

**K181782 Medline Reinforced Epidural Catheter**Mar 4, 2019  
244 days to decisionK181782 · Product code: **BSO** · AnesthesiologySource: <https://www.510kdatabase.net/k181782/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Conduction, Anesthetic (BSO)
Date received	Jul 3, 2018
Decision date	Mar 4, 2019
Days to decision	244 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medline Industries, Inc.</b>
Location	Mchenry, IL, US
Contact	Claire Pigman
510(k) history	238 submissions · 234 cleared · 1977-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k181782/> 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 24, 2026