

Medical Devices

Creating and commercializing a cutting-edge medical device is demanding work. Remaining compliant in the face of ever increasing regulation makes that job even more difficult.

We leverage the vast experience gleaned from guiding companies of all sizes through the hard-to-navigate health care environment. We ask the right questions and explore uncovered possibilities to put your product in the best position for success.

- Coverage and reimbursement
- Financing, distribution, and other agreements
- Good Clinical Practice (GCP) compliance
- Humanitarian device exemptions
- Import/export
- In vitro diagnostics
- Intellectual property
- Investigational Device Exemptions (IDEs)
- Litigation and internal investigations
- Post-market compliance and enforcement actions
- Pre-IDE submissions
- Premarket notification and pre-market approvals — 510(k)s and PMAs
- Product safety and liability
- Quality system regulation
- Reclassification petitions

Contacts

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Industries

Life Sciences and Health
Care

- Requests for designation

Representative experience

Obtaining the FDA's designation of expedited review status for a novel orthopedic and women's health device and negotiated favorable review timetables.

Managing PMA and Advisory Panel Process and helped secure approval for novel, noninvasive screening test using advanced DNA technology to detect colorectal cancer.

Drafting one of the first de novo submissions and obtaining multiple downclassifications and subsequent premarket clearances for capsule endoscopy imaging systems.

Assisting a client seeking FDA regulatory marketing approval of the first ever Retinal Prosthesis System intended for partial restoration of vision.

Conducting due diligence for investment bank on medical device startup, uncovering several important FDA issues that substantially altered the target's valuation.

Assisting a client that manufactures an artificial heart to reverse Medicare's national noncoverage decision and secure more advantageous Medicare reimbursements.

Advising a U.S.-based national trade association that provides educational assistance to medical technology companies in Chinese medical device reimbursement rules and policies.

Acting for a global medical device manufacturer on various claims regarding cardiac devices.

Advising a U.S. medical devices company on regulatory and commercial issues regarding the market launch of a gene test in 24 European countries.

Coordinating the defense of a manufacturer of implantable class III medical devices against alleged liability in clinical trials after a voluntary worldwide product withdrawal.

Assisting a medical device manufacturer with investigations by the authorities, criminal investigations, and civil proceedings

arising out of the over-exposure of patients to radiations.

Representing a leading medical device manufacturer in a global recall of one of its products in more than 70 countries.

Counsel to a publicly traded company in a voluntary disclosure to USDOJ and the SEC regarding alleged payments to win public tenders and in follow-on litigation.

Represented a global technology company in the creation of a joint venture to develop next-generation surgical robots.

We have prepared more companies for panel meetings in the last two years than almost any other law firm or consultant in the country.

Latest thinking and events

Hogan Lovells Publications

Proposed measures in Germany for safeguarding supply of critical goods in the combat against COVID-19

Hogan Lovells Publications

Highlights from the DigiHealth Leaders Conference: interoperability, reimbursement, patient centricity, and forging effective partnerships

Hogan Lovells Publications

DHSC Update on Medicines and Medical Devices Contingency Planning

Sponsorships and Speaking Engagements

Applying FDA's Rules in the New World of Online Marketing and Crowdsourcing

Hogan Lovells Publications

A bit more Sunshine: CMS proposes collecting additional information on manufacturer payments under the Open Payments program

Health Alert

Blog Post

The European Commission releases an updated Borderline Manual for medical devices