

Take Your Medtech Innovation From China to the US and European markets.

A dedicated pathway for Chinese MedTech companies. Combining decades of China business expertise with world-class regulatory insight and clinical execution excellence in the world's largest MedTech markets.




UNITED STATES
FDA pathway




CHINA
Your Innovation


EUROPE
CE/MDR pathway

YOUR DEDICATED TEAM



Yuval Binur
Business Advisor - China Operations

Decades of on-the-ground experience in China.
Yuval serves as your strategic partner and primary point of contact understanding your market, your goals, and how to position your product for Western expansion.



Meditrial
Regulatory & Clinical Execution

Global medtech CRO with proven U.S. & EU expertise.
Meditrial delivers regulatory strategy, clinical trials, and approval pathways from FDA submissions and Breakthrough Designation to CE marking under EU MDR.

How we accelerate your global market access

Three pillars that turn your China-built innovation into a U.S. and EU success



REGULATORY STRATEGY

FDA & EU MDR pathway selection, Breakthrough Designation, predicate analysis, and submission management.



CLINICAL EXECUTION

Trial design, site selection, and management of pivotal studies meeting U.S. and European evidence standards.



MARKET ACCESS

U.S. / EU entity setup, distribution strategy, reimbursement guidance, and commercial launch support.

Your journey with us to access global market

From first conversation to global growth

1

DISCOVER

Free consultation & market fit assessment

2

STRATEGIZE

U.S. first, EU first, or both — tailored plan

3

EXECUTE

Regulatory submissions & clinical trials

4

APPROVE

FDA clearance & CE marking secured

5

LAUNCH

Market entry & commercial growth

Ready to expand your innovation to global markets?

Schedule a confidential consultation with Yuval Binur to explore your fastest path to U.S. and European approval.



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