FDA 510(k) Premarket Notification

K073705 MODEL TW SB, SCALP VEIN SET

CLEARED

Mar 24, 2008
84 days to decision

K073705 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k073705/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Dec 31, 2007
Decision date	Mar 24, 2008
Days to decision	84 days
Third-party review	Yes
Summary / Statement	Statement
Other names	MODEL TW LB, STERILE HYPODERMIC NEEDLE

APPLICANT

Company	**Shandong Qiaopai Group Co., Ltd.**
Location	Zibo, CN
Contact	MARK JOB
510(k) history	1 submissions · 1 cleared · 2008-2008