究開発部門

医師募集 ： 治験担当責任者（Medical Doctor）

職務内容 【概要】

研究開発部門にて医学的・科学的見地から、医薬品開発計画や臨床試験実施における提言をする業務です。

部下を持つポジションではなく、部員として部長にレポートします。

【業務内容】

〇治験実施計画書（プロトコール）の作成

〇治験薬概要書、治験総括報告書のレビュー

〇重篤な有害事象レポートのレビュー

〇Key Opinion Leader（KOL）との関係構築

〇メディカルアドバイザー業務

登録資格 【登録資格】

-必須-

〇日本の医師免許証を有していること

〇日本での臨床経験 5 年以上

-尚可-

〇臨床医として治験参加などの新薬開発経験があれば尚可

〇専門領域不問、ただし、オンコロジー領域であれば尚可

【英語】

ビジネスレベル以上 （TOEIC 750 以上）

会話：流暢

読み書き：上級以上

＊グローバルプロジェクトの電話会議、メール。

プロトコールなどの英文ドキュメントの作成等

【人物像】

医薬品開発は社内関連部署や KOL との協力が不可欠であることから、受身ではなく積極的に仕事に取り組み、社内外関係者とうまくコミュニケーションができて信頼関係を構築できる方

待遇　【給与・賞与】

経験・能力を考慮の上、当社規定により算出します

【諸手当】

通勤手当、住宅手当、帰宅旅費手当、等

【昇給・賞与】

昇給　年1回、賞与　年１回

【休日・休暇】

完全週休2日制（土・日）、祝日、年末年始（12/29～1/4）、フレキシブル休日、5/1、年次有給休暇、慶弔休暇　他

【福利厚生】

各種社会保険完備、社宅制度、退職金制度、財形貯蓄制度、共済会　等

勤務地　東京本社（初台）

備考　資格　：　日本の医師免許を持つ日本人医師（外国の医師資格は不可）

年齢　：　30 代後半～50 才くらいまで

性別　：　不問

企業経験：　2-3 年

役職レベル：　経験スキルによりますが、MD である場合は基本的に管理職での採用で考えています。

募集理由　：欠員

【応募にあたって】

・応募時必須書類：職務経歴書（英文・和文）

※志望動機と現職での具体的な成果を必ず明記してください

・選考は 2-3 回を予定しております。また英語による面接がございます。

（遠方の場合は電話面接も検討いたします）

↑トップ

サノフィパスツールワクチン事業部

メディカル・マネジャー(Medical Manager)(パスツール) NEW !

職務内容 The Medical Manager (MM) is a medical/science based role whose main objectives are to:

1.capture, digest, update, summarize, and provide the vaccine related medical and scientific information to internal/external stakeholders (scientific communities, KOLs, regulatory authorities) with impact in the value proposition of the company assets.

2.exhibit a high degree of scientific and medical knowledge, i.e. a robust knowledge of diseases, vaccinology, product development, clinical trials, safety to drive the value proposition of the company assets.

3.organize and lead the medical advisory board of key opinion leaders to bring the answers and insight on clinical / medical questions for the company

4.exhibit a high degree of knowledge of healthcare and payer environment, medical practice and regulatory frameworks, public health priorities, and vaccine guideline development

5.facilitate timely, appropriate, balanced, credible, robust, long-term scientific exchanges with influential external experts and decision-makers to enhance the understanding of the scientific and medical value of our products with no sharing of product promotional information

6.drive and lead the clinical program of which medical function is responsible (post marketing and a product registered in other countries but new to local)

7.drive and lead the discussion/negotiation with regulatory authorities such as Ministry of Health.

8.develop robust evidence generation plans to enhance the

value of our products in a real-world setting
9.demonstrate operational understanding and risk management by ensuring excellence in execution of all governance processes; including Medical review and approval of promotional and non-promotional materials
10.develop insights that deepen our understanding of the needs of patients, consumers, regulators, payers, and healthcare providers and proactively identifies unmet needs
11.provide the training and education of medically related topics to internal customers including MSLs, and commercial teams
12.set the medical strategy with Medical Director and medical team

In addition, the MM is expected to build strong internal relationships with colleagues in Commercial Operations, while maintaining full compliance with all relevant company, industry, legal, and regulatory requirements. The insights gained by the MM while engaging with KOLs are to be shared with the broader organization in order to contribute to the enhancement of messages, plans, and strategy.

The MM will leverage existing educational slide decks developed internally and will follow all applicable internal procedures for the review and approval of any education material he/she adapts.

【Key Accountabilities】
■Drive Engages external stakeholders(Key Opinion Leaders; KOLs) on medical and scientific information exhibiting excellent scientific and clinical knowledge
■Drive and lead the clinical program of which medical function is responsible
■Supports evidence-generation activities by aiding in data acquisition and identifying opportunities for further

data collection

■Collaborates routinely and effectively with internal stakeholders

登録資格 [Education/experience]

M.D (plus Ph.D. is preferred)

Medical speciality in infectious diseases, vaccinology, immunology, public health, or paediatrics preferred.

TOTAL RELATED EXPERIENCE

・at least 3 years in clinical practice

・research experience in related areas is preferred

・experience in industry at least three years is preferred

[SKILLS/KNOWLEDGE]

The minimum required level (Basic–Competent – Proficient–Master) of each of the skills listed below to be determined.

■Technical competencies

Scientific Excellence: Proficient

Clinical Excellence: Competent

Operational Excellence & Risk Management: Proficient

Public Health Value–driven Access: Proficient

Lifecycle management: Competent

■Strategic/communication competencies

Business Insights: Proficient

Business Development: Basic

Communication Excellence: Master

[Launguage]

Japanese language proficiency: required

English proficiency: required

TOEIC >850

High level English laungage skills are required including speaking,writing, discussing, and presenting

待遇　【給与・賞与】

経験・能力を考慮の上、当社規定により算出します

【諸手当】

通勤手当、住宅手当、等

【昇給・賞与】

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【福利厚生】

各種社会保険完備、社宅制度、退職金制度、財形貯蓄制度、共済会　等

勤務地　東京本社（初台）

備考　【募集部署】

サノフィパスツールメディカル部メディカル・マネジャー

【レポートライン】

Country Medical Head, Sanofi Pateur

【年齢】

30 代～40 代前半

【その他】

●国内・海外ともに学会も含めて出張あり

●日本の医師免許保持者のみとなります

【入社時期】

ASAP

【応募にあたって】

・応募時必須書類：職務経歴書（英文・和文）

※志望動機と現職での具体的な成果を必ず明記してください

・選考は 3 回を予定しております。

※2・3 次面接は英語面接となります。

【Context of the Job】

○MM is a highly-trained professionals with strong scientific and medical backgrounds and technical expertise in the areas of vaccinology, product knowledge, product development, immunology and clinical development.

○MM may need to lead the clinical program in which medical function is in charge of, acting as a leader in a local program or as a local leader in global programs.

○MM may need to handle partnership business from medical perspectives, including clinical program and KOL handling.

○Liaised with MSL team, MM handles KOL's complex and controversial opinions; addresses and defends vaccine portfolio and corporate policy/ decisions, which may be controversial

○MM adhere to and work within clear compliance regulations and ensures the information shared with stakeholders is non-promotional in nature

○MM may need to handle adverse events, including addressing and defending corporate policy.

○MM has to work closely with Commercial partners while minding regulatory boundaries for non-promotion

○MM is required to capture and report information and insights appropriately, using available mechanisms and tools

○MM is expected to model core company values routinely.