

**K905069 IODOSORB GEL**Jan 31, 1991  
83 days to decisionK905069 · Product code: **KOZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k905069/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Beads, Hydrophilic, For Wound Exudate Absorption (KOZ)
Date received	Nov 9, 1990
Decision date	Jan 31, 1991
Days to decision	83 days
Third-party review	No

**APPLICANT**

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Company	<b>Perstorp AB C/O Robert Joel Slomoff</b>
Location	Potomac, MD, US
Contact	SLOMOFF
510(k) history	2 submissions · 1 cleared · 1987-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905069/>; 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 July 7, 2026