FDA 510(k) Premarket Notification

K081774 SOLX 790 TITANIUM SAPPHIRE LASER

CLEARED

Sep 12, 2008
81 days to decision

K081774 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k081774/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jun 23, 2008
Decision date	Sep 12, 2008
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Solx, Inc.**
Location	Waltham, MA, US
Contact	DOUG ADAMS
510(k) history	1 submissions · 1 cleared · 2008-2008