



# **Biological dose calculation on dual-layered DECT images for CIRT**

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II. Method

- Dual energy CT (DECT)

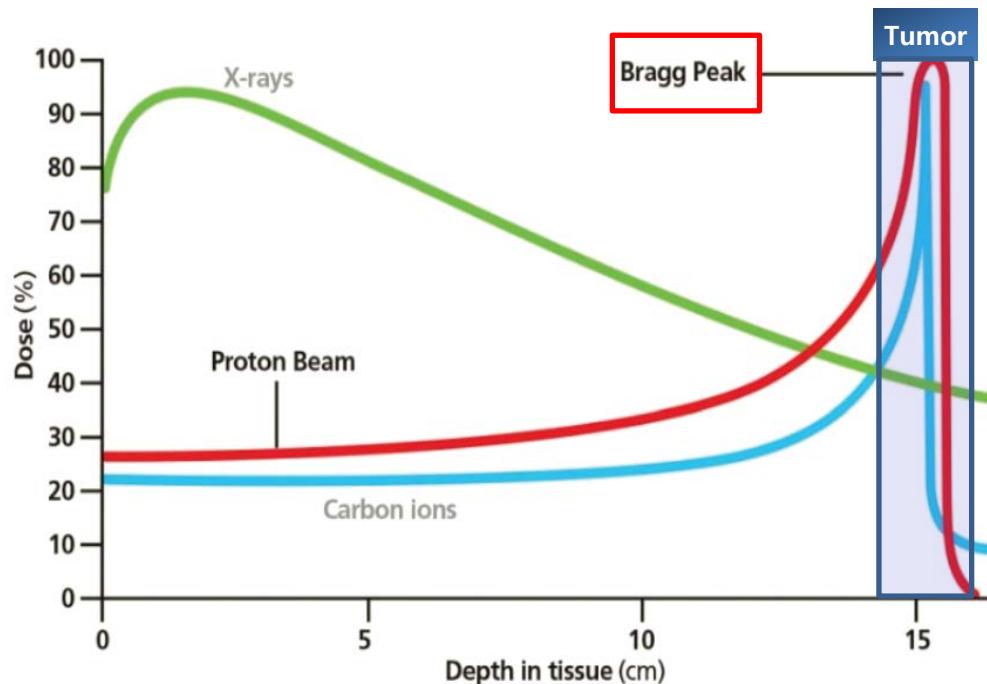
- Relative Biological Effectiveness calculation

III. Results

IV. Discussion & Conclusion

# I. Introduction

## Ion therapy



B. Mustapha, et al. 2016

### Bragg peak

- Reduction of exposure to normal tissues

### Ion therapy

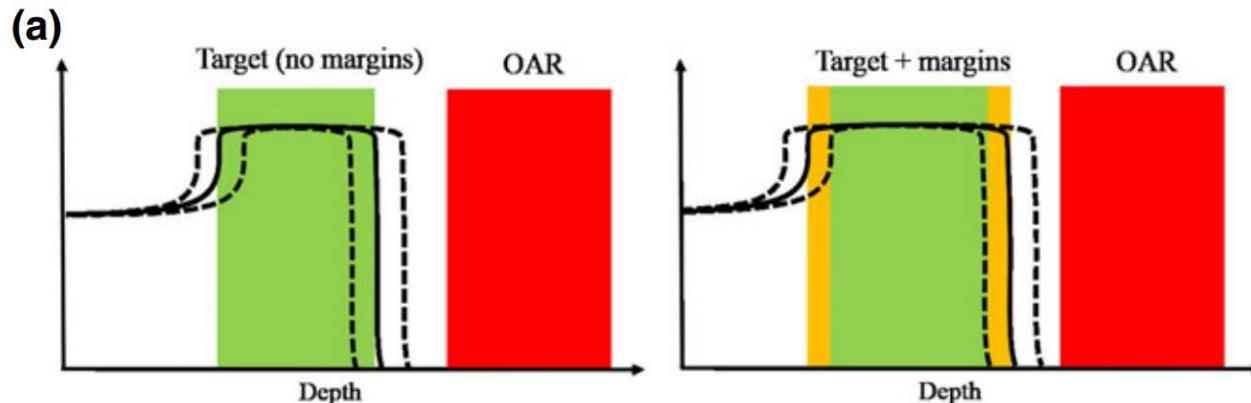
- proton, Carbon,  $\alpha$ -particles, ions

# I. Introduction

## Proton therapy

Range Uncertainty exists (2012, paganetti)

- Organ motion, setup and anatomical variation, dose calculation approximation, biological consideration
- Range margin : 3.5 % + 1 mm (MGH)
  - ex) 20 cm range field → 8 mm margin

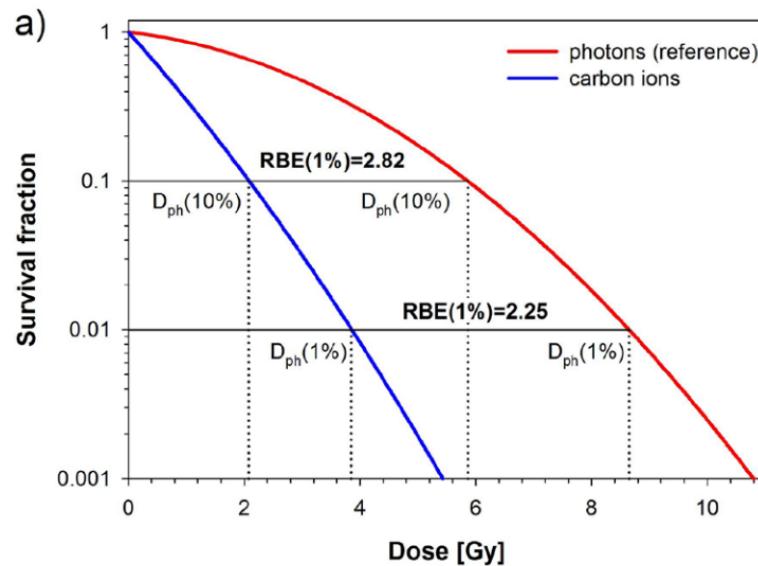


# I. Introduction

## Carbon ion therapy

Difference with proton therapy

- larger **Relative Biological Effectiveness (RBE)**



$$RBE = \frac{D_{ph}}{D_{ion}} \Big|_{\text{isoeffect}}$$

☞ **Biological dose (BD) = RBE × Physical dose**

Karger, et al. 2017

Carbon Ion Treatment planning is based on **biological dose**

# I. Introduction

For successful carbon ion treatment?

**Accurate Dose calculation**

Monte Carlo  
simulation

Reduction  
of range  
uncertainty



**Dual energy CT (DECT)**

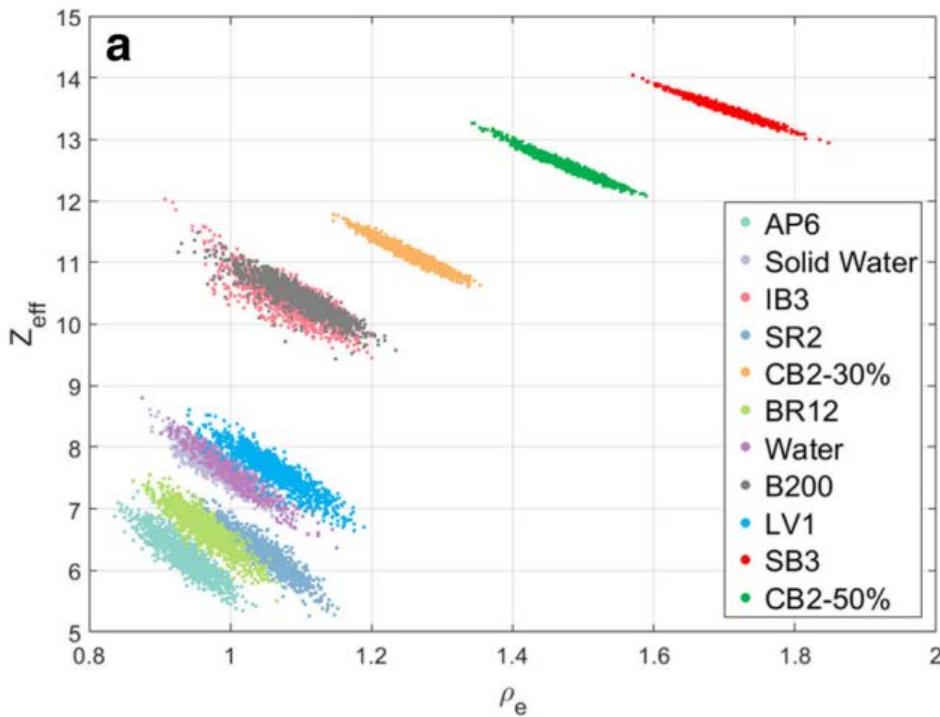
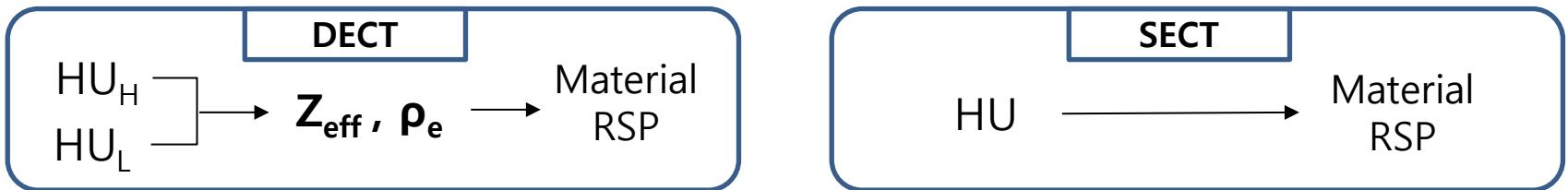
# I. Introduction

## Research purpose

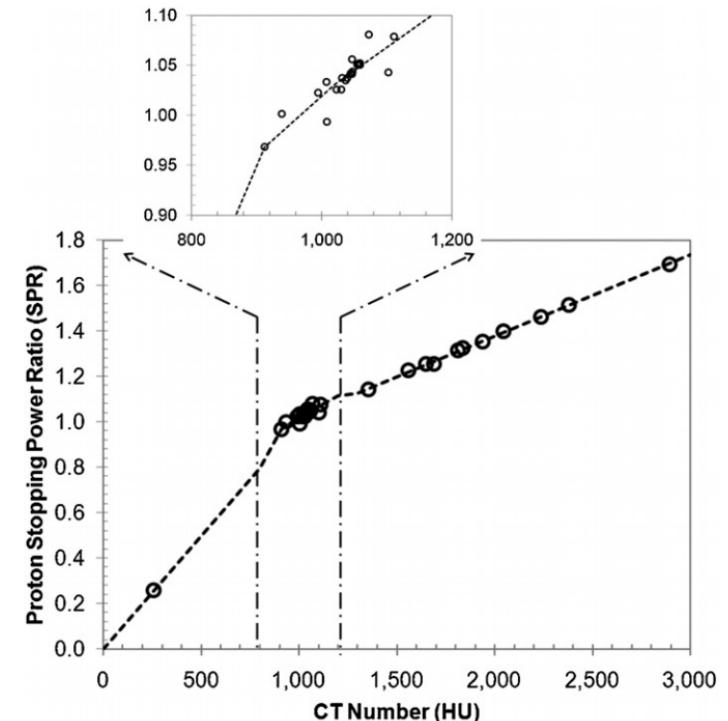
- Calculation of biological dose in CIRT using DECT image using Monte Carlo simulation
- Comparison of DECT-based and SECT-based dose calculation results

## II. Method

### DECT vs SECT



Vaniqui, et al. 2017



## II. Method

### Dual-layered DECT

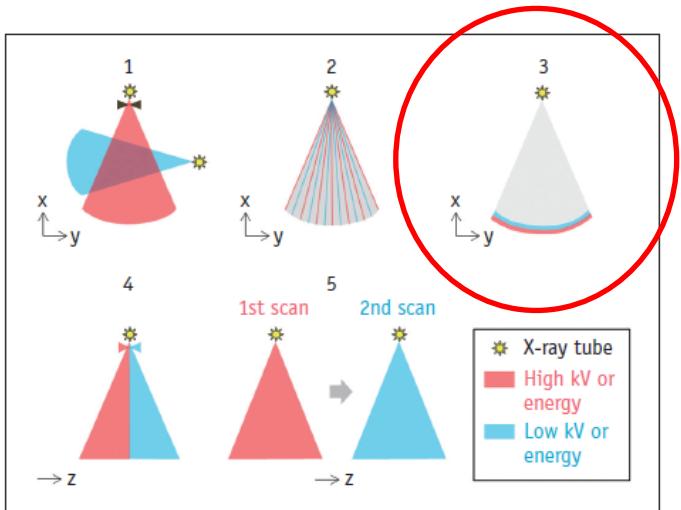


Fig. 1. Illustration of five different methods of dual-energy CT data acquisition. 1 = dual tubes with or without beam filtration, 2 = rapid voltage switching with single tube, 3 = dual-layer detector with single tube, 4 = single tube with split filter, 5 = single tube with sequential dual scans

Goo et al, *Korean J Radiol*, 2017

Philips, IQon spectral CT



### Advantages

- Real time
- offers Virtual Monochromatic Image
- Retrospective DECT analysis (offers  $Z_{eff}$ ,  $\rho_e$ )

### Disadvantages

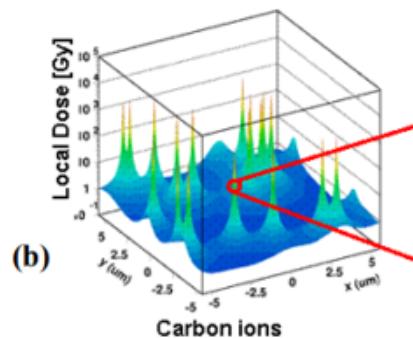
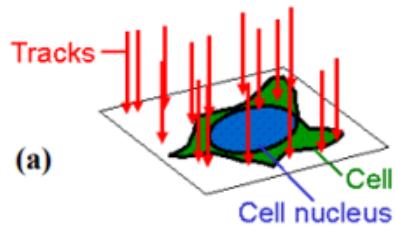
- lower sensitivity to optical photon
- cross-talk between the two detector layers

## II. Method

### RBE Calculation model

#### Local Effect Model (LEM)

$$\overline{N(D)} = \int \frac{-\ln S(d(x,y,z))}{V} dV.$$



#### Microdosimetric Kinetic model (MKM)

$$\overline{N(D)} = -\ln S = (\alpha_0 + \beta z_{1D}^*)D + \beta D^2$$

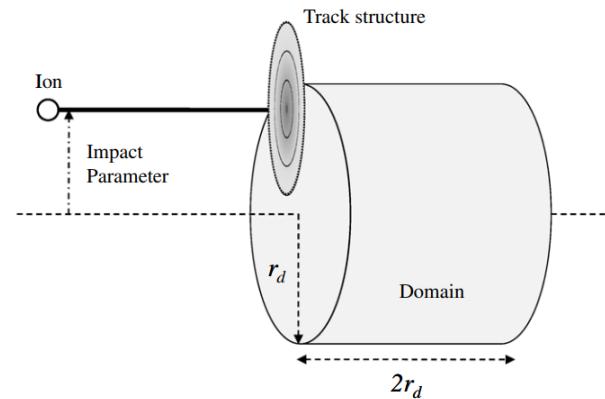


Figure 1. Schematic of an incident ion with respect to a cylindrical sensitive volume.

## II. Method

### MKM model

**Modified MKM(mMKM)**<sub>(2010, inaniwa)</sub>

$$\overline{N(D)} = -\ln S = (\alpha_0 + \beta z_{1D}^*)D + \beta D^2$$

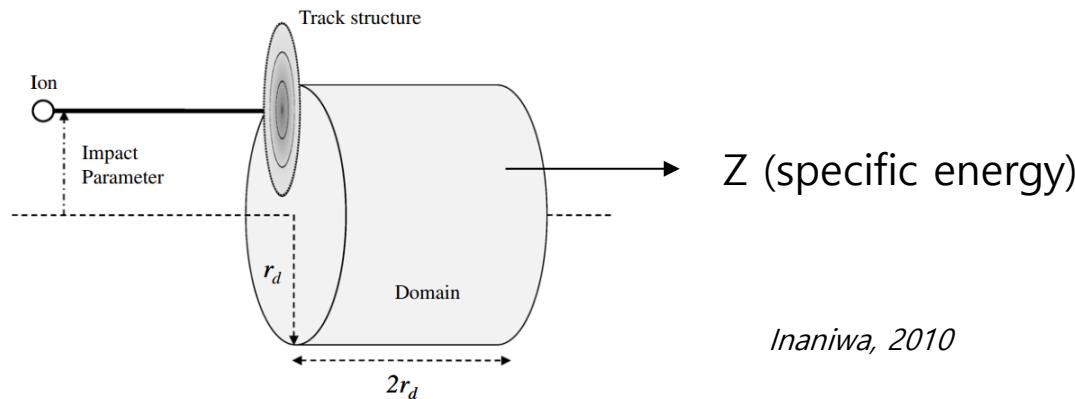
$$\alpha_0 = 0.172, \beta = 0.0615 \text{ (constant)}$$

$z_{1D}^*$  : Dose mean saturation corrected specific energy in domain

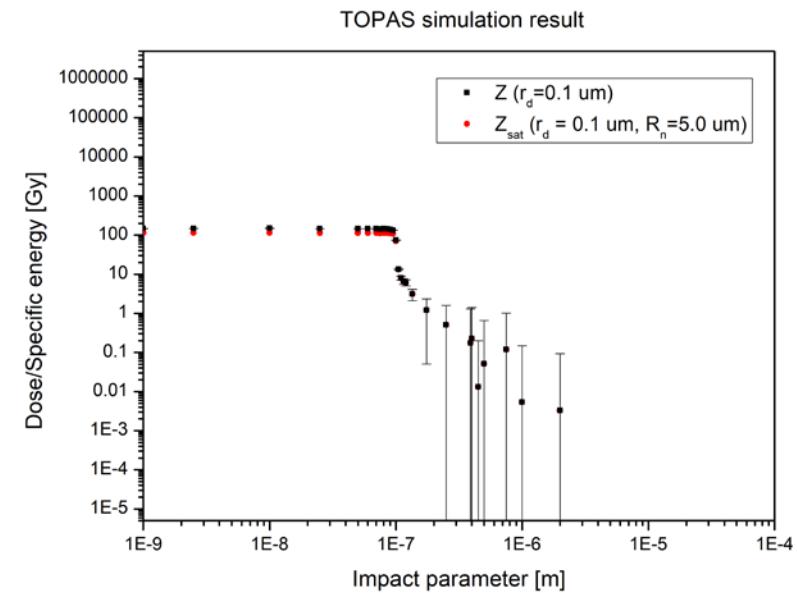
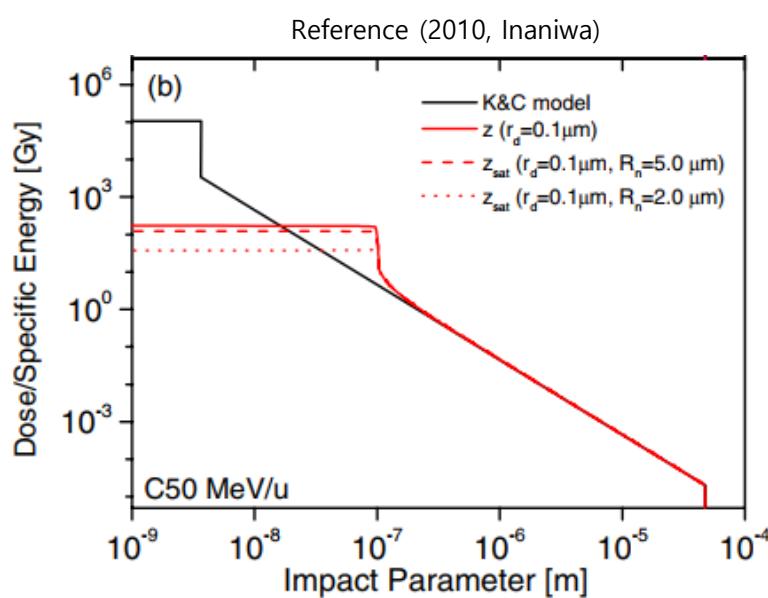
|              | Reference                          | This study  |
|--------------|------------------------------------|---|
| <b>Micro</b> | Track structure model              | Monte Carlo simulation<br><b>(TOPAS, geant4-dna)</b>  |
| <b>Macro</b> | Monte Carlo simulation<br>(Geant4) | Monte Carlo simulation<br><b>(TOPAS, QGSP_BIC_HP)</b> |

## II. Method

### Microdosimetric simulation

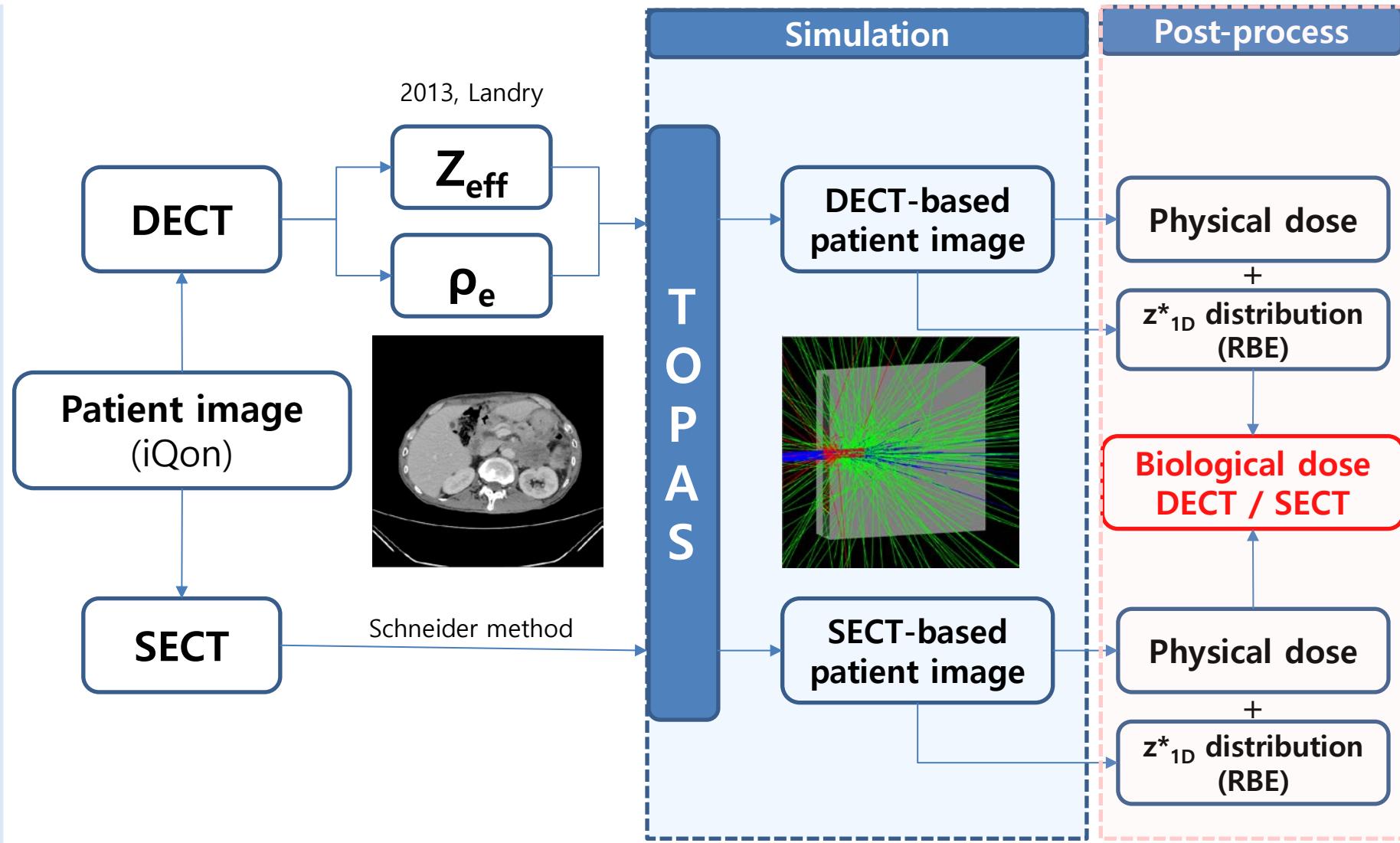


**Figure 1.** Schematic of an incident ion with respect to a cylindrical sensitive volume.



## II. Method

### Schematic diagram of B.D calculation process

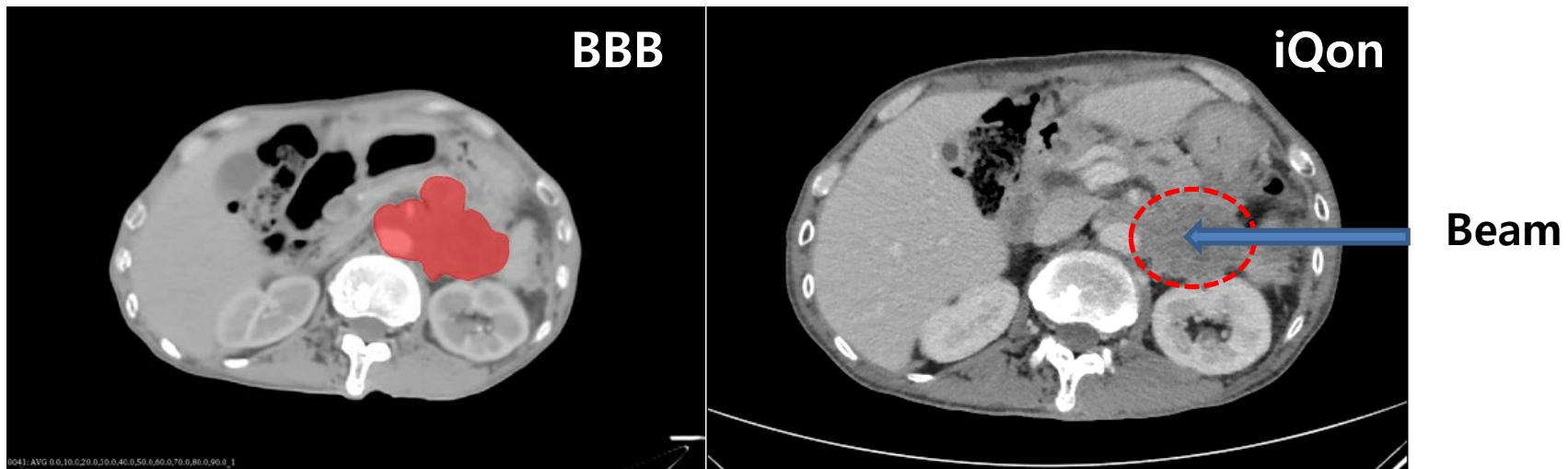




## II. Method

### TOPAS simulation

- ① Abdomen (soft tissue) : pancreas + partial abdomen

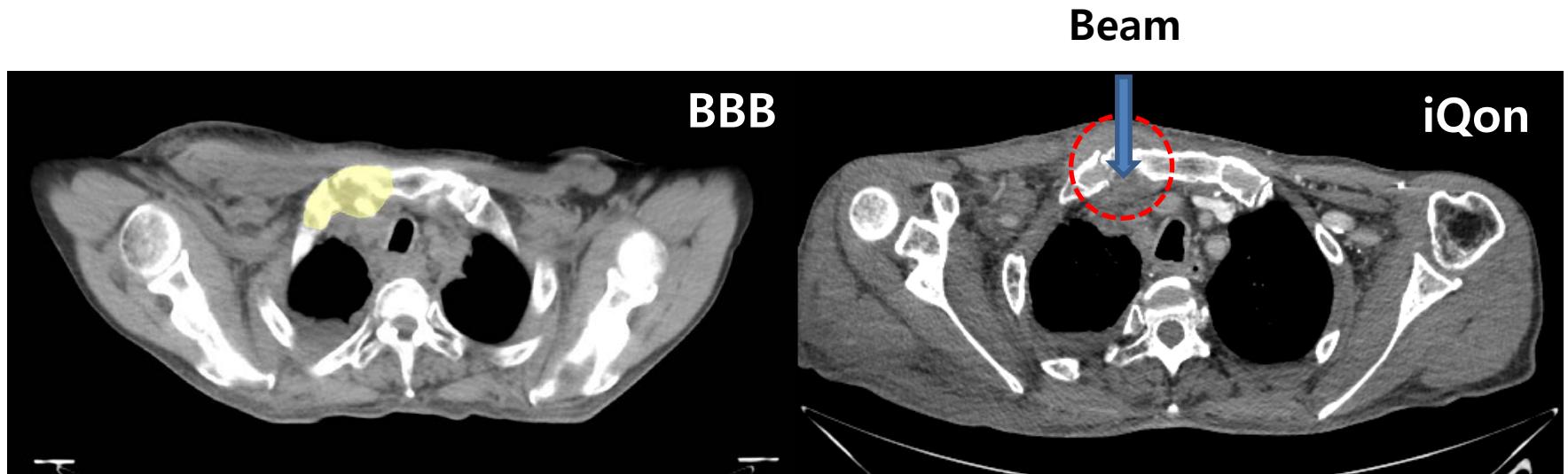


Beam specification : monoenergetic , 180 MeV/u ,  $\sigma = 3$  mm  
History :  $10^5$  / Scoring z\*1D simultaneously

## II. Method

### TOPAS simulation

- ② Head and Neck (bone) : sternum

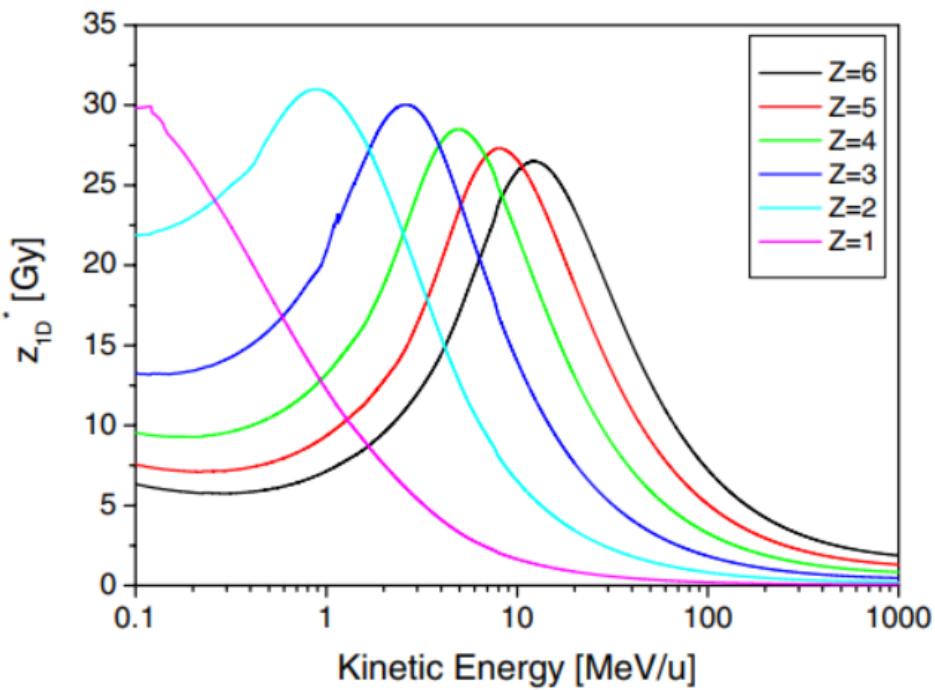


Beam specification : monoenergetic , 130 MeV/u ,  $\sigma = 3$  mm  
History :  $10^5$  / Scoring z\*1D simultaneously

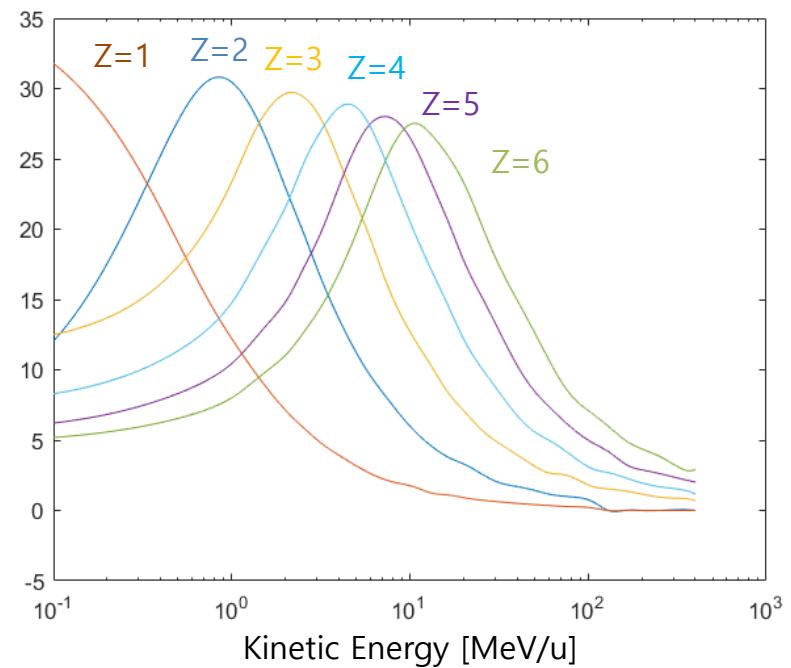
### III. Results

#### Specific energy ( $z^*_{1D}$ ) table

Inaniwa (2010)



TOPAS microdosimetric simulation

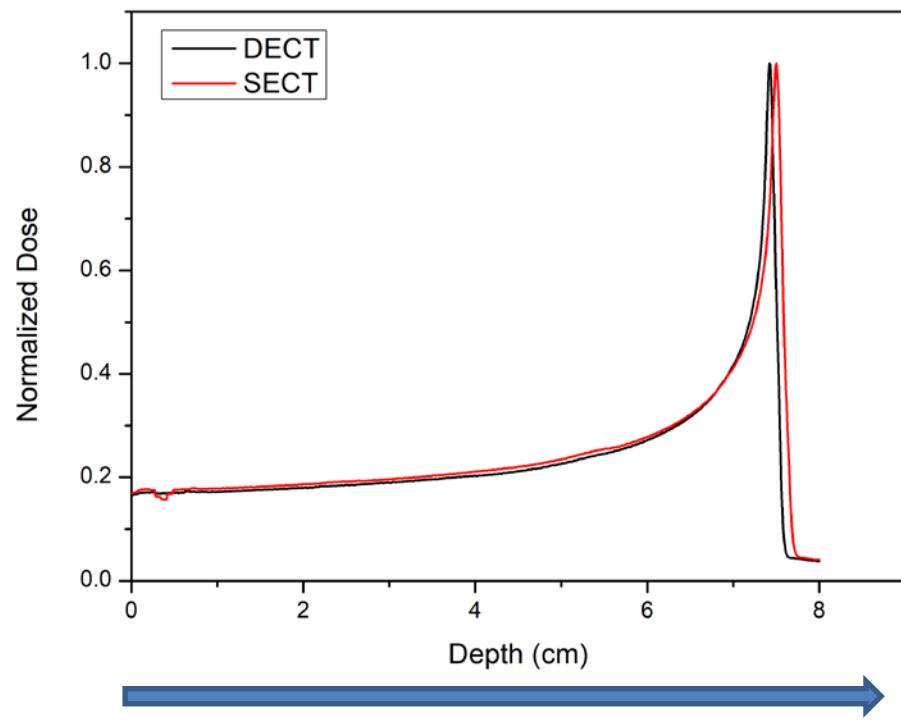
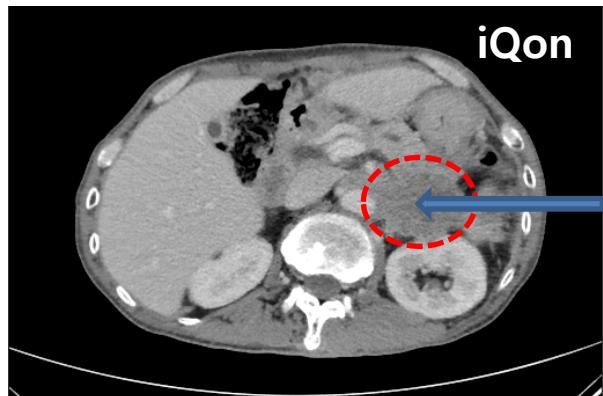


Analytical model based calculation

Monte carlo calculation

### III. Results

#### Physical dose distribution

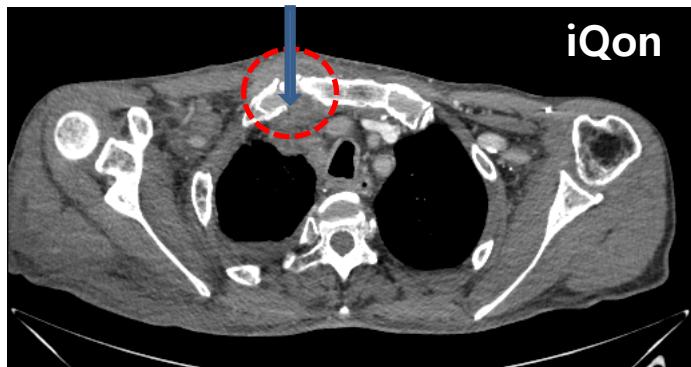


Range( $R_{80}$ ) difference : 0.8 mm, 1.1 %

※  $R_{80}$  : depth of the 80% distal end of the Bragg peak maximum

### III. Results

#### Physical dose distribution

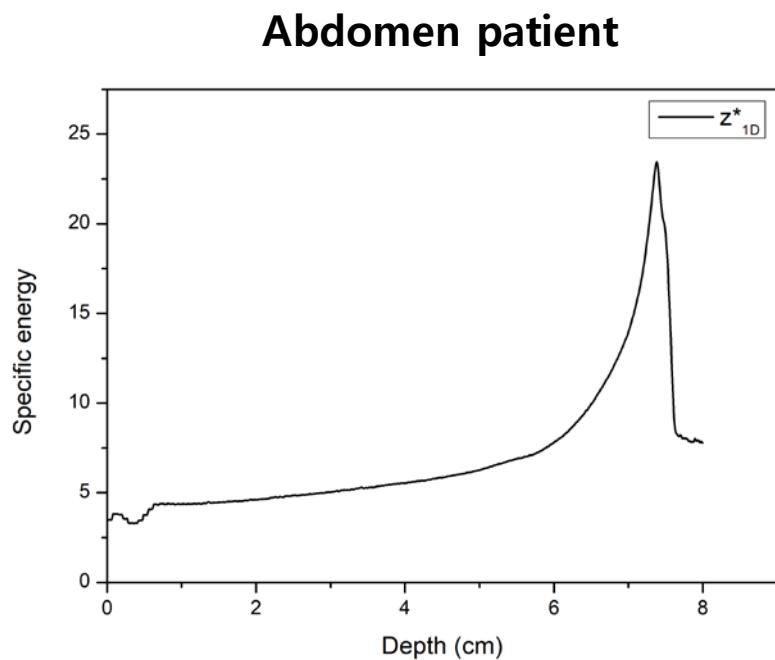


For head and neck patient

Range( $R_{80}$ ) difference : mm

### III. Results

#### $z^*_{1D}$ distribution



**H&N patient**

For head and neck patient

Common to DECT / SECT

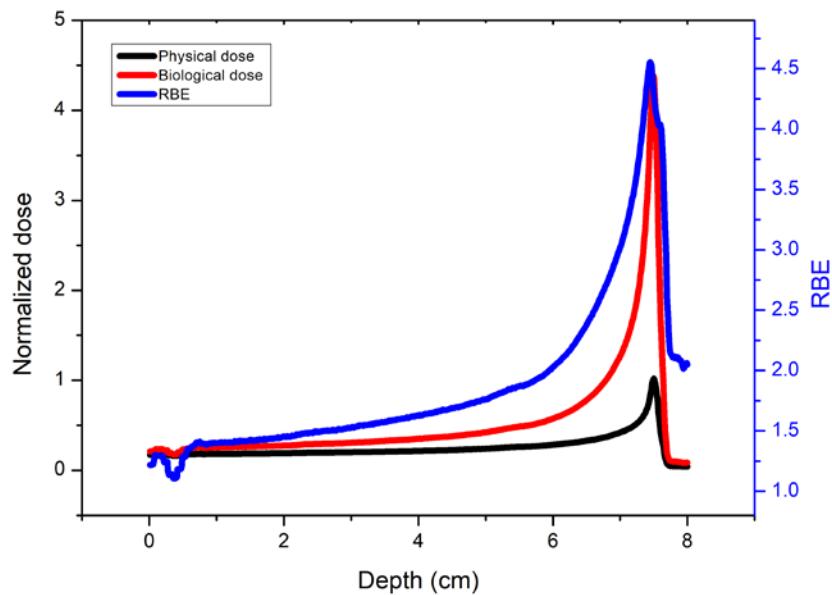
# III. Results

## Biological dose distribution

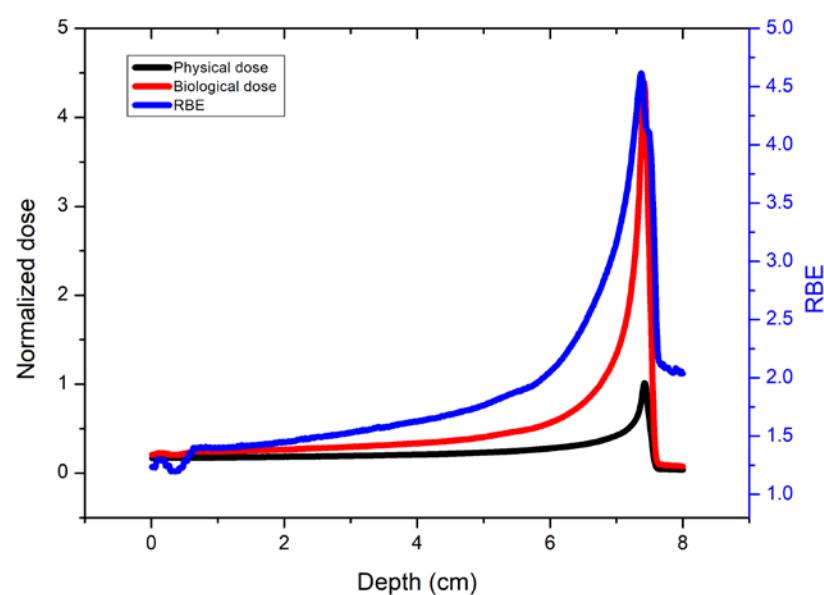
**Normalized by biological dose : 4.4 Gy (RBE)**

For pancreatic cancer patient, Carbon-ion Radiotherapy, 2014

SECT



DECT



**Normalized physical dose difference : 0.5 %**

# III. Results

## Biological dose distribution

### Normalized by biological dose : Gy (RBE)

For sternum cancer patient, Carbon-ion Radiotherapy, 2014

For head and neck patient

For head and neck patient

Normalized physical dose difference : %

## 4. Discussion & Conclusion

- By applying DECT, which can reduce the intrinsic uncertainty of SECT-based calculation, DECT-based biological dose calculations for patients were performed through Monte Carlo simulations.
- In addition, SECT-based dose calculation was performed and the results were compared in two patient case.
- In the soft tissue of the abdomen, the range difference was 0.8 mm (1.1 %) and in the H&N patients containing bone, the range was xx mm (yy %).
- The difference between the physical dose at the target internal point was calculated based on the DECT and SECT-based biological dose calculation results, and it was found to be 0.5% in soft tissue and zz% in tissue containing bone.
- If DECT is used in place of SECT through further verification, it is expected that correction of the range uncertainty will be greater in the area containing bone than in the area containing a lot of soft tissue.

Thank you for attention