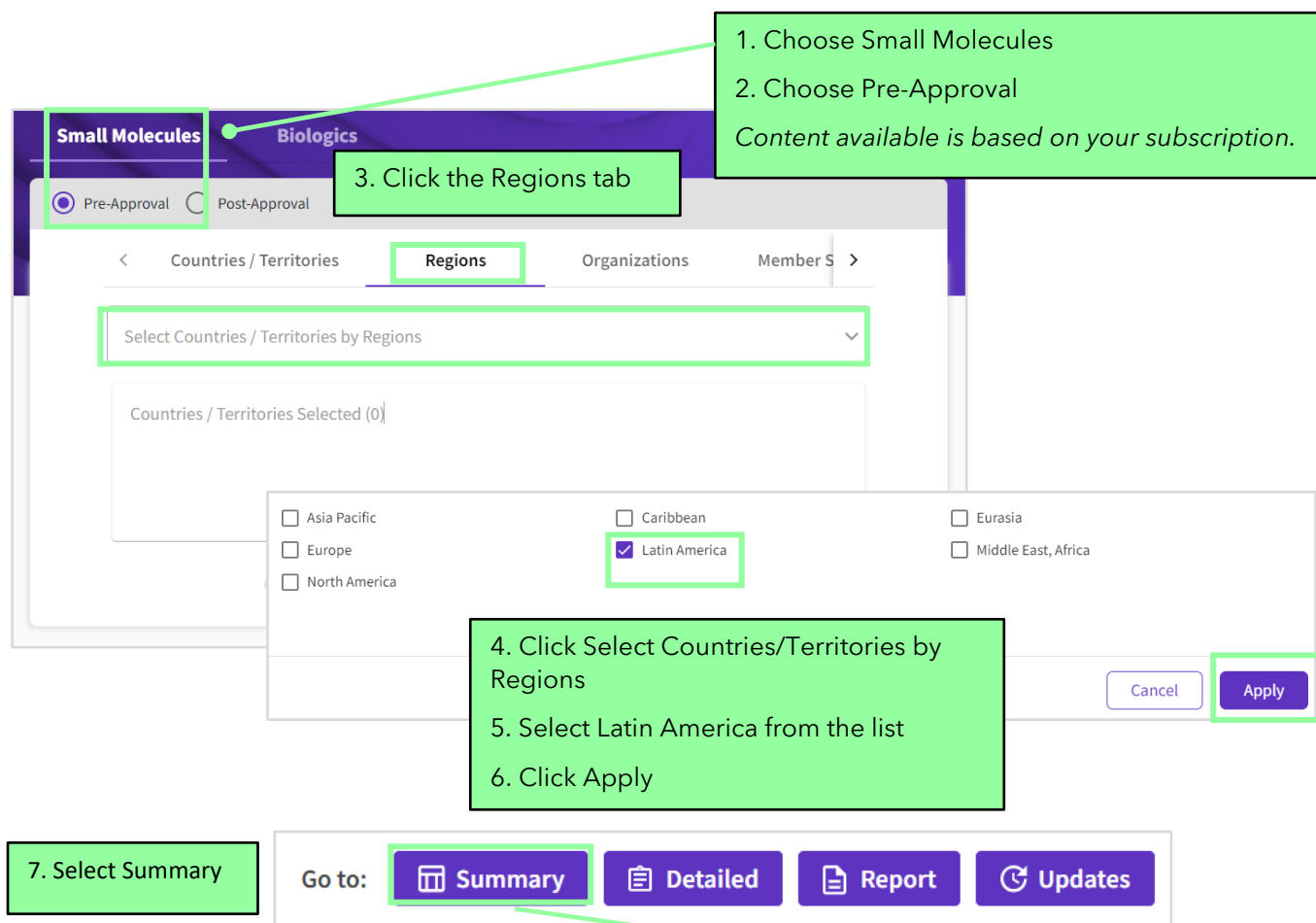


Quickly compare key information with Summary requirements

Example: You want to register your small molecule drug in all Latin American countries, and you need to compare the required licenses and certificates for each country.

Please note: This example only covers Pre-Marketing Small Molecules. For Post-marketing Small Molecules, and Pre- and Post-marketing Biologics the Summary tables are all different and applicable to that content set. Be sure to explore all the Summary tables that are available with your subscription.



The screenshot shows the Cortellis CMC Intelligence interface. At the top, there are tabs for 'Small Molecules' and 'Biologics'. Below these are radio buttons for 'Pre-Approval' (selected) and 'Post-Approval'. A navigation bar includes 'Countries / Territories', 'Regions' (highlighted), 'Organizations', and 'Member S'. A dropdown menu is open for 'Select Countries / Territories by Regions'. Below this, a list of regions is shown with checkboxes: Asia Pacific, Europe, North America, Caribbean, Latin America (checked), Eurasia, and Middle East, Africa. At the bottom, there are buttons for 'Cancel' and 'Apply'. A 'Go to:' section contains buttons for 'Summary' (highlighted), 'Detailed', 'Report', and 'Updates'.

1. Choose Small Molecules
2. Choose Pre-Approval
3. Click the Regions tab
4. Click Select Countries/Territories by Regions
5. Select Latin America from the list
6. Click Apply
7. Select Summary

Content available is based on your subscription.



Summary

Compare and contrast core requirements for your selections to identify differences and similarities for your global operations.

In the left-hand column, select the requirements you'd like to compare. For our example we'll focus on Licences and Certificates located under Marketing Application Requirements.

Clarivate

Beth Wise

Cortellis CMC Intelligence | Small Molecules | Pre-Approval

Export to Excel

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Cortellis

Physicochemical and biological properties

Accelerated Stability - Drug Substance

Long-term Stability - Drug Substance

Accelerated Stability - Drug Product

Long-term Stability - Drug Product

Marketing Authorization Application Requirements

Licenses and certificates

Accelerated registration procedure

Pediatric formulations

GMP certificate

Recognition of SRA or WHO PQ decision

Medicines Procurement

Licenses and certificates

Export

Compare key requirements across the selected

My selection	Application form	Proof of payment	Brand name clearance	Free sale certificate	GMP certificate	Submission of QOS	Business license	Manufacturing license	SM
Argentina	✓	✓	✗	✗	✓	✓	✗	✓	
Bolivia	✓	✓	✗	Not specified	✓	✓	✓	✓	
Brazil	✓	✓	✗	Not specified	✓	✓	✓	✓	
Colombia	✓	✓	Not specified	✓	✓	✓	✓	✓	N
Costa Rica	✓	✓	✓	✓	✓	✓	✓	✓	N
Guatemala	✓	✓	Not specified	✓	✓	✓	✓	✓	N

Select from the table of contents

Discover additional content by selecting Detailed or Report from the left hand menu bar or see the latests Updates or set Alerts.

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