At Astellas, we believe that nurturing exceptional relationships with our employees delivers exceptional business results.

Everyone at Astellas has a responsibility for creating a brighter future for patients around the world. From the first moment, Astellas will inspire you to put this ethos into practice – with a positive, agile company culture and with well-defined ethical principles, values, and systems.

Everything we do is led by our company values of integrity, being patient centric, taking ownership, delivering results, and communicating openly. These values are essential to Astellas' relationship with its employees and now is an exciting time to join us as we continue to evolve as a cutting-edge, value driven life sciences innovator.

Position: Intelligence Lead, RAPV Strategy
Department: RAPV Strategy
Line Manager: Head, RAPV Intelligence
Location: Beijing

Purpose and Scope

The RAPV Intelligence Lead, is responsible for gathering and disseminating regulatory intelligence and trends, delivering situational analyses, and making recommendations that encompass both the present regulatory landscape and the anticipated/desired future healthcare environment to bolster global and regional regulatory policy advocacy endeavors.

Responsibilities and Accountabilities

- **Monitor and disseminate regulatory intelligence**: Track upcoming legislation, analyze impacts, and share insights via blogs, alerts, presentations, webinars, and meetings.
- **Manage internal processes for guidelines**: Oversee handling of draft/final regulatory guidelines and coordinate input for consultancy in alignment with company strategy.
- **Maintain regulatory information systems**: Administer internal websites, databases, and electronic distribution tools for Regulatory Intelligence & Policy.

- **Provide expertise and representation**: Offer internal consultancy on regulatory interpretation, contribute to cross-functional initiatives, and represent the company in industry groups and public-private partnerships.
- **Support global policy advocacy**: Help develop company positions and action plans for regulatory and policy engagement worldwide.

Required Qualifications

- Health Science Diploma or equivalent.

Preferred Qualifications

- Significant experience within Regulatory Affairs, pharmaceutical industry, or related experience.
- Scientific knowledge and regulatory experience applicable to the role and solid basis in scientific approach, an ability to deal with technical information from a variety of disciplines.
- Experience in information searching and associated technology.
- Skilled networker and strong communication and collaboration skills.
- High integrity to maintain confidential and proprietary information.