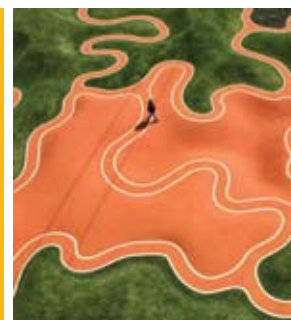


IDRAC[®] INDIA MODULE

A SINGLE, AUTHORITATIVE SOURCE OF INDIAN REGULATORY REQUIREMENTS



WHAT SETS IDRAC APART?

- An unrivaled depth of global information
- Covers both established and emerging economies
- Unique, proprietary regulatory intelligence
- Compiled by Thomson Reuters regulatory experts
- Every entry is curated manually

HOW YOU CAN BENEFIT

- Optimize your regulatory strategy and minimize your risk
- Save time and cost
- Get your product to the market on time and keep it there
- Gain a competitive advantage

UNSURPASSED DEPTH OF COVERAGE

The India module contains 38 explanatory documents written by local experts and based on *IDRAC* proprietary specifications, covering all topics relevant to pharmaceutical industry activities in India, including:

- Authorities/organizations
- Legal framework
- Prescription and supply requirements
- Format and content of applications
- Assembly and submission of applications
- Marketing authorization procedures
- Fees
- Product information
- Clinical research
- Quality assurance
- Pharmacovigilance and risk management
- Import/export
- Advertising/promotion
- Pricing and reimbursement

Additionally, the India module includes a “How to market” section for generics, medical devices, herbal products, drug for pediatric use and Ayurvedic, Siddha and Unani Medicines.

Reference texts cover the Constitution of India, acts, rules, orders, policies, guidelines and notifications from 1940 onwards, as well as relevant forms.

REQUEST A TRIAL

Contact us today to take an IDRAC trial and see for yourself how it will improve your regulatory affairs intelligence. Contact us at scientific.thomsonreuters.com/contact

