FDA 510(k) Premarket Notification

CLEARED

K260122 Lancing Device (PressGo, LDE3, LDE3AST, LDE5, EasyStep)

Jul 6, 2026
172 days to decision

K260122 · Product code: **QRL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k260122/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Multiple Use Blood Lancet For Single Patient Use Only (QRL)
Date received	Jan 15, 2026
Decision date	Jul 6, 2026
Days to decision	172 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**SteriLance Medical (Suzhou), Inc.**
Location	Suzhou, CN
Contact	Susan Sun
510(k) history	10 submissions · 10 cleared · 2016-2026