Beta Bionics

Investor Presentation

September 2026

Disclaimer

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Statements in this presentation that are not statements of historical fact are forward-looking statements. Such forward-looking statements include, without limitation, statements regarding: expectations of Beta Bionics, Inc. (the "Company," "we," "our" or "us") regarding the potential benefits, effectiveness, and safety of the iLet, Mint, 3D Intelligence algorithm and other product candidates; our expectations with regard to the design, results and timing of our research and development efforts, clinical trials and real world use or experience, including the timing and availability of data from such trials and real world use; our regulatory development plans for the iLet, Mint, iLet 3D, Mint 3D and bi-hormonal system and other product candidates, including the timing for potential regulatory approvals; the markets and market opportunities for the iLet, Mint and other product candidates, if approved; the timing, likelihood or success of our business strategy, including commercialization and our multi-channel reimbursement strategy, as well as plans and objectives of management for future operations including market and indication expansion; and our anticipated growth and other measures of future operating results and financial performance including 2026 full year guidance regarding revenue, new patient starts through pharmacy, and gross margin. In some cases, you can identify forward-looking statements by terms such as "expect," "should," "plan," "anticipate," "could," "intend," "target," "project," "believe," "estimates," "potential" or "continue" or the negative of these terms or other similar expressions. These forward-looking statements are based on the beliefs of the management of the Company as well as assumptions made by and information currently available to the Company. These statements involve substantial known and unknown risks, uncertainties and other factors that may cause our actual results, timing of results, levels of activity, performance, or achievements to be materially different from the information expressed or implied by these forward-looking statements.

Other factors that may cause the Company's actual results to differ from current expectations are discussed in the Company's filings with the Securities and Exchange Commission (the "SEC"), including in its Annual Report on Form 10-K for the year ended December 31, 2025, filed with the SEC on February 24, 2026, as updated by its Quarterly Report on Form 10-Q for the quarter ended June 30, 2026, filed with the SEC on July 29, 2026, and other filings that the Company may make from time to time with the SEC the section titled "Risk Factors" in the Company's Form 10-Q for the quarter ended June 30, 2026, filed with the SEC on June 30, 2026. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date this presentation is given. Except as required by law, the Company undertakes no obligation to publicly update any forward-looking statements, whether as a result of new information, future events or otherwise.

We operate in a very competitive and rapidly changing environment, and new risks emerge from time to time. It is not possible for our management to predict all risks, nor can we assess the impact of all factors on our business or the extent to which any factor, or combination of factors, may cause actual results to differ materially and adversely from those anticipated or implied in the forward-looking statements. We may not actually achieve the plans, intentions, or expectations disclosed in our forward-looking statements, and you should not place undue reliance on our forward-looking statements. Actual results or events could differ materially from the plans, intentions and expectations disclosed in the forward-looking statements we make. The forward-looking statements in this presentation represent our views as of the date of this presentation. We anticipate that subsequent events and developments will cause our views to change. However, while we may elect to update these forward-looking statements at some point in the future, we have no current intention of doing so except to the extent required by applicable law. Except as required by law, neither we nor any other person assumes responsibility for the accuracy and completeness of the forward-looking statements in this presentation and the accompanying oral commentary. You should, therefore, not rely on these forward-looking statements as representing our views as of any date subsequent to the date of this presentation.

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This presentation discusses certain product candidates that have not yet received marketing authorization or clearance by the U.S. Food and Drug Administration. No representation is made as to the safety, effectiveness or likelihood of marketing authorization or clearance of these product candidates.

In addition to the financial information presented in accordance with U.S. generally accepted accounting principles (GAAP), this presentation and accompanying oral presentation includes certain non-GAAP financial measures. These measures are presented for supplemental informational purposes only, have limitations as analytical tools and should not be considered in isolation or as a substitute for financial information presented in accordance with GAAP. Our calculations of each of these measures may differ from the calculations of measures with the same or similar titles by other companies and therefore comparability may be limited. Because of these limitations, when evaluating our performance, you should consider each of these non-GAAP financial measures alongside other financial performance measures, including the most directly comparable financial measure calculated in accordance with GAAP and our other GAAP results. A reconciliation of non-GAAP measures to the most directly comparable GAAP measures are included in the Appendix to this presentation and can be found in our filings with the SEC.

We have not provided the forward-looking GAAP equivalents for certain forward-looking non-GAAP measures presented in this presentation or the accompanying oral presentation, or a GAAP reconciliation, as a result of the uncertainty regarding, and the potential variability of, reconciling items such as stock-based compensation expense. Accordingly, a reconciliation of these non-GAAP guidance metrics to their corresponding forward-looking GAAP equivalents is not available without unreasonable effort. However, it is important to note that material changes to reconciling items could have a significant effect on future GAAP results.

Our mission is to **simplify and alleviate** the burden of managing diabetes with **life-changing solutions.**

Beta Bionics

Market-Leading Innovators

Beta Bionics has the only fully adaptive insulin-dosing algorithm, and most advanced system available in the U.S.[1]

Key Operating Highlights

$32.0M
Q2'26
Net Sales

59.0%
Q2'26
Gross Margin

69%
New Patient Starts from MDI in Q2'26

High 30s%
New Patient Starts reimbursed through pharmacy in Q2'26

Q1'27
Targeted Mint™ patch pump full commercial launch in the U.S.

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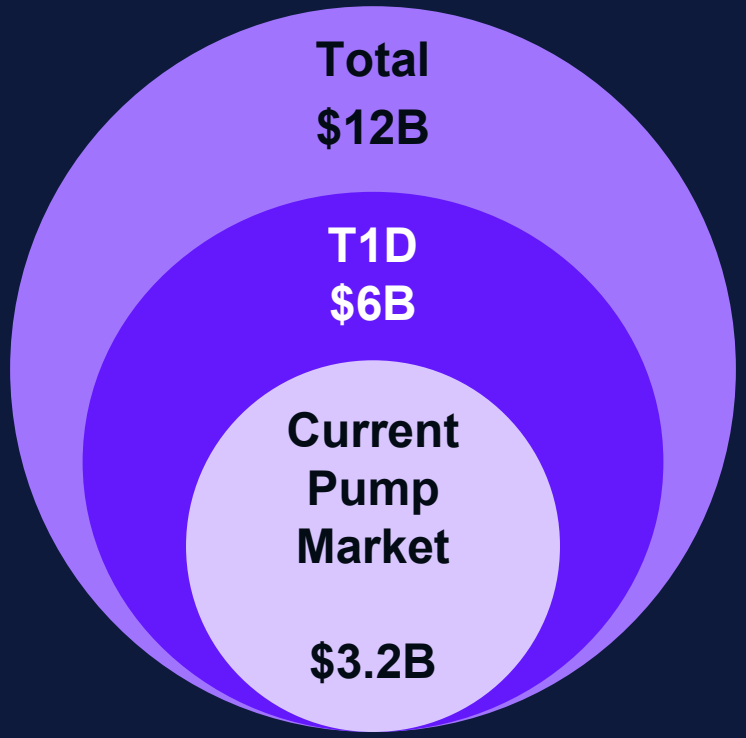
Note: Data as of Q2 2026. Multiple Daily Injections (MDI). Year End (YE).

1. Cook, K., et al. (2025). Defining the Automated Insulin Delivery Continuum: User Engagement and System Adaptation. ADCES 2025. Poster Presentation, 2025 Aug 9

Large & Underpenetrated Insulin Pump Market

Total Addressable Market for Insulin Delivery in the U.S.



Type 1 Diabetes

1.9M
People in the U.S.

~1.2M
On multiple daily injections (MDI)

>0.7M
On pump therapy

~39%
Pump penetration

Type 2 Diabetes (Insulin-Intensive)

1.9M
People in the U.S.

>1.7M
On multiple daily injections (MDI)

<0.2M
On pump therapy

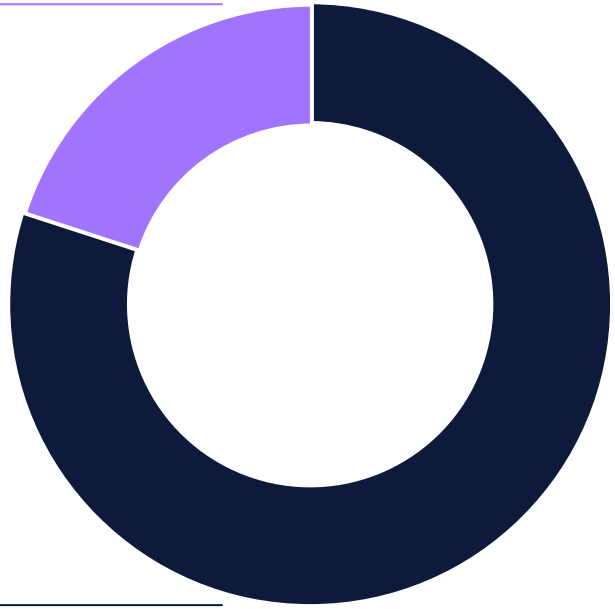
<10%
Pump penetration

Source: Internal estimates informed by data from the CDC, Seagrove Partners, and publicly-reported 2025 product sales data.
Note: Type 1 Diabetes (T1D).

Poor Clinical Outcomes for People with T1D in the U.S.

~80% not at treatment goal, defined by the ADA as HbA1c <7%

~20%
of people with T1D are **at ADA goal**, but **many are overburdened** with their diabetes management



~80%
of people with T1D are **not at ADA goal**[1,2]

~380K
People with T1D

- Meeting goal & content
- Meeting goal & overburdened

~1.5M
People with T1D

- Not meeting goal & overburdened

Source: Internal estimates informed by data from the CDC, Seagrove Partners, and publicly-reported 2025 product sales data.
Note: Type 1 Diabetes (T1D). American Diabetes Association (ADA).
1. Foster et al. Diabetes Technol Ther. 2019;21:66-72.
2. Pettus et al., Diabetes Care. 2019;42:2220–2227.

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Significant Unmet Market Needs Remain

Pump complexity is a major obstacle to adoption

Complexity of Pumps

User

Adjusts everyday as needed

Carb Counting for Meal Doses

Manual Correction Doses

Temporary Basal Rate

Physician

Adjusts every 3 months

Insulin-to-Carb Ratio

Correction Factor

Basal Rate

Obstacles to Pump Use

- Burdensome for users to manage everyday
- Difficult for physicians to setup and manage, particularly primary care providers
- Costly for users to purchase pump and monthly supplies

T1D Unmet Needs

~80%

Not at ADA treatment goal[1,2]

~39%

Insulin pump therapy penetration

Penetration remains low and treatment outcomes are poor

Source: Internal estimates informed by data from the CDC, Seagrove Partners, and publicly-reported 2025 product sales data.
Note: Type 1 Diabetes (T1D). Type 2 Diabetes (T2D), American Diabetes Association (ADA).

1. Foster et al. Diabetes Technol Ther. 2019;21:66-72.
2. Pettus et al., Diabetes Care. 2019;42:2220–2227.

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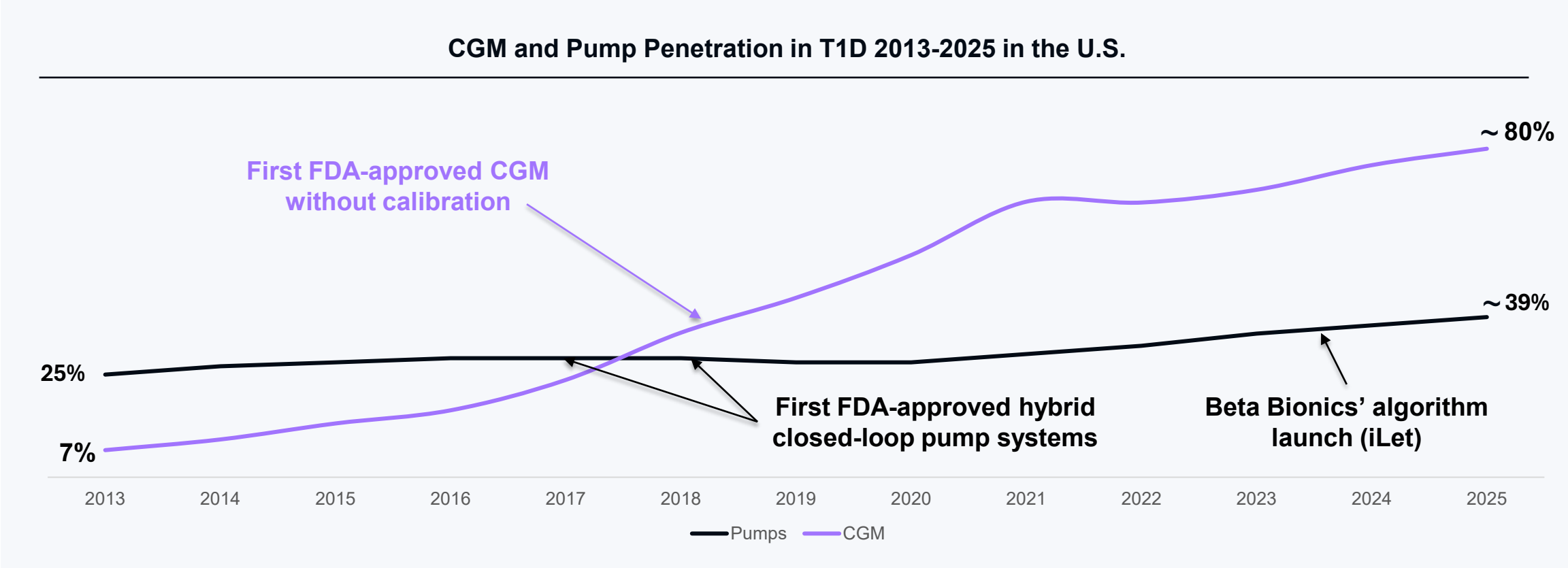
The Market Responds to Major Ease-of-Use Innovations

No fingerstick calibration was the key innovation for CGMs; Beta Bionics' fully adaptive insulin-dosing algorithm may be that innovation for pumps



Source: Internal estimates informed by data from the CDC, Seagrove Partners, and publicly-reported 2025 product sales data.
Note: Continuous Glucose Monitor (CGM), U.S. Food & Drug Administration (FDA), Type 1 Diabetes (T1D).

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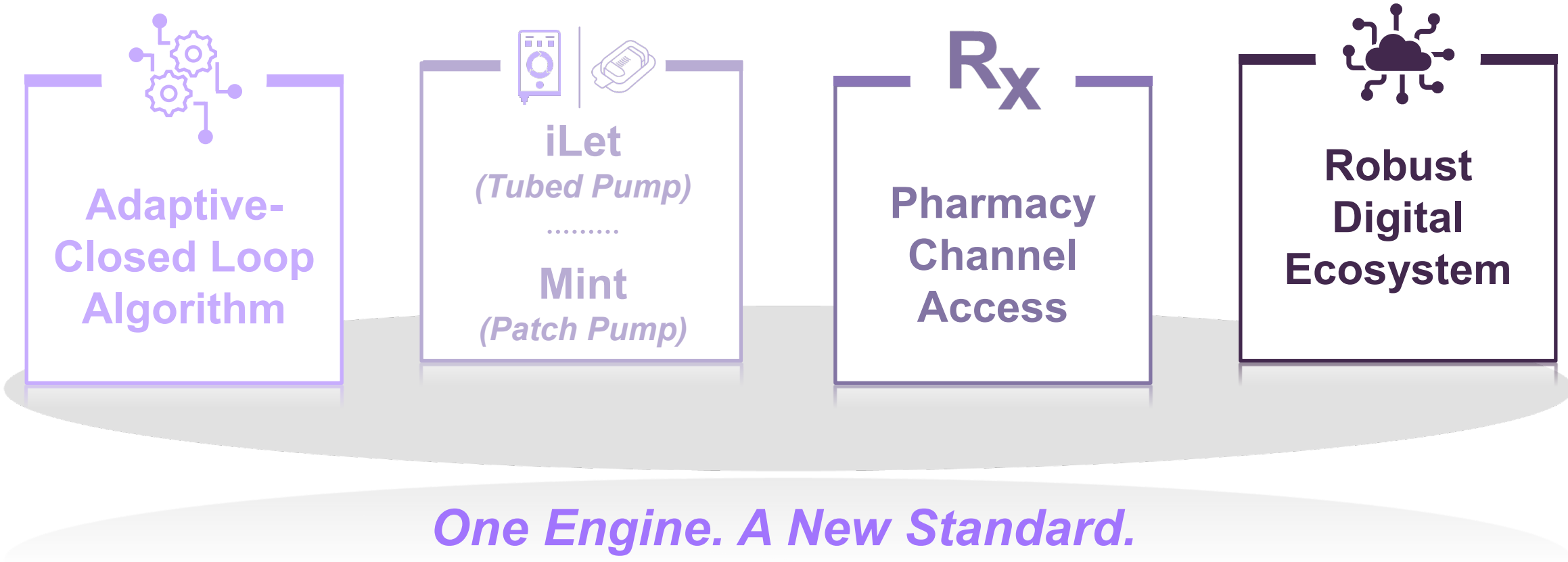
We believe we can do more for people with diabetes.

Most Advanced Pump Platform in the U.S.[1]



One Engine. A New Standard.

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Note: Mint is not yet available for sale.

1. Cook, K., et al. (2025). Defining the Automated Insulin Delivery Continuum: User Engagement and System Adaptation. ADCES 2025. Poster Presentation, 2025 Aug 9

First and Only Fully Adaptive Closed-Loop Algorithm

Generates similar or better clinical outcomes, with substantially less work for the user and provider[1,2]

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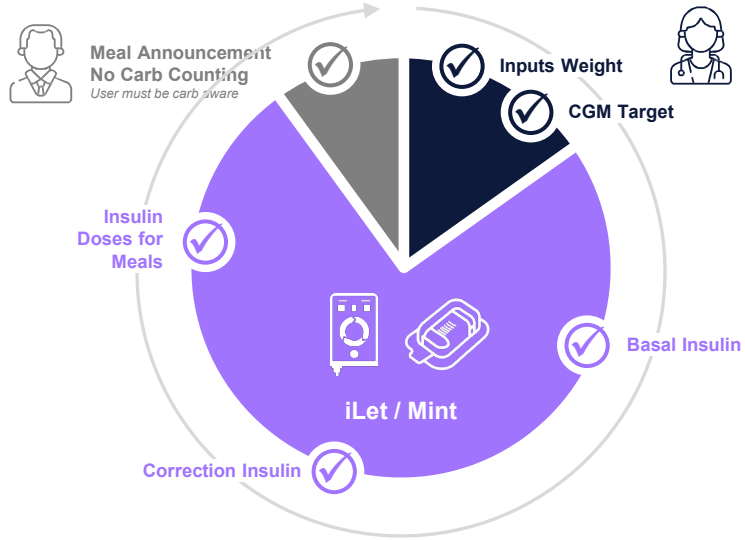
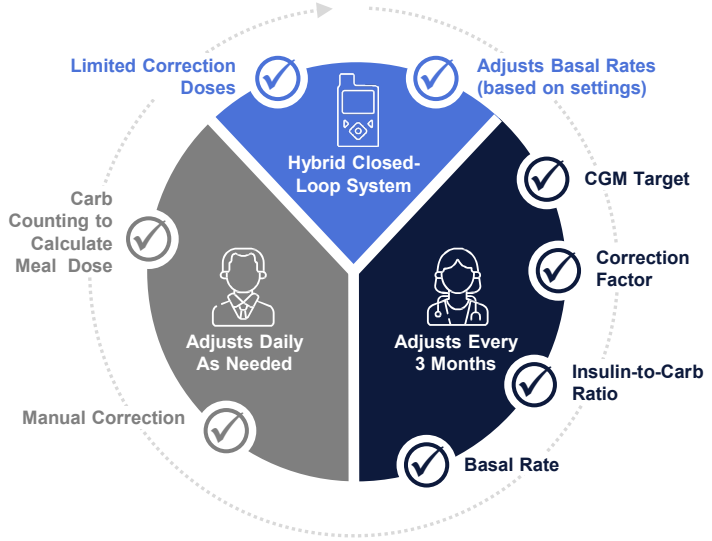
Adaptive Closed-Loop Algorithms



- **Initialized with only the user's body weight**
- **No adjustment of settings** by the provider required after initialization
- **User only announces meals;** no carb counting*
- **Algorithm learns and adapts** to the user's insulin requirements

Pump Competitors

Hybrid Closed-Loop System



- **Initialized with numerous inputs** determined by the provider
- **Requires quarterly adjustment of settings** by the provider
- **User continuously makes daily decisions** about their diabetes
- **Algorithm responds** to ongoing inputs from the user and provider

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Note: Continuous glucose monitor (CGM). *User must be carb aware

1. Russell, S.J., et al. (2022) A Multicenter Randomized Trial of a Bionic Pancreas in Type 1 Diabetes. The New England Journal of Medicine. 387:1161–1172.
2. Damiano et al. (2024) Glycemic and CGM Outcomes are nearly independent of the frequency of user engagement with the iLet in the Bionic Pancreas Pivotal Trial and real-world setting. https://www.betabionics.com/wp-content/uploads/iLet-Glycemic-and-CGM-outcomes-vs-user-engagement.pdf

3D Intelligence™, Beta Bionics' Next-Gen Algorithm

Three selectable algorithm experiences offer greater personalization while maintaining the core simplicity and automation of the current Beta Bionics algorithm

Original

The currently available, fully adaptive algorithm*
Proven real-world experience across more than 40,000 users

Optimized

New algorithm experience with enhanced learning
Builds on and incorporates real-world learnings from the currently available, fully adaptive algorithm to optimize performance

Conservative

New, alternative algorithm experience
Designed to provide a more measured response when correcting high glucose levels

iLet 3D™

Expected commercial launch by the end of 2026

Mint 3D™

Expected commercial launch in Q1 2027

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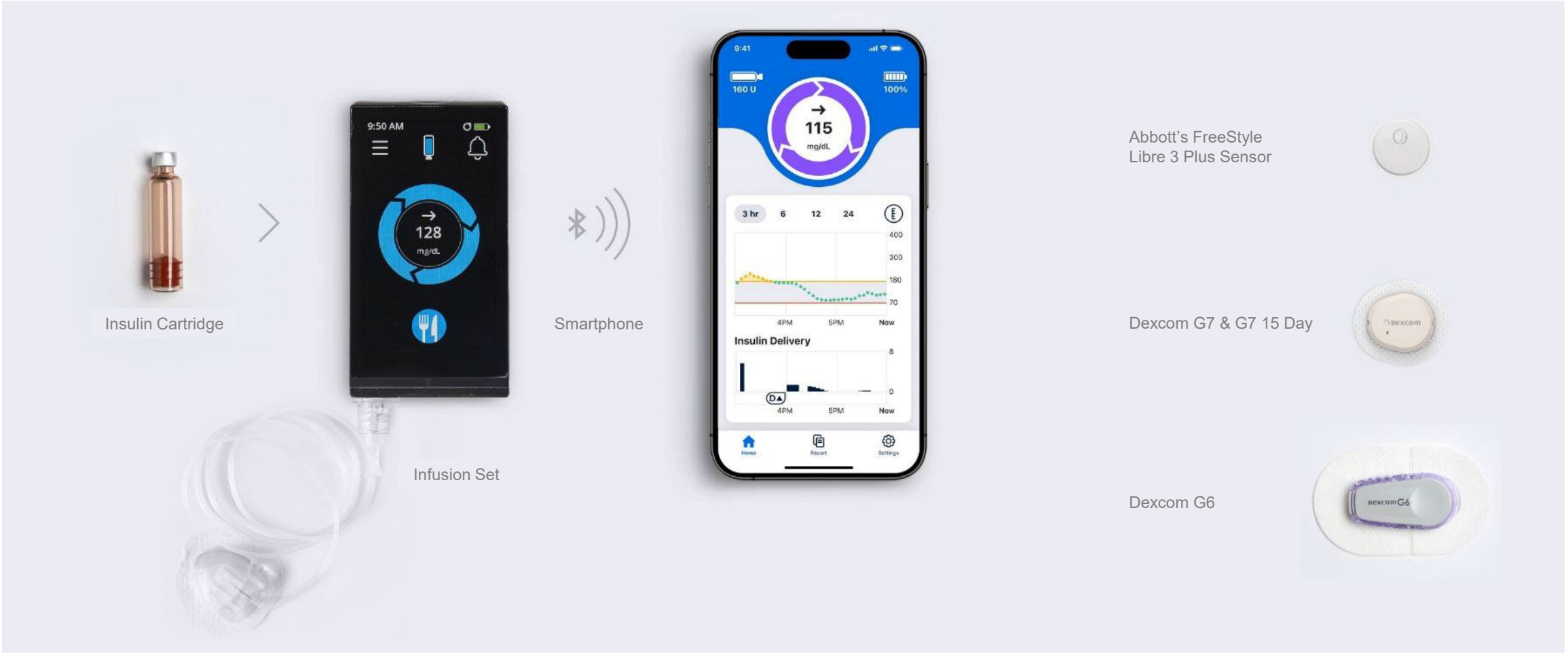
*Minor differences in adaptation compared to previous algorithm are not anticipated to have a significant impact on outcomes.
Note: Commercialization of iLet 3D and Mint 3D is subject to regulatory clearance of 3D Intelligence by the FDA, and the devices are not available for sale. Actual product features may vary.

iLet Bionic Pancreas

Sleek tubed pump design with control screen on pump and option to use pre-filled Fiasp PumpCart



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Mint Patch Pump

FDA Cleared in September 2026; Full Commercial Launch in the U.S. expected in Q1 2027

New, innovative reusable-disposable architecture

Reusable Controller

- Designed for 2 years of use
- iOS and Android smartphone controlled
- Never requires charging
- Firmware-over-the-air updates
- Waterproof for up to 5ft deep for 60 minutes (IPX8)

Disposable Cartridge

- 200U insulin reservoir
- 3-day wear time; additional 12-hour grace period
- Disposable batteries
- No phone required for patch change
- Waterproof for up to 5ft deep for 60 minutes (IPX8)

Powered by fully adaptive algorithm

Seamless patch change process

Compatible with industry-leading CGMs & smartphones

Note: The images shown are for illustrative purposes only. Actual product specifications may vary. Mint is not yet available for sale. Continuous glucose monitor (CGM).

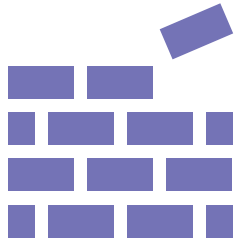
Targeting Mint Full U.S. Commercialization in Q1 2027

mint™ by Beta Bionics

Potential Business Advantages

Expands Addressable Market

Reimbursed Exclusively in the Pharmacy Channel



Cost Effective to Manufacture at Scale

Limits Environmental Waste

Note: Mint is not yet available for sale.

Providing Flexible Reimbursement

We believe iLet is the first durable pump to access the pharmacy channel; Mint will be exclusively available in the pharmacy channel

	DME (Medical)	Pharmacy
	Traditional reimbursement channel for durable insulin pumps	*Pay-as-you-go model with advantages for users, providers, payors, and Beta Bionics*
Illustrative Economics for User and Payor		
User (out of pocket)		
Upfront cost	~$500 to ~$2,000	~$0 to ~$100
Monthly cost	~$35	~$25 or less
Commitment	Typically 4 years	None
Payor		
Pump	~$3,500 up front	~$450 / month
Supplies	~$70 / month	

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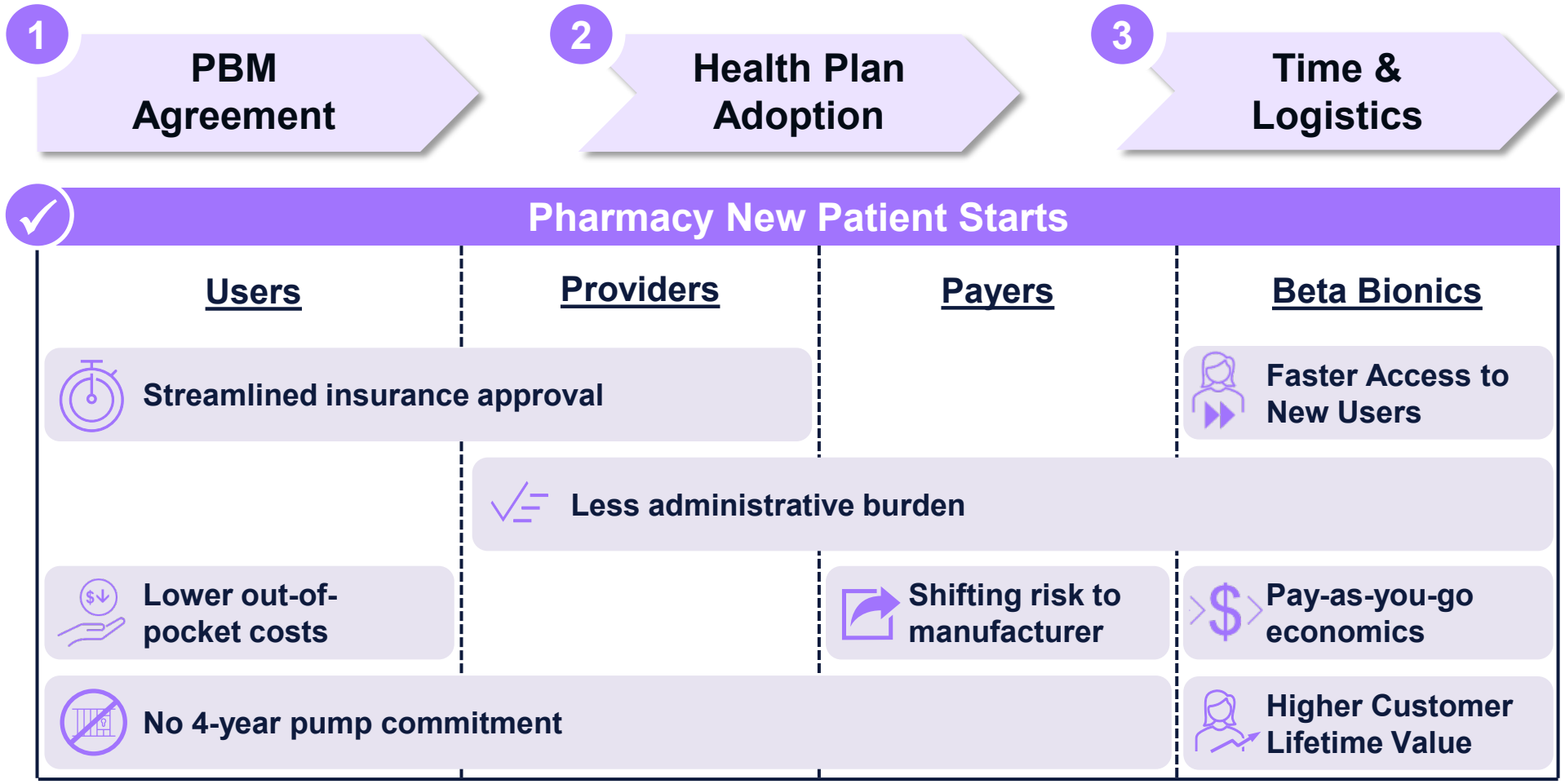
Note: The information presented on this slide is for illustrative purposes only and expresses estimated dollar amounts. Durable Medical Equipment (DME).

Pharmacy Channel Strategy

Three-step process delivers benefits to users, healthcare providers, payers, and Beta Bionics



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Note: Pharmacy Benefit Manager (PBM)

Robust Digital Ecosystem

For our users, their caregivers, and their healthcare providers

iLet and Mint Apps *(iOS & Android)*

- Easy-to-use interface
- Automatic data uploads to the cloud

Bionic Circle App *(iOS & Android)*

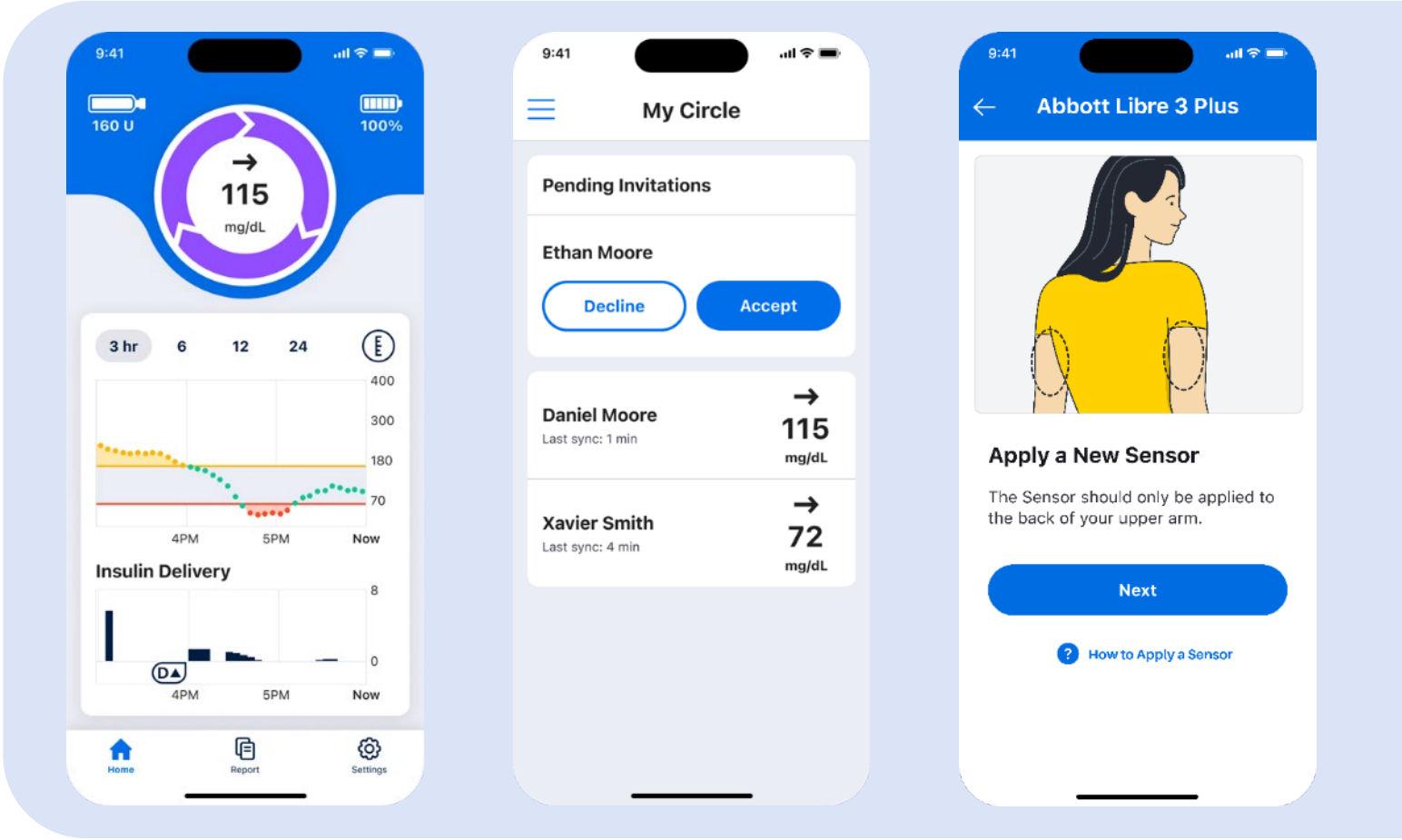
- Remote monitoring experience for family members, friends, and caregivers

Bionic Portal, Report & Insights

- Web-based portal for healthcare providers with patient reports and actionable insights

CGM Compatibility

- iLet and Mint are both compatible with Dexcom G7, Dexcom G7 15 Day, and Abbott's FreeStyle Libre 3 Plus



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Positioning for success

Differentiation Where it Matters Most

		Fully Adaptive Algorithm	Tubeless Hybrid Closed-loop System	Tubed Hybrid Closed-loop System	Smart Insulin Pens
User Decision-Making Framework		Beta Bionics	Insulet[2]	MiniMed and Tandem[3,4]	Multiple
Algorithm(s)	Algorithm(s) initialized with only body weight	✓	✗	✗	✗
	Algorithm(s) determines 100% of insulin doses	✓	✗	✗	✗
	No carb counting[5]	✓	✗	✗	✗
Form Factor	Tubed / durable pump	✓	✗	✓	✗
	Prefilled cartridges (for tubed pump)	✓	✗	✗	✗
	Patch pump (FDA-cleared)	✓	✓	✗	✗
CGM Integration	Dexcom G7 & G7 15 Day	✓	✓	✓ *(Tandem only)*	✗
	Abbott Freestyle Libre 3 Plus	✓	✓	✓	✗
Reimbursement	DME (medical) channel	✓	✗	✓	✗
	Pay-as-you-go pharmacy channel	✓	✓	✓	✓

Beta Bionics

Note: Continuous Glucose Monitor (CGM), Durable medical equipment (DME). Mint is not yet available for sale.

1. Beta Bionics. iLet Bionic Pancreas User Guide. 2022.
2. Insulet. Omnipod 5 User Guide. 2022.
3. MiniMed 780g System User Guide. 2020.
4. Tandem Diabetes Care. t:slimx2 Insulin Pump with Control IQ Technology User Guide. 2023.
5. User must be carb aware.

Superior Pivotal Trial Outcomes vs. Standard of Care

HbA1c % Improvement with iLet[1]		HbA1c % Improvement with SoC[1]
- 0.6	All Users	- 0.0
- 0.7	Adults	- 0.1
- 0.6	Children	- 0.0
- 0.9	At least 7% Baseline HbA1c	- 0.1
- 1.0	Multiple Daily Injections (MDI)	- 0.0
- 0.5	Pumps	- 0.0
- 0.3	Hybrid Closed-Loop[2]	- 0.1

iLet delivered greater reduction in HbA1c vs. competitive hybrid-closed loop systems[3]

Source: Russell, S.J., et al. (2022) A Multicenter Randomized Trial of a Bionic Pancreas in Type 1 Diabetes. The New England Journal of Medicine. 387:1161–1172., Data on File.
Note: Standard of Care (SoC).
1. Data reflects percentage point improvements in HbA1c measurements at 13 weeks as compared to baseline.
2. Hybrid-closed loop systems included t:slim X2 with Control-IQ, and MiniMed 670G or 770G
3. Analyses of subgroups represent post hoc analyses of the pivotal trial data. The pivotal trial was not designed to evaluate the effect of the iLet Bionic Pancreas in the listed subgroups.

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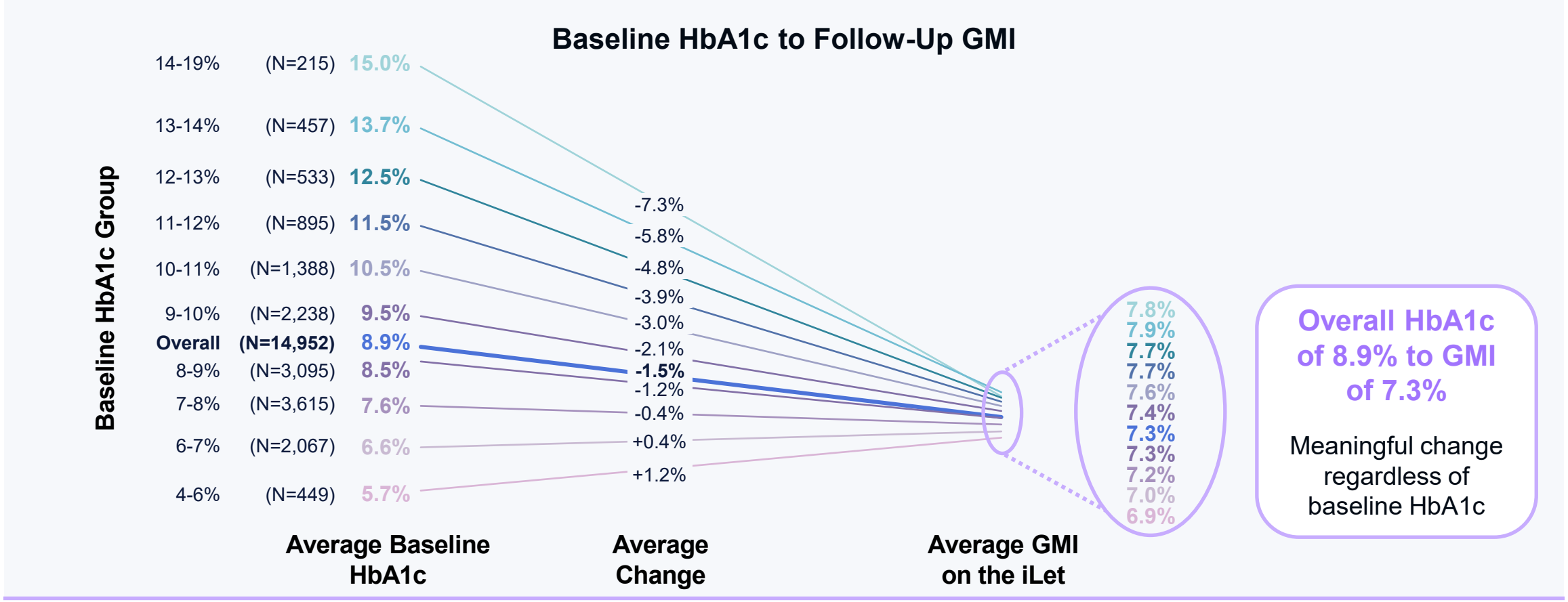
Real World Results Are Consistent with the Pivotal Trial



>84% of users meeting ADA hypoglycemia goals of <4% time below 70 mg/dL and <1% time below 54 mg/dL[1]

Source: The postmarket surveillance data presented is observational data. These data do not come from a prospective study designed to evaluate the effect of the iLet Bionic Pancreas in subgroups by baseline HbA1c. Real-world data comes from the first two years (5/19/2023–5/18/2025) of iLet usage among iLet users with baseline HbA1c data and who uploaded data to Beta Bionics, Inc., including users who do not meet all criteria in the Indications for use. Individual user results may vary from the average values shown here.

Note: Glucose Management Indicator (GMI). American Diabetes Association (ADA).

1. American Diabetes Association Professional Practice Committee. 6. Glycemic Goals and Hypoglycemia: Standards of Care in Diabetes-2024. Diabetes Care. 2024;47:S111-S125

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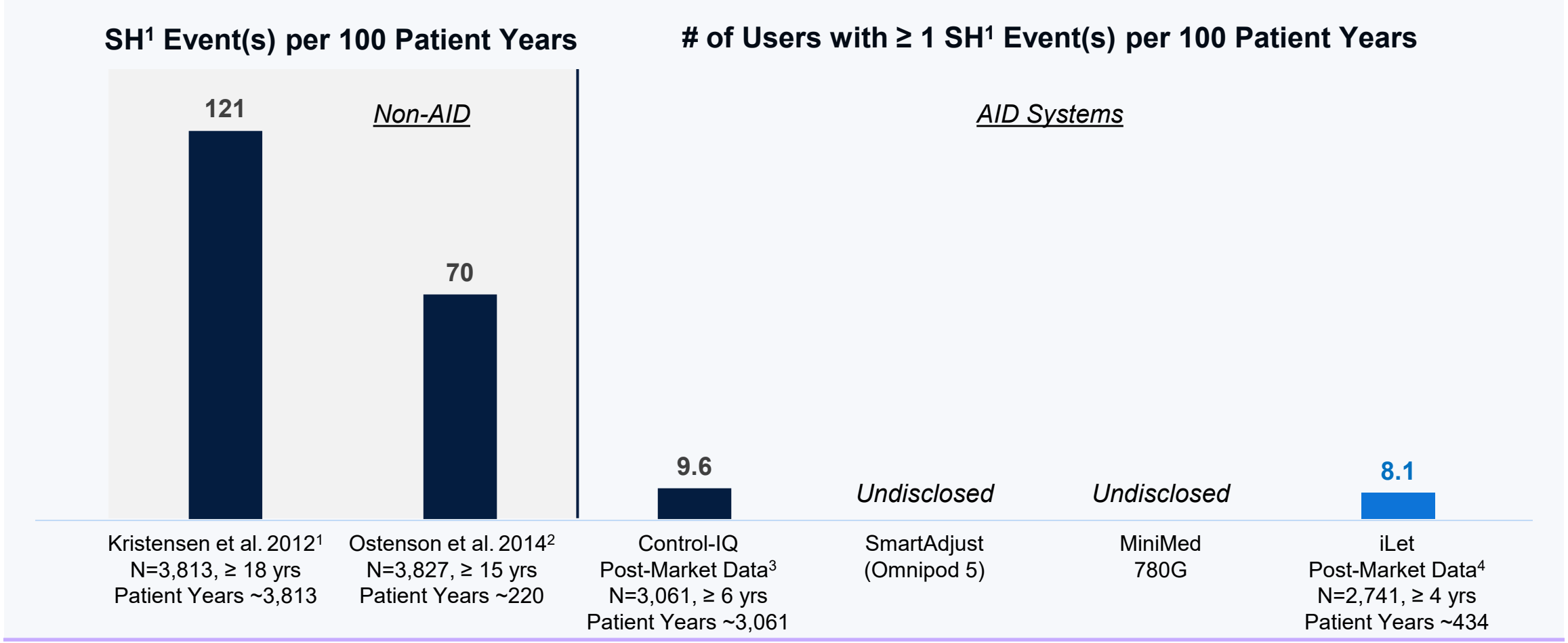
Real World Severe Hypoglycemia in Non-AID vs. AID Systems



Note: Severe Hypoglycemia (SH) defined as hypoglycemia with cognitive impairment requiring the assistance of a third party for treatment. Automated Insulin Delivery (AID).
Data presented are not based on head-to-head studies. As a result, cross-study comparisons cannot be made, and caution is advised when attempting to extrapolate these data to new patients.

1. Kristensen et al. (2012) Insulin analogues and severe hypoglycaemia in type 1 diabetes. Diabetes Res Clin Pract; 96:17-23.
2. Ostenson et al. (2014) Self-reported non-severe hypoglycaemic events in Europe. Diabetic Medicine; 31:92-101.
3. Graham et al. (2024) Real-World Use of Control-IQ Technology Is Associated with a Lower Rate of Severe Hypoglycemia and Diabetic Ketoacidosis Than Historical Data: Results of the Control-IQ Observational (CLIO) Prospective Study. Diabetes Technol Therapy; 26: 24-32.
4. Data on file. The iLet Bionic Pancreas is indicated for people with type 1 diabetes ages 6 years and older.

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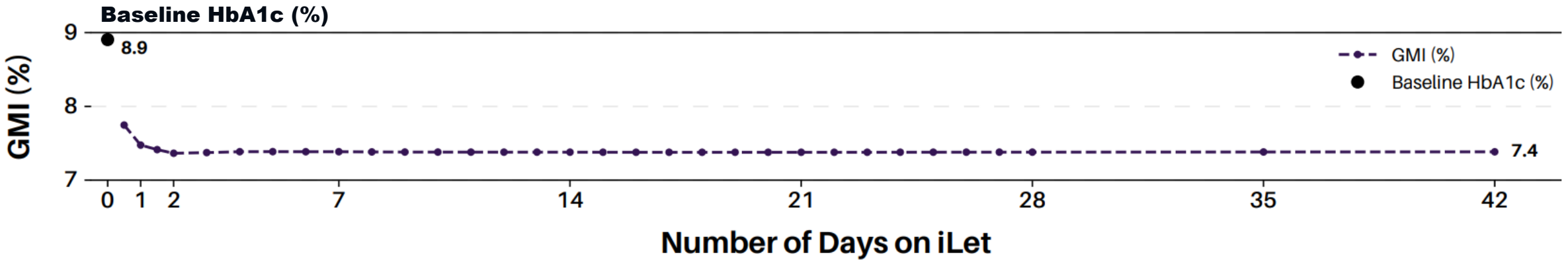
Beta Bionics' Adaptive Algorithm "Warms-Up" in ~2 Days

Time to Steady-State GMI

(N=30,077 sessions[1])



Average baseline HbA1c of 8.9% to follow-up GMI on iLet of 7.4%
<u>just 2 days after initiation</u>

Source: The real-world data presented are observational data. These data do not come from a prospective study designed to assess safety and efficacy of the iLet by duration of iLet use. Data comes from the first 2 years (5/18/2023-5/19/2025) of real-world iLet usage among those people who uploaded CGM data to Beta Bionics, Inc., including users who do not meet all criteria in the Indications for use. Ninety percent of users had a baseline HbA1c available. Individual user results may vary from the average values shown above.

Note: Glucose Management Indicator (GMI). Continuous Glucose Monitor (CGM).

1. Most iLets had only one session. A session reflects each initial use of an iLet or after a factory reset.

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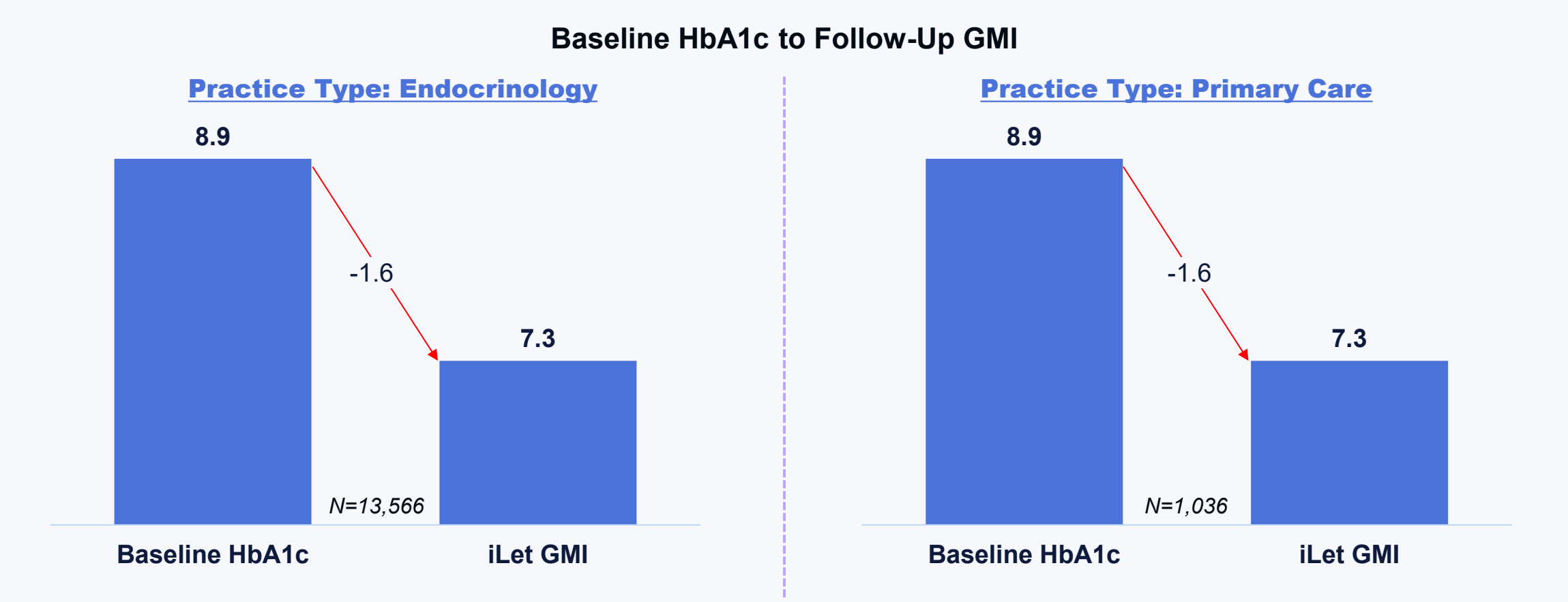
Similar Improvement in GMI Regardless of Provider Type



Similar improvement in GMI with >84% of users meeting ADA hypoglycemia goals, regardless of provider practice type

Source: The real-world data presented is observational data. These data do not come from a prospective study designed to evaluate the effect of the iLet Bionic Pancreas in subgroups by provider practice type. Real-world data comes from the first two years (5/19/2023–5/18/2025) of iLet usage among iLet users with baseline HbA1c data and provider practice type data from Symphony Health Solutions and the Centers for Medicare & Medicaid Services' National Plan and Provider Enumeration System, and who uploaded data to Beta Bionics, Inc., including users who do not meet all criteria in the Indications for use. Individual user results may vary from the average values shown here.

Note: Glucose Management Indicator (GMI). Primary Care Provider (PCP). American Diabetes Association (ADA).

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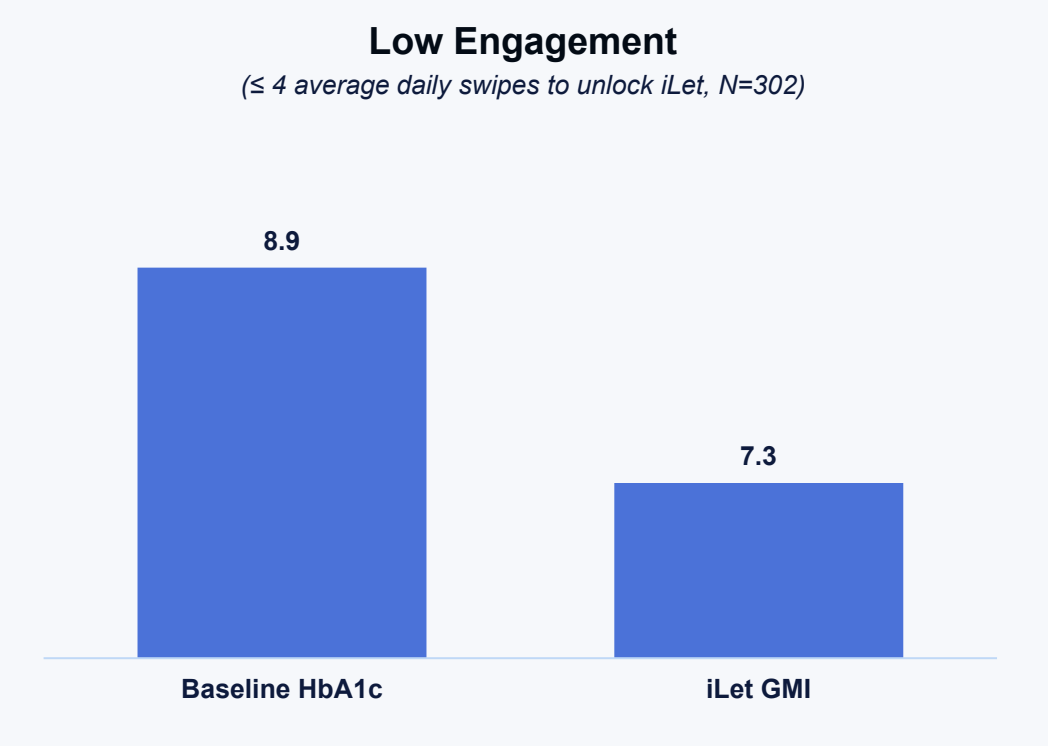
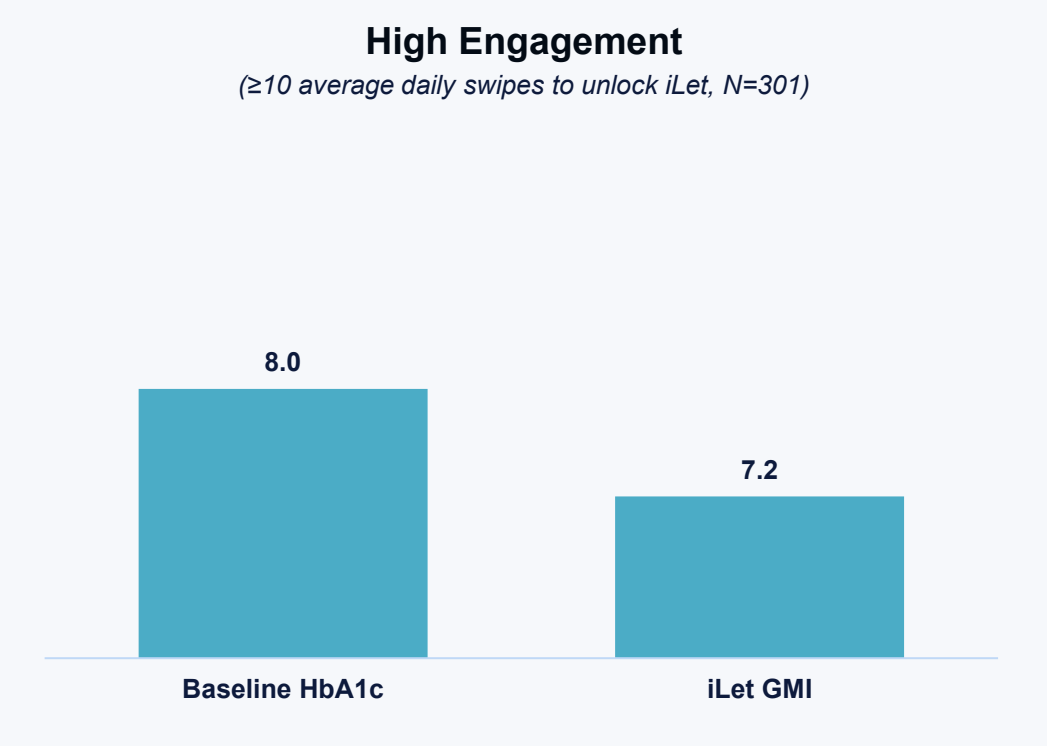
Clinical Results Are Independent of User Engagement

Historically, low engagement with pumps meant poor outcomes – not with iLet

iLet Real-world Data: Change from Baseline HbA1c to Mean GMI by Level of Engagement in Adults (≥ 18 years old)





Source: The real-world data presented is observational data. These data do not come from a prospective study designed to evaluate the effect of the iLet Bionic Pancreas in subgroups by baseline HbA1c or user engagement. Real-world data comes from the first year (5/19/2023–5/18/2024) of iLet usage among those people with baseline HbA1c data and who uploaded data to Beta Bionics, Inc., including users who do not meet all criteria in the Indications for Use. Individual user results may vary from the average values shown here.

Note: Glucose Management Indicator (GMI).

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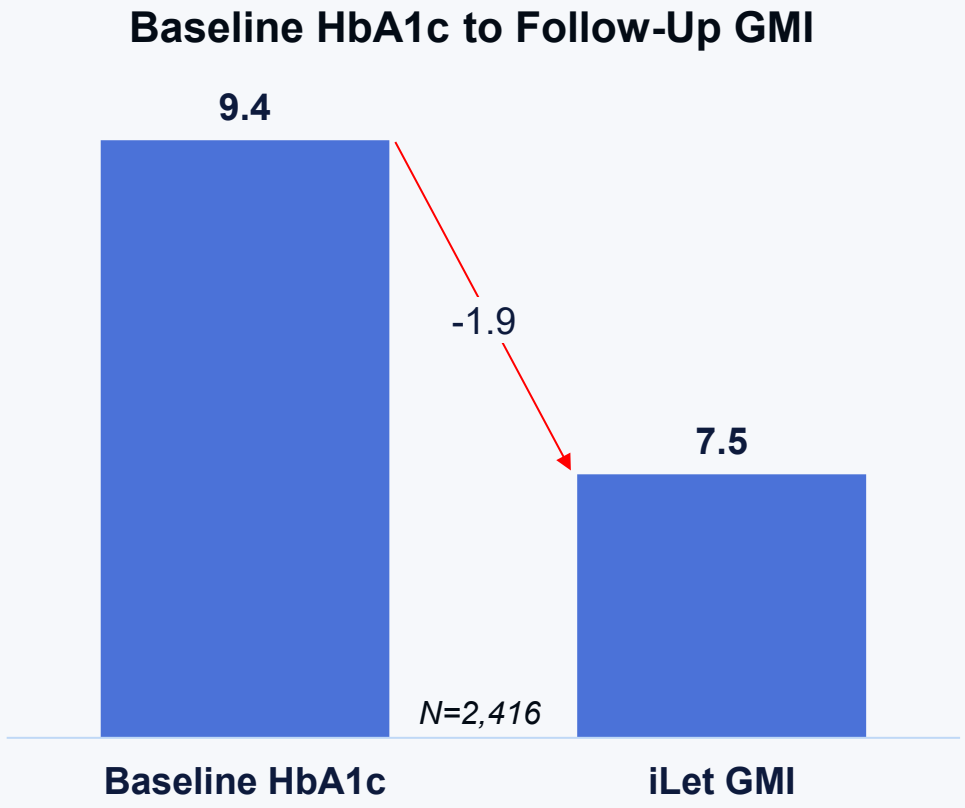
Use of iLet in a “Fully-Closed Loop” Manner



- **~16% of iLet users** (with available data) **use the pump in a “fully-closed loop” manner**[1]
- On average, **“fully-closed loop” iLet users announced ~1 meal every 3 days**
- “Fully-closed loop” users achieved **~62% time in range, on average**
- ~83% of “fully-closed loop” iLet users meeting ADA hypoglycemia goals
- **530 iLet users have not announced a single meal** over the past 21 days
  - ❑ **Improvement from baseline A1c of 9.4% to a follow-up GMI of 7.4%**
  - ❑ ~86% meeting ADA hypoglycemia goals

Source: The real-world data presented is observational data. These data do not come from a prospective study designed to evaluate the effect of the iLet Bionic Pancreas in subgroups by frequency of meal announcements. Real-world data comes from the 21-day period ending 5/18/2025 of iLet usage among those people with baseline HbA1c data and who uploaded data to Beta Bionics, Inc., including users who do not meet all criteria in the Indications for use. Individual user results may vary from the average values shown here.

Note: Glucose Management Indicator (GMI). American Diabetes Association (ADA).

1. Fully-closed loop iLet users defined as users who announced <1 meal per day on average over the 21-day period ending May 18, 2025

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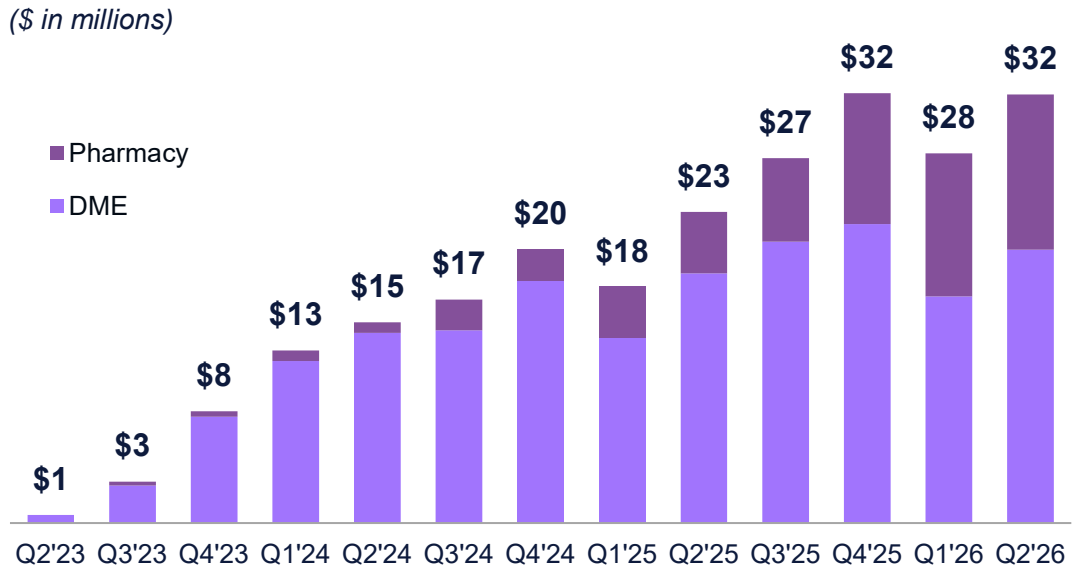
Where we are going next

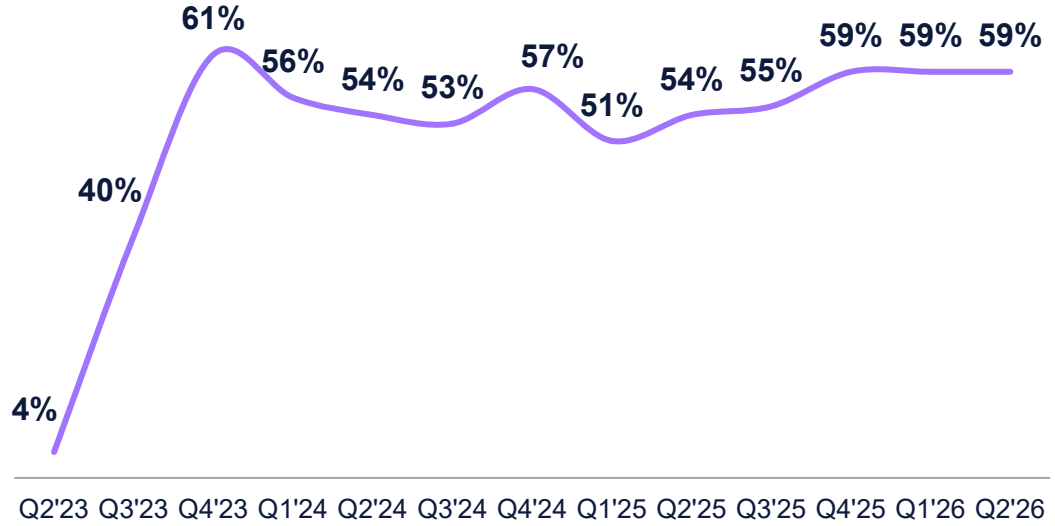
Build On iLet's Highly Successful Commercial Launch

Launched in May 2023, iLet is expanding the pump market and taking market share

Revenue



Gross Margin



High 30s %

New patient starts reimbursed through pharmacy in Q2'26

~69%

New patient starts coming from multiple daily injections in Q2'26

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Beta Bionics Innovation Pipeline

Innovations designed to expand our addressable market and enhance the user experience

Next-Gen Algorithm: 3D Intelligence

- Designed with three algorithm experiences (Original, Optimized, Conservative) to enable a more customizable treatment approach
- 510(k) application submitted to FDA
- Expect to commercialize in the U.S. by the end of 2026 as iLet 3D[1] and in Q1 2027 as Mint 3D[1]

Type 2 Diabetes (T2D) Indication Expansion

- Pivotal trial enrollment initiated in July 2026
- Expect to expand the algorithm's indications for use to adults with T2D in the U.S. **around mid-2027[1]**
- Expanded algorithm indication[1] is expected to enable sales of both iLet and Mint to adults with T2D in the U.S.[2]

Fully-Closed Loop (FCL) Algorithm

- Intend to initiate a fully-closed loop pivotal trial in adults with type 1 and type 2 diabetes in the coming quarters

Bihormonal System

- Completed first-in-human Phase 2a feasibility trial in Q4'25
- Additional Phase 2a feasibility trial initiated in Q1'26
- Path through Phase 2b and Phase 3 trials to support NDA (glucagon) and 510(k) (algorithm and pump) filings

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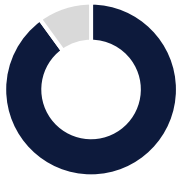
Note: U.S. Food & Drug Administration (FDA). Investigational Device Exemption (IDE), New Drug Application (NDA).
1. Subject to regulatory clearance
2. The Company estimates that greater than 30% of its new patient starts in Q2'26 were from off-label use by T2D patients.

Expansion in Primary Care & Type 2 Diabetes

Largely untapped segments of the diabetes market that iLet may be uniquely positioned to address

Primary Care

~50%
Of the 1.9M people with T1D in the U.S. are treated by PCPs



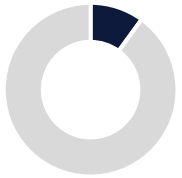
~90%
Of people with T2D in the U.S. are treated by PCPs

<10%
Pump penetration in primary care in the U.S.

Type 2 Diabetes (IIT)[1]

1.9M
People with T2D in the U.S. who require intensive insulin (IIT)

<0.2M
People with T2D IIT in the U.S. use an insulin pump



<10%
Pump penetration in T2D IIT population in the U.S.

Our Path to Win

- **Simplicity combined with efficacy**
- **Similar glycemic outcomes relative to usual care for PCPs**
- **Recently approved patch form factor**

Our Approach

- **Pharmacy reimbursement** reduces administrative burden for PCPs and streamlines approval process for people with T2D
- Label expansion in T2D expected in mid-2027, subject to regulatory clearance
- **Change institution-level protocols that drive PCP prescribing patterns** at the clinic level

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Source: Internal estimates informed by data from the CDC, Seagrove Partners, The Diabetes Forum: 2021 Diabetes Blue Book, and publicly-reported 2024 product sales data.
Note: Type 1 Diabetes (T1D). Type 2 Diabetes (T2D), Insulin-Intensive Therapy (IIT), Primary Care Provider (PCP).
1. Oser SM et al. Diabetes 2024; 73(Supplement_1):957-P.

Bihormonal Pump in Development

Potential to revolutionize diabetes care

First-of-its-kind bihormonal configuration of the iLet

- Intended to utilize adaptive closed-loop algorithms that autonomously determine and deliver 100% of all insulin and glucagon doses
- Exclusive collaboration and license agreement with Xeris Pharmaceuticals for a pump-compatible, shelf-stable, glucagon formulation

Clear path to market

- **Phase 2a (current stage):** Feasibility trials to stress test the system's capabilities and iterate accordingly.
    - Completed first-in-human Phase 2a feasibility trial in Q4'25; additional Phase 2a feasibility trial initiated in Q1'26
- **Phase 2b:** Robust pivotal-enabling feasibility trial
- **Phase 3:** Concurrent pivotal trials to support regulatory filings for the glucagon asset (NDA), algorithm (510(k)), and pump (510(k))

Potentially transformative treatment outcomes and burden reduction

- In 6 pre-pivotal outpatient clinical trials, participants achieved **reduced hypoglycemia and increased time in range relative to both standard-of-care treatment and our insulin-only bionic pancreas** configuration[1,2,3,4,5]
- Potential to substantially reduce fear of hypoglycemia and regimen burden which could lead to **reduction in the challenges associated with diabetes management**[6,7]

1. Russell et al. (2014) New England Journal of Medicine, 371:313-325
2. Russell et al. (2016) The Lancet Diabetes and Endocrinology, 4:233-243
3. El-Khatib et al. (2017) The Lancet, 389:369-380
4. Balliro et al. (2017) 77th Scientific Sessions of the American Diabetes Association, 1062-P
5. Sherwood et al. (2018) 78th Scientific Sessions of the American Diabetes Association, 299-OR
6. Weissberg-Benchell et al. (2017) Diabetes Technology & Therapeutics, 12: 723-729
7. Weissberg-Benchell et al. (2016) Journal of Diabetes Science & Technology 4: 840-4

Full Year 2026 Guidance

Previous	New	
$131-136M	$121-126M	Total Revenue
37-39%	37-39%	New Patient Starts through Pharmacy
58.5-59.5%	58.5-59.5%	Gross Margin

Assumptions & Drivers

- **Anticipated revenue deferral of up to $10 million in Q4 2026** as new prospective iLet users await the launch of Mint
- **Stable utilization and retention across DME and pharmacy channels**
- Contribution from **at least 20 new sales territories** onboarded in the first half of 2026
- **Execution of additional formulary agreements and deeper adoption of iLet at the individual health plan level** to drive increases in pharmacy new patient starts
- **Continued cost discipline and improved leverage of manufacturing overhead** at greater scale

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Note: Previous guidance as of July 2026. New guidance as of September 2026.

Strategic Outlook

Innovating for today and the future

Lay the Foundation

2023 to 2024 Achievements

- iLet launch (Black & White)
- CGM integrations (Dexcom G6 & G7, Abbott Freestyle Libre 3 Plus)
- Formulary agreements (e.g., Express Scripts & CVS Caremark)
- Collaboration and licensing agreement with Xeris Pharmaceuticals
- Color iLet launch
- Expansion to 43 sales territories

Execute Relentlessly

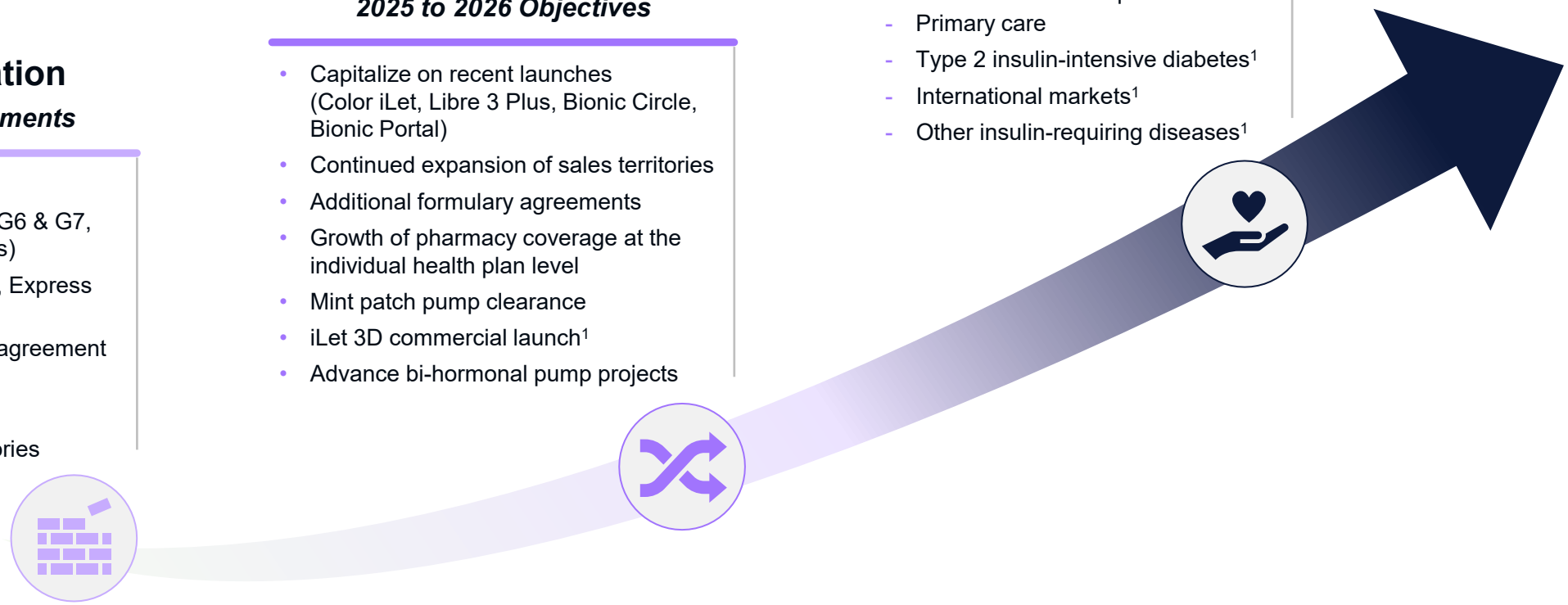
2025 to 2026 Objectives

- Capitalize on recent launches (Color iLet, Libre 3 Plus, Bionic Circle, Bionic Portal)
- Continued expansion of sales territories
- Additional formulary agreements
- Growth of pharmacy coverage at the individual health plan level
- Mint patch pump clearance
- iLet 3D commercial launch[1]
- Advance bi-hormonal pump projects

Revolutionize Care

Future Objectives

- Mint 3D full commercial launch[1]
- Bi-hormonal system commercialization[1]
- Expansion of sales territories
- Growth of pharmacy channel mix
- Market and indication expansion
  - Primary care
  - Type 2 insulin-intensive diabetes[1]
  - International markets[1]
  - Other insulin-requiring diseases[1]



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1. Subject to regulatory clearance

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Appendix

Quarterly Financials & Reconciliation to GAAP Financials

($ in millions)	Q3 2024	Q4 2024	Q1 2025	Q2 2025	Q3 2025	Q4 2025	Q1 2026	Q2 2026
Net Sales	**$16.7**	**$20.4**	**$17.6**	**$23.2**	**$27.3**	**$32.1**	**$27.6**	**$32.0**
Gross Profit	**$8.9**	**$11.7**	**$9.0**	**$12.5**	**$15.1**	**$18.9**	**$16.4**	**$18.9**
% Margin	*53%*	*57%*	*51%*	*54%*	*55%*	*59%*	*59%*	*59%*
Total Operating Expenses	**$19.9**	**$24.7**	**$27.6**	**$32.4**	**$32.2**	**$35.1**	**$40.7**	**$44.5**
Adjusted EBITDA[1]	**($8.7)**	**($11.3)**	**($15.5)**	**($14.5)**	**($12.2)**	**($10.5)**	**($17.7)**	**($17.7)**
Stock-Based Compensation Expense	(2.0)	(1.6)	(2.8)	(4.8)	(4.5)	(4.3)	(5.4)	(7.0)
Change in Fair Value of Warrant Liabilit	0.4	(6.0)	(12.5)	0.0	0.0	0.0	0.0	0.0
Depreciation Expense	(0.3)	(0.2)	(0.3)	(0.3)	(0.4)	(0.5)	(0.6)	(0.7)
Interest Income	0.8	1.0	2.4	3.0	2.8	2.7	2.4	2.2
Income Tax Expense (Benefit)	0.0	0.0	0.0	(0.0)	0.0	0.0	0.0	(0.0)
Litigation Settlement & Other Related Expense	0.0	0.0	0.0	(0.2)	0.0	(0.2)	0.0	(0.1)
Quality System Remediation[2]	0.0	0.0	0.0	0.0	0.0	(0.6)	(0.6)	(0.1)
Net Loss	**($9.7)**	**($18.1)**	**($28.7)**	**($16.9)**	**($14.2)**	**($13.5)**	**($21.9)**	**($23.4)**

1. Adjusted EBITDA addbacks include stock-based compensation expense, depreciation and amortization expense, interest income, provision for state taxes, litigation settlement expense, quality system remediation, and change in fair value of warrant liabilities.
2. Amounts presented under “Quality system remediation” for the three months ended December 31, 2025, March 31, 2026, and June 30, 2026 reflect the same category of expenses previously labeled “Other non-recurring” in our Form 10-K for the year ended December 31, 2025. These expenses relate to our one-time remediation efforts in response to the Form 483 and the Warning Letter, including contractor support and the various updates to our quality system to meet FDA expectations.

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