

New Product Medical Lead

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Company: Novartis Farmacéutica

Location: Indonesia

Category: other-general

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 therapy 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 therapy area; develop strategic engagement plan(s) for country customer-facing medical activities and events, and ensure timely execution of planned medical affairs activities in an efficient and compliant way. 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. 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. 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. 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. Responsible for risk identification and assessment, mitigation planning as well as implementation and monitoring of appropriate internal controls within the area of responsibilities. Reporting of technical complaints / adverse events / special case

scenarios related to Novartis products within 24 hours of receipt -Distribution of marketing samples (where applicable) Major accountabilities: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 therapy 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 therapy area; develop strategic engagement plan(s) for country customer-facing medical activities and events, and ensure timely execution of planned medical affairs activities in an efficient and compliant way.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.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.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.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.Responsible for risk identification and assessment, mitigation planning as well as implementation and monitoring of appropriate internal controls within the area of responsibilities.Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt -Distribution of marketing samples (where applicable)Key performance indicators:Works within Ethics and Compliance policies -Achievement of annual targets for medical activitiesMinimum Requirements:Work Experience:Operations Management and Execution.Project Management.Collaborating across boundaries.Skills:Medical Governance.Third-Party Sponsored Trials.Medical Education and Scientific Engagement.Health Economics and Market Access.Non-Interventional Studies (NIS) / Epidemiology Studies.Medical Science and Disease Area Knowledge.STEAM - Applied Science, Technology, Engineering, Arts, Math.Clinical Trial Set-up, Management & Conduct.Clinical Trial Design, Data & Reporting.Preclinical Safety.Medical Safety.Languages :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> 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> Division International Business Unit Pharmaceuticals Location Indonesia Site Jakarta Company / Legal Entity ID03 (FCRS = ID003) PT Novartis Indonesia Functional Area Research & Development Job Type Full time Employment Type Regular Shift Work No 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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