

**K101309 EXELINT ALUMINUM HUB BLUNT NEEDLES**Oct 20, 2010  
163 days to decision

K101309 · Product code: LHI · General Hospital

Source: <https://www.510kdatabase.net/k101309/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	May 10, 2010
Decision date	Oct 20, 2010
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Exelint International, Co.</b>
Location	Los Angeles, CA, US
Contact	ARMAND HAMID
510(k) history	7 submissions · 7 cleared · 2000-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101309/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 25, 2026