Manufacturing and Environmental Regulatory Policy Officer

What inspires you to become Medicines for Europe's Manufacturing and Environmental Regulatory Policy Officer?

*Do you want to play an **important role in ensuring access to off-patent medicines in Europe?***
*Are you familiar with the **pharmaceutical sector?***
*Are Medicines for Europe's five pillars: **Patients • Quality • Value • Sustainability • Partnership** also your values?*
*If so, you might be our **Manufacturing and Environmental Regulatory Policy Officer.***

Why work for Medicines for Europe?

- Since 1993, Medicines for Europe has represented **pharmaceutical companies** supplying the **largest share of prescription medicines** across Europe.
- They are the voice of the **generic, biosimilar and value added medicines industries.**
- Leading partner for political change at an EU level and hence **better healthcare.**
- Increasing the health and wellbeing of all Europeans through **better access** to high quality medicines.
- The portfolios of Medicines for Europe members cover **80%** of therapy areas.
- Safeguarding the **sustainability** of Europe's healthcare systems for future generations.

https://youtu.be/nDQ4THE7mPo

Your Role

You will work closely with the **Policy and Regulatory Operations Manager in shaping legislature initiatives with manufacturing whilst maintaining an environmental focus**, whilst supporting both internal and external outreach.

1. Project and Workstream Support

- **Collaborate with the Policy and Regulatory Operations Manager** on manufacturing, environmental legislation, and anti-microbial resistance projects.
- **Provide crucial support** for ongoing Pharmaceutical Legislation initiatives.

2. Policy Monitoring and Advocacy

- **Monitor, analyse, and advise** on key policy, regulatory, and scientific areas.
- Research and stay abreast of policy developments, **contributing insights to regulatory and scientific discourse.**
- **Take ownership for coordinating committees, working groups, and task forces,** including agenda, minutes, reports, and briefing notes.
- Draft responses to **public consultations,** develop **position papers,** and conduct **surveys** among association members.

- **Collaborate with external partners** such as **the European Medicines Agency, Heads of Medicines Agencies, CMDh, and European Trade Associations,** advocating Medicines for Europe's stance.

3. Conference and Communication Contributions

- **Contribute to organizing conference sessions, including program development** for various meetings and events pertinent to your role.
- Assist in developing **communication and educational materials.**

Your Profile

- Master's degree in **Life Sciences or EU Law or Healthcare Policy.**
- Candidates **with or without experience** considered.
- Understanding of **European Pharmaceutical Framework, Market Dynamics and Policies at national level.**
- Proficient in **English.**
- Strong **communication, writing and presentation** skills.
- Comfortable in a **multicultural** environment
- Proficient in **Microsoft.**
- **Strategic thinking and creativity** to convey complex ideas clearly.
- Quick learner with **political savvy.**
- Excellent **project management** skills, with a proactive, problem-solving mindset.

Offer

- **Competitive Salary** + Meal vouchers (8€), Net remote work allowance, Hospitalisation insurance and Pension fund.
- Up to **2 days teleworking/week.**
- The possibility to **represent a dynamic European industry** and interact with diverse stakeholders.
- **European travel,** (20% of your time)
- The opportunity to take **ownership** and have **impact** as well as being part of a **motivated and multicultural team.**

Hiring process

- 1st and 2nd interview with the **Policy and Regulatory Operations Manager, the Deputy Director General, and the Head of Human Resources.**
- Decision after **10 working days.**

Join **Medicines for Europe,** be at the **forefront of shaping pharmaceutical policies** and make a **meaningful impact on healthcare and the environment.** If you're dynamic, proactive and ready to contribute, **apply now.**

GDPR Compliance

As a data subject, you have a number of rights. You can:

- access and obtain a copy of your data on request.

- require the association to change incorrect or incomplete data.
- require the association to delete or stop processing your data, for example where the data is no longer necessary for the purposes of processing.