

**K980760 KLS-MARTIN MMF SCREW**Dec 21, 1999  
662 days to decisionK980760 · Product code: **DZL** · Dental  
Source: <https://www.510kdatabase.net/k980760/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Intraosseous (DZL)
Date received	Feb 27, 1998
Decision date	Dec 21, 1999
Days to decision	662 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>KLS-Martin L.P.</b>
Location	Jacksonville, FL, US
Contact	PATRICK J LAMB
Website	<a href="https://www.klsmartin.com">https://www.klsmartin.com</a>
510(k) history	78 submissions · 78 cleared · 1994-2026

KLS-Martin L.P. is a surgical device manufacturer based in Jacksonville, US. The company specializes in surgical innovation across orthopedic, dental, and neurology device categories. KLS-Martin has received FDA 510(k) clearances from total submissions since its first clearance in 1994. The company maintains active regulatory status, with its latest FDA 510(k) clearance in 2026. Core product areas include orthopedic implants and fixation systems, dental implants and surgical instruments, and neurosurgical devices including cranial implants and expansion systems. Notable r...

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