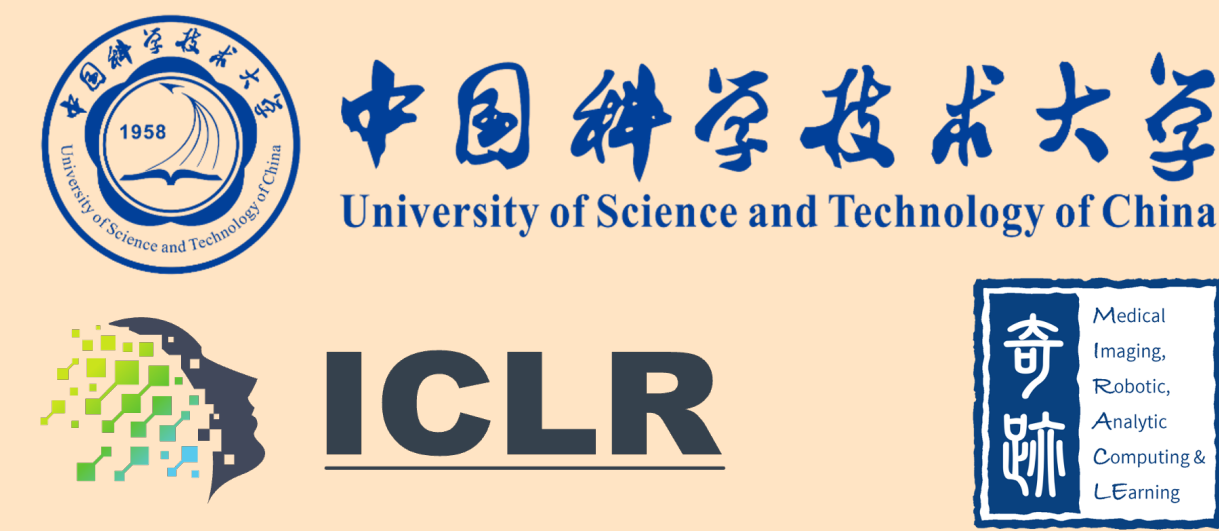


Self-Supervised Diffusion MRI Denoising via Iterative and Stable Refinement

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Background

MRI, including diffusion MRI (dMRI), serves as a “microscope” for anatomical structures.

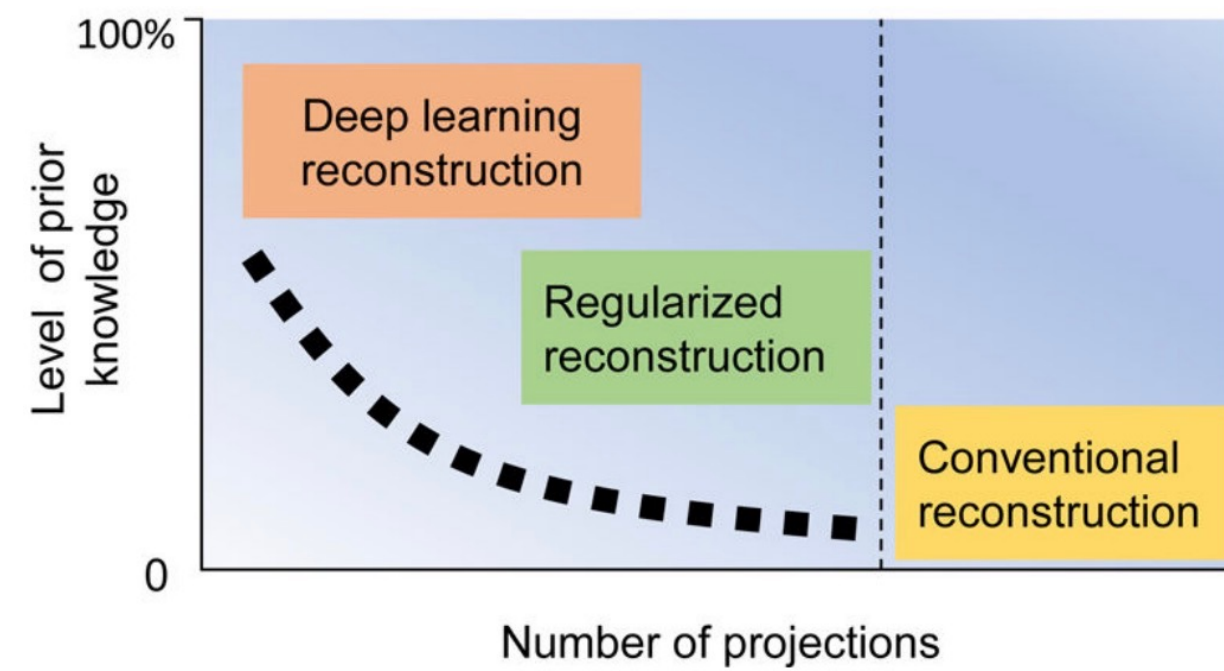


Figure 1. Different image-reconstruction schemes in the context of prior knowledge and projection sampling [1].

dMRI routinely mitigates the influence of low SNR scans by compromising temporal or spatial resolution. However, these fail to meet clinical demands for both efficiency and precision.

Noise2Noise [2] performs image restoration solely by observing corrupted measurements:

$$\arg \min_{\theta} \left\{ \mathbb{E} \|f_{\theta}(x') - x\|^2 \right\} \approx \arg \min_{\theta} \left\{ \mathbb{E} \|f_{\theta}(x') - y\|^2 + \mathbb{E} \|x - y\|^2 \right\}, \quad (1)$$

DDPM [3, 4]. In the forward process, noise with specified levels is introduced to the data,

$$q(x_t|x_0) := \mathcal{N}(x_t; \sqrt{\alpha_t}x_0, (1 - \alpha_t)\mathbf{I}), \quad (2)$$

where $\bar{\alpha}_t := \prod_{s=1}^t \alpha_s$, $\alpha_t := 1 - \beta_t$ and β_t is a variance of noise level. For the reverse process,

$$q(x_{t-1}|x_t, x_0) := \mathcal{N}(x_{t-1}; \tilde{\mu}_t(x_t, x_0), \sigma_t^2 \mathbf{I}), \quad (3)$$

where $\tilde{\mu}_t(x_t, x_0) := \frac{\sqrt{\alpha_{t-1}}\beta_t}{1 - \bar{\alpha}_t}x_0 + \frac{\sqrt{\alpha_t}(1 - \bar{\alpha}_{t-1})}{1 - \bar{\alpha}_t}x_t$ and $\sigma_t^2 := \frac{1 - \bar{\alpha}_{t-1}}{1 - \bar{\alpha}_t}\beta_t$.

Key Features in our methods

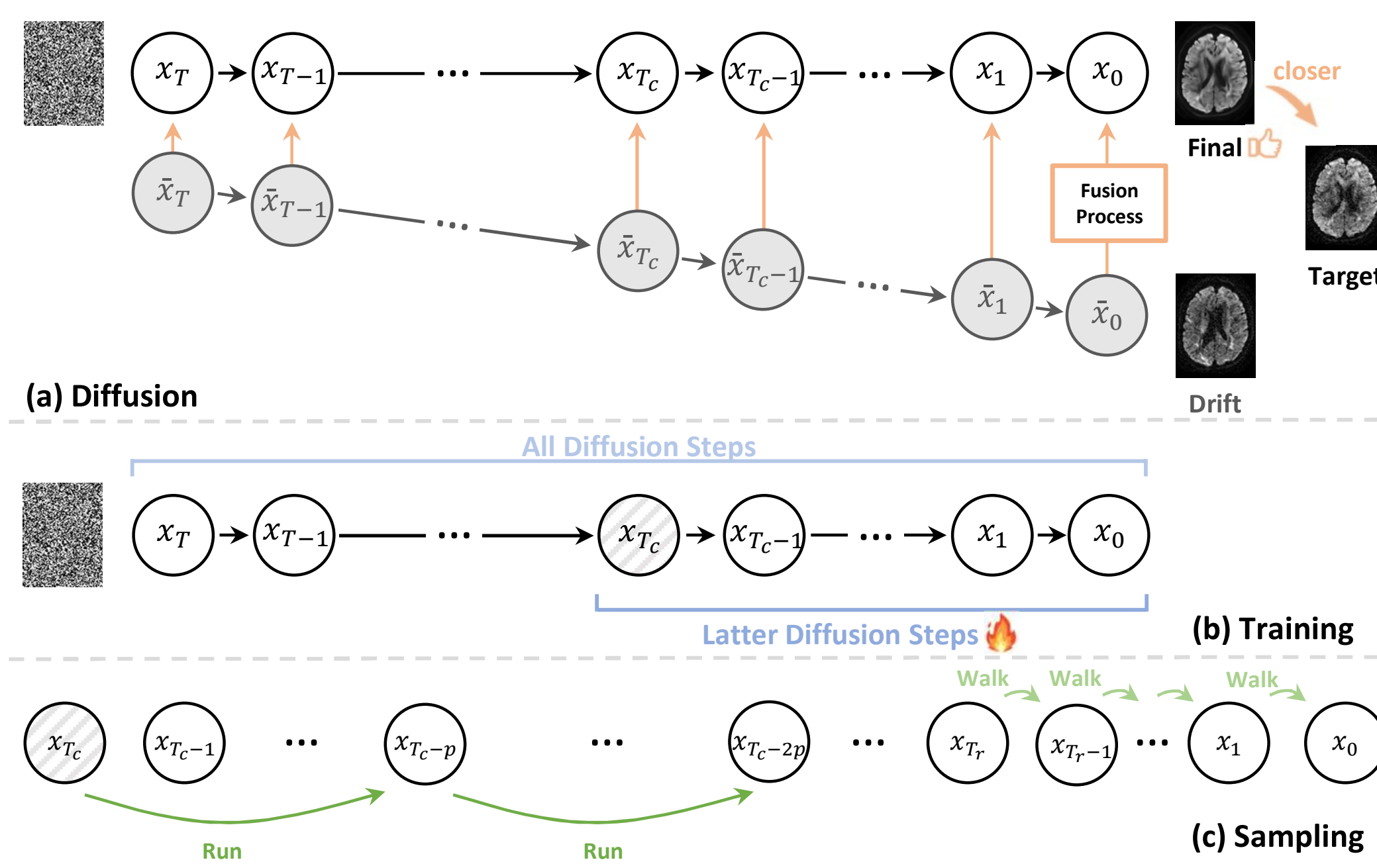


Figure 2. (a) Fusion process aligns $\{\bar{x}_t\}_1^T$ to $\{x_t\}_1^T$ and avoids drift (“Drift” means drifted results, “Final” means the denoised version of “Target”); (b) Training the latter diffusion steps imposes restrictions on the diversity of diffusion models and decreases uncertainty; (c) Run-Walk accelerated sampling accelerates the entire sampling process.

Methods

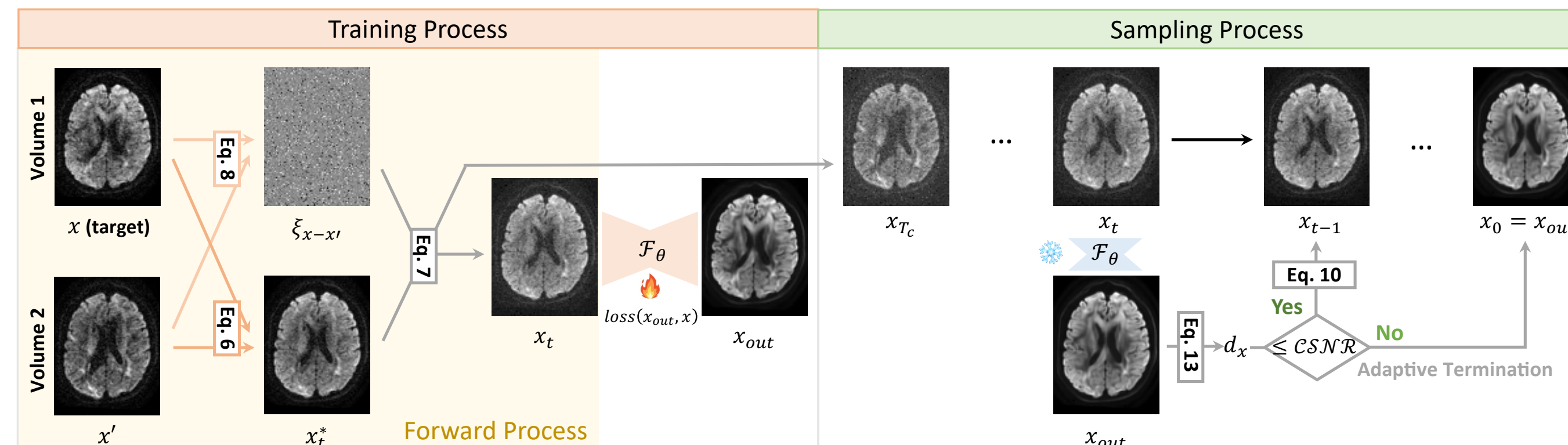


Figure 3. Overview of our single-stage Di-Fusion (See following for details). The training process does not involve any extra model training apart from $\mathcal{F}_{\theta}$, and the sampling process offers adaptive and controllable results.

Input 4D data: $X \in \mathbb{R}^{w \times h \times d \times l}$, $x = X_{*,*,i,j}$, $x' = X_{*,*,i,j-1}$

Eq.6: $x_t^* = \lambda_1^t x + \lambda_2^t x'$ (Fusion process interpolates between two endpoints)

Eq.8: $\xi_{x-x'} = \text{mess}((x - x') - \mu_{x-x'})$ (Di- process characterizes real-world noise)

Eq.7: $x_t = \sqrt{\alpha_t}x_t^* + \sqrt{1 - \bar{\alpha}_t}\xi_{x-x'}$ (Diffusion forward process)

Training target: $L_{\text{simple}}(\theta) := \mathbb{E}_{t, x_t^*, \xi_{x-x'}} [\|x - \mathcal{F}_{\theta}(\sqrt{\alpha_t}x_t^* + \sqrt{1 - \bar{\alpha}_t}\xi_{x-x'}, t)\|^2]$

Latter diffusion steps: $x_{T_c} \rightarrow x_0, T_c \leq T$.

Eq.13: $d_x = \|x - x_{out}\|^2 \times b_x$, here b_x accounts for the ratio of brain tissue to the entire image.

Eq.10: Diffusion sampling process. Our sampling schedule in Figure 2 (c).

Experiments

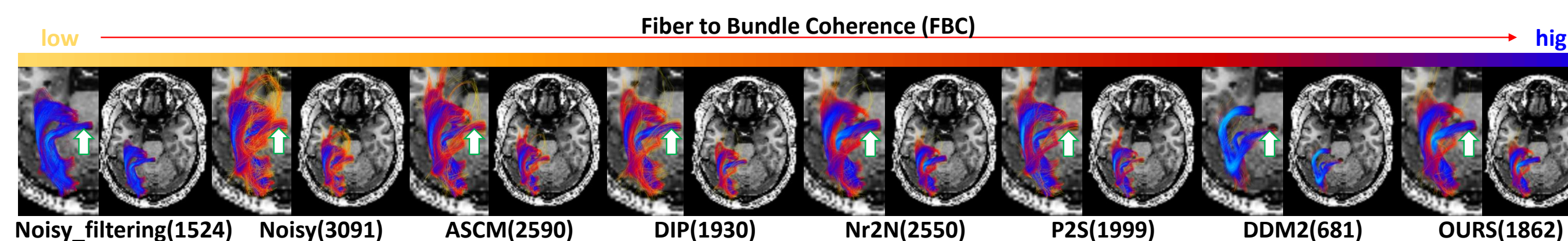


Figure 4. Density map of FBC projected on the streamlines of the OR bundles. The numbers in parentheses represent the number of streamlines. OURS generates the minimal number of streamlines while maintaining high FBCs (consider “Noisy_filtering” as references for high FBCs).

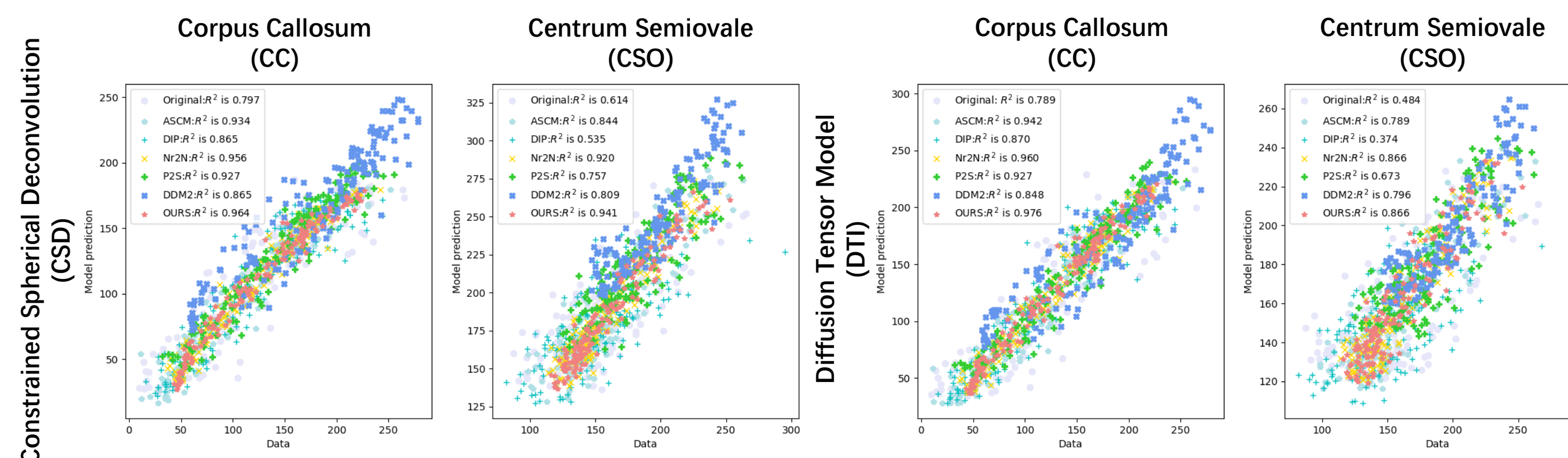


Figure 5. Scatter plots of the microstructure model predictions against input data. The top-left of each plot shows the quantitative R^2 metric computed from each model fit on the corresponding data. OURS data points are more concentrated (higher R^2), which means OURS aids in the characterization of the microstructure.

Metric	Noisy	ASCM	MPPCA	Nr2N	P2S	DDM2	OURS
PSNR (FA)	23.47	22.63	26.37	29.47	24.18	26.77	30.79
PSNR (MD)	31.79	21.81	35.28	38.42	32.16	37.53	40.26
PSNR (RD)	33.56	24.70	36.35	38.48	30.62	37.39	40.35
PSNR (AD)	26.39	20.89	29.63	34.87	33.27	33.21	35.30
SSIM (FA)	87.60	88.97	90.48	93.54	90.90	90.41	94.50
SSIM (MD)	96.35	81.88	97.51	98.51	97.08	98.72	99.31
SSIM (RD)	96.37	89.21	97.80	98.65	96.32	98.48	99.23
SSIM (AD)	92.47	81.02	95.34	97.12	96.77	96.10	97.63

Table 1. Quantitative results for DTI diffusion signal estimates. PSNR (dB) and SSIM (%) are reported. FA (fractional anisotropy), MD (mean diffusivity), RD (radial diffusivity), and AD (axial diffusivity). Best results are in bold. **OURS** provides better evidence for the diffusion signal estimates.

Method	Simulation 1		Simulation 2		Simulation 3		Simulation 4		Simulation 5	
	SSIM	PSNR	SSIM	PSNR	SSIM	PSNR	SSIM	PSNR	SSIM	PSNR
Noisy	11.38	13.72	20.65	16.09	37.41	20.38	52.02	23.76	64.62	26.63
P2S	24.53	11.20	43.94	17.63	65.34	24.91	78.61	30.26	86.13	33.87
Nr2N	23.72 _{0.93}	16.78 _{0.22}	50.83_{2.27}	22.13_{0.94}	71.77_{6.94}	26.80_{1.69}	70.32_{7.99}	26.40 _{3.17}	87.31_{4.70}	32.82_{0.72}
DDM2	26.02_{2.63}	17.92_{1.06}	49.57 _{15.6}	21.76 _{1.64}	59.64 _{8.47}	23.25 _{4.05}	77.96 _{2.16}	29.14_{1.01}	81.43 _{2.23}	31.35 _{2.23}
OURS	39.05_{1.42}	19.91_{0.32}	62.02_{1.11}	23.60_{0.05}	77.36_{0.61}	26.96_{0.37}	83.43_{1.39}	28.69_{0.67}	89.52_{0.18}	30.63 _{0.27}

Table 2. Quantitative results on simulated data. PSNR (dB) and SSIM (%) are reported. Numbers are presented as mean value with standard deviation. Best results are in bold. **OURS** holds a tremendous potential for generalization and applicability for its stable performance and better performance under high noise intensity

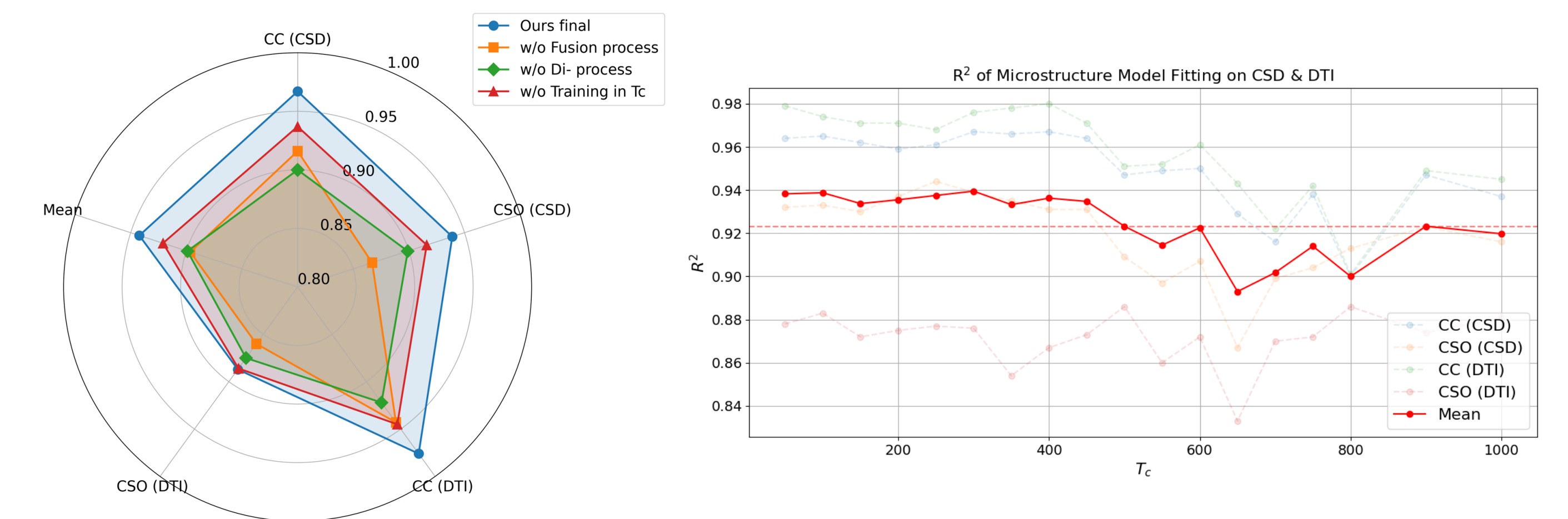


Figure 6. Ablation of our methods.

Figure 7. T_c vs. R^2 . When $T_c < 500$, the performance is consistent.

Limitations: (i) Long inference duration. (ii) Possible hallucinations. (iii) Noise-artifact conflation [5] and additive noise assumptions [6]. While our assumption enables tractable solutions, it may limit effectiveness for signal-dependent noise (better characterized as artifacts).

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