FDA 510(k) Premarket Notification

K861705 CORONA 3000 (DERMATOLOGIC ULTRAVIOLET LAMP)

CLEARED

May 16, 1986
11 days to decision

K861705 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k861705/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	May 5, 1986
Decision date	May 16, 1986
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Vita-Tan Corp.**
Location	Kansas City, MO, US
Contact	DONALD DEWIT
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k861705/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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