

**K001132 BONE BIOPSY NEEDLE**May 1, 2000  
21 days to decisionK001132 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k001132/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Apr 10, 2000
Decision date	May 1, 2000
Days to decision	21 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Promex, Inc.</b>
Location	Indianapolis, IN, US
Contact	JOSEPH L MARK
510(k) history	18 submissions · 18 cleared · 1994-2002

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