FDA 510(k) Premarket Notification

K060094 ARCADIA MEDICAL SILICONE WIRE REINFORCED ENDOTRACHEAL TUBES

CLEARED

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