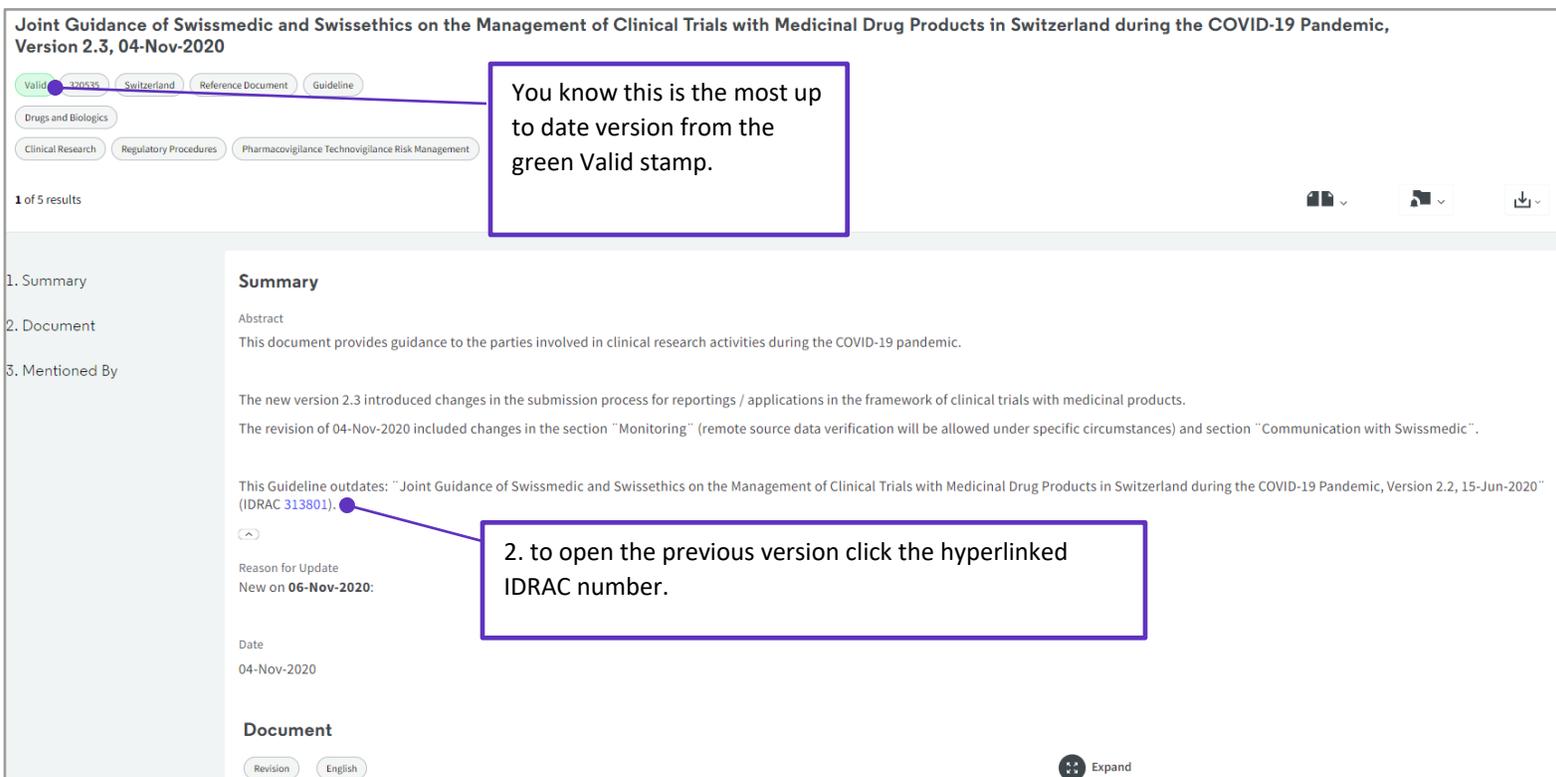


Cortellis Regulatory Intelligence

Consulting previous versions of a regulatory document

Cortellis includes current and outdated document versions, allowing you to easily navigate between them, compare them, go back in history, and identify changes.

1. Information on version management is given in the **Cortellis abstract** under the Summary. If you can't see all details, click **the arrow to show more**. If a previous version exists, a corresponding sentence will be available, such as "This guideline outdates...".



The screenshot shows a document titled "Joint Guidance of Swissmedic and Swissethics on the Management of Clinical Trials with Medicinal Drug Products in Switzerland during the COVID-19 Pandemic, Version 2.3, 04-Nov-2020". The document is marked as "Valid" with a green stamp. A callout box points to this stamp, stating: "You know this is the most up to date version from the green Valid stamp." Below the document title, there are tabs for "Valid", "Switzerland", "Reference Document", and "Guideline". There are also filters for "Drugs and Biologics", "Clinical Research", "Regulatory Procedures", and "Pharmacovigilance Technovigilance Risk Management". The document is listed as "1 of 5 results". The "Summary" section contains the following text: "Abstract: This document provides guidance to the parties involved in clinical research activities during the COVID-19 pandemic. The new version 2.3 introduced changes in the submission process for reportings / applications in the framework of clinical trials with medicinal products. The revision of 04-Nov-2020 included changes in the section 'Monitoring' (remote source data verification will be allowed under specific circumstances) and section 'Communication with Swissmedic'. This Guideline outdates: 'Joint Guidance of Swissmedic and Swissethics on the Management of Clinical Trials with Medicinal Drug Products in Switzerland during the COVID-19 Pandemic, Version 2.2, 15-Jun-2020' (IDRAC 313801)." A callout box points to the hyperlinked IDRAC number, stating: "2. to open the previous version click the hyperlinked IDRAC number." The "Reason for Update" is "New on 06-Nov-2020" and the "Date" is "04-Nov-2020". The "Document" section shows "Revision" and "English" tabs. An "Expand" button is visible at the bottom right.

2. The previous version will have a **red outdated stamp** appearing in the top left-hand corner of the report.

Outdated: Joint Guidance of Swissmedic and Swissethics on the Management of Clinical Trials with Medicinal Drug Products in Switzerland during the COVID-19 Pandemic, Version 2.2, 15-Jun-2020

Outdated 315801 Switzerland Reference Document Guideline

Drugs and Biologics

Clinical Research Regulatory Procedures Pharmacovigilance Technovigilance Risk Management

Red Outdated stamp.

As part of the abstract, Cortellis editors also highlight the key changes against the previous version.

In this new version a new section on "Resumption of clinical trials activities following the COVID-19 pandemic" has been added and the section "Monitoring" has been updated.

This Guideline outdates: "Joint Guidance of Swissmedic and Swissethics on the Management of Clinical Trials with Medicinal Drug Products in Switzerland during the COVID-19 Pandemic, Version 2.1, 07-Apr-2020 (IDRAC

Reason for Update

Content Update on **06-Nov-2020**: This document is outdated by: "Joint Guidance of Swissmedic and Swissethics on the Management of Clinical Trials with Medicinal Drug Products in Switzerland during the COVID-19 Pandemic, Version 2.3, 04-Nov-2020" (IDRAC 320535).

New on **16-Jun-2020**: [Link back to the valid document here.](#)

Date

15-Jun-2020

To compare versions, please consult the **Cortellis Side By Side Viewer** guide.

3. After running a search, you have the option to Include Outdated documents in your results list by toggling this button on. The default is for Outdated documents to not be included in search results.

Refine Search

Side by Side Viewer

Showing 1-10 of 60,456 results

Include Outdated My Regions

Customize Columns Sorted by Relevance

Summary	Title	Abstract	Reason for Update	Country/Region	Language(s)	Product Cat
May-2015 V EU EN RD	121897 - EMA Leaflet: Pharmacovigilance - May-2015	This leaflet provides information on pharmacovigilance, more specifically addressing these topics:- Benefit-risk	N/A	European Union	English	Drugs and I
Jun-2006 O UK EN RD	56375 - Outdated: MHRA: Statutory Pharmacovigilance Inspection - Summary of	This document provides information to assist companies with the preparation of	Please refer to the current version of Statutory Pharmacovigilance Inspection - Summary of Pharmacovigilance (SPS)	United Kingdom	English	Drugs and I

Identify Valid from Outdated in a result list from the green and red icons.

For more information contact Customer Service at **LS Product Support**.