30-12-2011

Submission of comments on '**Commission Guidelines on Good Distribution Practice of Medicinal Products for Human Use**'

Comments from:

Name of organisation or individual
Irén dr. Kocsi

Please note that these comments and the identity of the sender will be published unless a specific justified objection is received.

When completed, this form should be sent to the European Medicines Agency electronically, in Word format (not PDF).

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1. General comments

Stakeholder number *(To be completed by the Agency)*	General comment (if any)	Outcome (if applicable) *(To be completed by the Agency)*
	n/a	

2. Specific comments on text

Line number(s) of the relevant text *(e.g. Lines 20-23)*	Stakeholder number *(To be completed by the Agency)*	Comment and rationale; proposed changes *(If changes to the wording are suggested, they should be highlighted using 'track changes')*	Outcome *(To be completed by the Agency)*
Chapter 5; Section 5.17		Comment: To explain the term of "other products" in the glossary would be beneficial. Proposed change (if any):	
Chapter 9; Section 9.12		Comment: To explain the term of "transportation hubs" in the glossary would be beneficial. Proposed change (if any):	
n/a		Comment: n/a Proposed change (if any):	

Please add more rows if needed.