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Press Office

Guidelines and concept papers

Adopted during the CHMP meeting 17-20 September 2012

The guidelines and concept papers which have been adopted during this meeting of the Committee for Medicinal Products for Human Use (CHMP) will be published shortly on the European Medicines Agency's website under [Regulatory/Human/Scientific guidelines](#). Documents for public consultation will also be available under [Document search/Public consultations](#).

Biologics Working Party (BWP)

Reference number	Document	Status
EMA/CHMP/BWP/457920/2012	Guideline on the use of bovine serum in the manufacture of human biological medicinal products	3-month public consultation

Blood Products Working Party (BPWP)

Reference number	Document	Status
EMA/CHMP/BPWP/153137/2011	Reflection paper on immune tolerance induction in haemophilia A patients with inhibitors	3-month public consultation¹

Cardiovascular Working Party (CVWP)

Reference number	Document	Status
EMA/CHMP/494506/2012	Paediatric addendum to CHMP guideline on clinical investigation of medicinal products in the treatment of lipid disorders	Adopted

¹ This document was adopted at the July 2012 CHMP meeting



Reference number	Document	Status
EMA/CHMP/229823/2012	Concept paper on the need for revision of the guideline of medical products used in weight control	3-month public consultation
CHMP/EWP/2986/03 Rev.1	Guideline on clinical investigation of medicinal products for the treatment of acute heart failure	6-month public consultation

Central Nervous System Working Party (CNSWP)

Reference number	Document	Status
EMA/CHMP/40072/2010	Guideline on clinical investigation of medicinal products, including depot preparations in the treatment of schizophrenia	Adopted
EMA/CHMP/771815/2011	Guideline on clinical investigation of medicinal products for the treatment of Multiple Sclerosis	6-month public consultation

Rheumatology/Immunology Working Party

Reference number	Document	Status
EMA/CHMP/520782/2012	Concept paper on the need for revision of the guideline on clinical investigation of medicinal products for the treatment of juvenile idiopathic arthritis	3-month public consultation