

MORE THAN 80% TIME IN RANGE WITH OPTIMIZED SETTINGS¹



Of real-world Omnipod® 5 users with time in range (TIR) below 70%

53% do not frequently use the lowest Target Glucose setting²

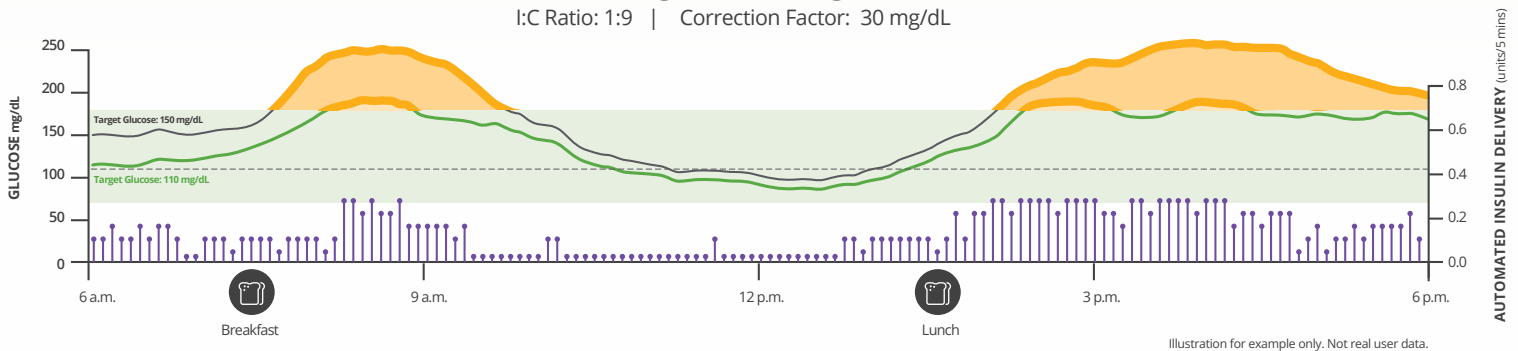
Lowering Target Glucose significantly improved TIR²



Real-world users showed a 12% time in range difference between the lowest and highest Target Glucose setting³

This was achieved with no clinically meaningful impact on time below range.³

Target Glucose
150 mg/dL vs. 110 mg/dL
I:C Ratio: 1:9 | Correction Factor: 30 mg/dL



Optimize settings



Target Glucose Setting: 110 mg/dL¹



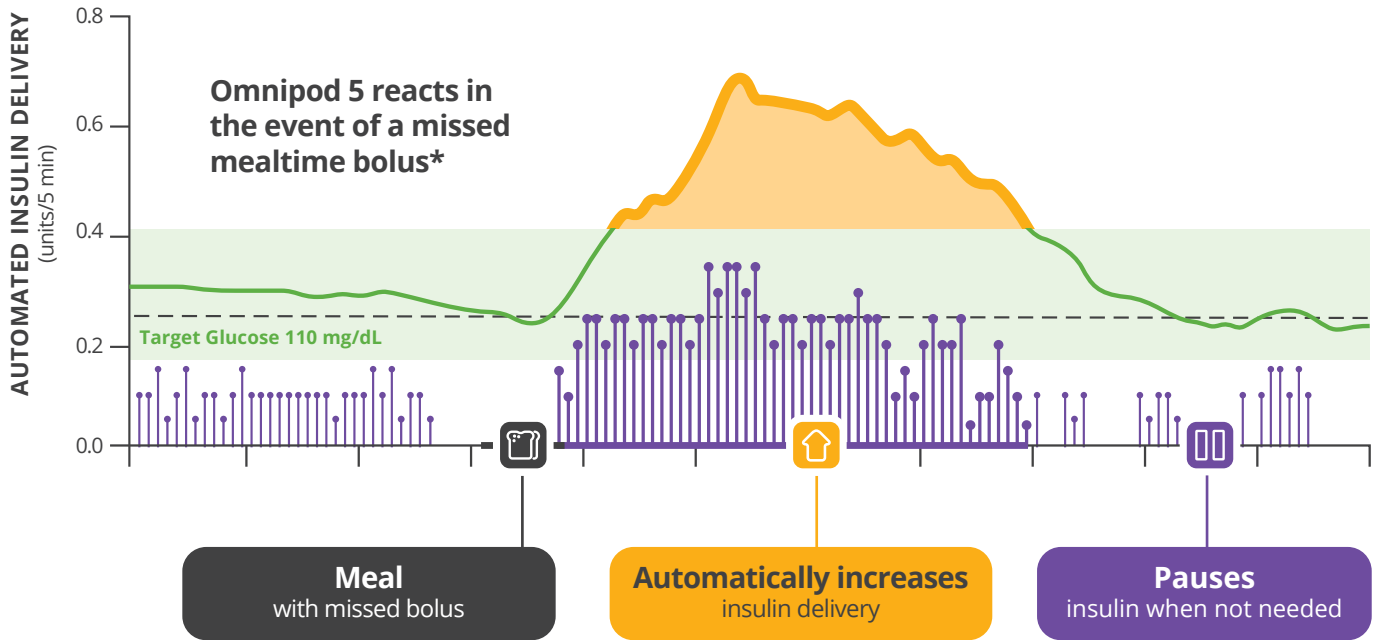
Insulin-to-carb ratio: I:C Ratio x TDI ≤ 350¹



Correction factor: ISF x TDI ≤ 1500 mg/dL¹

1. Retrospective RWE data on file. 2025. Results shown for users with optimized settings including sufficient CGM data (≥75% of days with ≥220 readings), ≥90% time in Automated Mode, ≥5 bolus/day and an average Target Glucose of 110-115 mg/dL. Optimized settings: ISF x TDI = 1500 mg/dL, I:C Ratio x TDI ≤ 350. RF-062025-00014.

Omnipod® 5 autocorrects for highs while helping to protect from lows.^{4,5}



Refer to Cari Berget's "Practical considerations for using the Omnipod® 5 Automated Insulin Delivery System: Clinical experience from the United States and Europe" for tips on optimization.

“Most people with T1D will benefit from AID therapy, and the **Omnipod 5 System has been found safe and effective in diverse clinical trial cohorts, such as those transitioning from multiple daily injection (MDI) therapy and those with high baseline HbA1c, including those who may not count carbohydrates or bolus consistently.**”
—Berget C, et al.

*Bolusing with the Omnipod 5 System is recommended for meals

References:

1. Retrospective RWE data on file. 2025. Results shown for users with optimized settings including sufficient CGM data (≥75% of days with ≥220 readings), ≥90% time in Automated Mode, ≥5 bolus/day and an average Target Glucose of 110-115 mg/dL. Optimized settings: ISF x TDI ≤1500, I:C Ratio x TDI ≤350. RF-062025-00014. 2. Forlenza G, et al. Presented at: ATTD; March 19-22, 2025; Amsterdam, Netherlands. Impact of Lowering Target Glucose Setting with the Omnipod 5 Automated Insulin Delivery System: Real-world data from 403 type 1 diabetes (T1D) users who transitioned from the 150 mg/dL to 110 mg/dL Target Glucose. Omnipod 5 results based on users with ≥90 days CGM data and ≥75% of days with ≥220 readings available. Insulet Data on File. 06.27.25. RF-062025-00038 3. Forlenza G, et al. Presented at: ATTD; March 19-22, 2025; Amsterdam, NL. Real-world data from 403 people with type 1 diabetes aged 2+ using the Omnipod 5 System who transitioned from the 150 mg/dL to 110 mg/dL Target Glucose. Each Target Glucose was used for a consecutive period of 14-90 days. Median time in Range 70-180 mg/dL improved 11.8% (p<0.05). Median time <70 mg/dL +0.23% (p<0.05). Omnipod 5 results based on users with ≥75% of days with ≥220 readings available. Insulet Data on file. 05.15.25. RF-042025-00013 4. Brown S, et al. Diabetes Care. 2021;44:1630-1640. Prospective pivotal trial in 240 participants with T1D aged 6 - 70 yrs [adults/adolescents (n= 128; aged 14-70 yrs) children (n=112; aged 6-13.9 yrs)]. Study included a 14-day standard therapy (ST) phase followed by a 3-month Omnipod 5 hybrid closed-loop phase. Mean time >180 mg/dL in adults/adolescents and children, ST vs. 3-mo Omnipod 5: 32.4% vs. 24.7%; 45.3% vs. 30.2%, P<0.0001, respectively. Mean time <70 mg/dL in adults/adolescents and children, ST vs. 3-mo Omnipod 5: 2.9% vs. 1.3%, P<0.0001; 2.2% vs. 1.8%, P=0.8153, respectively. Results measured by CGM. Study funded by Insulet. 5. Sherr JL, et al. Diabetes Care. 2022. 45(8):1907-1910. Prospective trial in 80 participants with T1D aged 2 - 5.9 yrs. Study included a 14-day standard therapy (ST) phase followed by a 3-month Omnipod 5 hybrid closed-loop (HCL) phase. Mean time >180 mg/dL in very young children (2 - 5.9yrs) as measured by CGM: ST = 39.4%, 3-mo Omnipod 5 = 29.5%, P<0.0001. Mean time <70 mg/dL in very young children (2-5.9 yrs) as measured by CGM: ST = 3.43%, 3-mo Omnipod 5 = 2.46%, P=0.0204. Results measured by CGM. Study funded by Insulet.



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The Omnipod 5 Automated Insulin Delivery System is a single hormone insulin delivery system intended to deliver U-100 insulin subcutaneously for the management of type 1 diabetes in persons aged 2 and older requiring insulin. The Omnipod 5 System is intended for single patient use. The Omnipod 5 System is indicated for use with NovoLog®/NovoRapid®, Humalog®, Trurapi®/Truvelog®/Insulin aspart Sano®, Kirsty®, and Admelog/Insulin lispro Sano U-100 insulin. Refer to the Omnipod® 5 Automated Insulin Delivery System User Guide and www.omnipod.com/safety for complete safety information including indications, contraindications, warnings, cautions, and instructions.

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Omnipod® 5 System Availability Guide



Please contact a Geffen Medical representative for more information