

SAP Product Data Submission Management for Life Sciences

## Comply with IDMP Requirements Timely and Efficiently



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# Quick Facts

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## Summary

According to European regulations on the Identification of Medicinal Products (IDMP), marketing authorization holders must submit a specific set of data of medicinal products to authorities. The SAP® Product Data Submission Management for Life Sciences application helps your company fulfill these requirements while reducing cost of compliance and improving operational excellence through high data quality, visibility, and consistency.

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## Objectives

- Fulfill upcoming IDMP regulations efficiently
  - Strengthen relationships with authorities through maximized credibility
  - Mitigate risk of delays due to questions or noncompliance
  - Establish a long-term, flexible strategy while the regulatory framework evolves
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## Solution

- Collection, consolidation, and organization of submission-relevant data for IDMP and for similar standards from many sources
  - Ability to secure high data quality
  - Accelerated creation of required reports
  - Simplified submissions to authorities in the right formats
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## Expected Benefits

- Reduced compliance costs
  - Improved R&D productivity as a result of high transparency on product-related data
  - High data quality through automated checks and alerts
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## Learn more

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