

K223372 Omnipod GO Insulin Delivery DeviceApr 24, 2023
171 days to decisionK223372 · Product code: **LZG** · Chemistry
Source: <https://www.510kdatabase.net/k223372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Insulin (LZG)
Date received	Nov 4, 2022
Decision date	Apr 24, 2023
Days to decision	171 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Insulet Corporation
Location	Bedford, MA, US
Contact	Lauren Davis
510(k) history	23 submissions · 23 cleared · 2005-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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