT ICDs

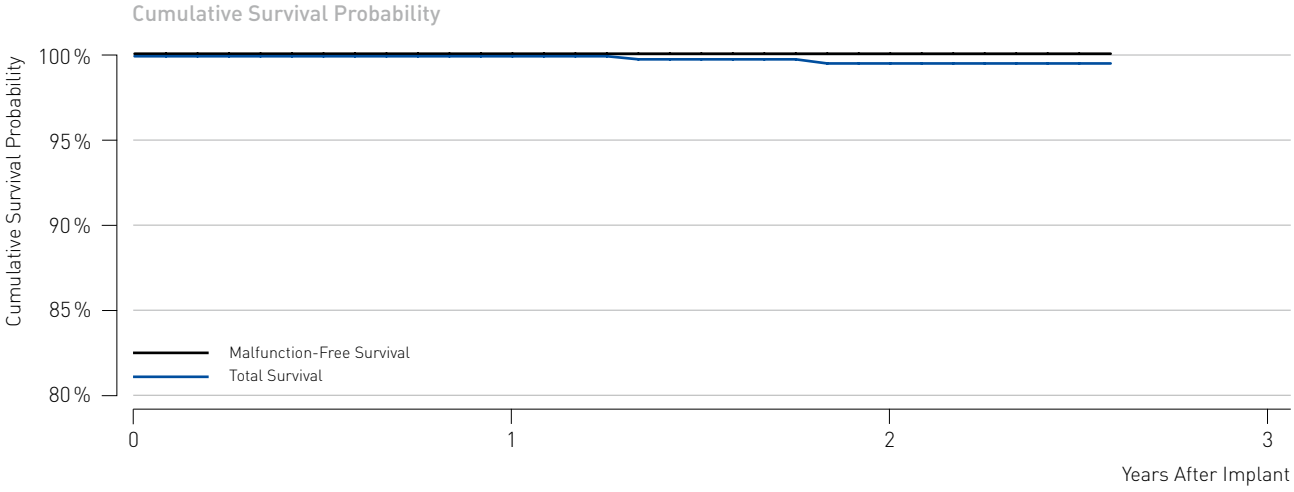
5.1 Single-Chamber ICDs

Ilesto 7

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VVIR
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	2 490
Registered U.S. Implants	1 270
Estimated Active U.S. Implants	1 120
U.S. Normal Battery Depletions	2

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	100.0	99.8
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

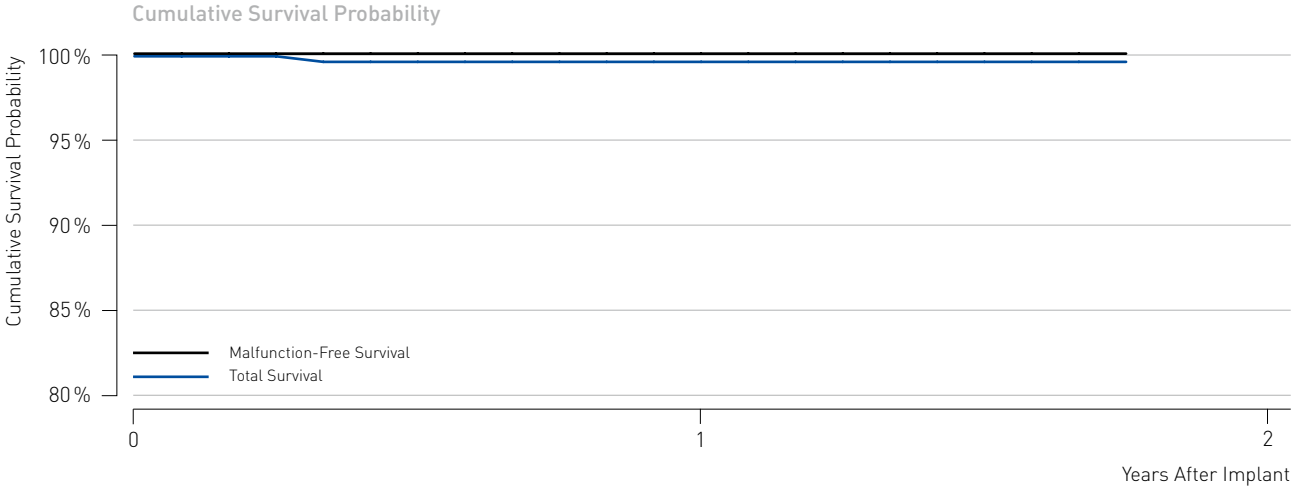
5.1 Single-Chamber ICDs

Ilesto 7 DF4

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VVIR
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	2420
Registered U.S. Implants	466
Estimated Active U.S. Implants	415
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	99.8
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

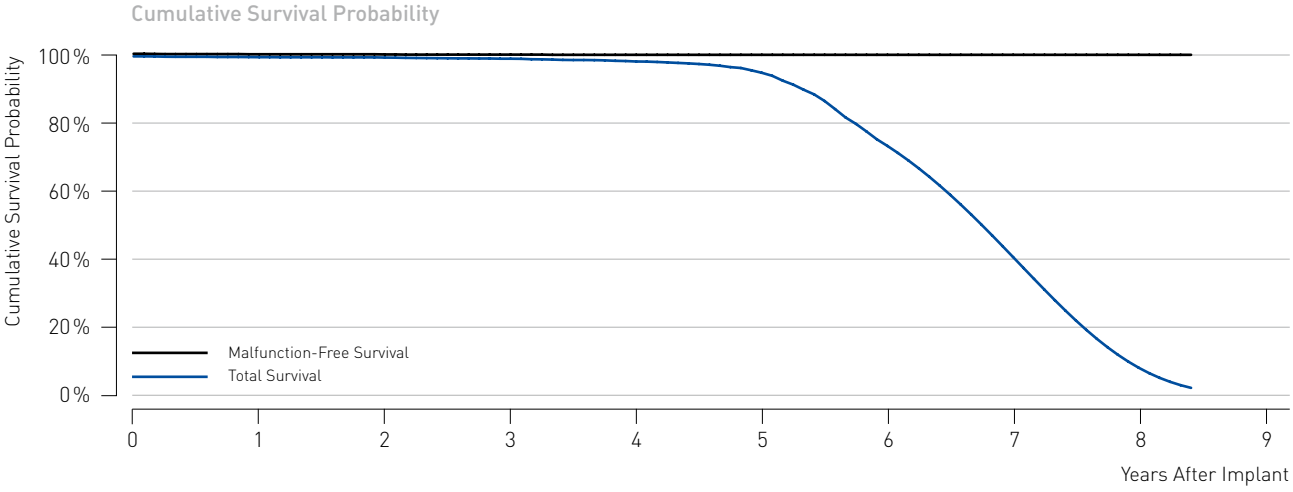
5.1 Single-Chamber ICDs

Lumax 340

Product Details

Product Versions	VR, VR-T
NBG Code(s)	VVE-VVIR
Maximum Energy [J]	40
U.S. Market Release	Feb 2007
CE Market Release	Feb 2007
Worldwide Distributed Devices	27 100
Registered U.S. Implants	3 990
Estimated Active U.S. Implants	1 210
U.S. Normal Battery Depletions	706

	Quantity	Rate
U.S. Confirmed Malfunctions	6	0.15%
▪ Therapy Compromised	4	0.10%
▪ Therapy Available	2	0.05%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.
Total Survival [%]	100.0	99.8	99.6	99.3	98.5	95.2	73.6	40.9	8.3
(95% Confidence Interval)		±0.2	±0.2	±0.3	±0.4	±0.8	±1.8	±1.2	±0.3
Malfunction-Free Survival [%]	100.0	99.9	99.9	99.9	99.8	99.8	99.8	99.8	99.8
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

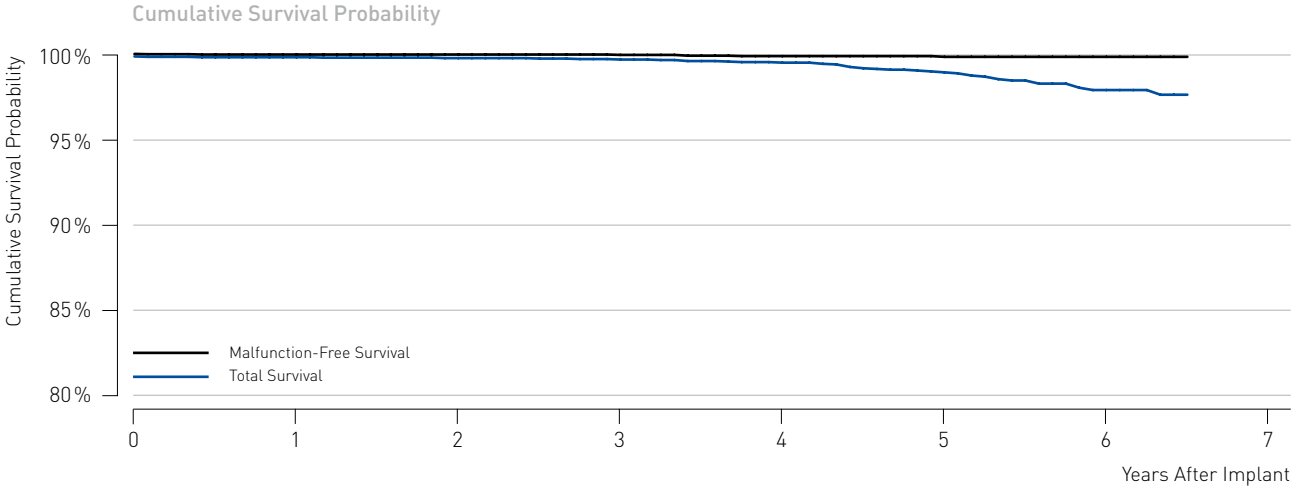
5.1 Single-Chamber ICDs

Lumax 540

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VVIR
Maximum Energy [J]	40
U.S. Market Release	May 2009
CE Market Release	Jun 2008
Worldwide Distributed Devices	19 700
Registered U.S. Implants	4 550
Estimated Active U.S. Implants	3 080
U.S. Normal Battery Depletions	36

	Quantity	Rate
U.S. Confirmed Malfunctions	7	0.15%
▪ Therapy Compromised	4	0.09%
▪ Therapy Available	3	0.07%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.
Total Survival [%]	100.0	99.9	99.9	99.8	99.6	99.1	98.1
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	99.9	99.8	99.8	99.8
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

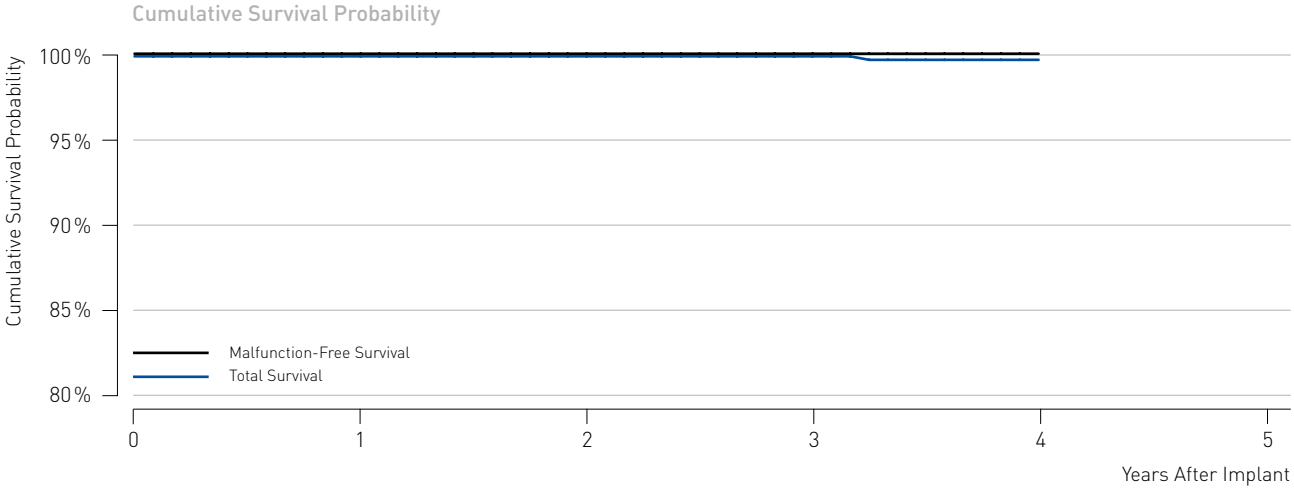
5.1 Single-Chamber ICDs

Lumax 740

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VVIR
Maximum Energy [J]	40
U.S. Market Release	Sep 2012
CE Market Release	Apr 2012
Worldwide Distributed Devices	4 770
Registered U.S. Implants	1 580
Estimated Active U.S. Implants	1 240
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	99.9
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1

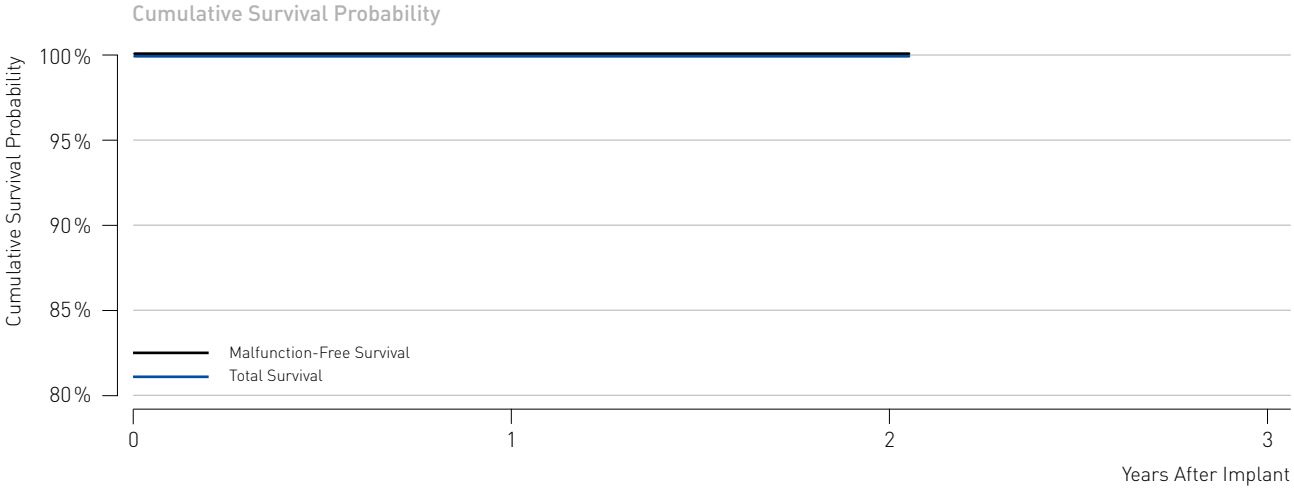
5.2 Dual-Chamber ICDs

Iforia 7

Product Details

Product Versions	DR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	939
Registered U.S. Implants	614
Estimated Active U.S. Implants	536
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

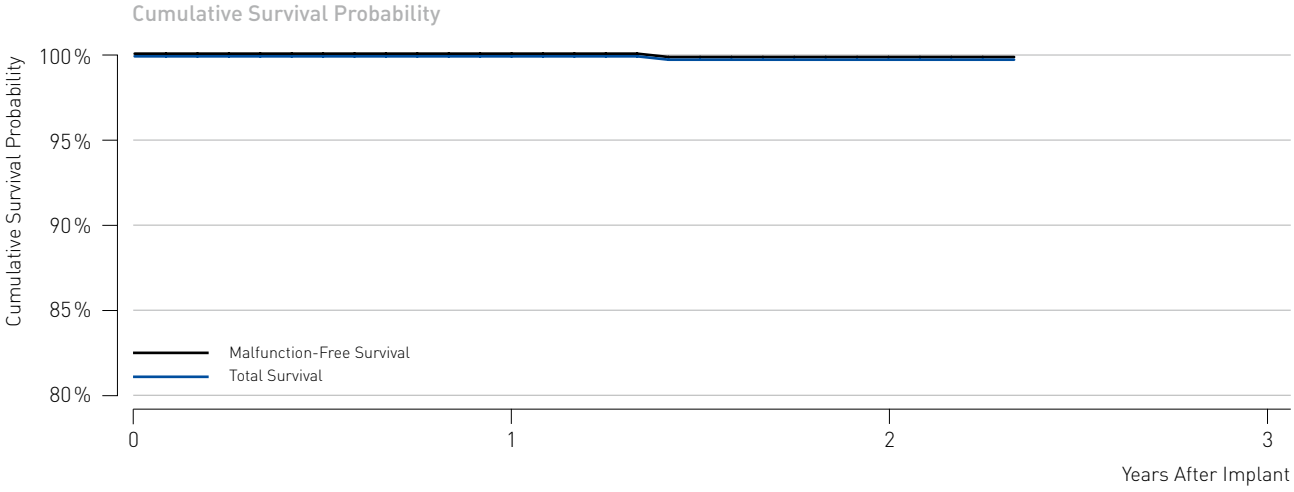
5.2 Dual-Chamber ICDs

Iforia 7 DX

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	2360
Registered U.S. Implants	1460
Estimated Active U.S. Implants	1300
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.14%
▪ Therapy Compromised	2	0.14%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	99.9	99.9	99.8
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	99.9	99.9	99.8
(95% Confidence Interval)	±0.1	±0.1	±0.1

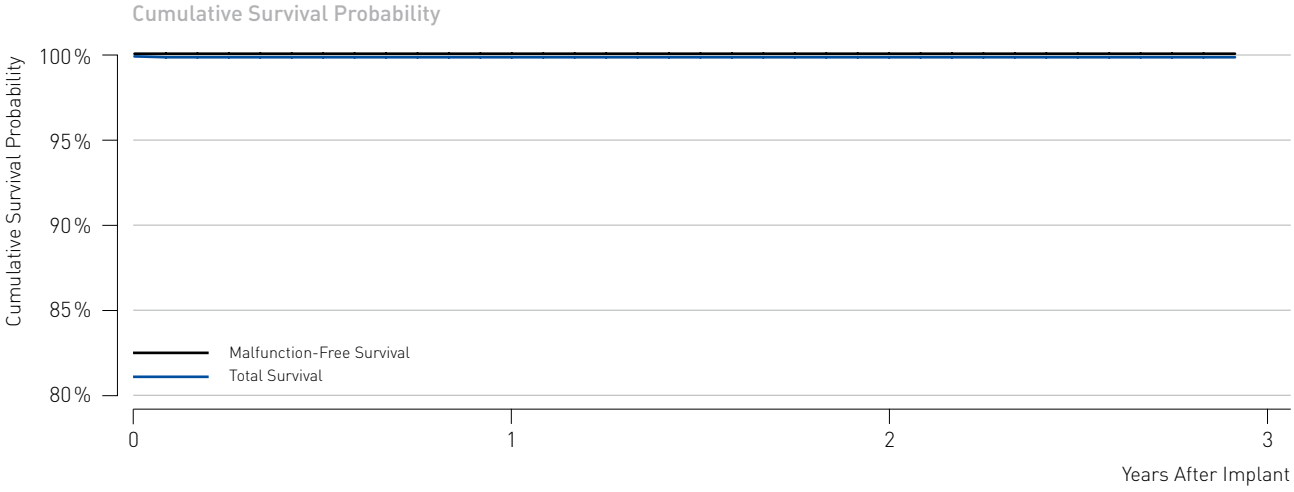
5.2 Dual-Chamber ICDs

Ilesto 7

Product Details

Product Versions	DR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	5 140
Registered U.S. Implants	3 490
Estimated Active U.S. Implants	2 970
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

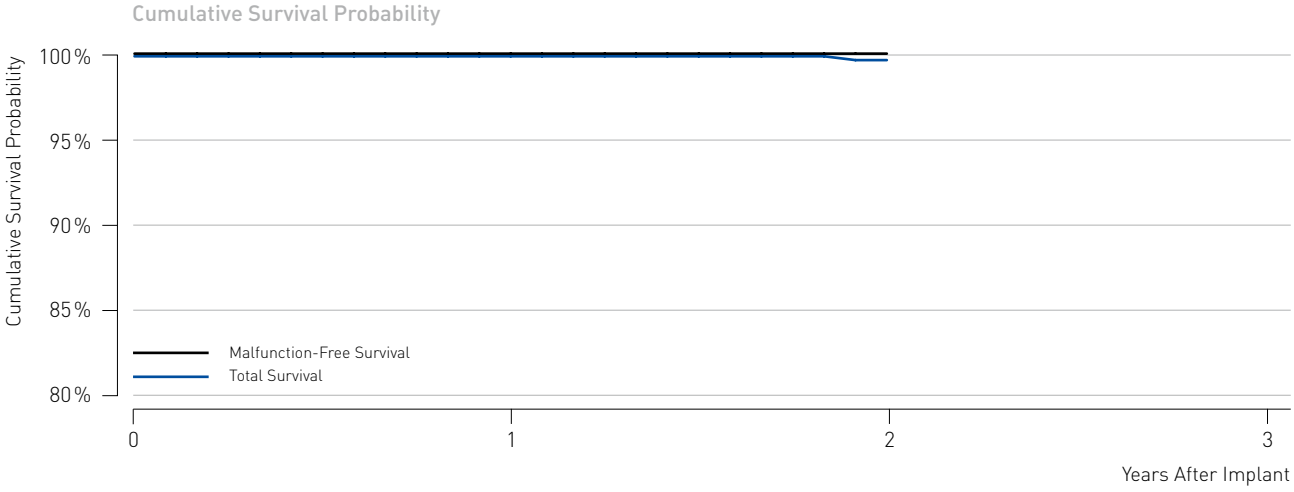
5.2 Dual-Chamber ICDs

Ilesto 7 DF4

Product Details

Product Versions	DR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Jul 2014
CE Market Release	Jun 2013
Worldwide Distributed Devices	3 780
Registered U.S. Implants	1 150
Estimated Active U.S. Implants	1 030
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	100.0	99.7
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

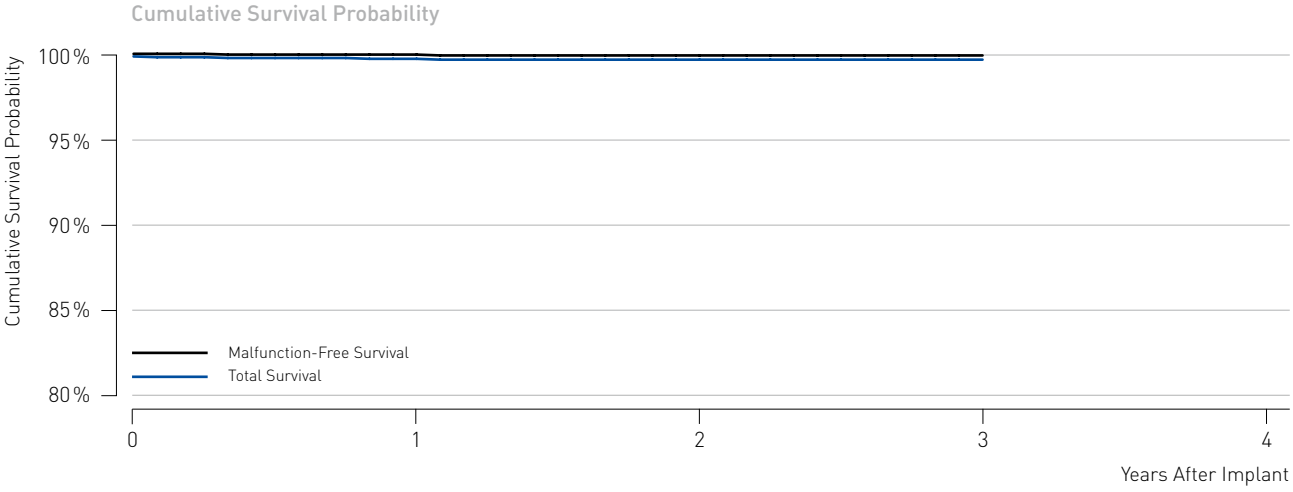
5.2 Dual-Chamber ICDs

Ilesto 7 DX

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	6 620
Registered U.S. Implants	4 720
Estimated Active U.S. Implants	4 160
U.S. Normal Battery Depletions	3

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.04%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	2	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.
Total Survival [%]	100.0	99.9	99.9	99.9
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1

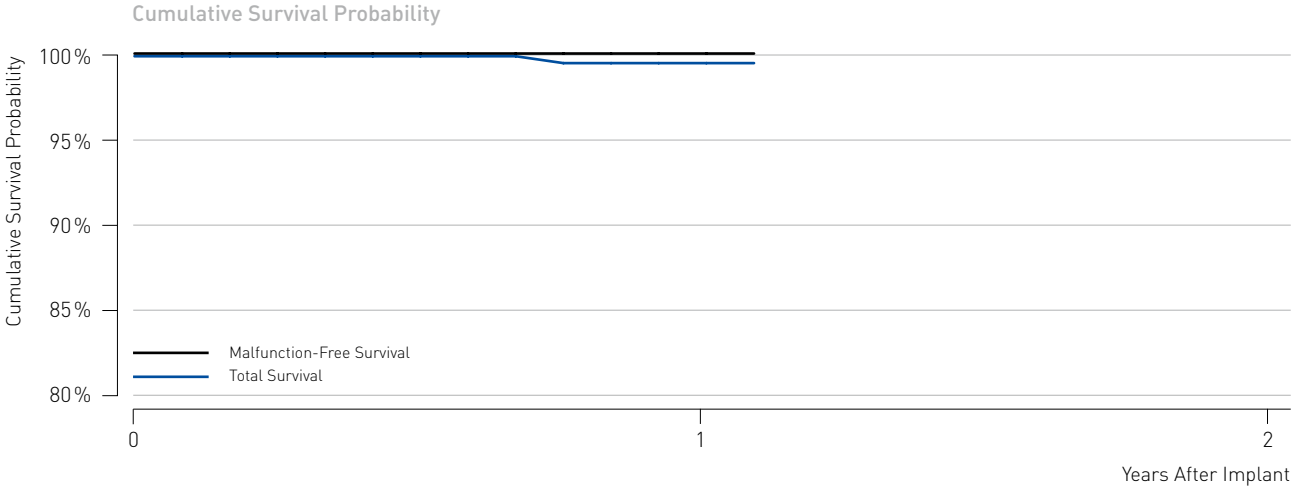
5.2 Dual-Chamber ICDs

Inventra 7 DX

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	45
U.S. Market Release	Mar 2015
CE Market Release	Dec 2014
Worldwide Distributed Devices	2 010
Registered U.S. Implants	1 440
Estimated Active U.S. Implants	1 370
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	99.8
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

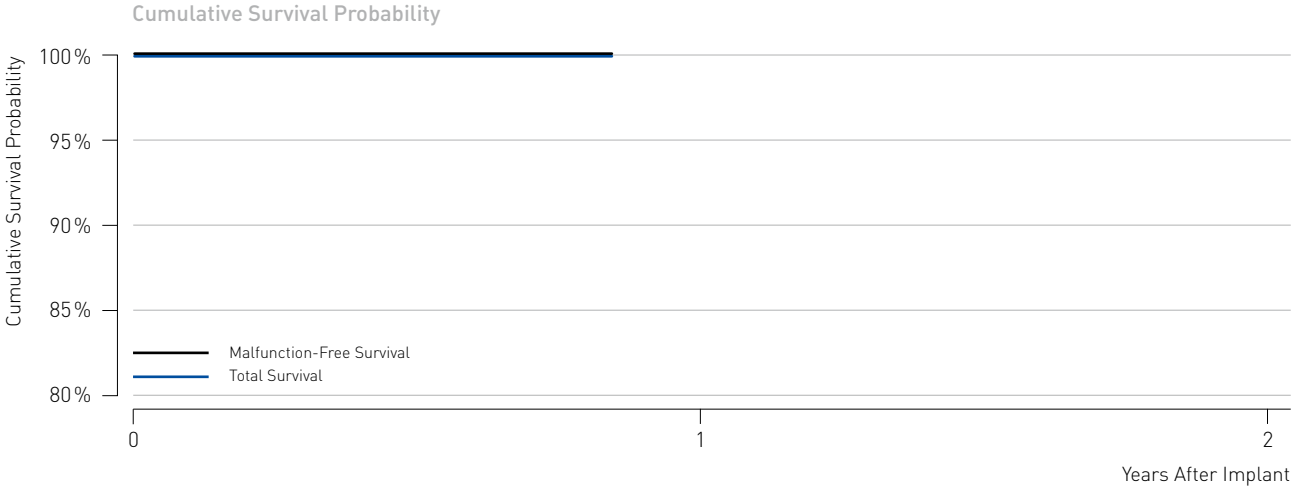
5.2 Dual-Chamber ICDs

Iperia 7 DF4

Product Details

Product Versions	DR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Dec 2015
CE Market Release	Dec 2014
Worldwide Distributed Devices	5 070
Registered U.S. Implants	1 680
Estimated Active U.S. Implants	1 650
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
■ Therapy Compromised	0	0.00%
■ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.
Total Survival [%]	100.0
(95% Confidence Interval)	±0.1
Malfunction-Free Survival [%]	100.0
(95% Confidence Interval)	±0.1

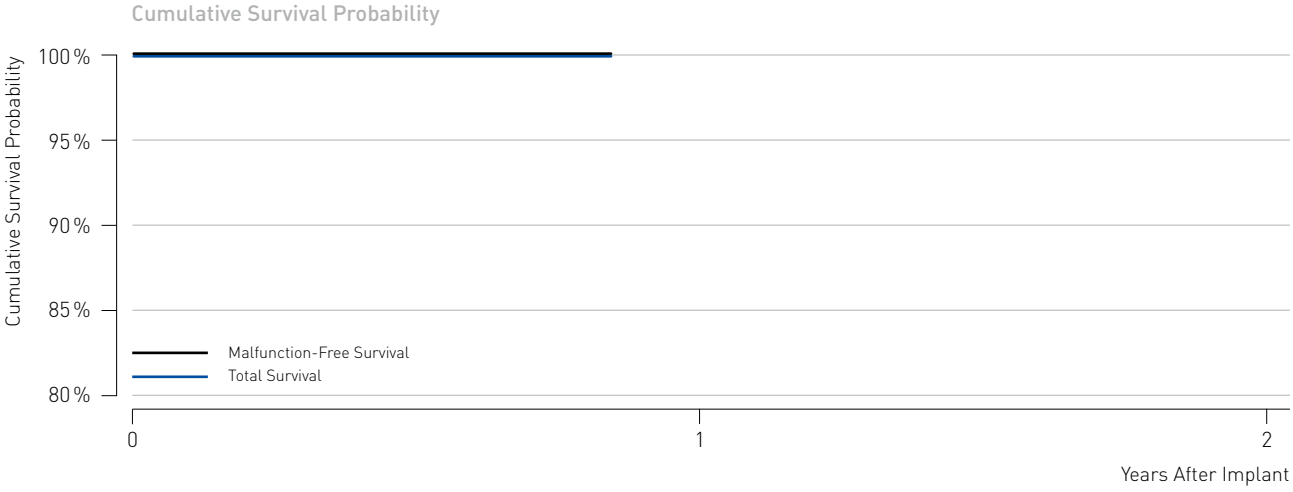
5.2 Dual-Chamber ICDs

Iperia 7 DX

Product Details

Product Versions	VR-T
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	Dec 2015
CE Market Release	Dec 2014
Worldwide Distributed Devices	4 670
Registered U.S. Implants	2 180
Estimated Active U.S. Implants	2 120
U.S. Normal Battery Depletions	0

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.
Total Survival [%]	100.0
(95% Confidence Interval)	±0.1
Malfunction-Free Survival [%]	100.0
(95% Confidence Interval)	±0.1

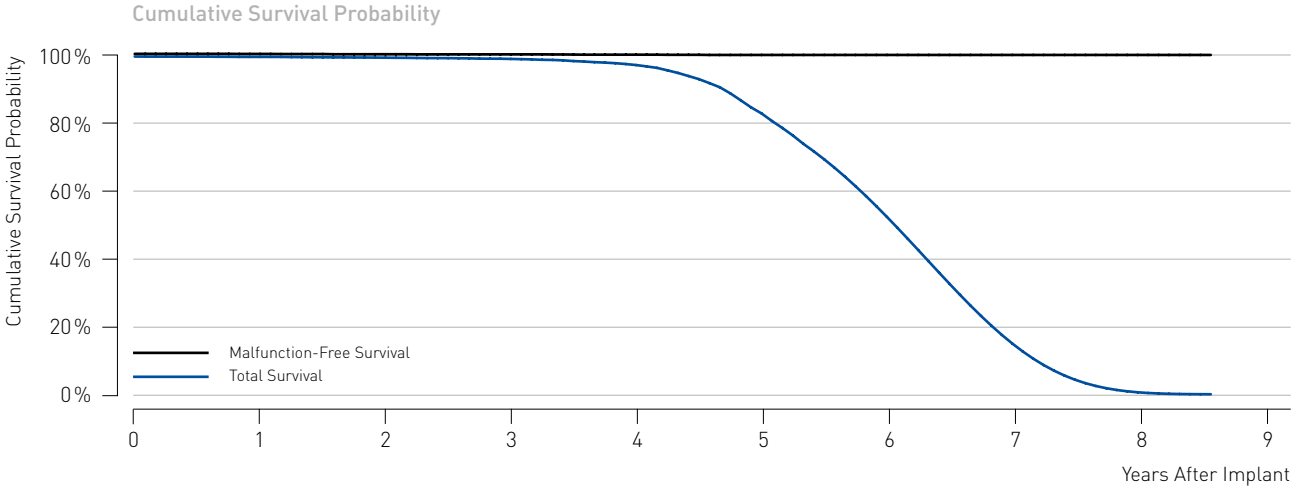
5.2 Dual-Chamber ICDs

Lumax 340

Product Details

Product Versions	DR, DR-T
NBG Code(s)	VVE-DDDR
Maximum Energy [J]	40
U.S. Market Release	Feb 2007
CE Market Release	Feb 2007
Worldwide Distributed Devices	26 400
Registered U.S. Implants	8 220
Estimated Active U.S. Implants	2 150
U.S. Normal Battery Depletions	1 788

	Quantity	Rate
U.S. Confirmed Malfunctions	10	0.12%
▪ Therapy Compromised	8	0.10%
▪ Therapy Available	2	0.02%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.
Total Survival [%]	100.0	99.9	99.7	99.3	97.5	83.2	52.5	15.0	0.6
(95% Confidence Interval)		±0.1	±0.1	±0.2	±0.4	±1.0	±1.1	±0.3	
Malfunction-Free Survival [%]	100.0	100.0	99.9	99.9	99.9	99.8	99.8	99.8	99.8
(95% Confidence Interval)			±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

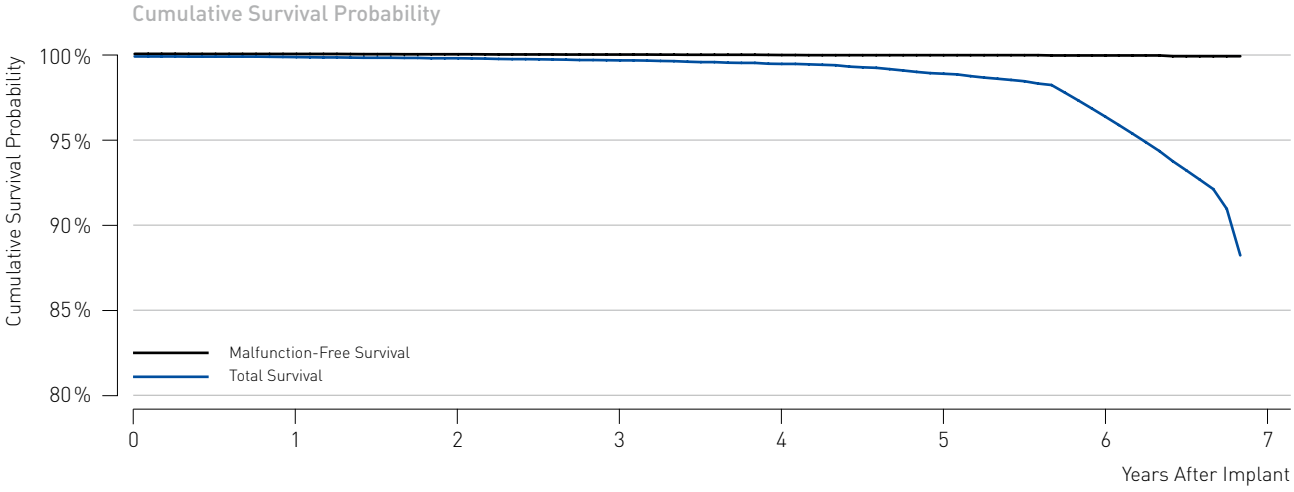
5.2 Dual-Chamber ICDs

Lumax 540

Product Details

Product Versions	DR-T
NBG Code(s)	VVE-DDDR
Maximum Energy [J]	40
U.S. Market Release	May 2009
CE Market Release	Jun 2008
Worldwide Distributed Devices	25 700
Registered U.S. Implants	11 600
Estimated Active U.S. Implants	7 490
U.S. Normal Battery Depletions	183

	Quantity	Rate
U.S. Confirmed Malfunctions	14	0.12%
▪ Therapy Compromised	8	0.07%
▪ Therapy Available	6	0.05%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.
Total Survival [%]	100.0	100.0	99.9	99.8	99.6	99.0	96.4
(95% Confidence Interval)			±0.1	±0.1	±0.1	±0.2	±0.5
Malfunction-Free Survival [%]	100.0	100.0	100.0	99.9	99.9	99.9	99.8
(95% Confidence Interval)				±0.1	±0.1	±0.1	±0.1

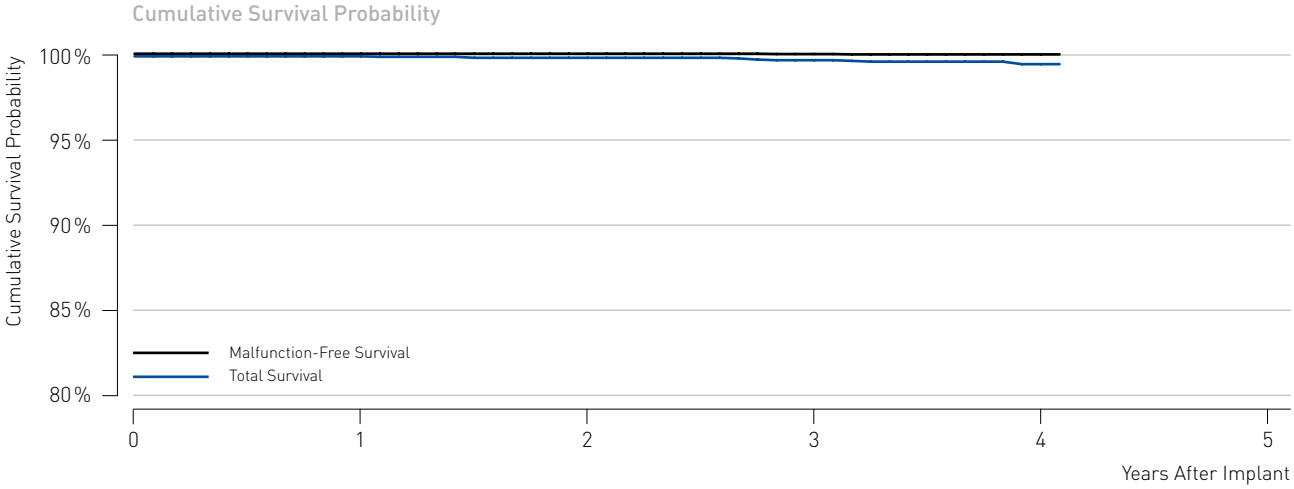
5.2 Dual-Chamber ICDs

Lumax 740

Product Details

Product Versions	DR-T
NBG Code(s)	VVE-DDDR
Maximum Energy [J]	40
U.S. Market Release	Sep 2012
CE Market Release	Apr 2012
Worldwide Distributed Devices	7 970
Registered U.S. Implants	3 820
Estimated Active U.S. Implants	3 040
U.S. Normal Battery Depletions	9

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.05%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	2	0.05%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	100.0	99.9	99.8	99.5
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0	99.9
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1	±0.1

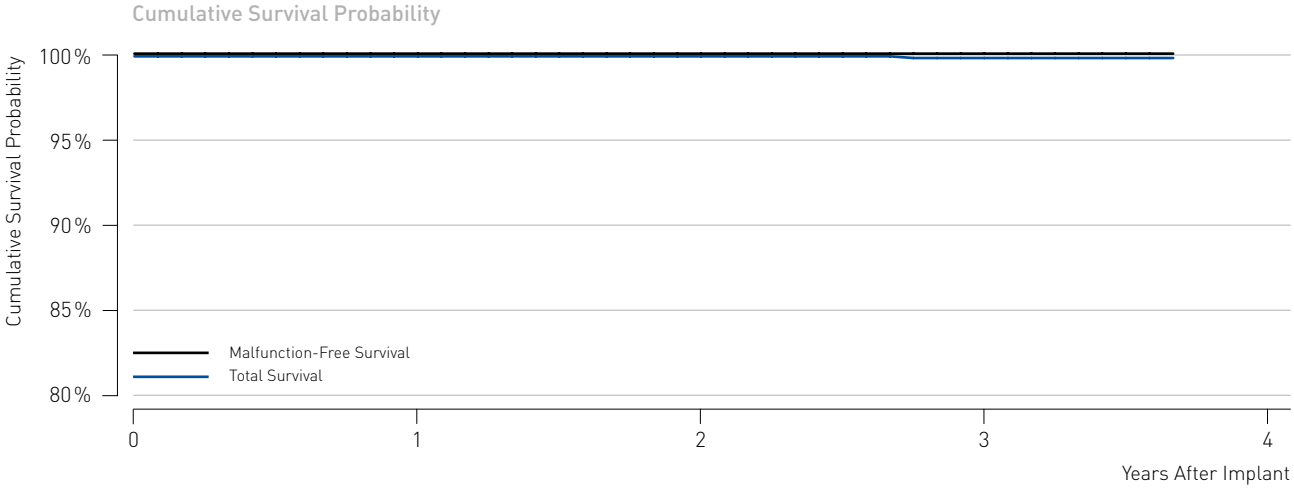
5.2 Dual-Chamber ICDs

Lumax 740 DX

Product Details

Product Versions	VR-T DX
NBG Code(s)	VVE-VDDR
Maximum Energy [J]	40
U.S. Market Release	May 2012
CE Market Release	Nov 2011
Worldwide Distributed Devices	4 560
Registered U.S. Implants	2 230
Estimated Active U.S. Implants	1 860
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.
Total Survival [%]	100.0	100.0	100.0	99.9
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1

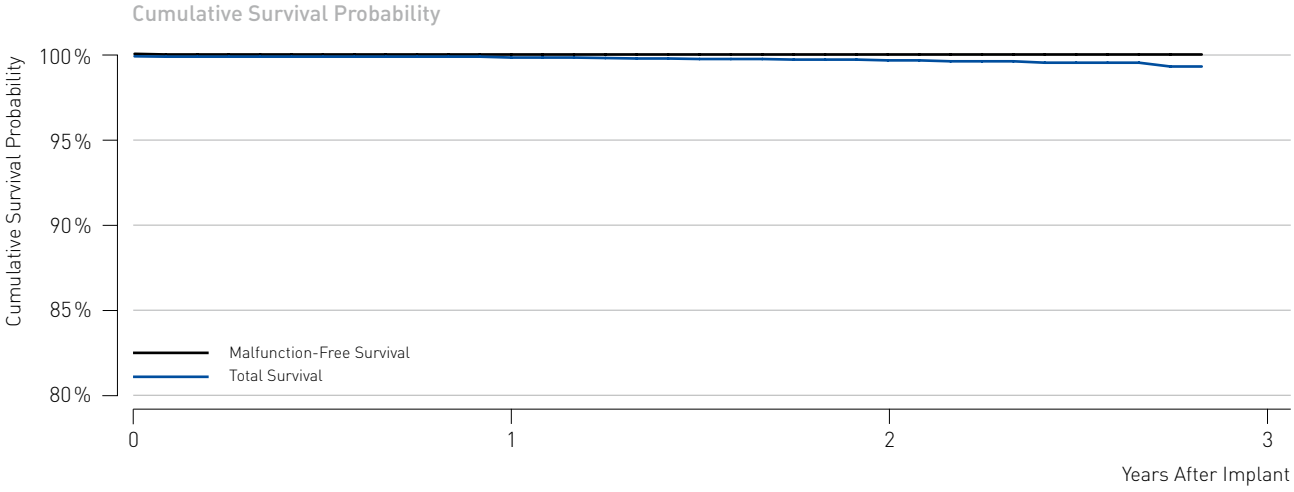
5.3 CRT ICDs

Ilesto 7

Product Details

Product Versions	HF-T
NBG Code(s)	VVE-DDDRV
Maximum Energy [J]	40
U.S. Market Release	Sep 2013
CE Market Release	Jun 2013
Worldwide Distributed Devices	5330
Registered U.S. Implants	3840
Estimated Active U.S. Implants	3060
U.S. Normal Battery Depletions	11

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.03%
▪ Therapy Compromised	1	0.03%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	99.9	99.7
(95% Confidence Interval)	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1	±0.1

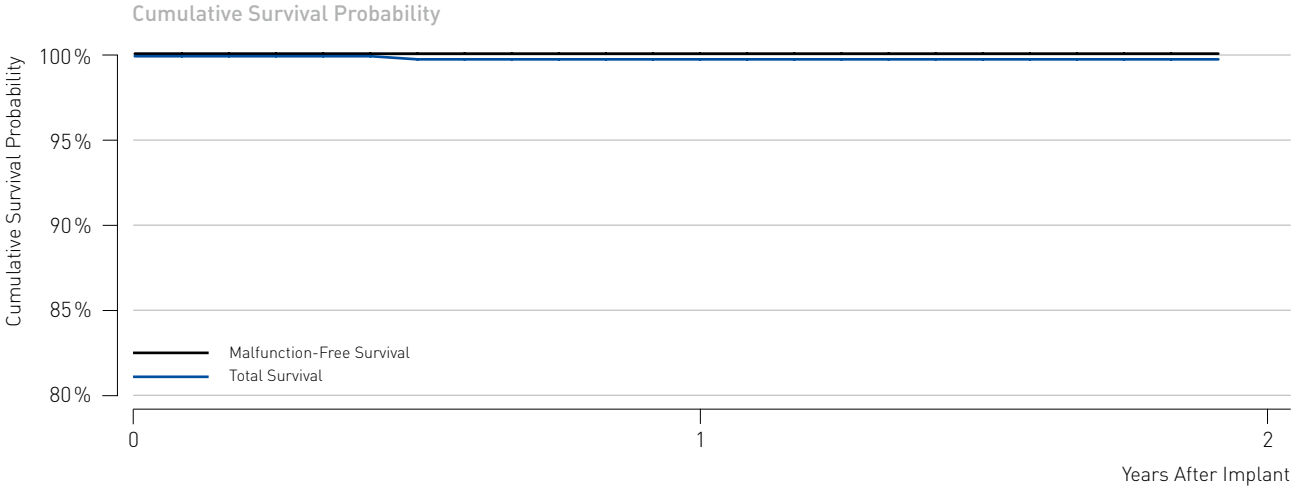
5.3 CRT ICDs

Ilesto 7 DF4

Product Details

Product Versions	HF-T
NBG Code(s)	VVE-DDDRV
Maximum Energy [J]	40
U.S. Market Release	Jul 2014
CE Market Release	Jun 2014
Worldwide Distributed Devices	2410
Registered U.S. Implants	967
Estimated Active U.S. Implants	831
U.S. Normal Battery Depletions	1

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.
Total Survival [%]	100.0	99.9
(95% Confidence Interval)	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0
(95% Confidence Interval)	±0.1	±0.1

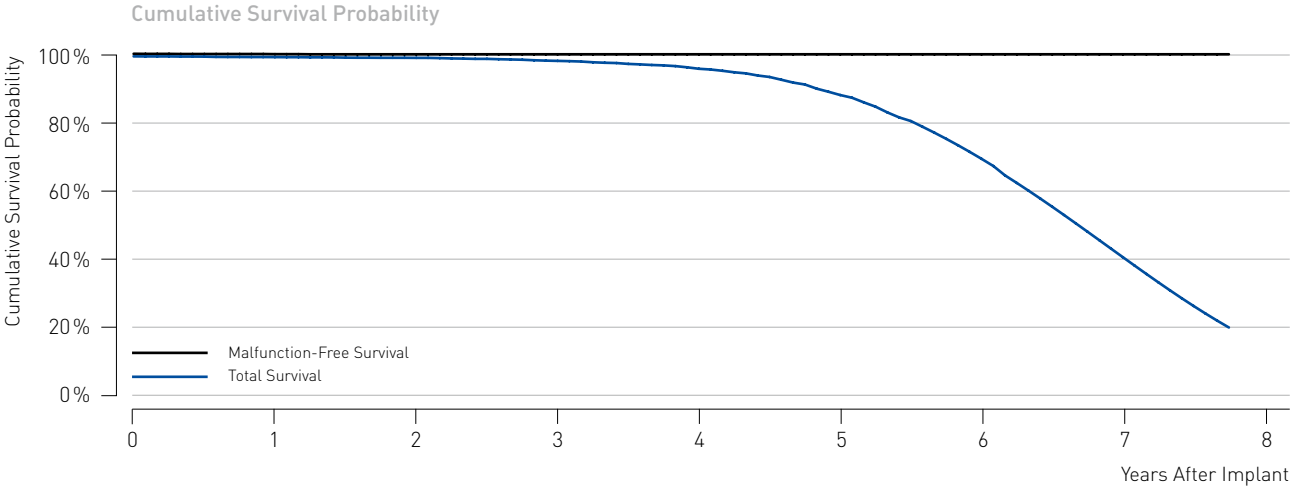
5.3 CRT ICDs

Lumax 340

Product Details

Product Versions	HF, HF-T
NBG Code(s)	VVE-DDDRV
Maximum Energy [J]	40
U.S. Market Release	Feb 2007
CE Market Release	Dec 2006
Worldwide Distributed Devices	20 700
Registered U.S. Implants	5 310
Estimated Active U.S. Implants	918
U.S. Normal Battery Depletions	1023

	Quantity	Rate
U.S. Confirmed Malfunctions	4	0.08%
▪ Therapy Compromised	2	0.04%
▪ Therapy Available	2	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.
Total Survival [%]	100.0	99.7	99.5	98.6	96.3	88.5	69.7	40.6
(95% Confidence Interval)	±0.1	±0.1	±0.2	±0.4	±0.6	±1.1	±1.7	±1.8
Malfunction-Free Survival [%]	100.0	99.9	99.9	99.9	99.9	99.9	99.9	99.9
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1

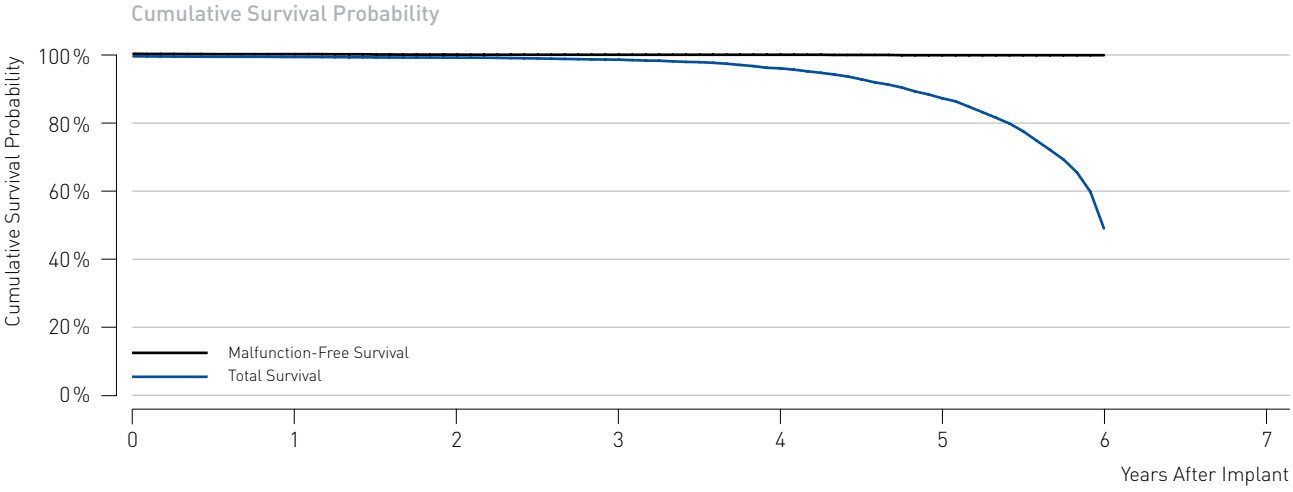
5.3 CRT ICDs

Lumax 540

Product Details

Product Versions	HF-T
NBG Code(s)	VVE-DDDRV
Maximum Energy [J]	40
U.S. Market Release	May 2009
CE Market Release	Jun 2008
Worldwide Distributed Devices	24 600
Registered U.S. Implants	8 660
Estimated Active U.S. Implants	3 610
U.S. Normal Battery Depletions	1 001

	Quantity	Rate
U.S. Confirmed Malfunctions	11	0.13%
▪ Therapy Compromised	5	0.06%
▪ Therapy Available	6	0.07%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.
Total Survival [%]	100.0	99.8	99.6	99	96.4	87.5	48.9
(95% Confidence Interval)		±0.1	±0.1	±0.2	±0.5	±1.0	±3.7
Malfunction-Free Survival [%]	100.0	100.0	99.9	99.9	99.9	99.8	99.8
(95% Confidence Interval)			±0.1	±0.1	±0.1	±0.1	±0.1

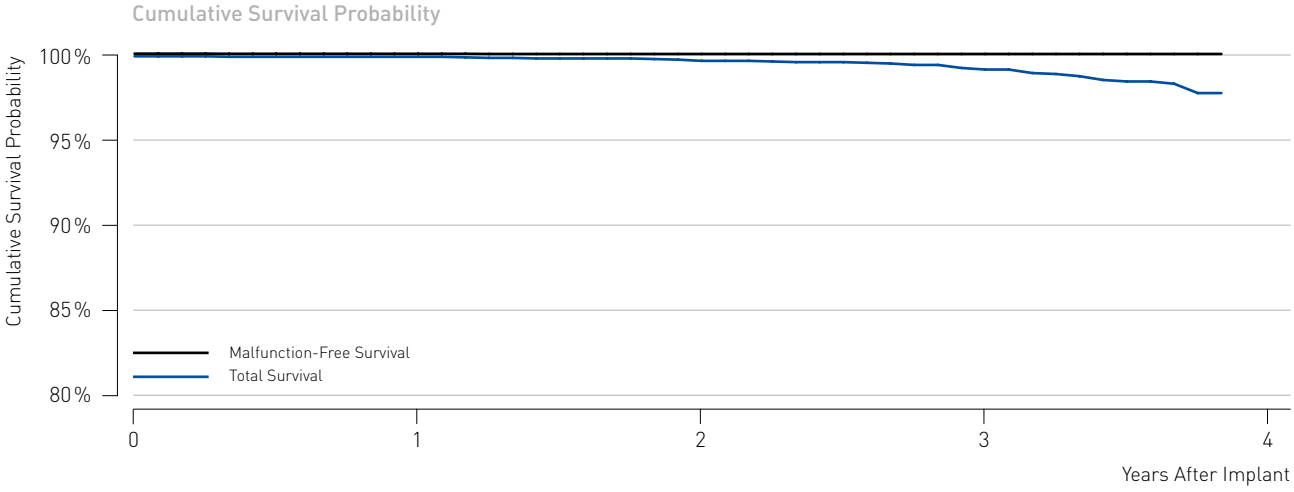
5.3 CRT ICDs

Lumax 740

Product Details

Product Versions	HF-T
NBG Code(s)	VVE-DDDRV
Maximum Energy [J]	40
U.S. Market Release	Sep 2012
CE Market Release	Apr 2012
Worldwide Distributed Devices	7030
Registered U.S. Implants	3410
Estimated Active U.S. Implants	2270
U.S. Normal Battery Depletions	34

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.06%
▪ Therapy Compromised	0	0.00%
▪ Therapy Available	2	0.06%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.
Total Survival [%]	100.0	100.0	99.7	99.2
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1
Malfunction-Free Survival [%]	100.0	100.0	99.9	99.9
(95% Confidence Interval)	±0.1	±0.1	±0.1	±0.1

6 Methodology for Lead Survival Estimates Based on Returned Product Analysis and Complaint Information

6.1 Cumulative Lead Survival Probability

This report has been prepared in accordance with ISO 5841-2:2014(E) applying actuarial analysis for the calculation of lead survival probabilities based on returned product analysis. Lead Survival Estimates given in this report are considered to be generally representative for worldwide performance of BIOTRONIK's pacing and ICD leads.

The Cumulative Survival Probability for leads is an estimate based on the percentage of devices that remain implanted and in service at various points of the product's service time in the absence of concurrent events such as morbidity. The Lead Survival Estimate over time is displayed in cumulative survival curves (Kaplan-Meier). The product's performance is evaluated in discrete one-month intervals. The survival probability for each month is given by the number of leads that remain implanted and active through this month divided by the number of leads that were actively implanted at the start of the interval. The cumulative survival probability for any period is given by multiplying all survival probabilities of previous months.

At the time of implantation, the cumulative lead survival probability is 100 %. Even though they are analyzed as part of our quality system monitoring, leads that are found to be out of specification prior to or during the implantation procedure are removed from the statistics as they do not contribute to a patient's risk of being subject to a device malfunction or replacement during the device's service time.

Because this report is provided to communicate information regarding product performance, it does not include data regarding medical complications such as erosion, infection or diaphragmatic stimulation.

Compared to pacemakers and ICDs, a considerable portion of leads with observed or suspected failures are not explanted and returned for laboratory analysis. This is primarily because it is much more difficult and risky to the patient to remove chronically implanted leads. In order to report a conservative measure of lead performance, unconfirmed reports of lead complications are therefore also included in the calculation of a lead's survival probability.

In order to be classified as a qualifying lead complication and thus contributing to the survival probability calculation the same way as a confirmed malfunction, the reported anomaly must have occurred at least 30 days post-implant. Otherwise, factors not related to the lead would likely be the root cause of the observed anomaly, (i.e., patientspecific conditions or implant techniques).

In order to minimize the effect of underreporting of lead malfunctions, BIOTRONIK additionally includes the long term performance post market study data if available.

6.2 Lead Data Acquisition

The following sections of this report provide performance data on BIOTRONIK's pacing and ICD leads and are based on the observation of BIOTRONIK's U.S. products through review of our device registration and tracking systems, and through the analyses of both, returned leads as well as reports of lead complications of non-returned leads. The prospective data from BIOTRONIK's post-approval studies is presented separately in chapters 8 and 9.

In order to be included in the population under observation a lead must be registered and implanted for at least one calendar day. The cut-off date for the data included in this report is December 31, 2016. The sample sizes of U.S. leads that are implanted and remain active as well as the total number of products distributed worldwide are provided for each lead family in this report.

Survival estimates are calculated for lead families having accumulated at least 10,000 cumulative implant months. Products no longer being distributed with less than 500 active implants may be excluded from this report.

ISO 5841-2:2014(E) describes a method for adjusting the device survival probability for underreported malfunctions and unrelated patient deaths that result in an overestimation of the device's survival probability. The factor for U.S. underreporting of malfunctions of pacing and ICD leads is unknown as currently no systematic data is available that reveals this factor. Consequently, this factor remains unaccounted for in this report. Patient mortality is artificially elevated if the reported rate from our registration and tracking systems is below the annual mortality in clinical studies.

6.3 Returned Product Analysis

Information for the lead sections of this report is taken from the analysis of returned products. The outcome of this analysis is the basis for the final classification of the cause for explantation of the lead. Additionally, reports of lead complications not confirmed by laboratory analysis are taken into consideration. Both, leads with confirmed malfunctions as well as unconfirmed lead complications decrease a lead's total survival probability.

Every lead and lead segment returned to BIOTRONIK is analyzed per our internal procedures and classified as within specification, damaged by external causes, or out of specification (malfunction) while implanted and in service.

Those leads found to be out of specification, are divided into the following categories as proposed by AdvaMed and ISO 5841-2:2014(E):

- Conductor Fracture – Conductor break with complete or intermittent loss of continuity that could interrupt current flow (e.g. fractured conductors)
- Crimps, Welds and Bonds – Any interruption in the conductor or lead body associated with a point of connection
- Insulation Breach – Any lead insulation breach
- Other – Includes specific proprietary lead mechanical attributes.

6.4 Lead Complications

A considerable portion of leads with observed or suspected failures are not explanted and returned for laboratory analysis. A clinical observation is considered a lead complication if a complaint, associated with at least one of the clinical manifestations listed below, is reported and where the non-returned lead is:

- Verified by medical records to have been implanted and in-service, and
- Reported to have been removed from service,
- Modified surgically or electrically to remedy the malfunction, or
- Left in service based on medical judgment.

Complications for leads implanted greater than 30 days are reported as Qualifying lead complications, whereas complications occurring during the first 30 days are reported as Acute Lead Observations.

In accordance with the latest AdvaMed guidelines and ISO 5841-2:2014(E) such clinical observations are classified in the following categories:

- Failure to Capture – Intermittent or complete failure to achieve cardiac stimulation at programmed output delivered outside of the cardiac refractory period. Sudden and significant increase in the pacing threshold value at which 2:1 safety margin can no longer be achieved.
- Failure to Sense – Intermittent or complete loss of sensing or failure to detect intended intrinsic cardiac signals during non-refractory periods at programmed sensitivity settings
- Oversensing – Misinterpretation of cardiac or non-cardiac events as cardiac depolarization
- Abnormal Pacing Impedance – Pacing impedance is typically considered abnormal if a measurement is < 200 ohms or > 3000 ohms
- Abnormal Defibrillation Impedance – Defibrillation impedance is typically considered abnormal if a measurement is < 20 ohms or > 200 ohms. Including high or low shock impedance when attempting to deliver a shock
- Insulation Breach – A disruption or break in lead insulation observed visually, electrically, or radiographically
- Conductor Fracture – A mechanical break within the lead conductor observed visually, electrically, or radiographically
- Lead Dislodgement – Radiographic, electrical or electrocardiographic evidence of electrode displacement from the original implant site or electrode displacement that adversely affects pacing and/or lead performance
- Extracardiac Stimulation – Clinical observation of inadvertent nerve/muscle stimulation other than cardiac muscle
- Cardiac Perforation – Penetration of the lead tip through the myocardium, clinically suspected and confirmed by chest x-ray, fluoroscopy, echocardiogram, or visual observation, which results in clinical symptoms, typically degradation of pacing/ICD lead electrical performance, chest pain, and tamponade
- Other – Specific proprietary attributes of a lead such as sensors which affect a lead's ability to perform as designed or remain in service

In order to report a conservative measure of lead performance, qualifying lead complications are also included in the calculation of a lead's survival probability.

Acute Lead Observations may be subject to a number of factors, including patient-specific anatomy, clinical conditions and/or varying implant conditions/ techniques. Therefore, acute lead observations are not included in lead survival probability.

6.5 Lead Product Performance Graphs and Data

The lead performance information is shown in each section in alphabetical order and by product name.

For each lead, the report provides:

- Product versions that contribute to the evaluation
- Types of leads
- Polarity
- Steroid
- CE and U.S. market release dates
- Worldwide quantity of products that have been distributed
- U.S. registered implants (number of products included in this report)
- Estimated active U.S. implants
- Number of U.S.qualifying complications
- Number of U.S. acute lead observations
- Number of U.S. confirmed malfunctions
- Number of U.S. leads or partial leads returned post-implant for analysis with a complaint

The survival plots provide:

Total Survival

The cumulative survival probability free of component malfunction or unconfirmed observation of an anomaly. Removals for clinical reasons unrelated to the device's performance (i.e., infections) are excluded.

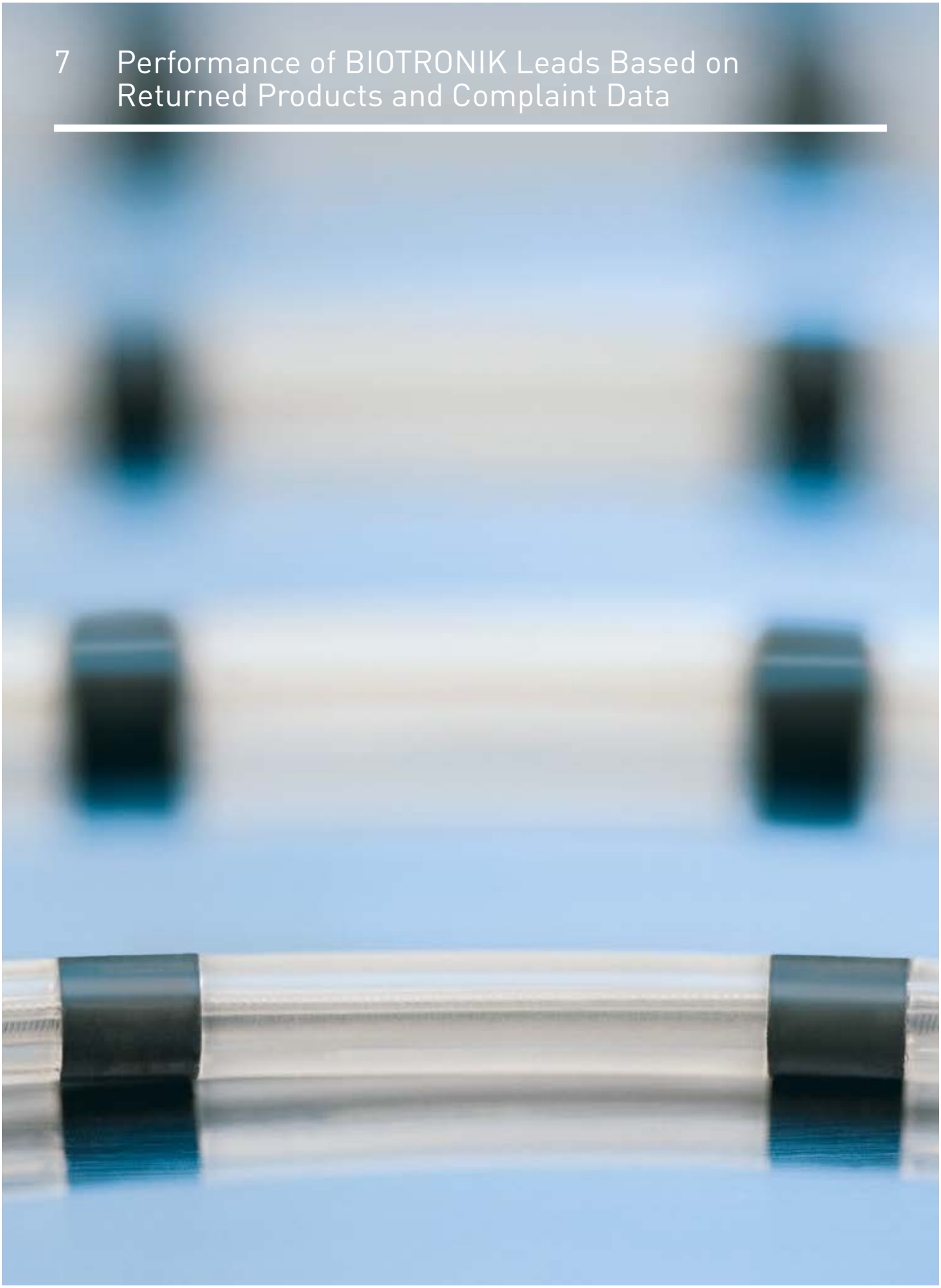
Products or subgroups of products may become subject to advisory notifications that can significantly impact the overall product performance.

Current advisories are listed in chapter 10 of this report, however to date, BIOTRONIK has never had a pacing or ICD lead safety advisory notification, therefore no summary of lead advisories is provided.

The cumulative survival data and the 95 % confidence intervals according to the Greenwood's formula[1] are shown in numerical form for the observed sample population.

1 Greenwood, M. The Natural Duration of Cancer. Reports on Public Health and Medical Subjects 33, London: Her Majesty's Stationery Office, 1–26, 1926

7 Performance of BIOTRONIK Leads Based on Returned Products and Complaint Data



7.1 Pacing Leads

7.2 ICD Leads

7.3 CRT Leads

7.1 Pacing Leads

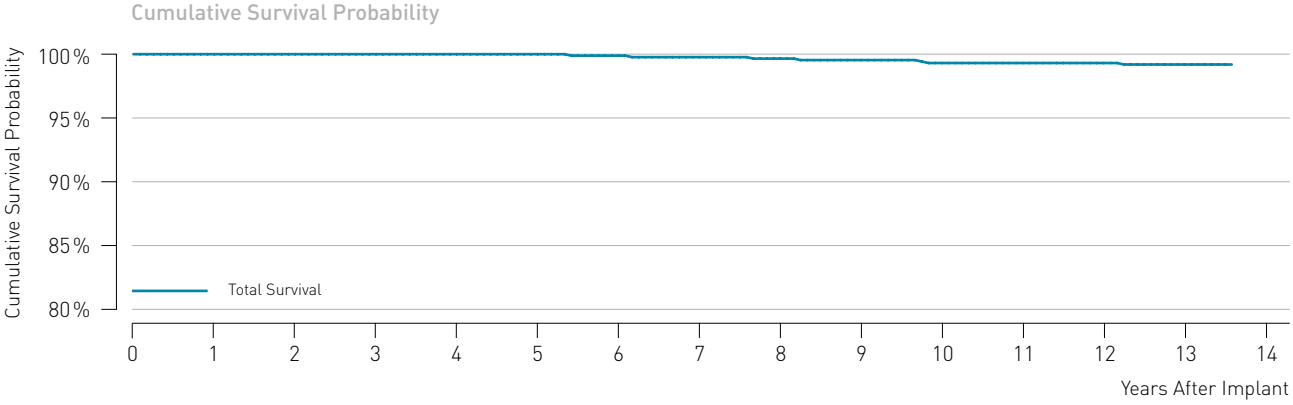
Arox

Product Details

Product Versions	53-BP, 60-BP
Lead Type	straight, passive fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	Sep 2002
CE Market Release	Jan 2002
Worldwide Distributed Devices	36 500
Registered U.S. Implants	8 550
Estimated Active U.S. Implants	4 510
U.S. Total Returned	19

	Quantity	Rate
U.S. Qualifying Complications	29	0.34%
▪ Abnormal pacing impedance	9	0.11%
▪ Conductor fracture	2	0.02%
▪ Failure to capture	14	0.16%
▪ Insulation breach	2	0.02%
▪ Other	2	0.02%

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.01%
▪ Insulation breach	1	0.01%
U.S. Acute Lead Observations	2	0.02%
▪ Lead dislodgement	2	0.02%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0	99.9	99.8	99.7	99.6	99.4	99.4	99.4	99.3
(95% Confidence Interval)							±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.2	±0.3

7.1 Pacing Leads

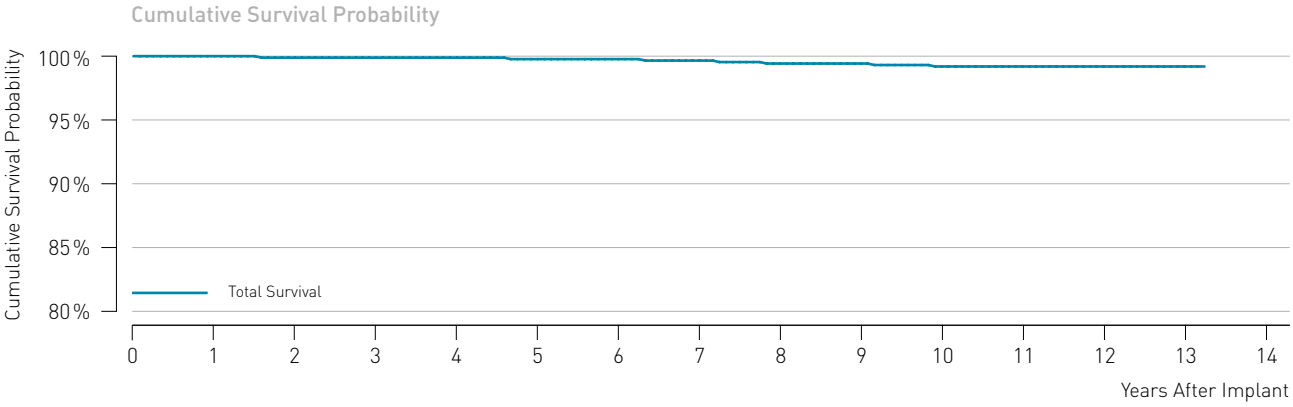
Arox J

Product Details

Product Versions	45-JBP, 53-JBP
Lead Type	J-shape, passive fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	Sep 2002
CE Market Release	Jan 2002
Worldwide Distributed Devices	8760
Registered U.S. Implants	3470
Estimated Active U.S. Implants	2090
U.S. Total Returned	8

	Quantity	Rate
U.S. Qualifying Complications	15	0.43%
▪ Abnormal pacing impedance	2	0.06%
▪ Conductor fracture	1	0.03%
▪ Failure to capture	9	0.26%
▪ Lead dislodgement	2	0.06%
▪ Oversensing	1	0.03%

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.
Total Survival [%]	100.0	100.0	99.9	99.9	99.9	99.8	99.8	99.7	99.5	99.5	99.3	99.3	99.3	99.3
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.3	±0.3	±0.4	±0.4	±0.4	±0.4

7.1 Pacing Leads

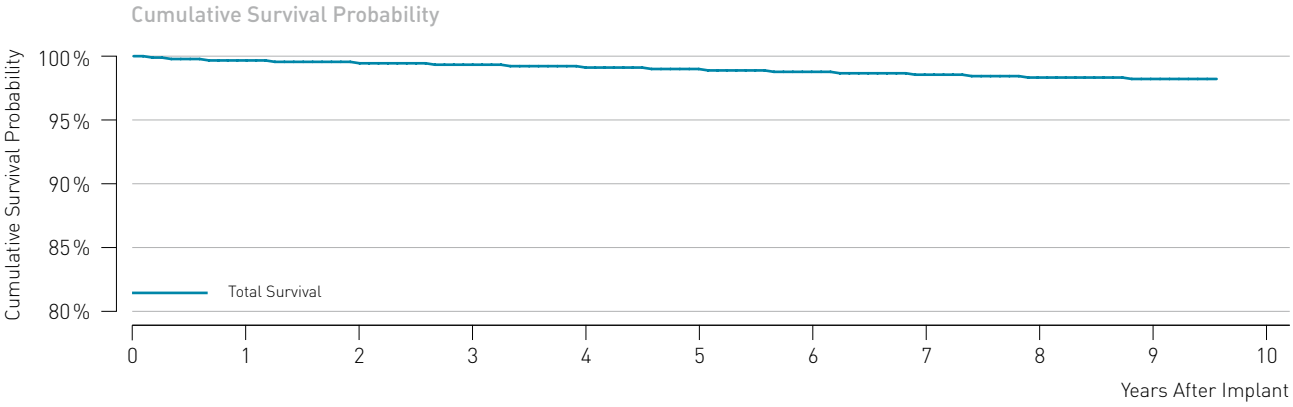
Dextrus

Product Details

Product Versions	4135, 4136, 4137	CE Market Release	May 2007
Lead Type	straight, active fixation	Worldwide Distributed Devices	487 000
		Registered U.S. Implants	382 000
Polarity	bipolar	Estimated Active U.S. Implants	276 000
Steroid	yes	U.S. Total Returned	2 076
U.S. Market Release	Apr 2007		

	Quantity	Rate
U.S. Qualifying Complications	2 536	0.66%
▪ Abnormal pacing impedance	191	0.05%
▪ Cardiac perforation	23	0.01%
▪ Conductor fracture	58	0.02%
▪ Extracardiac stimulation	17	0.00%
▪ Failure to capture	705	0.18%
▪ Failure to sense	103	0.03%
▪ Insulation breach	52	0.01%
▪ Lead dislodgement	454	0.12%
▪ Oversensing	445	0.12%
▪ Other	488	0.13%

	Quantity	Rate
U.S. Confirmed Malfunctions	254	0.07%
▪ Conductor fracture	99	0.03%
▪ Insulation breach	151	0.04%
▪ Other	4	0.00%
U.S. Acute Lead Observations	1 475	0.39%
▪ Abnormal pacing impedance	27	0.01%
▪ Cardiac perforation	59	0.02%
▪ Extracardiac stimulation	14	0.00%
▪ Failure to capture	202	0.05%
▪ Failure to sense	57	0.01%
▪ Insulation breach	9	0.00%
▪ Lead dislodgement	588	0.15%
▪ Oversensing	36	0.01%
▪ Other	483	0.13%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.
Total Survival [%]	100.0	99.7	99.5	99.4	99.2	99.1	98.9	98.7	98.5	98.4
(95% Confidence Interval)								±0.1	±0.1	±0.1

7.1 Pacing Leads

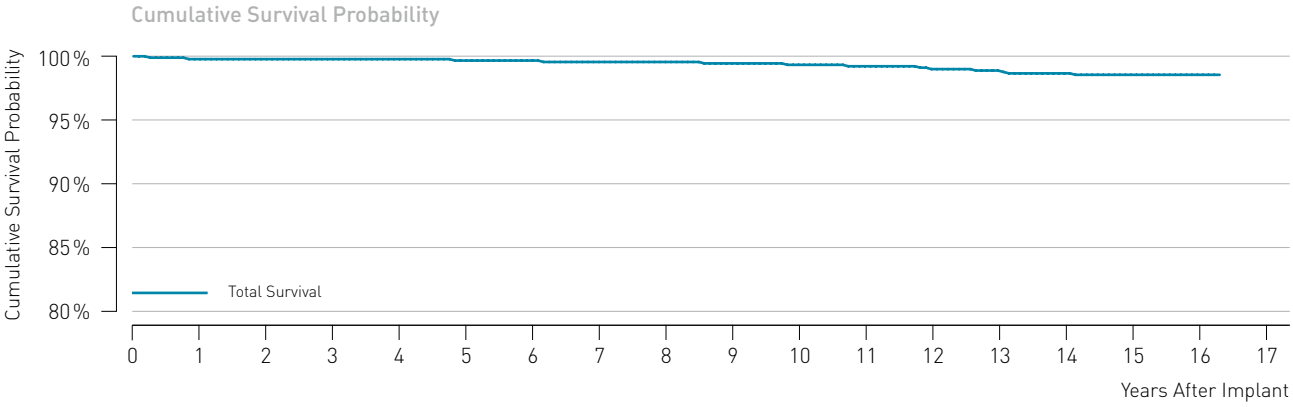
Elox

Product Details

Product Versions	45-BP, 53-BP, 60-BP
Lead Type	straight, active fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	May 2000
CE Market Release	May 2000
Worldwide Distributed Devices	36 000
Registered U.S. Implants	11 000
Estimated Active U.S. Implants	3 700
U.S. Total Returned	56

	Quantity	Rate
U.S. Qualifying Complications	59	0.54%
▪ Abnormal pacing impedance	3	0.03%
▪ Conductor fracture	2	0.02%
▪ Extracardiac stimulation	1	0.01%
▪ Failure to capture	17	0.15%
▪ Failure to sense	11	0.10%
▪ Insulation breach	4	0.04%
▪ Lead dislodgement	3	0.03%
▪ Oversensing	13	0.12%
▪ Other	5	0.05%

	Quantity	Rate
U.S. Confirmed Malfunctions	7	0.06%
▪ Conductor fracture	4	0.04%
▪ Insulation breach	3	0.03%
U.S. Acute Lead Observations	8	0.07%
▪ Failure to capture	4	0.04%
▪ Failure to sense	1	0.01%
▪ Oversensing	2	0.02%
▪ Other	1	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.	16 yr.
Total Survival [%]	100.0	99.8	99.8	99.8	99.8	99.7	99.7	99.6	99.6	99.5	99.4	99.3	99.1	99.0	98.8	98.7	98.7
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.3	±0.3	±0.3	±0.3	±0.3

7.1 Pacing Leads

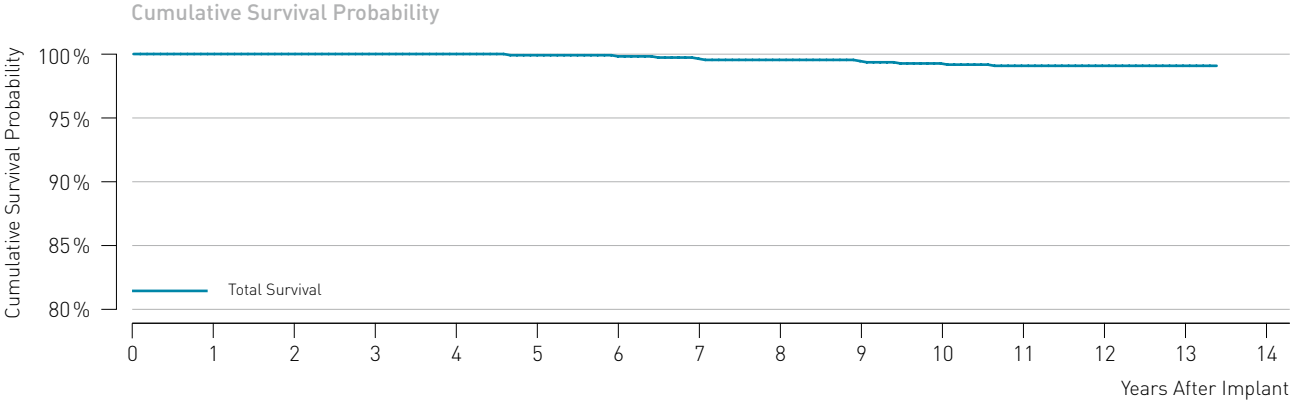
Elox P

Product Details

Product Versions	45-BP, 53-BP, 60-BP
Lead Type	straight, active fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	May 2003
CE Market Release	Feb 2003
Worldwide Distributed Devices	21 900
Registered U.S. Implants	3 030
Estimated Active U.S. Implants	1 330
U.S. Total Returned	19

	Quantity	Rate
U.S. Qualifying Complications	15	0.50%
▪ Abnormal pacing impedance	1	0.03%
▪ Failure to capture	8	0.26%
▪ Failure to sense	1	0.03%
▪ Insulation breach	2	0.07%
▪ Lead dislodgement	1	0.03%
▪ Oversensing	1	0.03%
▪ Other	1	0.03%

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.03%
▪ Insulation breach	1	0.03%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	100.0	99.9	99.8	99.6	99.5	99.4	99.2	99.0	99.0	99.0
(95% Confidence Interval)						±0.1	±0.2	±0.3	±0.3	±0.4	±0.4	±0.5	±0.5	±0.5

7.1 Pacing Leads

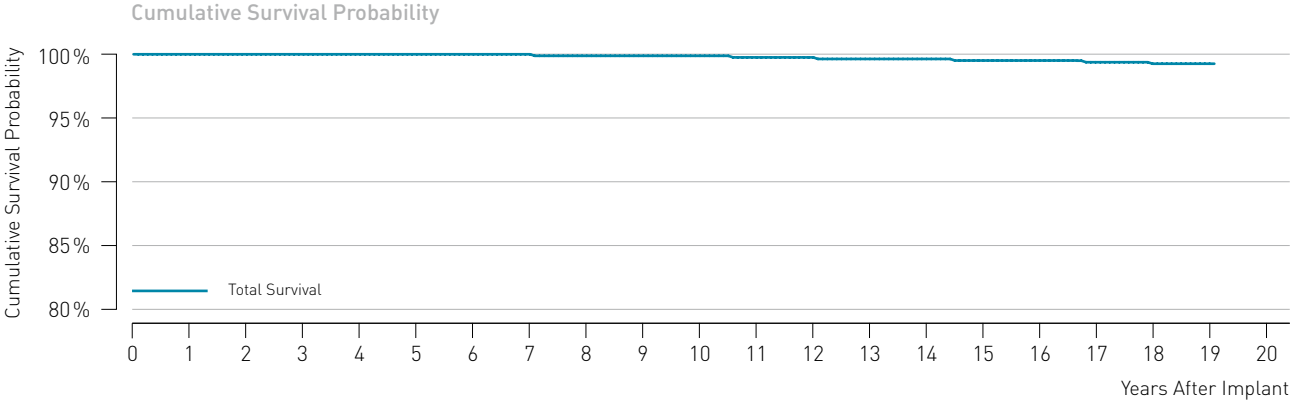
Polyrox

Product Details

Product Versions	60-UP, 53-BP, 53/15-BP, 60-BP, 60/15-BP
Lead Type	straight, passive fixation
Polarity	unipolar/bipolar
Steroid	no
U.S. Market Release	Mar 1997
CE Market Release	Jul 1996
Worldwide Distributed Devices	351 000
Registered U.S. Implants	15 100
Estimated Active U.S. Implants	4 530
U.S. Total Returned	26

	Quantity	Rate
U.S. Qualifying Complications	21	0.14%
▪ Abnormal pacing impedance	2	0.01%
▪ Conductor fracture	2	0.01%
▪ Failure to capture	13	0.09%
▪ Insulation breach	2	0.01%
▪ Lead dislodgement	1	0.01%
▪ Oversensing	1	0.01%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.01%
▪ Insulation breach	2	0.01%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.	16 yr.	17 yr.	18 yr.	19 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	99.9	99.9	99.9	99.8	99.8	99.7	99.7	99.6	99.6	99.5	99.4	99.4
(95% Confidence Interval)									±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.4	±0.4

7.1 Pacing Leads

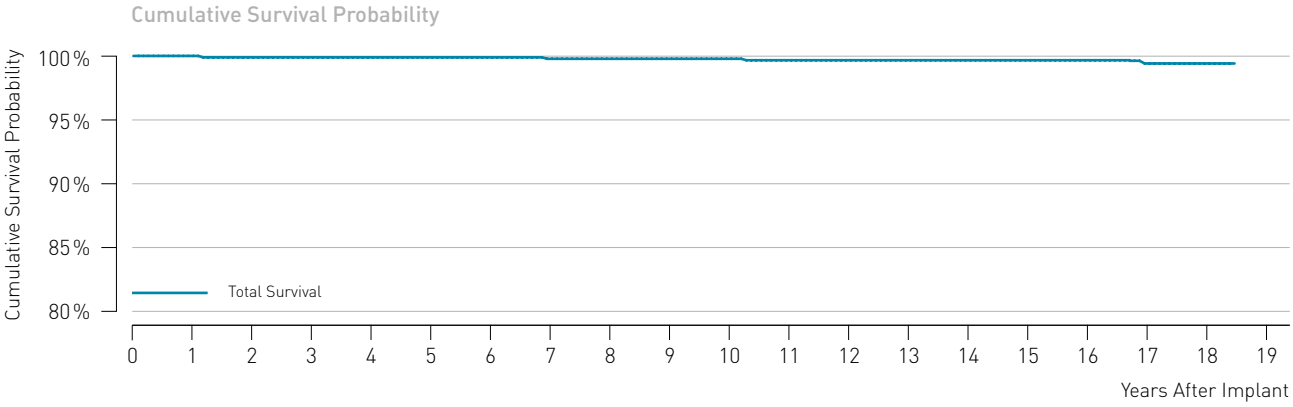
Polyrox J

Product Details

Product Versions	45-JBP, 53-JBP, 53-JUP
Lead Type	J-shape, passive fixation
Polarity	unipolar/bipolar
Steroid	no
U.S. Market Release	Mar 1997
CE Market Release	Jul 1996
Worldwide Distributed Devices	45 900
Registered U.S. Implants	3 730
Estimated Active U.S. Implants	1 200
U.S. Total Returned	6

	Quantity	Rate
U.S. Qualifying Complications	7	0.19%
▪ Abnormal pacing impedance	1	0.03%
▪ Failure to capture	1	0.03%
▪ Failure to sense	2	0.05%
▪ Lead dislodgement	1	0.03%
▪ Other	2	0.05%

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
U.S. Acute Lead Observations	1	0.03%
▪ Failure to capture	1	0.03%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.	16 yr.	17 yr.	18 yr.
Total Survival [%]	100.0	100.0	99.9	99.9	99.9	99.9	99.9	99.8	99.8	99.8	99.8	99.7	99.7	99.7	99.7	99.7	99.7	99.5	99.5
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.5	±0.5

7.1 Pacing Leads

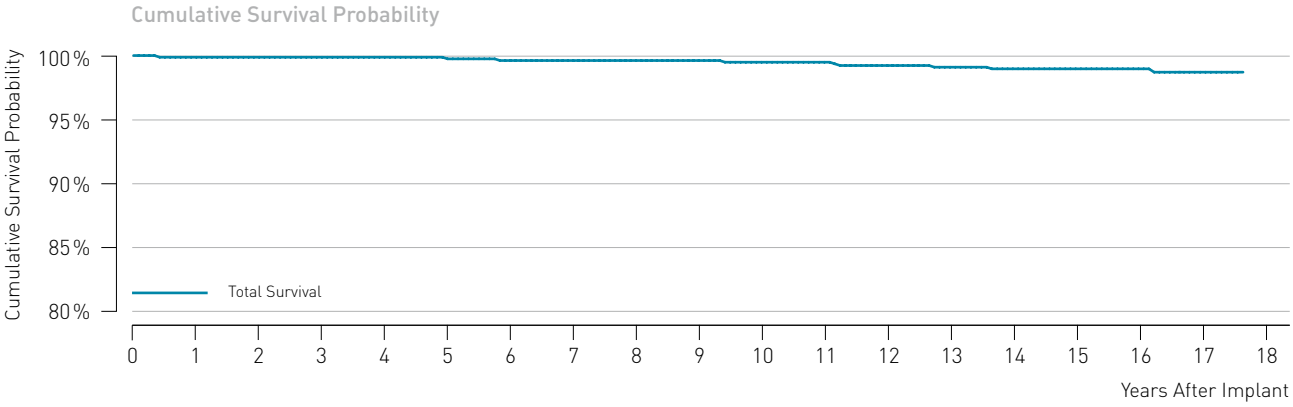
Retrox J

Product Details

Product Versions	45-JBP, 53-JBP
Lead Type	J-shape, active fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	Aug 1998
CE Market Release	Mar 1997
Worldwide Distributed Devices	14 000
Registered U.S. Implants	4 250
Estimated Active U.S. Implants	1 270
U.S. Total Returned	15

	Quantity	Rate
U.S. Qualifying Complications	17	0.40%
▪ Abnormal pacing impedance	2	0.05%
▪ Failure to capture	7	0.16%
▪ Failure to sense	2	0.05%
▪ Lead dislodgement	3	0.07%
▪ Oversensing	2	0.05%
▪ Other	1	0.02%

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
U.S. Acute Lead Observations	1	0.02%
▪ Failure to capture	1	0.02%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.	16 yr.	17 yr.
Total Survival [%]	100.0	99.9	99.9	99.9	99.9	99.8	99.7	99.7	99.7	99.7	99.6	99.6	99.4	99.3	99.2	99.2	99.2	99.0
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.2	±0.3	±0.4	±0.4	±0.4	±0.4	±0.4	±0.5

7.1 Pacing Leads

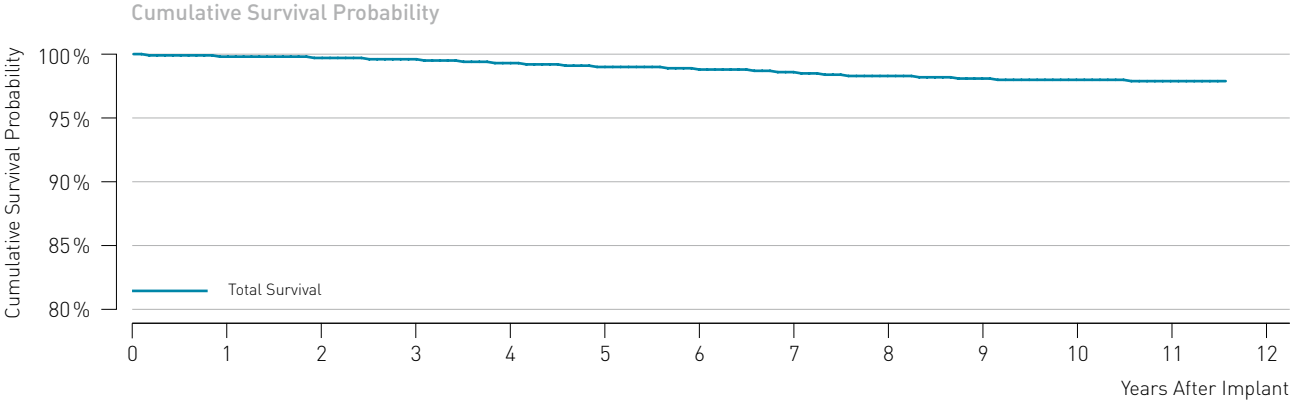
Selox JT

Product Details

Product Versions	45, 53
Lead Type	J-shape, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Nov 2004
CE Market Release	Nov 2004
Worldwide Distributed Devices	143 000
Registered U.S. Implants	15 900
Estimated Active U.S. Implants	12 000
U.S. Total Returned	104

	Quantity	Rate
U.S. Qualifying Complications	153	0.96%
▪ Abnormal pacing impedance	15	0.08%
▪ Cardiac perforation	1	0.01%
▪ Conductor fracture	5	0.03%
▪ Extracardiac stimulation	1	0.01%
▪ Failure to capture	72	0.45%
▪ Failure to sense	7	0.04%
▪ Insulation breach	8	0.05%
▪ Lead dislodgement	26	0.16%
▪ Oversensing	2	0.01%
▪ Other	16	0.10%

	Quantity	Rate
U.S. Confirmed Malfunctions	8	0.05%
▪ Insulation breach	8	0.05%
U.S. Acute Lead Observations	40	0.25%
▪ Failure to capture	5	0.03%
▪ Lead dislodgement	32	0.20%
▪ Other	3	0.02%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.
Total Survival [%]	100.0	99.8	99.7	99.6	99.3	99	98.8	98.6	98.3	98.1	98	97.9
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.3	±0.3	±0.4	±0.4

7.1 Pacing Leads

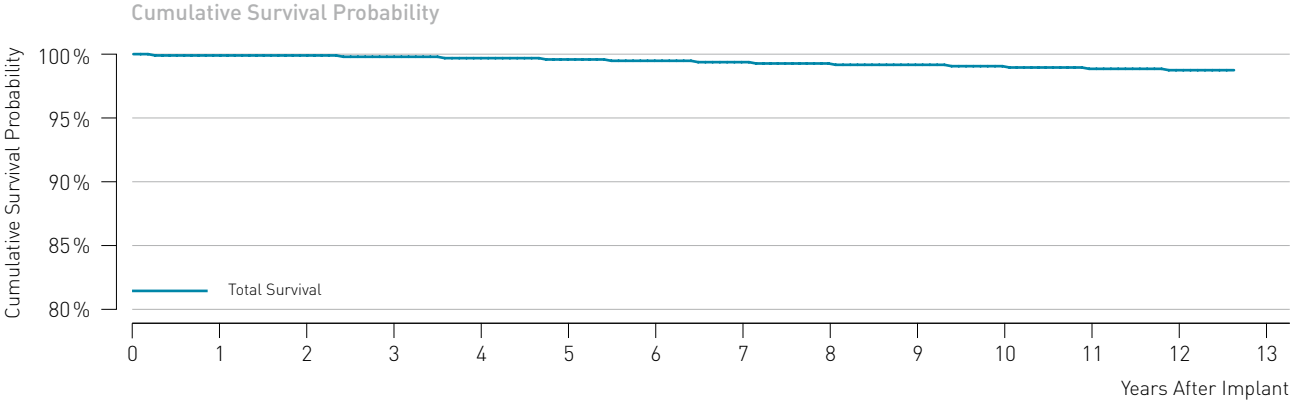
Selox SR

Product Details

Product Versions	45, 53, 60
Lead Type	straight, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Mar 2004
CE Market Release	Feb 2004
Worldwide Distributed Devices	171 000
Registered U.S. Implants	14 400
Estimated Active U.S. Implants	7 190
U.S. Total Returned	59

	Quantity	Rate
U.S. Qualifying Complications	90	0.63%
▪ Abnormal pacing impedance	3	0.02%
▪ Conductor fracture	5	0.03%
▪ Extracardiac stimulation	2	0.01%
▪ Failure to capture	37	0.26%
▪ Failure to sense	1	0.01%
▪ Insulation breach	6	0.04%
▪ Lead dislodgement	13	0.09%
▪ Oversensing	10	0.07%
▪ Other	13	0.09%

	Quantity	Rate
U.S. Confirmed Malfunctions	10	0.07%
▪ Insulation breach	10	0.07%
U.S. Acute Lead Observations	20	0.15%
▪ Cardiac perforation	1	0.01%
▪ Failure to capture	11	0.08%
▪ Insulation breach	1	0.01%
▪ Lead dislodgement	8	0.06%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.
Total Survival [%]	100.0	99.9	99.9	99.8	99.7	99.6	99.5	99.4	99.3	99.2	99.1	98.9	98.8
(95% Confidence Interval)			±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2

7.1 Pacing Leads

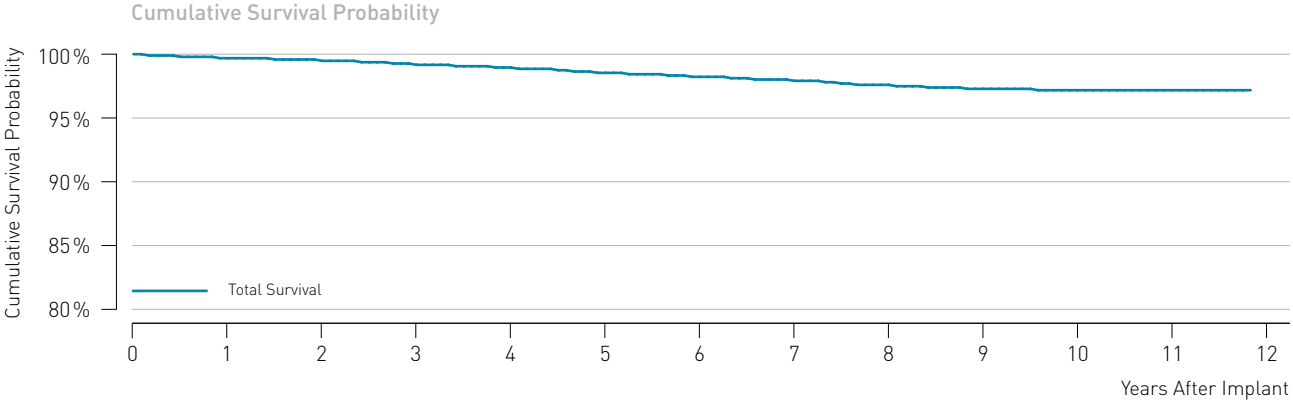
Selox ST

Product Details

Product Versions	53, 60
Lead Type	straight, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Nov 2004
CE Market Release	Nov 2004
Worldwide Distributed Devices	367 000
Registered U.S. Implants	30 900
Estimated Active U.S. Implants	22 400
U.S. Total Returned	152

	Quantity	Rate
U.S. Qualifying Complications	420	1.36%
▪ Abnormal pacing impedance	99	0.32%
▪ Cardiac perforation	3	0.01%
▪ Conductor fracture	36	0.12%
▪ Extracardiac stimulation	7	0.02%
▪ Failure to capture	203	0.66%
▪ Insulation breach	32	0.10%
▪ Lead dislodgement	16	0.05%
▪ Oversensing	5	0.02%
▪ Other	19	0.06%

	Quantity	Rate
U.S. Confirmed Malfunctions	14	0.05%
▪ Conductor fracture	1	0.00%
▪ Crimps, welds and bonds	1	0.00%
▪ Insulation breach	12	0.04%
U.S. Acute Lead Observations	42	0.14%
▪ Abnormal pacing impedance	1	0.00%
▪ Failure to capture	16	0.05%
▪ Lead dislodgement	19	0.06%
▪ Other	6	0.02%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.
Total Survival [%]	100.0	99.7	99.5	99.2	99	98.6	98.3	98.0	97.7	97.4	97.3	97.3
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.3	±0.3	±0.3

7.1 Pacing Leads

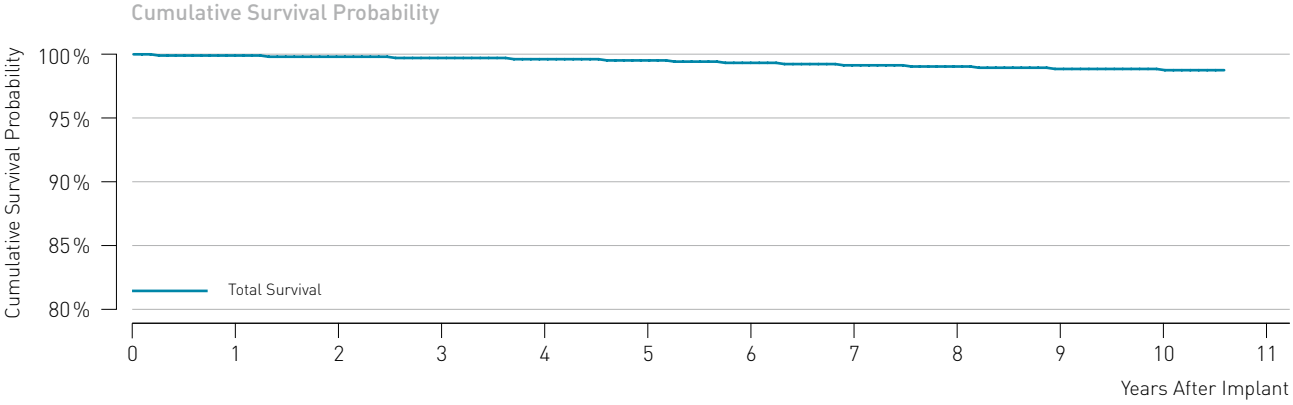
Setrox S

Product Details

Product Versions	45, 53, 60
Lead Type	straight, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Apr 2006
CE Market Release	Mar 2006
Worldwide Distributed Devices	656 000
Registered U.S. Implants	242 000
Estimated Active U.S. Implants	201 000
U.S. Total Returned	1 386

	Quantity	Rate
U.S. Qualifying Complications	951	0.39%
▪ Abnormal pacing impedance	66	0.03%
▪ Cardiac perforation	8	0.00%
▪ Conductor fracture	35	0.01%
▪ Extracardiac stimulation	9	0.00%
▪ Failure to capture	322	0.13%
▪ Failure to sense	27	0.01%
▪ Insulation breach	53	0.02%
▪ Lead dislodgement	244	0.10%
▪ Oversensing	97	0.04%
▪ Other	90	0.04%

	Quantity	Rate
U.S. Confirmed Malfunctions	112	0.05%
▪ Conductor fracture	43	0.02%
▪ Insulation breach	68	0.03%
▪ Other	1	0.00%
U.S. Acute Lead Observations	259	0.11%
▪ Abnormal pacing impedance	1	0.00%
▪ Cardiac perforation	19	0.01%
▪ Failure to capture	36	0.01%
▪ Failure to sense	2	0.00%
▪ Insulation breach	4	0.00%
▪ Lead dislodgement	182	0.08%
▪ Other	15	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.
Total Survival [%]	100.0	99.9	99.8	99.7	99.6	99.5	99.3	99.1	99.0	98.9	98.8
(95% Confidence Interval)								±0.1	±0.1	±0.1	±0.1

7.1 Pacing Leads

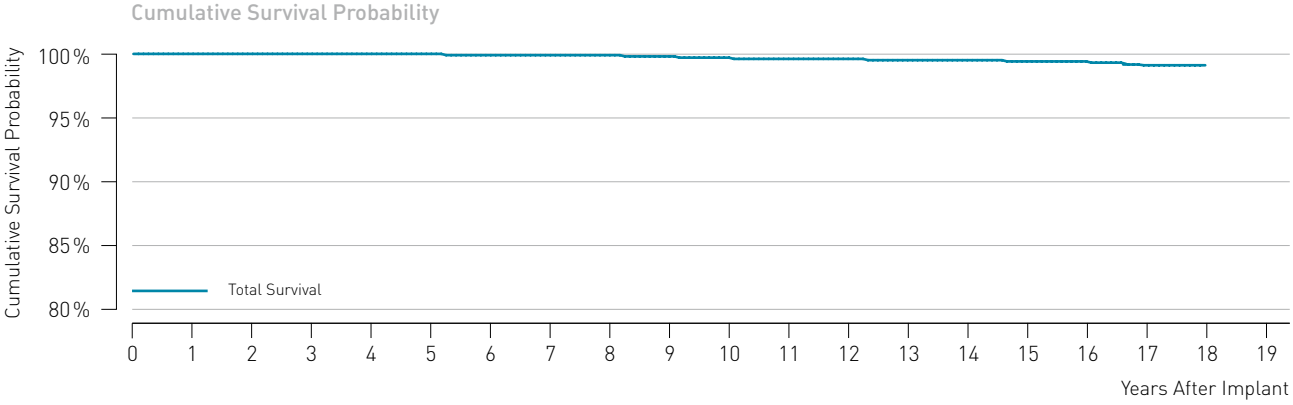
Synox

Product Details

Product Versions	60-UP, 53-BP, 60-BP
Lead Type	straight, passive fixation
Polarity	unipolar/bipolar
Steroid	no
U.S. Market Release	Sep 1998
CE Market Release	Jul 1996
Worldwide Distributed Devices	169 000
Registered U.S. Implants	17 600
Estimated Active U.S. Implants	6 210
U.S. Total Returned	57

	Quantity	Rate
U.S. Qualifying Complications	41	0.23%
▪ Abnormal pacing impedance	3	0.02%
▪ Conductor fracture	2	0.01%
▪ Extracardiac stimulation	1	0.01%
▪ Failure to capture	19	0.11%
▪ Failure to sense	1	0.01%
▪ Insulation breach	6	0.03%
▪ Lead dislodgement	1	0.01%
▪ Oversensing	2	0.01%
▪ Other	6	0.03%

	Quantity	Rate
U.S. Confirmed Malfunctions	5	0.03%
▪ Conductor fracture	2	0.01%
▪ Insulation breach	3	0.02%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.	16 yr.	17 yr.	18 yr.
Total Survival [%]	100.0	100.0	100.0	100.0	100.0	100.0	99.9	99.9	99.9	99.8	99.7	99.6	99.6	99.5	99.5	99.4	99.4	99.1	99.1
(95% Confidence Interval)								±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.4	±0.4

7.1 Pacing Leads

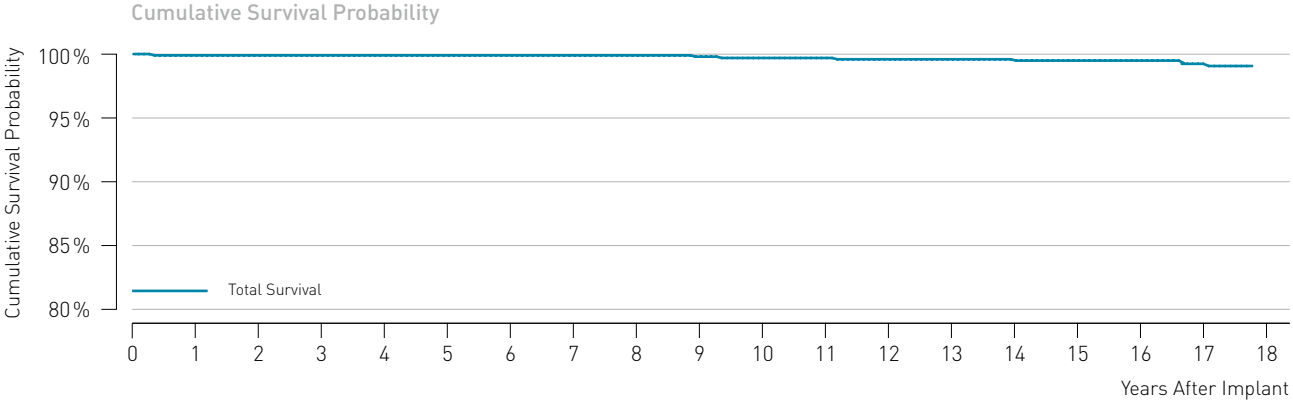
Synox J

Product Details

Product Versions	45-JBP, 53-JBP
Lead Type	J-shape, passive fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	Sep 1998
CE Market Release	Jul 1996
Worldwide Distributed Devices	81 400
Registered U.S. Implants	8 170
Estimated Active U.S. Implants	3 320
U.S. Total Returned	27

	Quantity	Rate
U.S. Qualifying Complications	21	0.26%
▪ Abnormal pacing impedance	1	0.01%
▪ Conductor fracture	2	0.02%
▪ Failure to capture	5	0.06%
▪ Failure to sense	4	0.05%
▪ Insulation breach	2	0.02%
▪ Lead dislodgement	2	0.02%
▪ Oversensing	3	0.04%
▪ Other	2	0.02%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.02%
▪ Insulation breach	1	0.01%
▪ Crimps, welds and bonds	1	0.01%
U.S. Acute Lead Observations	2	0.02%
▪ Failure to capture	1	0.01%
▪ Oversensing	1	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.	16 yr.	17 yr.
Total Survival [%]	100.0	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.9	99.8	99.7	99.7	99.6	99.6	99.5	99.5	99.5	99.3
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.1	±0.2	±0.2	±0.2	±0.2	±0.2	±0.2	±0.4

7.1 Pacing Leads

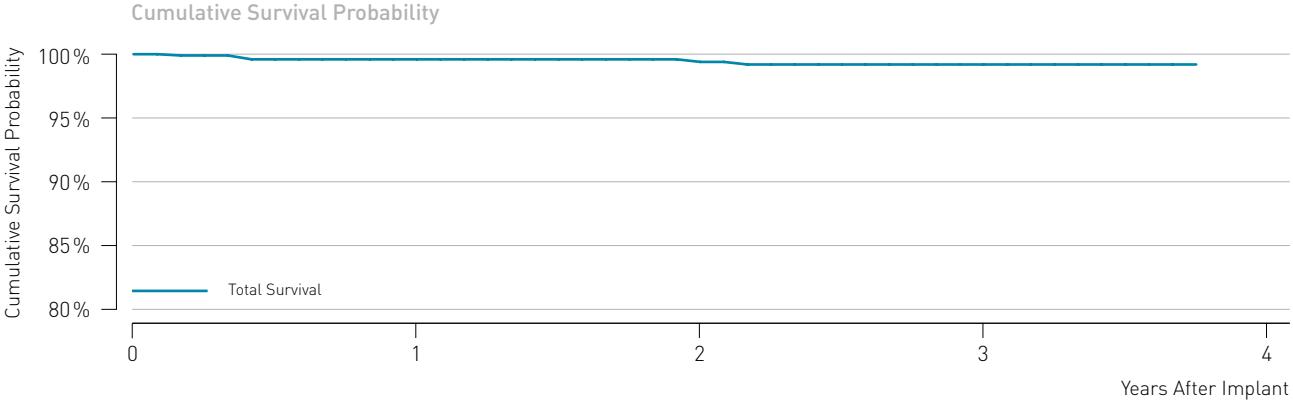
Tilda JT

Product Details

Product Versions	45, 53
Lead Type	J-shape, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Feb 2012
CE Market Release	Sep 2012
Worldwide Distributed Devices	14 700
Registered U.S. Implants	736
Estimated Active U.S. Implants	717
U.S. Total Returned	0

	Quantity	Rate
U.S. Qualifying Complications	5	0.68%
▪ Abnormal pacing impedance	1	0.14%
▪ Failure to capture	1	0.14%
▪ Lead dislodgement	3	0.41%

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
U.S. Acute Lead Observations	1	0.14%
▪ Lead dislodgement	1	0.14%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.
Total Survival [%]	100.0	99.6	99.4	99.2
(95% Confidence Interval)		±0.5	±0.6	±0.7

7.1 Pacing Leads

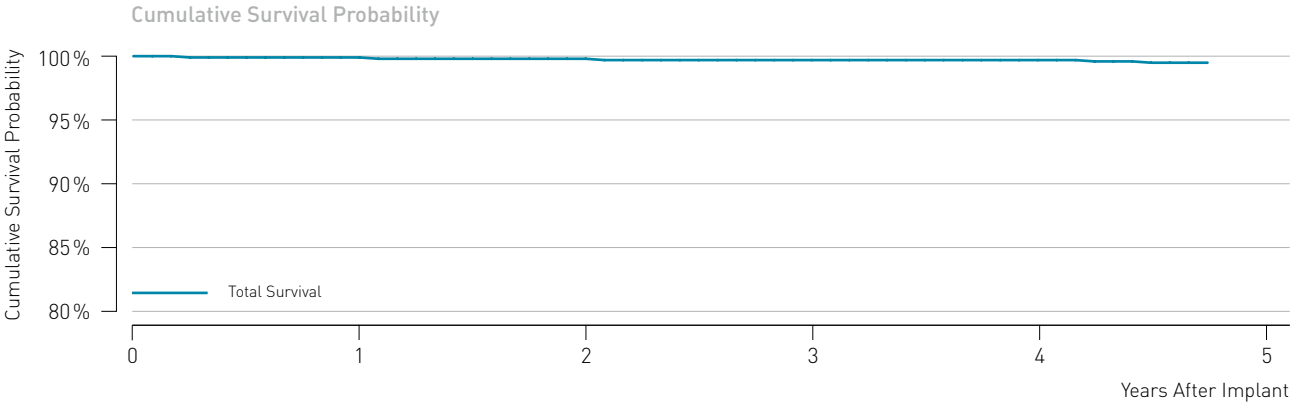
Tilda R

Product Details

Product Versions	45, 53, 60
Lead Type	straight, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Dec 2011
CE Market Release	Aug 2011
Worldwide Distributed Devices	39 400
Registered U.S. Implants	9 340
Estimated Active U.S. Implants	9 000
U.S. Total Returned	14

	Quantity	Rate
U.S. Qualifying Complications	26	0.28%
▪ Abnormal pacing impedance	1	0.01%
▪ Conductor fracture	3	0.03%
▪ Extracardiac stimulation	1	0.01%
▪ Failure to capture	7	0.07%
▪ Insulation breach	2	0.02%
▪ Lead dislodgement	9	0.10%
▪ Oversensing	1	0.01%
▪ Other	2	0.02%

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.01%
▪ Conductor fracture	1	0.01%
U.S. Acute Lead Observations	8	0.09%
▪ Lead dislodgement	7	0.07%
▪ Other	1	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	99.9	99.8	99.7	99.7
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.1

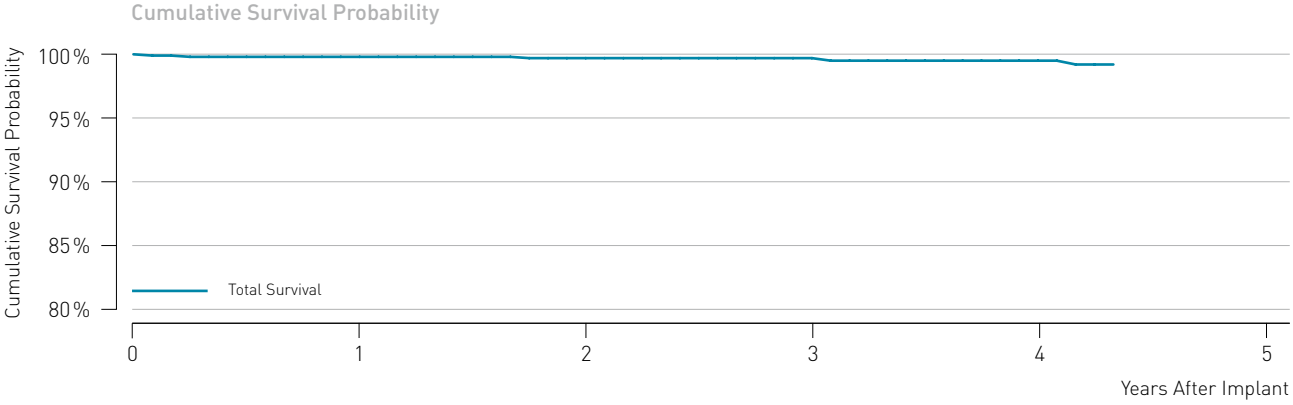
7.1 Pacing Leads

Tilda T

Product Details

Product Versions	53, 60
Lead Type	straight, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Dec 2011
CE Market Release	Aug 2011
Worldwide Distributed Devices	20 000
Registered U.S. Implants	1 280
Estimated Active U.S. Implants	1 240
U.S. Total Returned	1

	Quantity	Rate		Quantity	Rate
U.S. Qualifying Complications	6	0.47%	U.S. Confirmed Malfunctions	0	0.00%
▪ Abnormal pacing impedance	2	0.16%	U.S. Acute Lead Observations	0	0.00%
▪ Insulation breach	1	0.08%			
▪ Lead dislodgement	3	0.24%			



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	99.8	99.7	99.7	99.5
(95% Confidence Interval)		±0.3	±0.3	±0.3	±0.4

7.2 ICD Leads

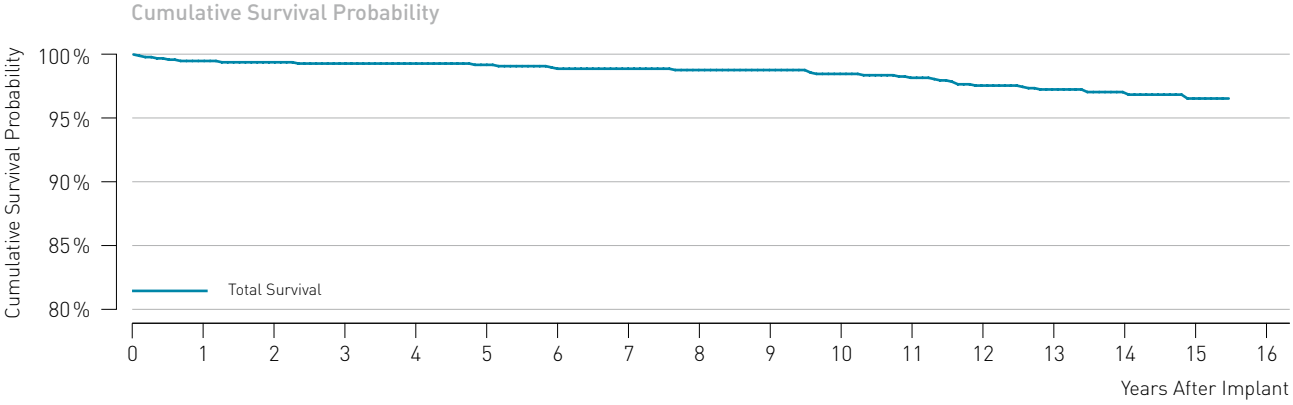
Kainox SL

Product Details

Product Versions	65, 75, 100
Lead Type	dual-coil, passive fixation
Polarity	bipolar
Steroid	no
U.S. Market Release	Nov 1998
CE Market Release	Sep 1997
Worldwide Distributed Devices	9 600
Registered U.S. Implants	2 500
Estimated Active U.S. Implants	874
U.S. Total Returned	18

	Quantity	Rate
U.S. Qualifying Complications	37	1.48%
▪ Abnormal defibrillation impedance	1	0.04%
▪ Abnormal pacing impedance	4	0.16%
▪ Conductor fracture	3	0.12%
▪ Failure to capture	8	0.32%
▪ Failure to sense	1	0.04%
▪ Insulation breach	3	0.12%
▪ Lead dislodgement	1	0.04%
▪ Oversensing	14	0.56%
▪ Other	2	0.08%

	Quantity	Rate
U.S. Confirmed Malfunctions	3	0.12%
▪ Insulation breach	3	0.12%
U.S. Acute Lead Observations	5	0.20%
▪ Failure to capture	3	0.12%
▪ Failure to sense	1	0.04%
▪ Oversensing	1	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.	12 yr.	13 yr.	14 yr.	15 yr.
Total Survival [%]	100.0	99.5	99.4	99.3	99.3	99.2	98.9	98.9	98.8	98.8	98.5	98.2	97.6	97.3	97.1	96.6
(95% Confidence Interval)		±0.3	±0.3	±0.4	±0.4	±0.4	±0.5	±0.5	±0.5	±0.5	±0.6	±0.7	±0.8	±0.9	±1.0	±1.2

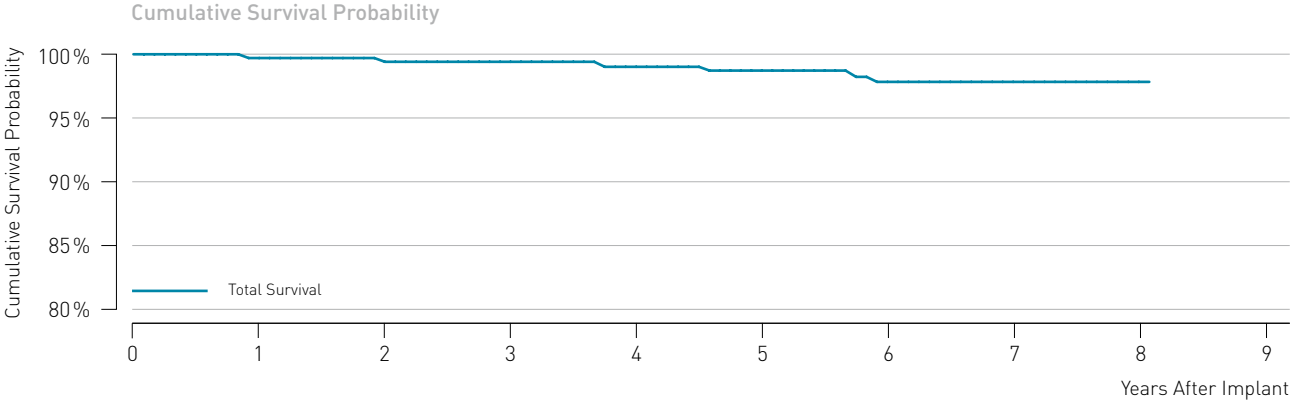
Kentrox RV

Product Details

Product Versions	65, 75, -Steroid
Lead Type	single-coil, passive fixation
Polarity	bipolar
Steroid	yes/no
U.S. Market Release	Mar 2002 / Oct 2004
CE Market Release	Jan 2001 / Dec 2004
Worldwide Distributed Devices	5490
Registered U.S. Implants	409
Estimated Active U.S. Implants	175
U.S. Total Returned	8

	Quantity	Rate
U.S. Qualifying Complications	7	1.71%
▪ Conductor fracture	1	0.24%
▪ Failure to capture	1	0.24%
▪ Insulation breach	1	0.24%
▪ Oversensing	4	0.98%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.49%
▪ Conductor fracture	1	0.24%
▪ Insulation breach	1	0.24%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.
Total Survival [%]	100.0	99.7	99.4	99.4	99	98.7	97.8	97.8	97.8
(95% Confidence Interval)		±0.6	±0.8	±0.8	±1.1	±1.3	±1.8	±1.8	±1.8

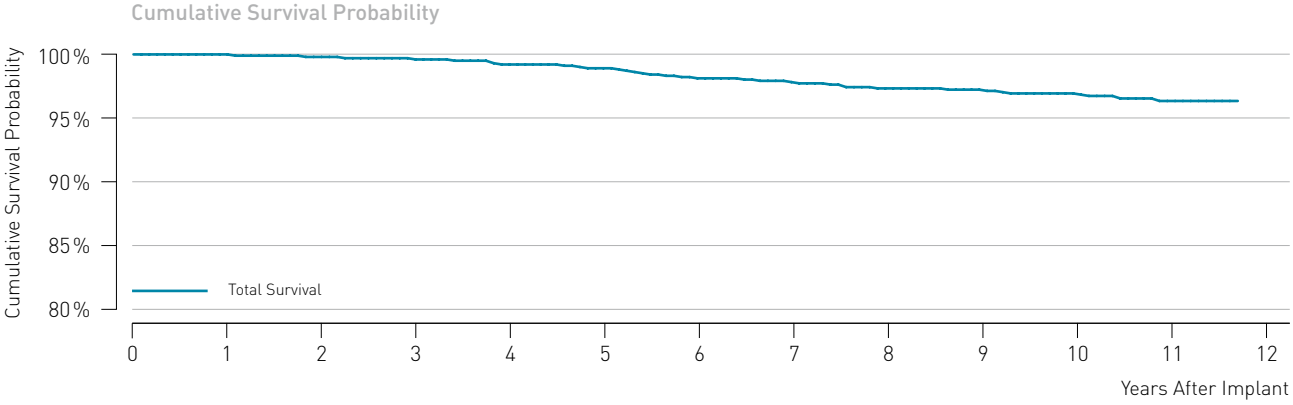
Kentrox SL-S

Product Details

Product Versions	65/16, 18 -Steroid
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes/no
U.S. Market Release	Oct 2004
CE Market Release	Jun 2004
Worldwide Distributed Devices	8 730
Registered U.S. Implants	2 440
Estimated Active U.S. Implants	1 270
U.S. Total Returned	41

	Quantity	Rate
U.S. Qualifying Complications	43	1.77%
▪ Abnormal defibrillation impedance	1	0.04%
▪ Abnormal pacing impedance	3	0.12%
▪ Conductor fracture	3	0.12%
▪ Failure to capture	3	0.12%
▪ Insulation breach	3	0.12%
▪ Lead dislodgement	3	0.12%
▪ Oversensing	25	1.03%
▪ Other	2	0.08%

	Quantity	Rate
U.S. Confirmed Malfunctions	14	0.57%
▪ Insulation breach	14	0.57%
U.S. Acute Lead Observations	2	0.08%
▪ Insulation breach	1	0.04%
▪ Oversensing	1	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	11 yr.
Total Survival [%]	100.0	100.0	99.8	99.6	99.2	98.9	98.1	97.8	97.3	97.2	96.9	96.3
(95% Confidence Interval)		±0.1	±0.2	±0.3	±0.4	±0.5	±0.6	±0.7	±0.8	±0.8	±0.9	±0.9

7.2 ICD Leads

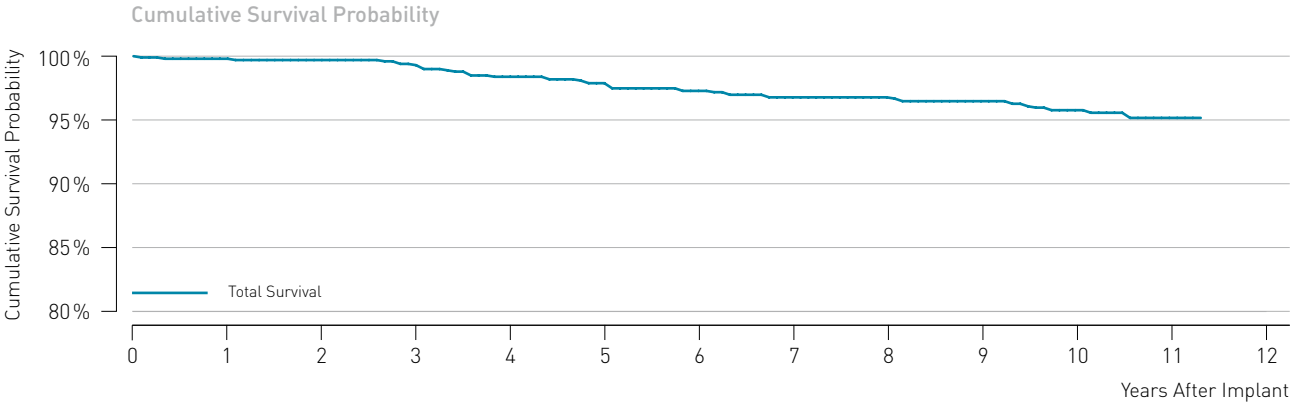
Kentrox SL

Product Details

Product Versions	65, 75, 100, -Steroid
Lead Type	dual coil, passive fixation
Polarity	bipolar
Steroid	yes/no
U.S. Market Release	Oct 2004
CE Market Release	Dec 2003 / Dec 2004
Worldwide Distributed Devices	8480
Registered U.S. Implants	1010
Estimated Active U.S. Implants	542
U.S. Total Returned	19

	Quantity	Rate
U.S. Qualifying Complications	27	2.68%
▪ Abnormal pacing impedance	3	0.30%
▪ Conductor fracture	2	0.20%
▪ Failure to capture	1	0.10%
▪ Insulation breach	6	0.60%
▪ Oversensing	13	1.29%
▪ Other	2	0.20%

	Quantity	Rate
U.S. Confirmed Malfunctions	5	0.50%
▪ Insulation breach	5	0.50%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.	12 yr.
Total Survival [%]	100.0	99.8	99.7	99.3	98.4	97.9	97.3	96.8	96.8	96.5	95.8	95.2
(95% Confidence Interval)		±0.3	±0.4	±0.6	±0.9	±1.0	±1.2	±1.3	±1.3	±1.4	±1.5	±1.7

7.2 ICD Leads

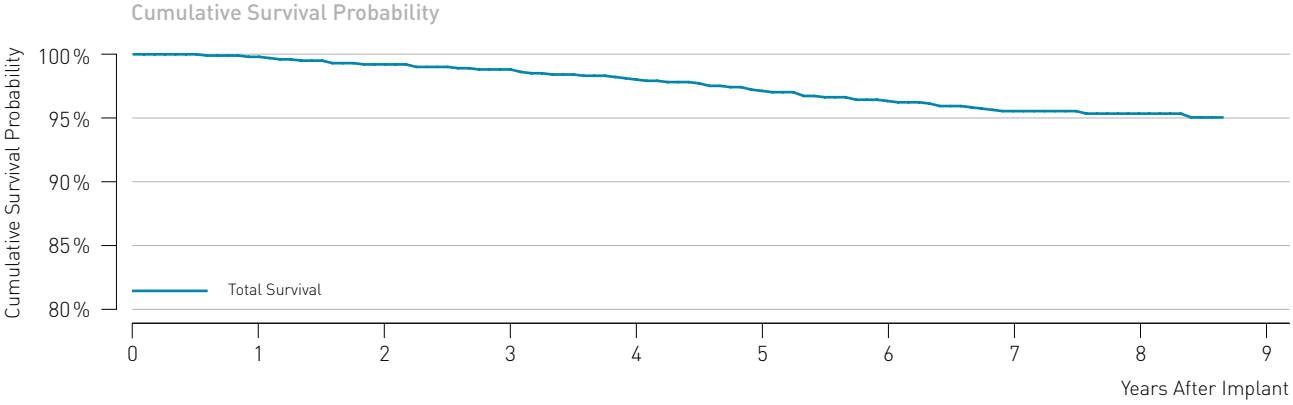
Linox S

Product Details

Product Versions	65, 75
Lead Type	single-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Feb 2007
CE Market Release	Mar 2007
Worldwide Distributed Devices	31 900
Registered U.S. Implants	2 500
Estimated Active U.S. Implants	1 750
U.S. Total Returned	62

	Quantity	Rate
U.S. Qualifying Complications	54	2.17%
▪ Abnormal defibrillation impedance	6	0.24%
▪ Abnormal pacing impedance	3	0.12%
▪ Conductor fracture	4	0.16%
▪ Failure to capture	7	0.28%
▪ Insulation breach	4	0.16%
▪ Oversensing	24	0.96%
▪ Other	6	0.24%

	Quantity	Rate
U.S. Confirmed Malfunctions	29	1.16%
▪ Conductor fracture	4	0.16%
▪ Insulation breach	25	1.00%
U.S. Acute Lead Observations	2	0.08%
▪ Lead dislodgement	1	0.04%
▪ Other	1	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.
Total Survival [%]	100.0	99.8	99.2	98.8	98.0	97.1	96.3	95.5	95.3
(95% Confidence Interval)		±0.2	±0.4	±0.5	±0.6	±0.7	±0.8	±1.0	±1.1

7.2 ICD Leads

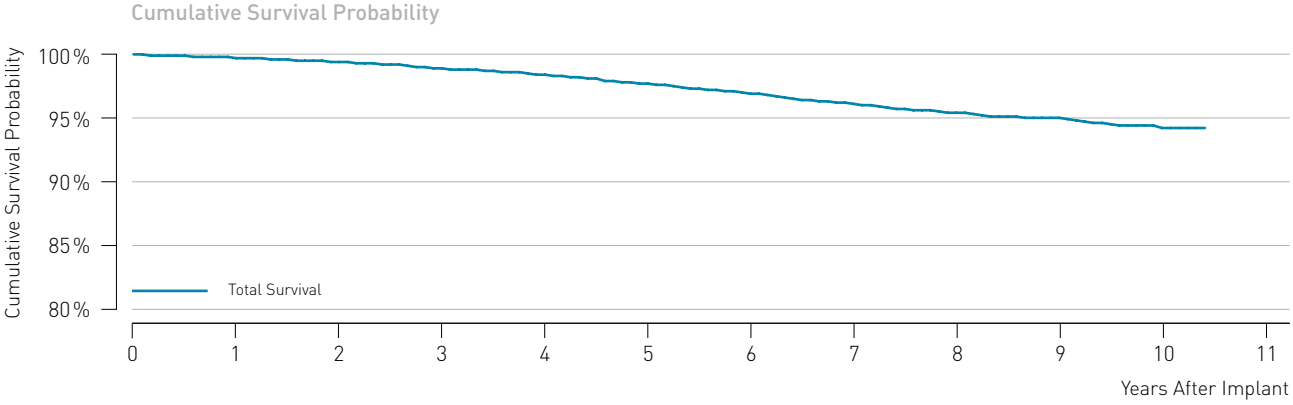
Linox SD

Product Details

Product Versions	60, 65, 75 / 16,18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Apr 2006
CE Market Release	Jun 2005
Worldwide Distributed Devices	55 100
Registered U.S. Implants	22 300
Estimated Active U.S. Implants	14 900
U.S. Total Returned	436

	Quantity	Rate
U.S. Qualifying Complications	561	2.52%
▪ Abnormal defibrillation impedance	37	0.17%
▪ Abnormal pacing impedance	43	0.19%
▪ Cardiac perforation	2	0.01%
▪ Conductor fracture	49	0.22%
▪ Failure to capture	57	0.26%
▪ Failure to sense	7	0.03%
▪ Insulation breach	52	0.23%
▪ Lead dislodgement	31	0.14%
▪ Oversensing	239	1.07%
▪ Other	44	0.20%

	Quantity	Rate
U.S. Confirmed Malfunctions	172	0.77%
▪ Conductor fracture	23	0.10%
▪ Insulation breach	148	0.67%
▪ Other	1	0.00%
U.S. Acute Lead Observations	11	0.05%
▪ Abnormal pacing impedance	1	0.00%
▪ Cardiac perforation	1	0.00%
▪ Failure to capture	1	0.00%
▪ Lead dislodgement	6	0.03%
▪ Oversensing	1	0.00%
▪ Other	1	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.	10 yr.
Total Survival [%]	100.0	99.7	99.4	98.9	98.4	97.7	96.9	96.1	95.4	95	94.2
(95% Confidence Interval)		±0.1	±0.1	±0.1	±0.2	±0.2	±0.3	±0.3	±0.3	±0.4	±0.6

7.2 ICD Leads

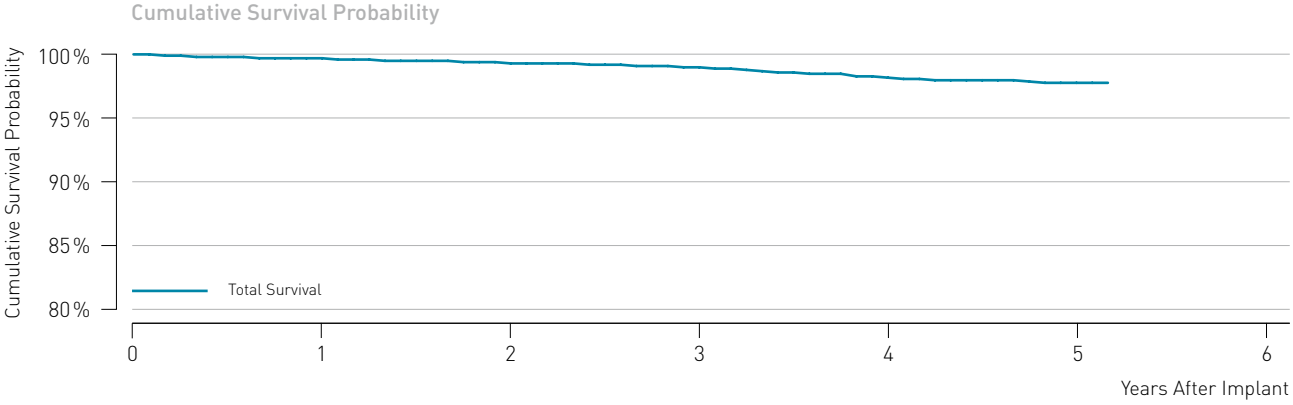
Linox^{smart} S

Product Details

Product Versions	60, 65, 75
Lead Type	single-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Aug 2011
CE Market Release	Dec 2010
Worldwide Distributed Devices	46 100
Registered U.S. Implants	7 460
Estimated Active U.S. Implants	6 460
U.S. Total Returned	117

	Quantity	Rate
U.S. Qualifying Complications	62	0.83%
▪ Abnormal defibrillation impedance	3	0.04%
▪ Abnormal pacing impedance	2	0.03%
▪ Cardiac perforation	1	0.01%
▪ Conductor fracture	3	0.04%
▪ Failure to capture	9	0.12%
▪ Failure to sense	2	0.03%
▪ Insulation breach	1	0.01%
▪ Lead dislodgement	13	0.17%
▪ Oversensing	24	0.32%
▪ Other	4	0.05%

	Quantity	Rate
U.S. Confirmed Malfunctions	27	0.36%
▪ Conductor fracture	4	0.05%
▪ Insulation breach	23	0.31%
U.S. Acute Lead Observations	11	0.15%
▪ Abnormal pacing impedance	1	0.01%
▪ Cardiac perforation	1	0.01%
▪ Lead dislodgement	8	0.11%
▪ Other	1	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	99.7	99.3	99.0	98.2	97.8
(95% Confidence Interval)		±0.1	±0.2	±0.3	±0.4	±0.6

7.2 ICD Leads

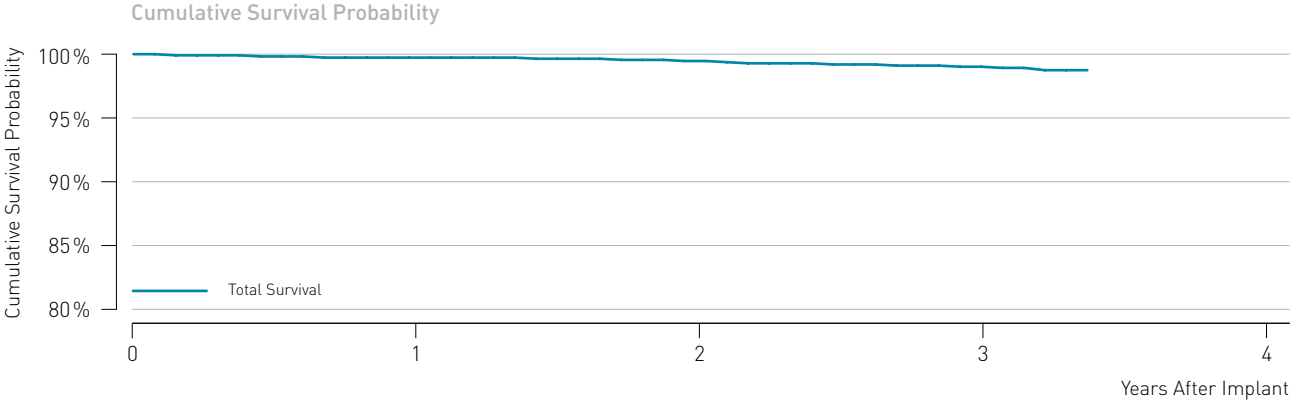
Linox^{smart} S DX

Product Details

Product Versions	65/15, 65/17
Lead Type	single-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Feb 2013
CE Market Release	Mar 2010
Worldwide Distributed Devices	33 400
Registered U.S. Implants	13 400
Estimated Active U.S. Implants	12 400
U.S. Total Returned	168

	Quantity	Rate
U.S. Qualifying Complications	44	0.33%
▪ Abnormal defibrillation impedance	2	0.02%
▪ Abnormal pacing impedance	1	0.01%
▪ Conductor fracture	3	0.02%
▪ Failure to capture	4	0.03%
▪ Failure to sense	2	0.02%
▪ Lead dislodgement	20	0.15%
▪ Oversensing	9	0.07%
▪ Other	3	0.02%

	Quantity	Rate
U.S. Confirmed Malfunctions	25	0.19%
▪ Conductor fracture	2	0.02%
▪ Insulation breach	23	0.17%
U.S. Acute Lead Observations	32	0.24%
▪ Cardiac perforation	3	0.02%
▪ Failure to capture	7	0.05%
▪ Lead dislodgement	15	0.11%
▪ Oversensing	2	0.02%
▪ Other	5	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.
Total Survival [%]	100.0	99.7	99.5	99.0
(95% Confidence Interval)		±0.1	±0.1	±0.3

7.2 ICD Leads

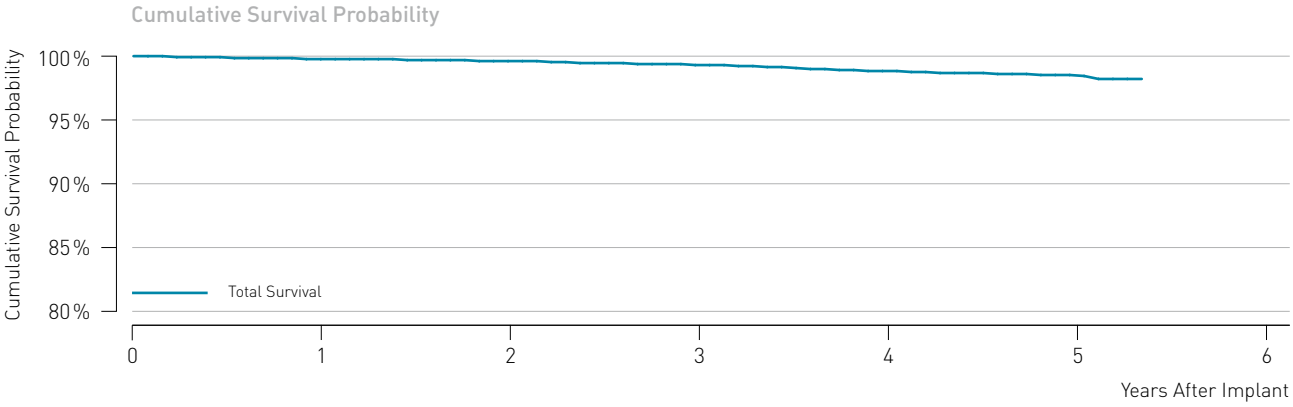
Linox^{smart} SD

Product Details

Product Versions	60/16, 65/16, 65/18, 75/18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jan 2011
CE Market Release	Oct 2009
Worldwide Distributed Devices	53 400
Registered U.S. Implants	13 100
Estimated Active U.S. Implants	11 100
U.S. Total Returned	192

	Quantity	Rate
U.S. Qualifying Complications	114	0.87%
▪ Abnormal defibrillation impedance	7	0.05%
▪ Abnormal pacing impedance	4	0.03%
▪ Conductor fracture	14	0.11%
▪ Failure to capture	10	0.08%
▪ Failure to sense	2	0.02%
▪ Insulation breach	6	0.05%
▪ Lead dislodgement	17	0.13%
▪ Oversensing	48	0.37%
▪ Other	6	0.05%

	Quantity	Rate
U.S. Confirmed Malfunctions	39	0.30%
▪ Conductor fracture	4	0.03%
▪ Insulation breach	35	0.27%
U.S. Acute Lead Observations	30	0.23%
▪ Abnormal defibrillation impedance	1	0.01%
▪ Cardiac perforation	2	0.02%
▪ Failure to capture	4	0.03%
▪ Insulation breach	1	0.01%
▪ Lead dislodgement	13	0.10%
▪ Oversensing	2	0.02%
▪ Other	7	0.05%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	99.7	99.5	99.2	98.7	98.2
(95% Confidence Interval)		±0.1	±0.1	±0.2	±0.2	±0.3

7.2 ICD Leads

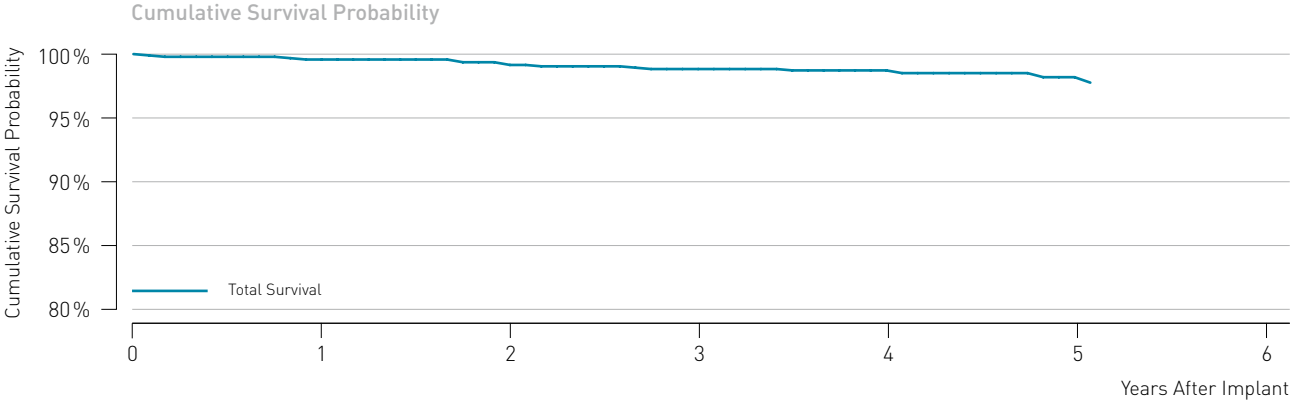
Linox^{smart} TD

Product Details

Product Versions	65/16, 65/18, 75/18
Lead Type	dual-coil, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jan 2011
CE Market Release	Oct 2009
Worldwide Distributed Devices	7 670
Registered U.S. Implants	1 280
Estimated Active U.S. Implants	1 080
U.S. Total Returned	18

	Quantity	Rate
U.S. Qualifying Complications	16	1.25%
▪ Abnormal defibrillation impedance	1	0.08%
▪ Abnormal pacing impedance	1	0.08%
▪ Conductor fracture	1	0.08%
▪ Failure to capture	2	0.16%
▪ Insulation breach	2	0.16%
▪ Lead dislodgement	4	0.31%
▪ Oversensing	5	0.39%

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
U.S. Acute Lead Observations	3	0.24%
▪ Lead dislodgement	3	0.24%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	99.6	99.2	98.9	98.8	98.3
(95% Confidence Interval)		±0.4	±0.5	±0.6	±0.7	±1.0

7.2 ICD Leads

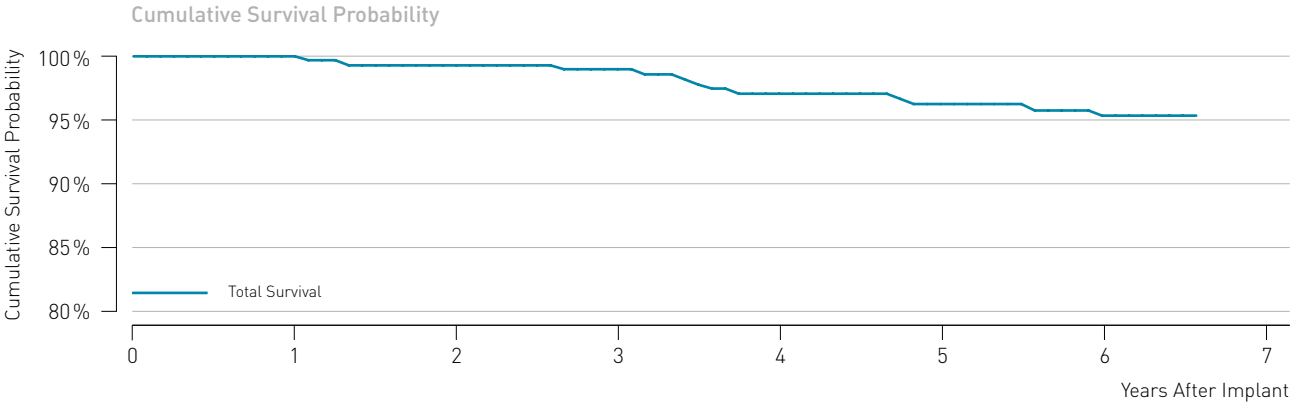
Linox T

Product Details

Product Versions	65, 75
Lead Type	single-coil, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Feb 2007
CE Market Release	Mar 2007
Worldwide Distributed Devices	2 280
Registered U.S. Implants	322
Estimated Active U.S. Implants	226
U.S. Total Returned	3

	Quantity	Rate
U.S. Qualifying Complications	13	4.04%
▪ Abnormal pacing impedance	2	0.62%
▪ Failure to capture	3	0.93%
▪ Insulation breach	1	0.31%
▪ Oversensing	6	1.86%
▪ Other	1	0.31%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.62%
▪ Insulation breach	2	0.62%
U.S. Acute Lead Observations	1	0.31%
▪ Other	1	0.31%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.
Total Survival [%]	100.0	100.0	99.3	99.0	97.1	96.3	95.4
(95% Confidence Interval)			±0.9	±1.2	±2.0	±2.3	±2.5

7.2 ICD Leads

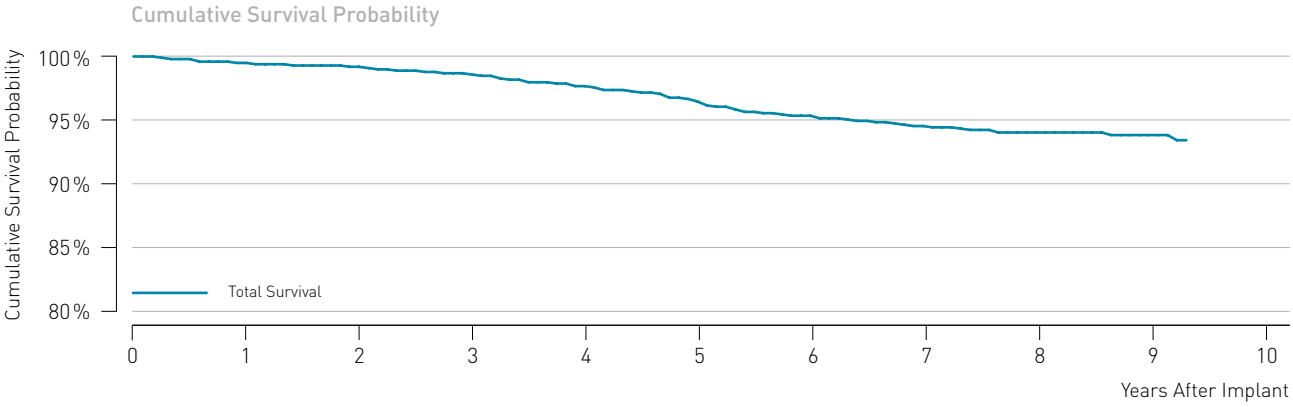
Linox TD

Product Details

Product Versions	65, 75, 100/16, 100/18
Lead Type	dual-coil, passive fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Oct 2006
CE Market Release	Oct 2006
Worldwide Distributed Devices	14 600
Registered U.S. Implants	3 060
Estimated Active U.S. Implants	2 090
U.S. Total Returned	72

	Quantity	Rate
U.S. Qualifying Complications	98	3.21%
▪ Abnormal defibrillation impedance	8	0.26%
▪ Abnormal pacing impedance	10	0.33%
▪ Conductor fracture	11	0.36%
▪ Failure to capture	15	0.49%
▪ Failure to sense	2	0.07%
▪ Insulation breach	13	0.43%
▪ Lead dislodgement	4	0.13%
▪ Oversensing	33	1.08%
▪ Other	2	0.07%

	Quantity	Rate
U.S. Confirmed Malfunctions	35	1.15%
▪ Conductor fracture	6	0.20%
▪ Insulation breach	29	0.95%
U.S. Acute Lead Observations	3	0.10%
▪ Failure to capture	1	0.03%
▪ Lead dislodgement	2	0.07%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.
Total Survival [%]	100.0	99.5	99.2	98.6	97.7	96.5	95.4	94.6	94.1	93.9
(95% Confidence Interval)		±0.3	±0.3	±0.5	±0.6	±0.7	±0.8	±0.9	±1.0	±1.1

7.2 ICD Leads

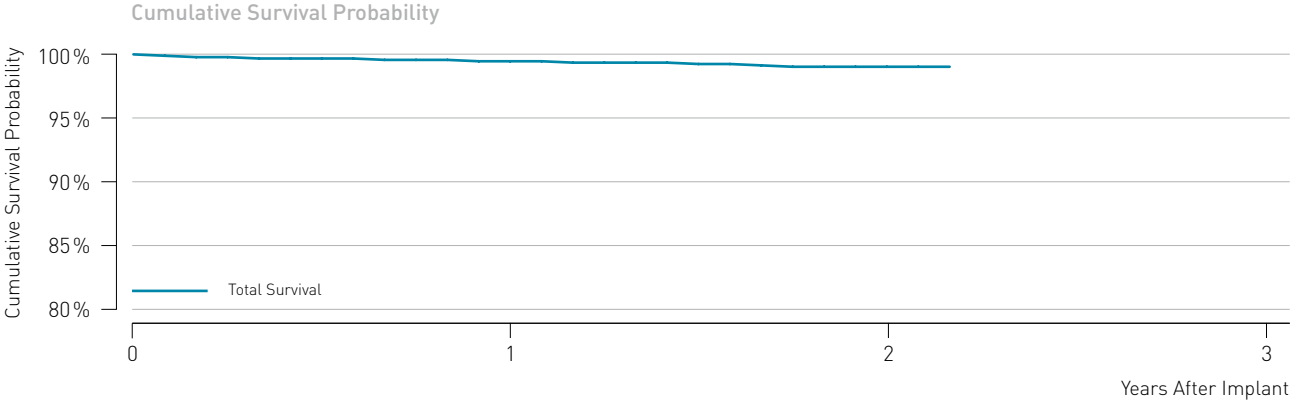
Protego S

Product Details

Product Versions	60, 65, 75
Lead Type	single-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jul 2014
CE Market Release	Feb 2014
Worldwide Distributed Devices	43 300
Registered U.S. Implants	5 780
Estimated Active U.S. Implants	5 450
U.S. Total Returned	33

	Quantity	Rate
U.S. Qualifying Complications	23	0.40%
▪ Cardiac perforation	1	0.02%
▪ Extracardiac stimulation	1	0.02%
▪ Failure to capture	3	0.05%
▪ Lead dislodgement	13	0.22%
▪ Oversensing	1	0.02%
▪ Other	4	0.07%

	Quantity	Rate
U.S. Confirmed Malfunctions	3	0.05%
▪ Conductor fracture	2	0.03%
▪ Insulation breach	1	0.02%
U.S. Acute Lead Observations	21	0.36%
▪ Cardiac perforation	1	0.02%
▪ Extracardiac stimulation	1	0.02%
▪ Failure to capture	3	0.05%
▪ Lead dislodgement	10	0.17%
▪ Other	6	0.10%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	99.5	99.1
(95% Confidence Interval)		±0.2	±0.5

7.2 ICD Leads

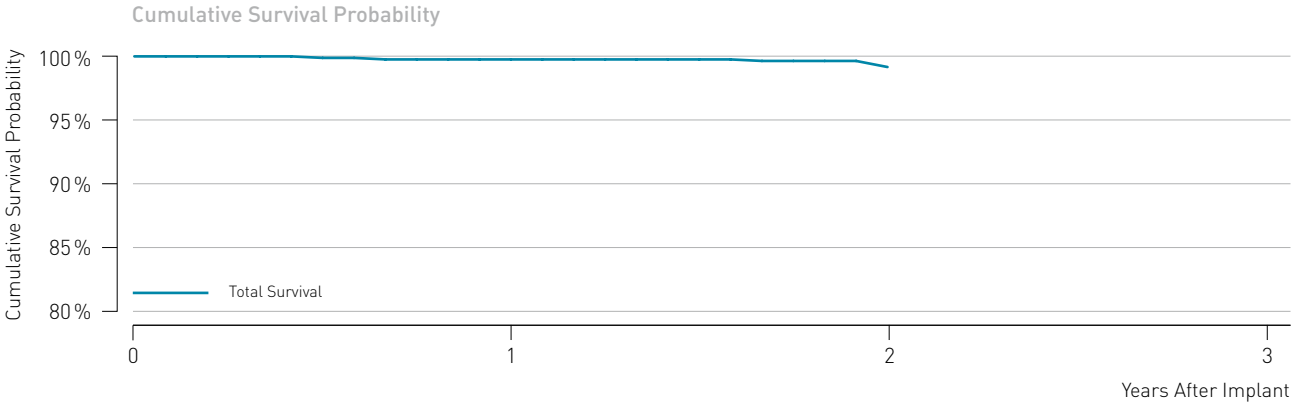
Protego SD

Product Details

Product Versions	60/16, 65/16, 65/18, 75/18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jul 2014
CE Market Release	May 2013
Worldwide Distributed Devices	16 700
Registered U.S. Implants	2 590
Estimated Active U.S. Implants	2 430
U.S. Total Returned	16

	Quantity	Rate
U.S. Qualifying Complications	4	0.15%
▪ Conductor fracture	1	0.04%
▪ Insulation breach	1	0.04%
▪ Lead dislodgement	2	0.08%

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.04%
▪ Insulation breach	1	0.04%
U.S. Acute Lead Observations	3	0.12%
▪ Lead dislodgement	2	0.08%
▪ Other	1	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	99.8	99.3
(95% Confidence Interval)		±0.2	±0.9

7.2 ICD Leads

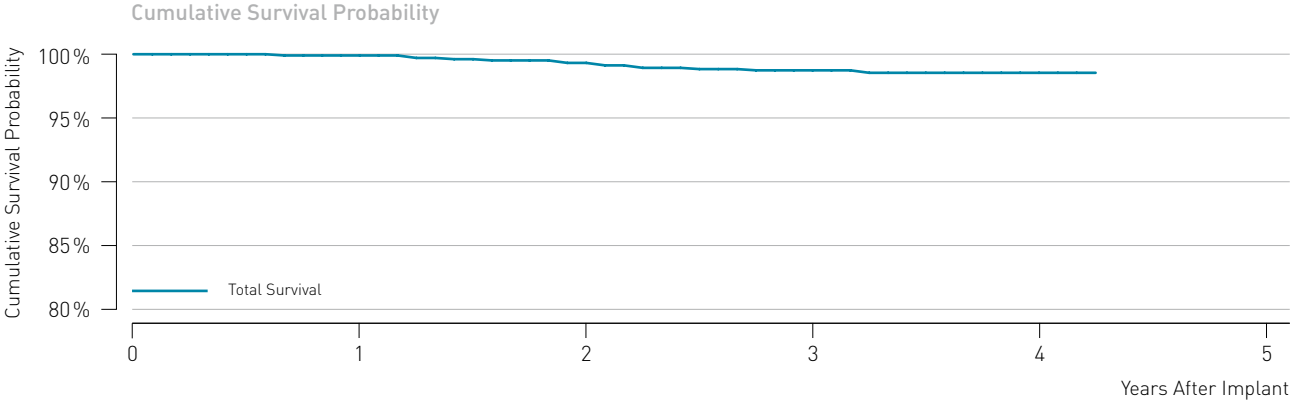
Vigila 2CR

Product Details

Product Versions	60/16, 65/18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Feb 2012
CE Market Release	Oct 2011
Worldwide Distributed Devices	3010
Registered U.S. Implants	799
Estimated Active U.S. Implants	732
U.S. Total Returned	11

	Quantity	Rate
U.S. Qualifying Complications	8	1.00%
▪ Conductor fracture	1	0.13%
▪ Lead dislodgement	3	0.38%
▪ Oversensing	4	0.50%

	Quantity	Rate
U.S. Confirmed Malfunctions	3	0.38%
▪ Insulation breach	3	0.38%
U.S. Acute Lead Observations	0	0.00%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	99.9	99.3	98.7	98.5
(95% Confidence Interval)		±0.3	±0.6	±0.8	±0.9

7.3 CRT Leads

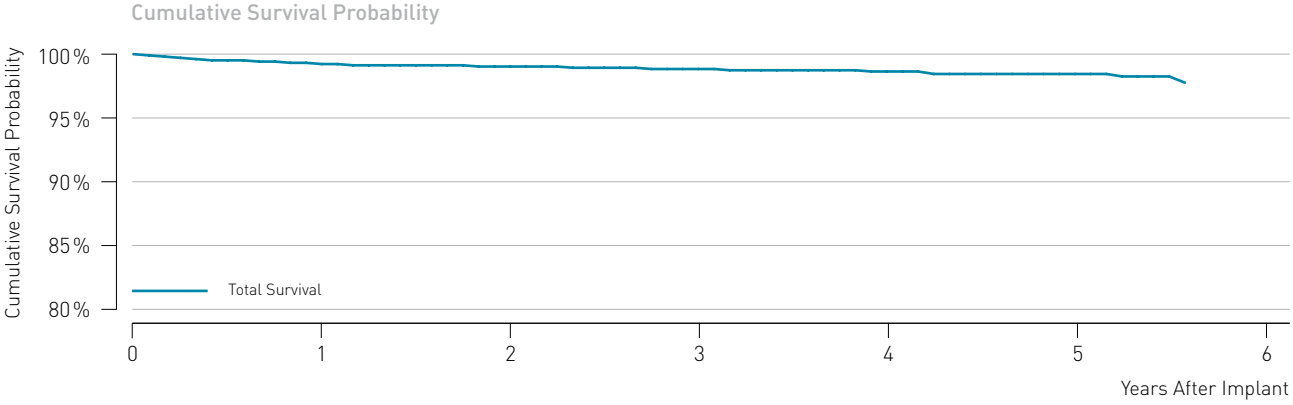
Corox

Product Details

Product Versions	OTW-L 75-BP, 85-BP
Lead Type	dual-curve fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jan 2011
CE Market Release	Dec 2009
Worldwide Distributed Devices	30 300
Registered U.S. Implants	5 880
Estimated Active U.S. Implants	4 920
U.S. Total Returned	56

	Quantity	Rate
U.S. Qualifying Complications	63	1.07%
▪ Conductor fracture	2	0.03%
▪ Extracardiac stimulation	12	0.20%
▪ Failure to capture	20	0.34%
▪ Lead dislodgement	24	0.41%
▪ Oversensing	1	0.02%
▪ Other	4	0.07%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.03%
▪ Conductor fracture	1	0.02%
▪ Insulation breach	1	0.02%
U.S. Acute Lead Observations	20	0.34%
▪ Extracardiac stimulation	6	0.10%
▪ Failure to capture	2	0.03%
▪ Lead dislodgement	9	0.15%
▪ Other	3	0.05%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	99.2	99	98.8	98.6	98.4
(95% Confidence Interval)		±0.2	±0.3	±0.3	±0.4	±0.4

7.3 CRT Leads

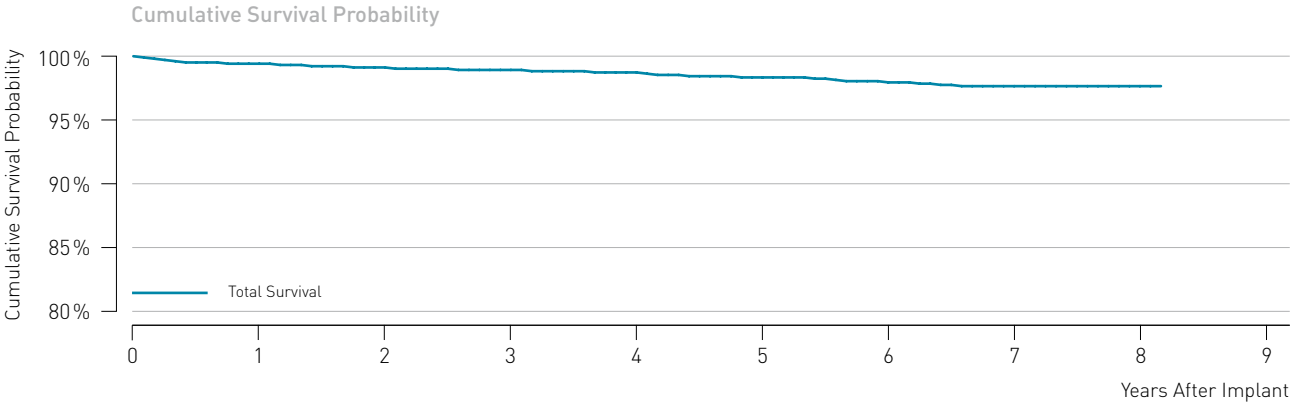
Corox

Product Details

Product Versions	OTW-S 75-BP, 85-BP
Lead Type	thread fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	May 2008
CE Market Release	Dec 2006
Worldwide Distributed Devices	25 700
Registered U.S. Implants	7 940
Estimated Active U.S. Implants	5 710
U.S. Total Returned	112

	Quantity	Rate
U.S. Qualifying Complications	97	1.22%
▪ Abnormal pacing impedance	2	0.03%
▪ Conductor fracture	2	0.03%
▪ Extracardiac stimulation	9	0.11%
▪ Failure to capture	22	0.28%
▪ Insulation breach	4	0.05%
▪ Lead dislodgement	42	0.53%
▪ Oversensing	1	0.01%
▪ Other	15	0.19%

	Quantity	Rate
U.S. Confirmed Malfunctions	11	0.14%
▪ Conductor fracture	6	0.08%
▪ Insulation breach	4	0.05%
▪ Other	1	0.01%
U.S. Acute Lead Observations	29	0.37%
▪ Cardiac perforation	1	0.01%
▪ Extracardiac stimulation	4	0.05%
▪ Failure to capture	5	0.06%
▪ Lead dislodgement	18	0.23%
▪ Other	1	0.01%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.
Total Survival [%]	100.0	99.4	99.1	98.9	98.7	98.3	97.9	97.6	97.6
(95% Confidence Interval)		±0.2	±0.2	±0.3	±0.3	±0.4	±0.4	±0.5	±0.5

7.3 CRT Leads

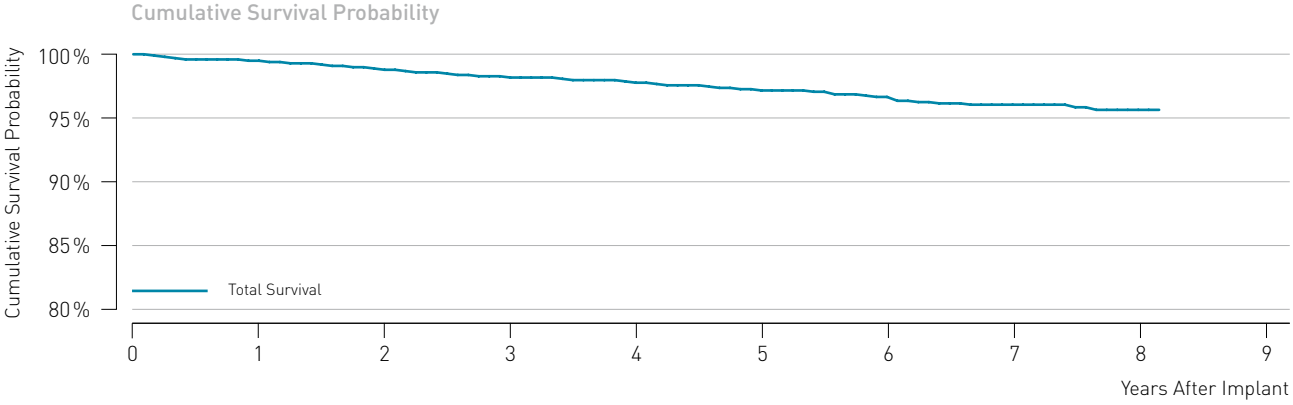
Corox

Product Details

Product Versions	OTW 75-BP Steroid, 85-BP Steroid
Lead Type	helix fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	May 2008
CE Market Release	Dec 2006
Worldwide Distributed Devices	27 700
Registered U.S. Implants	4 100
Estimated Active U.S. Implants	2 710
U.S. Total Returned	71

	Quantity	Rate
U.S. Qualifying Complications	84	2.05%
▪ Abnormal pacing impedance	2	0.05%
▪ Conductor fracture	3	0.07%
▪ Extracardiac stimulation	8	0.20%
▪ Failure to capture	28	0.68%
▪ Insulation breach	2	0.05%
▪ Lead dislodgement	30	0.73%
▪ Oversensing	2	0.05%
▪ Other	9	0.22%

	Quantity	Rate
U.S. Confirmed Malfunctions	16	0.39%
▪ Conductor fracture	15	0.37%
▪ Insulation breach	1	0.02%
U.S. Acute Lead Observations	8	0.20%
▪ Lead dislodgement	6	0.15%
▪ Other	2	0.05%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.
Total Survival [%]	100.0	99.5	98.8	98.2	97.8	97.2	96.7	96.1	95.7
(95% Confidence Interval)		±0.2	±0.4	±0.5	±0.5	±0.6	±0.7	±0.8	±1.0

7.3 CRT Leads

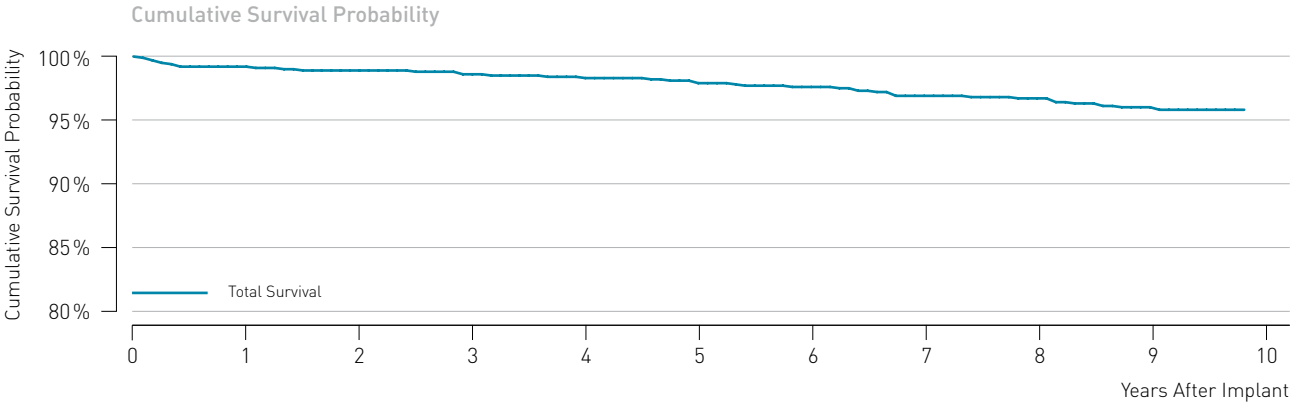
Corox

Product Details

Product Versions	OTW 75-UP Steroid, 85-UP Steroid
Lead Type	helix fixation
Polarity	unipolar
Steroid	yes
U.S. Market Release	Aug 2006
CE Market Release	Apr 2004
Worldwide Distributed Devices	10 400
Registered U.S. Implants	1 430
Estimated Active U.S. Implants	703
U.S. Total Returned	26

	Quantity	Rate
U.S. Qualifying Complications	37	2.60%
▪ Extracardiac stimulation	7	0.49%
▪ Failure to capture	13	0.91%
▪ Insulation breach	2	0.14%
▪ Lead dislodgement	9	0.63%
▪ Oversensing	1	0.07%
▪ Other	5	0.35%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.14%
▪ Insulation breach	2	0.14%
U.S. Acute Lead Observations	4	0.28%
▪ Failure to capture	3	0.21%
▪ Lead dislodgement	1	0.07%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.	6 yr.	7 yr.	8 yr.	9 yr.
Total Survival [%]	100.0	99.2	98.9	98.6	98.3	97.9	97.6	96.9	96.7	96.0
(95% Confidence Interval)		±0.5	±0.6	±0.7	±0.7	±0.8	±0.9	±1.1	±1.1	±1.3

8 Methodology for Lead Survival Estimates Based on Clinical Studies

8.1 Introduction

All leads and lead segments returned to BIOTRONIK are thoroughly analyzed to determine whether or not they meet BIOTRONIK's long term quality standards.

Although analysis of returned product is an excellent method for gaining insight into lead failure mechanisms, this data relies on the return of explanted leads. For the majority of complications the lead is not received for analysis as challenging clinical environments may not allow for the return, e.g. the extraction of an implanted lead may not be possible.

BIOTRONIK includes all reported chronic complications in the calculation of the survival estimates as described in chapter 6, i.e. reports with returned and without returned products.

However, BIOTRONIK can only report events in the survival estimates if those events were reported to BIOTRONIK. In order to eliminate possible biased survival estimates due to underreporting, BIOTRONIK performs clinical surveillance studies with active follow-up's under FDA guidance yielding the most reliable lead performance data.

In the following chapter BIOTRONIK shows – in addition to the survival data based on returned product analysis and chronic complication information – the lead performance data from clinical trials. These studies are designed to record clinical observations representative of the total clinical experience.

8.2 BIOTRONIK's Clinical Studies

8.2.1 GALAXY and CELESTIAL

BIOTRONIK's GALAXY and CELESTIAL Registries are prospective, non-randomized, observational studies. The key purpose of these registries is to confirm the long-term safety and reliability of BIOTRONIK leads as used in conjunction with a BIOTRONIK ICD (GALAXY) or CRT (CELESTIAL) system. All devices in the registries are legally marketed and available to physicians according to approved FDA indications for use. GALAXY and CELESTIAL Registries are registered on clinicaltrials.gov under NCT00836589 and NCT00810264 respectively.

The evaluation of safety for GALAXY is based on the analysis of BIOTRONIK Linox ICD lead-related adverse events. The evaluation of safety for CELESTIAL is based on the analysis of BIOTRONIK Corox LV pacing lead-related adverse events. However, many CELESTIAL patients also have a Linox ICD lead implanted and the Linox clinical studies data in this report represents combined data from the GALAXY and CELESTIAL registries. Both registries are designed to continue for a 5 year follow-up duration per patient. To ensure a sufficiently large and representative source of data, participating clinical sites must meet pre-specified selection criteria.

Every effort is made to ensure participants are representative of the range of clinical environments in which BIOTRONIK's cardiac rhythm products are used. Patients will be seen for routine follow-up visits per their institutional standard of care until they can no longer be followed (e.g., death, lost to follow-up, etc.). However, to ensure regular patient status assessments are completed, follow-up windows consistent with typical care practices have been established with a minimum of once every six months follow-up requirement.

During each follow up at a study center the following steps are required during the follow-up visit:

- Interrogate programmed parameters
- Determine lead electrical parameters
- Evaluate device diagnostics and programmed parameters to ensure the device is providing appropriate therapy
- Determine if there are any reportable lead-related, pulse generator-related or implant procedure-related adverse events. If there are, complete an adverse event electronic case report form (eCRF)
- Complete all appropriate eCRFs

This active surveillance model ensures a robust dataset for effectively monitoring product performance.

Patient Enrolment Criteria

To support the objectives of these registries, participants are required to meet the following inclusion criteria prior to enrollment:

- Successfully implanted BIOTRONIK ICD (GALAXY) or BIOTRONIK CRT (CELESTIAL) system, including the study lead
- Able to understand the nature of the registry and give informed consent
- Available for follow-up visits on a regular basis at the study site
- Age greater than or equal to 18 years

Each site must inform BIOTRONIK whenever a lead complication has occurred or when a patient is no longer participating.

All leads that experience a complication and are subsequently explanted and returned to BIOTRONIK undergo root-cause analyses. Product performance is analyzed as a function of time using the survival analysis method. Root causes for any failures, regardless of the incidence rates, are investigated.

8.2 BIOTRONIK's Clinical Studies

8.2.2 SIELLO Clinical Study

BIOTRONIK's SIELLO Clinical Study is a prospective, non-randomized, combined Pre-Market Study and Post-Approval Registry designed to demonstrate the safety and effectiveness of the Siello pacing lead as used in conjunction with any market-released BIOTRONIK pacemaker device. The SIELLO Clinical Study is registered on clinicaltrials.gov under NCT01791127.

For the Pre-Market Study, the evaluation of safety is based on the analysis of Siello lead-related adverse events through a follow-up time of 12 months post-implant, while the evaluation of effectiveness is based on analysis of the success rate of the implanted system including one or two Siello leads to sense and deliver pacing at 12 months post-implant.

For the Post-Approval Registry, the evaluation of long-term safety is based on the analysis of Siello lead-related adverse events through a follow-up time of 5 years post-implant. To ensure a sufficiently large and representative source of data, participating clinical sites must meet pre-specified selection criteria.

Every effort is made to ensure participants are representative of the range of clinical environments in which BIOTRONIK's cardiac rhythm products are used. Patients will be seen for routine follow-up visits until they can no longer be followed (e.g., death, lost to follow-up, etc.). However, to ensure regular patient status assessments are completed, a study follow-up schedule consistent with typical care practices has been established, with required follow-ups at 3 and 6 months post-implant and every 6 months thereafter.

During each study follow-up visit the following steps are required:

- Interrogate programmed parameters
- Record electrical parameters of the implanted leads. Perform all pacing threshold measurements at 0.4 ms or 0.5 ms pulse width when feasible
- Evaluate device diagnostics, electrical parameters and programmed parameters to ensure the device is correctly pacing and sensing
- Determine if there are any lead-related, pulse generator-related or procedure related adverse events. If any are recorded, complete the Adverse Event eCRF
- Complete all appropriate eCRFs

This active surveillance model ensures a robust dataset for effectively monitoring product performance.

Patient Enrolment Criteria

To support the objectives of the study, participants are required to meet the following inclusion criteria prior to enrollment:

- Candidate for de novo implantation of a market-released BIOTRONIK pacemaker system, including one or two Siello leads. Candidate meets recommendation for pacemaker system implant put forth by guidelines of relevant professional societies
- Able to understand the nature of the study and provide informed consent
- Available for follow-up visits on a regular basis at the investigational site for the expected 5 years of follow-up
- Age greater than or equal to 18 years

Each site must inform BIOTRONIK whenever a lead complication has occurred or when a patient is no longer participating.

All leads that experience a complication and are subsequently explanted and returned to BIOTRONIK undergo root-cause analyses. Product performance is analyzed as a function of time using the survival analysis method. Root causes for any failures, regardless of the incidence rates, are investigated.

8.2 BIOTRONIK's Clinical Studies

8.2.3 Protego Post-Approval Registry

BIOTRONIK's Protego Post-Approval Registry is a prospective, single-arm, non-randomized registry to confirm the long-term safety and reliability of the Protego DF4 ICD lead when used in conjunction with a BIOTRONIK DF4 compatible ICD or CRT-D pulse generator. The Protego DF4 Post-Approval Registry is registered on clinicaltrials.gov under NCT02243696.

The evaluation of safety will be based on the analysis of Protego DF4 lead related or header related adverse events through a follow-up time of 5 years post-implant. To ensure a sufficiently large and representative source of data, participating clinical sites must meet pre-specified selection criteria.

Every effort is made to ensure participants are representative of the range of clinical environments in which BIOTRONIK's cardiac rhythm products are used. Patients will be seen for routine follow-up visits per their institutional standard of care until they can no longer be followed (e.g., death, lost to follow-up, etc.). However, to ensure regular patient status assessments are completed, a study follow-up schedule consistent with typical care practices has been established, with required follow-ups at 3 and 6 months post-implant, and every 6 months thereafter.

During each study visit, the following are required:

- Interrogate programmed parameters
- Record electrical parameters of the implanted leads. Perform all pacing threshold measurements at 0.4 ms or 0.5 ms pulse width when feasible
- Evaluate device diagnostics and programmed parameters to ensure the device is providing appropriate therapy
- Determine if there are any lead-related, pulse generator-related or procedure related adverse events. If any are recorded, complete the Adverse Event eCRF
- Complete all appropriate eCRFs

This active surveillance model ensures a robust dataset for effectively monitoring product performance.

Patient Enrolment Criteria

To support the objectives of the study, participants are required to meet the following inclusion criteria prior to enrollment:

- Candidate for implant or successfully implanted with a BIOTRONIK ICD or BIOTRONIK CRT-D system, including a Protego lead
- Able to understand the nature of the registry and give informed consent
- Available for follow-up visits on a regular basis at the study site
- Age greater than or equal to 18 years

Each site must inform BIOTRONIK whenever a lead complication has occurred or when a patient is no longer participating.

All leads that experience a complication and are subsequently explanted and returned to BIOTRONIK undergo root-cause analyses. Product performance is analyzed as a function of time using the survival analysis method. Root causes for any failures, regardless of the incidence rates, are investigated.

8.3 Lead Complications

The data presented characterizes chronic lead performance by estimating lead-related complication free survival probabilities. Following industry practice, for analysis purposes, the complication criteria, which align with the AdvaMed 'Industry Guidance for Uniform Reporting of Clinical Performance of Cardiac Rhythm Management Pulse Generators and Leads', are defined below.

8.3.1 GALAXY and CELESTIAL

All reported lead-related adverse events within the GALAXY and CELESTIAL Registries are classified by the reporting investigator and are adjudicated by an independent event adjudication committee. A lead related complication is considered to have occurred if a clinical observation happens after successful implantation, is classified with at least one of the following event classifications and at least one of the following clinical actions is made. Any Clinical Event without a related Clinical Action is not considered a Qualifying Complication. Events with an onset date 30 days or less after the implant are acute observations and are listed separately.

Event Classifications

- Failure to capture
- Failure to sense/undersensing
- Oversensing
- Abnormal pacing impedance (based on lead model, but normal range is typically 200- 2,000 ohms)
- Abnormal defibrillation impedance (based on lead model, but normal range is 25 – 150 ohms)
- Insulation breach
- Conductor fracture, confirmed electrically, visually or radiographically
- Extracardiac stimulation
- Cardiac perforation
- Lead dislodgement

Clinical Actions

- Lead surgically abandoned/capped
- Lead electrically abandoned
- Lead explanted
- Lead replaced
- Lead conductor taken out of service
- Lead use continued based on medical judgment despite a known clinical performance issue
- Other lead-related surgery

8.3 Lead Complications

8.3.2 SIELLO

All reported lead-related adverse events within the SIELLO Clinical Study are classified by the reporting investigator and are adjudicated by an independent event adjudication committee. A lead related complication is considered to have occurred if a clinical observation happens after successful implantation and is classified with at least one of the following event classifications. Events with an onset date 30 days or less after the implant are acute observations and are listed separately.

Event Classifications

- Failure to capture
- Failure to sense/undersensing
- Oversensing
- Abnormal pacing impedance (based on lead model, but normal range is typically 200–2,000 ohms)
- Insulation breach
- Conductor fracture, confirmed electrically, visually or radiographically
- Extracardiac stimulation
- Cardiac perforation
- Lead dislodgement

8.3.3 Protego

All reported lead-related adverse events within the Protego Registry are classified by the reporting investigator and are adjudicated by an independent event adjudication committee. A lead related complication is considered to have occurred if a clinical observation happens after successful implantation and is classified with at least one of the following event classifications. Events with an onset date 30 days or less after the implant are acute observations and are listed separately.

Event Classification

- Failure to capture
- Failure to sense/undersensing
- Oversensing
- Abnormal pacing impedance (based on lead model, but normal range is typically 200–2,000 ohms)
- Abnormal defibrillation impedance (based on lead model, but normal range is 25–150 ohms)
- Insulation breach
- Conductor fracture, confirmed electrically, visually or radiographically
- Extracardiac stimulation
- Cardiac perforation
- Lead dislodgement

8.4 Lead Product Performance Graphs and Data

The clinical data presented on the following page is intended to show the long term clinical performance of leads based on clinical studies. The same analysis methods as described in chapter 6 are applied.

Returned Product Analysis Results

Although the returned product analysis data is not used to generate the survival estimates for the clinical data, it provides valuable insight into the causes of lead malfunction. Following the same approach as for complaint data, a malfunction is reported as described in section 6.3 of this report.

9 Performance of BIOTRONIK Leads Based on Clinical Studies

9.1 Performance of Pacing Leads

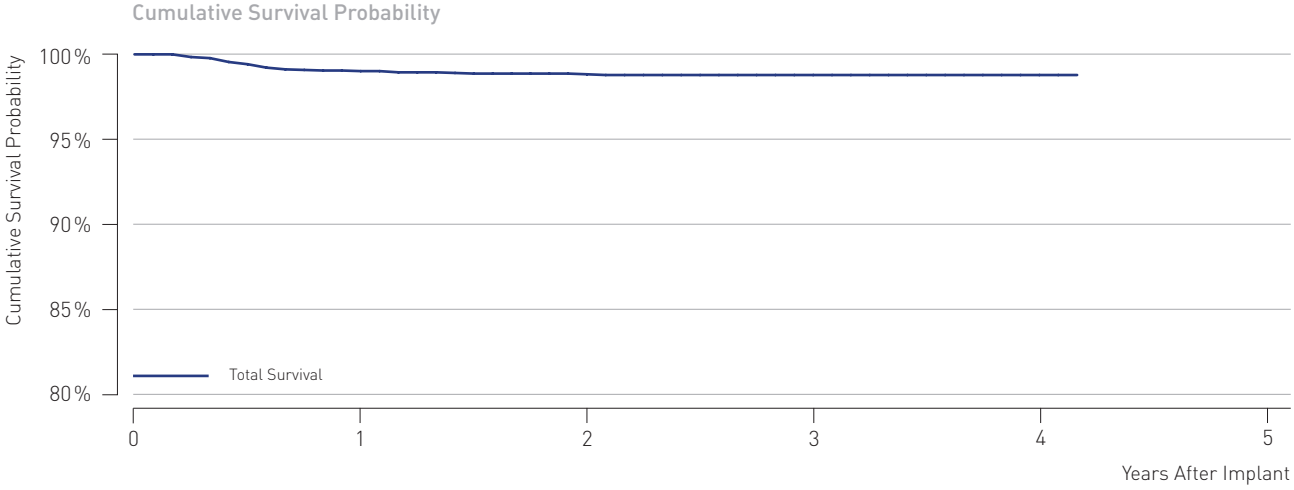
Siello S / Solia S Study Data

Product Details

Product Versions	45, 53, 60
Lead Type	straight, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jun 2016
CE Market Release	Jul 2009
Worldwide Distributed Devices	674 000
Registered U.S. Implants	3 236

	Quantity	Rate
U.S. Qualifying Complications	36	1.11%
▪ Cardiac perforation	1	0.03%
▪ Conductor fracture	1	0.03%
▪ Failure to capture	16	0.49%
▪ Failure to sense (undersensing)	8	0.25%
▪ Lead dislodgement	8	0.25%
▪ Oversensing	1	0.03%
▪ Other	1	0.03%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.06%
▪ Conductor fracture	1	0.03%
▪ Insulation breach	1	0.03%
U.S. Acute Lead Observations	26	0.80%
▪ Cardiac perforation	8	0.25%
▪ Extracardiac stimulation	2	0.06%
▪ Failure to capture	6	0.19%
▪ Failure to sense (undersensing)	5	0.15%
▪ Lead dislodgement	5	0.15%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	99.0	98.8	98.8	98.8
(95% Confidence Interval)		±0.4	±0.4	±0.4	±0.4

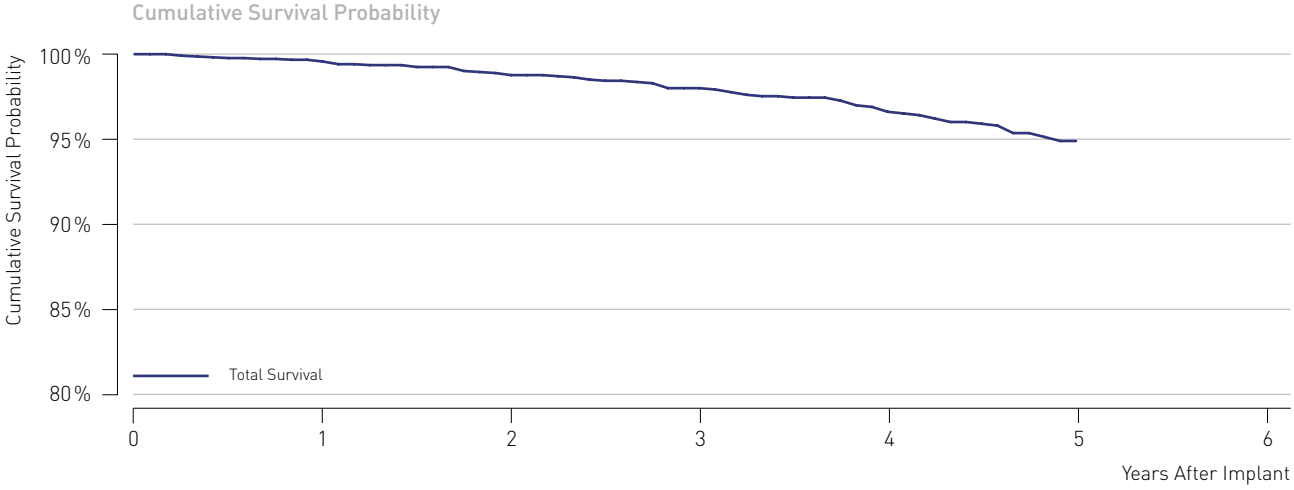
Linox SD Study Data

Product Details

Product Versions	60/16, 65/16, 65/18, 75/18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Apr 2006
U.S. Study Begin	Aug 2008
Worldwide Distributed Devices	55 100
Registered U.S. Implants	2 272

	Quantity	Rate
U.S. Qualifying Complications	67	2.95%
▪ Abnormal defibrillation impedance	4	0.18%
▪ Abnormal pacing impedance	9	0.40%
▪ Cardiac perforation	1	0.04%
▪ Conductor fracture	10	0.44%
▪ Failure to capture	7	0.31%
▪ Failure to sense	1	0.04%
▪ Failure to sense (undersensing)	2	0.09%
▪ Insulation breach	13	0.57%
▪ Lead dislodgement	3	0.13%
▪ Oversensing	17	0.75%

	Quantity	Rate
U.S. Confirmed Malfunctions	22	0.97%
▪ Conductor fracture	3	0.13%
▪ Insulation breach	19	0.84%
U.S. Acute Lead Observations	9	0.40%
▪ Cardiac perforation	4	0.18%
▪ Conductor fracture	1	0.04%
▪ Failure to capture	2	0.09%
▪ Lead dislodgement	1	0.04%
▪ Other	1	0.04%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	99.6	98.8	98.0	96.7	95.0
(95% Confidence Interval)		±0.3	±0.5	±0.7	±0.9	±1.2

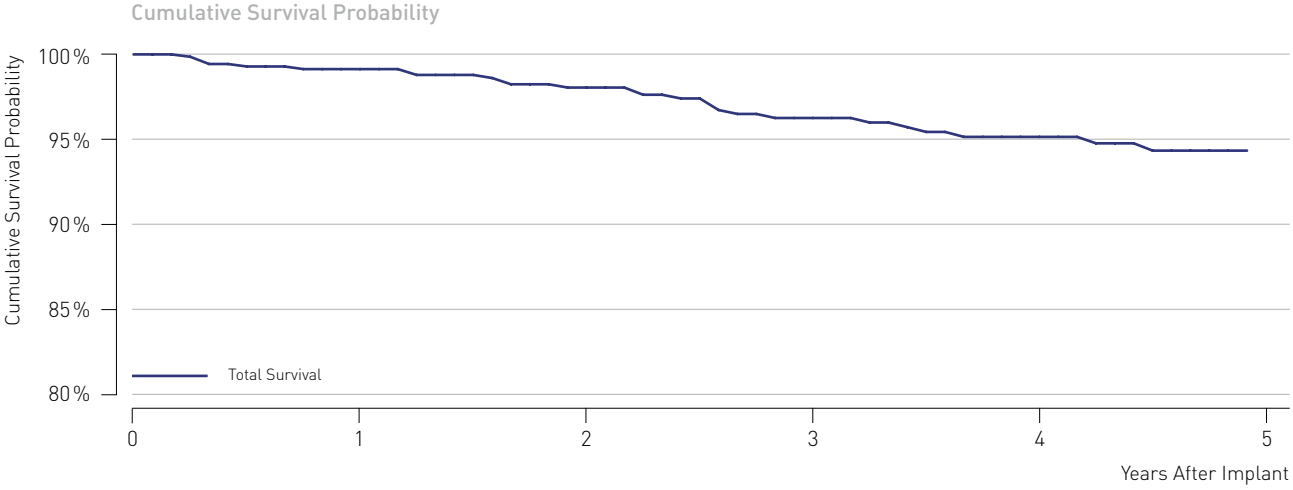
Linox^{smart} SD Study Data

Product Details

Product Versions	60/16, 65/16, 65/18, 75/18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jan 2011
CE Market Release	Oct 2009
Worldwide Distributed Devices	53 400
Registered U.S. Implants	736

	Quantity	Rate
U.S. Qualifying Complications	27	3.67%
▪ Abnormal defibrillation impedance	2	0.27%
▪ Abnormal pacing impedance	1	0.14%
▪ Conductor fracture	3	0.41%
▪ Failure to capture	2	0.27%
▪ Insulation breach	4	0.54%
▪ Lead dislodgement	6	0.82%
▪ Oversensing	9	1.22%

	Quantity	Rate
U.S. Confirmed Malfunctions	7	0.95%
▪ Insulation breach	7	0.95%
U.S. Acute Lead Observations	2	0.27%
▪ Lead dislodgement	2	0.27%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.
Total Survival [%]	100.0	99.1	98.1	96.3	95.2
(95% Confidence Interval)		±0.8	±1.1	±1.7	±2.0

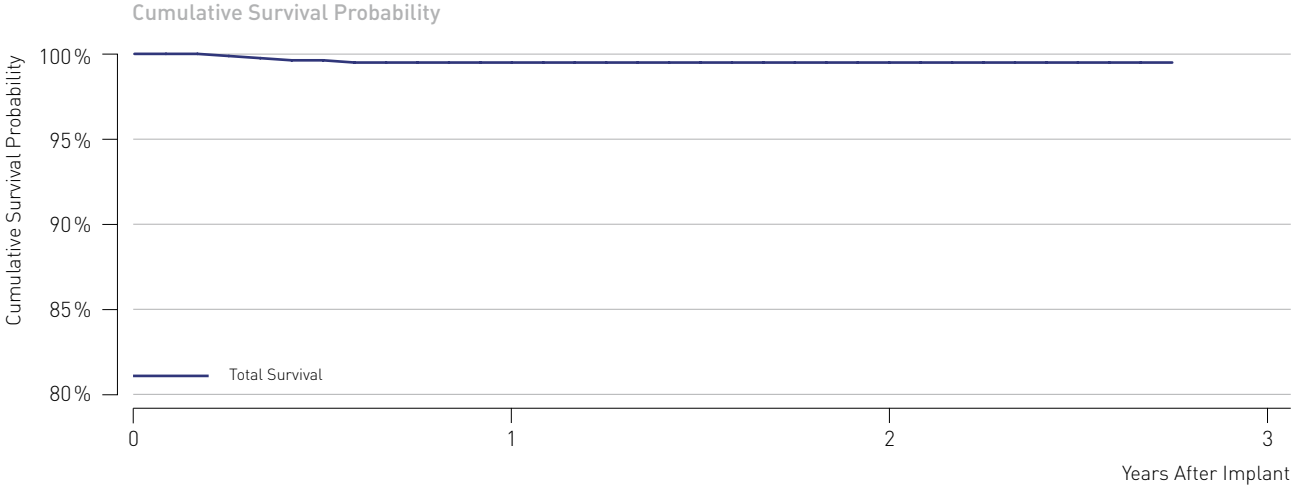
Protego S Study Data

Product Details

Product Versions	60, 65, 75
Lead Type	single-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jul 2014
CE Market Release	Feb 2014
Worldwide Distributed Devices	43 300
Registered U.S. Implants	1 085

	Quantity	Rate
U.S. Qualifying Complications	4	0.37%
▪ Conductor fracture	1	0.09%
▪ Failure to sense (undersensing)	1	0.09%
▪ Lead dislodgement	2	0.18%

	Quantity	Rate
U.S. Confirmed Malfunctions	2	0.18%
▪ Conductor fracture	2	0.18%
U.S. Acute Lead Observations	4	0.37%
▪ Cardiac perforation	3	0.28%
▪ Lead dislodgement	1	0.09%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	99.6	99.6
(95% Confidence Interval)		±0.4	±0.4

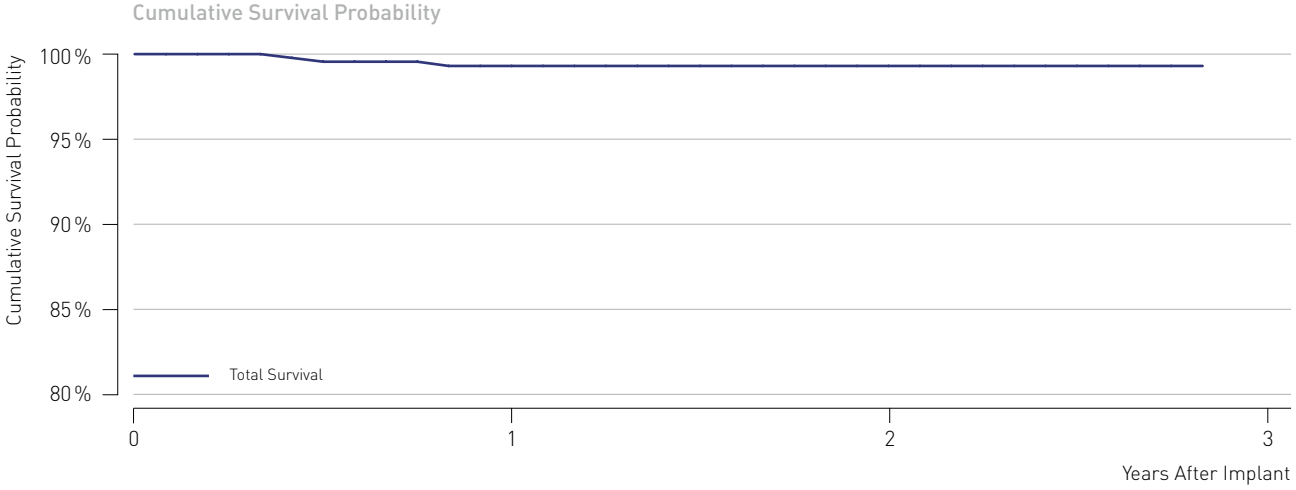
Protego SD Study Data

Product Details

Product Versions	60/16, 65/16, 65/18, 75/18
Lead Type	dual-coil, active fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jul 2014
CE Market Release	May 2013
Worldwide Distributed Devices	16 700
Registered U.S. Implants	533

	Quantity	Rate
U.S. Qualifying Complications	3	0.56%
▪ Abnormal pacing impedance	1	0.19%
▪ Conductor fracture	1	0.19%
▪ Failure to capture	1	0.19%

	Quantity	Rate
U.S. Confirmed Malfunctions	0	0.00%
U.S. Acute Lead Observations	2	0.38%
▪ Lead dislodgement	2	0.38%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.
Total Survival [%]	100.0	99.4	99.4
(95% Confidence Interval)		±0.8	±0.8

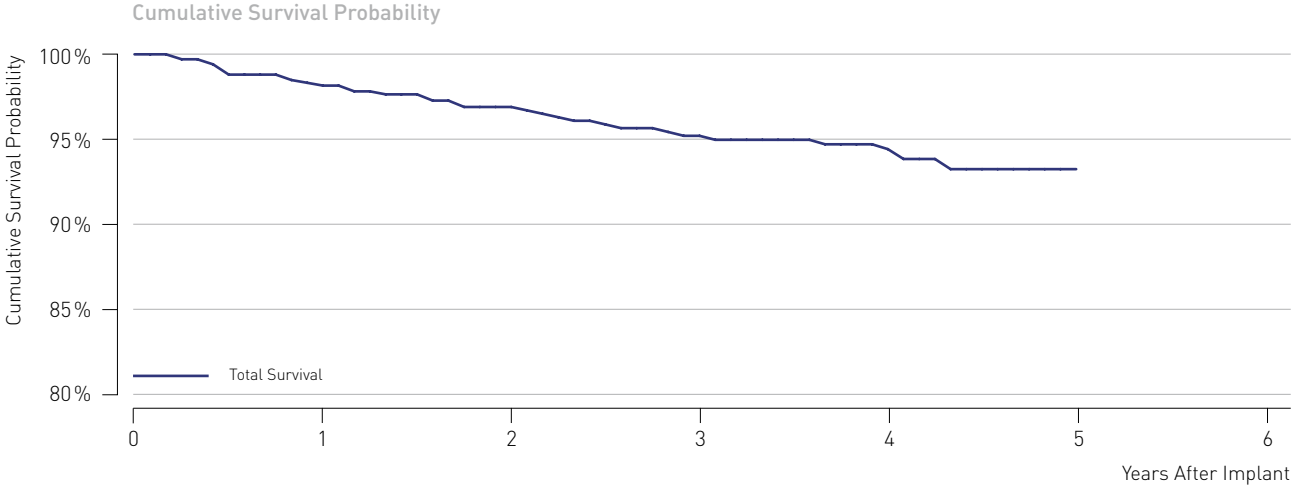
Corox Study Data

Product Details

Product Versions	OTW 75-BP Steroid, 85-BP Steroid
Lead Type	helix fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	May 2008
CE Market Release	Dec 2006
Worldwide Distributed Devices	27 700
Registered U.S. Implants	696

	Quantity	Rate
U.S. Qualifying Complications	34	4.89%
▪ Abnormal pacing impedance	5	0.72%
▪ Conductor fracture	5	0.72%
▪ Extracardiac stimulation	3	0.43%
▪ Failure to capture	5	0.72%
▪ Lead dislodgement	16	2.30%

	Quantity	Rate
U.S. Confirmed Malfunctions	6	0.86%
▪ Conductor fracture	6	0.86%
U.S. Acute Lead Observations	4	0.57%
▪ Extracardiac stimulation	1	0.14%
▪ Lead dislodgement	3	0.43%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	98.2	96.9	95.2	94.4	93.3
(95% Confidence Interval)		±1.1	±1.4	±1.8	±2.0	±2.3

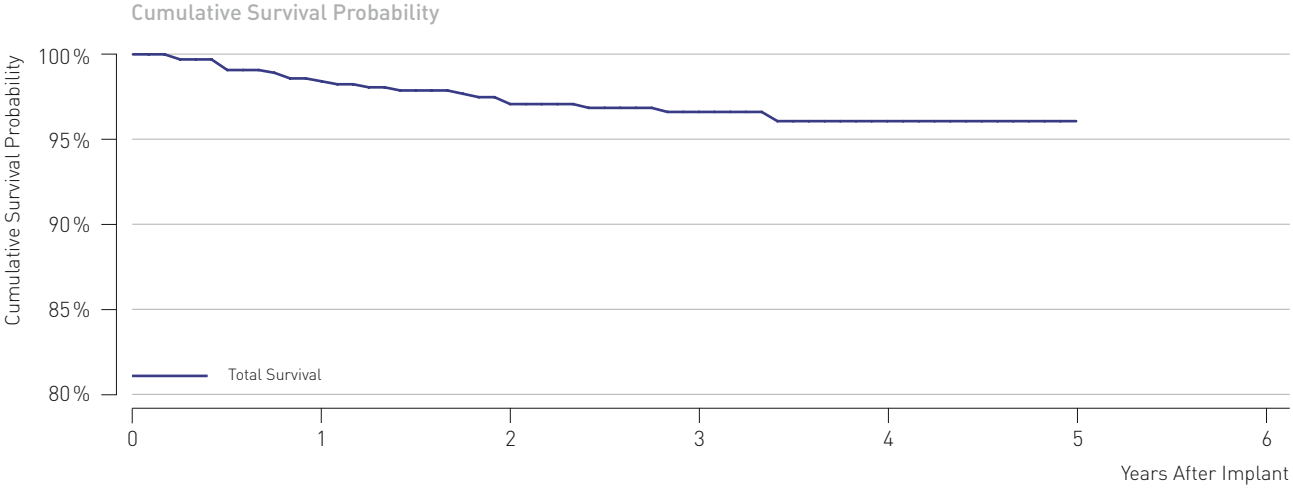
9.3 Performance of Corox Pacing Leads

Corox Study Data

Product Details

Product Versions	OTW-L 75-BP, Corox OTW-L 85-BP
Lead Type	dual-curve fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	Jan 2011
CE Market Release	Dec 2009
Worldwide Distributed Devices	30 300
Registered U.S. Implants	698

	Quantity	Rate		Quantity	Rate
U.S. Qualifying Complications	21	3.01%	U.S. Confirmed Malfunctions	0	0.00%
▪ Extracardiac stimulation	4	0.57%	U.S. Acute Lead Observations	4	0.57%
▪ Failure to capture	7	1.00%	▪ Extracardiac stimulation	3	0.43%
▪ Lead dislodgement	10	1.43%	▪ Lead dislodgement	1	0.14%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	98.5	97.1	96.7	96.2	96.2
(95% Confidence Interval)		±0.1	±1.4	±1.5	±1.7	±1.7

9.3 Performance of Corox Pacing Leads

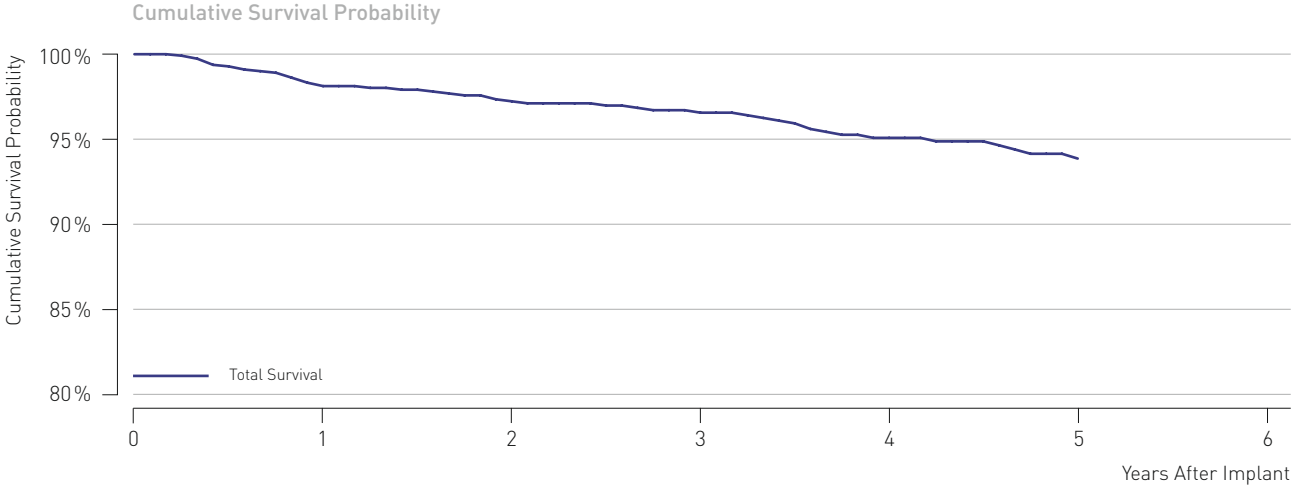
Corox Study Data

Product Details

Product Versions	OTW-S 75-BP, 85-BP
Lead Type	thread fixation
Polarity	bipolar
Steroid	yes
U.S. Market Release	May 2008
CE Market Release	Dec 2006
Worldwide Distributed Devices	25 700
Registered U.S. Implants	1 141

	Quantity	Rate
U.S. Qualifying Complications	47	4.12%
▪ Abnormal pacing impedance	10	0.88%
▪ Extracardiac stimulation	9	0.79%
▪ Failure to capture	9	0.79%
▪ Lead dislodgement	19	1.67%

	Quantity	Rate
U.S. Confirmed Malfunctions	1	0.09%
▪ Insulation breach	1	0.09%
U.S. Acute Lead Observations	6	0.53%
▪ Extracardiac stimulation	1	0.09%
▪ Failure to capture	1	0.09%
▪ Lead dislodgement	4	0.35%



Cumulative Survival Probability after	Impl.	1 yr.	2 yr.	3 yr.	4 yr.	5 yr.
Total Survival [%]	100.0	98.1	97.2	96.6	95.1	93.8
(95% Confidence Interval)		±0.8	±1.0	±1.2	±1.5	±1.9

10 Advisories

Stratos LV-T

Potentially defective low voltage capacitors
84 devices world-wide, none in the United States

Status Update

As of July 2017

- No reports of malfunctioning pacemakers associated with this capacitor lot were received.
- No reports of deaths or serious injuries were received associated with this advisory.
- An extended test program yielded no malfunctions of the capacitors under question.

Original communication July 2006

A limited number of our Stratos LV-T pacemakers may exhibit a device malfunction resulting in increased battery current and in loss of output, very likely simultaneously on all channels.

BIOTRONIK has identified a specific batch of low voltage capacitors from a single component supplier as the root cause. Please be assured that the capacitors from this batch are no longer being used in any BIOTRONIK device, and that no other pacemaker or ICD manufactured by BIOTRONIK is affected.

To date, no field reports related to this phenomenon were received.

The anomaly is limited to 84 devices world-wide. BIOTRONIK has taken immediate action to retrieve all potentially affected devices not yet delivered to hospitals. According to our records, 14 devices were delivered to hospitals and may have been implanted.

Based on preliminary data received from the component supplier, the projected rate of occurrence is expected to less than 1 out of 1000 devices.

Our records show that you are the implanting and/or follow-up physician for one or more patients with an affected device. We recommend the following actions:

- If Home Monitoring is activated, please check the Cardio Report and the pacing impedance history. A rapid and significant increase of the pacing impedance on one or more channels may indicate a possible loss of output. In this case we recommend to schedule a patient follow-up as soon as possible.
- If Home Monitoring is not activated or pacing impedance data from Cardio Reports are not available we recommend to schedule a patient follow-up as soon as possible.
- During a follow-up session please perform the "Battery Lead Telemetry" test. A significantly increased pacing impedance on one or more channels or a significantly increased battery current may indicate a possible loss of output.
- If no anomaly is detected please activate Home Monitoring. You may use regular follow-up intervals if Home Monitoring is used for remote monitoring between follow-ups.
- In any case we recommend to replace the device of a pacemaker-dependent patient without underlying intrinsic rhythm.

We have tried to be as specific as possible with our recommendations. As always, individual circumstances and medical judgement determine decisions about patient care, the urgency of follow-ups and possible replacement of devices.

If a Stratos LV-T is replaced for the reason explained in this notification, BIOTRONIK will provide a replacement device consistent with the terms of our warranty. For activation of the warranty, please return the explanted device to BIOTRONIK within 30 days of the explantation.

11 X-Ray Identifiers for Pacemakers and ICDs

Pacemaker/ICD Product Versions	X-Ray ID
Cylos DR, DR-T, VR	RZ
Cylos 990 DR, DR-T, VR	FV
Evia DR, DR–T, SR, SR-T, HF, HF-T	SF
Estella SR, SR-T, DR, DR-T	SF
Entovis DR, DR-T, SR, SR-T	SF
Eluna 8 DR, Eluna 8 DR-T	SF
Etrinsa 8 VR-T, DR-T, HF-T	BIO SF
Iforia 7 VR-T DX, DR-T	NT
Ilesto 7 DR-T, HF-T, VR-T DX, VR-T, DR-T DF4	NT
Inventra 7 VR-T SX	AH
Iperia 7 VR-T, DR-T, VR-T DX, VR-T DF4, HF-T DF4	NH
Lumax 340 DR-T, HF-T, VR-T	HR
Lumax 540 DR-T, HF-T, VR-T	SH
Lumax 740 DR-T, HF-T, VR-T, VR-T DX	RH
Philos DR, D, SLR, SR, S	LE
Philos DR-T	VV
Philos II DR, D, S, SLR, SR	ET
Philos II DR-T	KP
Stratos LV, LV-T	SV
Talos DR, D, SLR, SR, S	PV
Protos DR/CLS, VR/CLS	EZ

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13 Contacting BIOTRONIK

Regarding This Report

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E-mail	PPR@biotronik.com
Address	BIOTRONIK, Inc. Attn: Compliance Department 6024 Jean Road Lake Oswego, OR 97035

Regarding Products

BIOTRONIK invites customers to call the following locations with suggestions, comments or specific questions related to BIOTRONIK products:

Worldwide CRM Product Support

Phone	+ 49 (0) 30 68905-1133
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Product Performance Report

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