

**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549**

FORM 8-K

**CURRENT REPORT
Pursuant to Section 13 or 15(d)
of the Securities Exchange Act of 1934
Date of Report (Date of earliest event reported): February 26, 2026**

PROCEPT BIOROBOTICS CORPORATION
(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

001-40797
(Commission
File Number)

26-0199180
(IRS Employer
Identification Number)

**150 Baytech Drive
San Jose, California 95134**
(Address of principal executive offices, including Zip Code)

Registrant's telephone number, including area code: (650) 232-7200

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

| Title of each class | Trading Symbol(s) | Name of each exchange on which registered |
|--|----------------------|--|
| Common Stock, \$0.00001 par value per share | PRCT | The Nasdaq Stock Market LLC |

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 7.01 Regulation FD Disclosure

PROCEPT BioRobotics Corporation (the “Company”) will host an Investor Day on February 26, 2026 at the NASDAQ Headquarters in New York City beginning at 8:00 a.m. Eastern Time. A copy of the Investor Day presentation is furnished as Exhibit 99.1 to this report and is incorporated herein by reference and constitutes a part of this report.

A live and an archived recording of the Investor Day will be available on the “Investors” section of the Company’s website at <https://ir.procept-biorobotics.com>.

The information included under Item 7.01 in this Current Report on Form 8-K, including Exhibit 99.1, shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that Section, nor shall it be deemed to be incorporated by reference into any filing of the Company under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

| <u>Exhibit No.</u> | <u>Description</u> |
|--------------------|---|
| 99.1 | Investor Day Presentation dated February 26, 2026 |
| 104 | Cover Page Interactive Data File, formatted in Inline XBRL. |

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

PROCEPT BIROBOTICS CORPORATION

Date: February 26, 2026

By: /s/ Alaleh Nouri
Alaleh Nouri
EVP, Chief Legal Officer and Secretary

Welcome to 2026 Analyst Day

Matt Bacso
VP, Investor Relations



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Safe Harbor Statement

This presentation and accompanying oral presentation contain "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, including the expected financial of PROCEPT BioRobotics Corporation (the "Company"). Words such as "anticipates," "believes," "expects," "intends," "projects," "anticipates," and "future" or similar expressions are intended to identify forward-looking statements. Any forward-looking statements made by us in this presentation speaks only as of the date on which it was made and are based on management's current expectations of events, assumptions, estimates, and beliefs, and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those set forth in or implied by: forward-looking statements. Factors that could cause actual results to differ materially from those described in the forward-looking statements include, among others: (i) the rate and degree of market acceptance of the AQUABEAM and HYDROS Robotic Systems and Aquablation therapy and descriptions of the Company's revenues, gross margin, profitability, operating expenses, or procedure growth installed base growth, (ii) the establishment and maintenance of consistent and favorable payment policies for Aquablation therapy, (iii) the rate of growth of the commercial sales and marketing organ and the ability to manage this anticipated growth, (iv) the impact on volumes of elective procedures performed by health care providers and hospital medical device budgets, (v) the effects of increased competition as well as innovations by new and existing competitors in the market for competitive treatments, (vi) the ability to obtain the required regulatory approvals and clearances to market and sell products in certain other countries, (vii) the development and protection of future innovation, (viii) dependence on a limited number of third-party suppliers for components of our products (ix) the maintenance of intellectual property rights and the ability to operate the business without infringing the intellectual property rights and proprietary technology of third parties, (x) the successful completion of clinical trials and (xi) the adoption of our technology for expanding or additional indications.

This presentation and the accompanying oral presentation also contain estimates and other statistical data made by independent parties and by us relating to market size and growth and other data in our industry. This data involves a number of assumptions and limitations, and you are cautioned not to give undue weight to such estimates. In addition, projections, assumptions, and estimates of our performance and the future performance of the markets in which we compete are necessarily subject to a high degree of uncertainty and risk.

Factors that could cause actual results to differ materially from those contemplated in this presentation can be found in the Risk Factors section of the Company's public filings with the Securities and Exchange Commission ("SEC"), including the Annual Report on Form 10-K filed with the SEC on February 26, 2026 and subsequent quarterly reports on Form 10-Q, available at www.sec.gov.

Because forward-looking statements are inherently subject to risks and uncertainties, you should not rely on these forward-looking statements as predictions of future events. All statements other than statements of historical fact are forward-looking statements. Except to the extent required by law, the Company undertakes no obligation to update or review any estimate, projection, or forward-looking statement. Actual results may differ from those set forth in this presentation due to the risks and uncertainties inherent in the Company's business. In light of the foregoing, investors are urged not to rely on any forward-looking statement or third-party data in reaching any conclusion or making any investment decision about any securities of the Company.

This presentation regarding the Company shall not constitute an offer to sell or the solicitation of an offer to buy any securities, nor shall there be any sale of these securities in any state or jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction. Sales and offers to sell PROCEPT BioRobotics securities will be made in accordance with the Securities Act of 1933, as amended, and applicable SEC regulations, including prospectus requirements.

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PROCEPT

Use of Non-GAAP Financial Information

In addition to financial information presented in accordance with U.S. generally accepted accounting principles ("GAAP"), this presentation and the accompanying oral statements include certain non-GAAP financial measures, which include non-GAAP Adjusted EBITDA. The Company defines Adjusted EBITDA as net income (loss) earnings before interest expense, taxes, depreciation and amortization and stock-based compensation expense. The Company believes that presenting Adjusted EBITDA provides useful supplemental information to investors about the Company in understanding and evaluating its operating results, enhancing the overall understanding of its past performance and future prospects, and allowing for greater transparency with respect to key financial metrics used by its management in financial and operational decision making. However, there are a number of limitations related to the use of non-GAAP measures and their nearest GAAP equivalents. For example, such measures may exclude significant expenses required by GAAP to be recognized in our financial statements and companies may calculate non-GAAP measures differently or may use other measures to calculate their financial performance, and therefore any non-GAAP measures the Company uses

not be directly comparable to similarly titled measures of other companies. Non-GAAP financial measures are not a substitute for or superior to measures of financial performance presented in accordance with GAAP and should not be considered as an alternative to any other performance measures derived in accordance with GAAP. Any non-GAAP measure is presented for supplemental informational purposes only and should not be considered a substitute for or superior to financial information presented in accordance with GAAP. A reconciliation of these measures to the most directly comparable GAAP measures is included at the end of this presentation.

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PROCEPT

PROCEPT
BIOROBOTICS®

The Next Chapter for PROCEPT

Larry L Wood
Chief Executive Officer



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A Large, Underpenetrated BPH Market with a Disciplined Path to Growth



- BPH is significantly undertreated
- Many patients fear the procedure more than their condition
- Delaying treatment significantly impacts QOL

1. Vuichoud C, Loughlin KR. Benign prostatic hyperplasia: epidemiology evaluation. *Can J Urol.* 2015 Oct;22 Suppl 1:1-6. PMID: 26497338.
2. Based on management estimates and data provided by AcuityMD, De Release, US market estimates, Q4 2024 – Q3 2025

PROCEPT Biorobotics Stands on a Strong Foundation

Game-changing Technology

Deep Clinical Evidence Base

The Next Chapter

- Accelerate Procedure Growth
- Drive Path to Profitability
- Advance Evidence and Innovation

We are now moving from establishing Aquablation® as a therapy to leading BPH treatment

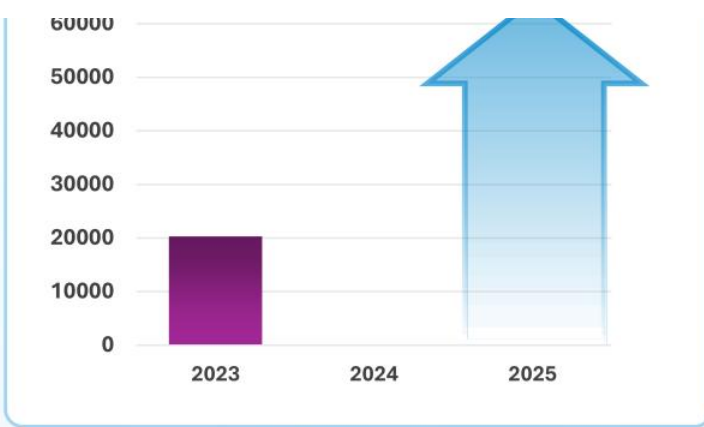
Data on File at PROCEPT BioRobotics

PROCEPT

Growing Procedures and Install Base Paired With Strong Reimbursements

Worldwide Aquablation Procedures

>125K patients treated worldwide



>900 global install base



Category I Reimbursement as of January 1, 2026

Aquablation is ready to move to the standard of care

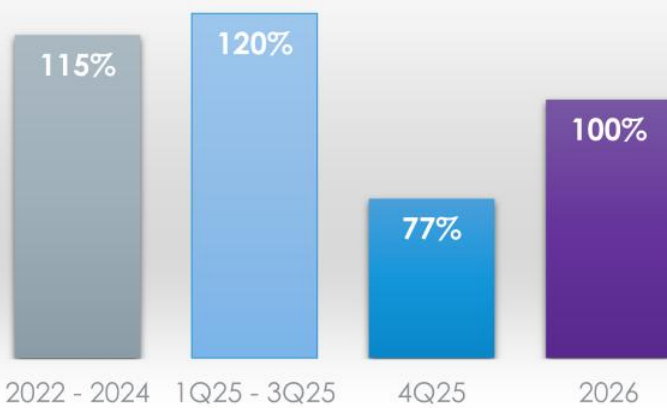
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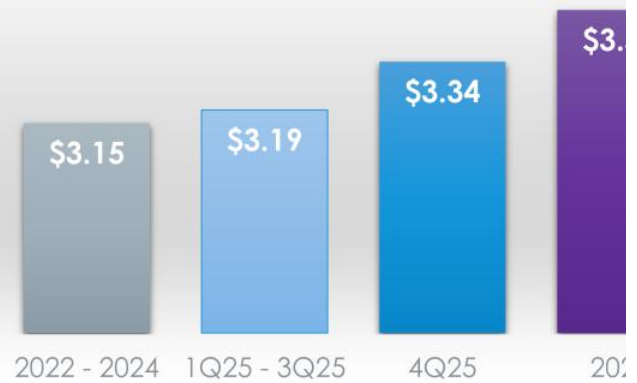


Benefits of Improved Organizational Discipline

% HP to Procedure



HP Pricing (\$000)

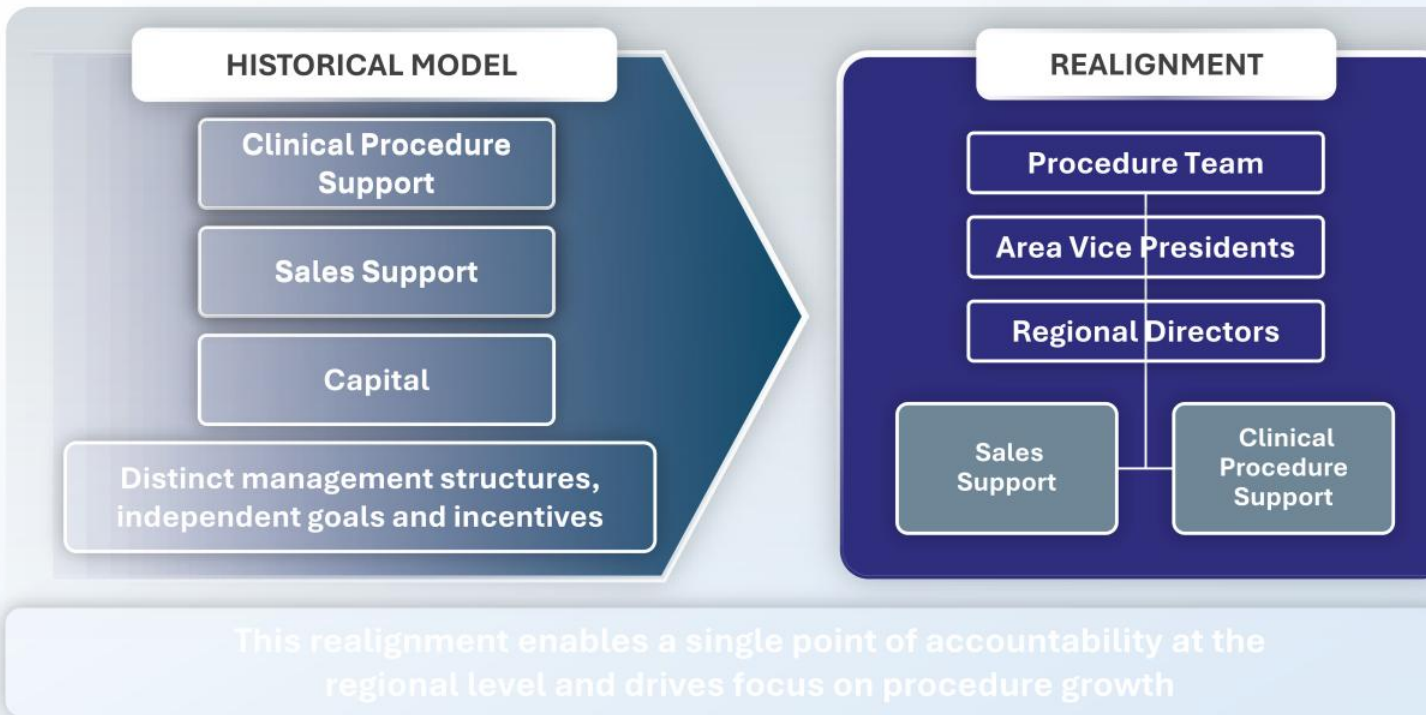


Two positive structural effects: Higher-than-expected ASP and improved quality and predictability of revenue

Data on File at PROCEPT BioRobotics

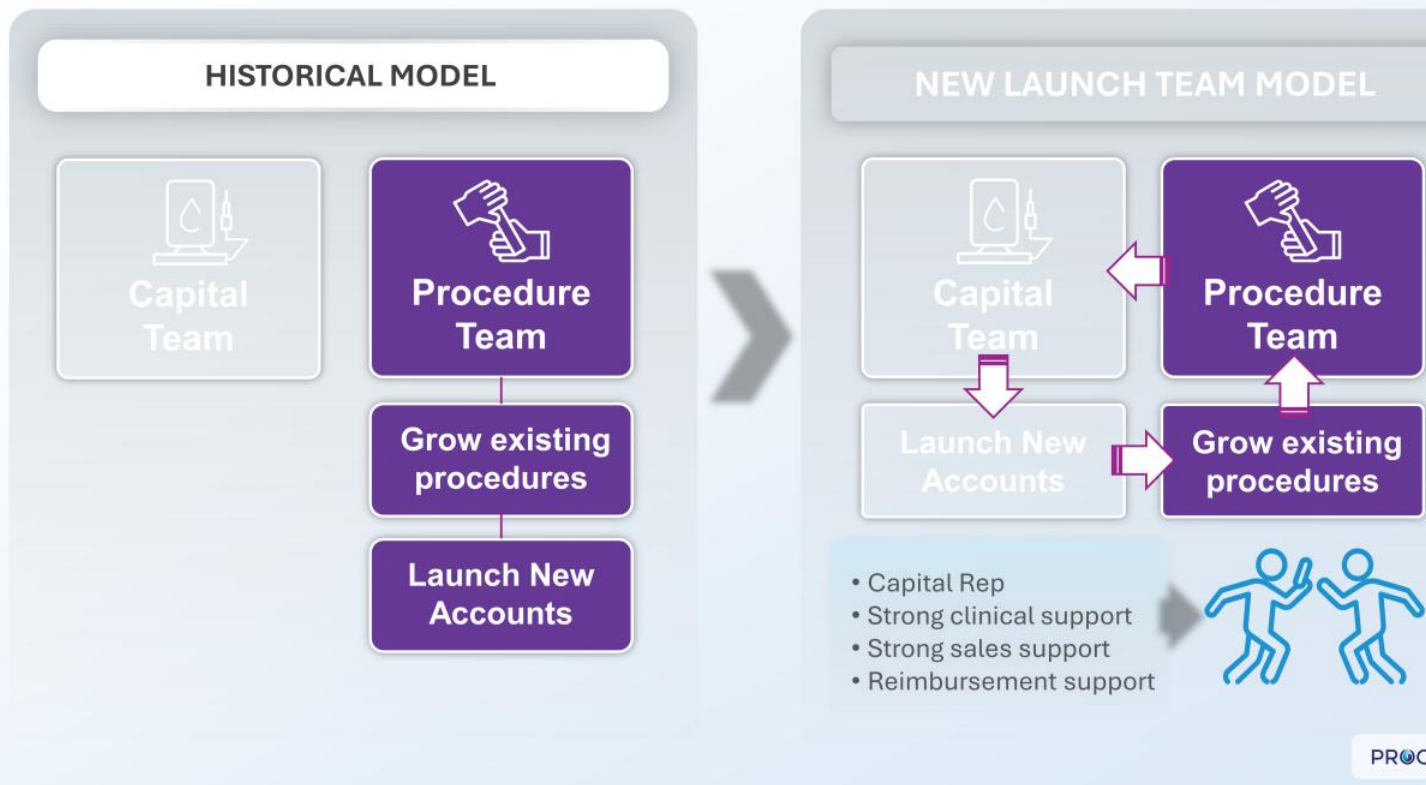
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Realignment of the Commercial Organization



PROC

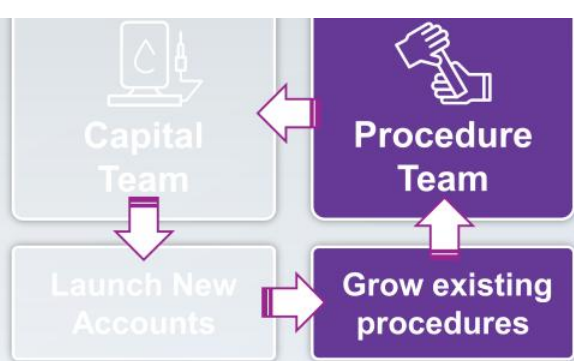
We Have Formed Dedicated Launch Teams



PROC

We Have Formed Dedicated Launch Teams





- Capital Rep
- Strong clinical support
- Strong sales support
- Reimbursement support



~50% reduction in time from PO to first ten cases

The net result is a dedicated launch team of experts who standardize the pathway for training, reimbursement, and patient education in new accounts

Data on File at PROCEPT BioRobotics

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ACCELERATED PROCEDURE

Greenfield Placements Remain Core to Capital Revenue



Alternative capital acquisition strategies enable us to expand access to additional customer groups

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Making the Clinical Case for Aquablation



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Supported by a Robust Body of Evidence

Five-year outcomes for Aquablation therapy compared to TURP: results from a double-blind, randomized trial in men with LUTS due to BPH

Peter J. Gillings, MD,¹ Neil Barber, MD,² Mohamed Bidair, MD,³ Paul Anderson, MD,⁴ Mark Sutton, MD,⁵ Tev Aho, MD,⁶ Eugene Kramolowsky, MD,⁷ Andrew Thomas, MD,⁸ Ronald P. Kaufman, Jr., MD,⁹ Gopal Badlani, MD,¹⁰ Mark Plante, MD,¹¹ Mihir Desai, MD,¹² Leo Doumanian, MD,¹³ Alexis E. Te, MD,¹⁴ Claus G. Roehrborn, MD¹⁵

Submitted 12/15/24; accepted 1/16/25; published online 2/10/25. This is an Open Access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited. DOI: 10.1097/JUQ.0000000000000000

OBJECTIVE: To report real-world experience of 4-year safety and efficacy outcomes of Aquablation procedure for the treatment of men with symptomatic benign prostatic hyperplasia (BPH).
METHODS: This is a prospective single-center, observational study evaluating the outcomes of robotic-assisted Aquablation therapy for moderate-to-severe BPH between December 2019 and December 2023. Patient-level data included age, prostate volume, IPSS score, peak urinary flow rate (Q_{max}), post-void residual (PVR) were evaluated at 1M, 6M, and 1Y. Primary outcomes included change in IPSS score, change in Q_{max}, change in PVR, preservation of antegrade ejaculation, and complications.
RESULTS: In this cohort of 330 men, mean prostate volume was 112.1 mL (range 38-335 mL) at baseline. International Prostate Symptom Score (IPSS) improved from a baseline of 21.8 (SD 8.4) to 6.9 (SD 2.9) at 4 years. Mean peak urinary flow rate (Q_{max}) also demonstrated improvement and increased from 6.4 mL/sec (SD 4.2) to 17.4 mL/sec (SD 5.5) at 4 years. At 1 year, mean prostate volume reduction was 45.5 mL (-41.9%). Postoperative antegrade ejaculation was preserved in 2,692/30 men (89.6%) of men. Complications included urinary tract infection within first month after procedure in 17 (4.1%), and bleeding requiring blood transfusion in 11 (3.3%). Thrombotic events (3.9%) required a second procedure including 2 for post-operative bleeding, 1 for a bladder neck disimpaction and 10 (3.0%) for transurethral resection of residual anterior tissue. We demonstrate Aquablation to not only be safe but also providing durable outcomes at 4 years for men with BPH. UROLOGY 194: 218-220, 2024. © 2024 Elsevier Inc. All rights are reserved, including those for text and data mining, AI training, and similar technologies.

OPEN

Aquablation Therapy in Large Prostates (80-150 mL) for Lower Urinary Tract Symptoms Due to Benign Prostatic Hyperplasia: Final WATER II 5-Year Clinical Trial Results

Nasem Bhojani,^{1*} Mo Bidair,² Eugene Kramolowsky,³ Mihir Desai,⁴ Leo Doumanian,⁵ Kevin C. Zorn,⁶ Dean Ertman,⁷ Ronald P. Kaufman Jr.,⁸ Gregg Eurs,⁹ Gopal Badlani,¹⁰ Mark Plante,¹¹ Edward Uchio,¹² Greg Gin,¹³ Ryan Paterson,¹⁴ Alan So,¹⁵ Claus Roehrborn,¹⁶ Jay Motola,¹⁷ Steven Kaplan,¹⁸ and Mitch Humphreys¹⁹

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OBJECTIVE: To report real-world experience of 5-year safety and efficacy outcomes of Aquablation procedure for the treatment of men with symptomatic benign prostatic hyperplasia (BPH).
METHODS: This is a prospective single-center, observational study evaluating the outcomes of robotic-assisted Aquablation therapy for moderate-to-severe BPH between December 2019 and December 2023. Patient-level data included age, prostate volume, IPSS score, peak urinary flow rate (Q_{max}), post-void residual (PVR) were evaluated at 1M, 6M, and 1Y. Primary outcomes included change in IPSS score, change in Q_{max}, change in PVR, preservation of antegrade ejaculation, and complications.
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THE JOURNAL OF UROLOGY

Available at www.sciencedirect.com
Journal homepage: www.europeanurology.com/efocus

Benign Prostatic Hyperplasia

WATER III: A Prospective, Partially Randomized Trial of Aquablation Therapy Versus Transurethral Laser Enucleation of the Prostate for Treatment of Lower Urinary Tract Symptoms

Manuel Kitter,^{1*} Johannes Stein,² Neil Barber,³ Jas Kabli,⁴ Rick Popert,⁵ Edward Bess,⁶ Robert Nüneth,⁷ Matthias Schmid,⁸ Simon Glöckner,⁹ Burkhard Ullrich,¹⁰ Arkadiusz Mierak,¹¹ Christian Gratzke¹²

Submitted 12/15/24; accepted 1/16/25; published online 2/10/25. This is an Open Access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited. DOI: 10.1097/JUQ.0000000000000000

OBJECTIVE: To report real-world experience of 4-year safety and efficacy outcomes of Aquablation procedure for the treatment of men with symptomatic benign prostatic hyperplasia (BPH).
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Prostatic Disease & Male Voiding

Aquablation at 4-years: Real World Data From the Largest Single-center Study With Associated Outcomes Follow-up

Olamide O. Omidele, Alexandra S. Singal, Reza Roshandel, Alexis E. Te, and Steven A. Kaplan

OBJECTIVE: To report real-world experience of 4-year safety and efficacy outcomes of Aquablation procedure for the treatment of men with symptomatic benign prostatic hyperplasia (BPH).
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Journal of Clinical Medicine

Article First Multi-Center All-Comers Study for the Aquablation Procedure

Thorsten Bach,^{1,*} Peter Gillings,² Albert El Hajj,³ Paul Anderson,⁴ and Neil Barber⁵

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MDPI

Original Article

Aquablation for benign prostatic hyperplasia: prostate size relevance and bleeding events across 6 years

Mario Klotz,^{1*} Sascha Frensch,² Sigi Brighi,³ Oliver Baku,⁴ Nouven Bregent,⁵ Saeed Ch Khan,⁶ Samir Ghomash, and Steven A. Kaplan⁷

Submitted 12/15/24; accepted 1/16/25; published online 2/10/25. This is an Open Access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited. DOI: 10.1097/JUQ.0000000000000000

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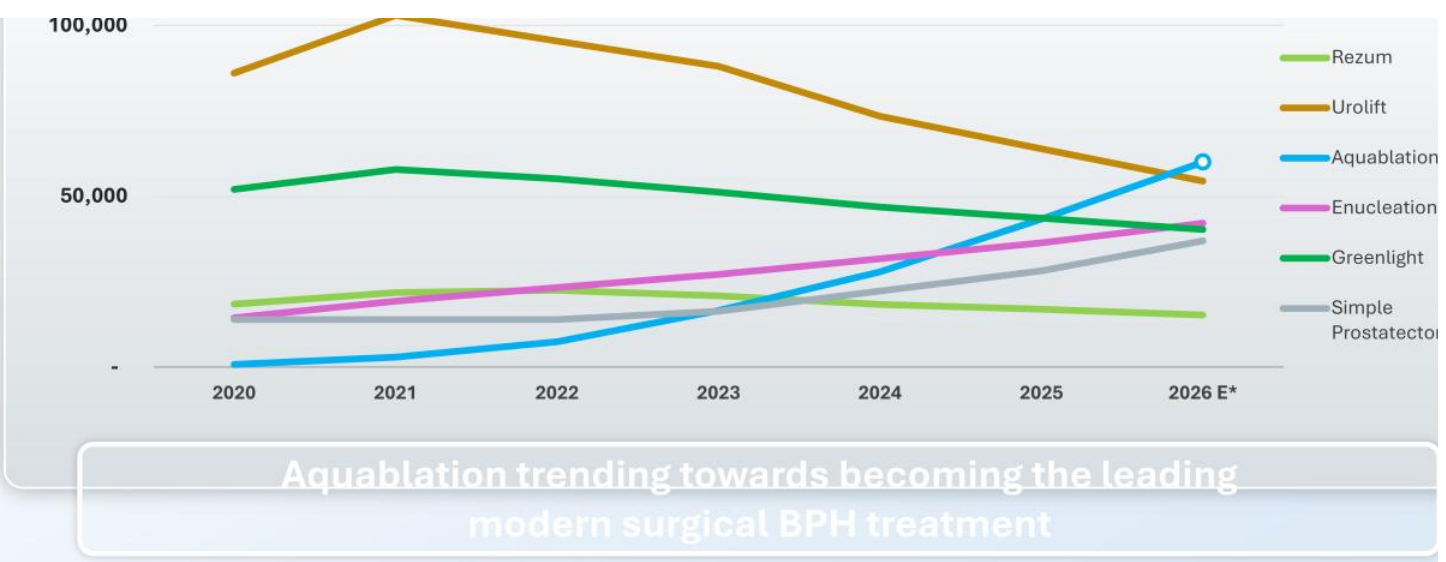
Data from thousands of patients published with ~250 PubMed citations

Aquablation is Growing Within the BPH Category

All BPH Volume Trends (Excluding TURP)

PROC

ACCELERATED PROCEDURE

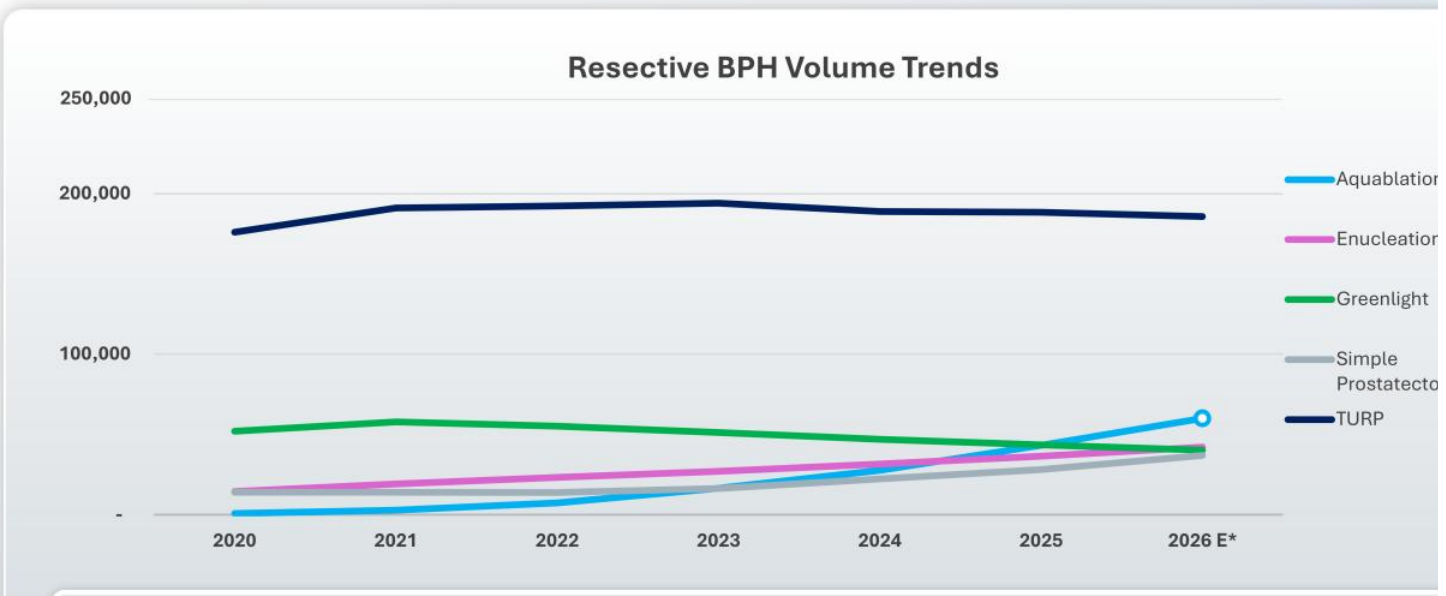


Aquablation trending towards becoming the leading modern surgical BPH treatment

Data on File at PROCEPT BioRobotics
 Based on management estimates and data provided by AcuityMD, Dec 19, 2026 DataRelease, US market estimates, Q4 2024 – Q3 2025. FY 2025 and 2026 volumes for competitive procedures are estimates based on historical data.



Aquablation is Closing the Gap with TURP



Aquablation is now the leading resective BPH treatment – second only to TURP

Data on File at PROCEPT BioRobotics
 Based on management estimates and data provided by AcuityMD, Dec 19, 2026 DataRelease, US market estimates, Q4 2024 – Q3 2025. FY 2025 and 2026 volumes for competitive procedures are estimates based on historical data.

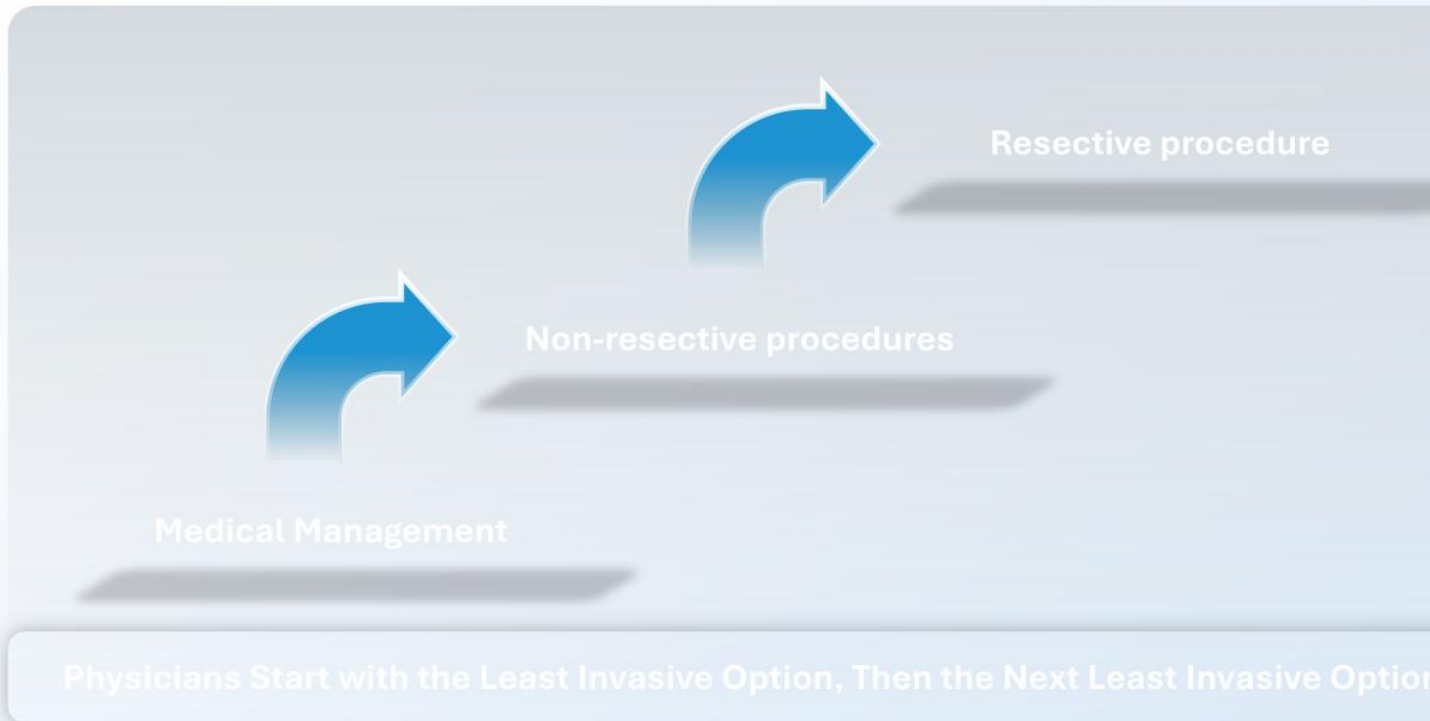


Aquablation Delivers a Complete Solution Across All Prostate Sizes

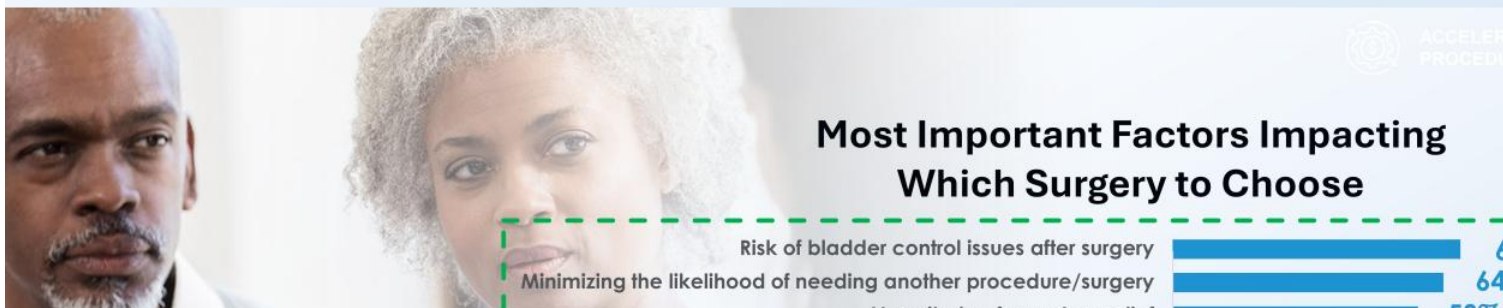


PVP = Photovaporization of Prostate; MIST: Minimally Invasive Surgical Technology; BPH size ranges: AUA Guidelines: Management of Lower Urinary Tract Symptoms Attributed to Benign Prostatic Hyperplasia (BPH): AUA Guideline Amendment 2023; Tanneru et al: An Indirect Comparison of Newer Minimally Invasive Treatments for Benign Prostatic Hyperplasia: A Network Meta-Analysis Model, Journal of Endourology, 2020

Physicians Often Take a Stepwise Approach to Treatment

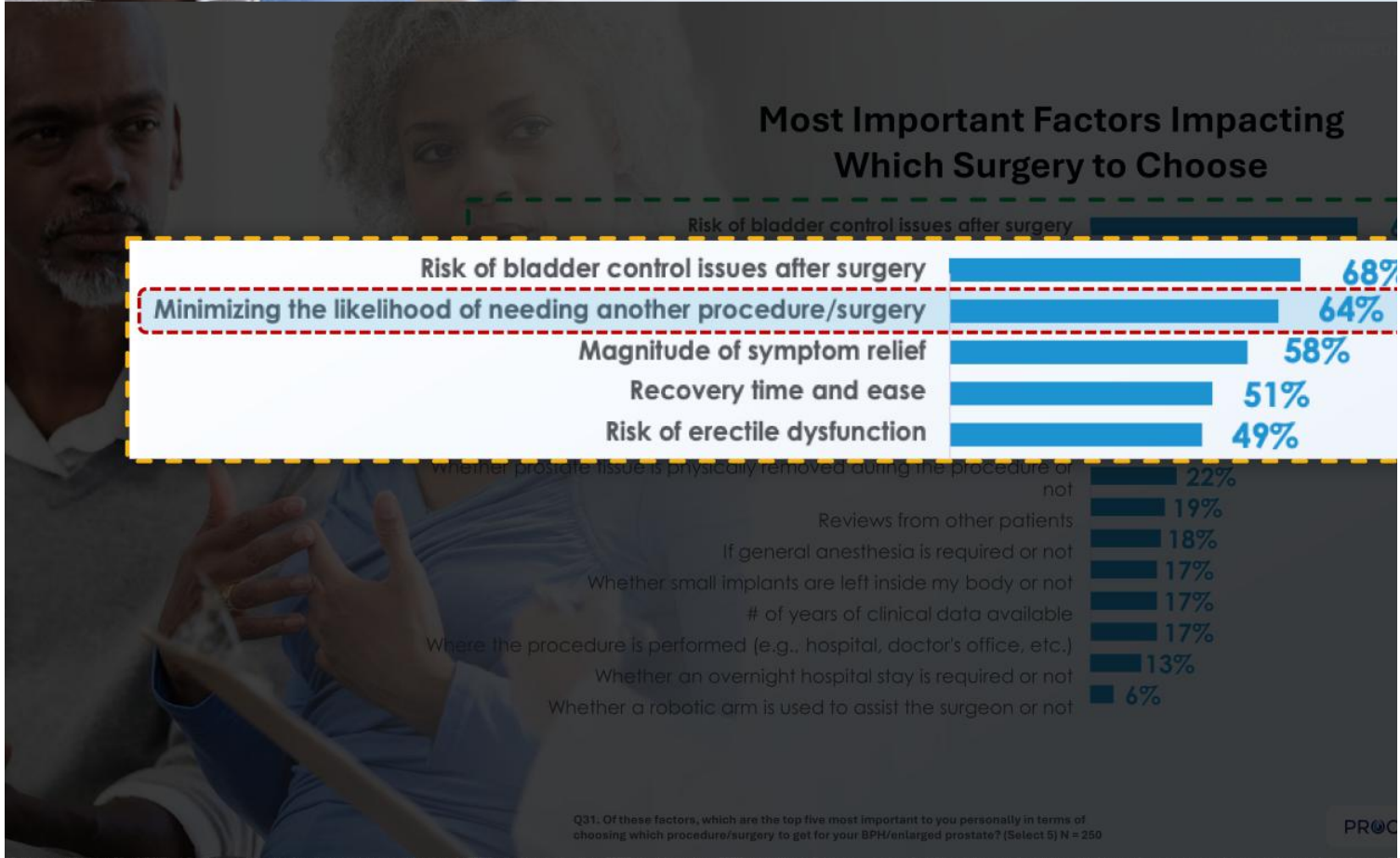
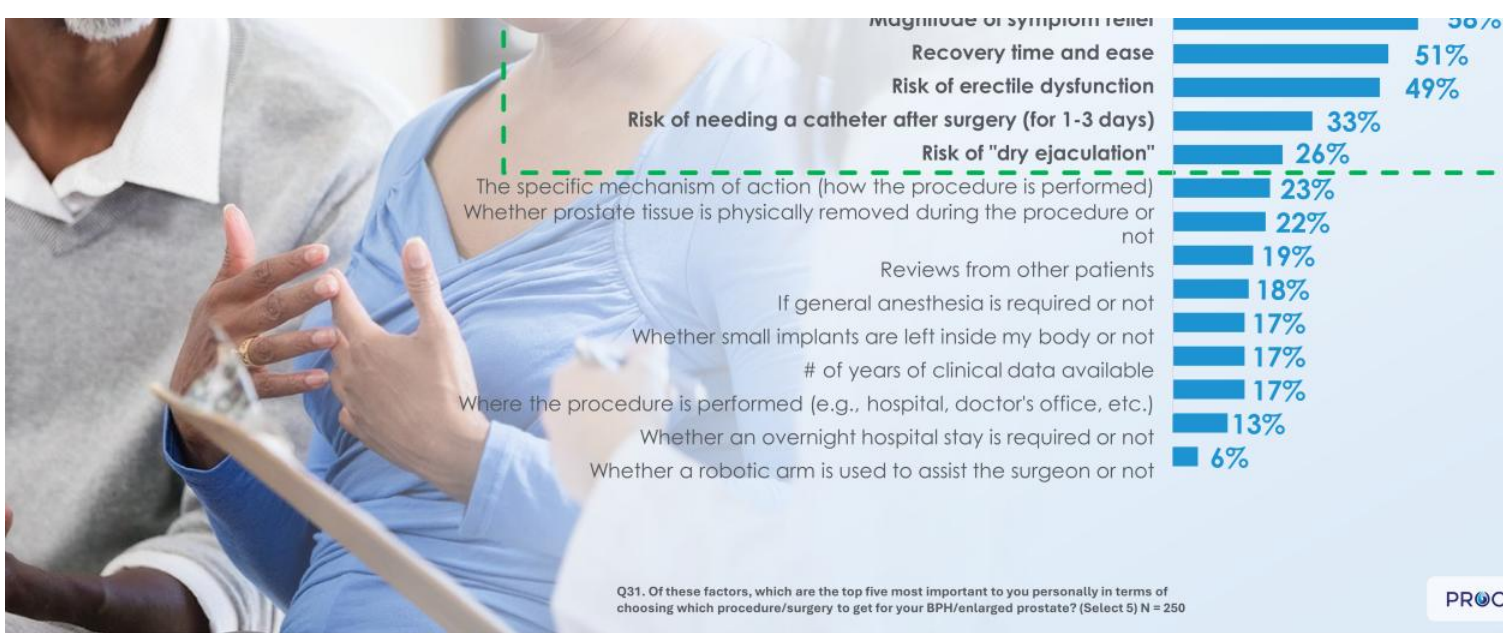


Market research data on file



Most Important Factors Impacting Which Surgery to Choose

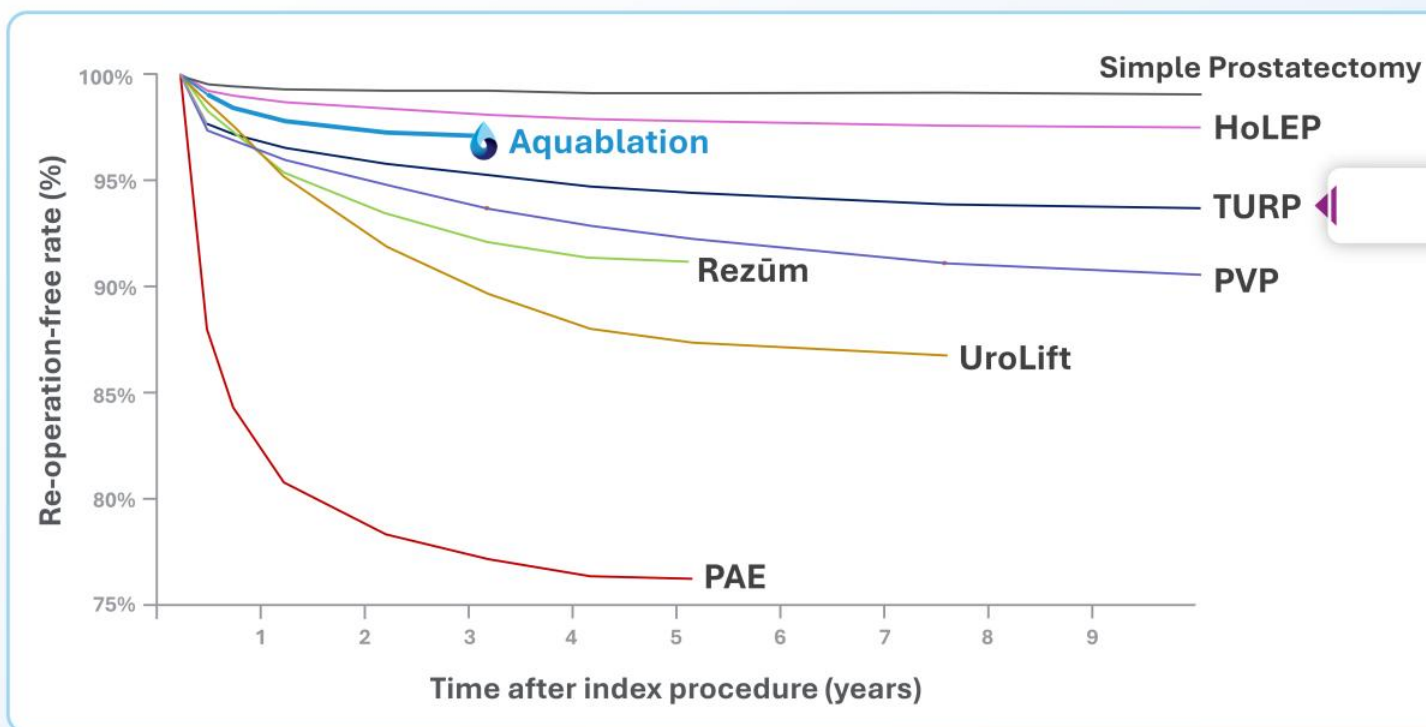
| | |
|--|-----|
| Risk of bladder control issues after surgery | 4% |
| Minimizing the likelihood of needing another procedure/surgery | 64% |
| Magnitude of symptom relief | 69% |



These procedures involve a 24F catheter in the urethra – no one wants that more than once

Patients deserve to have a procedure that offers a complete solution with the best opportunity to avoid a second intervention

Patients Want One Procedure Only



Rubin B, et al. National review of re-operation rates for modern benign prostatic hyperplasia procedures using a live claims database. Abstract IP03-06. Study Design: Retrospective observational study utilizing the TriNetX database (2004–2024). AUA 2025 from University of Vermont



Our Innovation Portfolio



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HYDROS® System: Next Generation Robotic Platform



HYDROS: The First AI-Enabled Robotic Platform for BPH

- **AI-Interpreted Live Ultrasound Guidance:** Personalized treatment planning combined with surgeon judgment
- **Robotic Resection:** Precise, reproducible tissue removal while protecting critical anatomy
- **Designed for Scalable Adoption:** Streamlined workflow that enables broader adoption across care settings

HYDROS is a feature-rich platform that enables a hybrid case support model



We expect HYDROS to represent the majority of the install base by the end of 2026



Global expansion of HYDROS underway – UK launch has begun



Continued pipeline investment

International represents an untapped opportunity and a more meaningful growth contributor over time

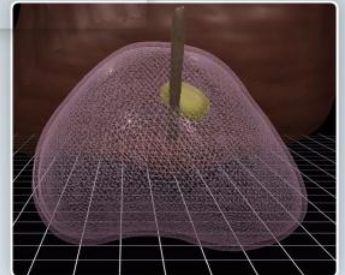
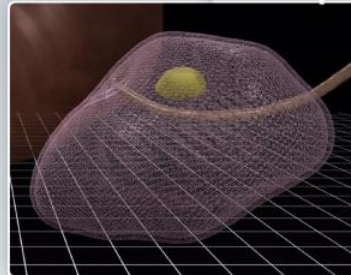
The Next Frontier: Prostate Cancer

Prostate Cancer

A Strategic Adjacency

- Technology
- Installed Base
- Commercial Infrastructure

Aquablation PCa



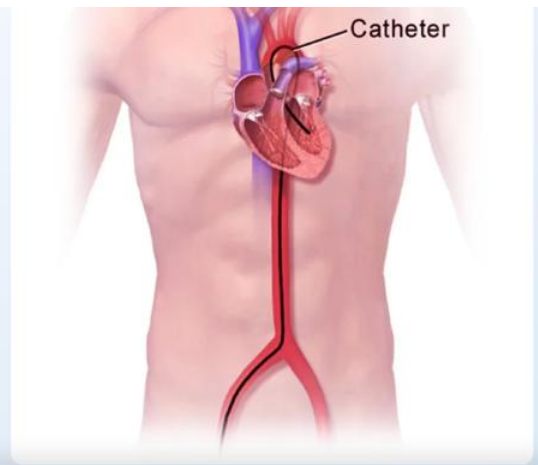
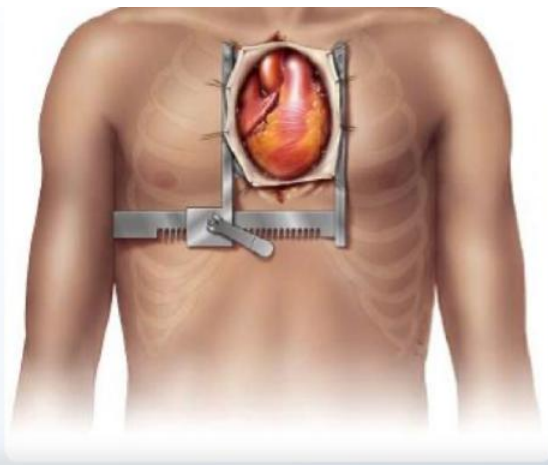
This is an Area That is Very Familiar To Me

Open Heart Surgery (AVR)



Transcatheter Valve Replacement (TAVR)





Watchful waiting was common until the suffering became unbearable

PROC

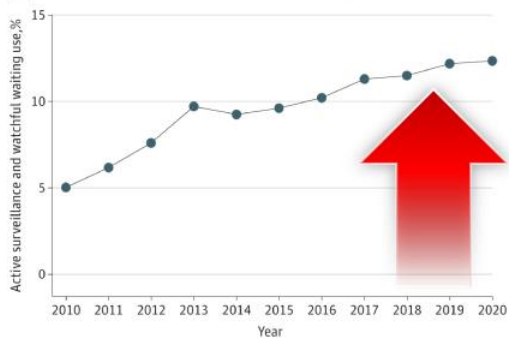


ADVANCE
AND IMPROVE

The fastest growing treatment for prostate cancer :

ACTIVE SURVEILLANCE!

A Overall use of active surveillance and watchful waiting



Radical Prostatectomy



Majority of men experience severe erectile dysfunction



25-33% rate of Urinary Incontinence at 1 year

>60% of patients ultimately get treated with a radical prostatectomy or radiation

Trends in Active Surveillance for Men With Intermediate-Risk Prostate Cancer. Marshall et al. JAMA 2024; Hamdy, F. C., Donovan, J. L., Lane, J. A., Metcalfe, C., Davis, M., Turner, E. L., Martin, R. M., et al. (2023). Fifteen-Year Outcomes after Monitoring, Surgery, or Radiotherapy for Prostate Cancer. New England Journal of Medicine, 388(17), 1547-1558. DOI: 10.1056/NEJMoa2214122. Donovan et al NEJM 2016; 2. Bridge et al Eur Urol Open Sci 2024

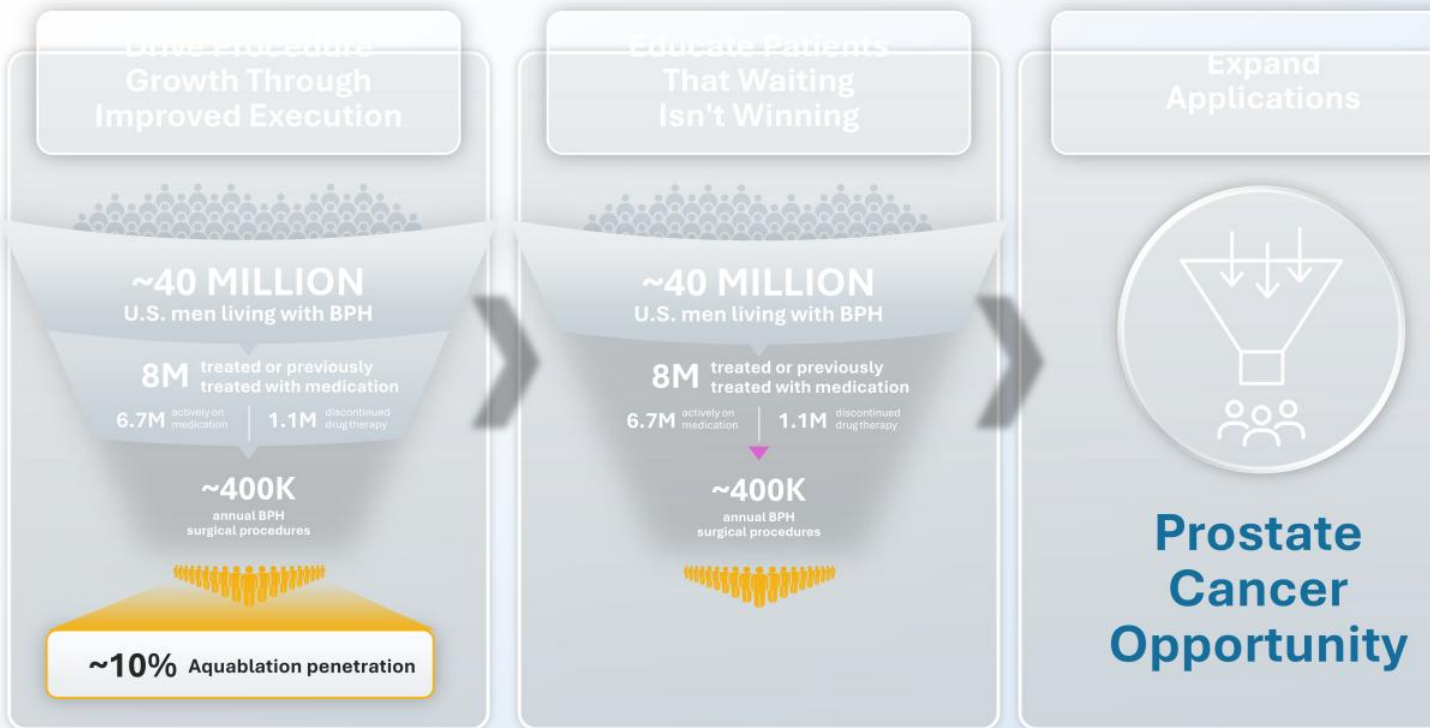
PROC



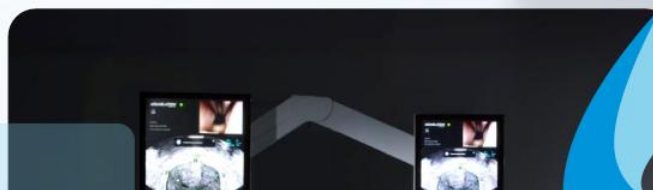
We believe Aquablation will play a meaningful role in the treatment of prostate cancer

WATER IV PCa
STUDY

Multiple Initiatives to Drive Differentiated Growth Across the Horizon



1. Vuichoud C, Loughlin KR. Benign prostatic hyperplasia: epidemiology, economics and evaluation. Can J Urol. 2015 Oct;22 Suppl 1:1-6. PMID: 26497338.
 2. Based on management estimates and data provided by AcuityMD, Dec 19, 2026 Data Release, US market estimates, Q4 2024 – Q3 2025



Unlocking Growth: A Closer Look

Pooja Sharma Rao
Chief Marketing and Strategy Officer



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Driving an Acceleration in Procedure Growth

The Next Chapter:

 **Accelerate Procedure Growth**



Drive Path to Profitability



Advance Evidence and Innovation



The Pathway to Growth

- Sales force realignment
- Dedicated launch teams
- **Share capture through strong clinical differentiation**
- **Targeted patient education and engagement programs**

The engine around BPH lays a foundation on which we can launch a prostate cancer application

We Believe This is an Important Inflection Point

>125K patients treated worldwide

>900 global install base



Category I Reimbursement

Five-year outcomes for Aquablation therapy compared to TURP: results from a double-blind, randomized trial in men with LLUTS due to BPH...
 Aquablation Therapy in Large Prostates (80-150 mL) for Lower Urinary Tract Symptoms Due to Benign Prostatic Hyperplasia: Final WATER 5-Year Clinical Trial Results...
 Aquablation at 4-years: Real World Data From the Largest Single-center Study With Associated Outcomes Follow-up...
 First Multi-Center All-Comers Study for the Aquablation Procedure...
 Aquablation for benign prostatic hyperplasia: real prostate size relevance and bleeding events across 6 years...
 Maturing body of evidence

Data on File at PROCEPT BioRobotics

PROCEPT

Our Approach Starts Intentionally with Building Belief in Aquablation with Clinicians and Practices



Patients need to encounter physicians and care teams who believe in the differentiation of Aquablation therapy

This sequencing reflects lessons learned

- Establish the therapy
- Become the Procedure of Choice
- Targeted patient education as an accelerant

PROCEPT

Our Immediate Focus: Improve Penetration of Aquablation into the Surgical Segment



Grounded in what

~40 MILLION

U.S. men living with BPH

8M treated or previously treated with medication

6.7M actively on medication

1.1M discontinued drug therapy

~400K annual BPH surgical procedures



~10% Aquablation penetration

patients want, Aquablation delivers a COMPLETE BPH solution, supported by a mature and growing body of clinical evidence

1. Vuichoud C, Loughlin KR. Benign prostatic hyperplasia: epidemiology and clinical evaluation. Can J Urol. 2015 Oct;22 Suppl 1:1-6. PMID: 26497338. 2. Based on management estimates and data provided by AcuityMD. Release, US market estimates, Q4 2024 - Q3 2025



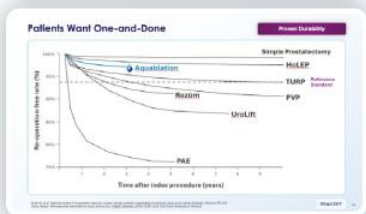
1

PATIENTS WANT: Complete Symptom Relief



2

PATIENTS WANT: Preserved Sexual and Urinary Function



3

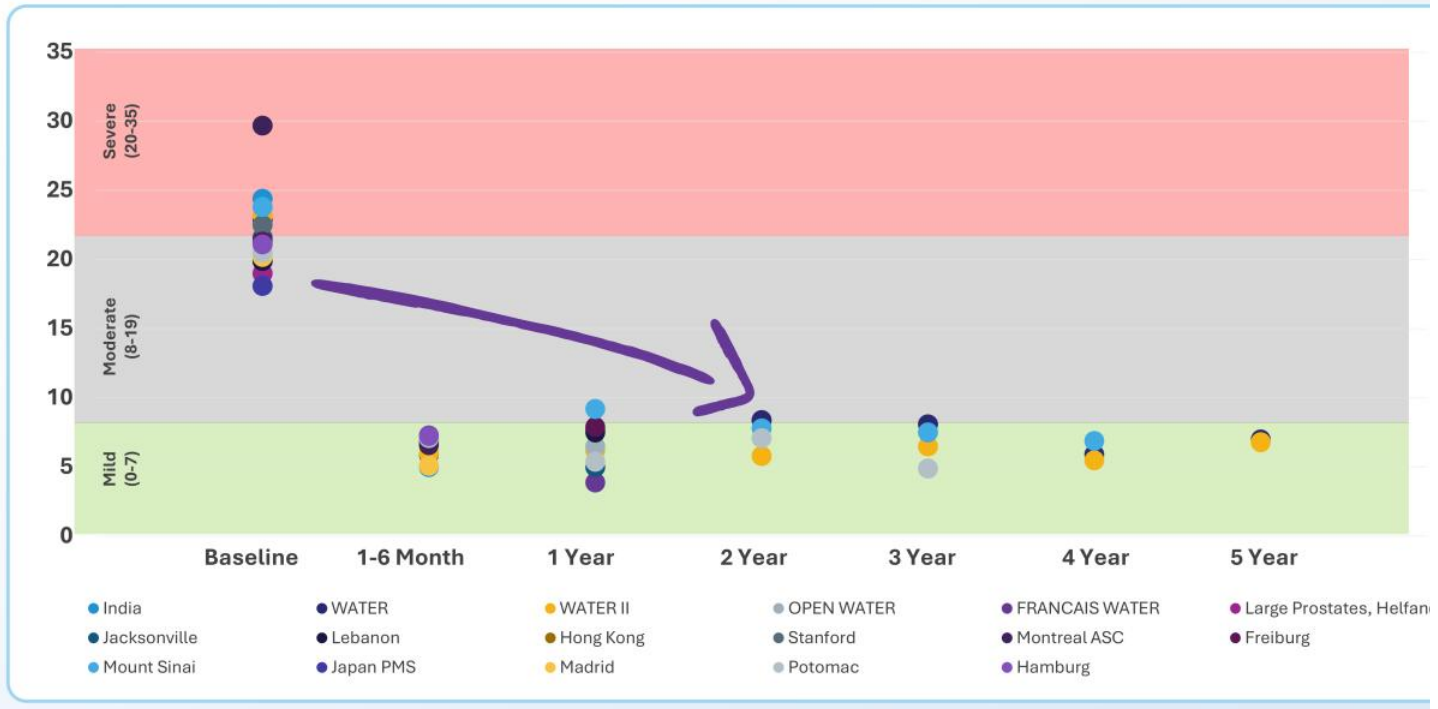
PATIENTS WANT: One Procedure Only

Aquablation is a COMPLETE Solution for BPH patients

Market research data on file

PROC

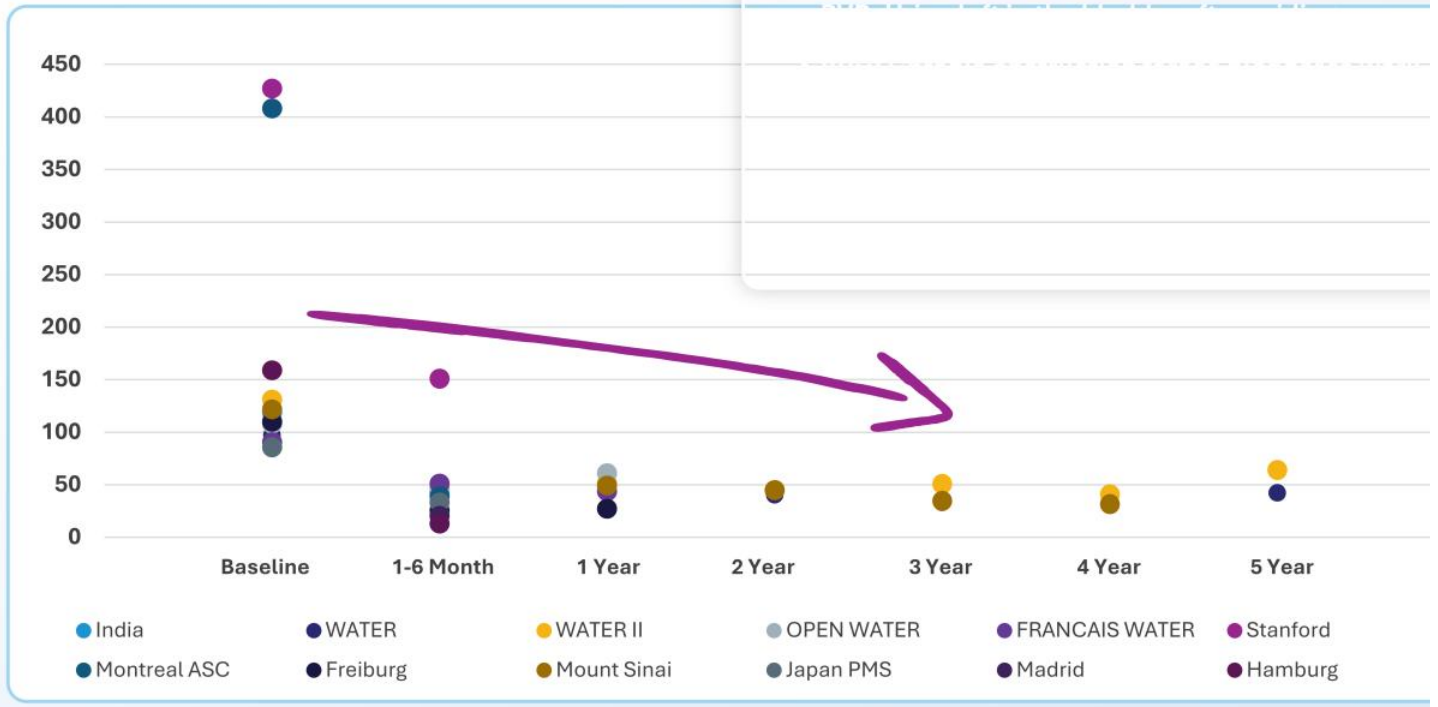
Symptom Reduction: International Prostate Symptom Score Results



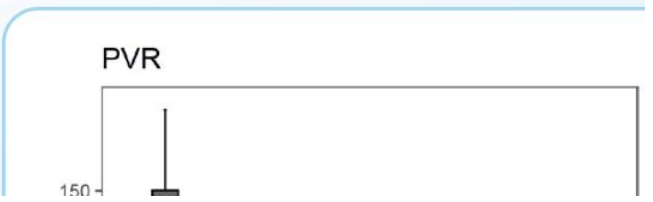
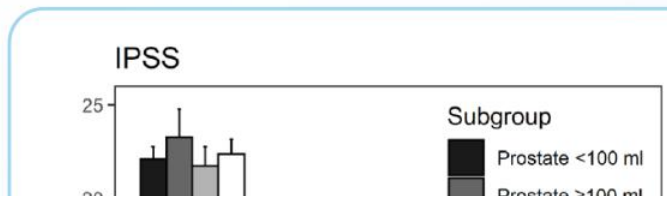
India, 20-118, n=47, Desai et al; WATER, 30-80ml, n=116, Gillig et al; Urodynamics, 30-80ml, n=43, Pimentel et al; WATER II, 80-150ml, n=101, Bhojani et al; OPEN WATER, 20-148ml, n=178, Bach et al; FRANCAIS WATER, 30-80ml, n=30, Misra et al; Focal Bladder Neck Caustery, 20-263ml, n=2,089, Elterman et al; HoLEP vs Aquablation Hematuria Risk, 56±25ml, n=167, Gloger et al; Very Large Prostates, 151-362ml, n=38, Helfand et al; Jacksonville, 27-223ml, n=55, Kasraeian et al; Lebanon, 13-148ml, n=59, Labban et al; Hong Kong AUR, 61±16ml, n=20, Yee et al; Stanford AUR and CUR, 29-250ml, n=113, Burton et al; UK Day Case, 22-120ml, n=40, Ng et al; Montreal ASC, 41-270ml, n=60, Zorn et al; Freiburg Aquablation vs HoLEP, 55±19ml, n=16, Michaelis et al; Mount Sinai, 38-330ml, n=330, Omidele et al; Hamburg, 20-154ml, n=118, Bach et al; Japan PMS, 33-242ml, n=103, Hinata et al; Madrid Aquablation vs HoLEP, 72±35ml, n=75, Quintas et al; Potomac Urology, 22-263ml, n=812, Marhamati et al; Aquablation 5 Year Trend, 20-657, n=36,555, Elterman et al

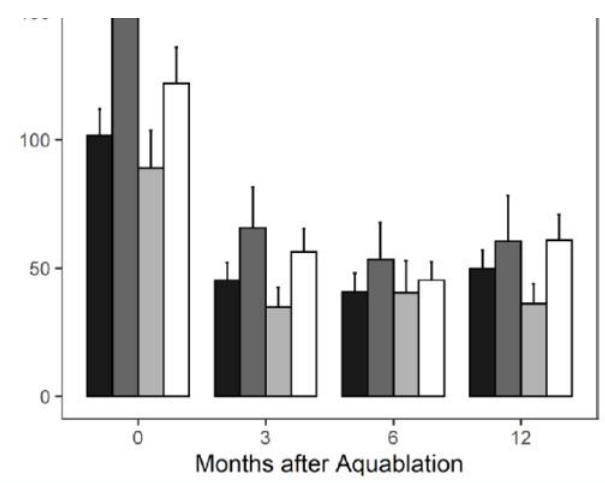
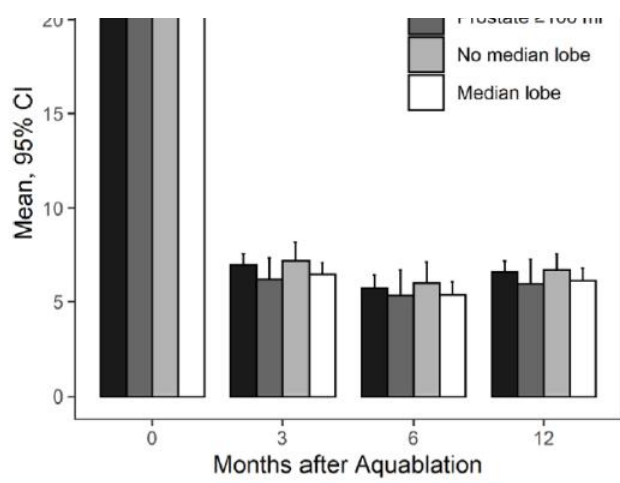


Post Void Residual Results



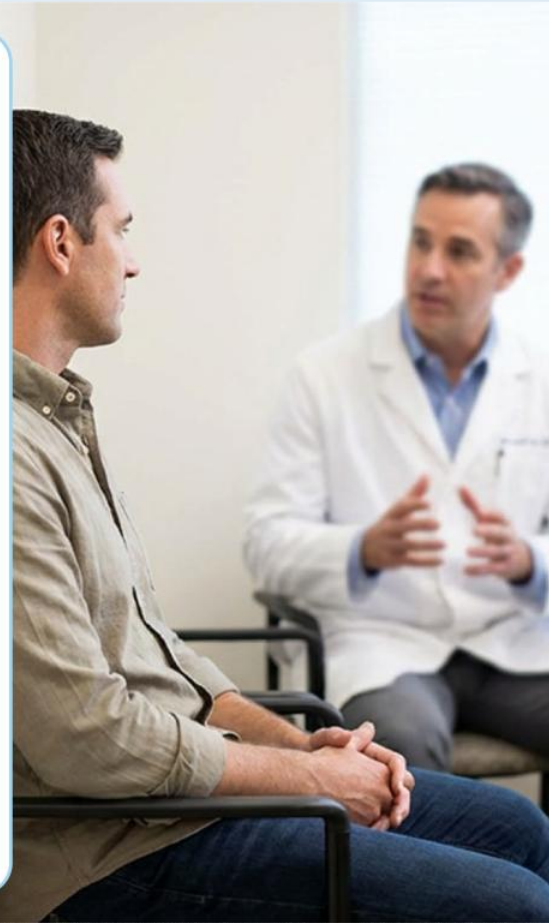
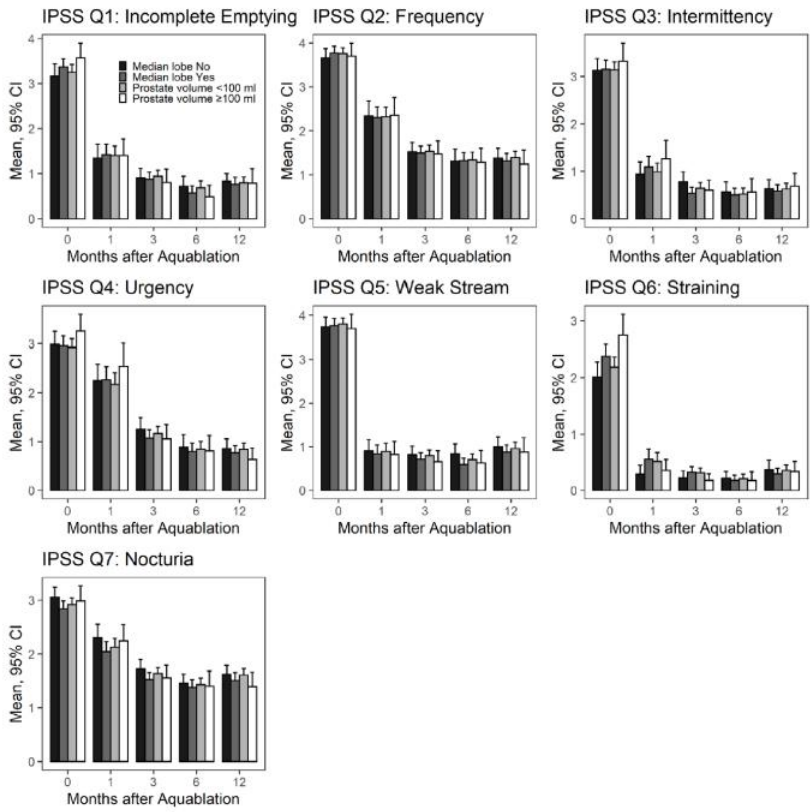
India, 20-118, n=47, Desai et al; WATER, 30-80ml, n=116, Gillig et al; Urodynamics, 30-80ml, n=43, Pimentel et al; WATER II, 80-150ml, n=101, Bhojani et al; OPEN WATER, 20-148ml, n=178, Bach et al; FRANCAIS WATER, 30-80ml, n=30, Misra et al; Focal Bladder Neck Caustery, 20-263ml, n=2,089, Elterman et al; HoLEP vs Aquablation Hematuria Risk, 56±25ml, n=167, Gloger et al; Very Large Prostates, 151-362ml, n=38, Helfand et al; Jacksonville, 27-223ml, n=55, Kasraeian et al; Lebanon, 13-148ml, n=59, Labban et al; Hong Kong AUR, 61±16ml, n=20, Yee et al; Stanford AUR and CUR, 29-250ml, n=113, Burton et al; UK Day Case, 22-120ml, n=40, Ng et al; Montreal ASC, 41-270ml, n=60, Zorn et al; Freiburg Aquablation vs HoLEP, 55±19ml, n=16, Michaelis et al; Mount Sinai, 38-330ml, n=330, Omidele et al; Hamburg, 20-154ml, n=118, Bach et al; Japan PMS, 33-242ml, n=103, Hinata et al; Madrid Aquablation vs HoLEP, 72±35ml, n=75, Quintas et al; Potomac Urology, 22-263ml, n=812, Marhamati et al; Aquablation 5 Year Trend, 20-657, n=36,555, Elterman et al





Elterman D, Gilling P, Boehrborn C et al, Meta-analysis with individual data of functional outcomes following Aquablation for lower urinary tract symptoms due to BPH in various prostate anatomies. *BMJ Surg Interv Health Technologies* 2021.

PROC



<1%
incontinence

<1%
ED

Aggregated data from 18 publications. References and analysis methodology in appendix.

<1%
incontinence

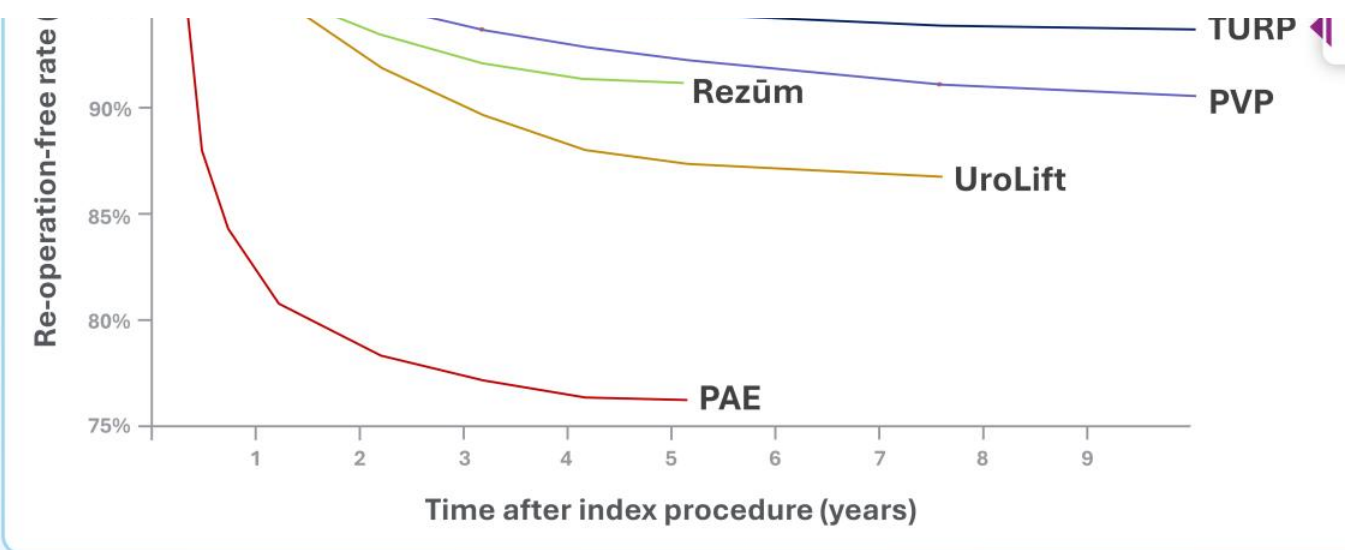
<1%
ED

Aggregated data from 18 publications. References and analysis methodology in appendix.; Bitar M, et al. Aquablation for benign prostatic hyperplasia: real-world prostate size relevance and bleeding events across 6 years. BJU Int. 2026;137(3):480-484. doi:10.1111/bju.70118



Patients Want One Procedure Only





Rubin B, et al. National review of re-operation rates for modern benign prostatic hyperplasia procedures using a live claims database. Abstract IP03-06. Study Design: Retrospective observational study utilizing the TriNetX database (2004–2024), AJA 2025 from University of Vermont

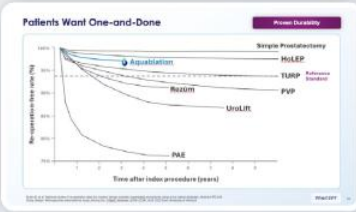
PROC



1 PATIENTS WANT: Complete Symptom Relief



2 PATIENTS WANT: Preserved Sexual and Urinary Function



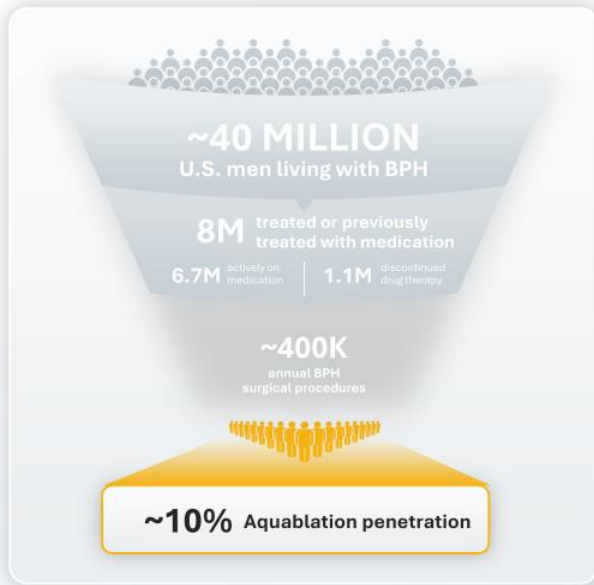
3 PATIENTS WANT: One Procedure Only

Aquablation is a COMPLETE Solution for BPH patients

Market research data on file

PROC

Improving Penetration of Aquablation Within the Surgical Segment



- 1 Misalignment:** In patient priorities and physician assumptions
- 2 Surgeon Comfort With Legacy Procedures**
- 3 Low Patient Awareness:** Minimal awareness of Aquablation as a differentiated option

We see both as opportunities and responsibilities to lead through education and execution

Data Analysis on File at PROCEPT BioRobotics

PROCEPT

A Complete BPH Solution Through the Lens of What Matters Most to Patients

| | Preservation of Continence | Durability | IPSS Symptom Reduction | Reduction of Post Void Residual Volume | Qmax Improvement | Recovery Time and Ease | Preservation of Erectile Function | Preservation of Ejaculatory Function |
|--|----------------------------|------------|------------------------|--|------------------|------------------------|-----------------------------------|--------------------------------------|
| Aquablation® Therapy | Green | Green | Green | Green | Green | Yellow | Green | Green |
| BPH Medications | Green | Grey | Red | Yellow | Green | Green | Green | Green |
| UroLift™ System | Green | Yellow | Yellow | Yellow | Yellow | Green | Green | Green |
| Rezūm Water Vapor Therapy™ | Green | Yellow | Yellow | Yellow | Red | Green | Green | Green |
| iTind™ | Green | Grey | Yellow | Yellow | Red | Green | Green | Green |
| Optilume® | Green | Grey | Yellow | Yellow | Yellow | Green | Green | Green |
| Transurethral resection of the prostate (TURP) | Green | Green | Green | Green | Yellow | Yellow | Yellow | Yellow |
| Laser Enucleation of the Prostate (LEP) | Yellow | Green | Green | Green | Red | Green | Yellow | Yellow |
| Greenlight™ Laser Therapy | Green | Yellow | Green | Green | Yellow | Yellow | Yellow | Yellow |
| Simple Prostatectomy | Green | Green | Green | Green | Red | Green | Yellow | Yellow |

PROCEPT

A Complete BPH Solution Through the Lens of What Matters Most to Patients



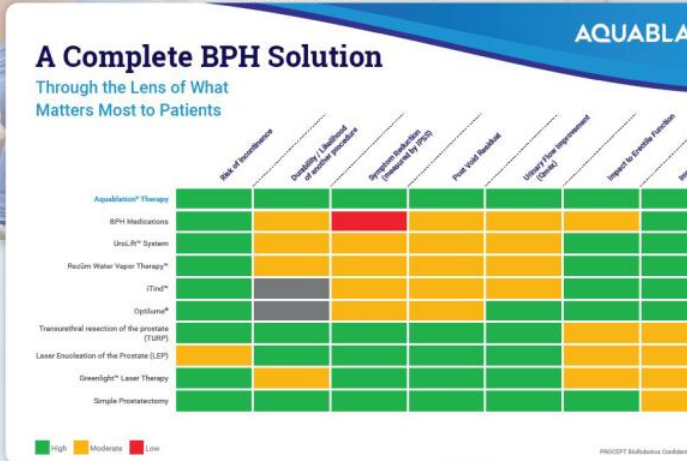


Operationalizing Our Clinical Differentiation

Standardize and strengthen the message

Operationalize that message:

- Selling tools
- Packaged research insights highlighting what BPH patients prioritize
- Peer-to-peer education programs



Improving Penetration of Aquablation Within the Surgical Segment



- 1 **Misalignment:** In patient priorities and physician assumptions
1-2% unaided awareness for BPH patients
- 2 **Surgeon Comfort With Legacy Procedures**
- 3 **Low Patient Awareness:** Minimal awareness of Aquablation as a differentiated option

Data Analysis on File at PROCEPT BioRobotics

PROCEPT

We Start With Patients Considering Alternative Surgical Options



OBJECTIVE:

Share capture within existing surgical market

APPROACH:

- In-practice education tools (treatment comparison guides, short videos for consult rooms)
- Digital ads for patients searching surgical terms
- Patient testimonials reinforcing clinical differentiation



Data Analysis on File at PROCEPT BioRobotics

PROCEPT

Then Expand to Dissatisfied Medically Managed Patients



Alpha Blockers

Tamsulosin:



Short-term side effects include:

• Dizziness, Fatigue, Flushing

Long-term side effects include:

• Sexual side effects continue

~40 MILLION
U.S. men living with BPH

8M treated or previously treated with medication
6.7M actively on medication | 1.1M discontinued drug therapy

~400K
annual BPH surgical procedures

~10% Aquablation penetration

Dizziness, fatigue, Ejaculatory changes

Sexual side effects continue



Combination Therapy

5-alpha reductase inhibitors (5-ARIs)

Finasteride:

AT131

Short-term side effects include:

- Erectile dysfunction

Long-term side effects include:

- Sexual adverse effects may persist

Contemporary studies suggest BPH medications may negatively affect mental and psychological status, sexual function, and overall health

Data Analysis on File at PROCEPT BioRobotics

Bortnick EM, Simma-Chiang V, Kaplan SA. Long-term Consequences of Medical Therapy for Benign Prostatic Hyperplasia. Rev Urol. 2019;21(4):154-157.

PROCEPT

Then Expand to Dissatisfied Medically Managed Patients

~40 MILLION
U.S. men living with BPH

8M treated or previously treated with medication
6.7M actively on medication | 1.1M discontinued drug therapy

~400K
annual BPH surgical procedures

~10% Aquablation penetration

OBJECTIVE:

Educate patients already frustrated with medication and under care of a urologist

APPROACH:

- Digital engagement (social media, search, educational email)
- Geo-targeted digital ads around Aquablation centers
- Symptom self-assessment tools
- Direct-to-consult scheduling links



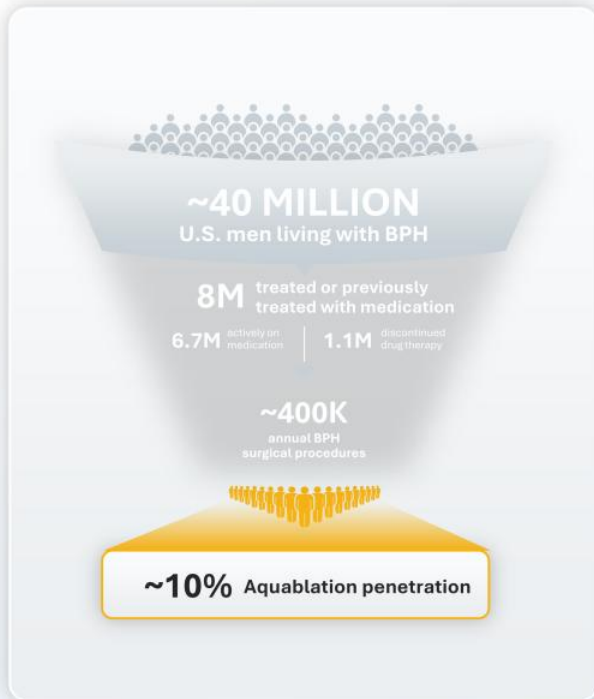
Google



Data Analysis on File at PROCEPT BioRobotics

PROCEPT

Then Expand to Dissatisfied Medically Managed Patients



~20%
Of BPH Patients

Request specific BPH procedures by name

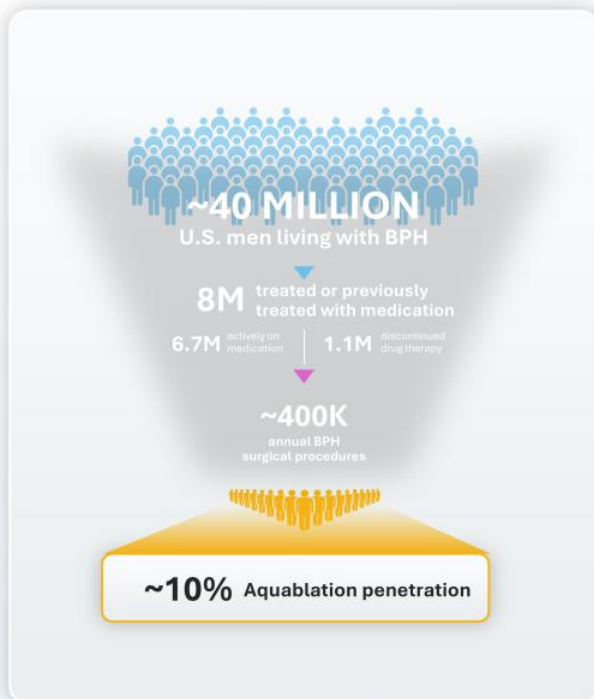
>70%
Of Urologists

Consider patient requests if efficacy is similar

Data Analysis on File at PROCEPT BioRobotics

PROCEPT

Long-Term, We Can Broaden Our Educational Efforts



OBJECTIVE:

Build broader awareness only after conversion pathways are validated

APPROACH:

- Broader therapy development campaigns
- Patient webinars & community education



MEN'S HEALTH
AWARENESS MONTH



Data Analysis on File at PROCEPT BioRobotics

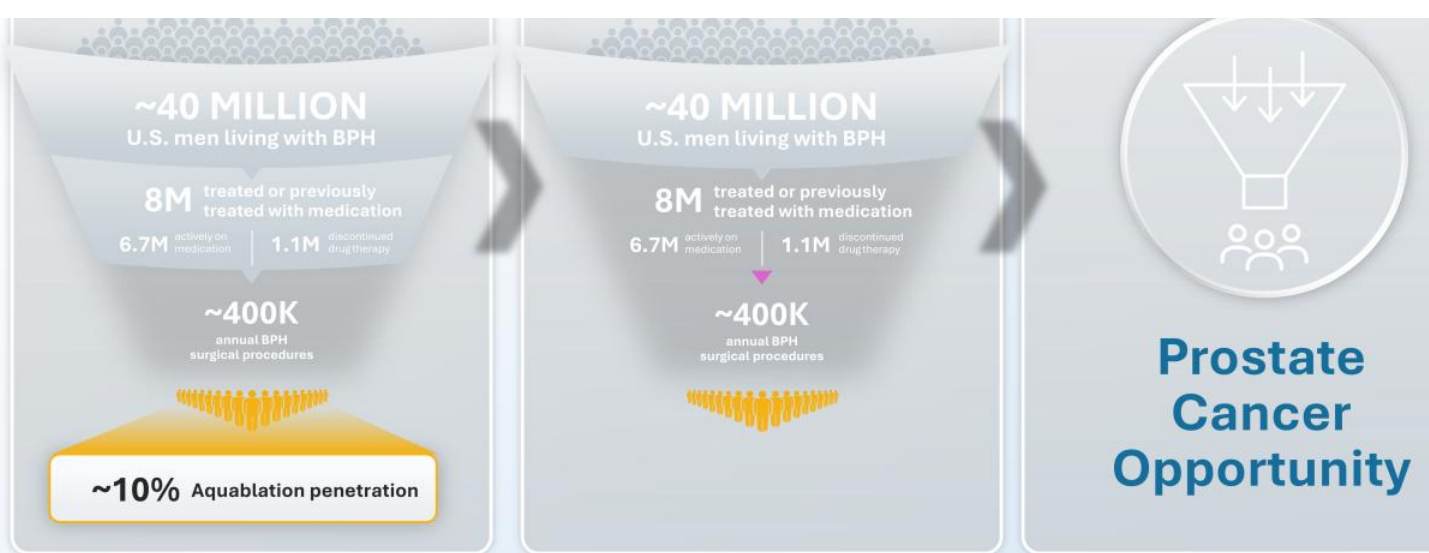
PROCEPT

Multiple Initiatives to Drive Differentiated Growth Across the Horizon

Drive Procedure Growth Through Improved Execution

Educate Patients That Waiting Isn't Winning

Expand Applications



1. Vuichoud C, Loughlin KR. Benign prostatic hyperplasia: epidemiology, economics and evaluation. Can J Urol. 2015 Oct;22 Suppl 1:1-6. PMID: 26497338.
 2. Based on management estimates and data provided by AcuityMD, Dec 19, 2026 Data Release, US market estimates, Q4 2024 – Q3 2025

PRO

PROCEPT
BIOROBOTICS®

Prostate Cancer: Our Next Frontier

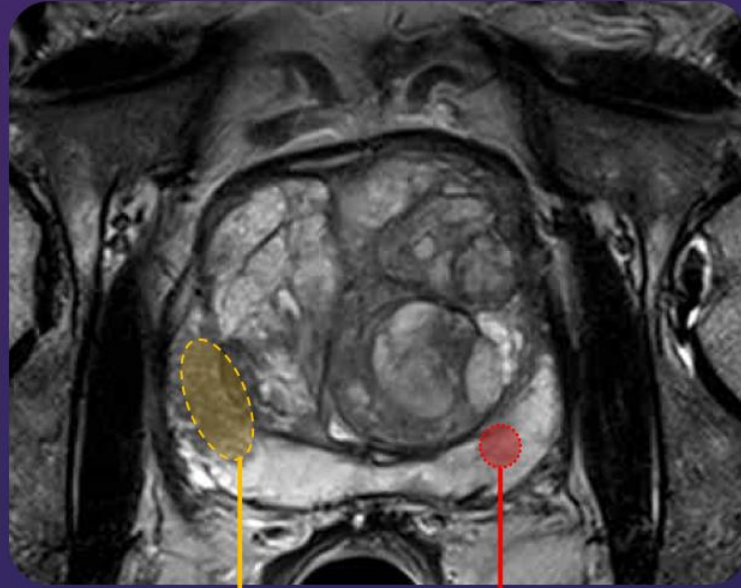
Barry Templin
Chief Technology Officer



Prostate Cancer Is A Multifocal Disease Requiring A Whole Gland Treatment

Prostate cancer is not a focal disease, and focal treatment is limited

>40% of patients with GG2 cancer on one side of the prostate had unidentified GG2 cancer on the other side at final pathology¹



Visible on MRI

Not visible on MRI

1. Johnson et al Cancer 2019

Methodical Approach in Evidence Development

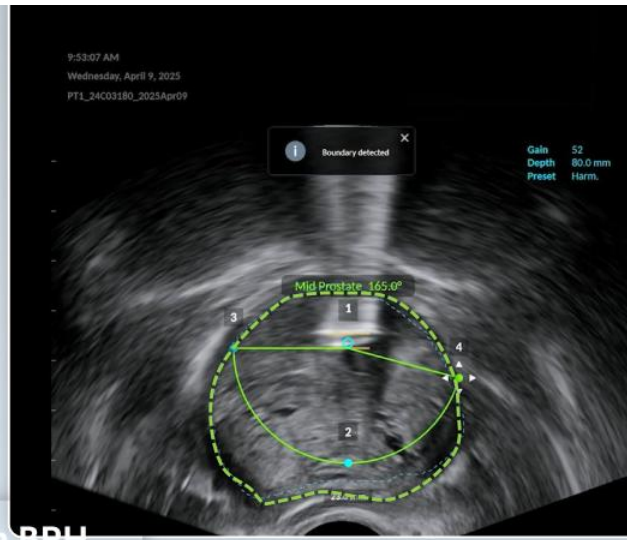
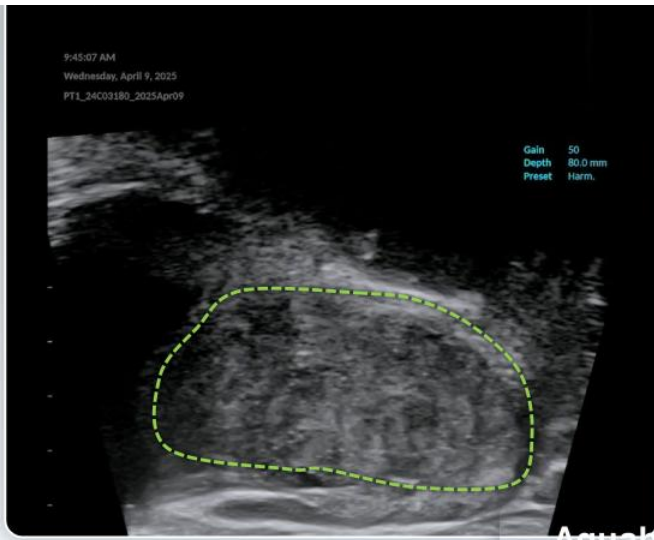


| | PRCT001.A | Endeavor Health Series | PRCT002 | Retrospective Study | PRCT001.B | WATER IV |
|------------------------|---------------------------------|-------------------------------------|--|--|---|-----------------------------|
| | Prospective, Feasibility | Retrospective, Foundational | Prospective, Foundational | Retrospective, Foundational | Prospective, Foundational | Prospective Randomized |
| Cancer Patient Profile | GG 1 – 2 | GG 1 | GG 1 – 2 | Salvage RP in previous Aquablation patients | GG 1-3 | GG 1-3 RCT vs RP |
| Cohort Size (n=) | 5 | 1,445 | 22 | 14 | 119 | 280 |
| Primary Research Aim | Safety and CTC characterization | Cancer progression characterization | Procedural safety & cancer progression precursor to WATER IV | Evidence that salvage RP is feasible following an Aquablation BPH or PCa procedure | Procedural development & cancer progression | Pivotal study efficacy & sa |

RP = Radical Prostatectomy CTC = circulating tumor cells GG = Grade Group RCT = Randomized Control Trial

A Typical BPH Surgical Technique: Planning



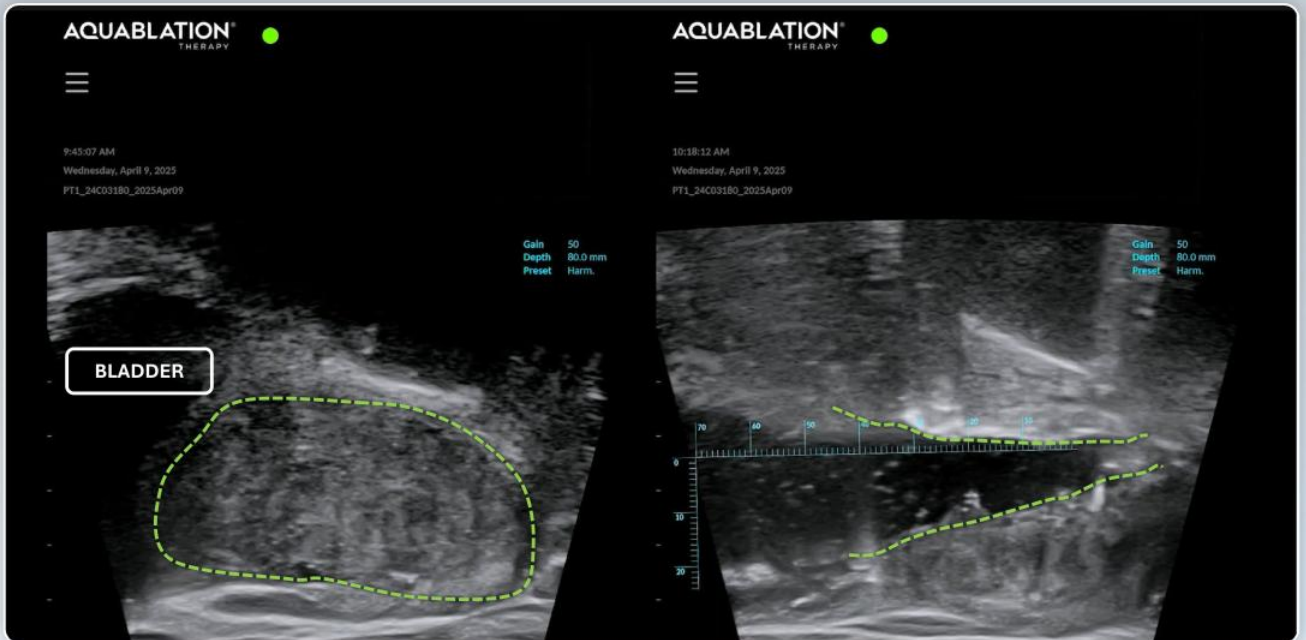


Aquablation BPH

PRO

BPH Resection With Aquablation Results in a Large Unobstructed Channel

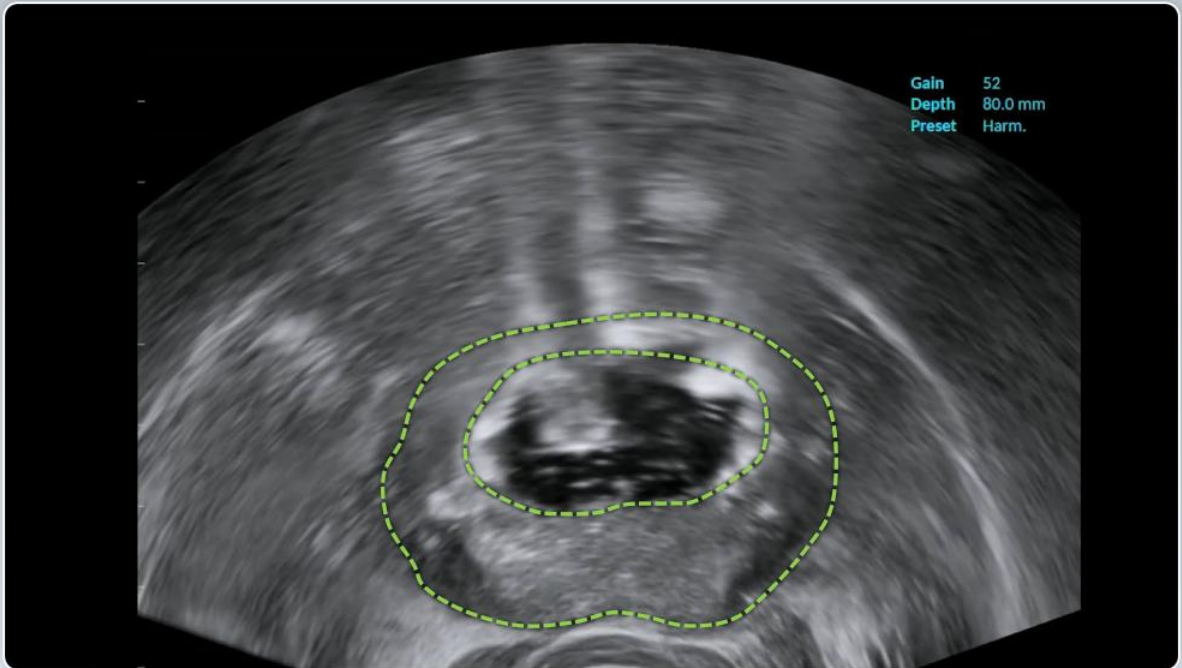
Sagittal view



PRO

BPH Resection With Aquablation Results in a Large Unobstructed Channel

Transverse view



PROC

Planning for a Prostate Cancer Case with Aquablation

MRI Details to Help Inform Surgical Approach

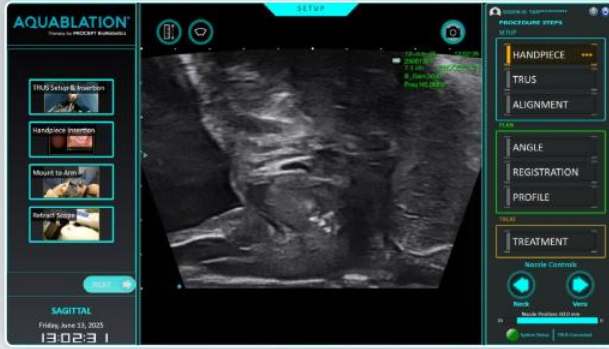
Aquablation PCa

The block contains two 3D models of a prostate gland, one showing a yellow spot representing a tumor. Below the models are two screenshots of the Aquablation software interface. The left screenshot shows the 'SAGITTAL' view with a dashed green line outlining the tumor. The right screenshot shows the 'TRANSVERSE' view with a dashed green line outlining the tumor. Both screenshots include a 'PROCEDURE STEPS' panel on the right with options for HANDPIECE, TRUS, ALIGNMENT, ANGLE, REGISTRATION, PROFILE, and TREATMENT. The 'TREATMENT' panel has 'Mark' and 'Verify' buttons. The bottom of each screenshot shows the time and date: Friday, June 25, 2025.

PROC

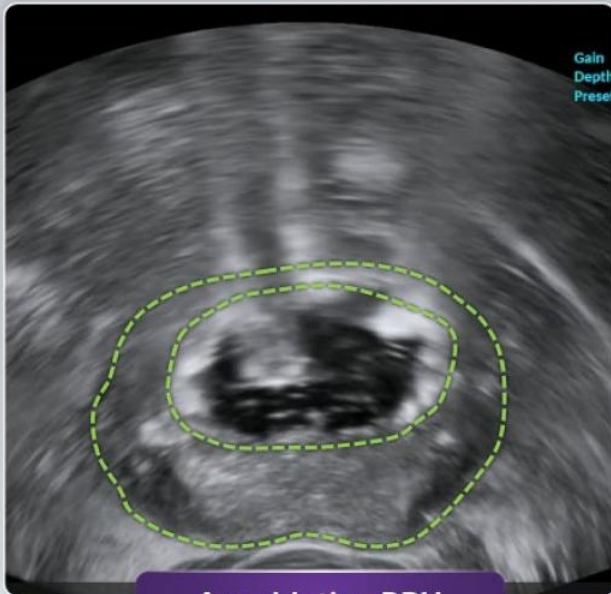
Aquablation for Prostate Cancer Achieves Whole Gland Resection While Sparing the Capsule

The block contains two screenshots of the Aquablation software interface. The left screenshot shows the 'SAGITTAL' view with a dashed green line outlining the tumor. The right screenshot shows the 'TRANSVERSE' view with a dashed green line outlining the tumor. Both screenshots include a 'PROCEDURE STEPS' panel on the right with options for HANDPIECE, TRUS, ALIGNMENT, ANGLE, REGISTRATION, PROFILE, and TREATMENT. The 'TREATMENT' panel has 'Mark' and 'Verify' buttons. The bottom of each screenshot shows the time and date: Friday, June 25, 2025.

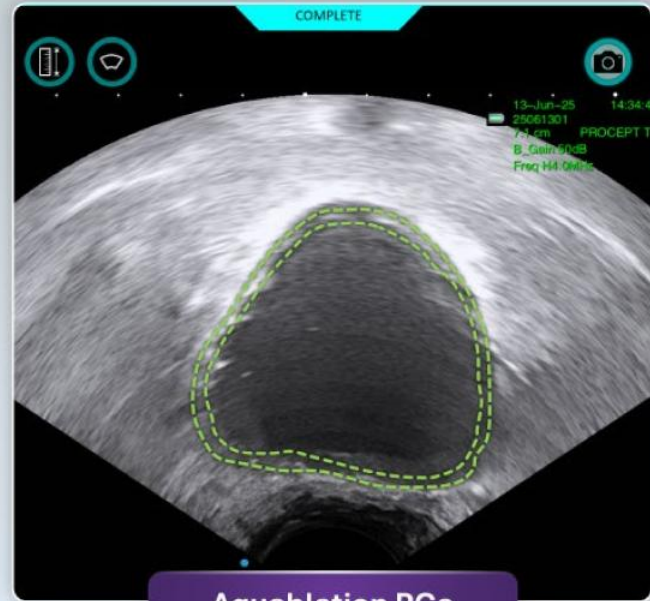


PROC

Aquablation's Versatility Can Deliver an Appropriate Procedure for both BPH and PCa Patients



Aquablation BPH

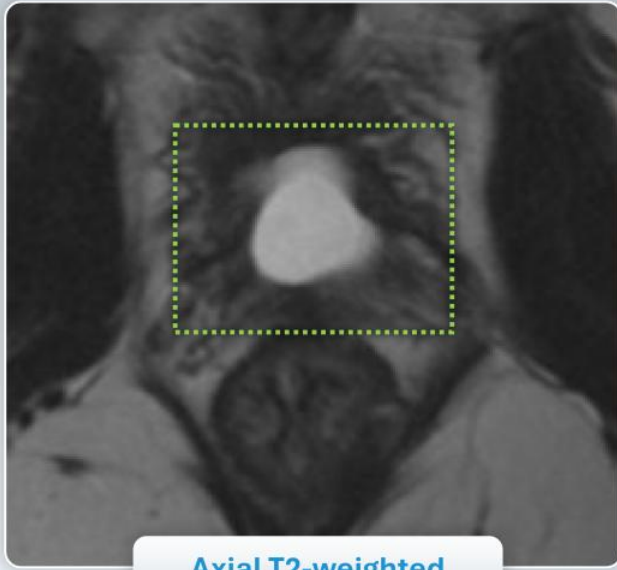


Aquablation PCa

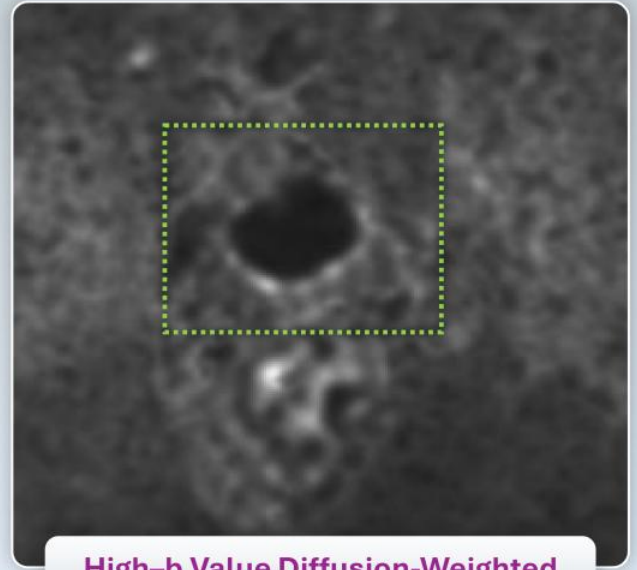
PROC

Post Aquablation Prostate Cancer Resection: 6M MRI

MRI Confirmation of Tissue Removal



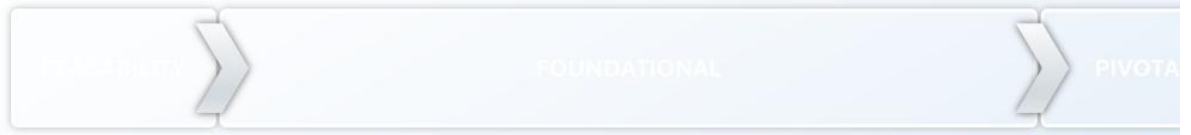
Axial T2-weighted



High-b Value Diffusion-Weighted

PROC

We Are Taking a Methodical Approach in Evidence Development



| | PRCT001.A | Endeavor Health Series | PRCT002 | Retrospective Study | PRCT001.B | WATER IV |
|------------------------|---------------------------------|-------------------------------------|--|--|---|-----------------------------|
| | Prospective, Feasibility | Retrospective, Foundational | Prospective, Foundational | Retrospective, Foundational | Prospective, Foundational | Prospective Randomized |
| Cancer Patient Profile | GG 1 – 2 | GG 1 | GG 1 – 2 | Salvage RP in previous Aquablation patients | GG 1-3 | GG 1-3 RCT vs RP |
| Cohort Size (n=) | 5 | 1,445 | 22 | 14 | 119 | 280 |
| Primary Research Aim | Safety and CTC characterization | Cancer progression characterization | Procedural safety & cancer progression precursor to WATER IV | Evidence that salvage RP is feasible following an Aquablation BPH or PCa procedure | Procedural development & cancer progression | Pivotal study efficacy & sa |

RP = Radical Prostatectomy

CTC = circulating tumor cells

GG = Grade Group

RCT = Randomized Control Trial

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We Are Taking a Methodical Approach in Evidence Development

PRCT001.A

PIVOTAL

| | Endeavor Health Series | PRCT002 | Retrospective Study | PRCT001.B | WATER IV |
|-----------------------------|-------------------------------------|--|--|---|---------------------------------|
| | Prospective, Foundational | Prospective, Foundational | Retrospective, Foundational | Prospective, Foundational | Prospective Randomized |
| Cancer Patient Population | GG 1 | GG 1-2 | Salvage RP in previous Aquablation patients | GG 1-3 | GG 1-3 |
| Cohort Size (n=) | 145 | 22 | 14 | 119 | 280 |
| Primary Research Objectives | Cancer progression characterization | Procedural safety & cancer progression precursor to WATER IV | Evidence that salvage RP is feasible following an Aquablation BPH or PCa procedure | Procedural development & cancer progression | Pivotal study efficacy & safety |

RP = Radical Prostatectomy, GG = Grade Group, RCT = Randomized Control Trial

Prospective, Feasibility

GG 1 – 2

5

Safety and CTC characterization

We Are Taking a Methodical Approach in Evidence Development

PRCT001.A

Prospective, Feasibility

GG 1 – 2

5

Safety and CTC characterization

Enrollment for PRCT001 Now Complete

PRCT001.B

Prospective, Foundational

GG 1-3

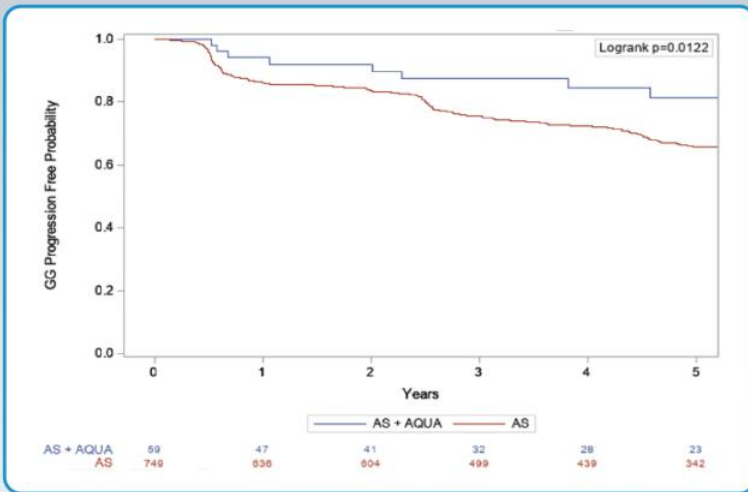
119

Procedural development & cancer progression

We Are Taking a Methodical Approach in Evidence Development

| PRCT001.A | Endeavor Health Series | PRCT001.B |
|------------------------------|-------------------------------------|---|
| Prospective, Feasibility | Retrospective, Foundational | Prospective, Foundational |
| GG 1 – 2 | GG 1 | GG 1-3 |
| Cohort Size (n=) | 1,445 | 119 |
| Primary Research / Objective | Cancer Progression characterization | Procedural development & cancer progression |

ENDEAVOR Health Series: Aquablation Significantly Slowed the Progression of Cancer



By removing a moderate amount of prostate tissue, Aquablation delayed the natural progression of cancer in this study

44%
Fewer upgrades at 5 years compared to Active surveillance

The PRCT001 Study: Demographics Reflected the Typical Profile of PCa Where Majority of Patients Had Cancer in the Peripheral Zone

| Patient Demographics | | Anatomy & Procedure Details | |
|----------------------|------------|-----------------------------|-------------|
| Age (years old) | 66.4 ± 6.7 | Prostate Vol. Avg | 65.3 ± 31.5 |

Grade-Group

| | |
|-----|--------------|
| GG1 | 36% (44/124) |
| GG2 | 61% (76/124) |
| GG3 | 3% (4/124) |

| | |
|---|----------------|
| Patients with MR visible lesion (≥PIRADS 3) | 77% (95/124) |
| Percent Patients with IPSS ≥ 8 | 100% (124/124) |
| IPSS | 17.4 ± 6.7 |

Cancer Location

| | |
|-------------------------------------|-------------|
| Transitional Zone | 32% (24/75) |
| Peripheral Zone | 57% (43/75) |
| Transitional Zone + Peripheral Zone | 7% (5/75) |
| Zone not specified | 4% (3/75) |
| LOS (days) | 1.3 ± 1.1 |
| Cath Duration (days) | 2.3 ± 2.2 |

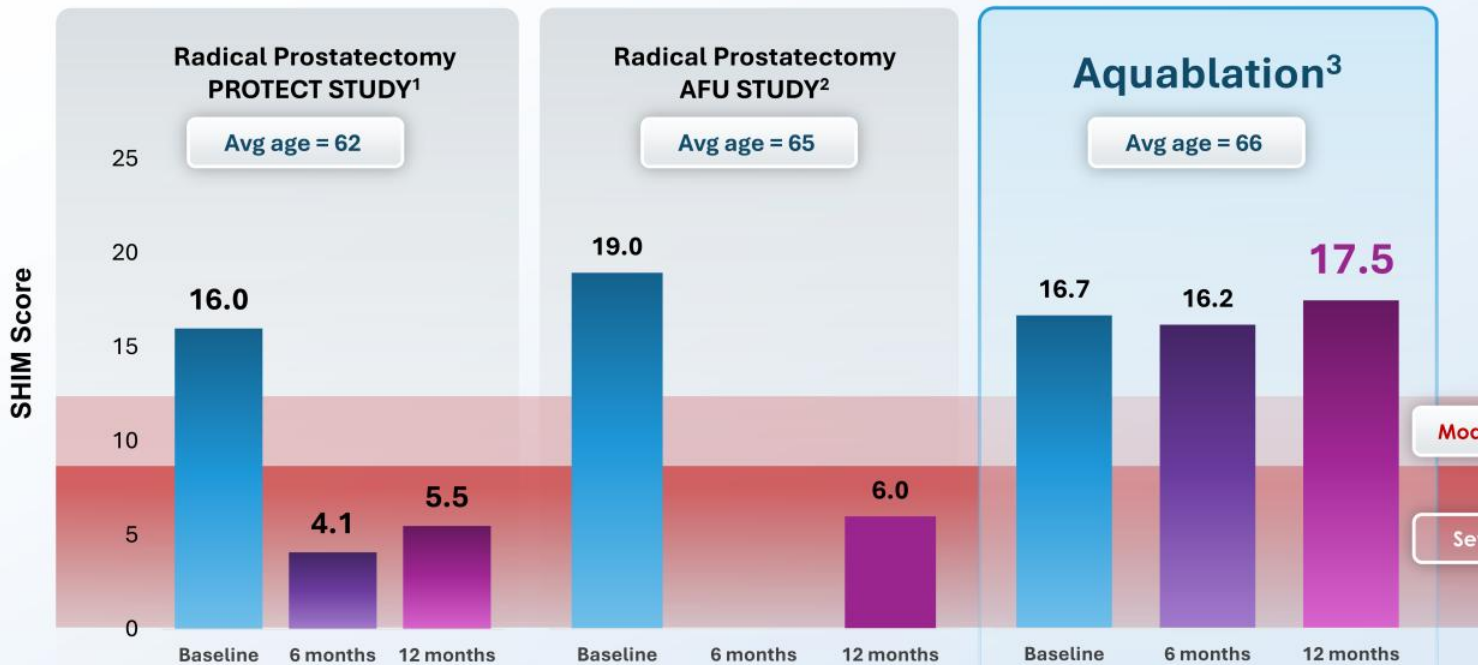
124 Prostate Cancer Patients Treated

90 Patients with 6 mo follow up

37 Patients with 12 mo follow up

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PRCT001 Study: Aquablation Patients Had Stable Erectile Function Through 12 Months

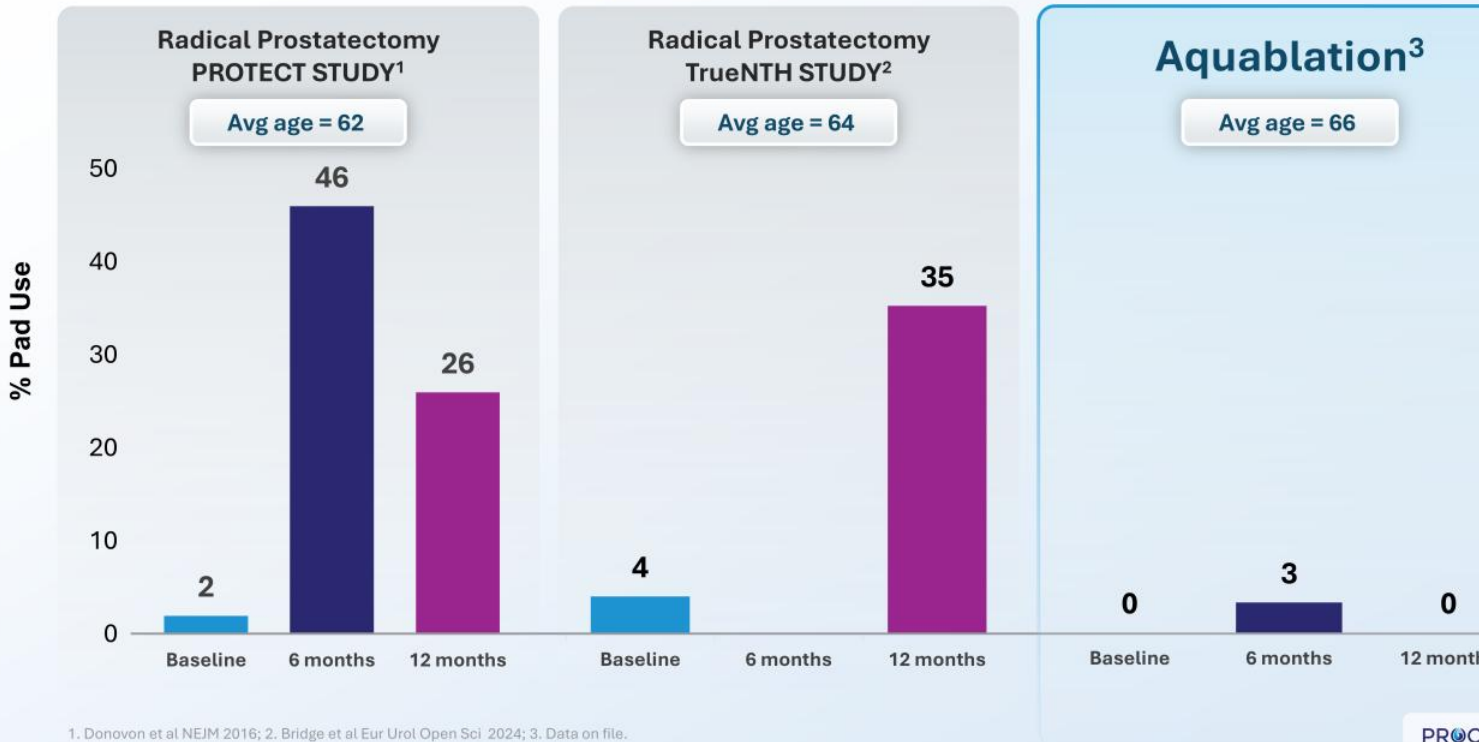


1. Donovan et al NEJM 2016; 2. Ploussard et al 2024; 3. Data on file

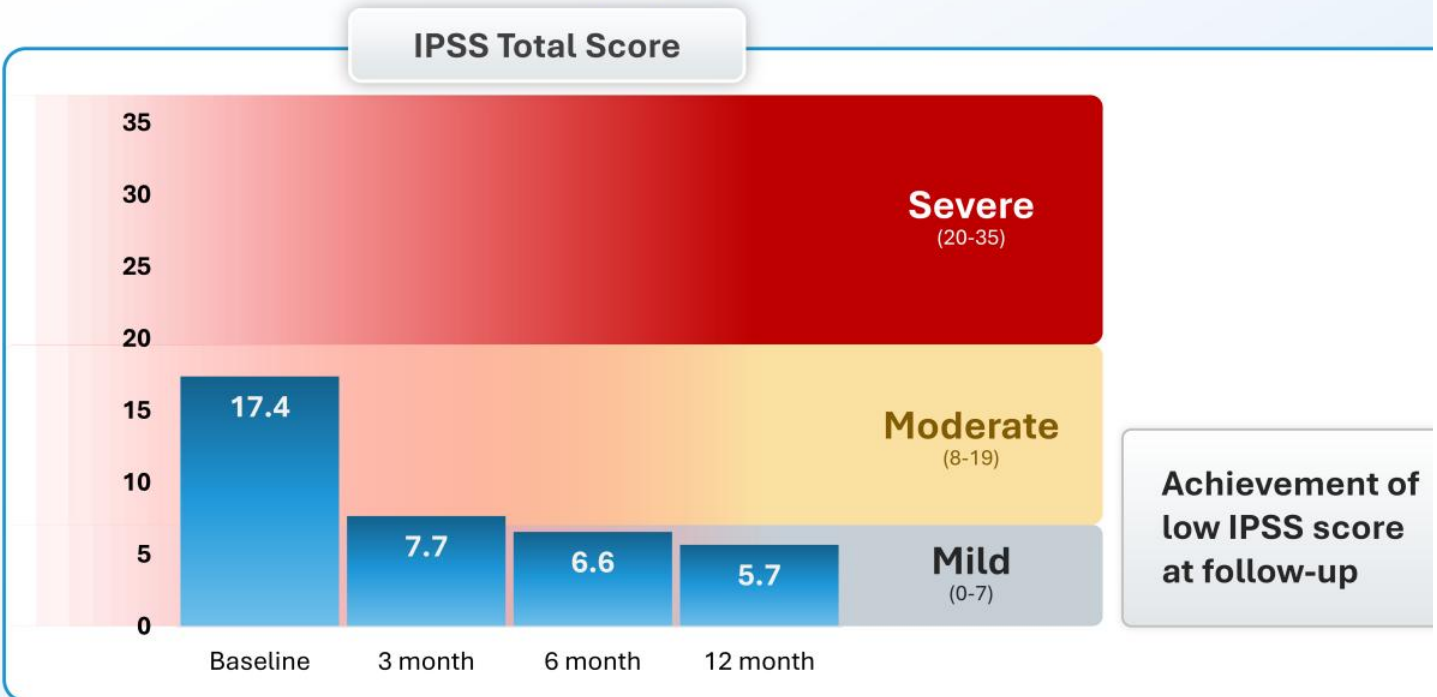
The Sexual Health Inventory for Men (SHIM) is a 5-item, validated questionnaire (also known as IIEF-5) used by urologists to assess erectile dysfunction (ED) severity, with total scores ranging from 1 to 25. A score of 21 or less suggests ED, categorized as: 22-25 (None), 17-21 (Mild), 12-16 (Mild-moderate), 8-11 (Moderate), and 1-7 (Severe).

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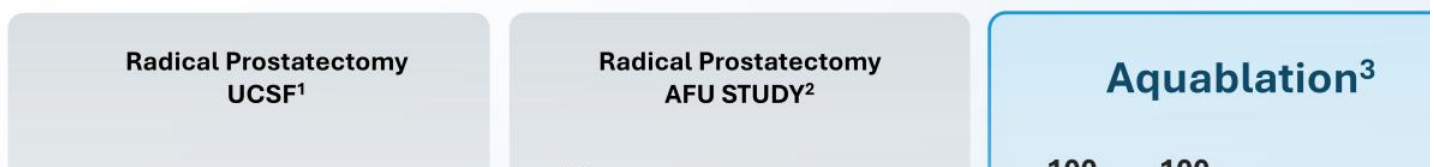
The PRCT001 Study: Aquablation Patients Had Low and Sustained Rates of Urinary Incontinence Out to 12 Months

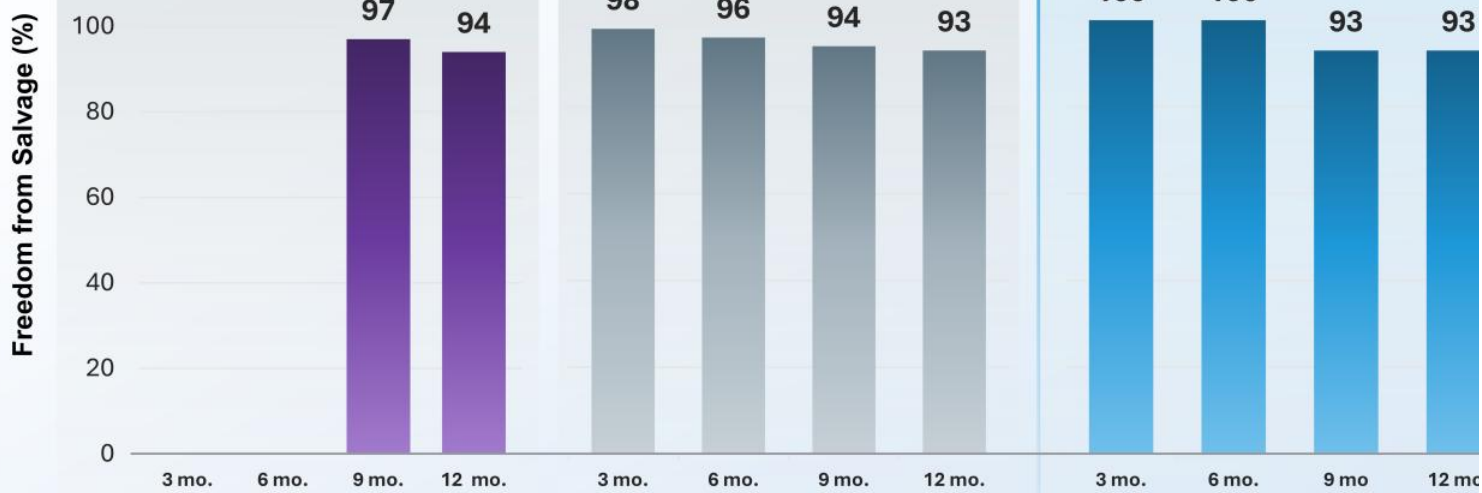


The PRCT001 Study: Aquablation Patients Had An Additional Benefit of BPH Symptom Reduction



The PRCT001 Study: Freedom from Additional (Salvage) Treatment in Aquablation vs Published Radical Prostatectomy Data

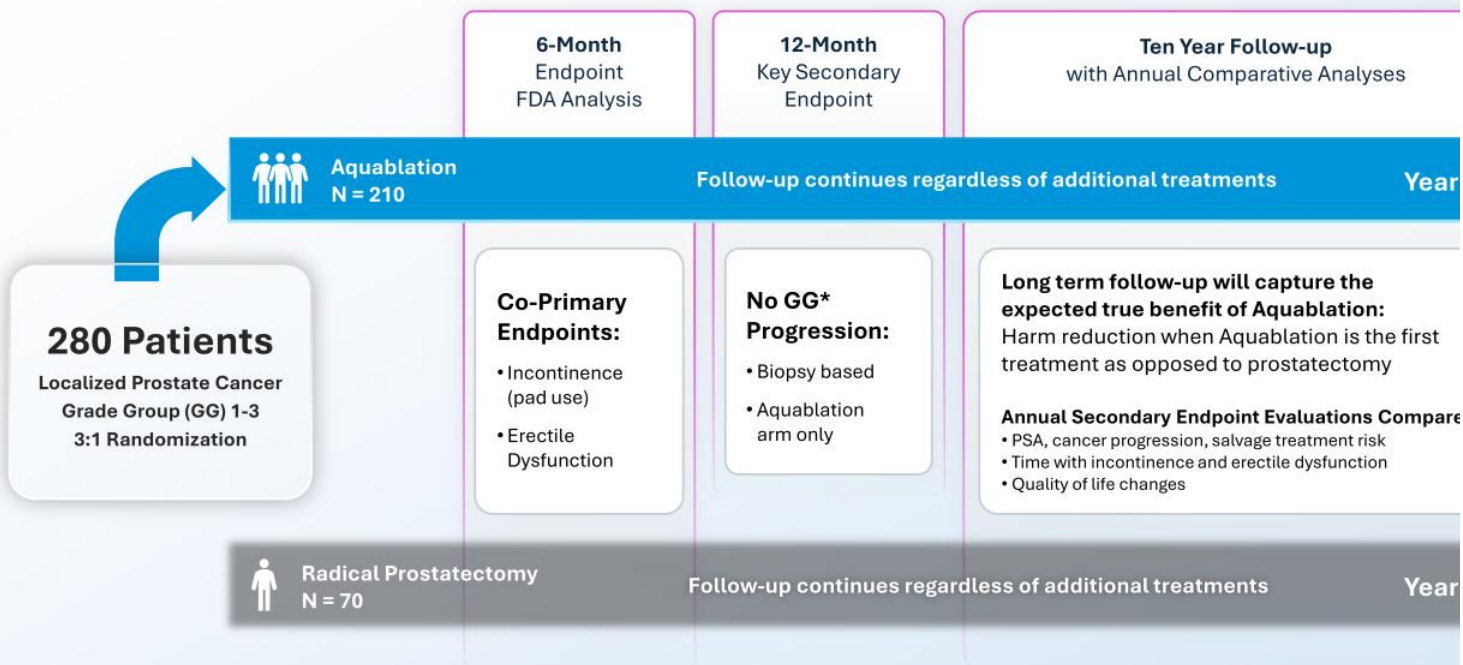




1. Szymaniak et al 2023; 2. Ploussard et al 2024; 3. Data on file

WATER IV PCa: Aquablation vs. Prostatectomy RCT

Trial design



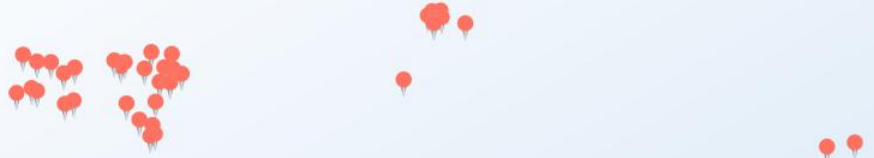
WATER IV CLINICAL TRIAL

One year of enrollment progress

Goal: 280
randomized
patients

31 global centers
recruiting as we
exited 2025

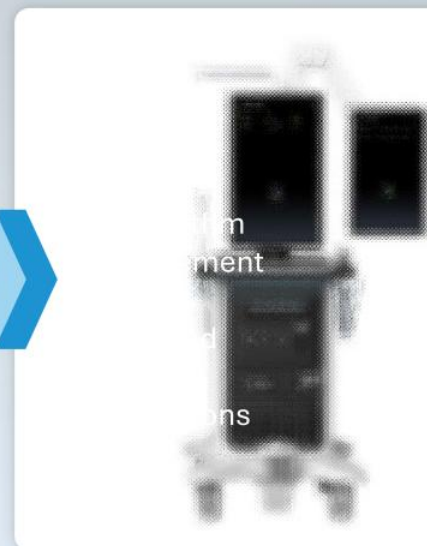
Averaging <60 days from
consent to treatment



On pace to complete all procedures by mid-2026
significantly faster than anticipated

PROC

Our Technology Enables Expanded Applications Beyond BPH



PROC

We Believe Aquablation Will Play an Important Role in
Treatment of Prostate Cancer

1

2

3

Prostate Cancer is a **multi-focal disease that requires a whole gland treatment**

Aquablation therapy can **treat the whole gland including the peripheral zone**

Growing body of clinical evidence supports Aquablation can become a frontline option



We are
the First
Company

To receive **FDA-IDE approval to enroll a randomized trial** comparing surgical therapy vs. radical prostatectomy

PROCEPT

PROCEPT
BIOROBOTICS®

Financial Update

Kevin Waters
Chief Financial Officer



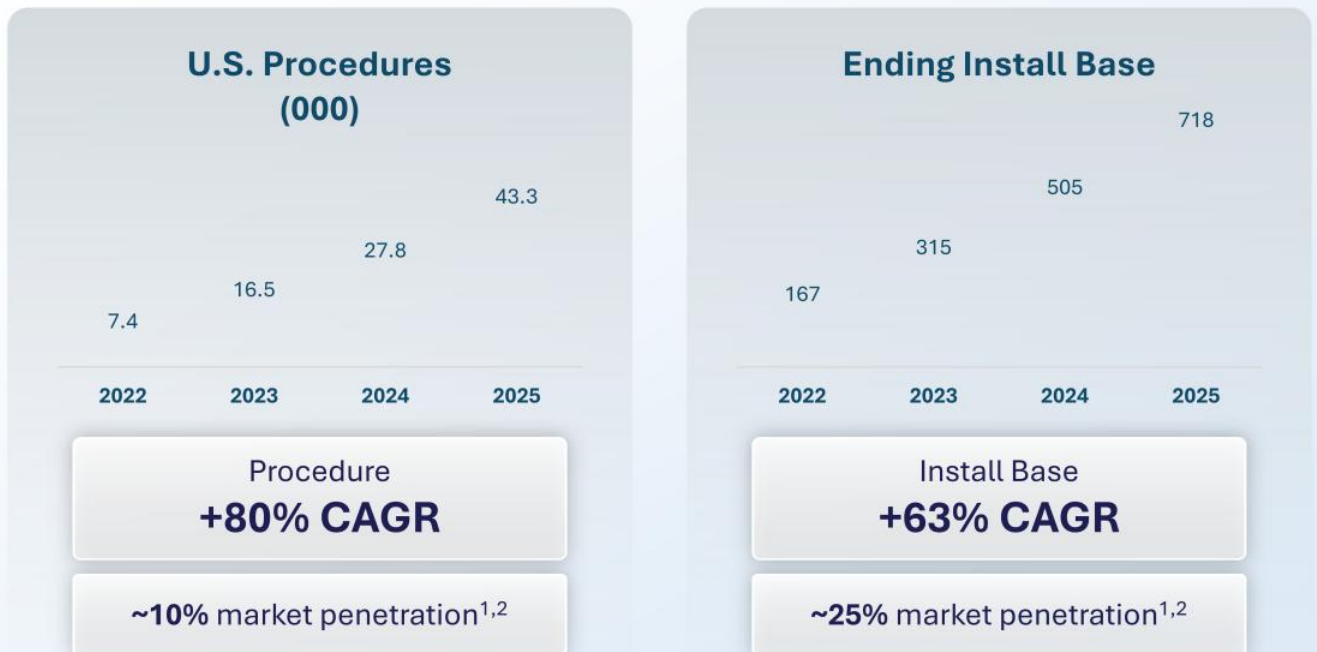
Strong track record of driving growth, expanding margins and scaling efficiently



*Adjusted EBITDA is a financial measure that is not prepared in accordance with generally accepted accounting principles in the United States (GAAP). For more information about the Company's use of non-GAAP financial measures, please see appendix

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Aquablation remains significantly underpenetrated in the U.S.



Market penetration estimates calculated based on market size from the following sources

- Vuichoud C, Loughlin KR. Benign prostatic hyperplasia: epidemiology, economics and evaluation. Can J Urol. 2015 Oct;22 Suppl 1:1-6. PMID: 26497338.
- Based on management estimates and data provided by AcuityMD, Dec 19, 2026 Data Release, US market estimates, Q4 2024 – Q3 2025

PROCEPT



Financial Outlook



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2026 – 2027: Key Financial Metrics



Strong Revenue Growth

- 2026 revenue guidance¹ of **\$390 to \$410 million**, representing growth of 27% to 33%
- **25% to 30% annual revenue growth in 2027¹** driven by increased procedure growth from new and existing accounts, and Hydros replacement cycle as legacy AquaBeam systems age



Expanding gross margins

- 2026 gross margin **65%¹**.
- Targeting 2027 gross margin range of **68% to 70%¹**
- **HP margin** will be tailwind to corporate gross margin expansion



Improved operating leverage

- 2026 adjusted **EBITDA^{1,2} loss** guidance of \$30 to \$17 million
- 2027 adjusted **EBITDA^{1,2} gain** of \$25 to \$30 million

1. 2026 + 2027 financial guidance issued on February 26, 2026

2. Adjusted EBITDA is a financial measure that is not prepared in accordance with generally accepted accounting principles in the United States (GAAP). For more information about the Company's use of non-GAAP financial measures, please see appendix

Scalable Revenue Growth Drivers



Strong Revenue Growth



Expanding gross margins



Improved operating leverage

Commercial execution to drive strong revenue growth

- Incentivize **organic, same facility procedure growth**
- Activate New Account **Launch Teams**
- Handpiece **Pricing Discipline**
- Initiate **Replacement Cycle**
- Explore **operating lease pilots** to expand into additional customer groups

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2026 Revenue Guidance*

\$390M to \$410M
Total Revenue

\$340M to \$359M
U.S. Revenue

\$50M to \$51M
Int'l Revenue

\$95M to \$100M
System Revenue

\$221M to \$235M
Handpiece + Other
Consumable Revenue

~\$24M
Service Revenue

\$3,500
Handpiece Price

60,000 to 64,000
Procedures

**HPs sold to align with
procedure volumes**

*2026 financial guidance issued on February 25, 2026

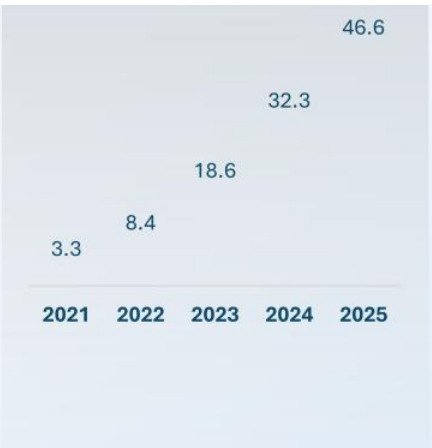
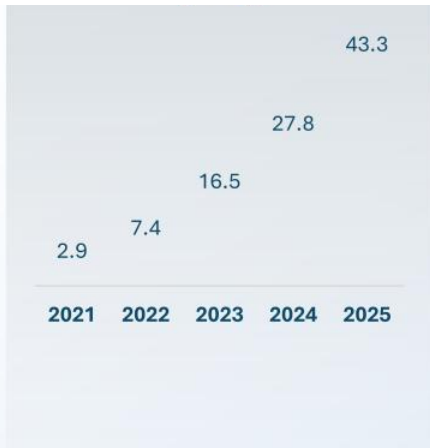
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2026 Guidance* Assumes 1:1 HP to Procedure ratio

**U.S. Procedures
(000)**

**U.S. Handpieces
(000)**

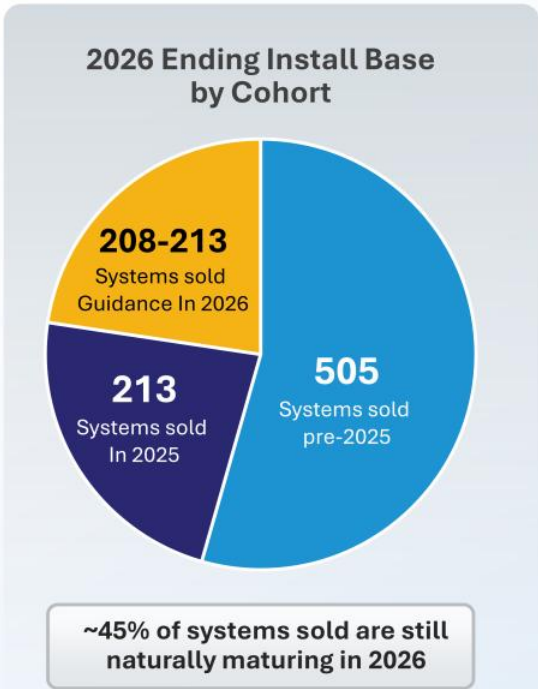
**% Handpieces to
Procedures**



Handpiece ordering should more closely track procedure volume going forward

*2026 financial guidance issued on February 25, 2026

2026 Procedure Contribution Breakdown to Achieve Low End of Guidance Range*



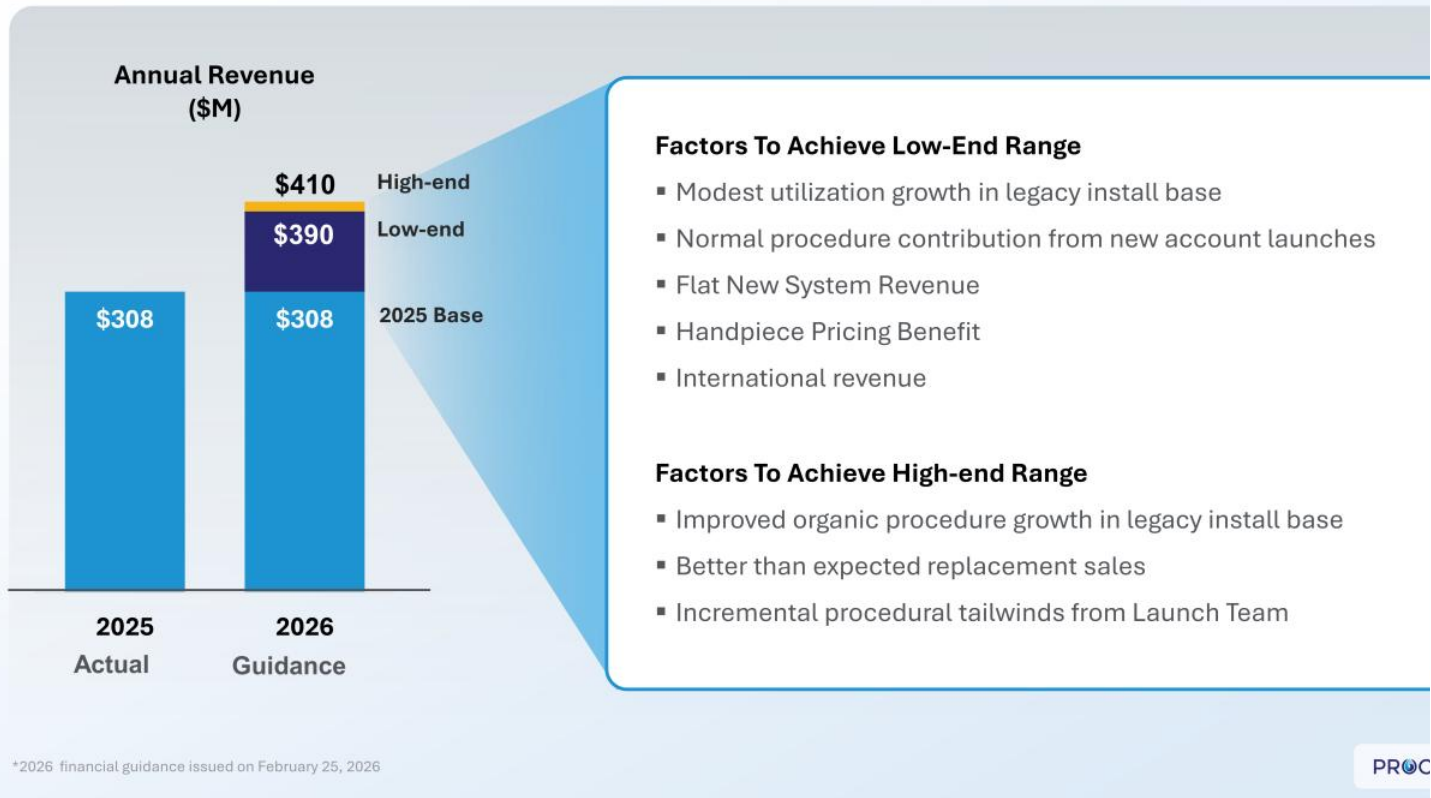
2025 Cohort will have full year contribution calendar 2026

With launches occurring throughout the year, the 2025 cohort contributes for only about half the year

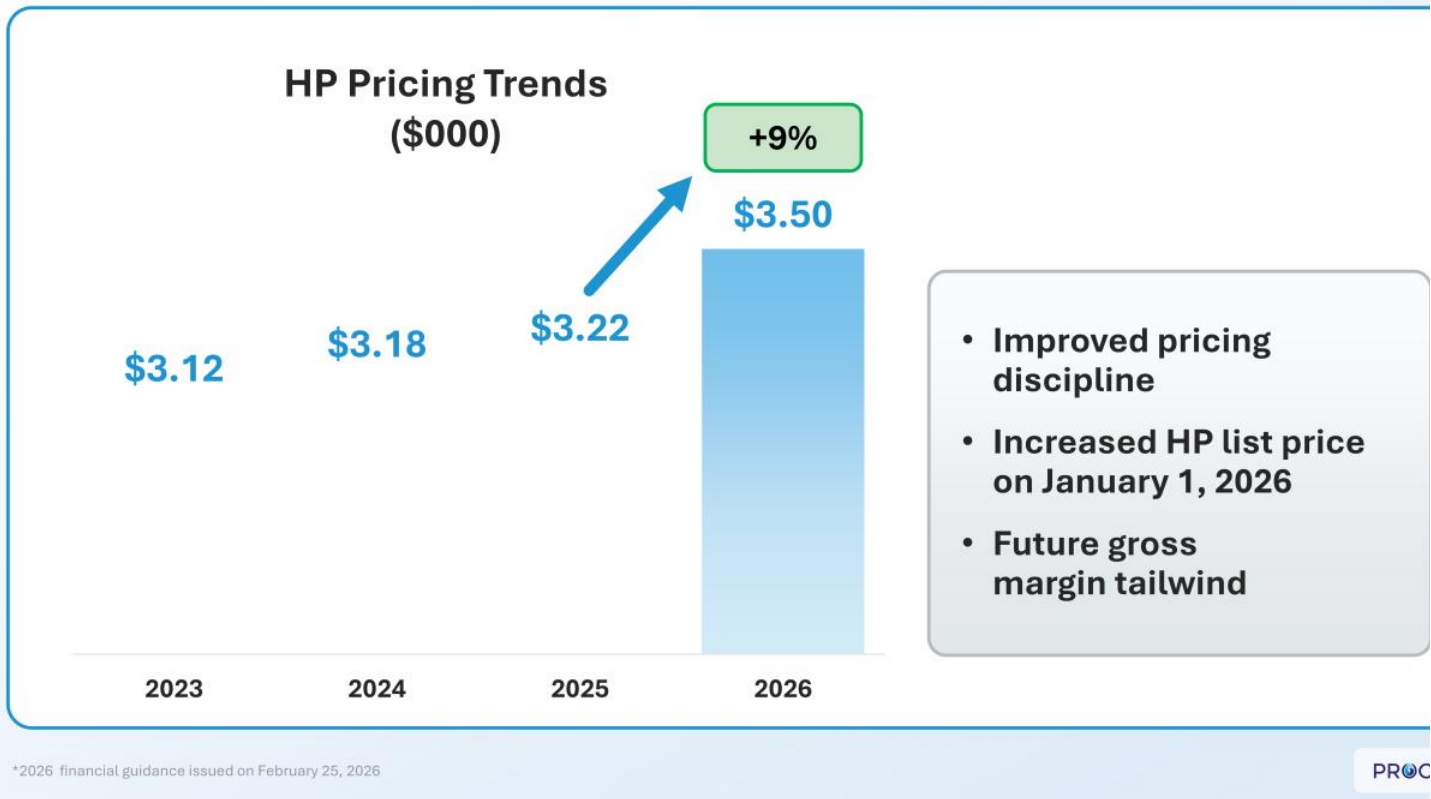
2026 guidance assumes only modest additional benefit from Launch Teams compared to prior years' launch contribution

*2026 financial guidance issued on February 25, 2026

2026 Revenue Build by Segment*

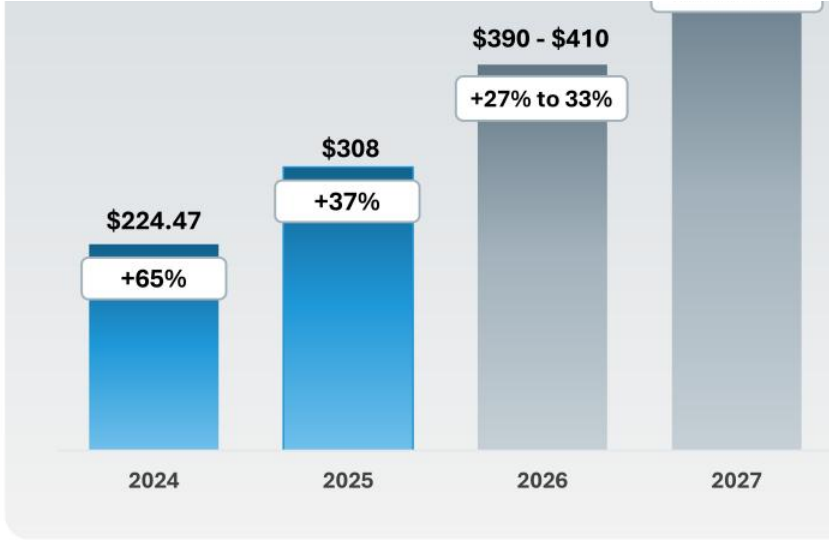


HP pricing will be meaningful revenue tailwind in 2026*



Global Revenue Profile*





2027 Revenue Drivers

- Growing U.S. Install Base
- Increased Replacement Revenue
- Stable handpiece & system pricing
- Continued penetration of UK + Japan market

*2026 + 2027 financial guidance issued on February 26, 2026

PR^{OC}

Long Term Profitability and EBITDA Expansion



PR^{OC}

Clear Path to Profitability



Strong Revenue Growth



Expanding gross margins



Improved operating leverage

Operational Efficiencies Driven by

- HP revenue mix will be meaningful tailwind to corporate gross margin expansion
- Disciplined HP pricing
- Targeting R&D as percent of sales ratio of mid-teens long-term
- Introduce hybrid clinical support model to scale sales force long-term

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How to Achieve 68% to 70% gross margin by 2027* ...

Handpiece

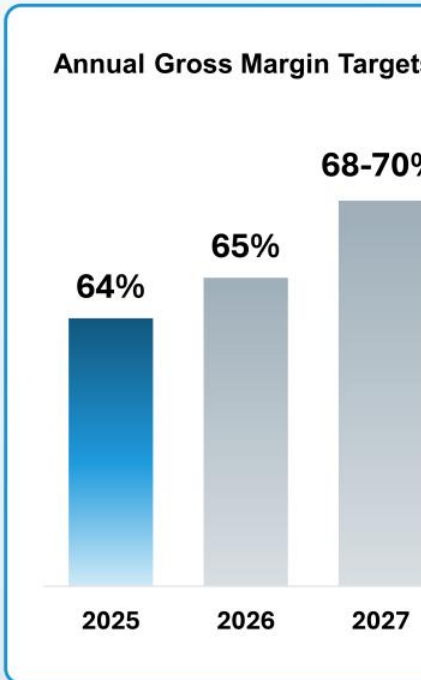
- Favorable Unit Margin
- HP revenue will be tailwind to multi-year gross margin expansion as percent of global revenue increases

Capital

- Declining mix of system revenue will reduce pressure on multi-year gross margin expansion
- Chinese ultrasound supplier is largest tariff exposure

Tariff Mitigation Strategy

Working with suppliers to onshore manufacturing
Tariff exposure expected to stabilize in 2027
 2025 - \$1.3M 2026E - \$5M to \$6M 2027E - \$6M



*2026 + 2027 financial guidance issued on February 26, 2026

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R&D spend will remain robust, but will be meaningful source of future operating leverage*

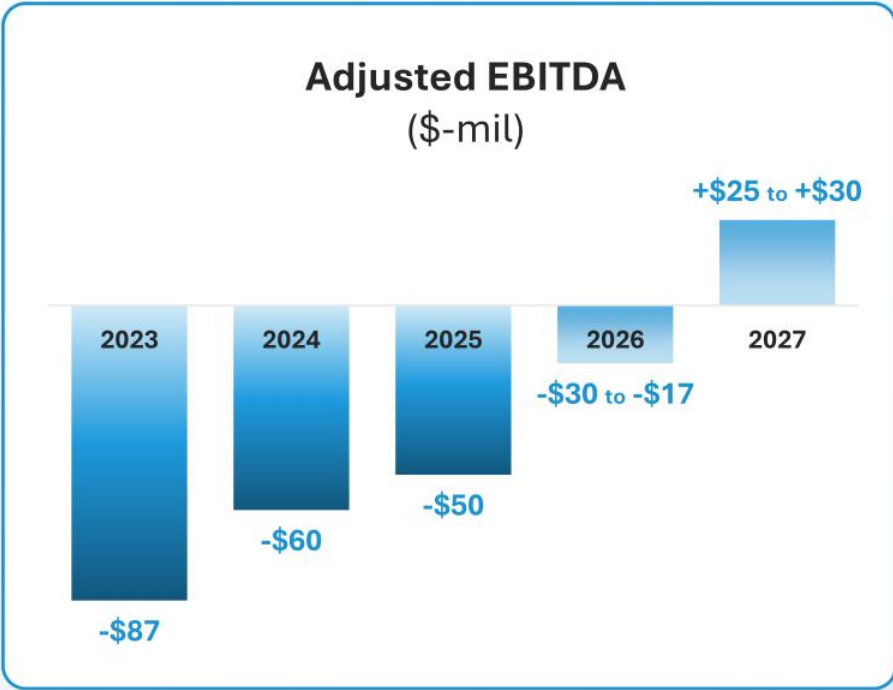
Annual R&D Spend

R&D as % of Sales

Invest in core strategic areas to accelerate innovation & support growth



Once positive, adjusted EBITDA^{1,2} is expected to achieve sustained positive momentum



SG&A Leverage Drivers

- Improved sales force productivity
- Initiate hybrid clinical support model
- SG&A is expected to drive leverage despite increased marketing investments
- 2026 Guidance Assumes positive EBITDA in 4Q26

1. 2026 + 2027 financial guidance issued on February 26, 2026
 2. Adjusted EBITDA is a financial measure that is not prepared in accordance with generally accepted accounting principles in the United States (GAAP). For more information about the Company's use of non-GAAP financial measures, please see appendix

Balance Sheet and Metrics

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Strong balance sheet supports scalable growth



Operating Cash Flow

- Continue to drive working capital improvements
- Meaningful opportunity to reduce Days Inventory on Hand and Day Sales Outstanding
- **Do not expect cash balance to drop below \$175 million***



Capital Spending (PPE)

- Procept requires **little CAPEX investment**
- Existing San Jose headquarters provides meaningful capacity to scale
- **3-4% target:** CAPEX as percent of sales*



Outstanding Debt

- \$52 Million bank loan with extremely attractive rate
- Debt maturity 4Q27
- Procept is positioned to be **cash-flow positive at maturity***

*2026 + 2027 financial guidance issued on February 26, 2026

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Moderated Q&A



THANK YOU

Appendix - Non-GAAP Reconciliations

PROCEPT BioRobotics Corporation
RECONCILIATION OF GAAP NET LOSS TO ADJUSTED EBITDA
(Unaudited, in thousands)

| | Three Months Ended | | Twelve Months Ended | |
|---|--------------------|-------------|---------------------|-------------|
| | December 31, | | December 31, | |
| | 2025 | 2024 | 2025 | 2024 |
| Net loss | \$ (29,845) | \$ (18,856) | \$ (95,572) | \$ (91,413) |
| Depreciation and amortization expense | 1,709 | 1,453 | 6,390 | 5,234 |
| Stock-based compensation expense | 10,842 | 9,085 | 47,603 | 31,840 |
| Interest (income) and interest expense, net | (1,719) | (2,017) | (8,632) | (6,711) |
| Adjusted EBITDA | \$ (19,013) | \$ (10,335) | \$ (50,211) | \$ (61,050) |

PROCEPT BioRobotics Corporation
RECONCILIATION OF GAAP NET LOSS TO ADJUSTED 2026 EBITDA Guidance
(Unaudited, in thousands)

| | For the Year Ending Decem | |
|---|---------------------------|------|
| | 2026 | |
| | LOW | HIGH |
| Net loss | \$ (91,500) | \$ |
| Depreciation and amortization expense | 7,500 | |
| Stock-based compensation expense | 59,000 | |
| Interest (income) and interest expense, net | (5,000) | |
| Adjusted EBITDA | \$ (30,000) | \$ |

PRO

Risk and Safety Information

All surgical treatments have inherent and associated side effects, some of which may lead to serious outcomes and may require intervention. Individual's outcomes may depend on a number of factors, including but not limited to patient characteristics, disease characteristics and/or surgeon behavior. The most common side effects are mild and transient and may include mild pain or difficulty when urinating, discomfort in the pelvis or penis, blood in the urine, inability to empty the bladder or a frequent and/or urgent need to urinate, and bladder or urinary tract infection. Other risks include but are not limited to: anesthesia risk; sexual dysfunction, including ejaculatory or erectile dysfunction; injury to the urethra, such as false passage or stricture, or to the rectum, including rectal incontinence/perforation; bladder or prostate capsule perforation; infection, including the potential transmission of blood borne pathogens; bleeding; incontinence; embolism; electric shock/burn; transurethral resection (TUR) syndrome; bladder neck contracture; bruising. No claim is made that the AquaBeam Robotic System or HYDROS Robotic System will cure any medical condition, or entirely eliminate the diseased entity. Repeat treatment or alternative therapies may sometimes be required.

Rx Only

Prostate Cancer Clinical Trial (WATER IV)

Caution: Aquablation therapy for the treatment of prostate cancer is limited by federal law to investigational use only. The safety and effectiveness of Aquablation therapy for the treatment of prostate cancer has not been established.

PRO

Clinical Data Summary Methodology

Studies with distinct patient cohorts evaluating Aquablation therapy were included, abstracts and unpublished data were excluded; Studies with fewer than 30 patients were excluded; Data was pulled from 18 Aquablation therapy publications up to January 2025; Listed below; Outcomes of interest were aggregated together, and subsequently weighted based on the studies sample size; Not all publications included all outcomes of interest, and thus were excluded from each respective analysis.

India 20-118, n=47, Deesai et al; WATER 30-80ml, n=116, Gilling et al; WATER II 80-150ml, n=101, Bhojani et al; OPEN WATER 20-148ml, n=178, Bach et al; FRANCAIS WA

30-80ml, n=30, Misrai et al; Very Large Prostates, 151-362ml, n=36, Helfand et al; Jacksonville, 27-223ml, n=55, Kasraeian et al Lebanon, 13-148ml, n=59, Labban et al; Sta AUR and CUR, 29-250ml, n=113, Burton et al; Montreal ASC, 41-270ml, n=60, Zorn et al; Mount Sinai, 38-330ml, n=330, Omidele et al; Hamburg, 20-154ml, n=118, Bach et al Japan PMS, 33-242ml, n=103, Hinata et al; Madrid Aquablation vs HoLEP, 72±35ml, n=75, Quintas et al; Israel, 31-138, n=50, Shvero et al; Italy, 43-81, n=109, Amparore et al Focal Bladder Neck Cauteary, 20-263ml, n=2,089, Elterman et al; HoLEP vs Aquablation Hematuria Risk, 56±25ml, n=167, Gloger et al

Clinical Methodology (Slide 44 & 45)

Categories reflect the top patient priorities identified through patient market research commissioned by PROCEPT BioRobotics and conducted by third-party research organizations. Note: IPSS, Qmax and PVR were all included as measures of the patient priority, magnitude of symptom relief.

References

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- Gilling P, et al. WATER trial: Aquablation vs TURP. J Urol. 2018;199:1252-1261.
- Sandhu JS, Bixler BR, Dahm P, et al. Management of lower urinary tract symptoms attributed to benign prostatic hyperplasia (BPH): AUA Guideline amendment 2023. J Urol. 2023;10.1097/JU.0000000000003698.

Category Criteria

TURP, as the standard-of-care, is considered the benchmark for the graphic representation.

Preservation of Continence: rate of persistent urinary incontinence ^{2,3,5,6,11,12}

Durability: the proportion of patients who go on to receive an additional surgical intervention. Real-world-evidence prioritized as source data, with minimum 3-year follow-up data available in third-party national healthcare database. Non-applicable to BPH medications, due to no definitive treatment. Other unrepresented procedures due to unavailability of 3-year RWE.¹

IPSS Symptom Reduction: reduction in International Prostate Symptom Score (IPSS) from baseline. According to AUA guideline, Management of Lower Urinary Tract Symptoms Attributed to Benign Prostatic Hyperplasia (2023)¹⁶ – “There is nearly universal agreement that [alpha blockers] are all relatively equally effective in terms of IPSS improvement, with an expected range of improvement of 5-8 points, compared to an expected effect of placebo from 2-4 points.” Intermediate group identifies reported symptom reduction values above the range of symptom improvement from medications, but not clinically proven to be comparable to TURP. ^{2,3,4,9,10,11,12}

Reduction of Post Void Residual Volume: reduction in post-void residual volume, in mL. ^{2,3,4,9,11,12,13}

Qmax Improvement: peak urinary flow, mL/s ^{2,3,4,9,10,11,12}

Recovery Time and Ease: No single outcome measure to represent post-operative recovery experience is widely available in the literature, or collected using consistent methodology, to apply as the criteria. Therefore, the procedures differentiated from the intermediate group (default) reflect the following:

BPH medications – no post-operative recovery period; Rezum – inferior patient recovery in the CLEAR trial¹⁴; iTind – additional procedure required to retrieve temporary implant; LEP – rates of post-operative incontinence^{2,5,6,7}; Simple prostatectomy – requires invasive surgical access.

Preservation of Erectile Function: erectile dysfunction measured according to International Index of Erectile Function (IIEF) after symptoms from immediate postoperative period expected to resolve. ^{2,5,6,8,9,11,12} Note: TURP as the reference point is categorized as low due to other treatments with lesser (more favorable) reported rates of erectile dysfunction.

Preservation of Ejaculatory Function: ejaculatory dysfunction measured according to Male Sexual Health Questionnaire for Ejaculatory Dysfunction (MSHQ-EJD) at 3 months follow-up. ^{2,5,6,9,11,12} Note: TURP as the reference point is categorized as low due to other treatments with lesser (more favorable) reported rates of ejaculatory dysfunction.

