Field Medical Advisor

Job ID
REQ-10036502
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Résumé

The Field Medical Advisor - -In line with overall product strategy- is responsible for supporting the design, implementation and execution of Medical Affairs plans for assigned Therapy Area, providing scientific information, helping design & organize clinical studies, building educational dialogue with KOLs and regulatory stakeholders.

About the Role

Major accountabilities:

- Lead/Support country medical affairs strategy in line with the global strategy, country insights and market conditions & secure implementation of planned Medical Affairs activities within the designated therapeutic area(s)
- Coordinate scientific meeting, symposia, congresses, Continuous Medical Education (CME) and other medical/scientific exchange and engagement activities which could bring additional value to the relevant therapeutic area; develop strategic engagement plan(s) for country customer-facing medical activities and events, and ensure execution of planned medical affairs
  activities in an efficient & compliant way, in a timely manner and within budget
- Lead/Support Pre-launch or launch activities in cross functional teams (Market Access, Commercial teams, Regulatory affairs, Brand team and others)
- Ensure medical enquiries are responded to in a high quality, timely manner, and in accordance with applicable standards; establish standard response documents as appropriate, for frequently asked questions
- Coordinate review and approval of medical materials and locally developed promotional materials; ensure medical materials provided from global or region for stakeholder engagement and events are tailored to local needs, and reviewed/approved per local/P3 guidelines
- Maintain an external focus, predominantly focus on field interactions/field related activities
- Establish and maintain peer to peer relationships with Medical Experts.. Support the medical community with up-to-date medical information, robust disease expertise and Novartis products' information.
- Contribute to mapping/profiling of MEs. Develop and implement ME engagement plans
- Support access for patients with unmet medical need to innovative treatments, ensure
- Managed access programs execution as per local policy
- Provide medical/scientific input into the development and execution of clinical trial or clinical research related activities, including initiation and oversight of clinical studies/clinical research within the respective therapeutic area.
- Support country strategy for Non Interventional Studies/Investigator Initiated Trial activities
- Clinical Development Support: Identify clinical investigators and facilitate participation at Novartis

sponsored clinical trials
- Support delivery of Pipeline: Enhance referrals and network, support clinical trials' initiation/ recruitment at the investigational sites, as part of a cross-functional team
- Ensure medical insights are provided to cross functional groups, including, but not restricted to: Pharmacovigilance, Regulatory affairs, Market Access, QA, Commercial teams, Brand team and others
- Build and maintain robust expertise in therapeutic area by continuous education, active monitoring of the scientific literature and by participation in national and international congresses and NVS scientific events
- Responsible for risk identification and assessment, mitigation planning as well as implementation and monitoring of appropriate internal controls within the area of responsibilities

Key performance indicators:

- Works within Ethics & Compliance policies
- Achievement of annual targets for medical activities

Minimum Requirements:

Education:

- Life Sciences or Science Degree. MD, PhD or MSc preferable
  Business degree (e.g. MBA) desired

Work Experience:

- Collaborating across boundaries.
- Operations Management and Execution.
- Project Management.

Skills:

- Building Construction.
- Clinical Practices.
- Clinical Research.
- Clinical Trials.
- Drug Development.
- Hazard Identification.
- Health Sciences.
- Immunology.
- Intensive Care UnIT (Icu).
- Internal Control.
- Internal Medicine.
- Job Description.
- Medical Information.
- Organization Skills.
- Patient Care.
- Stakeholder Engagement.
- Tcp/Ip Protocols.
- Utilization Management (Um).

Languages :

- English.

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International
Business Unit
Innovative Medicines
Emplacement
République de Chypre
Site
Cyprus
Company / Legal Entity
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Functional Area
Recherche & Développement
Job Type
Full time
Employment Type
Regular
Shift Work
No
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Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.

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