PHASE

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VitaTek

REGULATORY

Wouldn't it be nice to have regulatory in-house?

Common Medical Device Regulatory Problems

PROBLEMS

- Hook you with small billable hours
- Charges every 30 minutes
- Solve one piece of a 10 piece puzzle
- Communication issues
- No manufacturing experience
- Never been in the operating room
- No sterilization experience
- No hydrophilic coating experience
- No die mold shop experience
- No marketing experience
- No sales experience
- No business experience
- No medical device success stories
- No international experience
- No tray machine experience

OUR SOLUTION

We are proud to offer solutions to all of these common industry problems through our vertical integration of Regulatory alongside all of our in-house services.

Our Services Inclusions

- ✓ Experienced in-house experts
- ✓ Regulatory pathway from Phase 0
- ✓ Regulatory market authorization prep
- ✓ Submission structures and outlines
- ✓ Regulatory documentation reviews
- ✓ 510k regulatory submission

Why Choose Us

VitaTek's unique Everything In-House model offers medical device innovators a better way to bring their product to market, faster.

- Expert In-House teams for every phase of development
- Clearly defined development roadmap and budget
- Weekly meetings to keep projects on track

▸ Learn more or book a call

www.vitatek.io