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Anatomical image guided fluorescence molecular tomography reconstruction using kernel method

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Abstract. Fluorescence molecular tomography (FMT) is an important *in vivo* imaging modality to visualize physiological and pathological processes in small animals. However, FMT reconstruction is ill-posed and ill-conditioned due to strong optical scattering in deep tissues, which results in poor spatial resolution. It is well known that FMT image quality can be improved substantially by applying the structural guidance in the FMT reconstruction. In this paper, a new approach to introducing anatomical information into the FMT reconstruction with a Laplacian-type regularization matrix, the proposed method introduces the anatomical guidance into the projection model of FMT. The primary advantage of the proposed method is that it does not require segmentation of targets in the anatomical images. Numerical simulations and phantom experiments have been performed to demonstrate the proposed approach's feasibility. Numerical simulation results indicate that the proposed kernel method can separate two FMT targets with an edge-to-edge distance of 1 mm and is robust to false positive guidance and inhomogeneity in the anatomical image. For the phantom experiments with two FMT targets, the kernel method has reconstructed both targets successfully, which further validates the proposed kernel method. We have compared the proposed kernel method with the soft prior method thoroughly and found that the kernel method without target segmentation is able to achieve similar anatomical guided results as the soft prior method.

Keywords: Fluorescence molecular tomography, anatomical guidance, kernel method.

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1 Introduction

Fluorescence molecular tomography (FMT) has been emerging as an optical imaging modality for many years. FMT, as an important molecular imaging tool, has a broad range of applications in biomedical studies from drug development in small animal models¹⁻⁸ to the clinical diagnosis in human⁹⁻¹¹. However, due to the strong scattering nature of optical photons in deep tissues and a limited number of measurements, the inverse problem of FMT is ill-posed and underdetermined, which results in low spatial resolution in FMT imaging, in particular for targets in deep turbid media.

Many approaches have been proposed to improve the FMT image quality, including multispectral wavelengths for both excitation and emission wavelengths, different illumination patterns,^{12,13} a large number of FMT measurements by using charge-coupled device(CCD) cameras,¹⁴⁻¹⁸ and improved FMT reconstruction algorithms-- especially the sparse enhancement FMT reconstruction for the sparse FMT targets¹⁹⁻²⁴. A region reconstruction method implemented with level set method was also introduced to improve the FMT image reconstruction.^{25, 26} A thorough review of FMT imaging in terms of instruments, methods, and applications was presented in Ref. 27.

Although numerous efforts have been implemented to improve FMT, its spatial resolution is still worse than those of other functional imaging modalities such as functional magnetic resonance imaging (fMRI), single-photon emission computed tomography (SPECT), and positron emission tomography (PET). To further improve the spatial resolution of FMT, structural guidance from other anatomical images has been introduced into the FMT.^{4,28-30} Davis *et al.* reported the magnetic resonance imaging (MRI)-coupled FMT implemented with the Laplacian-type regularization.^{3,5,10,28,31} Schulz *et al.* reported a hybrid system for simultaneous FMT and X-ray computed tomography,^{7,8,29,32} Stuker *et al.* reported combined MRI and FMT system using single photon avalanche diode detectors.³³ Recently, microscopic positron emission tomography (microPET), with a spatial resolution up to 1 mm, has been used to guide FMT imaging.^{30,34} More recently, tri-modality³⁵ and even pentamodality, tomographic imaging systems were also investigated.³⁶

One of major challenges in the multimodality FMT system is how to utilize anatomical information properly and easily in the FMT reconstruction. Soft prior method is a widely accepted approach, which allows variations within the regions. Local Laplace and weighted

segments have also been introduced to FMT reconstruction.^{29,32} It has been demonstrated that the combination of Laplace with weighted segments performed best in terms of quantification and localization. However, both methods require image segmentation, which is time-consuming and prone to human error. To eliminate the need for direct prior image segmentation, Holt et al. reported a direct regularization method, in which the anatomical image gray-scale values are introduced into a regularization operator³⁷. Similarly, our proposed kernel method also eliminates the need of anatomical image segmentation. The major difference is that our approach does not need the regularization operator, which allows us to have maximum flexibility to implement this method.

In this paper, inspired by the kernel method in PET image reconstruction,³⁸ we introduce the kernel-based image reconstruction as a new approach to incorporating anatomical guidance into FMT. Compared with the Laplacian-type regularization methods, the proposed kernel method does not require the target region segmentation. Furthermore, as demonstrated by the numerical simulations in this paper, the proposed kernel method is robust to the false positive guidance and inhomogeneity in the anatomical images.

In the kernel method, the fluorophore concentration at a node *i* is defined as a function of a set of features, f_i , which is directly extracted from the voxel intensities of the corresponding anatomical 3D images. Then, the kernelized FMT image model is incorporated into the forward model of FMT. Due to the simplicity of this model, we can combine it with any FMT reconstruction algorithm. In this study, we used a kernelized projection model with the majorization-minimization (MM) approach.^{39,40}

The rest of this paper is organized as follows. In Section 2, we describe the FMT forward model, the regularized reconstruction method of FMT, and the proposed kernel-based

reconstruction algorithm. In Section 3, numerical simulations and experimental results are presented. Finally, we conclude the paper with discussions in Section 4.

2 Methods

2.1 Forward model and reconstruction algorithms of FMT

Light propagation in tissues is dominated by optical scattering and can be modeled by the diffusion equation.⁴¹ For FMT in the continuous wave (CW) domain, the light propagation model in 3D is described by a set of coupled differential equations which are given below:^{42,43}

$$\begin{cases} -\nabla \cdot [D_{ex}(\mathbf{r}) \nabla \Phi_{ex}(\mathbf{r})] + \mu_{\alpha,ex} \Phi_{ex}(\mathbf{r}) = \delta_s(\mathbf{r} - \mathbf{r}_s) \\ \mathbf{n} \cdot [D_{ex}(\mathbf{r}) \nabla \Phi_{ex}(\mathbf{r})] + \alpha_{ex} \Phi_{ex}(\mathbf{r}) = 0 \\ -\nabla \cdot [D_{em}(\mathbf{r}) \nabla \Phi_{em}(\mathbf{r})] + \mu_{\alpha,em} \Phi_{em}(\mathbf{r}) = \Phi_{ex}(\mathbf{r}) \mathbf{x}(\mathbf{r}) \\ \mathbf{n} \cdot [D_{em}(\mathbf{r}) \nabla \Phi_{em}(\mathbf{r})] + \alpha_{em} \Phi_{em}(\mathbf{r}) = 0 \end{cases}$$
(1)

where ∇ denotes the gradient operator, $D(\mathbf{r}) = \{3[\mu'_s(\mathbf{r}) + \mu_a(\mathbf{r})]\}^{-1}$ is the diffusion coefficient, $\mu_a(\mathbf{r})$ is the absorption coefficient, $\mu'_s(\mathbf{r})$ is the reduced scattering coefficient, $\Phi(\mathbf{r})$ is the photon fluence at the location \mathbf{r} , $\delta_s(\mathbf{r} - \mathbf{r}_s)$ is Dirac delta function defining point source, \mathbf{x} is the unknown to be reconstructed which is related to the fluorescent dye concentration and the quantum yield at each node,²³ \mathbf{n} is the outward unit normal vector of the boundary, and α is the Robin boundary coefficient. In Eq. (1), subscripts of ex and em mean corresponding terms at the excitation and emission wavelengths, respectively. Eq. (1) can be solved by the finite element method (FEM) based on a finite element mesh and is linearized to the following equation:

$$K_{ex}\Phi_{ex} = \delta_s(\boldsymbol{r} - \boldsymbol{r}_s), \qquad K_{em}\Phi_{em} = \Phi_{ex}\boldsymbol{x}$$
(2)

where K_{ex} and K_{em} are the stiffness matrices at the excitation and emission wavelengths, respectively. With the conjugate gradient approach,¹⁶ the above equations can be described as:⁴²

$$Ax = b \tag{3}$$

where $A \in \mathbb{R}^{\{N_m \times N_n\}}$ is the system matrix, $x \in \mathbb{R}^{\{N_n \times 1\}}$ is the unknown related to fluorophore distribution or the FMT image to be reconstructed, $b \in \mathbb{R}^{\{N_m \times 1\}}$ is the measurement vector, N_n is the finite element node number, and N_m is the number of measurement.

Because of the ill-conditioned and ill-posed nature, Eq. (3) is usually solved as a regularized least square problem with the non-negativity constraint:

$$\mathbf{x} = \underset{\mathbf{x}, \mathbf{x} \ge 0}{\operatorname{argmin}} \Phi(\mathbf{x}) =: ||\mathbf{A}\mathbf{x} - \mathbf{b}||_{2}^{2} + \lambda ||\mathbf{x}||_{1} \}$$
(4)

where λ is L_1 the regularization parameter.

In this study, for the case without anatomical guidance, Eq. (4) is solved by the MM approach that updates the FMT image iteratively to minimize the mismatch between the model predictions and the measurements.^{39, 40, 43}

2.2 Soft prior method

When structured priors are present, the objective function of the FMT with Laplacian regularization will be:

$$\mathbf{x} = \underset{\mathbf{x}, \mathbf{x} \ge 0}{\operatorname{argmin}} \Phi(\mathbf{x}) =: ||\mathbf{A}\mathbf{x} - \mathbf{b}||_{2}^{2} + \lambda ||\mathbf{L}\mathbf{x}||_{2}^{2} \}$$
(5)

In soft prior method, regularization matrix L is defined as:¹⁰

$$L_{ij} = \begin{cases} 1, & \text{for } i = j \\ -\frac{1}{N}, & \text{if } i \text{ and } j \text{ are in the same region} \\ 0, & \text{otherwise} \end{cases}$$
(7)

Where N is number of nodes in that region. In Eq. (5), regularization term $||Lx||_2^2$ can be treated as special case of $||Ax - b||_2^2$ when b=0. Then it can be solved by the MM approach as described in Refs. 39, 40 and 43.

2.3 Kernel based anatomically-aided reconstruction algorithm

The anatomically-aided FMT reconstruction algorithms usually incorporate the anatomical guidance as a regularization matrix, which enhances the smoothness within the anatomical regions and also allows sharp transition between the different regions.³⁷ In this paper, we introduce the kernel method which includes the anatomical guidance into the projection model of FMT. The fluorophore distribution at the node *i* is defined with a kernel function as^{38,44,45}

$$\boldsymbol{x}_i = \sum_j \alpha_j \kappa(\boldsymbol{f}_i, \boldsymbol{f}_j) \tag{7}$$

where f_i and f_j are the anatomical feature vectors corresponding to the finite element nodes of *i* and *j*, respectively. These anatomical feature vectors are directly extracted from the corresponding voxels in the 3D anatomical images for each finite element node. The finite element mesh and the anatomical images should be co-registered. In some reported multimodality FMT systems, accurate co-registrations were reported,^{7,33} which makes the proposed kernel method to be implemented easily. It is also worth pointing out that voxels corresponding to finite element nodes on the surface of the mesh and outside of the mesh are excluded from the feature vector extraction. The length of the feature vectors depends on the voxels number. For example, for a voxel number of $3 \times 3 \times 3$, the length of the feature vector is 27.

In Eq. (7), κ is the kernel function. There are a variety of choices of the kernel function κ .^{46,47} Here we use the radial Gaussian kernel:⁴⁸

$$\kappa(\boldsymbol{f}_{i}, \boldsymbol{f}_{j}) = exp(\frac{-\left|\left|f_{i} - f_{j}\right|\right|^{2}}{\sigma^{2}})$$
(8)

where the parameter σ controls the edge sensitivity and yields more accurate results when $\sigma = 1.^{49}$ For computational efficiency, a *k*-nearest neighbor (*knn*) search is carried out for each feature vector corresponding to each finite element node using the *knnsearch* function in MATLAB. The search is carried out according to the Euclidean distance between the feature

vectors, not the physical distance between the finite element nodes in the Cartesian coordinate. Only those elements corresponding to the *k*-nearest neighbors are stored in the kernel matrix and the rest of them are set to be 0. This will result in the following definition of the kernel matrix:

$$K_{ij} = \begin{cases} \kappa(\boldsymbol{f}_i, \boldsymbol{f}_j), \ \boldsymbol{f}_j \in knn \text{ of } \boldsymbol{f}_i \\ 0, & \text{otherwise} \end{cases}$$
(9)

Thus, the kernel matrix is a sparse symmetric $N_n \times N_n$ matrix. The kernel matrix is normalized in this study for higher image quality:³⁸

$$\overline{\mathbf{K}} = diag^{-1}[\mathbf{K}\mathbf{1}_N]\mathbf{K} \tag{10}$$

here $\mathbf{1}_N$ is a vector of all ones. Eq. (7) can be written in a matrix-vector form as:

$$\boldsymbol{x} = \boldsymbol{\overline{K}}\boldsymbol{\alpha} \tag{11}$$

where the vector $\boldsymbol{\alpha}$ is a new unknown vector referred as the coefficient image. By substituting Eq. (11) into Eq. (3), the kernelized projection model of FMT can be written as

$$A\overline{K}\alpha = b \tag{12}$$

Combining the kernelized projection model of Eq. (12) with the objective function of Eq. (4) leads to the following objective function:

$$\boldsymbol{\alpha} = \operatorname{argmin} \Phi(\boldsymbol{\alpha}) =: ||\boldsymbol{A}\overline{\boldsymbol{K}}\boldsymbol{\alpha} - \boldsymbol{b}||_{2}^{2} + \lambda ||\boldsymbol{\alpha}||_{1} \}$$
(13)

Because the reconstructed images are already regularized by the kernels, we set the regularization parameter in Eq. (13) to zero in this study,³⁸ and solved by the MM approach.^{39,40} Once α is obtained we can easily obtain the final fluorophore distribution image by the linear transformation $\mathbf{x} = \overline{\mathbf{K}} \alpha$.

2.4 Numerical simulation setup

2.4.1 Cylindrical simulation phantom

In this simulation, we used a cylindrical phantom with a diameter of 22 mm and a height of 80 mm. Cylindrical targets with a diameter of 1.4 mm and a length of 20 mm were embedded 20 mm below the top surface of the phantom. In the coordinate system, the base of the cylinder was a circle on the x-y plane centering the origin of the coordinate system and the height was along the *z* axis. In this simulation, two targets were embedded at (-1.7, 5.56) and (1.7, 5.56) in the x-y plane with an edge-to-edge distance of 2 mm as shown in Fig. 1(a).

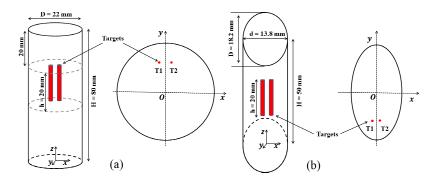


Fig. 1 Numerical simulation phantom geometry of (a) the cylindrical phantom with target locations at T1 (-1.7, 5.56) and T2 (1.7, 5.56) and (b) the elliptic cylindrical phantom with target locations at T1 (-1.2, -5.0) and T2 (1.2, -5.0).

In this and following numerical simulations, the phantom tissue optical properties were set to be $\mu_a = 0.012 \text{ mm}^{-1}$ and $\mu'_s = 0.83 \text{ mm}^{-1}$ at both the excitation wavelength (650 nm) and the emission wavelength (700 nm). We assigned the fluorophore concentration to be 1 in the target regions and 0 in the background regions.

The numerical phantom was discretized with a 3-dimensional (3D) tetrahedral finite element mesh with 29,989 nodes and 155,310 elements. Numerical FMT measurement data were generated by Eq. (3) with a line pattern laser projected on the phantom surface. The line laser had a width of 1 mm and a length of 50 mm. We had 30 excitation positions of the line laser to cover the whole surface.⁵⁰ For each line laser excitation, the 9,280 surface nodes on the side of the cylinder were used as the measurement detectors. Then, we added 30% Gaussian noise to the numerical FMT measurement data.

The 3D CT images with 220×220×801 voxels were generated with a grid size of 0.1 mm. The intensities of target regions and the background were set to be 0.24 and 0.06, respectively, which were close to the CT data in the phantom experiments. We added 15% white Gaussian noise to the numerical CT images.

2.4.2 Elliptic Cylindrical simulation phantom

In this simulation, we used an elliptic cylindrical phantom with a horizontal semi-axis of 6.9 mm, vertical semi-axis of 9.2 mm and a height of 50 mm. Cylindrical targets with a diameter of 1.4 mm and a length of 20 mm were embedded 20 mm below the top surface of the phantom. In the coordinate system, the base of the cylinder was an ellipse on the x-y plane centering the origin of the coordinate system and the height was along the z-axis. Two targets were embedded at (-1.2,-5.0) and (1.2,-5.0) in the x-y plane with an edge-to-edge distance of 1 mm as shown in Fig. 1(b). The numerical phantom was discretized with a three-dimensional (3D) tetrahedral finite element mesh with 32,882 nodes and 191,359 elements. Numerical FMT measurement data were generated by Eq. (3) with a line pattern laser projected on the phantom surface. The line laser had a width of 1 mm and a length of 50 mm. We had 30 excitation positions of the line laser to cover the whole surface.⁵⁰ For each line laser excitation, the 6,013 surface nodes on the side surface of the cylinder were used as the measurement detectors. Then, we added 30% Gaussian noise to the numerical FMT measurement data.

The transverse sections of the CT images were generated using the "*phantom*" command in MATLAB (as shown in Fig. 5(a) in the result section). The region outside of the ellipse was

trimmed. Then we stacked the "*phantom*" images to generate 3D CT images with 234×176×501 voxels. Two targets with a diameter of 1.4 mm and a length of 20 mm were added in the 3D CT phantom. The intensity of targets is 0.99, which is 1% less than the intensity of edges of the ellipse. This is because the tumor with CT contrast agent injection has contrast as high as bones in CT images.²⁹ As in the first simulation, we added 15% Gaussian noise to the numerical CT images.

2.4.3 Numerical simulation using MRI images of a rat brain

The ultimate goal of the proposed kernel method is its application in anatomical image (such as CT or MRI) guided FMT for small animal studies. To validate the feasibility of the proposed method using *in vivo* anatomical guidance with heterogeneous structures, we used MRI image of a rat brain as the anatomical guidance. MRI imaging was performed with a Bruker Biospec 7 Tesla (7T) small-animal scanner (Bruker BioSpin MRI, Ettlingen, Germany). A 72 mm internal diameter linear resonator was used for radio frequency (RF) transmission, and a four-channel rat brain phased array surface coil was used for signal reception. The rat brain was imaged coronally with a fast-spin echo sequence (RARE; axial: TE/TR = 8 ms/750 ms; FOV = 40×40 mm²; MTX= 256×256 ; ST/SI = 1 mm/1 mm; ETL = 4). Data were acquired and reconstructed using ParaVision 5.1 software (Bruker BioSpin MRI). The experiment was conducted under a protocol approved by the University of California, Davis, Animal Use and Care Committee (Davis, CA). A male athymic nude rat, purchased from Harlan Laboratories (Hayward, CA), was inoculated with 3×10mm⁶ U87 MG cells/10 µL intracranially. The rat was administered 0.5 mmol/kg of the small molecule gadolinium chelate, gadoteridol (Bracco Imaging) via bolus i.v. injection prior to T1w imaging.

From the MRI images, we used the open-source software, *iso2mesh*, to generated a 3D finite element mesh with 181,686 tetrahedral elements, and 41,427 nodes.⁵¹ We segmented the tumor in the MRI images as the FMT target region. Similar to the numerical phantom studies, numerical FMT measurement data was generated by Eq. (3) with a line pattern laser projected on the rat brain surface. We had 30 excitation positions of the line laser to cover the whole rat brain surface.⁵⁰ For each line laser excitation, the 20,055 surface nodes were used as the measurement detectors. The tissue optical properties were set to μ_a =0.012 mm⁻¹ and μ'_s =0.83 mm⁻¹ at both the excitation wavelength (650 nm) and the emission wavelength (700 nm). We assigned the fluorophore concentration to be 1 in the target regions and 0 in the background regions. Like the numerical simulation studies described above, we added 30% Gaussian noise to the numerical FMT measurement data.

In the kernel method, we extracted the feature vectors from the MRI images easily because the finite element mesh was generated from the same MRI images. Then we generated the kernel matrix using the Eqs. (8) and (9) to incorporate anatomical information from MRI images into the FMT reconstruction by minimizing the kernelized objective function as described in Eq. (13).

The voxel number for each corresponding node and the number of nearest neighbors in *knnsearch* are important parameters in generating the kernel matrix K and have significant effects on the kernel reconstruction method. In this paper we studied 3 different voxel numbers, $3\times3\times3$, $5\times5\times5$, and $7\times7\times7$. The lengths of feature vectors were 27, 125, and 343 respectively. For *knnsearch*, different values of *k* (16, 32, 64, 128, 256), the number of nearest neighbors, were also studied.

During the FMT reconstruction, for the kernel method, we used the MRI images without segmentation as the anatomical guidance to generate the K matrix. For the soft prior method, we

used the segmented images to generate the soft priors matrix without adding any segmentation error.

2.5 Phantom experimental setup

To validate our algorithm, we conducted an agar phantom experiment. In this experiment, we used a cylindrical phantom with a diameter of 22 mm and a length of 80 mm. The phantom was composed of 1% intralipid, 2% agar, 20 μ M bovine hemoglobin (H2625, Sigma-Aldrich Inc., St. Louis, MO) and water. We embedded two capillary tubes with a length of 20 mm and a diameter of 1.4 mm as targets, in which 20 μ M Sulfo-Cyanine5 dye (Lumiprobe Corporation, Hallandale Beach, FL) was injected. The geometry of the experimental phantom with two targets is shown in Fig 2, where the two red bars indicate two FMT targets. The edge-to-edge distance of the two targets was 2.94 mm.

The phantom geometry was discretized with a 3D tetrahedral finite element mesh with 37,333 nodes and 199,881 elements. During the FMT imaging, a line laser (1 mm wide and 50 mm long) at the wavelength of 643 nm scanned the surface of the phantom sequentially with 30 excitation positions that were distributed uniformly on the phantom surface. For each line laser excitation position, an emission picture at the wavelength of 720 nm was taken. All 9,384 surface nodes on the side surface of the cylinder were used as the detector nodes, and the measurements were obtained from the acquired emission pictures. Details of the conical mirror based FMT imaging system were described in Ref. 52. The phantom optical properties were $\mu_a = 0.012 \text{ mm}^{-1}$, $\mu'_s = 0.83 \text{ mm}^{-1}$ at both 643 nm and 720 nm wavelengths.

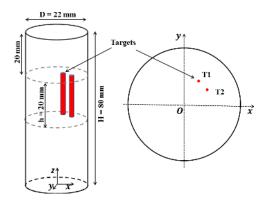


Fig. 2 The geometry of the phantom experiment with target locations at T1 (1.72, 4.71) and T2 (5.01, 1.87).

We scanned the phantom with our lab-made microCT imaging system with 180 projections⁵³ and reconstructed the 3D CT images of the phantom with an isotropic voxel size of 0.15 mm. The micro-CT system consisted of an X-ray source and a flat panel detector placed opposite to each other on a micro-CT gantry that rotated around the bed where the phantom was placed. The source-to-isocenter distance was 205.34 mm, and the source-to-detector distance was 246.2 mm. The detector had a 49.2 mm by 49.2 mm sensing area consisting of 1024 by 1024 pixel sensors with 48 µm pixel spacing. The X-ray tube was operated at a current of 0.5 mA and a voltage of 50 kVp. A filtered backprojection algorithm was used to reconstruct the micro-CT images with a Shepp-Logan filter. The obtained CT images are shown in Fig. 9(a), from which we calculated the targets' size and position. Because Sulfo-Cyanine5 dye does not have CT contrast, only the capillary tubes were observed in the reconstructed microCT images, and the fluorescence dye (target) regions were filled by pixels having the same CT contrast as the capillary tubes as guidance in the kernel method.

2.6 FMT image evaluation criteria

According to our previous studies, the combinations of 4 metrics listed below can evaluate the quality of the reconstructed FMT images very well. Their detailed definitions can be found in

Refs. 39 and 40. Briefly, the Volume Ratio (VR) measures the ratio between the true region of interest (ROI) and the reconstructed region of interest (rROI). The Dice similarity coefficient (Dice) measures the location accuracy of the reconstructed target. Ideally, VR and Dice coefficients should be 1. The Contrast-to-Noise Ratio (CNR) measures how well the reconstructed target is distinguished from its background. The higher the CNR coefficient is, the better the reconstructed image. The Mean Square Error (MSE) is the difference between the measurements and the model predictions. The MSE closer to zero is better.

3 Results

3. 1 Simulation Results

3.1.1 Numerical simulation with two FMT targets

In this simulation, we had two capillary tube targets embedded inside the cylindrical background phantom with an edge-to-edge distance of 2 mm as described in Fig. 1(a). For comparison, we have reconstructed FMT images with the soft prior method. The ground truth FMT images, simulated CT images and the reconstructed FMT images with the soft prior method are plotted in Fig. 3. All the FMT reconstructions in this paper were conducted in 3D and the reconstructed FMT images are shown by slices along the z-axis with equal distance. Then, we performed the reconstruction with the proposed kernel based FMT reconstruction algorithm. To investigate how the parameters in the kernel method affect the FMT reconstruction, we studied three different voxel numbers ($3 \times 3 \times 3$, $5 \times 5 \times 5$, and $7 \times 7 \times 7$) and three different nearest neighbors (k=16, 32, 64) with 9 combinations of the kernel based FMT reconstructions. The reconstructed FMT images are plotted in Fig. 4, in which each column indicates different voxel numbers and each row

indicates different numbers of nearest neighbors. For all 9 cases, the two targets have been reconstructed and separated successfully as indicated by Fig. 4.

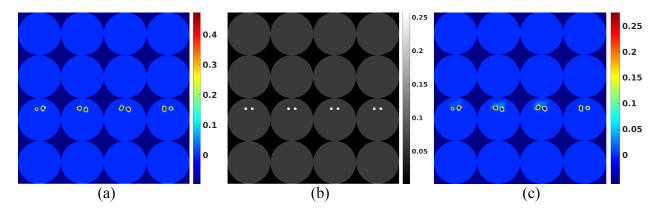


Fig. 3 For the numerical simulation of two targets: (a) the ground truth images,

(b) simulated anatomical guidance images, and (c) the reconstructed FMT images with the soft prior method. The

distance between slices along z-axis is 5.33 mm.

To evaluate the simulation results quantitatively, we calculated image quality metrics such as VR, Dice, CNR and MSE for the FMT reconstruction with the soft prior mehtod, and the 9 FMT reconstructions with the kernel method, as shown in Table 1. For the kernel method, when the voxel number is fixed, we have better FMT reconstruction quality as the nearest neighbor k increases. One example is that the Dice increased from 0.02 to 0.23 as k increased from 16 to 64 for the voxel number of $3\times3\times3$. Similarly, for the fixed nearest neighbor k, we found that the FMT image quality becomes better with larger voxel number. The best FMT reconstruction result was obtained with k=64 and a voxel number of $7\times7\times7$, which is highlighted in Table 1. From Table 1, we see that the soft prior method performed better than the kernel method in this simulation when the target regions were known accurately in the anatomical guidance.

Table 1 For the cylindrical phantom simulation of 2 targets, the calculated VR, Dice, CNR and MSE with the kernel method for different numbers of nearest neighbor *k* and different voxel numbers and the soft prior method.

k	Voxel number	VR	Dice	CNR	MSE	
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16	3×3×3	0.124	0.002	18.264	3.88e-4
16	5×5×5	0.223	0.159	21.351	2.96e-4
16	7×7×7	0.387	0.306	19.554	3.44e-4
32	3×3×3	0.127	0.006	19.688	3.41e-4
32	5×5×5	0.243	0.187	22.713	2.51e-4
32	7×7×7	0.451	0.430	22.099	2.66e-4
64	3×3×3	0.139	0.23	20.029	3.29e-4
64	5×5×5	0.347	0.323	23.956	2.22e-4
64	7×7×7	0.639	0.596	24.111	2.21e-4
	Soft prior	0.952	0.964	32.355	1.67e-4

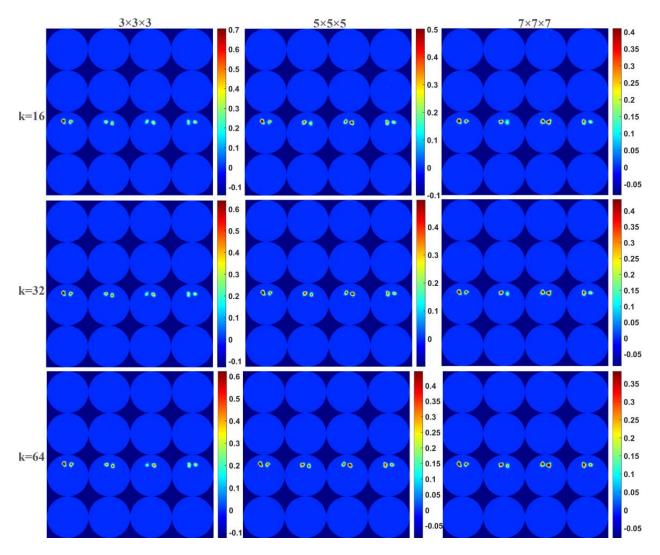


Fig. 4 Reconstruction FMT images for the cylindrical phantom simulation of 2 targets by the kernel method with different nearest neighbor k as indicated by each row and different voxel numbers indicated by each column. The distance between slices along z-axis is 5.33 mm.

3.1.2 Elliptic cylindrical phantom simulation with two FMT targets

In this simulation, we had two capillary tube targets embedded inside the elliptic cylindrical background phantom with an edge-to-edge distance of 1 mm as described in Fig. 1(b). For comparison, we have also reconstructed FMT images with the soft prior method. The ground truth FMT images, the simulate CT images, and the reconstructed FMT images with the soft prior method are plotted in Fig. 5. Then, we performed the reconstruction with the proposed kernel based FMT reconstruction algorithm. To investigate how the parameters in the kernel method affect the FMT reconstruction, we studied 3 different voxel numbers ($3\times3\times3$, $5\times5\times5$, and $7\times7\times7$) and 3 different nearest neighbors (k = 64, 128, 256) with 9 combinations of the kernel based FMT reconstructions. The reconstructed FMT images with the kernel method are plotted in Fig. 6, in which each column indicates different voxel numbers and each row indicates different numbers of nearest neighbors. For all 9 cases, the two targets have been reconstructed and separated successfully as indicated by Fig. 6.

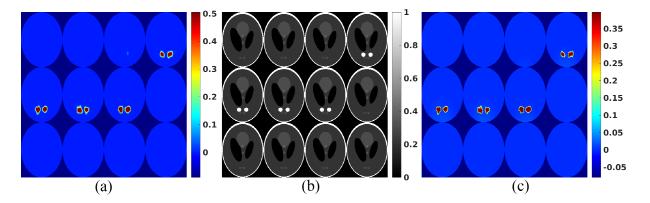


Fig. 5 For the simulation of elliptic cylindrical phantom with 2 FMT targets, (a) the ground truth images and (b) simulated CT images (c) the reconstructed FMT images with the soft prior method. The distance between slices along z-axis is 4.54 mm.

To evaluate the simulation results quantitatively, we calculated quantitative image quality metrics for the FMT reconstruction with the soft priors method and the 9 FMT reconstructions with the kernel method, as shown in Table 2. From Table 2, when the voxel number is fixed, we have better FMT reconstruction quality as the nearest neighbor k increases. One example is that the Dice increased from 0.469 to 0.803 as k increased from 64 to 256 for the voxel number of $3\times3\times3$. Similarly, for the fixed nearest neighbor k, we found that the FMT image quality becomes better with larger voxel number in this simulation setup up. The best FMT reconstruction result was obtained with k = 256 with $7\times7\times7$ voxel size, which is highlighted in Table 2.

Table 2 For the numerical simulation of elliptic cylindrical phantom with 2 FMT targets, the calculated VR, Dice,CNR and MSE with the kernel method for different numbers of nearest neighbor k and different voxel numbers, andwith the soft prior method.

k	Voxel number	VR	Dice	CNR	MSE
64	3×3×3	0.366	0.469	18.971	6.22e-4
64	5×5×5	0.318	0.419	19.452	6.16e-4
64	7×7×7	0.519	0.635	22.262	4.82e-4
128	3×3×3	0.516	0.604	20.811	5.16e-4
128	5×5×5	0.481	0.590	21.113	5.39e-4
128	7×7×7	0.519	0.653	23.072	4.57e-4
256	3×3×3	0.713	0.803	25.028	1.35e-4
256	5×5×5	0.668	0.782	25.286	1.46e-4
256	7×7×7	0.757	0.845	26.108	1.21e-4
	Soft prior	1.046	0.934	26.661	1.73e-4

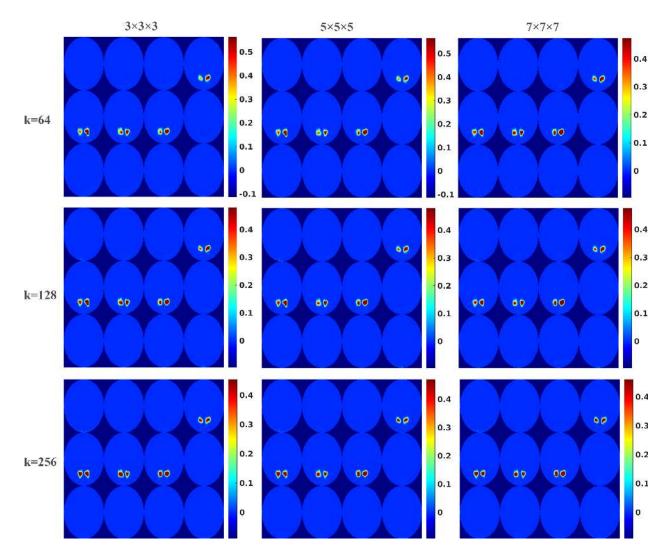


Fig. 6 Reconstructed FMT images for the elliptic cylindrical phantom simulation of 2 targets by the kernel method with different nearest neighbor k as indicated by each row and different voxel numbers indicated by each column. The distance between slices along z-axis is 4.54 mm.

3.1.3 Numerical simulation with false target size in the numerical anatomical CT images

The numerical phantom geometry of this simulation study same as the second simulation is plotted in Fig. 1b. However, the diameter of the right target in the simulated anatomical guidance CT images (Fig. 7b) was enlarged intentionally from 1.4 mm to 2.8 mm to study how the false target size affects the FMT reconstruction with the proposed kernel method. The enlarged target

was moved to the right side for 0.7 mm, so that the edge-to-edge distance of the two targets was still 1mm in the simulated CT images as shown in Fig. 7(b). For comparison, we have performed the reconstruction with the soft prior method and with the kernel method of 3 different nearest neighbor k and 3 different voxel numbers as in the above section. Among all the reconstructions with the kernel method, unlike the previous simulation, we found that the reconstruction with the nearest neighbor of k=256 and the voxel number of $3 \times 3 \times 3$ had the least error with an MSE of 8.54e-4, whereas the soft prior method had the MSE of 1.31e-3. Fig. 7 plots the ground truth images (Fig. 7a), the reconstructed FMT images with soft prior method (Fig. 7c), and the reconstructed FMT images by the kernel method with the nearest neighbor of k = 256 and the voxel number of $3 \times 3 \times 3$ (Fig.7d). As indicated in Fig.7c, the two targets were barely separated with the soft prior method. Fig.7d is a representative reconstructed FMT image with the kernel method and indicates that the image quality is much better than that of Fig. 7c as demonstrated by the CNR of 12.214 for Fig. 7c and 17.543 for Fig. 7d. The calculated image quality metrics are listed in Table 3.

Table 3 For the numerical simulation with a false target size, the calculated VR, Dice, CNR and MSE with the kernel method of different numbers of nearest neighbor k and different voxel numbers, and with the soft prior method.

k	Voxel number	VR	Dice	CNR	MSE
256	3×3×3	0.394	0.548	17.543	8.54e-4
256	5×5×5	0.281	0.416	16.816	8.91e-4
256	7×7×7	0.409	0.537	16.728	9.06e-4
	Soft prior	0.466	0.527	12.214	1.31e-3

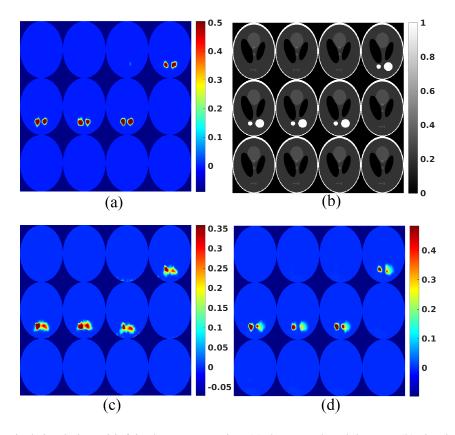


Fig. 7 For numerical simulation with false larger target size, (a) the ground truth images, (b) simulated CT images with the false enlarged target. (c) the reconstructed FMT images with the soft prior method, and (d) the reconstructed FMT images by the kernel method with the nearest neighbor of k = 256 and the voxel number of $3 \times 3 \times 3$. The distance between slices along z-axis is 4.54 mm.

3.1.4 Numerical simulation using MRI images of a rat brain

To validate the proposed kernel based FMT image reconstruction algorithm with guidance from realistic anatomical images, we conducted this simulation study using *in vivo* rat brain MRI images as shown in Fig. 8(a). Details of the MRI images and the simulation setup described in Section. 2.4.3. The contrast of the brain tumor to its surrounding tissues in the MRI images is 2:1 approximately. First, we reconstructed the FMT images using the soft prior method, in which only two regions were considered. We have obtained very good FMT results from the soft prior

method as shown in Fig. 8(b). Fig. 8(c) plots the reconstructed FMT images obtained by the kernel method using the MRI images as the anatomical guidance directly without segmentation. In the kernel method, we set the nearest neighbor of k = 256 and the voxel number of $3 \times 3 \times 3$. From Fig. 8, we can see that the kernel method has reconstructed the target very well with comparable results from the soft prior method. The VR, Dice, CNR and MSE are 0.529, 0.626, and 23.007 and 7.14e-4 for the kernel method, and 0.966, 0.974, 42.622, and 3.12e-4 for the soft prior method, respectively.

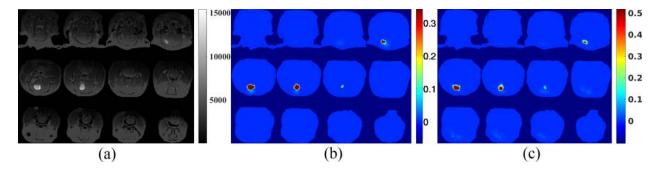


Fig. 8 Numerical simulation with the rat brain MRI images: (a) MRI images; the FMT reconstructed images with (b) the soft prior method and (c) the kernel method with k = 256 and the voxel number of $3 \times 3 \times 3$. The distance between slices along z-axis is 2.45 mm.

3.2 Phantom experimental results

3.2.1 Reconstruction with homogeneous background in CT images

The phantom's geometry is plotted in Fig. 2. As described in the numerical simulation section, we have performed the FMT reconstruction of this phantom experiment with the soft prior method and with kernel method of 15 different cases with 5 different nearest neighbor *k* (16, 32, 64, 128, 256) and 3 different voxel numbers $(3 \times 3 \times 3, 5 \times 5 \times 5, 7 \times 7 \times 7)$. The reconstructed FMT

images along with anatomical CT images are plotted in Fig. 9. The kernel based reconstruction results (Fig. 9(d)) are as good as the results from the soft prior method (Fig. 9(c)) when the homogeneous anatomical images were used as the guidance. For comparison, we have also reconstructed the target without anatomical guidance as shown in Fig. 9(b), from which we see that the two targets were reconstructed with large position errors. To analyze the reconstructed FMT images quantitatively, we have calculated the VR, Dice, and CNR as listed in Table 4 for each case, where the microCT images were referred as the ground truth images when we calculated the image quality metrics. The MSE has not been calculated because we do not know the exact fluorescent dye concentration. From Table 4, we know that the kernel method can achieve good reconstruction results with the nearest neighbor of 64 and the voxel number of $5 \times 5 \times 5$, in which the VR, Dice, and CNR are 0.714, 0.643, and 25.849, respectively. The VR, Dice, and CNR are 0.717, 0.740, and 29.846 for the FMT reconstruction with the soft prior method. These similar image quality metrics indicate that the kernel method is as good as the soft prior method for FMT target reconstruction with the homogeneous background.

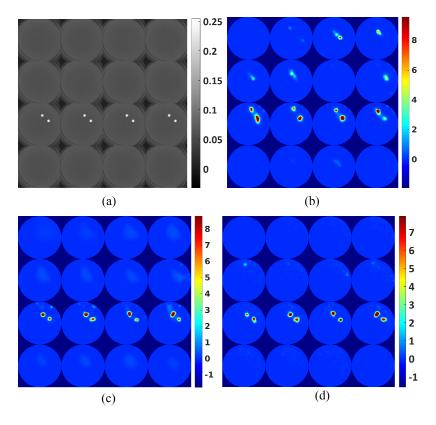


Fig 9 (a) Original CT images. The reconstructed FMT images (b) without priors, (c) with the soft prior method and the homogenous background, (d) with the kernel method using original CT images as guidance with k = 64 and the voxel number of 5×5×5. The distance between slices along z-axis is 5.33 mm.

 Table 4 The calculated VR, Dice and CNR for the phantom experiments without prior, with the kernel method, and

with the soft prior method.

	Homogeneous			Inhomogeneous			
	VR	Dice	CNR	VR	Dice	CNR	
Soft prior	0.729	0.757	30.312	0.677	0.728	29.704	
Kernel method	0.714	0.643	25.849	0.672	0.648	28.250	
No prior	0.752	0.025	2.569				

3.2.2 Reconstruction with inhomogeneous background in CT images

To further validate the proposed method in a more complex anatomical images, we added some artificial features in the physical CT image we obtained. As shown in Fig. 10(a), the darkest big cylinder has an intensity of less than 50% of the background. The other two big cylinders have

an intensity of 50% more than the background intensity. We also added another three small cylinders at the random locations with different intensities. Two of them have an intensity of 5 times more than the background, which is slightly higher than the targets' intensity. Those bright inclusions mimic bones in the CT images or fat and blood in the MRI images. For the FMT reconstruction with the soft prior method, we had 6 regions: two targets, three big cylinder artificial features and the background. Unlike the homogeneous background case, here we obtained the best kernel method based FMT images with the nearest neighbor of 256 and the voxel number of $3 \times 3 \times 3$, in which the VR, Dice, and CNR are 0.672, 0.648, and 28.250, respectively. These image quality metrics are slightly lower than those of the soft prior method, which are 0.717, 0.740, and 29.846 respectively. These results demonstrated that the kernel method is able to achieve comparable results with as the soft prior method when there are inhomogeneous inclusions in the anatomical guidance images.

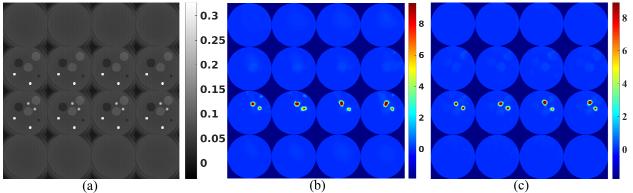


Fig 10 (a) CT images with artificial features. (b) Reconstructed FMT images with the soft prior method and the inhomogenous background in the CT images. (c) Reconstructed FMT images with the kernel method using the CT images with artificial features as the guidance with k = 256 and the voxel number of $3 \times 3 \times 3$. The distance between slices along z-axis is 5.33 mm.

4 Discussions and Conclusion

In this paper, we have introduced the kernel method as a new approach to including the anatomical guidance in the FMT image reconstruction, in which a kernel matrix having the anatomical priors is created and incorporated into the projection model of FMT. It is worth noting that we used the forward model without the kernel method to generate the FMT measurements for numerical simulations. Numerical simulations and phantom experiments have been performed to demonstrate that the proposed kernel method has reconstructed the FMT targets successfully and have comparable results as the soft prior method.

Compared with conventional Laplacian-type regularization method to include anatomical priors such as the soft priors, the kernel method has the advantage of easy implementation, in which we do not need to segment the target and background regions in the anatomical images. This advantage is more significant for some cases in which the targets are not easily differentiated and segmented. This may result in a concern of the misguidance from the false positive regions. To address this issue, we have performed one numerical simulation with a false target size (from 1.4 mm to 2.8 mm in diameters) as described in section 3.1.3. Our results indicate the false target size guidance has some effects when two targets are very close. However, the kernel method performs better than the soft prior method. Another advantage of the kernel method is that we do not have to search for the optimum regularization parameter, which is searched with the L-curve method in conventional regularization methods.

To generate the kernel matrix, three parameters must be set before the FMT image reconstruction. The first parameter is the Gaussian kernel coefficient σ . According to previous studies, $^{38,49}\sigma=1$ yields best results. The second parameter is the number of nearest neighbor's *k*. From the results of both numerical simulations and phantom experiments, we found that the

reconstructed image quality is better with a larger number of k. It also depends on the features of anatomical images. As indicated in the numerical simulations with cylindrical geometry and the phantom experiment with homogeneous background, for anatomical images with fewer features, we can obtain good reconstruction results considering only 64 nearest neighbors in the kernel matrix. However, for the anatomical images with rich features, such as elliptic cylindrical simulation, the simulation with MRI data and the phantom experiment with inhomogeneous background, we have to consider more nearest neighbors in the kernel matrix. We also found that for the simulations and experiments with the kernel method of k=64, we can obtain better reconstruction results with k=256 at a price of longer time to form the kernel matrix. The third parameter is the voxel number. As demonstrated in the first and second simulations, for the anatomical images without any false information, the quality of reconstructed FMT images increases slightly as the voxel number increases for a fixed number of nearest neighbors k. However, for the numerical simulations with the false target size in anatomical images, the kernel method with the smaller number of voxels performed better than the kernel method with the larger number of voxels. As shown in Table 3, Dice and CNR are achieved highest with the voxel number of $3 \times 3 \times 3$ for the reconstruction with the kernel method when k=256. MSE also reached the lowest for the case of voxel number $3 \times 3 \times 3$. VR coefficient is not informative in this case because the incorrect bigger size of the target in the anatomical images introduce higher volume ratio to the reconstruction image. Similarly, the numerical simulation with MR images and the phantom experiment with artificial features further demonstrate this trend by obtaining the best results with of voxel number of $3 \times 3 \times 3$.

The kernel matrix was generated before the FMT reconstruction with the matrix generation time depending on the voxel size and the nearest neighbor number. Table 5 lists the K matrix

generation time for the elliptic cylindrical simulation with two targets. For the best image quality setup with the nearest neighbor of 256 and the voxel number of $3 \times 3 \times 3$, the *K* matrix generation time was 20.09 seconds on a cluster with 12 nodes (2.8 GHz each node) and 128 GB memory. This is slightly longer than the time spends on generating the regularization matrix for the soft prior method, which was 12.28 seconds in this simulation with 5 regions. We acknowledge that, the time spent on the generating the soft prior matrix refers to the time for generating the matrix *L* from the region labeled vector. Since kernel matrices are sparse, multiplications involved in reconstruction processes also do not introduce significant computation time. In this study, the kernel method based FMT reconstruction converged in no more than 10 iterations which were around 5 seconds in total.

Table 5 Time to generate the kernel matrix K with different k and voxel sizes (in seconds)

Voxel size	<i>k</i> =16	<i>k</i> =32	k=64	k=128	<i>k</i> =256
3×3×3	3.36	3.78	5.46	8.88	20.09
5×5×5	11.32	12.78	16.81	23.58	38.79
7×7×7	28.89	34.59	43.05	57.30	96.42

The gray-scale values in the anatomical guidance images are included in the kernel matrix so that these values affect the kernel method. As shown in the rat brain case, when the target has distinct contrast to other background regions, the kernel method performed very well. And we have also found that the kernel method is robust to the inhomogeneous inclusions in the anatomical guidance images when these inclusions have lower gray-scale values than the values in the target region. However, it is fine to have inclusions with larger gray-scales values when the inclusions are not close to the target as shown in Fig. 10a.

In summary, we have introduced a kernel method based FMT reconstruction algorithm as a new approach to include the anatomical guidance. Numerical simulations prove that this method is robust in overcoming incorrect anatomical guidance. Phantom experiments further validate that the proposed method can improve the FMT reconstruction quality and does not increase the reconstruction time. In the future, we will apply the proposed kernel method to *in vivo* experiments on the hybrid systems.

Disclosures

No conflicts of interest, financial or otherwise, are declared by the authors.

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Biographies and photographs of the authors are not available.

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Fig. 1 Numerical simulation phantom geometry of (a) the cylindrical phantom with target locations at T1 (-1.7, 5.56) and T2 (1.7, 5.56) and (b) the elliptic cylindrical phantom with target locations at T1 (-1.2, -5.0) and T2 (1.2, -5.0).

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