<Date of submission>

Submission of comments on Guideline on the acceptability of names for human medicinal products processed through the centralised procedure' (EMA/CHMP/287710/2014, Revision 7)

Comments from:

| Name of organisation or individual |
| --- |
|  |

*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).*

1. General comments

| Stakeholder number  *(To be completed by the Agency)* | General comment (if any) | Outcome (if applicable)  *(To be completed by the Agency)* |
| --- | --- | --- |
|  |  |  |

1. 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)* |
| --- | --- | --- | --- |
|  |  | Comment:  Proposed change (if any): |  |
|  |  | Comment:  Proposed change (if any): |  |
|  |  | Comment:  Proposed change (if any): |  |

Please add more rows if needed.