Rigorous Uncertainty Estimation for MRI Reconstruction

Ke Wang^{1,2}, Anastasios Angelopoulos¹, Alfredo De Goyeneche¹, Amit Kohli¹, Efrat Shimron¹, Stella Yu^{1,2}, Jitendra Malik¹, and Michael Lustig¹ ¹Electrical Engineering and Computer Sciences, University of California, Berkeley, Berkeley, CA, United States, ²International Computer Science Institute, University of California, Berkeley, Berkeley, CA, United States

Synopsis

Deep-learning (DL)-based MRI reconstructions have shown great potential to reduce scan time while maintaining diagnostic image quality. However, their adoption has been plagued with fears that the models will hallucinate or eliminate important anatomical features. To address this issue, we develop a framework to identify when and where a reconstruction model is producing potentially misleading results. Specifically, our framework produces confidence intervals at each pixel of a reconstruction image such that 95% of these intervals contain the true pixel value with high probability. In-vivo 2D knee and brain reconstruction results demonstrate the effectiveness of our proposed uncertainty estimation framework.

Introduction

Deep learning (DL)-based reconstruction methods have shown great potential for efficient image reconstruction from undersampled k-space measurements¹⁻⁵. However, a substantial risk in DL-based reconstruction is hallucination or elimination of important anatomical features from the image⁶. To address this concern, we seek uncertainty estimates which tell us when we can trust a reconstruction. While existing⁷⁻¹¹ methods have shown promising results in estimating uncertainty maps, they do not come with statistical guarantees. Furthermore, they often require computational overhead (such as running multiple reconstructions) and/or modifications to the reconstruction network (e.g. nonstandard training procedures) that may sabotage its accuracy. In this work, we propose a simple and rigorous uncertainty estimation framework that works without modifying or retraining the reconstruction network (Figure 1). Our technique provides a rigorous finitesample statistical guarantee. Our key contribution is the development of a new form of Risk-Controlling Prediction Set (RCPS)¹² tailored to MRI reconstruction that outputs image-valued confidence intervals containing at least $(1 - \gamma)$ (e.g., 95%) of the ground truth pixel values. Our in-vivo knee and brain results probe the quality of our uncertainty estimation model, which allows us to identify specific regions where the model performs poorly.

Methods

Our method trains an uncertainty estimation network, then calibrates that network to achieve a rigorous guarantee. We will now detail these two subroutines.

1. Training the uncertainty estimation network

Given a pre-trained reconstruction network, e.g., MoDL⁴, the uncertainty estimation network predicts the absolute residual error for that network (Figure 2a). The pre-trained network G_w takes the zero-filled reconstruction and maps it to $\hat{x_i}$, an estimate of the ground truth image x_i . Our uncertainty estimation network f_{θ} is trained to output an estimate \hat{err}_i of the magnitude of the residual error $|\mathbf{x}_i - \mathbf{x}_i|$. In practice, the input to f_{θ} is actually several concatenated features from each iteration of G_w . After training is complete, we can now map new, unseen under-sampled inputs to the reconstructed images and uncertainty estimates in one forward pass. However, note that we have no guarantee that $e_{rr_i}^{\wedge}$ does well at estimating the pixel-wise error, so we will need to calibrate it.

2. Calibration of the heuristic uncertainty estimates

Once the uncertainty estimation network is trained, we aim to calibrate its output using Risk-controlling Prediction Sets¹² (Figure 2b) to achieve a statistical guarantee. We first select a subset of the validation set to form the calibration set $(x_i, x_i, err_i), i = 1, 2, 3, 4, ..., N$ (typically $N \ge 1000$). Then, we calibrate a global scalar α from the calibration set to ensure that, on average, at least $(1 - \gamma)$ of all pixels from the reference are within its confidence intervals

 $I_i^{(m,n)} = [x_i^{(m,n)} - \alpha \cdot \hat{\operatorname{err}}_i^{(m,n)}, x_i^{(m,n)} + \alpha \cdot \hat{\operatorname{err}}_i^{(m,n)}], \text{ for all pixel locations } (m,n) \text{ in an image of size } M \times N. \text{ For example, choosing } \gamma = 0.05 \text{ and } \delta = 0.1 \text{ will result}$ in 95% of the pixels being contained in their intervals with 90% probability. The detailed calibration procedure is described as follows. For a given image x_i, we first define the loss

$$L(\alpha)_i = \frac{|(m,n): \mathbf{x}_i^{(m,n)} \notin I_i^{(m,n)}|}{MN}$$

as the fraction of pixels not included in their respective intervals. We compute the empirical risk over the calibration dataset and use the Upper Confidence Bound (UCB)^{15,16} procedure from¹² with the WSR bound from¹⁵ to choose the smallest α that gives a RCPS,

$$\mathbb{P}[R^{+}(\alpha) \ge R(\alpha)] \ge (1-\delta),$$

where δ here is the desired violation rate (e.g., δ =0.1). In short, the method involves computing the UCB $R^{+}(\alpha)$ using a pointwise concentration inequality, then picking $\alpha = \min \left\{ \alpha : R^{+}(\alpha') < \gamma, \forall \alpha' > \alpha \right\}$. Deploying this choice of α guarantees risk-control; we defer the proof of this fact to¹².

Datasets and experimental setups

We evaluated the proposed framework on both 2D knee and brain fastMRI¹³ datasets. First, MoDL was trained for both anatomies using 5120 different slices. Then, we trained the uncertainty estimation network using the same training set, with a range of acceleration factors. Finally, we calibrated the heuristic uncertainty estimates using a calibration set of 1000 slices, while the validation set contained 2000 slices. We compared the heuristic uncertainty estimation results with the absolute residual errors. To evaluate the calibration procedure, we randomly split the validation set 2000 times. Each time, we calibrated a α_i , j = 1, 2, 3, ..., 2000and evaluated the empirical risk $\hat{R_i}$ on the rest of the validation set (evaluation set). We presented the histogram of the empirical risks to evaluate the empirical violation rate $\hat{\delta}$.

Results

Figure 3 shows the uncertainty estimation results for the knee and brain datasets. The results show strong agreement between the uncertainty estimates and the blurred residual error.

Figure 4 visualizes the textures and the corresponding uncertainty estimates. Zoomed-in details indicate that higher uncertainty appears in the regions where the reconstructed images did not successfully recover the fine textures and details.

Figure 5 shows the empirical risk distribution given different splits of calibration/evaluation sets. Histograms show that the empirical violation rate δ hits nearly exactly δ for both γ , which demonstrates the tightness and validity of our calibration procedure.

Conclusions

This work presented a rigorous uncertainty estimation framework, which can provide precise uncertainty estimates backed by a finite-sample guarantee. Without any constraints on the reconstruction model, our framework acts as a plug-and-play module, and may significantly improve the accuracy of the diagnosis and clinical interpretation of DL-based reconstructions.

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Figures



Figure 1. Overview of the proposed model-specific rigorous uncertainty estimation framework for general DL-based reconstruction models. After training $f_{ heta}$, our networks output the heuristic uncertainty estimates alongside the reconstructed image in one forward pass. By developing a new form of Risk-Controlling Prediction Set to calibrate the uncertainty estimates, our calibrated uncertainty estimates provide guaranteed confidence intervals that contain at least $(1 - \gamma)$ (e.g., 95%) of the ground truth pixel values.



Figure 2. Detailed subroutines for the proposed framework. a) we first train an uncertainty estimation network f_{θ} to predict the pixel-wise residual of a pre-trained reconstruction model G_w , where we name the output as heuristic uncertainty estimates. b) After training, we calibrate the uncertainty estimates to form finitesample confidence intervals, which ensures that on average, $(1-\gamma)$ of pixels are covered within the confidence interval with high probability regardless of the distribution of the training data.



Figure 3. Representative uncertainty estimation comparisons of the uncertainty estimates and the absolute residual error for both knee (Sequence: Proton density) and brain (Sequence: T1w post-contrast) reconstructions. Sampling masks are four times variable density masks and five times random masks for each dataset, respectively. We overlaid the MoDL reconstructed images and the calibrated uncertainty estimates for better visualization. Colorbar along with the overlaid image indicates the guaranteed confidence interval with respect to the maximum value of the image.



Figure 4. Visualization of textures and the corresponding uncertainty estimates from two representative images. As can be seen in the zoomed-in details, the reconstructions of the green-outlined patches are highly similar to the ground truth ones, while those of the yellow-outlined patches are of lower quality, since some of the high-frequency details are missing or blurred out. This is reflected by the overlaid calibrated uncertainty estimates, where the yellow-outlined patches have much higher uncertainty levels than the green ones.



Figure 5. Empirical risk distribution under 2000 random split of calibration/evaluation sets for brain and knee datasets. Each split of the calibration set outputs an lphaand the corresponding empirical risk \hat{R} , which roughly describes the number of pixels violating the desired risk/confidence level. Given a desired violation rate, the empirical violation rate $\hat{\delta}$ indicates how frequently the desired risk/confidence levels are violated. Comparisons of two desired risk/confidence levels $\gamma = 0.1, 0.05$ are presented.

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