Accelerated advisors

Expert regulatory guidance and practical compliance support at every stage.

Get a Market Pathway.

Fit for Your Business.

The World is Yours

Your pathway to market.

Understanding the different and frequently changing requirements of each country you intend to market in can become daunting. Let our experts help you determine the most appropriate pathway and provide the support you need to prepare accurate and timely submissions.

Approval Experience Matters

Approvals around the globe.

Our Advisors are recognized experts in helping companies plan, prepare and submit regulatory approvals and registering new products in every corner of the world.

Hands-on

- Classification determinations
- Documentation requirements
- Market registrations
- U.S. + International submissions
- Renewal strategies

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acceleratedadvisors.com/approvals

Global expertise

- FDA 21 CFR 820 and Parts 110 +111, 210 + 211, 1270 + 1271, 11
- ISO 13485:2016, ISO 14971
- EU MDR, MDSAP
- Product registration Americas, EMEA, APEC

Experienced, qualified, applied

- All Advisors former senior industry management
- Each averaging 30+ years of real-world application
- Current certifications in respective disciplines

Flexible engagement options

- Hourly with accurate estimates
- Project based, short term onsite or online
- Service subscriptions, our most requested option

Move Forward with Confidence.

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