Invoice attestation for Import of registered finished Drugs

Process Description

Name of Process	Permission for Import of registered finished Drugs
Name of a business process	Apply for grant of permission to import of registered finished Drugs
Related laws, rules, and regulations	The Drug Regulatory Authority of Pakistan Act, 2012 The Drugs Act, 1976 The Drugs (Import & Export) Rules, 1976
Process participant	Importer DRAP
Input and criteria to enter/ begin the business process	Importer is licensed with Provincial Authorities (Drug Sale License) Having valid indent from importer having registration. Product is registered with DRAP
Activities and associated documentary requirements	Initiate Process for grant of permission to imported finished Drugs 1. Importer prepares the application form (on Form 1). Importer attaches supporting documents required as per the checklist. 2. Importer submits the application to DRAP office. 3. DRAP receives and scrutinizes the application form and supporting documents as per checklist. • If application is not compliant; DRAP returns the application with observation notes. 4. If application is compliant, DRAP verifies the documents. 5. If verification is successful, DRAP takes final decision and endorses/stamps the invoice. 6. Applicant receives the stamped invoice from DRAP. 7. If case is not approved, DRAP rejects the application and informs the applicant.
Output criteria to exit the business process	Stamped/Endorsed Invoice for import clearance of finished drugs. Validity: For particular consignment Legal Reference: Drug Regulatory Authority of Pakistan Act, 2012, The Drugs Act, 1976 & The Drugs (Import & Export) Rules, 1976
Time required to complete this business process	Minimum = 1 days Maximum = 2 days
Fee	No Fee