Director Regulatory Policy & Intelligence

Job ID
REQ-10011630
Juil 03, 2024
Royaume-Uni de Grande-Bretagne et d'Irl. du Nord

Résumé

Our Development Team is guided by our purpose: to reimagine medicine to improve and extend people's lives. To do this, we are optimizing and strengthening our processes and ways of working. We are investing in new technologies and building specific therapeutic area and platform depth and capabilities – all to bring our medicines to patients even faster. We are seeking key talent, like you, to join us and help give people with disease and their families a brighter future to look forward to. Apply today and welcome to where we thrive together! The Role: This role offers hybrid working, requiring 3 days per week in our White City, London office. As Director, Regulatory Policy and Intelligence you will facilitate the application of regional regulatory requirements and intelligence to provide strategic direction and product advocacy in global development. This will ensure Regulatory Affairs enterprise Objectives are met, and enable regulatory compliance.

About the Role

Major accountabilities:

- You will provide regular communications and briefings to Global and Regional Policy Heads on relevant global regulatory policy issues, participating in internal product team meetings to provide regulatory policy context for business decisions.
- Provide strategic regulatory advice to RA colleagues on drug development projects and registration, and marketed products in preparation for HA meetings. This follows analysis of the impact of important emerging regulatory policies and new requirements on Novartis projects and business.
- Prepare and coordinate internal stakeholder feedback on proposed laws, regulations and guidance; represent and ensure Novartis' position is considered by trade associations.
- Serve as representative in external organizations such as trade associations, consortia, and/or Public Private Partnerships.
- Identify and escalate internally key information with stakeholders, collaborating with Corporate Affairs to shape legislative proposals.
- Coordinate and lead internal cross functional implementation of regulatory guidelines in new and emerging areas.

Your experience:

- Bachelor's or Master's Degree, science, health policy or legal based.

- Experience in regulatory affairs, and or Drug/ biologics development.
- Demonstrable success in a regulatory or health policy role, with strong understanding of the regulatory and legislative environment.
- Strong interpersonal, communication and negotiation skills. Ability to enable teams to think strategically.
- Experience of working in a complex, cross functional environment, within either industry or HA setting.
- Fluency in English.

Why Novartis: Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together?:https://www.novartis.com/about/strategy/people-and-culture

Commitment to Diversity & Inclusion:

Novartis is committed to building an outstanding, inclusive work environment and diverse team's representative of the patients and communities we serve.

Join our Novartis Network:

Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: https://talentnetwork.novartis.com/network

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Development
Business Unit
Innovative Medicines
Emplacement
Royaume-Uni de Grande-Bretagne et d'Irl. du Nord
Site
London (The Westworks)
Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.
Functional Area
Recherche & Développement
Job Type
Full time

Employment Type
Regular
Shift Work
No
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