Performance schedule for the course Methodology of clinical research as a part of the Postgraduate Study “Medicine based on evidence”


Lectures

Prof. Eduard Vrdoljak, MD, PhD

P1 (09 00 - 09 45) – Drug development

P2 (09 50 - 10 35) – Planning of clinical trials

Ass.prof. Tihana Boraska Jelavic, MD, PhD

P3 (10 40 - 11 25) – The recruitment process and enrolment of subjects into clinical trials

Pause (11 25 - 11 55)

Seminars

Ass.prof. Marijo Boban, MD, PhD

S1 (11 55 - 12 40) – Safety procedures during clinical trials

S2 (12 45 – 13 30) – Safety basics during clinical trials

Recommended literature:


List of participants responsible for preparing the seminar:

ADAMOVIĆ IVAN

BLOSS HEINZ-GEORG


Ass.prof. Tomislav Omrcen, MD, PhD
S3 (09 00 - 09 45) – Monitoring in clinical trials

S4 (09 50 - 10 35) – Audit in clinical trials

S5 (10 40 - 11 25) – Inspection in clinical trials

Recommended literature:


2. Gobel C, Baier D, Rufhus B, Hundt F. GCP inspections in Germany and Europe following the implementation of the Directive 2001/20/EC. GMS German Medical Science 2009;7, ISSN 1612-3174


List of participants responsible for preparing the seminar:

GOEBEL HOLGER
MAYER DIRK
POHLIG CHRISTIAN

Pause (11 25 - 11 55)

Ass.prof. Branka Petric Mise, MD, PhD

S6 (11 55 - 12 40) – Clinical trial results

S7 (12 45 – 13 30) – Legal aspects of clinical trials

Recommended literature:


List of participants responsible for preparing the seminar:
FRANKE CHRISTIAN

ZINTL KONSTANTIN