

**K911202 ATRAUMATIC STANDARD SPROTTE NEEDLE**May 4, 1992  
412 days to decisionK911202 · Product code: **BSP** · Anesthesiology  
Source: <https://www.510kdatabase.net/k911202/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Conduction, Anesthetic (w/wo Introducer) (BSP)
Date received	Mar 19, 1991
Decision date	May 4, 1992
Days to decision	412 days
Third-party review	No

**APPLICANT**

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Company	<b>Pajunk GmbH</b>
Location	Washington, DC, US
Contact	BRUCE MACKLER
510(k) history	19 submissions · 19 cleared · 1991-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911202/>; 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