

K980166 LIGHTSOURCE OR ILLUMINATORApr 1, 1998
75 days to decisionK980166 · Product code: **HET** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k980166/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Jan 16, 1998
Decision date	Apr 1, 1998
Days to decision	75 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Cuda Products Co.
Location	Jacksonville, FL, US
Contact	KIM REEK
510(k) history	17 submissions · 17 cleared · 1988-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k980166/> Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026