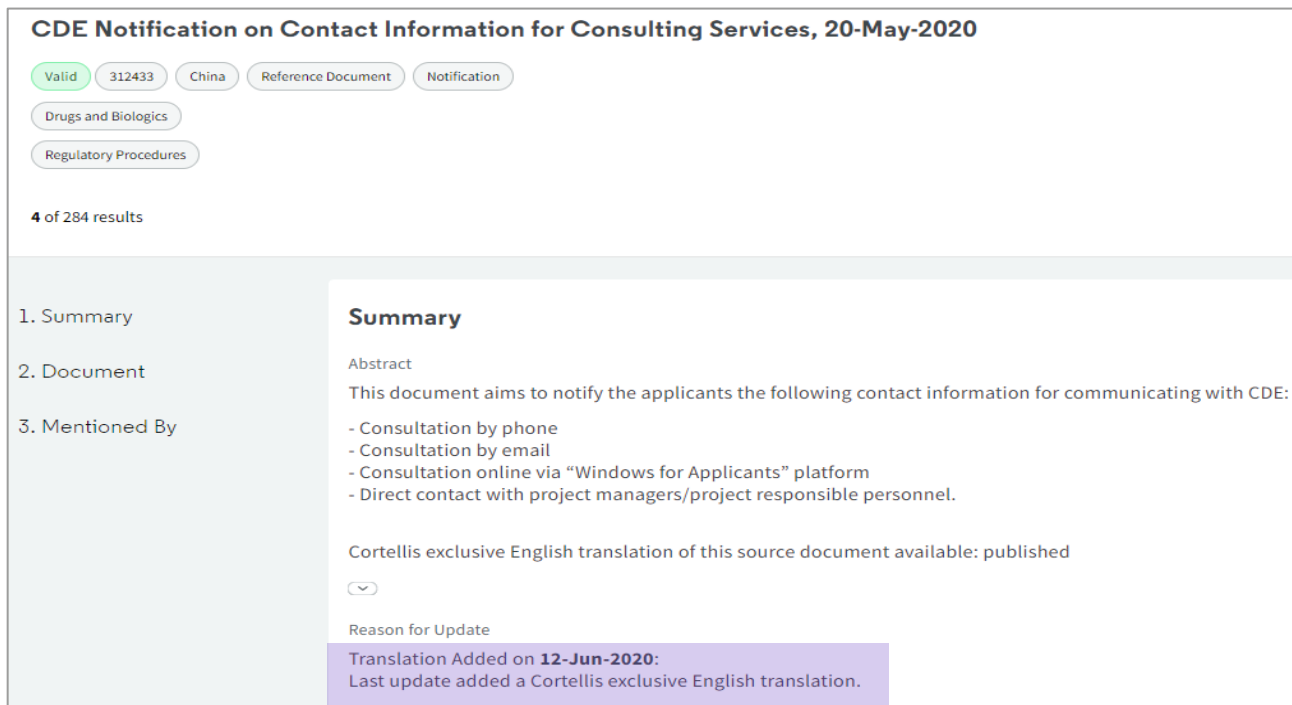


Cortellis Regulatory Intelligence

Identify multilingual documents and exclusive English translations

This guide shows how to identify regulatory documents covered in various languages in *Cortellis*. In addition to official translations published by the regulatory authorities, *Cortellis* provides exclusive English translations prepared by a professional agency.

1. All documents in Cortellis include **English titles** and value-added **English abstracts** written by Cortellis experts, available in the Summary section of the document page. The **Reason for Update** field allows you to easily find out if a **Cortellis exclusive English translation** is available and when it was added.



CDE Notification on Contact Information for Consulting Services, 20-May-2020

Valid 312433 China Reference Document Notification

Drugs and Biologics

Regulatory Procedures

4 of 284 results

1. Summary

2. Document

3. Mentioned By

Summary

Abstract

This document aims to notify the applicants the following contact information for communicating with CDE:

- Consultation by phone
- Consultation by email
- Consultation online via “Windows for Applicants” platform
- Direct contact with project managers/project responsible personnel.

Cortellis exclusive English translation of this source document available: published

Reason for Update

Translation Added on **12-Jun-2020**:
Last update added a Cortellis exclusive English translation.

- The document labels indicate the languages available, in this case **English** and **Chinese**. Within the pdf document you have direct access to all language versions provided via the table of contents on the left.

Document

None English Chinese Expand

1 von 6 100%

File 1

- Internal English version
 - Notice of CDE on the Contact Information of Business Consultation Services
 - 1. Telephone
 - 2. Email
 - 3. Applicants' Window
 - 4. Project management personnel
- Original file
 - 药审中心关于业务咨询服务联络方式的通知
 - 一、电话咨询
 - 二、邮件联系
 - 三、申请人之窗
 - 四、联系项目管理 人员

Notice of CDE on the Contact Information of Business Consultation Services

Released on 20 May, 2020

Considering the new normal of COVID-19 pandemic prevention and control, in order to ensure the quality and efficiency of business consultation services and facilitate applicants' effective contact and communication with our centre (Centre for Drug Evaluation, NMPA (CDE)), applicants may contact us by the following means, and the relevant specific matters are hereby announced as follows:

1. Telephone

In order to comprehensively implement the joint prevention and control measures for COVID-19 pandemic, minimise the flow and gathering of people, block the spread channels of the pandemic, and effectively protect applicants' life safety and health, starting from 3 February, 2020, CDE has suspended on-site consultation services, and applicants may dial the following phone numbers for consultation.

Consultation category	Tel.	Date	Working hours
Consultations about	010-85242306	Tuesdays and Thursdays	9:00-11:30, 13:30-

- The Regulatory homepage Search options as well as Refine Search on the results page allow you to filter documents by language. Click **All other filters**, then click **Languages**, select the **language(s) of interest** and click **Apply**.

Search

Document title, topic, country, reference number... Search Advanced search

Medical Devices Specialty

Regulatory Version

2 Languages

Search

Document title, topic, country, ref...

Filter

Country/Region Topic Document

English (194312) French (9288) Spanish (6265) German (5204)

Portuguese (4770) Chinese (4405) Japanese (3591) Korean (2948)

Italian (2639) Dutch (1831) Danish (1823) Swedish (1796) Arabic (1770)

Turkish (1520) Greek (1147) Czech (1073) Russian (846) Thai (809)

Polish (731) Finnish (679) Romanian (670) Slovak (614) Hebrew (601)

All other filters Reset Filters

Cancel Apply

- Multilingual documents can also easily be identified from the **labels on the results page** in the Summary and the Languages columns.

<input checked="" type="checkbox"/>	Summary	Title
<input checked="" type="checkbox"/>	20-Nov-2020 V CH DE,EN,FR,IT RD	321401 - Swissmedic Communication: Validity of GMP Certificates During the COVID-19 Pandemic - 20-Nov-2020
<input checked="" type="checkbox"/>	19-Nov-2020 V CO ES RD	321396 - Guide for the Marketing Authorisation Application for Radiopharmaceutical Products (Draft) - 19-Nov-2020

Abstract	Reason for Update	Country/Region	Language(s)
In this Communication the Swissmedic informs that the GMP certificates issued by Swissmedic and based on a routine GMP ...	New on 23-Nov-2020 :	Switzerland	German English ...
INVIMA calls for public consultation all MAHs and importers who wish to contribute on the guidelines and requirements for applying ...	New on 23-Nov-2020 :	Colombia	Spanish

Cortellis English exclusive translations of local documents are available for **notifications, guidelines, regulations, laws, degrees, orders and announcements, among others.**

Translations for **Drugs and Biologics** are provided for **Brazil, China, Israel, Japan, Mexico, Russia, South Korea and Taiwan.** Coverage for Medical Devices and IVDs includes **Argentina, Austria, Brazil, China, France, Germany, Israel, Italy, Mexico, Russia, South Korea, Spain, Switzerland, Japan, Taiwan and Turkey.**

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