

### 3.2.P.5. Control of Drug Product [{Drug Product Name}, {Dosage Form}]

## 3. VALIDATION OF ANALYTICAL PROCEDURES [{DRUG PRODUCT NAME}, {DOSAGE FORM}]

Analytical validation information, including experimental data, for the analytical procedures listed in 3.2.P.5.2. should be provided.

Reference ICH guidances Q2A, Q2B, and Q6B.

Note to the Author: Please replace text in <<blue>> with hyperlinks to the relevant document as submitted with this section.

Validation information demonstrating that the analytical procedures provided in P.5.2 and used to test drug product are suitable for their intended purpose is provided in this section. Validation information for each analytical method included in P.5.2 for which validation is required is provided in the following order.

**Table 1: Validation reports for Analytical methods used to release {Drug Product Name}**

Test	Analytical Procedure	Attachment
		<<Validation Report>>
		<<Validation Report>>
		<<Validation Report>>