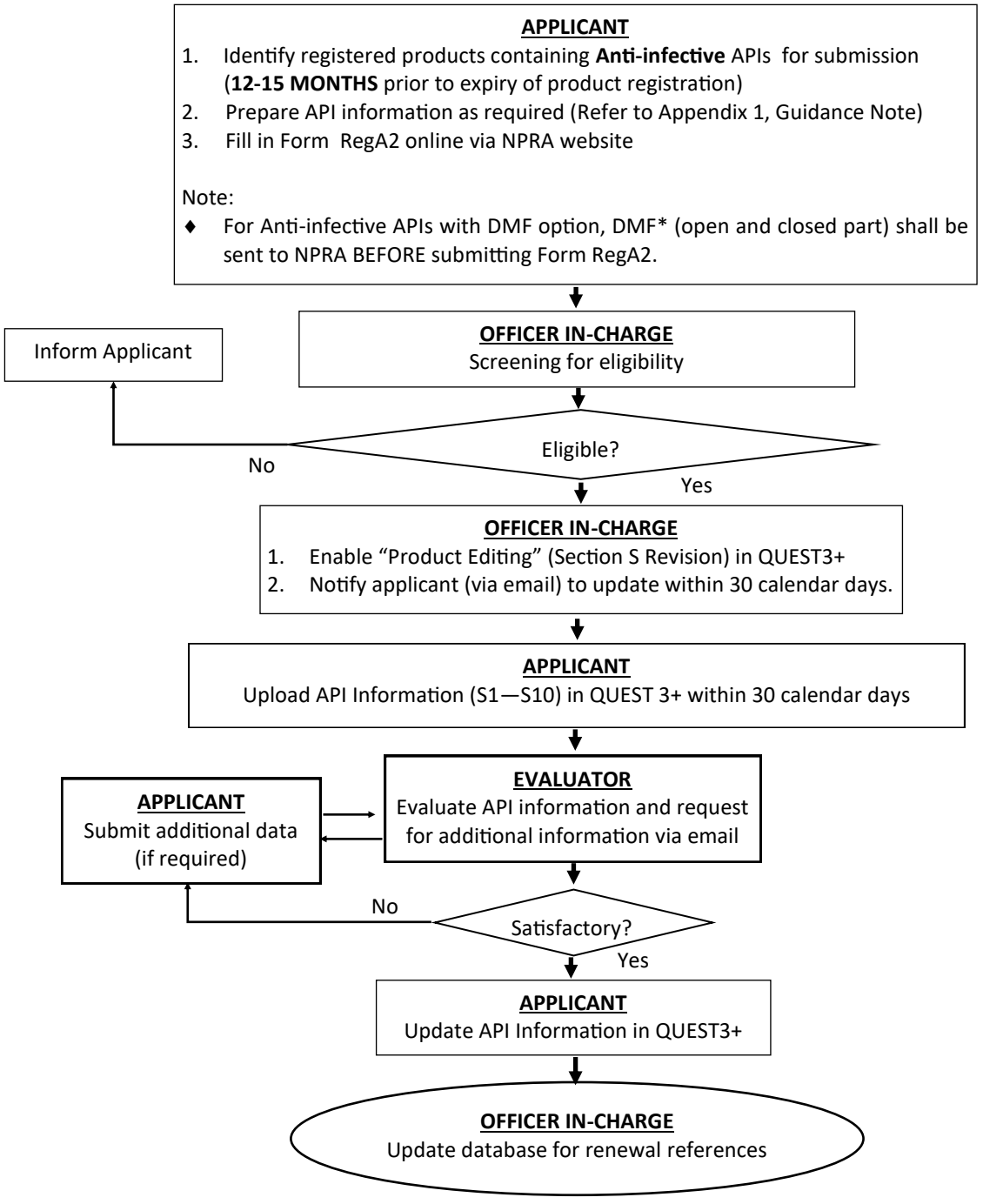
ADMINISTRATIVE PROCEDURE FOR REGULATORY CONTROL OF ACTIVE PHARMACEUTICAL INGREDIENT (API) IN REGISTERED PRODUCT CONTAINING ANTI INFECTIVE API



Footnote:

1. *CD copy of DMF (open and closed part) with a Letter of Access and Cover Letter should be sent to: *Head of New Drug Product Section/ *Head of Generic Medicines Section (*refer to product category)
2. **For registered products not containing anti-infective APIs, part II S information shall be kept by the PRH. It is not necessary to upload to Quest 3+ system.**