

K153670 Genteel Lancing DeviceMay 12, 2016
143 days to decisionK153670 · Product code: **FMK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k153670/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Dec 21, 2015
Decision date	May 12, 2016
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Genteel, LLC
Location	Midland, TX, US
Contact	Christopher Jacobs
510(k) history	1 submissions · 1 cleared · 2016-2016

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