

**K101417 BABYLANCE HEEL INCISION DEVICE , MODELS  
BLP50, BLP200, BLN50, BLN200**Oct 19, 2010  
152 days to decisionK101417 · Product code: **FMK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k101417/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	May 20, 2010
Decision date	Oct 19, 2010
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medipurpose Pte. , Ltd.</b>
Location	Alpharetta, GA, US
Contact	JULIE STEPHENS
510(k) history	5 submissions · 5 cleared · 2010-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101417/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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