

LEACHABLES AND EXTRACTABLES HANDBOOK



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Equivalence Testing for Comparability BioPharm International

The United States Pharmacopeia (USP) chapter <1033> (2) indicates the preference for equivalence testing over significance testing: This is a standard statistical approach used to demonstrate conformance to expectation and is called an equivalence test. It should not be confused with the practice of performing a significance test, such as a t-test, which seeks to establish a difference from

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Method Validation Essentials Limit of Blank Limit of

Figure 2: Limit of blank (LOB), limit of detection (LOD), and limit of quantitation (LOQ) based on the blank.

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