

**K800312 150 DRIVER/REAMER**Feb 22, 1980  
10 days to decisionK800312 · Product code: **KIJ** · Orthopedic  
Source: <https://www.510kdatabase.net/k800312/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Orthopedic, Dc-powered Motor And Accessory/attachment (KIJ)
Date received	Feb 12, 1980
Decision date	Feb 22, 1980
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Black &amp; Decker(Tm)</b>
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1980-1982

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k800312/>; 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 23, 2026