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Gastric cancer – How to improve results with a multimodal approach?

H. Wilke, H.-J. Meyer, M. Stahl. *Kliniken Essen-Mitte, Germany*

Since more than 20 years, attempts were made to improve the unchanged poor prognosis of gastric cancer (GC) patients by combining surgery (S), chemotherapy (CTx) and radiotherapy (RTx). Such approaches include preoperative CTx ± RTx, intraoperative RTx or postoperative CTx ± RTx. Until now, postoperative (adjuvant) CTx or CTx/RTx could not show that they really improve the results as compared to S alone. This is at least true for trials conducted in Western countries. However, this doesn't mean that postop. approaches are generally inactive, because in most trials CTx regimens were used which nowadays would be regarded as only moderately active. Moreover the patient populations accrued in these trials were heterogeneous and surgical approaches as well as surgical quality controls frequently not well defined. Intraoperative treatments such as IORT or i.p. CTx have also not yet demonstrated that they may contribute to a better outcome. Currently the most promising multimodal approach is preoperative CTx or CTx/RTx. Most published trials dealing with this issue were phase II trials in patients with clinically staged "potentially" resectable or clinically staged locally advanced (LAD) GC. These trials show that preoperative CTx is feasible, does not increase perioperative mortality and at least in LAD appears to improve R0 resection rates and survival. However well designed clinically trials based on an appropriate staging (endoscopic ultrasound, laparoscopy) are still lacking but urgently needed in order to better define the possible role of preop. CTx in this situation. In patients whose tumor was defined as unresectable during an explorative laparotomy, preop. CTx clearly demonstrated its efficacy. An R0 resection rate of approx. 40–50% and long-term survival of 15–20% were reported after preop. CTx. Also of note were the first reports of preop. CTx/RTx in proximal (cardia) gastric cancer indicating that this might become another step ahead in the management of this challenging tumor.

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Pancreatic cancer – Can we do better?

M. Wagner, C. Kulli, M.W. Büchler. *Department of Visceral and Transplantation Surgery, Inselspital, 3010 Bern, Switzerland*

Pancreatic cancer is the third leading neoplasm of the gastrointestinal system and represents one of the most aggressive human malignancies. Due to the late onset of clinical symptoms and the lack of accurate screening tests, the majority of patients are no more suitable for resection at time of diagnosis due to the presence of metastases or major infiltrations of the retroperitoneum. Oncological treatment modalities have failed so far to improve long-term results. Therefore, radical resection remains the only therapy with a chance for cure. For the surgical treatment of pancreatic head cancer, the classical Whipple operation is still the standard procedure but during the last two decades, the pylorus-preserving duodenopancreatectomy has been evolved as a more conservative procedure in order to omit the consequences of partial gastrectomy. For cancer of the pancreatic body and tail, distal pancreatectomy or total pancreatectomy may be indicated. With the advances in surgical technique and intensive care, the mortality rate has markedly decreased during the last two decades and averages around 5% in experienced centres. In contrast to these improvements, advances in long-term results are less obvious. A 5-year survival rate around 10% after radical resection of cancer of the pancreatic head is still considered the standard achievement and results for cancer of the pancreatic body and tail are even worse. So far, adjuvant treatment modalities have failed to demonstrate a significant benefit in a recent prospective and randomised trial. Moreover, more radical methods like regional pancreatectomy and resection with extended lymph node dissection could not improve long-term survival compared to the standard types of resection. For further advances in the treatment of pancreatic cancer, prospectively randomised trials are needed to compare these extended surgical procedures with the standard types of resection.

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Multimodal treatment in colon cancer: Recent advances and future prospects

M. Ducreux¹. *¹ Gastroenterology Unit, Gustave Roussy Inst., Villejuif, France*

Multimodal treatment is more and more frequent in the treatment of colon cancer, especially in adjuvant setting.

Adjuvant treatment: In approximately half of the patients treated surgically for colon cancer, incurable tumour recurrence can be expected. This has led to the development of adjuvant cytotoxic therapy. The role of 5FU + levamisole in stage III colon cancer was demonstrated in 1990. 5 years later, the combination of leucovorin + 5FU was proved to be effective with a significant reduction of mortality. The role of adjuvant systemic chemotherapy in lower-risk groups remains uncertain. Another way of multimodal adjuvant treatment of colon cancer is portal chemotherapy. After promising results, a meta-analysis and recent large studies have shown that the efficacy of intraportal chemotherapy was less than the efficacy of systemic chemotherapy. Question of its efficacy as a complementary method used in parallel with systemic chemotherapy is not solved. The future of multimodal adjuvant treatment of colon cancer can be broadly split into five categories: evaluation and reproducibility of surgical procedures, novel cytotoxic drugs: raltitrexed, oxaliplatin or irinotecan, specific or non specific immunotherapy, inhibitors of angiogenesis, gene therapy.

Metastatic disease: Recent results show that the resection of previously unresectable hepatic metastases became possible in up to 16% of patients after chemotherapy with oxaliplatin plus 5FU and leucovorin. Of the patients who had successful resections, 40% were alive at 5 years. After cytoreductive surgery of peritoneal carcinomatosis, intraoperative chemotherapy or chemohyperthermia and postoperative intraperitoneal chemotherapy have given promising results.

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Preoperative radiotherapy is better than postoperative in rectal cancer

L. Pahlman. *Department of Surgery, Colorectal unit, Uppsala University, Uppsala, Sweden*

Due to the high local recurrence rate in rectal cancer surgery, adjuvant treatment with radiotherapy has been proposed. However, the local recurrence rates differ substantially in literature, which might be one reason that adjuvant treatment is not obvious to all surgeons. This difference can be a matter of selection bias, different criteria for curative surgery, different follow-up routine, and/or the skill of the surgeon. Indisputable data from all randomized trials where surgery alone has been compared with surgery plus radiotherapy, either given pre- or postoperatively, is the average local recurrence rate of 29% in the surgery alone arm.

With pre- or postoperative radiotherapy the local recurrence rate is more or less halved. However, preoperative irradiation is more dose-efficient than the postoperative one, indicating that a higher dose has to be used if postoperative radiotherapy is delivered with an increased risk of damaging the normal surrounding tissues in the pelvis. This damage of the normal tissue also has an impact on sphincter function as well as on postoperative small bowel obstruction.

Due to the fact that preoperative radiotherapy is more effective on tumour cell kill at a lower dose than for postoperative treatment and with less toxic effects on normal tissues, preoperative radiotherapy should be used, if radiotherapy for rectal cancer is considered.

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A big bullet for a big tumour? – Locally advanced rectal cancer

K.M. Tveit. *Department of Oncology, Ullevaal University Hospital, Oslo, Norway*

Approximately 10% of primary rectal cancers are fixed to pelvic structures, making them unsuitable for primary surgery. Moreover, a considerable number of patients primarily operated for cure develop local recurrences which in most cases are located outside the rectum and fixed to the pelvic wall. These patients represent a real challenge to surgeons, oncologists and others. Treatment varies from simple symptom palliation via palliative radio- and chemotherapy to an aggressive curative approach including several treatment modalities.

Cure is possible to obtain in selected groups of patients, using a multidisciplinary approach. Thus, several reports conclude with 5-year survival rates of 40% and 20%, in primary and recurrent cases, respectively, provided a "radical" resection could be performed. In these studies preoperative external radiotherapy at a dose of 45–54 Gy (1.8–2.0 Gy/fraction) was combined with extensive surgery, in many cases also involving prostate, bladder and internal genitals. Less documentation exist regarding quality of life of these patients.

Chemotherapy in addition to radiotherapy and surgery is employed in many institutions, concomitantly with the irradiation and/or postoperatively,