

Job Title:	Senior Regulatory Affairs Officer	Work Location:	Greater Malé Region
Division:	Compliance	Employment Type:	Full-time
Deadline:	10th June 2026, before 12:00 PM		

Key Responsibilities

- Prepare, review, and submit comprehensive product registration and licensing applications to ensure compliance with all relevant regulatory frameworks.
- Maintain and update all regulatory dossiers, product documentation, and technical files to guarantee they reflect current global guidelines.
- Strategically track and manage lifecycle milestones, including registration timelines, upcoming renewals, and agency approvals to avoid business disruption.
- Lead and coordinate cross-functional responses to complex inquiries and information requests from regulatory authorities.
- Proactively monitor the validity and status of all operational permits, establishment licenses, and official certificates.
- Establish and maintain robust regulatory databases and physical or digital filing systems to ensure data integrity and easy accessibility.
- Provide critical support during regulatory agency inspections and internal compliance audits, ensuring all documentation is organized and audit ready.
- Perform any other duties as may be assigned by the Management.

Minimum Qualifications and Experience

- Bachelor’s degree (MQA Level 7 or 8) in Pharmaceutical Sciences, Biomedical Sciences, or a related discipline, with one (1) to three (3) years of relevant professional experience;
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- Diploma (MQA Level 5 or 6) in Pharmaceutical Sciences, Biomedical Sciences, or a related discipline, with four (4) to six (6) years of relevant professional experience.
- Experience in managing product registration activities, preparing regulatory documentation, and ensuring compliance with applicable regulatory requirements.

Salary & Benefits

- Remuneration between MVR 19,000 – MVR 20,200 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

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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 Senior Regulatory Affairs Officer**

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

For queries, please contact Human Resources at hr@statepharma.mv