

**K900082 MAXX 470 CORE BIOPSY GUN**Jun 27, 1990  
173 days to decisionK900082 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k900082/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Jan 5, 1990
Decision date	Jun 27, 1990
Days to decision	173 days
Third-party review	No

**APPLICANT**

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Company	<b>Dlp, Inc.</b>
Location	Mchenry, IL, US
Contact	CHET VANHOF
510(k) history	56 submissions · 56 cleared · 1979-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900082/>; 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 July 2, 2026