

**K973810 DRG LARGE SCREW SYSTEM**Jan 5, 1998  
90 days to decisionK973810 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k973810/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Oct 7, 1997
Decision date	Jan 5, 1998
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Dolphin Medical Technologies, Inc.</b>
Location	Sarasota, FL, US
Contact	MARK G URBANSKI
510(k) history	2 submissions · 1 cleared · 1998-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k973810/>, 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 21, 2026