FDA 510(k) Premarket Notification

K132778 GRAB 'N GO PLUS, GRAB 'N GO DIGITAL

CLEARED

Jul 7, 2014
305 days to decision

K132778 · Product code: **ECX** · Anesthesiology
Source: https://www.510kdatabase.net/k132778/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cylinder, Compressed Gas, And Valve (ECX)
Date received	Sep 5, 2013
Decision date	Jul 7, 2014
Days to decision	305 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Praxair Healthcare Services**
Location	Marlborough, MA, US
Contact	TINA WU, PH.D.
510(k) history	1 submissions · 1 cleared · 2014-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k132778/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 24, 2026