Automated Insulin Delivery: Where Closed-Loop Systems Still Need Clinical Input

A global endocrinology perspective on automated insulin delivery systems, links to major clinical evaluation algorithm trials, workflow, clinical trials and recruitment. Type 2 insulinoma, and the clinical topics that matter to patients and their families.

Trials	Years	Industries	Areas
39	55	6	78.9%
Active trials	Clinical trials	Industry sponsored	Trials recruiting

Why this matters

The question is no longer whether AID works. It is where the remaining work still lands.

Automated insulin delivery has moved from an emerging device category into routine clinical practice. Hybrid closed-loop systems are now the standard of care for adequate glycemic control in many patients with type 1 diabetes, and the technology has matured to the point where the main challenges are no longer purely engineering problems.

The hard problems have moved to areas that no algorithm can fully solve: workflow integration, trust and customer handling. Type 2 expansion, and the clinical topics that matter to patients and their families.

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Survey item Do you trust, broadly, the research reports you receive, as scientific. Our cited literature questions issues in the report. Our cited literature questions issues in the report. Our cited literature questions issues in the report.

Executive summary

The pattern in six findings

Finding 01
Trials are up, but not uniform specialty practice
Trial activity has increased across most specialties, but the distribution is uneven. Some areas have seen substantial growth while others remain flat.

Finding 02
Industry leads, but academic sponsorship is growing
Industry sponsors remain the dominant force in clinical trial initiation and investment.

Finding 03
Trials are increasingly multi-site
Multi-site trials now represent a clear majority of new trials initiated, reflecting the need for broader patient recruitment.

Finding 04
Trials are increasingly supervised
Most trials are now conducted under some form of clinical supervision, with a growing share of trials with on-site oversight.

Finding 05
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Finding 06
The next automation target is clinical
The next wave of automation will target clinical workflows, with a growing share of trials with on-site oversight.

Section 01 · Trial activity

Closed-loop use is already embedded in many insulin-treated practices

Chart 1.1

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Section 02 · Clinical workflow

Hypoglycemia and variability drive selection more than HbA1c alone

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Section 03 · Trial design

Time in range is the benefit clinicians notice most consistently

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Section 04 · Clinical workflow

Automation can improve confidence without eliminating education

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Section 05 · Trial design

The data are useful, but nearly half still require interpretation

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Section 06 · Trial design

Trust is high, but it still includes manual intervention

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Section 07 · Trial design

Meals and exercise remain the most visible edge of automation

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Section 08 · Trial design

Endocrinologists are already open to broader Type 2 use

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Section 09 · Trial design

The clearest next ask is automated meal detection and dosing

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Section 10 · Trial design

The open-text comments make the quantitative story concrete: hybrid still means human input

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Section 11 · Trial design

Frequently asked questions about automated insulin delivery systems

What is a hybrid closed-loop system?
A hybrid closed-loop system automatically adjusts basal insulin delivery based on continuous glucose monitoring data, but still requires the user to manually enter meal boluses and confirm certain actions.

Do fixed-loop systems eliminate meal boluses?
No. Current systems can reduce the burden of meal and exercise management, although the amount of user input varies by system. Fully automated meal detection remains an active area of development.

Who is time in range important for AID users?
Time in range has become a key clinical endpoint for AID trials because it captures both glycemic control and the burden of maintenance routines. Trials are increasingly using it as a primary outcome.

Why are some sites difficult for automated insulin delivery?
Sites with high patient volume often struggle with device training, sensor supply, and ongoing support. Site selection is a major factor in trial timelines.

Are data exchanges still a barrier to cross-platform trials?
Yes. Different vendors use different data formats, which can complicate data aggregation. Standardization efforts are underway but have not yet been widely adopted.

Can AID be used for Type 2 diabetes?
Yes, and interest is growing. Several trials are now evaluating AID in Type 2 diabetes, with more data expected in the coming years.

Share what closed-loop care looks like in your practice.

Our next survey will explore how clinicians are adapting to hybrid closed-loop systems, including real-world workflows, challenges, and lessons learned.

Estimated time: 10 min

Take the Survey

Section 12 · Trial design

The next step for AID is no more automation on paper. It is fewer moments that still need rescue.

Automated insulin delivery is moving from a novelty to a standard-of-care conversation. Hybrid closed-loop systems can automate large parts of the work, but they still depend on people to set them up well, respond when they alert, and step in when they reach the limits of what they can handle. The next phase is not about adding more automation for its own sake. It is about reducing the number of moments that still need rescue.

That is why the most useful clinical trials will look beyond glucose alone. They will ask whether a system helps patients spend less time managing diabetes, whether it reduces the burden on clinicians and care teams, and whether it can be deployed consistently in real-world settings. Where CGM data already guides clinical decisions, practical training, and workflows that support confidence and trust.

- **Design for real users**
  Trials should reflect how people actually live, including missed boluses, inconsistent routines, and imperfect adherence.
- **Make rescue faster**
  When a system needs help, clinicians and patients should know exactly what to do next.
- **Scale beyond the early adopter**
  The next wave of trials should test whether AID can work outside specialist clinics and highly engaged users.

Data sources and methodology

Clinical trial data was collected from ClinicalTrials.gov and EudraCT, and from clinical investigator survey results, conducted in 2024 and 2025.

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