Grant of Drug Registration (Pharmaceutical drug for human use) (For complete Process Description, Checklist and relevant Forms Click Here) Manufacturer / Applicant **DRAP** Shortcoming letter issued Fill Drug (Pharmaceutical) Registration Application Non-Form 5F Compliant Receive Application Dossier Submit Fee Challan Technical Evaluation and Verification by Pharma ceutical Evaluation Cell Compliant Submit Application Dossier New Drug/Subsequent Generic (product Specific Inspection if Generic Drugs required) Registration Not approved Deferred **Board Meeting** Expert opinion (if Receive Rejection Rejection letter required)

Firm submits application within 60 days against decision of DRB to the

Receive Registration

Letter

Appellate Board.

Approved

Issue Registration Letter (Subject

to price fixation by Federal

Government)

Imported Drugs

Dosage form specific

inspection (if required)

Recommended

Locally Manufactured

Not

Recommended