

K841300 ANTRUM NEEDLE-WOLF TYPEJun 5, 1984
67 days to decision

K841300 · Orthopedic

Source: <https://www.510kdatabase.net/k841300/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Mar 30, 1984
Decision date	Jun 5, 1984
Days to decision	67 days
Third-party review	No

APPLICANT

Company	Popper & Sons, Inc.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1984-1991

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Device record: <https://www.510kdatabase.net/k841300/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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