Medical Director Renal

Job ID
REQ-10028702
Nov 18, 2024
USA

About the Role

Major Accountabilities

- Lead scientific communication and publication for the renal portfolio: including multi-channel engagement plan.
- Drive and implement a diverse US renal council.
- Internal training lead (field).
- FUSE lead/reviewer.
- Represent the US medical voice at internal forums. Partner closely with US and Global stakeholders to provide US Medical input.
- Medical affairs renal portfolio safety POC.
- Mechanistic study US lead: site management, protocol compliance, etc.
- Lupus nephritis renal lead and cross therapeutic area POC.
- REMS lead.
- MSL POC for complement mediated kidney diseases.
- Serves as disease area medical expert for internal stakeholders from different line functions as well as external customers, including health care professionals, and patient advocacy groups.
- Provides medical scientific input for brand/program documents, including integrated disease area plans, Medical Information documents, Drug Safety reporting documents, etc. Ensures design and execution of all medical activities according to P3 compliance guidelines.

Education (minimum/desirable):

- Bachelors or equivalent 4-Year University Degree required.
- Doctorate level degree (MD, PharmD, DO, or PhD in Health Sciences or related field) or Nurse Practitioner (NP)/ Physician's Assistant (PA) degree with relevant clinical experience required.
- Medical Degree (MD) or equivalent preferred.

Experience required:

- 5 years' experience in progressively senior roles within clinical development and/or medical affairs roles in the biotech or pharmaceutical industry or academic institution/clinical practice.

Additional requirements

- US and European travel required. (20-25% annually, up to 30% seasonally).
- Location is flexible, but ideally in East Hanover, NJ.

The pay range for this position at commencement of employment is expected to be between $245,600.00 and

$368,400.00 per year; however, while salary ranges are effective from 1/1/24 through 12/31/24, fluctuations in the job market may necessitate adjustments to pay ranges during this period. Further, final pay determinations will depend on various factors, including, but not limited to geographical location, experience level, knowledge, skills and abilities. The total compensation package for this position may also include other elements, including a sign-on bonus, restricted stock units, and discretionary awards in addition to a full range of medical, financial, and/or other benefits (including 401(k) eligibility and various paid time off benefits, such as vacation, sick time, and parental leave), dependent on the position offered. Details of participation in these benefit plans will be provided if an employee receives an offer of employment. If hired, employee will be in an “at-will position” and the Company reserves the right to modify base salary (as well as any other discretionary payment or compensation program) at any time, including for reasons related to individual performance, Company or individual department/team performance, and market factors.

Role Requirements

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

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Division
US
Business Unit
Innovative Medicines
Location
USA
Site
Remote Position (USA)
Company / Legal Entity
U014 (FCRS = US014) Novartis Pharmaceuticals Corporation
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No
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