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4 Concept paper on the need for revision of the guideline  $\dots$  30 June 2015  $\dots$  9  $\dots$  10 The proposed guideline will replace "Guideline on the Requirements to t he Chemical and  $\dots$  The new Regulation (EU) No. 536/2014 on clinical 22 trials on medicinal products for human use, repealing Directive 2001/20/EC will become applicable not

#### Concept paper on the need for Revision of the Guideline on ...

Evidence and consensus based guideline for the management of delirium, analgesia, and sedation in intensive care medicine. Revision 2015 (DAS-Guideline 2015) – short version S3-Leitlinie Analgesie, Sedierung und Delirmanagement in der Intensivmedizin.

#### Evidence and consensus based guideline for the management ...

EMA/CHMP/287710/2014 – Rev. 6 Page 3/15 Executive summary Based on the experience gathered by the Name Review Group (NRG) since the last revision of the guideline in December 2007, it became apparent that some areas of the guideline would benefit from

#### Guideline on the acceptability of names for human ...

Genetic toxicology Guidance Document Aug 31, 2015 17 542 4.1 Issues specific to in vitro TGs. 543 4.1.1 Cells. 544 In the revision to the in vitro TGs there is new guidance concerning the characterization and 545 handling/culturing of cells that are used in the individual tests.

#### Guidance Document on Revisions to OECD Genetic Toxicology ...

Health Guidelines Revision Committee. The Health Guidelines Revision Committee (HGRC) is a select multidisciplinary consensus body of about 100 clinicians, administrators, architects, engineers, and representatives from authorities having jurisdiction that is convened to revise and update the FGI Guidelines documents.

#### **Health Guidelines Revision Committee - FGI**

This guideline is a revision of the clinical practice guideline, "Diagnosis and Management of Bronchiolitis," published by the American Academy of Pediatrics in 2006. The guideline applies to

children from 1 through 23 months of age. Other exclusions are noted. Each key action statement indicates level of evidence, benefit-harm relationship, and level of recommendation.

# Clinical Practice Guideline: The Diagnosis, Management ...

A transcript of the conference proceedings and a copy of all written comments will be maintained in Docket ID No. EPA-HQ-OAR-2015-0310, which will remain open until October 27, 2015, for the purpose of receiving additional comments after the conference and the public hearing on the proposed revisions to the Guideline.

#### Federal Register :: Revision to the Guideline on Air ...

After the publication of the Annex 15 draft at the beginning of February 2014, the EMA made a move towards the revision of its process validation guideline. The final document was published on 27 February 2014. For a long time now, the EMA had already announced this revision in a concept paper. The objective of the revision was to integrate modern GMP aspects:

### **EMA publishes New Process Validation Guideline**

The revision of the guideline has allowed clarifying some inconsistencies, to revise the decision tree, to harmonize with Q3B and to address some editorial issues. The Attachment 2 of this guideline has been revised under Step 4 without further public consultation on 25 October 2006 (Q3A(R2)).

# **Quality Guidelines: ICH**

J Lab Med 2015; 39(1): 26–69 On 19 September 2014, the current version of the "Guideline of the German Medical Association on Quality Assurance in Medical Laboratory Examinations" was published. It featured an introduction by the German Medical Association. Revision of the "Guideline of the German Medical Association on Quality Assurance in

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