K093109 NOVOTWIST NEEDLE, MODELS 30G * 8MM (1/3) 32 G TIP * 5 MM(1/5')

CLEARED

Jun 18, 2010
260 days to decision

K093109 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k093109/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 1, 2009
Decision date	Jun 18, 2010
Days to decision	260 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Novo Nordisk, Inc.**
Location	Princeton, NJ, US
Contact	CINDY CAO
510(k) history	14 submissions · 14 cleared · 2005-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k093109/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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