COLOFOL

Instructions for participating departments

Country:

Name of department:

Address:

Department – ID:

Contact-person:

1. Preoperative informed consent for preoperative CEA and intraoperative blood and tissue sampling.

2. Postoperative data-reporting of all colorectal cancers with stage II and III disease under the local patient ID. Do **not** submit data until informed consent are obtained or declined after 4 weeks.

3. Written informed consent (to be kept in the department and with a copy in the patient file)

4. CEA after one month

5. Data-reporting of perioperative data 30 days after surgery (after the 4 week-CEA-test). Submit data. Randomisation-group and patient COLOFOL number will be reported back from the system

6. Eventually data-reporting after interval visits. Submit these data every time

7. Data-reporting after each planned follow-up visit. Submit these data

8. 60 months final status data to be reported

9. If you want to look at your own data, please:

   A: Use the patient-ID to look for all the data for the individual patients

   B: Go to the feedback-page and chose “Download data”. Then you can receive a file with all of your departments data, and a description.