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|  | Annex 3 to the Procedure for Conducting Clinical Trials of Medicinal Products and Expert Evaluation of Materials Pertinent to Clinical Trials  |

(Section VII, 7.1 (7.1.2))

**Cover letter**

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| To the application for conducting clinical trial of medicinal product to be submitted to the central executive body: |  |
| To the application for getting approval of the Ethics Committee at HCS pertinent to the clinical trial of medicinal product: |  |

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(applicant)

submits for expert evaluation/ethical and legal aspects assessment

*(underline the necessary)*

the materials pertinent to clinical trial for getting conclusion/approval to conduct clinical trial in Ukraine:

 *(underline the necessary)*

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(full title of clinical trial)

ID of clinical trial protocol:

Investigational medicinal product (name):

Sponsor:

Other important information, if any: clinical trial characteristics, e.g., first-in-man study of new active substance, unusual investigational medicinal products, special trial subject groups, unusual design of clinical trial, trial subject fee or compensation terms for participation in clinical trial.

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| (applicant) | (full name in block letters) | (date) | (signature) |

*{annex in wording of the MoH Ukraine Order* [*№ 523 as of 12.07.2012*](http://zakon3.rada.gov.ua/laws/show/z1235-12/paran391#n391)*,* [*№ 966 as of 18.12.2014*](http://zakon3.rada.gov.ua/laws/show/z0062-15/paran15#n15)*}*