

**K060094 ARCADIA MEDICAL SILICONE WIRE REINFORCED
ENDOTRACHEAL TUBES**

May 8, 2006
116 days to decision

K060094 · Product code: **BTR** · Anesthesiology
Source: <https://www.510kdatabase.net/k060094/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Tracheal (w/wo Connector) (BTR)
Date received	Jan 12, 2006
Decision date	May 8, 2006
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Arcadia Medical Corporation
Location	Apollo Beach, FL, US
Contact	MARK FOOTE
510(k) history	2 submissions · 2 cleared · 2003-2006

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k060094/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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