

Accessing and launching the PDF Side by Side Viewer

The Cortellis Side by Side Viewer enables you to select, view, scroll and search within two pdf documents of your choice. This powerful tool can assist you to evaluate and compare two pdf documents side by side. For example:

- Two related documents, such as a previous and a current version
- Regulatory Summaries on two topics
- Regulatory Summaries for two different countries/regions
- Two product approval documents
- Comparing an English Machine Translation with the native language version of a source document

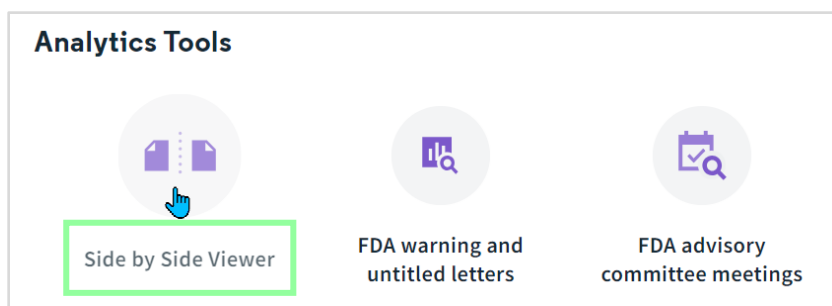
Accessing and launching the PDF Side by Side Viewer

The viewer can be accessed and launched from multiple places in your Cortellis Regulatory journey:

- Regulatory Intelligence homepage
- Regulatory results page
- Regulatory document or report page
- Side-by-side document list and history
- Machine Translation Document icon in a source document

From the Regulatory Intelligence homepage

Scroll down the Regulatory Intelligence homepage to the Analytics Tools and click on **Side by Side Viewer**.



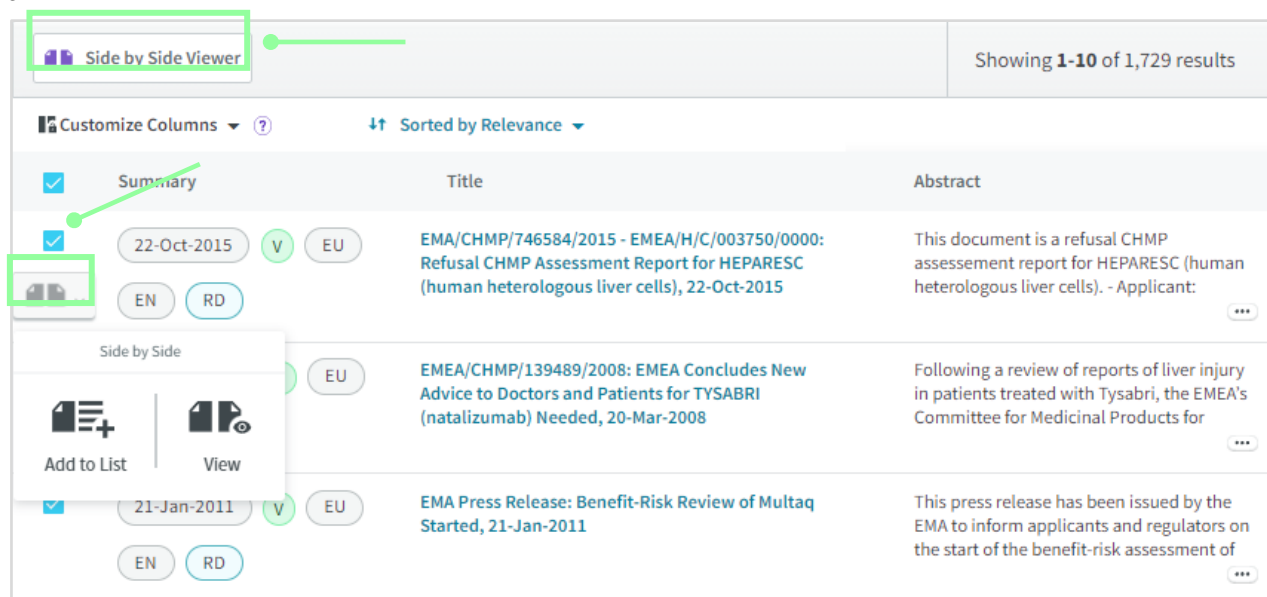
From the regulatory results page

On the results page click on the **Side by Side Viewer** button at the top left to launch the tool.

If you hover over any of the document tick boxes on the left, the Side by Side Viewer button will also appear. You will have two choices: **Add to List** or **View**.

If you click **View**, the pdf viewer will be launched with the related document displayed on the left-hand side. You can then load a second document to view both side by side.

Add to List will add the related document to a side by side list. See the next section about your side by side list.



The screenshot shows the regulatory results page. At the top left, the **Side by Side Viewer** button is highlighted with a green box. Below it, a table of results is displayed. The first row shows a document titled "EMA/CHMP/746584/2015 - EMEA/H/C/003750/0000: Refusal CHMP Assessment Report for HEPARESC (human heterologous liver cells), 22-Oct-2015". The second row shows a document titled "EMA/CHMP/139489/2008: EMEA Concludes New Advice to Doctors and Patients for TYSABRI (natalizumab) Needed, 20-Mar-2008". The third row shows a document titled "EMA Press Release: Benefit-Risk Review of Multaq Started, 21-Jan-2011". A green arrow points to the **Side by Side** button in the top right corner of the table. A green box highlights the **Add to List** and **View** buttons in the bottom right corner of the table.

From the regulatory document or report page

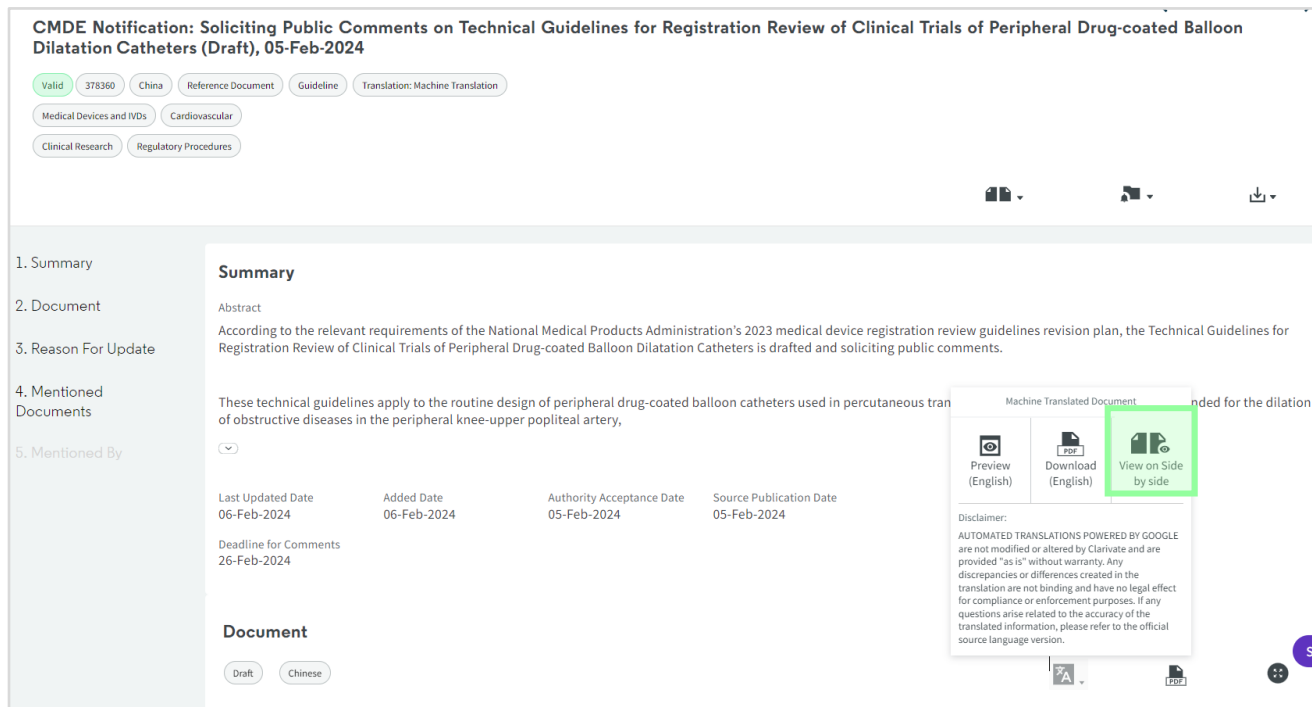
The **Add to List** and **View** options are also available on the document page if you click the **Side by Side Viewer** button at the top right of the Summary section. Click on either option to add the report to your side by side list or to launch the pdf viewer, respectively.



The screenshot shows the regulatory document page. At the top, the **Clinical Research** section is highlighted. Below it, a list of documents is displayed. The first document is titled "EMA/CHMP/746584/2015 - EMEA/H/C/003750/0000: Refusal CHMP Assessment Report for HEPARESC (human heterologous liver cells), 22-Oct-2015". The second document is titled "EMA/CHMP/139489/2008: EMEA Concludes New Advice to Doctors and Patients for TYSABRI (natalizumab) Needed, 20-Mar-2008". The third document is titled "EMA Press Release: Benefit-Risk Review of Multaq Started, 21-Jan-2011". A green arrow points to the **Side by Side** button in the top right corner of the document page. A green box highlights the **Add to List** and **View** buttons in the bottom right corner of the document page.

From the Machine Translation Document icon in a source document

Documents that are presented in their native language, will have an English Machine Translation version. Click the Icon to view options and select View on Side by side to open the viewer and compare the native language version with the Machine Translation in English in the viewer.



CMDE Notification: Soliciting Public Comments on Technical Guidelines for Registration Review of Clinical Trials of Peripheral Drug-coated Balloon Dilatation Catheters (Draft), 05-Feb-2024

Valid 378360 China Reference Document Guideline Translation: Machine Translation

Medical Devices and IVDs Cardiovascular Clinical Research Regulatory Procedures

1. Summary

2. Document

3. Reason For Update

4. Mentioned Documents

5. Mentioned By

Summary

Abstract

According to the relevant requirements of the National Medical Products Administration's 2023 medical device registration review guidelines revision plan, the Technical Guidelines for Registration Review of Clinical Trials of Peripheral Drug-coated Balloon Dilatation Catheters is drafted and soliciting public comments.

These technical guidelines apply to the routine design of peripheral drug-coated balloon catheters used in percutaneous tran of obstructive diseases in the peripheral knee-upper popliteal artery,

Last Updated Date 06-Feb-2024 Added Date 06-Feb-2024 Authority Acceptance Date 05-Feb-2024 Source Publication Date 05-Feb-2024

Deadline for Comments 26-Feb-2024

Document

Draft Chinese

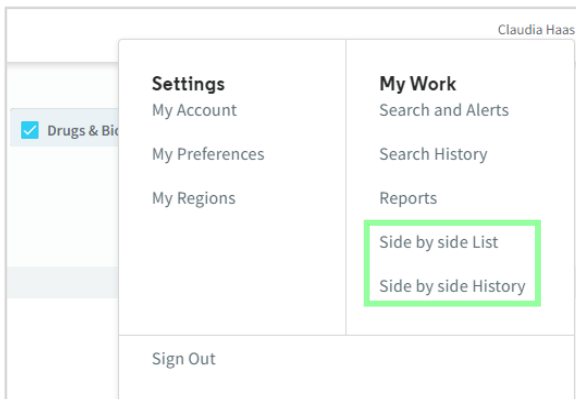
Machine Translated Document

Preview (English) Download (English) View on Side by side

Disclaimer: AUTOMATED TRANSLATIONS POWERED BY GOOGLE are not modified or altered by Clarivate and are provided "as is" without warranty. Any discrepancies or differences created in the translation are not binding and have no legal effect for compliance or enforcement purposes. If any questions arise related to the accuracy of the translated information, please refer to the official source language version.

From the Side by side document List and History

To access the list and the history of your side by side documents, **hover over your name** at the top right of any Cortellis page. Under My Work you will see links to access your **Side by side List** and your **Side by side History**.



Claudia Haas

Settings

My Account

My Preferences

My Regions

Sign Out

My Work

Search and Alerts

Search History


Reports

Side by side List

Side by side History

Click on **Side by side List** to see all documents that you've added from a regulatory results or report page. Select any two documents and click on **Side by side document viewer** on the top right of the page to launch the viewer. Both documents will be displayed in the pdf viewer side by side.

[< Back](#) | [Forward >](#) | My Searches and Alerts

 Side by side document viewer
 ✖ Delete Item


[Saved Searches](#) | [Saved Reports](#) | [Search History](#) | [Side by Side Document List](#) | [Side by Side History](#) | [Saved Predictions](#)

The side by side documents list contains Cortellis documents only.

<input checked="" type="checkbox"/>	Content	Title	Abstract	Date	Region	IDRAC Number
<input checked="" type="checkbox"/>	Regulatory	Clinical Research	This Cortellis Regulatory Summary document summarizes the regulatory requirements for Clinical Research. Main aspects include regulatory framework, planning and conducting a clinical trial, CTA/IND review by competent authority/-ies, ethical review, observational studies, Show more	20-Oct-2021	Ukraine	92217
<input checked="" type="checkbox"/>	Regulatory	Clinical Research	This Cortellis Regulatory Summary document summarizes the regulatory requirements for Clinical Research. Main aspects include regulatory framework, planning and conducting a clinical trial, CTA/IND review by competent authority/-ies, ethical review, observational studies, Show more	20-Oct-2021	United Arab Emirates	102923

Click on **Side by side History** to see a record of the last 50 pairs of documents you viewed side by side in the tool. You can review any of these pairs of documents in the viewer. Please note that we update all documents to the most recent version. So for example, if a Regulatory Summary was updated since you last viewed it, we will display the latest version of this document. Also note that we do not keep details of documents that are stored on your computer when you viewed them in the side by side tool.

[< Back](#) | [Forward >](#) | My Searches and Alerts

 Side by side document viewer
 ✖ Delete Item

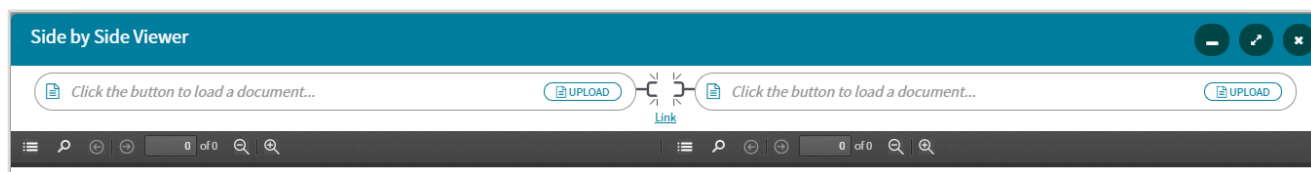
[Saved Searches](#) | [Saved Reports](#) | [Search History](#) | [Side by Side Document List](#) | [Side by Side History](#) | [Saved Predictions](#)

The side by side history can include a maximum of 50 files.

<input type="checkbox"/>	Content	Title	Abstract	Date	Region	IDRAC Number
<input type="checkbox"/>	Regulatory	Pharmacovigilance and Risk Management	This Regulatory Summary details the requirements for Pharmacovigilance & Risk Management in Venezuela. Main aspects include legal framework, definitions, reporting requirements, responsibilities, preparation of PSURs (IPS) and requirements for Risk Management Show more	20-Oct-2021	Venezuela	116124
	Regulatory	Guide: Periodic Safety Update Reports (PSURs) – Guide for the Industry, Aug-2014	This document is intended to give a practical tool that facilitates the elaboration of Periodic Safety Update Reports (PSURs), clearly oriented to MAH 'Marketing Authorization Holders' of medicinal products. It contains guidelines on the information PSURs must contain and t Show more	20-Oct-2021	Venezuela	294633
<input type="checkbox"/>	Regulatory	Clinical Research	This Cortellis Regulatory Summary document summarizes the regulatory requirements for Clinical Research. Main aspects include regulatory framework, planning and conducting a clinical trial, CTA/IND review by competent authority/-ies, ethical review, observational studies, Show more	20-Oct-2021	United Arab Emirates	102923
	Regulatory	Clinical Research	This Cortellis Regulatory Summary document summarizes the regulatory requirements for Clinical Research. Main aspects include regulatory framework, planning and conducting a clinical trial, CTA/IND review by competent authority/-ies, ethical review, observational studies, Show more	20-Oct-2021	Ukraine	92217

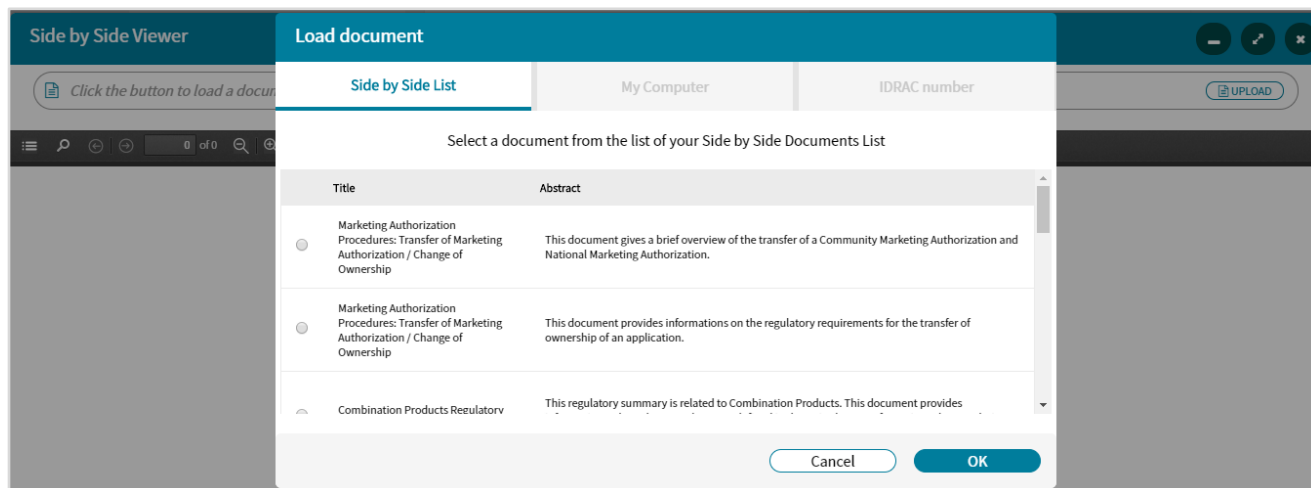
Using the PDF Side by Side Viewer

Once in the Side by Side viewer, click on one of the Upload buttons to upload documents from your side by side list or history, from your computer or by entering an IDRAC number of a Cortellis document.

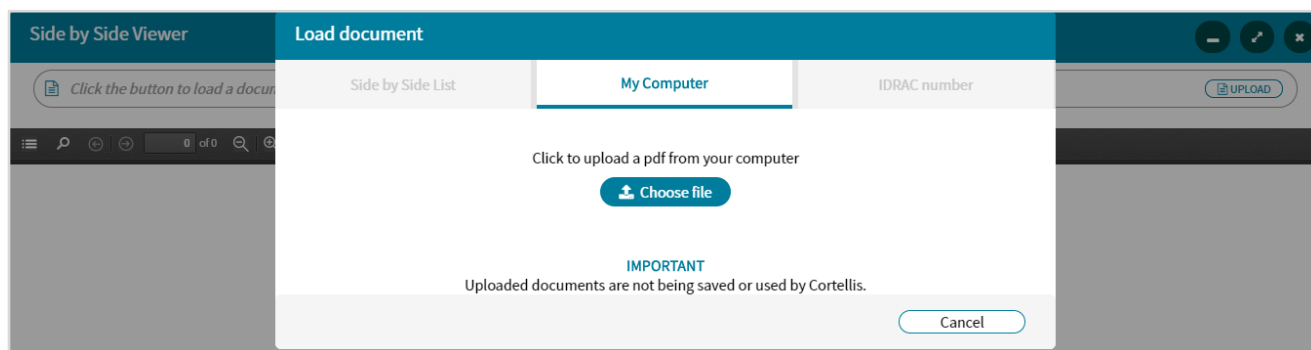


Loading documents into the viewer

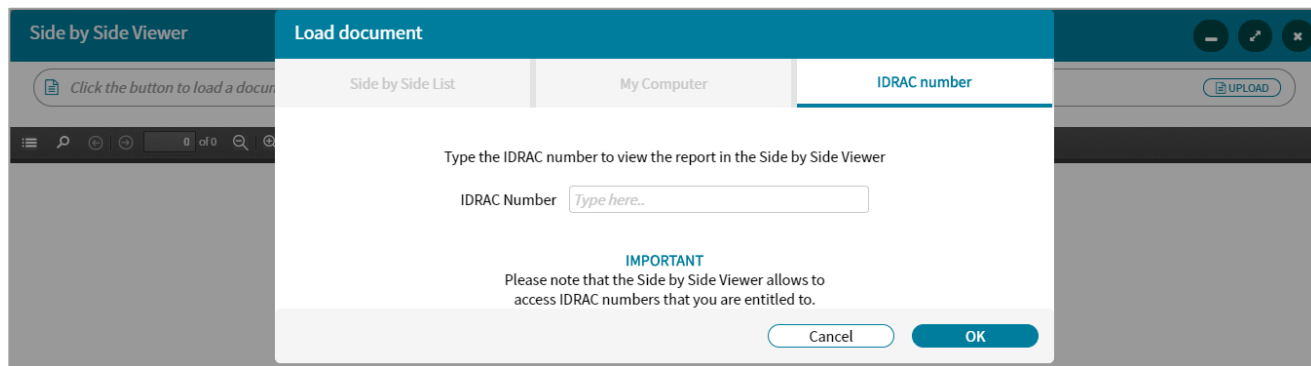
If you've added documents to the Side by side List previously, they will automatically be listed. Select **the document of interest** and click **OK** to load it. The selected pdf document will be displayed in the relevant panel with its title appearing in the box next to the load button. Click on the other **Load** button to add a second document.



To upload a document from your own files, click on **My Computer** and select your pdf of interest, which will then appear in the relevant panel of the viewer.

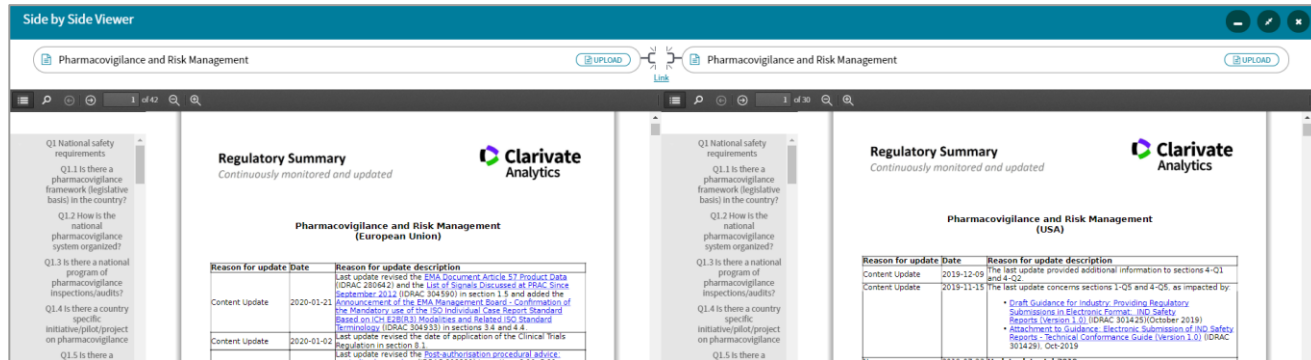


Alternatively, you can enter an **IDRAC number** to upload a Cortellis document to the viewer.

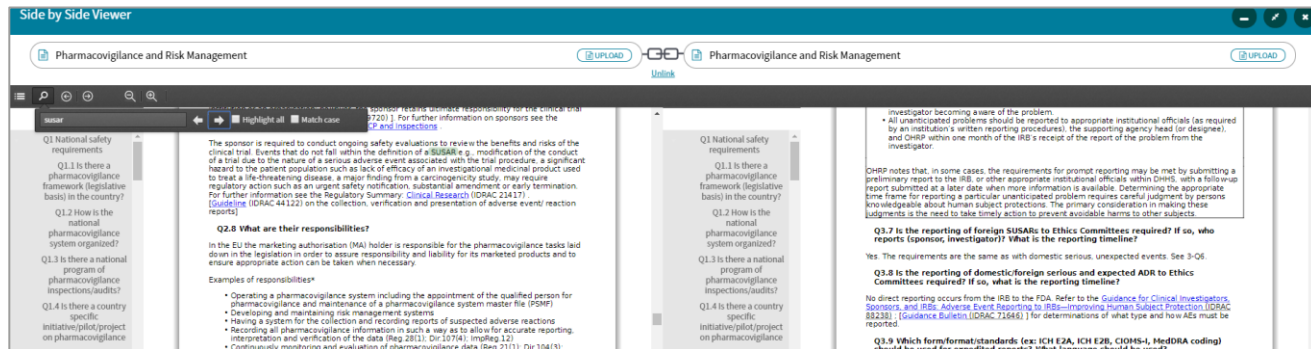


Using the viewer tools








When two documents have been uploaded to the viewer, they can be unlinked or linked to each other. If unlinked (default setting) you will see a toolbar at the top left corner of each document. From here you can scroll or search within each document independently. Click the Link button between the two documents to link to them.



If the documents are linked, there will be a joint toolbar at the top left corner and all options work in parallel. For example, you can search simultaneously in both documents and you can scroll or paginate synchronously in both documents.



The following actions can be executed from the toolbar:

	Show/hide bookmark9(s) or table of content(s)		Search within the document(s)
	Previous page(s)		Next page(s)
	Go to a specific page number		Zoom In to the document(s)
	Zoom Out of the document(s)		

You can fully minimize the viewer and continue working in Cortellis. For instance, run a search and add more documents to your side by side list. You can at any time get back to the viewer by maximizing it.

Side by Side Viewer



For more information contact Customer Service at [LS Product Support](#).