

Kinmed Publishes 2026 India Medical Device Registration Buyer Guide

Kinmed's 2026 buyer guide explains India's CDSCO registration, risk classes, import licensing, documentation and sourcing planning.

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International medical device procurement works best when registration, documentation and local representation are aligned before the purchase order is issued.”

Kinmed Editorial Team

resource helps international buyers and suppliers understand CDSCO registration, risk classification, import licensing and practical sourcing considerations for India.

[Kinmed](#) has published its “[CDSCO Medical Device Registration: India Buyer Guide 2026](#),” a practical resource for international procurement teams, importers and medical device suppliers planning to source products for the Indian market.

The guide is now available through the Kinmed Knowledge

Hub and explains the medical device registration pathway administered by India's Central Drugs Standard Control Organisation (CDSCO). It is designed to help readers understand the key issues that can affect procurement planning before a purchase order is placed.

The resource covers India's risk-based device classification system, including Classes A, B, C and D; the different filing authorities for lower- and higher-risk devices; the role of the SUGAM online system; and the importance of the Authorised Indian Agent for overseas manufacturers.

For procurement teams, the guide highlights the value of confirming registration status, documentation requirements and importer arrangements early in the sourcing process. It also outlines typical planning timelines and explains why CE marking or FDA clearance may support dossier preparation but does not replace the separate CDSCO import-license process.

The article further addresses practical considerations for international suppliers, including technical-file preparation, local representation, post-market responsibilities and the increasing importance of local-content questions in certain tenders. These factors can affect product selection, quotation preparation, delivery planning and market-entry schedules.

“India is a significant sourcing and destination market, but medical device procurement requires

early alignment between product documentation, registration strategy and local execution,” said the Kinmed Editorial Team. “This guide gives buyers a clearer starting point for evaluating those requirements.”

The guide is intended for informational purposes and does not constitute legal or regulatory advice. Buyers and suppliers should confirm current requirements directly with CDSCO and their appointed Authorised Indian Agent before submitting an application or placing an order.

Read the full guide:

<https://chinakinmed.com/en/article/cdsco-india-medical-device-registration-guide>

About Kinmed

Founded in 2009, Kinmed is a global B2B partner for quality-assured, compliant medical devices and equipment. The company supports healthcare providers and procurement teams with product sourcing, documentation coordination and international logistics across multiple markets.

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