

Using an ultrasound scanner (AU-590 Esaote-Biomedica), fine needles (Spinal-Icogamma, Ethanoject-TSK 22G), following the usual technique of alcoholization, varying amount of sterile alcohol (0.5–10 cc) at 95° were introduced into the lesions, for the total of 804 alcoholizations.

The response was evaluated by means of fine needle echo-guided biopsy (FNAB) carried out at the end of treatment: CR = necrobiosis and absence of CTM; PR = necrobiosis and rare, poorly-conserved, atypical CTM.

After the starting of PEI treatment (mean 4–9 months), 10 pts underwent surgery with exeresis of a total of 15 lesions. 21/24 pts responded to the treatment; of 33 treated lesions, we observed 6 CR (18.1%), 22 PR (66.6%), 3 NC and 2 local PRO. Histological confirmation of the cytologic response was obtained in 14/15 lesions undergone surgical exeresis. 10/24 pts died after a mean S of 13.5 months, in 9 cases due to extrahepatic disease progression.

Follow-up of the 14 alive pts is 17.5 months than S in 10 operated pts is 20.6 months, in 14 unoperated pts is 12.5 months. Overall survival is 15.8 months.

The comparison between our results and S reported in literature (6–12 months) comprehending any degree of hepatic involvement, is not possible since the criteria of admission are not homogeneous.

From a retrospective evaluation of 30 dead pts, followed in our service between '84 and '90 with liver metastases eligible for PEI but treated only with conventional CT, we observed a mean of S of 13.2 months; the mean follow up of our PEI-treated pts (17.5) has already surmounted that value, thus confirming the real efficacy of PEL.

The good result of mean follow-up is undoubtedly supported by the presence of pts undergone surgical exeresis. For this reason we calculated separately the mean S of unoperated pts, getting a result (13.2 months) next to the mean S of the survey group and expected to improve, since it regards still living pts.

In conclusion, we observed a real efficacy of PEI upon liver metastases, followed when possible by surgical exeresis; anyway PEI is a valid choice also in unoperable pts as palliative treatment in local control of disease.

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RANDOMIZED TRIAL OF INTRAOPERATIVE RADIOTHERAPY FOR COLORECTAL CANCER

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To evaluate the value of intraoperative radiotherapy (IORT) in colorectal cancer, a prospective randomized trial is carried out. For colon cancer, IORT plus surgery vs surgery alone is investigated. For rectal cancer, preoperative irradiation to the primary tumor was added to the treatment scheme in the experimental group. IORT was given in dose of 20 Gy, electron beam energy ranged from 8 to 15 MeV. After randomization and exclusion of unresectable and metastatic cases, the group consists of 42 patients. The mean age of patients was 56 years, 73% were males. Most part of the tumors presented B2 cases. There was no postoperative mortality. Approximately 1/2 of patients did not have postoperative complications. After a median follow-up period of 7 months (range 1–15) no patients in the experimental group had the disease recurrence. In the control group, 2 patients developed loco-regional failure.

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PALLIATIVE CYTOTOXIC THERAPY OF METASTATIC COLORECTAL CANCER IN GENERAL PRACTICE, A MODEL DESCRIBING THE COOPERATION BETWEEN THE MEDICAL ONCOLOGIST AND GENERAL PRACTITIONER

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Colorectal cancer is the most frequent diagnosed cancer in the western world. Less than 50% of the patients will today be cured of their disease. Thus, many patients will be in need of palliative treatment. The combination of 5-FU and Leucovorin has increased our ability to reduce tumour related symptoms by shrinking the tumour/metastases and not only mask them, e.g. with analgesics. However, it will not be feasible to administer such palliative oncologic treatment for this large patient group in the hospitals. In order to be able to offer this treatment to more patients, we have developed a program instructing general practitioners

how to administer the drugs and how to evaluate the effect and side-effects of the treatment. The patients are first seen at the outpatient unit at the Department of Oncology. The patient's physician is then contacted by telephone and asked if he/she can administer the treatment according to a written guideline which will be sent. The patient is sent the necessary prescriptions for the drugs and equipment. 5-FU is administered as a bolus injection day 1 and 2, and Leucovorin is given 30 minutes later the same days (according to a Nordic recommendation). The treatment is repeated every 2 weeks. Since the start of this project in early 1994, 21 patients have been treated by general practitioners. Altogether 340 courses of therapy have been administered. Ninety percent of the doctors have reported a positive experience both with regard to patient care and the practical side of carrying through the treatment. On the basis of the practical simplicity of this treatment procedure and the positive reports of general practitioners, this model of cooperation is recommended in order to give more patients with advanced metastatic colorectal cancer the possibility of symptom reducing anticancer palliative chemotherapy.

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5-FLUOROURACIL (5-FU) PLUS LEUCOVORIN VERSUS 5-FU LEUCOVORIN PLUS INTERFERON-A GIVEN AS ADJUVANT CHEMOTHERAPY IN COLON CANCER

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In the present study adjuvant chemotherapy of two drug combinations was given in colon cancer patients.

Material and treatment: One hundred and five patients were included. 98 were evaluable. The patients were randomized in two groups, A and B. There were 48 men and 50 women, 67 with stage Duke's C and 31 B2. Primary tumor was excised and histology was adenocarcinomas. Group A patients were treated with 5-fluorouracil 400 mg/m² and Leucovorin 30 mg/m² daily for 5 days every 4 weeks for 6 months. Group B patients had the same treatment plus interferon- α 5 mIU every 2d day for 6 months. Toxicity was common with mucositis and diarrhea and flu-like syndrome in Group B patients. **Results:** Median follow up was 38 months (6–58). Twenty-five patients recurred (25.5%). From group A the recurrence was 26.4% and from group B 24.4%. No statistical difference was observed. The median disease free survival was 14 months in either of the group. **Conclusions:** The combination of 5-Fu and leucovorin as adjuvant therapy in colon cancer seems to be effective and the addition of interferon- α did not reduce significantly the tumor recurrence.

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ISOLATED PELVIC PERFUSION FOR TREATMENT OF NON-RESECTABLE PELVIC RECURRENCES

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Twenty patients affected by non-resectable pelvic recurrence of primary pelvic malignant neoplasm were treated by Isolated Pelvic Perfusion. All patients were before treated by surgery and/or systemic chemotherapy and/or radiotherapy for their primary pelvic tumor. Patients underwent to laparotomy and IPP for 90 minutes at 40°C with high doses of drugs. Ten cases of rectal adenocarcinoma have been treated with Mitomycin-C (MM) and 5-FU and Mitoxantrone, in six cases of cervical carcinoma we employed a combination of MMC and Cis-Platin. One case of spindle cell anal carcinoma was treated with MMC, one case of penis carcinoma with Methotrexate, one case of endometrial adenocarcinoma with Cis-Platin and finally one case of melanoma of the vulva with L-PAM. Three CR and seven PR were observed; three patients had a stable disease and two other patients were no responders and died for progression in few months. One patient is lost to follow-up and four patients recently treated are not evaluable. The advantage in responsiveness and the low incidence of complications suggest a possible indication of this technique as a first approach for pelvic recurrences.

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