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# Example of Regulatory Strategy Job Description

Our growing company is searching for experienced candidates for the position of regulatory strategy. If you are looking for an exciting place to work, please take a look at the list of qualifications below.

## Responsibilities for regulatory strategy

* Participate in translating directives and regulations into practical and pragmatic business requirements and solutions for our clients
* Intervene as regulatory Subject Matter Expert on Regulatory Strategy assignments
* Participate to regulatory authorisation related assignments for Management Companies, Professionals of the Financial Sector or Banks
* Supporting executive decision making within the Regulatory affairs division, the Management Board by providing accurate insights and analysis to take strategic decisions within the regulatory space
* Serve as global regulatory lead accountable for all regulatory aspects of assigned project(s), including development and implementation of the global regulatory development plan for the project
* Represent GRA on cross-functional teams, including product development teams, study execution teams, clinical development task-forces and commercial sub-teams
* Serve as point of contact with regulatory authorities, contractors and corporate partners, where relevant, for specific projects
* Prepare, review and/or approve regulatory submission documents and Agency communications
* Conduct due diligence activities for potential partnerships and collaborations, as needed
* Contribute to the continuous improvement of existing department processes and strategies, providing recommendations in their area of expertise

## Qualifications for regulatory strategy

* Proficiency or aptitude in common database and data analytics languages
* Lead/Participate in Global Regulatory Teams (GRTs) Provide leadership to the global regulatory teams on critical project issues
* Assure priority market and non-priority market regulatory issues are addressed in development plans
* Provide strategic regulatory input to key development documents, including clinical protocols, clinical and nonclinical reports and summary documents, statistical analysis plans, DMC charters
* Review and approve key regulatory documents, including IB, IND, IMPDs, CTAs, RMP, Pediatric plans
* Support the preparation of, and participate in / lead (as appropriate), key HA interactions