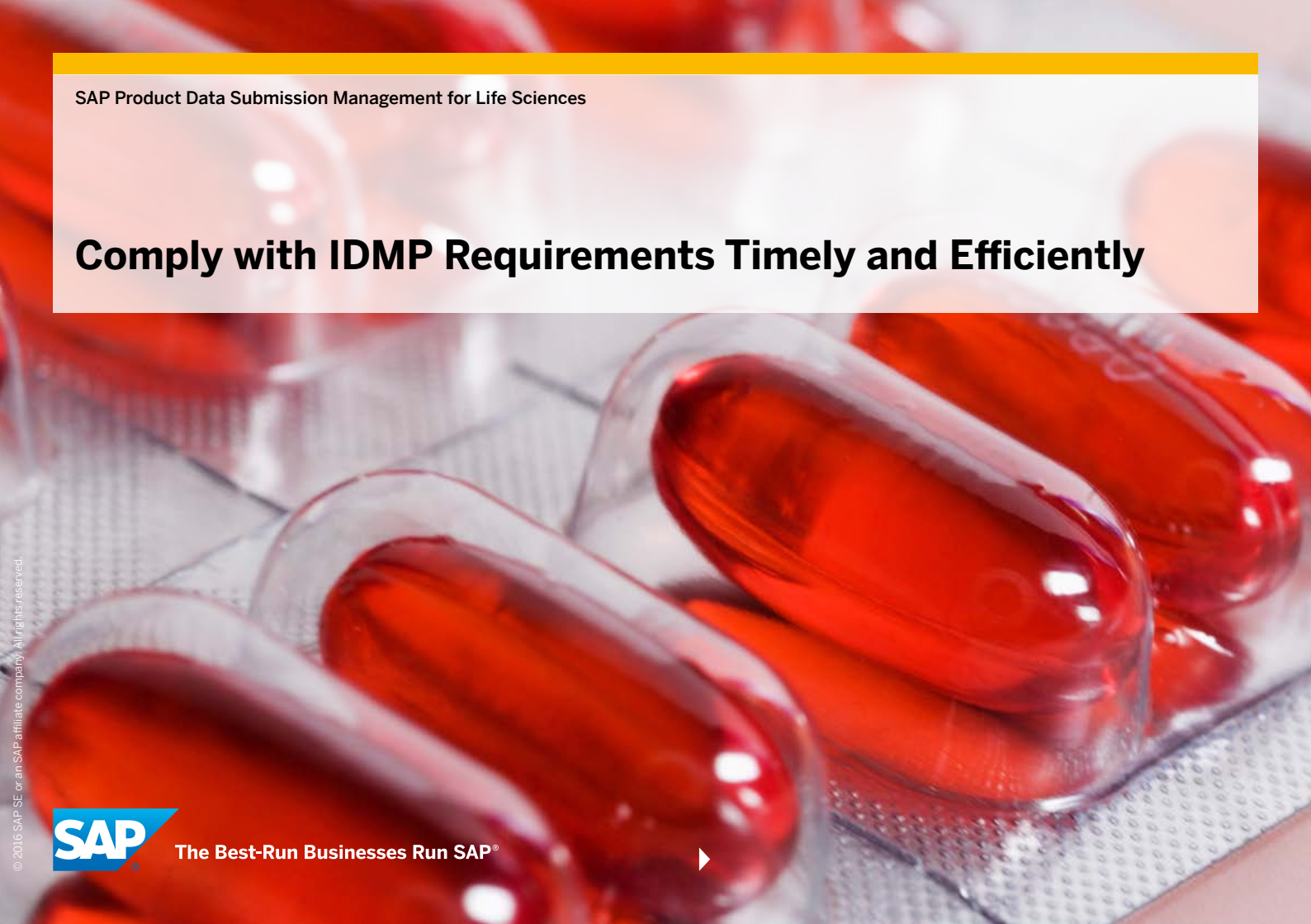
SAP Product Data Submission Management for Life Sciences

Comply with IDMP Requirements Timely and Efficiently



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Quick Facts

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.

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

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

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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