

K073705 MODEL TW SB, SCALP VEIN SETMar 24, 2008
84 days to decisionK073705 · Product code: **FMI** · General HospitalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073705/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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