

K040641 ULTRACINCH ABLATION DEVICE, ULTRACINCH ACCESSORY PACK, MODELS UC-8, UC-9, UC-10, UC-11, UC-12, UC-13, UC-14, UC-ACC-1

May 5, 2004
56 days to decision

K040641 · Product code: **NTB** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k040641/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Ablation, Ultrasound And Accessories (NTB)
Date received	Mar 10, 2004
Decision date	May 5, 2004
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Epicor Medical, Inc.
Location	Sunnyvale, CA, US
Contact	KATHI GUERRANT
510(k) history	2 submissions · 2 cleared · 2004-2004

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k040641/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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