

**K811679 SILASTIC TENDON PASSER H.P.**Jul 10, 1981  
25 days to decisionK811679 · Product code: **HWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k811679/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Passer (HWQ)
Date received	Jun 15, 1981
Decision date	Jul 10, 1981
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Dow Corning Corp. Healthcare Industries Materials</b>
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1976-1985

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Device record: <https://www.510kdatabase.net/k811679/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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