

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

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:

(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)

(full title of clinical trial)

ID of clinical trial protocol:

Investigational medicinal product (name):

Clinical trial 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.

(applicant)

(full name in block letters)

(date)

(signature)

{annex in wording of the MoH Ukraine Order № 523 as of 12.07.2012, № 966 as of 18.12.2014}