

# Global Submission Planning at Bristol-Myers Squibb

BMS is transforming their operations with an authoritative source for regulatory information, including new processes for scheduling submissions and developing submission content plans.

**HEADQUARTERS**

BMS decides to submit a new application or expand a registered indication or formulation.

**UNIFIED RIM PLATFORM**

**AFFILIATES**

The new processes balance headquarters' knowledge and need for control with that of affiliates to improve global visibility and operating efficiency.

## Global Submission Plan

Headquarters creates a global submission planning record describing the business objective.



The Global Regulatory Team maintains a calendar of all BMS submissions globally.

Global Regulatory Team evaluates feasibility of affiliates' dates and establishes cross-functional alignment with clinical, CMC, manufacturing, regulatory operations and others on planned submission dates.



## Country Planning Records

Country-specific planning records capture affiliates' input into the corporate planning process.

Proposed dates reviewed by international strategy leads in headquarters.



- 1 Cross functional governance committee evaluates whether teams can deliver dossier by proposed date(s) before approving plan.
- 2 Governance-approved dates are entered into Global Submissions Plan and notifications sent to all involved.

**HIGH LEVEL SCHEDULING AND PLANNING**

GRANULAR CONTENT PLANNING BEGINS 12 MONTHS AHEAD OF THE APPROVED COMPLETION DATE

## Submission Documents

A dashboard shows completion status of each module as documents are authored, reviewed, and approved.

## Archived Dossier

The final published output is archived within the global RIM platform and linked to the original submission planning record.

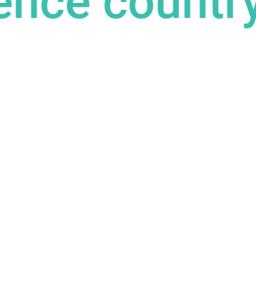
## Submission Content Plan

The RIM system auto-generates a submission content plan based on pre-defined parameters.

Personalized dashboards show planned dates for relevant submissions by month, country, or indication.



Tracking baseline, target, and approval dates for each component provides metrics for continuous process improvement.



Final content plan with approved documents provides basis for local dossiers in each reference country.