

Deep learning-based 3D in vivo dose reconstruction with EPID for magnetic resonance-linear accelerators

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Background

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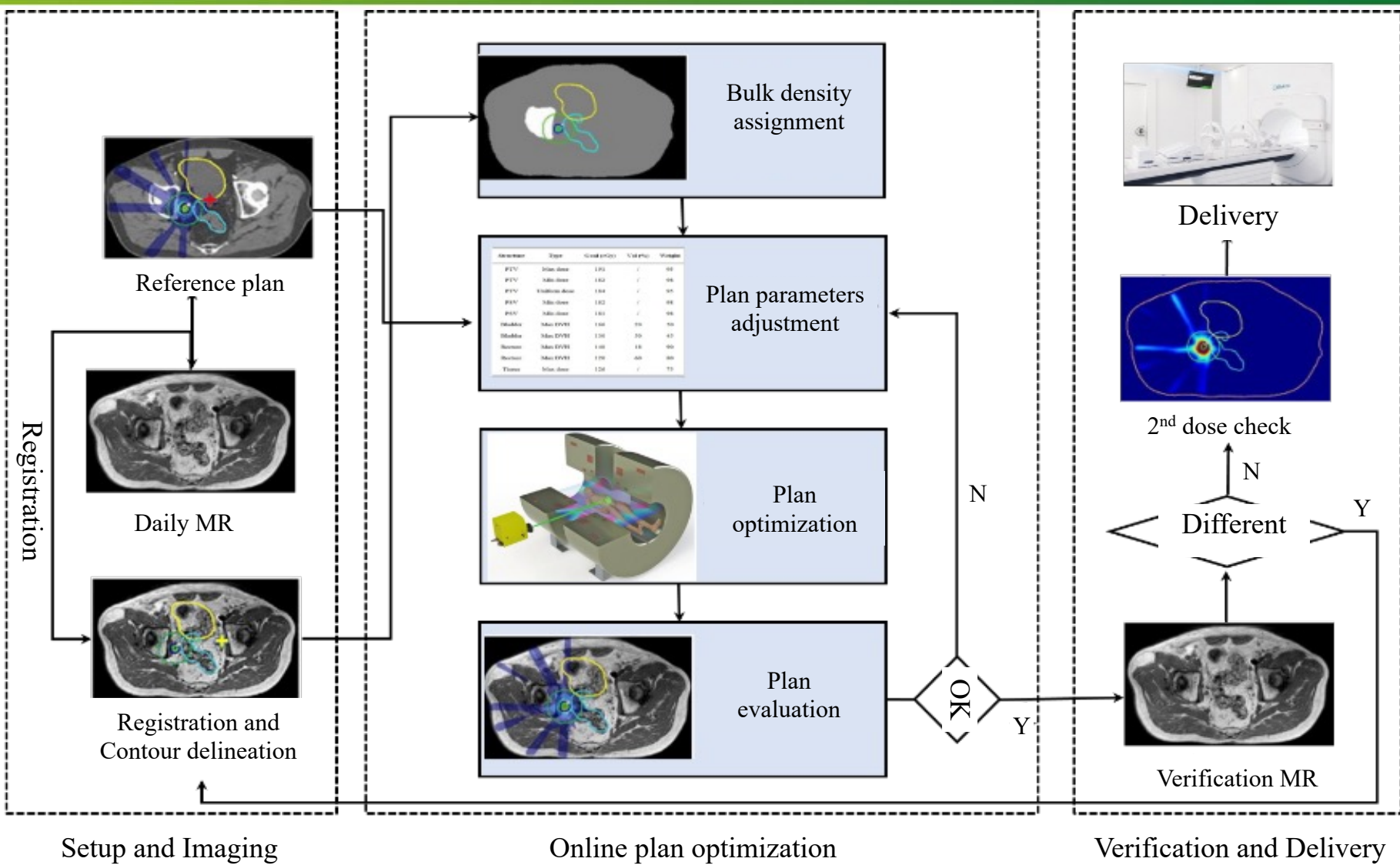


- MR-Linac
 - High soft-tissue contrast
 - No imaging dose
 - Imaging at different temporal scales
 - Functional imaging
- Adaptive radiotherapy
 - Online ART
 - Real-time ART
 - Biology guided ART



Online ART workflow

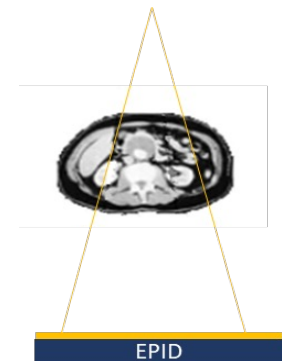
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Patient specific quality assurance (QA) for MR-Linac



- Measurement based QA for reference plan (Used in clinic)
 - Reference plan is not delivered
 - Patient modeling and dose calculation is not included
- Independent dose check for adaptive plan (Used in clinic)
 - No delivery information is included
- EPID-based in vivo dosimetry (Desired in clinic)
 - End-to-end dose verification, including verification of patient geometry and setup, synthetic CT generation or density assignment, dose calculation from TPS, plan transfer and delivery
 - Real-time dose monitoring



Unity EPID

SAD = 143.5 cm

SDD = 263.5 cm

Scale factor = 1.84

EPID size: 41 cm × 41 cm

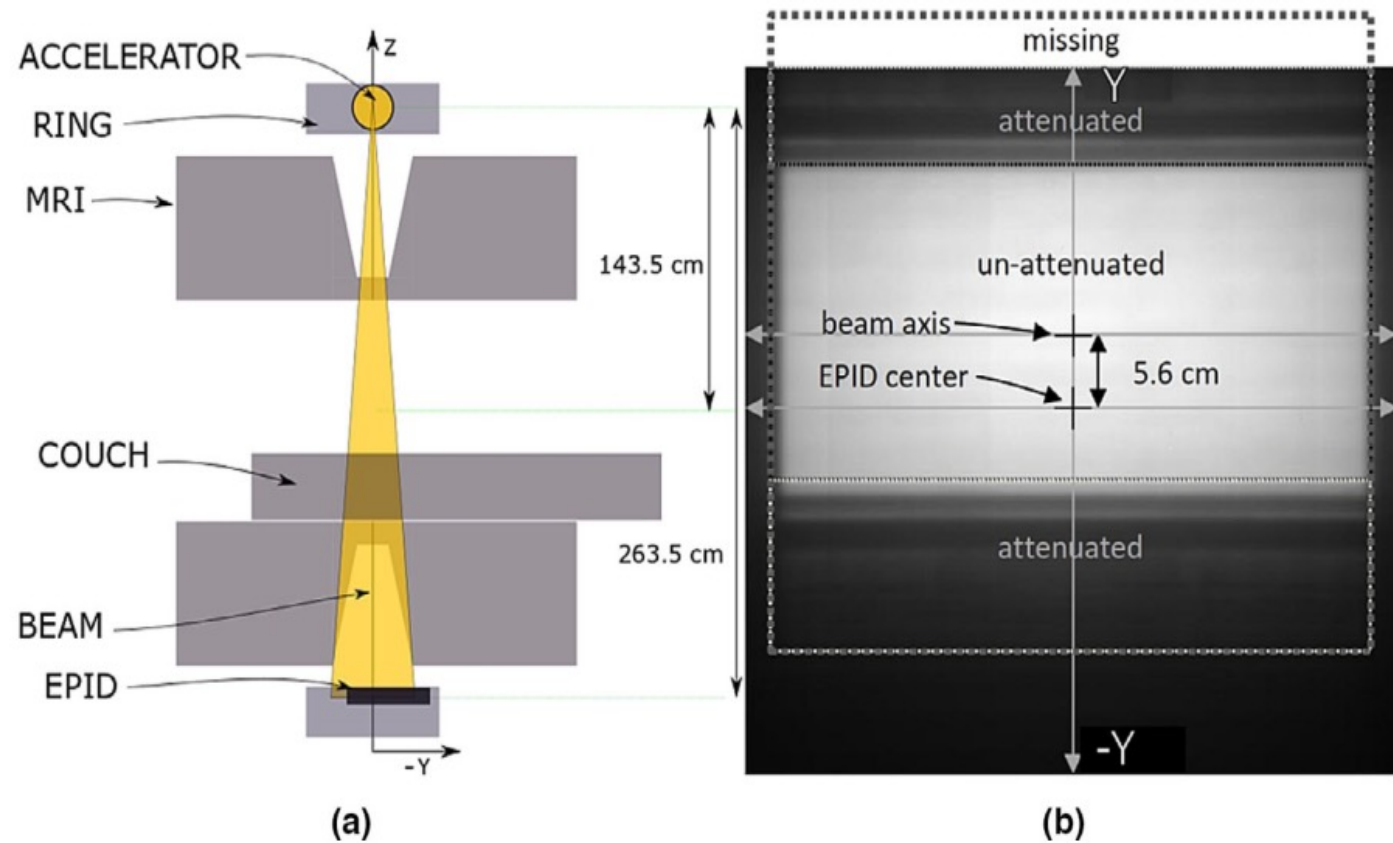
Dimension: 1024 × 1024

Resolution: 0.4 mm × 0.4 mm

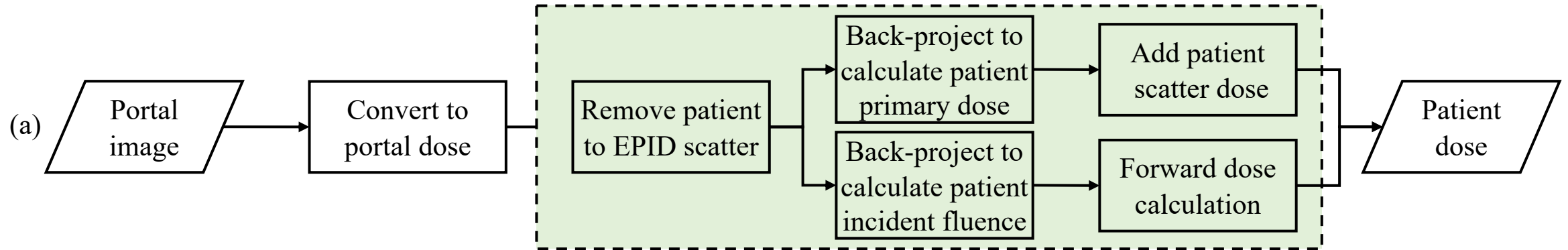
Unattenuated region:

X: [-11 cm, 11 cm]

Y: [-4.8 cm, 4.8 cm]



Conventional in vivo dose reconstruction workflow



- Back projection algorithm is not accurate for inhomogeneous region, magnetic field effect can not be considered
- Forward calculation with Monte Carlo has low efficiency, is challenged for real-time dose monitoring

Accuracy of BP algorithm in low density region

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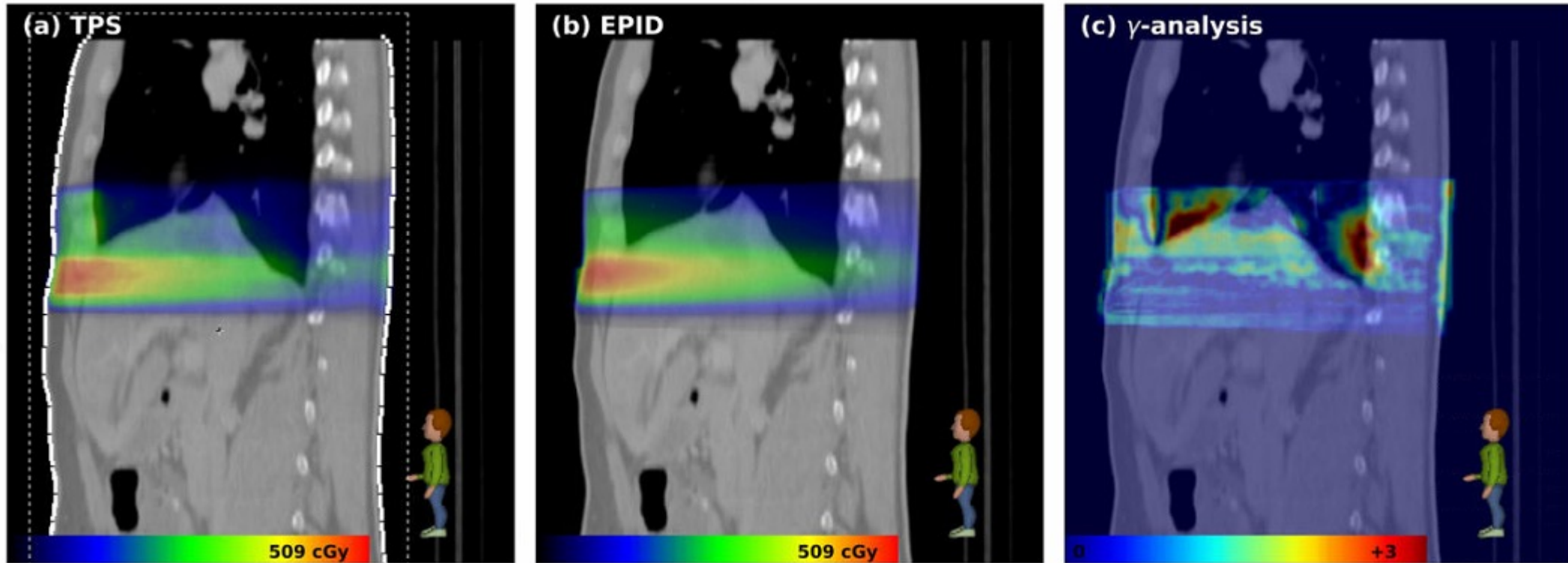
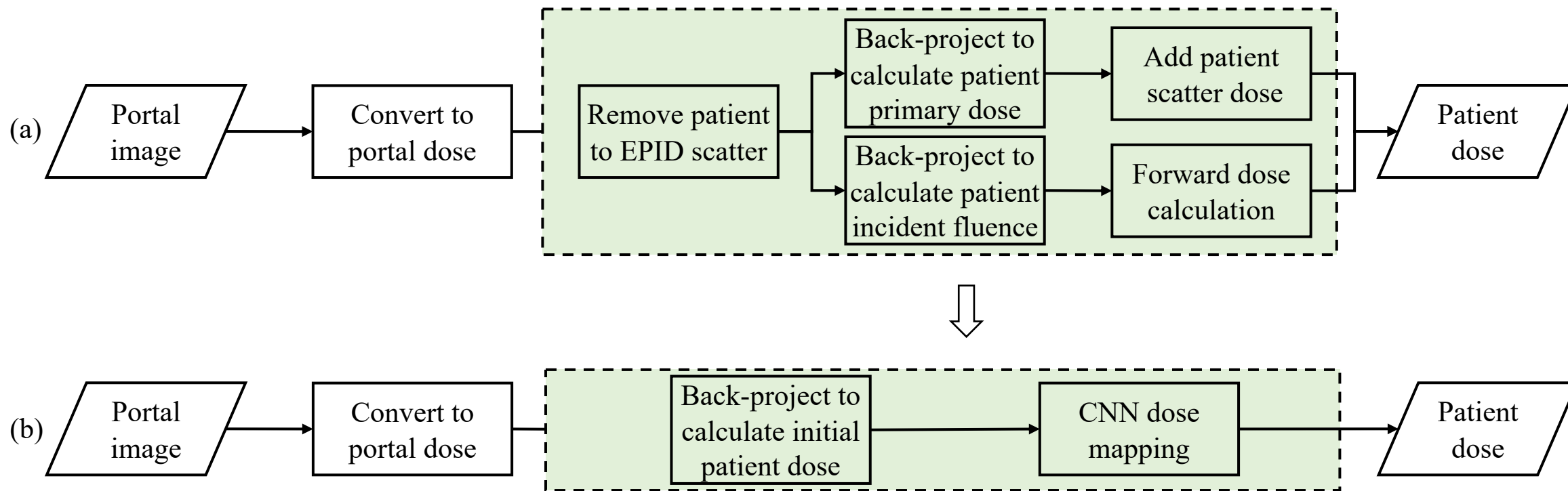


Fig. 4. (a) TPS dose, (b) *in vivo* EPID dose and (c) γ distributions corresponding to the field delivered at gantry angle 0° of the liver plan with the worst agreement. The agreement worsens for the parts of the beam traversing low density lung tissue.

Deep learning-based 3D in vivo dose reconstruction

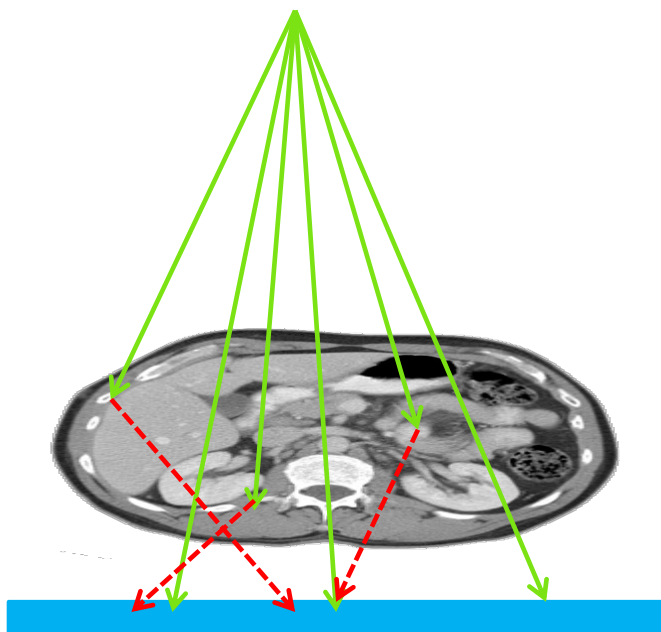


- No complex patient scatter modeling and commissioning step is needed
- All scatter effects, beam hardening effects, heterogeneity effects, and magnetic field-induced EREs were assumed to be captured by the CNN model

Deep learning-based 3D in vivo dose reconstruction

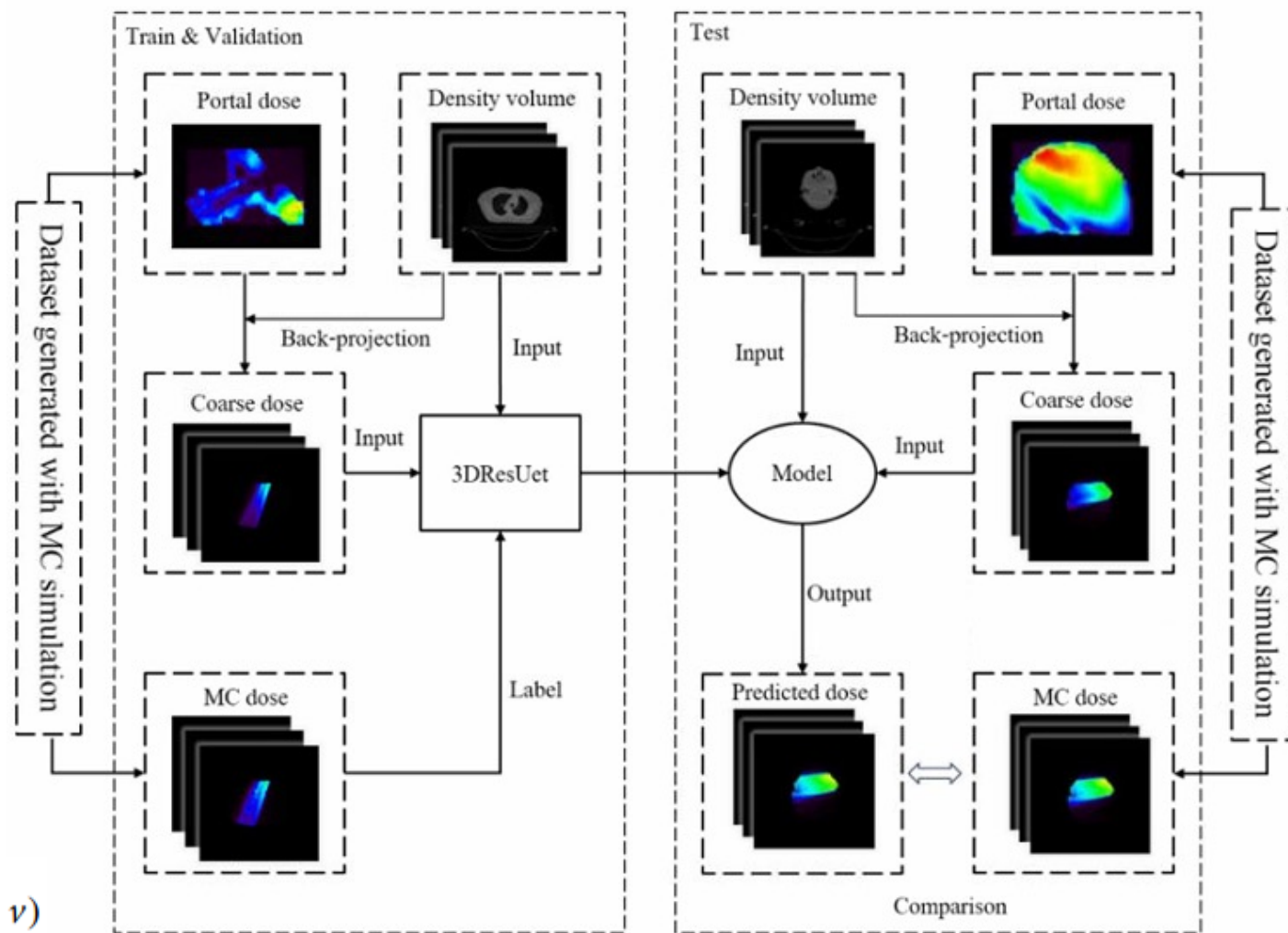


- Use Monte Carlo simulation to acquire 3D patient dose and 2D portal dose simultaneously



- Back projection to get coarse dose

$$d^{ini}(u, v, r) = (e^{-\mu\hat{r}} - e^{-\beta\hat{r}}) / (e^{-\mu\hat{r}_{EPID}} - e^{-\beta\hat{r}_{EPID}}) \frac{r_{EPID}^2}{r^2} \varphi(u, v)$$



Monte Carlo simulation with magnetic field

- gDPM ——— developed by Xun Jia (JHU)
 - Based on fast MC code-DPM
 - GPU acceleration
 - Separate photon and electron transportation
 - ~60-80 times efficiency improvement with CPU version
- Extend charged particle transportation in magnetic field
 - First order approximation



Xun Jia, et al. *Phys. Med. Biol.* 2011.

$$\Delta \mathbf{u} = \frac{q \cdot s}{m_0 \gamma v_0^2} [\mathbf{v}_0 \times \mathbf{B}_0]$$

$$\mathbf{u}(s) = \mathbf{u}_0 + \Delta \mathbf{u}$$

$$R = \frac{m_0 \gamma v_0^2}{q |\mathbf{v}_0 \times \mathbf{B}_0|}$$

$$\Delta \mathbf{u} = \frac{s}{R} \frac{\mathbf{u}_0 \times \hat{\mathbf{B}}_0}{|\mathbf{u}_0 \times \hat{\mathbf{B}}_0|}$$

$$\delta = \frac{s}{R} \ll 1$$

$$s = \min(R \cdot \delta, s_{vox}, s_{hard}, s_{ele})$$

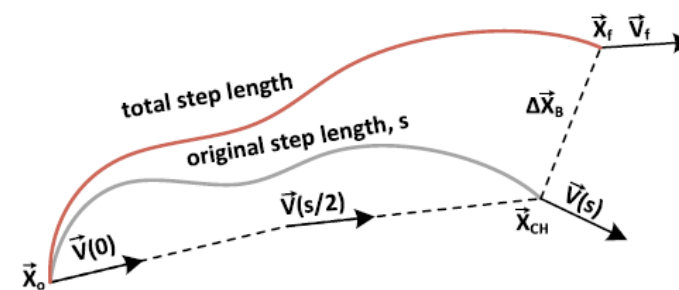
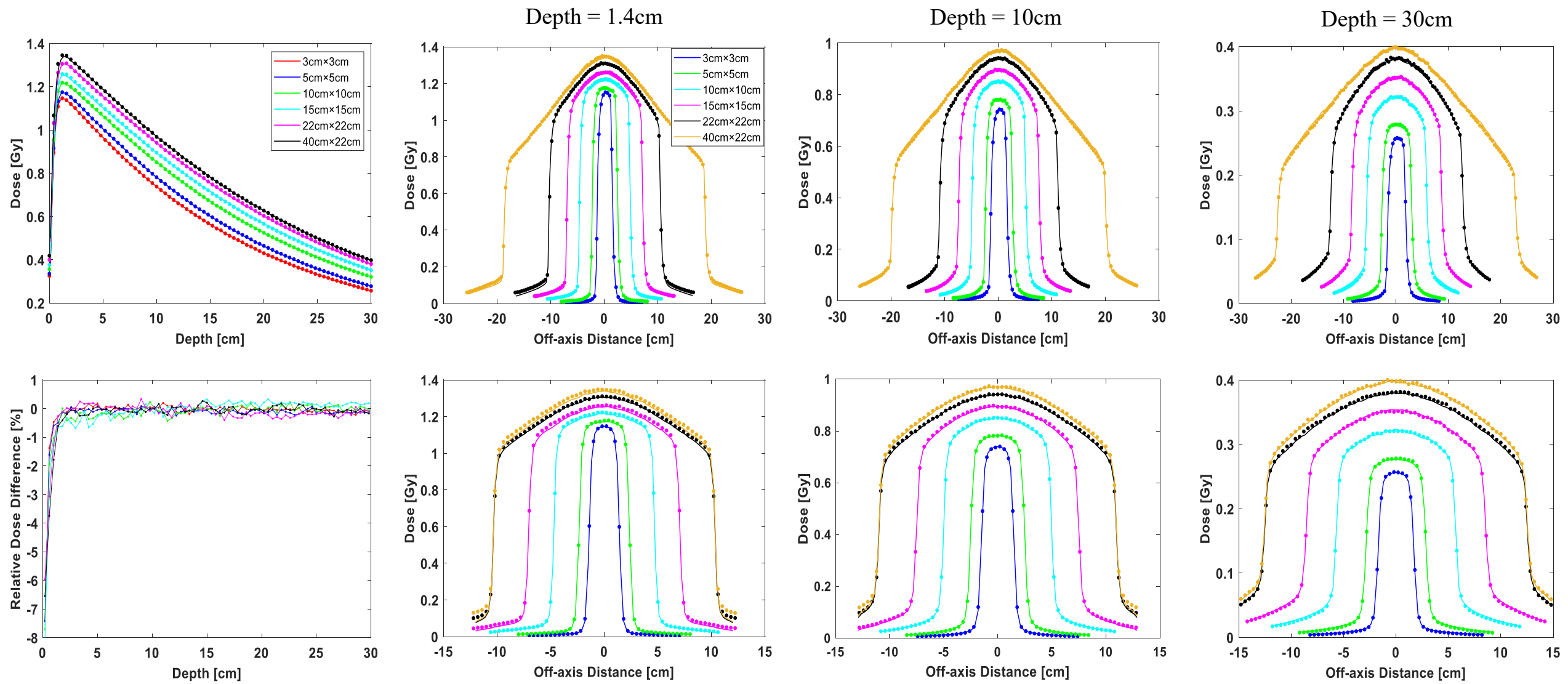


FIG. 1. Simplified EGSnrc PRESTA-II step in the presence of a magnetic field. The particle is initially at \vec{x}_0 with velocity $\vec{v}(0)$, it is then transported a step length, s , to \vec{x}_{CH} by the CH algorithm which samples direction of motion at an intermediate, $\vec{v}(s/2)$, and final, $\vec{v}(s)$. $\Delta \vec{x}_B$ and $\Delta \vec{u}_B$ are calculated using Eqs. (2) and (6) to obtain the final position, \vec{x}_f , and velocity, \vec{v}_f .

EGSnrc Manual.

Monte Carlo beam modeling for Unity with 1.5T

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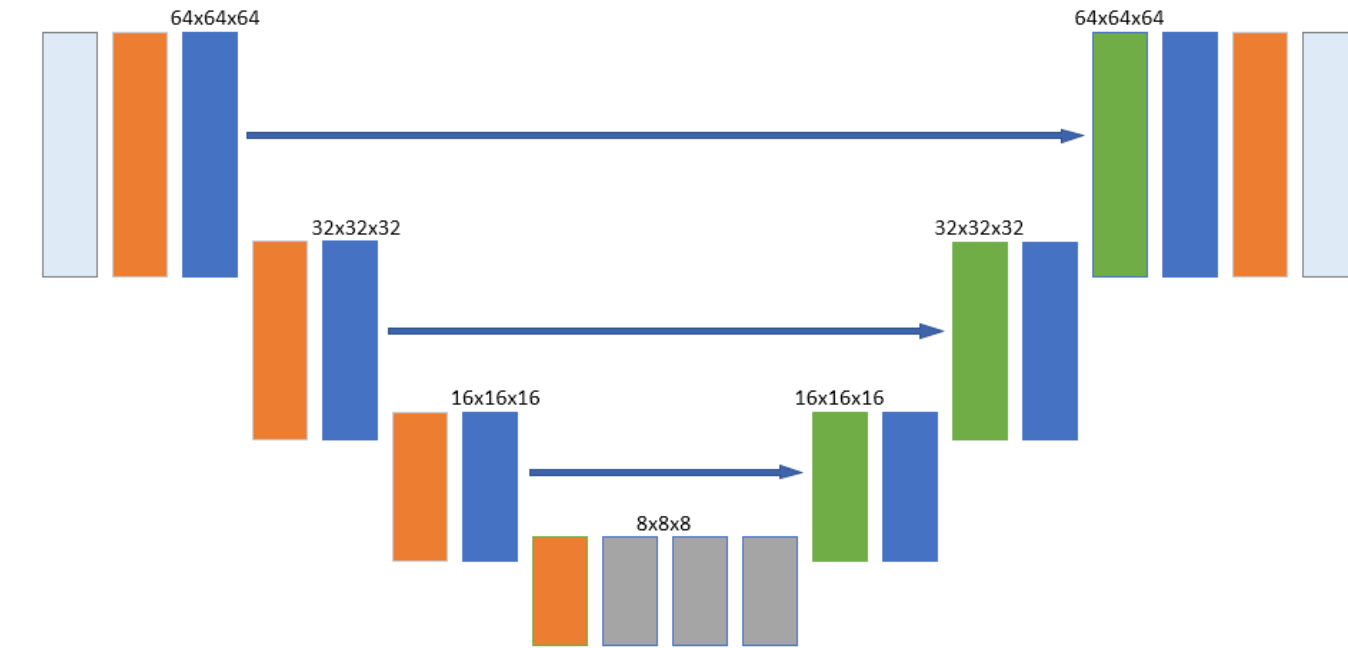


Dataset and augmentation

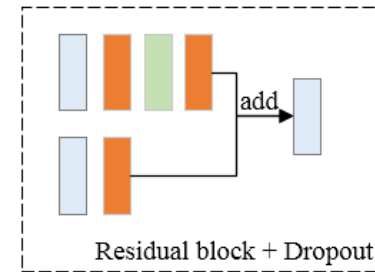
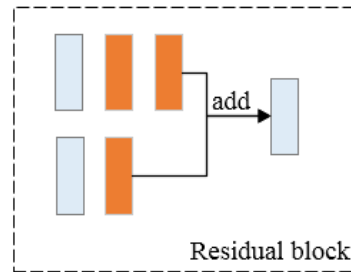


- Dataset
 - 21 brain cases, 46 NPC cases, 15 lung cases, 14 rectum cases
 - training and validation set (78 cases), test set (18 cases)
 - 576 original treatment beams
 - Augmentation
 - rotate the original beam angles by 10° – 15° for 2–3 times
 - 1841 and 121 beams for training and validation
 - all volume dose and portal dose were recalculated for each beam
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Network structure



- Input/Output
- Conv3D
- Residual block
- Conv3DTranspose
- Residual block + Dropout
- Dropout
- Concatenate



Dose reconstruction result

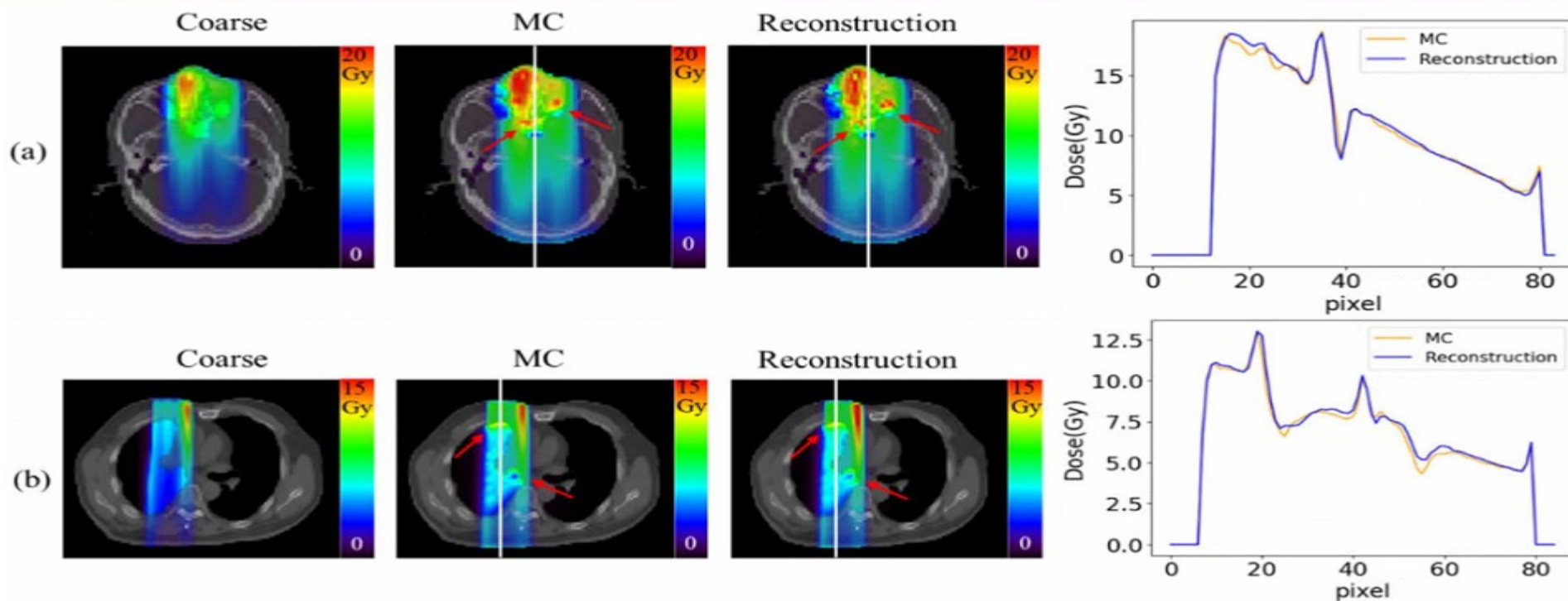
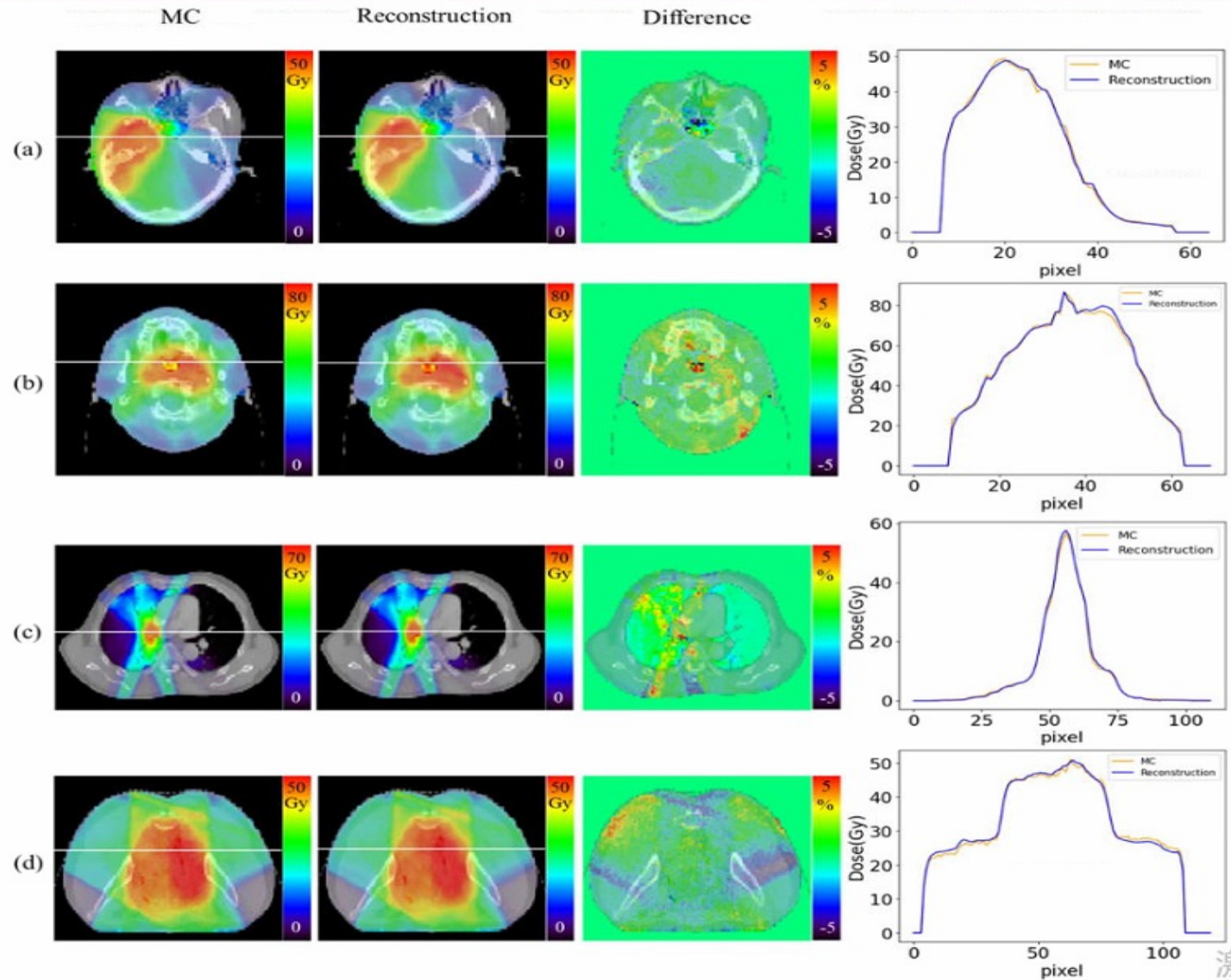


Table 1. Averaged γ -pass rates and MAE parameters for 18 tested patients (mean \pm SD).

Site		Brian (4)	nasopharynx (8)	Lung (3)	Rectum (3)
γ -pass (%)	Dose > 0%, 3%/2 mm	97.42 \pm 2.66	98.53 \pm 0.95	99.41 \pm 0.46	98.63 \pm 1.01
	Dose > 20%, 3%/2 mm	95.48 \pm 3.31	97.20 \pm 1.42	95.35 \pm 0.57	95.29 \pm 2.89
	Dose > 50%, 3%/2 mm	94.32 \pm 3.77	95.10 \pm 2.01	90.40 \pm 2.71	95.83 \pm 1.56
	Dose > 0%, 2%/2 mm	94.02 \pm 6.21	96.46 \pm 1.93	98.72 \pm 0.88	96.72 \pm 1.73
MAE (%)	0.82 \pm 0.36	0.88 \pm 0.21	0.41 \pm 0.19	0.67 \pm 0.09	

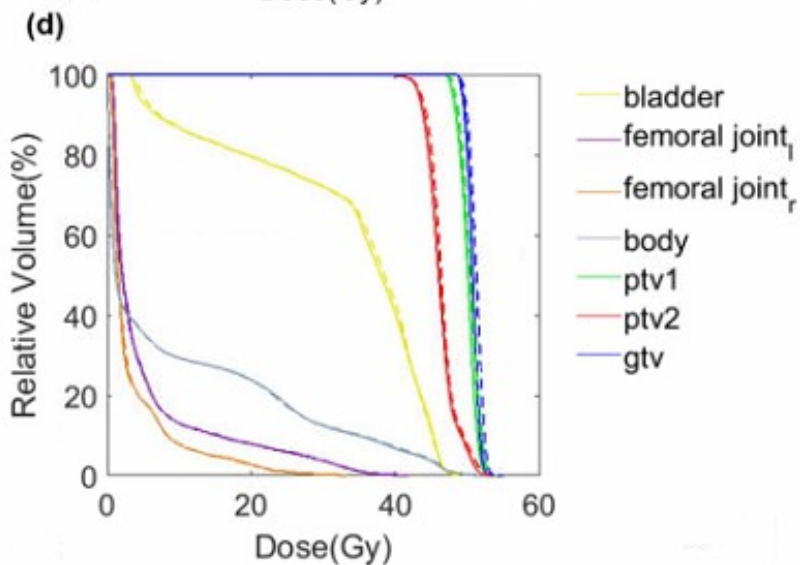
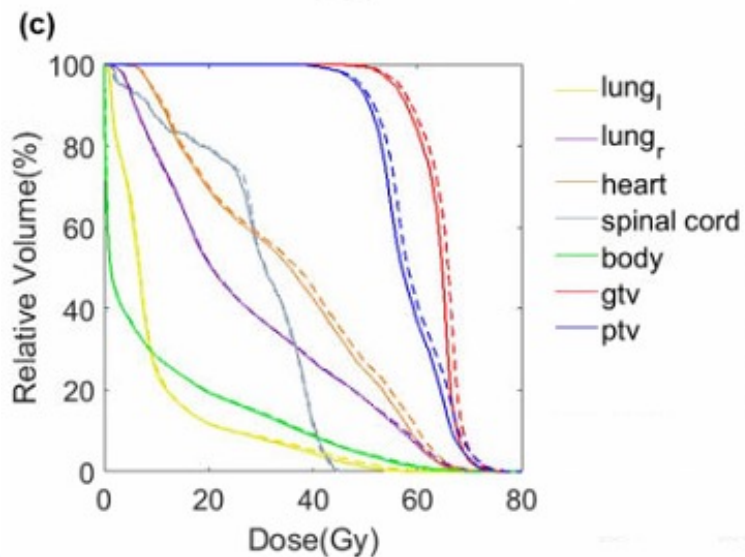
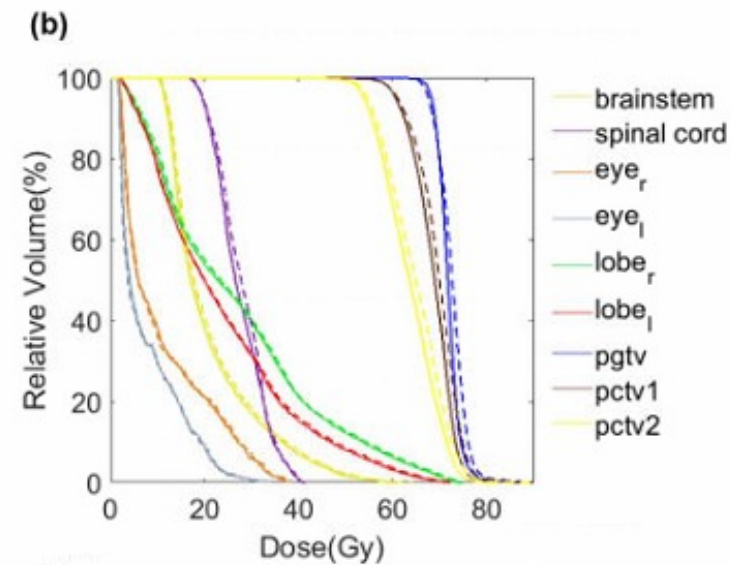
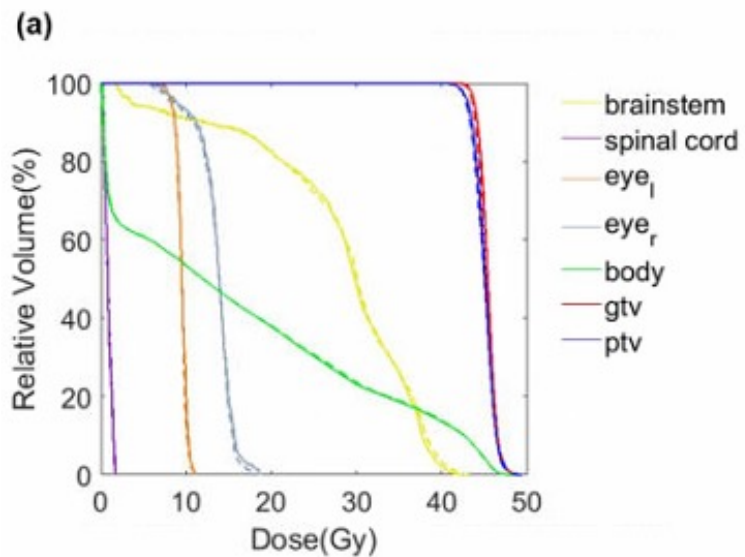
Dose reconstruction result

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Dose reconstruction result

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Conclusion

- Proposed a CNN-based 3D in vivo dose reconstruction method
 - Physical processes were learned through accurate MC data-driven model training
 - Simplicity in dose reconstruction and model commission
 - Training data can be generated with clinic TPS by adding a virtual EPID structure
 - Limitations
 - No validation was conducted for real measured EPID images
 - Field truncation by real EPID was not considered
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Thanks !
