Lead, TA Medical (General Medicines)

Job ID
REQ-10004442

Apr 26, 2024

Japan

Summary

革新的なエビデンスや上市準備、市販後ソリューションに重点を置いた戦略的・運用TAのグローバル・メディカル・アフェアーズ・プログラムの開発と実施を支援し、患者、臨床、アクセス、医療システムの価値に関する戦略的な上市前および上市後の医療活動のニーズに対応し、提供するメディカル・アフェアーズ・プランニングやメディカル・サイエンス・エンゲージメント戦略の実行などを行う。設計中およびプログラムのエンドツーエンドの実行に沿って、医学的/科学的/証拠生成のインプットを提供します。統合エビデンス計画(IEP)/機能別プログラムの開発と実行にインプットを提供し、優先的な上市ポートフォリオの価値提案と医薬品の影響を最大化します。

About the Role

Role Purpose:
The Lead, TA Medical or Associate, TA Medical is accountable for building responsible brand or disease area medical strategy and ensuring its execution in collaboration with MSLs, MSEs and other partners with support from Head, TA Medical, Senior leads, TA Medical and/or Leads,

TA Medical. The Lead, TA Medical or Associate, TA Medical is the key project representative to commercial and GDD within Japan, and to global medical outside of Japan.
Four TAs - Haematology, Solid Tumour, Speciality Medicine and General Medicine.

Major Accountabilities

The Lead, TA Medical or Associate, TA Medical is accountable for all of the below items with support from Head, TA Medical, Senior leads, TA Medical and/or Leads, TA Medical.

- Bring clarity to the medical strategy and the tactics in the therapeutic area by their own medical/scientific knowledge/interpretation/judgement and deep understanding of external voice/environment as medical/scientific/clinical research expert.
  - Develop outcome-focusing TA Medical strategy aligned with TA strategy from both local and global points of view
  - Act as "Medical lead" in the agile project
  - Drive the actionable insight cycle by giving regular clear direction to MSLs on activities, messaging, targets, etc., and utilize collected insights to further refine the medical tactics
  - Work with PMO to shape the project work breakdown structure for the TA and to identify/secure required resources and budget of the teams
  - Input local needs into global strategy and collaborate with global teams on global project
  - Develop omnichannel engagement (OCE) plan by conducting omnichannel data analysis in collaboration with OCE capability pool and Medical Science group
- Lead/execute the evidence generation strategy in the responsible therapeutic areas as a medical/scientific/clinical research expert through scientific publications and innovative medical solutions in close collaboration with internal and external stakeholders.
  - Plan/lead/execute primary data collection (PDC) and secondary use data (SUD) studies including retrospective researches, prospective researches, biomarker researches, translational researches, database studies, etc. in collaboration with ME&E Dept.
  - Plan/execute Integrated Evidence generation Plan (IEP) by leading cross functional team
- Provide training for the organization he/she belongs to by leveraging highly advanced medical/scientific/clinical research expertise in the therapeutic area.
- Work as Patient Engagement Liaison (PEL).

Background

Education:

- Master's degree or equivalent experiences. Advanced science degree (MD, PhD, PharmD, MPH etc) strongly preferred

Languages:

- Japanese, Intermediary English

Experience/Professio
nal requirement:

- At Least 3-year agile project management not in a software development but in a business context
- Preferred (not essential): Knowledge of healthcare industry
- 3-year experience corresponding to MSE/SciComm or Marketing/JPH/JPCH

Competency
- Agile Project Management
- Strategic and logical thinking skill
- Scientific hypothesis construction skill
- Communication skill with external specialists
- Research publication skill
- Deep knowledge of TA
- Ability to synthesize complex requirements into clear specifications
- Strong relationship management and natural collaborator
- Cost Management skills

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部門
International

部門
Pharmaceuticals

国
Japan

勤務地
Head Office (Japan) (Pharmaceuticals)

Company / Legal Entity
JP05 (FCRS = JP005) Novartis Pharma K.K.

Functional Area
Research & Development

職種
Full time

雇用形態
Regular

Shift Work
No

Apply to Job.

利便性と合理的配慮

ノバルティス は 障害 を 持 つ 個人 と 協力 し、合理的配慮 を 提供 することをお 約束 します。健康状態 や 障害 を 理由 に 採用 プロセス のいかなる 部分 においても、あるいは 職務 の 必須事項 を 果 たすた めに 合理的配慮 が 必要 な 場合 は midcareer-r.japan@novartis.com 宛 てに 電子 メール をお 送 りください。その 際 ご 依頼内容、ご 連絡先、求人票 の 番号 を 明 してください。

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