

K170768 Mini NeedleJul 10, 2017
118 days to decisionK170768 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k170768/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 14, 2017
Decision date	Jul 10, 2017
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ocuject, LLC
Location	Newport Beach, CA, US
Contact	Rebecca K. Pine
510(k) history	8 submissions · 8 cleared · 2017-2024

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