

**K250425 Ultraviolet Phototherapy Device**Apr 7, 2025  
52 days to decisionK250425 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k250425/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 14, 2025
Decision date	Apr 7, 2025
Days to decision	52 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>APK Technology Co., Ltd.</b>
Location	No. 2 Road(S); Shanghai, CN
Contact	Caifang Wang
510(k) history	5 submissions · 5 cleared · 2009-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Manton Business and Technology Services</b>
Contact	Melody Huang

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k250425/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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