

**K841624 PROBE**Aug 27, 1984  
130 days to decisionK841624 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k841624/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Apr 19, 1984
Decision date	Aug 27, 1984
Days to decision	130 days
Third-party review	No

**APPLICANT**

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Company	<b>Plastafil, Inc.</b>
Location	Walker, MI, US
510(k) history	13 submissions · 13 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k841624/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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