## MedAustron

# WP2: <br> 11Carbon PET-aided hadron therapy 

Research objectives
ESR positions
Secondments
L. Penescu, MedAustron

On behalf of WP2

## WP2 Scope. Presentation summary.

Scope of WP2: 11Carbon PET-aided hadion therapy
$\checkmark$ Production and mass separation of ${ }^{11} \mathrm{CO}^{+}$.
(ESR11)
$\checkmark$ Development of charge breeding scheme required for acceleration.
(ESR3)
$\checkmark$ Full acceleration and treatment $-{ }^{11} \mathrm{C}$ hadron therapy test and planning.
$\checkmark$ Further development of bioconjugates suitable for imaging and treatment of the ovarian cancer. New bifunctional fluorescent and radioactive bioligand. Tests with 11C as chelate.
(ESR15)
$\checkmark$ Development of multimodal imaging methodologies for the treatment planning.
Methodology and preclinical techniques. Biological models.
(ESR12)

## PARTNERS

| ESR11: University of Leuven (Recrun | Enrolment; 36 mo ) | + CNAO (Secondment; 10 mo ) |
| :---: | :---: | :---: |
| ESR3: CERN (Recruiting; 36 mo ) | + Chalmers University of Technology (Enrolment) | + MedAustron (Secondment; 5 mo ) |
| ESR9: CNAO (Recruiting; 36 mo ) | + University of Pavia (Enrolment) | + CERN, HUG (Secondment; $3+3 \mathrm{mo}$ ) |
| ESR15: HUG (Recruiting; 36 mo ) | + EPFL (Enrolment) | + AAA (Secondment; 3 mo ) |
| ESR12: HUG (Recruiting; 36 mo ) | + University of Geneva (Enrolment) | + |

Presentation SUMMMARY
> Delivering a beam for treatment at an existing facility (MedAustron, CNAO, HIT).
$>$ Possibilities for mixing/replacing ${ }^{12} \mathrm{C}$ with ${ }^{11} \mathrm{C}$, for treatment and PET imaging.
$>$ Open questions (to be addressed by the ESRs).

# PIMMS-based accelerators: HIT, CNAO, MedAustron 

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PIMMS: Proton-Ion Medical Machine Study, CERN 2000-006


## The 12C beam path at MedAustron



## Requesting a beam

Treatment plan = a sequence of cycle codes

- Particle type
- Ion Source.
- Beam size
- Variant.
- Irradiation room
- Version.
- MEBT degrader used
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Mask


Applied magnet current

000.T00.000.001.0000 (Protons; Variant T)



## Delivering 2 beam species

1 beam species, 1 plane


- 3D matrix of intensities defined by treatment plan
- Point by point scanning of slices
- A spill can cover a full slice
- From high E to low E


## 1 beam species, 2 planes (H and V)

- Treatment plan with dose splitting between planes.
- Different penetration depths (=energies) for the 2 planes.
- Full scan in 1 plane, followed by full scan in $2^{\text {nd }}$ plane.
- Scan in 1 plane as above.
(hysteresis of cw magnets)
- Scan in $2^{\text {nd }}$ plane following same strategy as above.


## 12C and/or 11C <br> $\checkmark$ Treatment <br> $\checkmark$ PET imaging

2 beam species, 1 (or 2) planes

- Treatment plan with dose splitting between species.
- Different penetration depths (=energies) for the 2 species.
- Full scan in 1 plane, followed by full scan in $2^{\text {nd }}$ plane.
- Scan in 1 plane as above.
- (hysteresis of cw magnets)
- Scan in $2^{\text {nd }}$ plane following same strategy as above.


## The accelerator side: commissioning a "cycle code"

## The accelerator is able to generate:

$\checkmark$ Number of ion species: 2
$\checkmark$ Number of different energies: 255
$\checkmark$ Number of beam sizes: 4
$\checkmark$ Number of intensities: 4
$\checkmark$ Number of extraction times: 8
> Beam combinations per beam line: 65280
> Gantry: different angles need to be considered
> Non-clinical research: extended energy range

PARAMETERS which are NOT part of cycle code, but part of design + commissioning:
$>$ Energy spread
$>$ Spill quality (stability: intensity, position)
$>$ Choices of beam optics (longitudinal and transversal)

DURATION OF A CYCLE - depends on accelerator performance:
$\rightarrow$ Injection time
$\Rightarrow$ Acceleration time
> Short hysteresis time
> Hardware configuration time

## Preparation and QA of a medical beam

## MedAustron ${ }^{\text {W }}$

(A) Same applied settings


Reproducible BEAM
(B) Reproducible machine performance

Understanding of:

- Hardware stability
- Hardware - Hardware interactions
- HW - Beam interactions
- Beam - Beam interactions
- Aging effects
- Maintenance effects
- Failure statistics
$\checkmark$ Spill formation: defined in the synchrotron
- Duration.
- Intensity uniformity.
- Position uniformity.
$\checkmark$ Spot formation: defined in the HEBT line
- Spot size
- Position at isocenter
$\checkmark$ Restrictions on injector: not critical
- Most of the injected beam limitations can be corrected in the synchrotron


## Question marks vs. ESR topics

1) How to produce the 11C with needed intensity, reproducibility and stability?
2) Where to inject the 11C along the beam path? (answer correlated to $1 \ldots$ )
3) PET-related questions...
4) Compact PET cyclotron $\rightarrow \mathrm{N} 2$ target $\rightarrow$ release of 11CO2 into ion source $\rightarrow$ beam path joining 12C
5) ISOL production $\rightarrow$ mass separation and charge breeding ( $\rightarrow$ post-acceleration?) $\rightarrow$ beam path joining 12C
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(ESR11)
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Methodology and preclinical techniques. Biological models.
(ESR12)

## TO HAVE IN MIND: The available intensity per spill is not a veto-condition.

Accelerator efficiency depends on all of the following:

- Intensity per spill
- Cycle time (acceleration time; hysteresis; re/configuration; injection of 11C?)
- Strategy for using effectively all accelerated particles.
- Dealing successfully with machine limitations at higher intensities.

