

<b>Job Title:</b>	Manager, Regulatory Affairs	<b>Work Location:</b>	Greater Malé Region
<b>Division:</b>	Compliance	<b>Employment Type:</b>	Full-time
<b>Deadline:</b>	<b>14<sup>th</sup> June 2026, before 12:00 PM</b>		

**Key Responsibilities**

- Develop and implement regulatory strategies to support product registrations, renewals, variations, and market expansion.
- Oversee all regulatory submissions to ensure compliance with applicable local and international regulatory requirements.
- Act as the primary liaison with regulatory authorities, managing high-level communications, negotiations, and issue resolution.
- Provide strategic guidance on regulatory requirements and ensure alignment with business and product development plans.
- Monitor regulatory changes and lead the interpretation and implementation of new or updated regulatory requirements within the organization.
- Lead and support regulatory inspections and audits, ensuring organizational readiness and effective handling of authority interactions.
- Supervise, guide, and develop regulatory affairs team members to ensure high-quality submissions and strong departmental performance.
- Advise senior management on regulatory risks, approval pathways, and compliance-related decisions impacting business operations.

**Minimum Qualifications and Experience**

- Master’s degree (MQA Level 9) in Pharmaceutical Sciences, Biomedical Sciences, or a related discipline, with one (1) to two (2) years of relevant professional experience;  
OR
- Bachelor’s Degree (MQA Level 7 or 8) in Pharmaceutical Sciences, Biomedical Sciences, or a related discipline, with three (3) to five (5) years of relevant professional experience.
- Demonstrated experience in team coordination or supervision, with strong capability in regulatory authority interactions and ensuring compliance with regulatory requirements.

**Salary & Benefits**

- Remuneration between MVR 26,500 – MVR 28,500 based on qualifications and experience
- Other benefits governed by applicable laws and the Corporation’s policies

**How to Apply:** Interested candidates are invited to submit the following documents via email to [hr@statepharma.mv](mailto:hr@statepharma.mv)

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| • Updated Curriculum Vitae (CV)               | • MQA accredited and attested educational certificates |
| • Copy of National ID Card                    | • Experience letters and service records               |
| • Recent passport-size photograph (soft copy) |  |

Please use the subject line: **Application for Manager, Regulatory Affairs**

Incomplete applications and applications received after the deadline will not be accepted.

For queries, please contact Human Resources at [hr@statepharma.mv](mailto:hr@statepharma.mv)