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Committee for Human Medicinal Products

ICH guideline Q4B Annex 10 on evaluation and recommendation of pharmacopoeial texts for use in the ICH regions on polyacrylamide gel electrophoresis - general chapter

Step 5

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1. Introduction

This annex is the result of the Q4B process for the Polyacrylamide Gel Electrophoresis General Chapter.

The proposed texts were submitted by the Pharmacopoeial Discussion Group (PDG).

2. Q4B outcome

2.1. Analytical procedures

The ICH Steering Committee, based on the evaluation by the Q4B Expert Working Group (EWG), recommends that the official pharmacopoeial texts, the section in Ph.Eur. 2.2.31. Electrophoresis entitled Sodium Dodecyl Sulphate Polyacrylamide Gel Electrophoresis (SDS-PAGE), JP General Information 23. SDS-Polyacrylamide Gel Electrophoresis, and USP <1056> Biotechnology-derived Articles – Polyacrylamide Gel Electrophoresis, can be used as interchangeable in the ICH regions.

2.2. Acceptance criteria

The texts evaluated did not contain acceptance criteria.

3. Timing of annex implementation

When this annex is implemented (incorporated into the regulatory process at ICH Step 5) in a region, it can be used in that region. Timing might differ for each region.

4. Considerations for implementation

4.1. General consideration

When sponsors or manufacturers change their existing methods to the implemented Q4B-evaluated pharmacopoeial texts that are referenced in Section 2.1 of this annex, any change notification, variation, and/or prior approval procedures should be handled in accordance with established regional regulatory mechanisms pertaining to compendial changes.

4.2. FDA consideration

Based on the recommendation above, and with reference to the conditions set forth in this annex, the pharmacopoeial texts referenced in Section 2.1 of this annex can be considered interchangeable. However, FDA might request that a company demonstrate that the chosen method is acceptable and suitable for a specific material or product, irrespective of the origin of the method.

4.3. EU consideration

For the European Union, regulatory authorities can accept the reference in a marketing authorisation application, renewal or variation application citing the use of the corresponding text from another pharmacopoeia as referenced in Section 2.1, in accordance with the conditions set out in this annex, as fulfilling the requirements for compliance with the Ph. Eur. Chapter 2.2.31. on the basis of the declaration of interchangeability made above.

4.4. MHLW consideration

The pharmacopoeial texts referenced in Section 2.1 of this annex can be used as interchangeable in accordance with the conditions set out in this annex. Details of implementation requirements will be provided in the notification by MHLW when this annex is implemented.

4.5. Health Canada consideration

In Canada any of the pharmacopoeial texts cited in Section 2.1 of this annex and used in accordance with the conditions set out in this annex can be considered interchangeable.

5. References used for the Q4B evaluation

5.1 The PDG Stage 5B sign-off document:

Japanese Pharmacopoeial Forum, Volume 9, number 1 (January 2000).

5.2 The pharmacopoeial references for Residue on Ignition/Sulphated Ash for this annex are:

5.2.1 European Pharmacopoeia (Ph. Eur.): 6th Edition (official in January 2008), Electrophoresis (reference 01/2008:20231).

5.2.2 Japanese Pharmacopoeia (JP): The JP General Information 23. SDS-Polyacrylamide Gel Electrophoresis as it appears in the Japanese Pharmacopoeia Fifteenth Edition (March 31, 2006, The Ministry of Health, Labour and Welfare Ministerial Notification No. 285).

5.2.3 United States Pharmacopeia (USP): <1056> Biotechnology-derived Articles – Polyacrylamide Gel Electrophoresis official in USP 32, May 1, 2009.