

**K960549 MITYVAC DIRECTIONAL VACUUM EXTRACTOR,  
DISPOSABLE**Aug 29, 1996  
203 days to decisionK960549 · Product code: **HDB** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k960549/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Extractor, Vacuum, Fetal (HDB)
Date received	Feb 8, 1996
Decision date	Aug 29, 1996
Days to decision	203 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Neward Enterprises, Inc.</b>
Location	Mchenry, IL, US
Contact	ALLYSON CARMACK
510(k) history	8 submissions · 8 cleared · 1984-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960549/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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