# Notification of an outcome of the national registration provided by the participating company to WHO-PQT

### **Details of pharmaceutical company using the Procedure**[[1]](#footnote-1)

Company: Click here to enter text.

Country: Click here to enter text.

Address: Click here to enter text.

Focal point: Click here to enter text.

Telephone number (please include codes): Click here to enter text.

E-mail: Click here to enter text.

### **Details of medicinal product (the Product) subject to the Procedure**

Name of the Product: Click here to enter text.

Active principle(s): Click here to enter text.

Strength: Click here to enter text.

Dosage form: Click here to enter text.

### **Course of the Procedure**

Country: Click here to enter text.

Regulatory authority: Click here to enter text.

Date of submission of the application: Click here to enter text.

Date of acceptance of the application (if differs from submission date): Click here to enter text.

Date of issuance of a decision: Click here to enter text.

Length of process interruption/clock-stop (if applicable)[[2]](#footnote-2): Click here to enter text.

### **Decision on registration**

Granted, rejected, withdrawn: Click here to enter text.

Registration number (if applicable): Click here to enter text.

Registration granted in line with the SRA decision or with deviations, please comment:

 Click here to enter text.

### **Compliance with the Procedure, other observations and recommendations**

In the course of the Procedure following deviations were observed and recorded:

 Click here to enter text.

Any other observations and recommendations: Click here to enter text.

For the company

Signature:

Name: Click here to enter text.

Title: Click here to enter text.

Place and date: Click here to enter text.

1. Collaborative Procedure in Assessment and Accelerated National Registration of Pharmaceutical Products Approved by Stringent Regulatory Authorities – pilot assisted the by WHO Prequalification Team [↑](#footnote-ref-1)
2. Time provided by NMRA to the applicant to complete data or respond regulatory questions. [↑](#footnote-ref-2)