



DiffusionBlend

Learning 3D Image Prior through Position-aware Diffusion Score Blending for 3D Computed Tomography Reconstruction

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Deep Learning for Inverse Problems



Background Methods

Experiments

References



Figure: Learn the data-driven prior by deep learning when the measurement is sparse ¹

- Inverse problems arise from a wide range of applications across many domains, including computational imaging, remote sensing, and so on.
- The goal is to reconstruct an unknown signal x_{true} given the observed measurements y of the form $y = A(x_{true}) + \epsilon$, where ϵ can be an additive noise.



¹ Shen, Liyue et al., Patient-specific reconstruction of volumetric computed tomography images from a single projection view via deep learning, Nature biomedical enfgineering

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- Methods
- Experiments
- References

- **Supervised** Approaches (assuming that the x_{clean} , y pair is available during training, train a network that maps y to x_{clean})
 - Need to retrain for a different inverse problem
 - Generalization capabilities may be limited in the presence of noise/modality shift
 - Need paired data for training
- Unsupervised Approaches (assuming that only x_{clean} is available during training)
 - Easily adapt to a new inverse problem in a zero-shot manner.
 - Do not need paired data for training.

Both approaches are widely reported in the literature [8].



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Diffusion (Score-based) Models





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Mathematical Formulation of Diffusion Process



Introduction Background

Methods

Experiments

References



Reverse Generative Diffusion SDE:

$$d\mathbf{x}_{t} = \left[-\frac{1}{2}\beta(t)\mathbf{x}_{t} - \beta(t)\nabla_{\mathbf{x}_{t}}\log q_{t}(\mathbf{x}_{t})\right] dt + \sqrt{\beta(t)} d\bar{\omega}_{t}$$

"Score Function"

Figure: The mathematical formulation of diffusion process [1]



- The solution of the stochastic differential equation can be utilized by the score function
- we can use a neural network to approximate it, such as $s_{\theta}(x_t) \approx \nabla_{x_t} \log p(x_t)$

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Solving Inverse Problem with Diffusion Models





Methods

Experiment

References





Figure: the flowchart of Score SDE [4]. Each $\hat{\mathbf{x}}_t$ is modified through optimization

To solve linear inverse problems with diffusion model priors, we can use

- **hard consistency**: modify \mathbf{x}_t with optimization, such as with the objective $\operatorname{argmin}_z \lambda || \hat{\mathbf{x}}_t z ||_2^2 + (1 \lambda) || \hat{\mathbf{y}}_t Az ||_2^2$ [4, 3]
- **soft consistency**: change $\nabla_{x_t} \log p(x_t)$ to $\nabla_{x_t} \log p(x_t|y)$ via Bayesian rule [1, 5]. We have $\nabla_{x_t} \log p(x_t|y) = \nabla_{x_t} \log p(y|x_t) + \nabla_{x_t} \log p(x_t)$, with $\nabla_{x_t} \log p(y|x_t)$ can be approximated through $\nabla_{x_t} \log p(y|\hat{x}_0(x_t))$

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Methods

Experiments

References



Beer's law: Each material has a characteristic linear attenuation coefficient μ for rays emitted at a given frequency. More precisely,

 $\frac{dI}{ds} = -\mu(x(s))I(s),$

where s is the arc length along the trajectory described by the ray.

- The forward process is given by a linear transformation called "Radon Transform", which has the formula $Rf(\rho, \theta) = \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} f(x, y) \,\delta(x \cos \theta + y \sin \theta \rho) \, dx \, dy$.
- Denote the matrix of Radon Transform be A, the measurement (projections) is given by Ax + n, where n is usually a Poisson-Gaussian noise, and in a high-dose setting, we can just ignore this term.



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- Methods
- Experiments
- References

To reconstruct the original signal from the projections, we need to invert the Radon Transform process

- The backprojection formula is given by $f(x, y) = \int_0^{\pi} p_{\theta}(x \cos \theta + y \sin \theta) d\theta$. The vanilla backprojection without filtering gives blurry reconstruction.
- After adding a ramp filter to the backprojection, we are able to reconstruct high-quality images with full-view projections (like 180 projections). $f(x, y) = \int_0^{\pi} R^{-1} (F(\rho) * h(\rho)) (x \cos \theta + y \sin \theta) d\theta$
- Nevertheless, the filtered backprojection fails to reconstruct high-quality images when the forward operator is ill-posed (i.e. when projections are sparse).



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Methods

- Experiments
- References

- Most existing diffusion-based methods for CT reconstruction use the scores of 2D slices [8]. There is no existing works on the usage of cross-slice 3D scores for CT reconstruction
- We observe that learning the distribution of the score of a combination of 2D slices can be effectively used for 3D CT reconstruction



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- Methods
- **Experiments**
- References

Specifically, let $\nabla \log p_x[i]$ be the score of the i_{th} 2D CT slice. Existing works model $\nabla \log p_x \approx \sum_i \nabla \log p_x[i]$, and add external regularizations such as total variation to restrict the variability of inter-slice changes.

- We propose to learn the cross slice score $\nabla \log p_x[i_i, ..., i_k]$
- However, combining each cross slice score into the score of the entire volume is not a trivial task. i.e. how to write $\nabla \log p_x$ into a formula of $\nabla \log p_x[i_i, ..., i_k]$?



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- Methods
- Experiments
- References

We propose a novel method do this by add conditional slice information to the original summation equation

- Instead of using $\nabla \log p_x \approx \sum_i \nabla \log p_x[i]$, we can rewrite this into $\nabla \log p_x \approx \sum_i \nabla \log p(x[i]|x[i-1], x[i+1])$ This equation is more accurate if we assume that x[i] are not independent of each other (but dependent on neighboring slices).
- We then train a diffusion network, that takes a batch of 2D noisy slices as input and then denoise the central slice. We call this method DiffusionBlend



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Methods

Experiments

References

However, training a conditional diffusion model on noisy slices discard the joint slice distribution that we can learn.

- If we can learn $\nabla \log p(x[i], x[i-1], x[i+1])$, then we can compute $\nabla \log p(x[i]|x[i-1], x[i+1])$ through marginal probability.
- However, if we directly sum up ∇ log p(x[i], x[i − 1], x[i + 1]) for each three blocks, we will lose the dependency information of each blocks. i.e. we learn no information about p(x[i]|x[i + 2])
- We propose a novel method that randomly partition the 3D volumes into different joint slices blocks, and compute the score of the partition at each reverse sampling time step.



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Methods

- **Experiments**
- References

Why Random Partitioning work?

- Even though we may loss the cross-block conditional information at time t, with a different partition at t-1, the border of the original block incorporates the information of adjacency slices. So that it approximates the score of $\nabla \log p((x[i], x[i-1], x[i+1]) \cup (x[i+2], x[i+3]))$
- By doing random paritioning and score computing many times, we are able to approximate the score of the entire 3D volume.



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Overview of Our Method

We propose to randomly partition the volume at the reverse inference step and then compute the score respective of the partition while adding data consistency.



Introduction Background

Methods

Experiments

References



Bowen Song*, Jason Hu*, Zhaoxu Luo, Jeff Fessler, Liyue Shen



- Methods
- Experiments
- References

One drop back of the naive random paritioning method is that it only learns the score of adjacency slices

- We can design a skip slice partitioning score that not only learns the score of adjacency slices, but the dependency of jumping slices (for example: dependency of slice 1, slice 4 and slice 7).
- The intuition of this approach is that the naive cross-slice score $\nabla \log p(x[i], x[i-1], x[i+1])$ with the partitioniong $(x[i], x[i-1], x[i+1]) \cup (x[i+2], x[i+3], x[i+4]...)$ can have border artifacts.



Bowen Song*, Jason Hu*, Zhaoxu Luo, Jeff Fessler, Liyue Shen



- Methods
- Experiments
- References

We propose a novel method to learn the score of the joint borders. For example, for a border of a partition $(x[i], x[i-1], x[i+1]) \cup (x[i+2], x[i+3], x[i+4]) \cup (x[i+5], x[i+6], x[i+7]))$, we can learn the joint border score $\nabla \log p(x[i+1], x[i+4], x[i+7])$, and then apply the previous algorithm to blend with this score during reverse sampling.

- We preserve more inter-slice information through this joint border score.
- We are able to remove artifacts on the border of the partition.



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Methods

Experiments

References

We also want to keep all different score computation done by one model. If we use different diffusion models, the same noise may not lead to the same output. To do this we propose to use a **relative encoding** method:



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Methods

Experiments

References

- We propose to condition the diffusion model on the partition type, specifically, the distance between two slices in one block from the partition
- For instance, a partition of
 - $(x[i], x[i-1], x[i+1]) \cup (x[i+2], x[i+3], x[i+4]) \cup (x[i+5], x[i+6], x[i+7]))$ has the code to be 1, and a partition of $(x[i], x[i-3], x[i+3]) \cup (x[i+1], x[i-2], x[i+4]) \cup (x[i+2], x[i-1], x[i+5]))$ has the code to be 3
- Fourier embedding is then computed for the code, and then feed into the diffusion model



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Methods

Experiments

References



Algorithm PseudoCode:

Require: $A, M, \zeta_i > 0, k, y$ Initialize $x_T \sim \mathcal{N}(0, \sigma_T^2 I)$ for t = T : 1 do Randomly select a partition $S = S_1 \cup \ldots \cup S_r$ (the partition can either be of adjacency slices or jumping slices) Compute the relative positional encoding PE_t For each *i* compute $\epsilon_{\theta}(x_t[:, :, S_i], PE_t)$ Compute $s = \nabla \log p(x_t)$ using (9) Compute $\hat{x}_t = \mathbb{E}[x_0|x_t]$ using Tweedie's formula Set $\hat{x}'_t = CG(A^*A, A^*y, \hat{x}_t)$ Sample x_{t-1} using \hat{x}'_t and *s* via DDIM sampling

end for

Return \boldsymbol{x} .



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- Methods
- Experiments
- References

- We perform experiments on two CT datasets: AAPM LDCT, and LIDC.
- For each dataset, we perform two tasks: Sparse-View CT reconstruction and Limited-Angle CT reconstruction.
- For Sparse-view CT reconstruction, we consider three experimental settings: 4 views, 6 views and 8 views.
- For LACT, we consider 90 degree limited angle CT reconstruction
- Results are evaluate on the entire of a patient in the validation set of both AAPM and LIDC



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Methods

Experiments

References

Algorithm 1 DiffusionBlend++

Require: $A, M, \zeta_i > 0, k, y$ Initialize $x_T \sim \mathcal{N}(0, \sigma_T^2 I)$ for t = T : 1 do Randomly select a partition $S = S_1 \cup \ldots \cup S_r$ (the partition can either be of adjacency slices or jumping slices) Compute the relative positional encoding PE_t For each *i* compute $\epsilon_{\theta}(x_t[:,:,S_i], PE_t)$ Compute $s = \nabla \log p(x_t)$ using (9) Compute $\hat{x}_t = \mathbb{E}[x_0|x_t]$ using Tweedie's formula Set $\hat{x}'_t = CG(A^*A, A^*y, \hat{x}_t)$ Sample x_{t-1} using \hat{x}'_t and *s* via DDIM sampling end for

Return \boldsymbol{x} .



Bowen Song*, Jason Hu*, Zhaoxu Luo, Jeff Fessler, Liyue Shen



Introduction

- Background
- Methods
- **Experiments**
- References

	AAPM and LIDC Datasets						Sparse-View CT Reconstruction on LIDC					
Method	AAPM Dataset		LIDC Dataset		Average		8 Views		6 Views		4 Views	
	PSNR ↑	SSIM↑	PSNR ↑	SSIM↑	PSNR ↑	SSIM↑	PSNR ↑	SSIM↑	PSNR ↑	SSIM↑	PSNR↑	SSIM↑
FBP	22.63	0.206	14.70	0.369	18.67	0.288	14.78	0.206	14.10	0.181	13.10	0.165
FBP-UNet	28.44	0.930	29.54	0.887	28.99	0.909	28.87	0.858	26.59	0.793	25.37	0.744
DiffusionMBIR	27.63	0.908	-	-	-	-	33.29	0.922	31.69	0.903	29.21	0.868
TPDM	28.12	0.833	25.78	0.804	22.29	0.735	-	-	-	-	-	-
DDS 2D	28.82	0.922	27.96	0.842	28.39	0.882	31.60	0.898	29.99	0.871	28.03	0.830
DDS	29.03	0.934	28.13	0.879	28.58	0.903	32.51	0.920	30.83	0.924	27.61	0.828
DiffusionBlend (Ours)	36.62	0.972	31.02	0.924	33.82	0.948	34.47	0.934	31.16	0.908	28.24	0.859
DiffusionBlend++ (Ours)	37.45	0.976	34.64	0.935	36.02	0.946	35.66	0.947	33.97	0.935	31.38	0.913

Table 2: Comprehensive comparison of quantitative results on Limited-Angle and Sparse-View CT Reconstruction on Coronal View for AAPM and LIDC datasets. Best results are in bold.



Bowen Song*, Jason Hu*, Zhaoxu Luo, Jeff Fessler, Liyue Shen

References I



Introduction Background

Experiments

References



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DiffusionBlend

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Thank You



