

**K103816 COHESION BONE CEMENT**Feb 4, 2011  
37 days to decisionK103816 · Product code: **NDN** · Orthopedic  
Source: <https://www.510kdatabase.net/k103816/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Cement, Bone, Vertebroplasty (NDN)
Date received	Dec 29, 2010
Decision date	Feb 4, 2011
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Teknimed Sas</b>
Location	Round Rock, TX, US
Contact	J. D WEBB
510(k) history	3 submissions · 3 cleared · 2011-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103816/> 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 26, 2026