

Medical Head Solid Tumors

Job ID
REQ-10034455
Ene 10, 2025
Italia

Resumen

~ Sviluppa e implementa programmi strategici e operativi di TA Global Medical Affairs, con particolare attenzione alle evidenze innovative e/o alla prontezza al lancio e/o alle soluzioni post-commercializzazione, compresa la pianificazione degli affari medici e l'esecuzione della strategia di coinvolgimento medico/scientifico che affronta e fornisce le esigenze strategiche delle attività mediche pre-lancio e lancio per i pazienti, cliniche, l'accesso e il valore ai sistemi sanitari

~ Fornisce competenze nello sviluppo e nell'esecuzione delle strategie generali, fornendo input durante la progettazione e lungo l'esecuzione end-to-end dei programmi

~Sviluppa ed esegue l'Integrated Evidence Plan (IEP)/programmi funzionali specifici per massimizzare la proposta di valore per il portafoglio di lanci prioritari e l'impatto dei nostri farmaci.

About the Role

Key Responsibilities:

Your responsibilities include, but are not limited to:

- Develop and implement the overall strategic direction for the Solid Tumors medical department in alignment with the goals and objectives of the broader oncology area and of the organization
- Continuously monitor industry trends, research advancements, and regulatory changes to adapt the strategy and ensure compliance with best practices
- Provides medical leadership, ensuring cross-functional alignment with the other teams inside and outside Medical Department to establish and execute strategic initiatives to improve patient outcomes and ensure the provision of cutting-edge Solid Tumors services.
- Supervise effective clinical development of products through accountability of the local medical plans for the Solid Tumors TA.
- Foster a culture of continuous learning, research, and innovation to drive advancements in Solid Tumors treatment options and care delivery.
- Lead scientific interactions with Top Medical Experts and Scientific Society of the Solid Tumors Area.
- Responsible for medical approval as per Doing Business Ethical (materials, events, grants as examples).
- Drive collaboration of the local medical team with Local, Regional and Global Teams in local Clinical Development activities and in Global and International Med Affairs activities
- Oversee together with Patient Safety local clinical trial adverse event reporting as well as clinical input and oversight into adherence to GCP together with SSO.
- Present highest ethical standards and contribute proactively to a credible reputation for Novartis CPO in the local Health Care and Medical community.
- Responsible for resource planning and management (FTEs and phase IV budget) within Solid Tumors

Therapeutic Area.

- May act as deputy to Medical Head Oncology

Essential Requirements:

- Medical Degree is required
- from 5 to 10 years of experience in Pharmaceutical Industry in roles of increasing responsibility, with significant senior leadership experience in Medical Affairs
- Experience in global or regional roles is desirable
- Demonstrated experience and ability to lead, engage and inspire science-driven teams and organizations of more than 10 people
- Strong cross functional collaborations skills
- Strong analytical and project management skills
- Experienced leading in a matrix environment
- Drive for innovation and continuous improvement
- Excellent communication skills
- Experience across multiple disease areas

Why Novartis? : Our purpose is to reimagine medicine to improve and extend people's lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <https://www.novartis.com/about/strategy/people-and-culture>

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International
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Italia

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Milano
Company / Legal Entity
IT08 (FCRS = IT008) Novartis Farma S.p.A.
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regolare
Shift Work
No
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