Heavy-Ion Therapy at GSI

Gerhard Kraft from the Gesellschaft für Schwerionenforschung, Darmstadt, describes how new approaches and a novel scanning technique to be implemented at GSI’s planned ion-therapy unit will offer biologically efficient, tumour conform treatment.

Beams of heavy charged particles beginning with protons represent the most advanced tools for external, subcutaneous therapy of deep-seated tumours in humans. Compared to electromagnetic radiation, they offer an improved distribution of the dose with depth owing to their heavier and more regular scattering; they travel in virtually straight lines and stop at a definite depth (or “range”). More importantly, the dose profile is inverted in that the energy deposition increases from a plateau with increasing penetration distance up to the Bragg maximum, and then cuts off sharply within a few millimetres. By combining overlapping Bragg peaks it is possible to achieve a homogeneous energy deposition over a defined volume (Fig. 1).

The lateral and range scattering decreases with the square of the atomic number so the precision of beam delivery increases as the atomic number increases. But the nuclear interaction rate also increases yielding lighter nuclear fragments having a longer range than the primary particles. An optimum is reached for ions around carbon where lateral scattering is small and nuclear fragmentation tolerable [1]. Moreover, nuclear fragmentation of ions heavier than protons and helium has strongly forward-peaked reaction kinematics. By measuring the γ-annihilation quanta of β+-radioactive ions using positron emission tomography (PET) one can therefore correlate the range distribution with the primary beam in order to localize on-line the beam inside the patient [2]. The increased biological efficiency of heavy ions providing high ionization density in certain types of tumours opens up new dimensions in the treatment of radio-resistant tumours.

HEAVY-ION THERAPY UNIT AT GSI

The creation of a heavy-ion therapy unit at GSI (HITAG) was proposed to the German government in May 1993 by the GSI together with the Heidelberg Radiological Clinic and the German Cancer Research Centre, both these institutes having a long tradition in conventional therapies, neutron therapy and the development of advanced treatment techniques. The proposed 13.2 MDM unit (some with 60% coming from government and other external sources including insurance companies) will consist of a dedicated therapy department and an annex housing a control room, waiting rooms and the like (diagnosis and treatment planning will be in Heidelberg).

The ion beam from the beam line is guided by two dipole magnets and a pair of focussing quadrupoles. Symmetry and achromatism of the beam line is important for safety because the final focus in the patient corresponds to a focus intermediate between the two dipole magnets. Scintillators used as a beam scratcher at the intermediate focus allow the beam to be controlled nondestructively. Other safety features include an asymmetric layout for the raster scanning magnets so that the beam passes above the patient when the magnets are not powered (in the case of a power failure, the beam will automatically move outside the patient). A high-speed position sensitive counter just in front of the patient will monitor on-line the beam position and compare it with the settings of the scanner magnets in less than a millisecond. Finally, the PET system mentioned in the text will monitor the location of the beam inside the patient.

The cave is under construction and most of the beam-line components have been ordered; construction of the unit will begin early in 1996. An extended experimental campaign to characterize carbon and oxygen beams both biologically and physically will allow a choice between the beams resting in the treatment cave, including exposure to anthropomorphic phantoms, will start in 1995 in preparation for the start of therapy in 1996. The aim is to treat about 100 patients a year suffering from local and inoperable cancers, mostly in the head and neck regions.

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Fig. 1 — The superimposition of Bragg curves to give a uniform dose over a spherical volume. For each Bragg curve, the ionization density rises from a plateau value as the ions slow down, until at the very end of the ion's range the ionic charge is reduced by electron pick-up and the ionization falls rapidly to zero.
tumours. In conventional therapy and therapy with ions, the rate at which tumour cells are killed is largely governed by the cells' capacity to repair themselves. The most important target for cell inactivation is the DNA molecule: it carries all the genetic information needed for cell functioning. Most of the radiation-induced damage to DNA can be repaired by the cells as long as the individual DNA lesions are separated. If local damage is increased, the chances of a correct repair are drastically reduced. Comparing the track structures for carbon ions and for protons with the size of the DNA molecule (Fig. 1) shows that carbon is biologically more efficient as it is more likely to produce correlated, unreparable damage within a single ion track than protons of the same energy.

Ideally, a beam for ion therapy should combine this high biological efficiency in the tumour volume with the creation of most of the treatable damage in the entrance channel in front of the tumour where healthy tissue is present. Recent experiments at GSI have demonstrated that carbon and oxygen ions are very suitable because the biological efficiency increases from a low value in the entrance channel to a four-times higher value at the end of the tumour. But in order to exploit this advantage of ion beams it is essential to adjust the beam so that the range of the particles is restricted to the target volume. This means that strictly conforming the irradiation is the most important requirement in using ions heavier than protons for radiotherapy.

Tumour Conform Irradiation

In all the particle therapies presently in operation the beam is distributed over the target volume using passive beam-forming elements. These are expensive and inadequate for charged-particle beams that can be manipulated using magnetic fields. Magnetic scanning has thus been introduced into ion therapy \[4\]. The target volume is dissected into slices of equal range and each slice is treated independently by scanning over the slice; once one slice has been treated, the beam energy and consequently the particle range is reduced and the next slice is treated.

Depending on the target geometry, parts of the tumour are pre-irradiated during the treatment of outer regions. The additional irradiation needed to achieve a homogeneous biological effect over the entire treatment volume must thus be distributed inhomogeneously over a given slice. Two strategies have been discussed, namely pixel scanning and raster scanning \[6\], involving discrete or continuous motion, respectively, of the beam. The differences are largely academic and essentially revolve around the issue of whether or not the beam needs to be turned off while moving from one point to the next. In practice, the beam diameter is several millimetres so that the distance between two pixels is small compared to the diameter, the beam need not be interrupted on moving from pixel to pixel (overexposure at a pixel can be largely compensated during irradiation at the next one). GSI has set up a digitized version of an intensity-controlled, raster scanning system (Fig. 3). The use of a transmission counter to control the writing velocity has been demonstrated to be a very elegant and efficient method for conform treatment and for coping with fluctuations in the beam intensity. In preliminary experiments, a spherical volume of water has been successfully irradiated to a constant dose (Fig. 4).

Computer Modelling

The extended region of high biological efficiency in the Bragg region for ions such as carbon and oxygen will be restricted to the target volume using the novel technique of fast magnetic scanning with active energy variation. While this will open up new opportunities in conformal tumour therapy, it involves a quantum leap in the complexity of beam control and treatment planning. This is because the dose at various points in the target volume comprises a mixture of doses originating from the plateau and Bragg peak ions having very different biological efficiencies, depending on their atomic number and beam energy. A model \[6\] for the biological action is essential as it is impossible to measure all the necessary data. It is only because of a deeper understanding of physical and biological properties of charged particles that it will be possible to use focussed ion beams in the same way as a surgeon uses a scalpel.

The GSI Therapy Unit

Tumour conform treatment has not been implemented up to now owing to the poor time-structure of extracted beams coming from accelerators designed for physics experiments. Indeed, heavy particles such as neon ions were found to give no significant advantages over protons. The passive methods of beam shaping adopted from X-ray therapy which were used did not allow the irradiation to completely conform to a tumour so the maximum dose necessary for controlling the tumour could not be specified. A new facility at GSI plans to eliminate these shortcomings in treating deep-seated, inoperable cancer tumours by using the novel technique of three-dimensional tumour conforming irradiation with the optimization of biological advantages using computer modelling. The Heavy-Ion Therapy Unit at GSI (HITAG; see insert), which was strongly endorsed by Dr. P. Krüger, Germany's science minister, during a visit to GSI last month, will use ions between carbon and oxygen at energies of 300-500 MeV per nucleon coming from the heavy-ion synchrotron SIS that can accelerate all particles from protons up to uranium ions to 1 GeV per nucleon or more. A long-standing research programme at GSI has already characterised the beam structure and properties and acquired an extensive understanding of radiobiological aspects.

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6. Scholz M. & Kraft G., in [3].

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**Fig. 2** — The tracks for protons (left) and carbon ions (right) compared with the dimensions of a DNA molecule shown schematically at the centre. The tracks were measured in water using 1 MeV per nucleon projectiles. Carbon ions have a much higher probability to produce breaks simultaneously in two opposite strands of DNA \[3\].