

**K913632 BATEMAN UPF-III BIPOLAR ENDOPROSTHESIS SYSTEM**Nov 13, 1991  
90 days to decisionK913632 · Product code: **KWY** · Orthopedic  
Source: <https://www.510kdatabase.net/k913632/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Hemi-, Femoral, Metal/polymer, Cemented Or Uncemented (KWY)
Date received	Aug 15, 1991
Decision date	Nov 13, 1991
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Kirschner Medical Corp.</b>
Location	Mchenry, IL, US
Contact	SAM SON
510(k) history	76 submissions · 57 cleared · 1983-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913632/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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