



RegulatoryID

Drive Informed Compliance Decisions with the Authoritative Transparency Reference Library

Researched, written, and verified by MedPro Systems' in-house team of pharmaceutical and medical device compliance professionals and attorneys, RegulatoryID offers a centralized, easy-to-use resource library of Federal, State, and International requirements, including:

New! US State Drug Price Reporting

- New Product Reports
- Periodic Reports (Quarterly & Annual)
- Price Increase Report (Pre & Post Price Increase)
- Upon Jurisdiction Request
- Disclosure to HCP
- Prescription Drug Affordability Boards

US Aggregate Spend Reporting

- Transparency Reporting
- Gift Bans and Limitation
- Field Rep Registration and Licensure
- Compliance Program Requirements

Global Aggregate Spend Reporting

- Centralized Disclosure Compendium of 50+ Countries
- Industry Association-Based Requirements (including EFPIA & MedTech Europe)
- National Laws, Industry Codes, Guidance Documents and Disclosure Templates

Stay Up-to-Date on New and Evolving Regulations with Proactive Notifications

Never miss a regulatory update with email alerts on industry news, evolving requirements, and deadline reminders sent directly to your inbox.



CMS Changes for Open Payments Data Collection Beginning January 1, 2023 for Reporting in CY 2024

The Centers for Medicare & Medicaid Services (CMS) finalized changes to the Open Payments Program in the [2022 Physician Fee Schedule](#), which took effect for data collection beginning on January 1, 2023 for reporting in CY 2024.

Learn more at [MedProSystems.com/RegulatoryID](https://www.MedProSystems.com/RegulatoryID)

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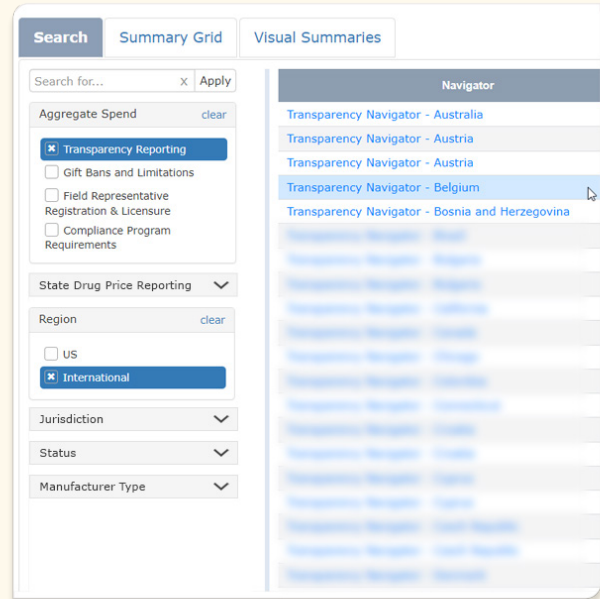


RegulatoryID™

Save Time by Searching and Sorting the Information you Need in Seconds

Gone is the effort of manually scraping disparate industry, legal, and government sources for the laws and regulations impacting your organization. Users can quickly search, filter, and highlight relevant information by:

- **Manufacturer Type:** Narrow requirements by pharmaceutical or medical device
- **Topic:** Filter topics by Aggregate Spend or State Drug Price Transparency Reporting
- **Region:** Sort content into US-based (Federal, State, and Local) or International requirements
- **Status:** Focus on requirements that are pending, current, archived or those that did not pass



Curate Results to Match your Preferred Level of Detail

View content in summary or detail to best fit the needs of your search:

- **Visual Summaries** provide a quick reference guide of high-level requirements across all jurisdictions in scope per topic.

Detailed Compilations (Navigators) outline requirements from disparate laws, regulations, and guidance documents in one convenient location, with links to authoritative sources for each topic and jurisdiction. As Navigators are updated, versions can be compared side by side with highlighted changes for quick review.



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