DESCRIPTION DE POSTE

Medical Advisor Endocrinology

Department	Medical Affairs
Supérieur immédiat	Global Medical Director Endocrinology
Gestion d'équipe	N/A

Position Summary

The Medical Advisor is responsible for the scientific integrity of the clinical development program and the translation of scientific information for tesamorelin into business concepts. The Medical Advisor will support the medical community and develop advocacy for the product. The Medical Advisor will also support the elaboration of abstracts and manuscripts for publication.

Key Responsibilities

- Responsibilities of the Medical Advisor Endocrinology are covering different therapeutic areas (including NASH and HIV) and also different life cycle management
- Provides deep medical/scientific expertise and medical affairs leadership and supports various functional areas.
- Provides expert product and therapeutic area medical expertise, as well as relevant scientific information in response to customer-stated needs.
- Develops customer advocacy to support brand business objectives and gathers insights to inform future brand strategies.
- Contributes to the brand strategy and develops a medical strategy to be executed as defined.
- Elaborates scientific rationale for medical and commercial projects and review new clinical study protocols and investigator-initiated research proposals.
- Participated to abstracts and manuscripts writing for publication.
- Manage clinical study(ies) and medical project(s).
- Supports the development of symposia and continuing medical education initiatives.
- Support clinical study feasibility, site mapping, identification and participation and provide support with resolving any pending issues with the study sites.
- Collaborate with the project manager to manage clinical studies and write clinical study documentation.
- Supports Regulatory Affairs and Pharmacovigilance department to establish program/product regulatory strategies that best support product medical and commercial objectives. Leads key cross-functional initiatives relating to Regulatory submissions.
- Be an internal scientific resource and trained commercial and medical colleagues.

Qualifications Required

- Doctoral degree (MD, PharmD, PhD) in a medical/scientific discipline
- At least 5 years in the pharmaceutical industry
- Experience in clinical research is an important asset

- Fluency in English, written and spoken, Spanish is an asset
- Good oral presentations skills, writing skills and aptitude for creating reports
- Autonomy and initiative
- Good team communication skills and team spirit
- Good planning and organizational skills
- Good leadership skills.
- Clinical trials/research experience a strong asset

Personal Qualities for all Staff

- Ability to communicate scientific and clinical information clearly and credibly verbally and in writing
- Interpersonal: ability to interact easily with all levels within the organization in a tactful, mature and flexible manner
- Must act ethically and with integrity
- High level of collaboration and influencing skills
- Ability to work independently and within cross-functional teams
- Excellent people leadership and organizational skills
- Ability to make sound and compliance guided judgments
- Must be a self-motivator with strong drive for result driven success
- Versatility and flexibility.

Current Incumbent: _____

Validated by : _____ Date : _____

Signature of direct supervisor: _____