

Semantic Segmentation with Incomplete Annotations

DeepVision Workshop



UNIVERSITY OF ICELAND

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IRCAD Strasbourg, Visible Patient
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Outline

1 Context

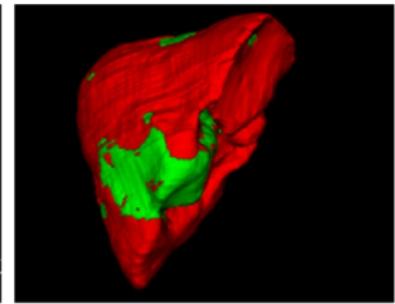
2 Semantic Segmentation with Incomplete Annotations

3 Experiments

4 Ongoing Works and Perspectives

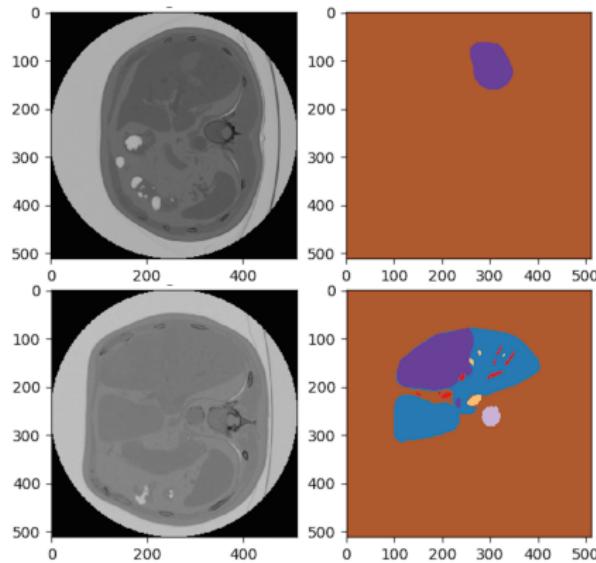
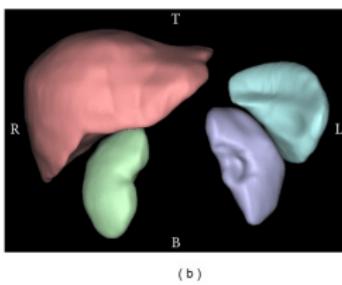
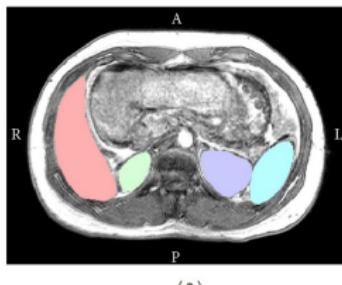
Context: Semantic Segmentation of Medical Images

- ▶ Semantic Segmentation: class label for each image pixel / voxel
- ▶ Deep ConvNets: tremendous success for visual recognition
- ▶ Semantic Segmentation of natural images: Fully Convolutional Networks (FCN), e.g. DeepLab [Chen et al., 2018]
 - ▶ Adapted FCN architectures for medical images, e.g. U-Net [Ronneberger et al., 2015]
 - ▶ FCN: base architecture for leading approaches in recent medical segmentation challenges, e.g. LITS'17 [Han, 2017, Li et al., 2017]



Datasets for Medical Image Semantic Segmentation

- ▶ ConvNets: large amount of data with clean annotations
- ▶ Annotation very costly for semantic segmentation: pixel-level labeling
 - ▶ Exacerbated in medical images: 3D data, highly qualified professionals needed, e.g. tumors (extreme appearance variations)

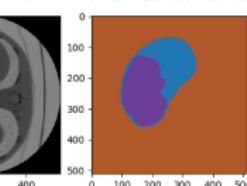
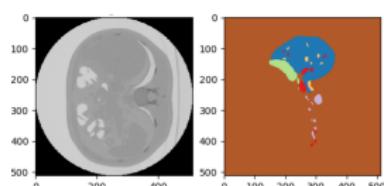
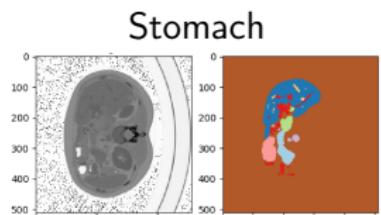
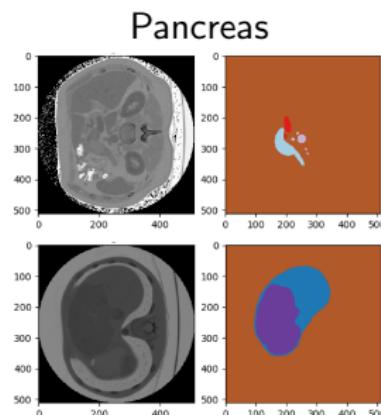
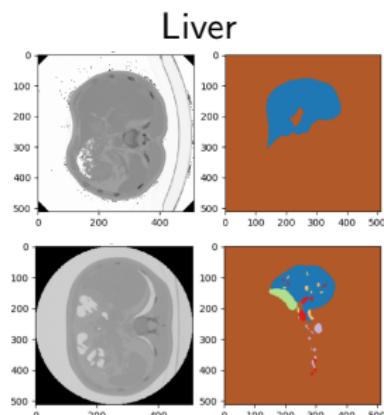


Semantic Segmentation of 3D CT-scans

- Internal dataset¹: ~ 1000 patients of $100 \times 512 \times 512$ images



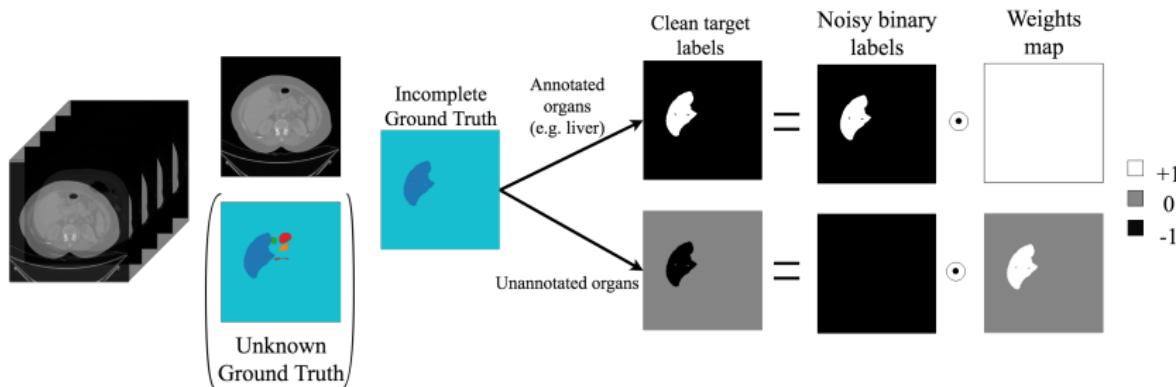
- 3D segmentation: focusing on 2D slices
⇒ independent training in each image



¹IRCAD: <https://www.ircad.fr/fr/>

Semantic Segmentation with Incomplete Annotations

- ▶ Large scale dataset, BUT:
 - ▶ Clinical experts: focus on a subset of organs
⇒ **Incomplete annotations** wrt full Ground Truth



- ▶ **How to train deep ConvNets in this context ?**
 - ▶ Organ(s) missing the whole volumes, but: organ segmented in volume ⇒ complete annotation for that class
 - ▶ **Core idea:** generating clean target labels from noisy input labels
 - ▶ Binary mask w_k for each class ⇒ ambiguous vs non-ambiguous pixels

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1 Context

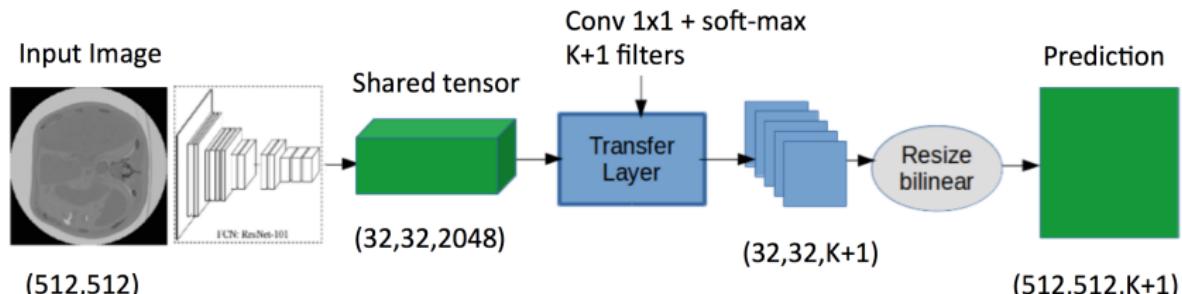
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Semantic Segmentation with Incomplete Annotations

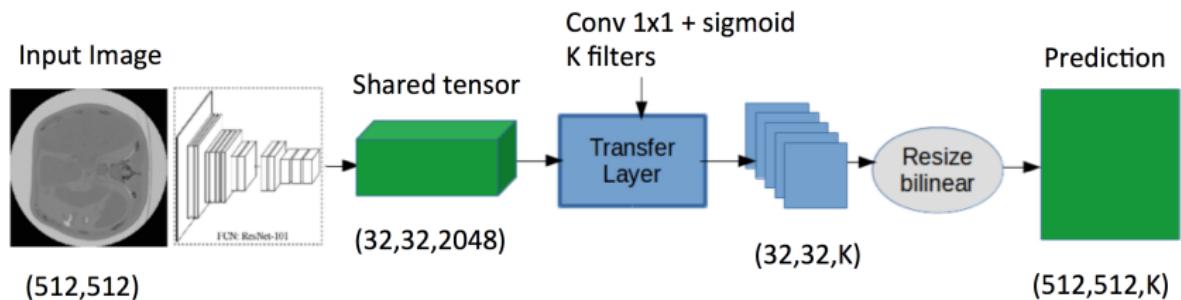
- Standard FCN not adapted to this context, e.g. DeepLab [Chen et al., 2018]



- Shared Fully Convolutional Layers, ResNet [He et al., 2016]
- Last tensor: 1×1 conv + soft-max \Rightarrow single class prediction
- Incomplete annotation: "background" \Leftrightarrow missing organ**
 \Rightarrow conflict with pixels with proper organ annotations during training

Semantic Segmentation with Incomplete Annotations

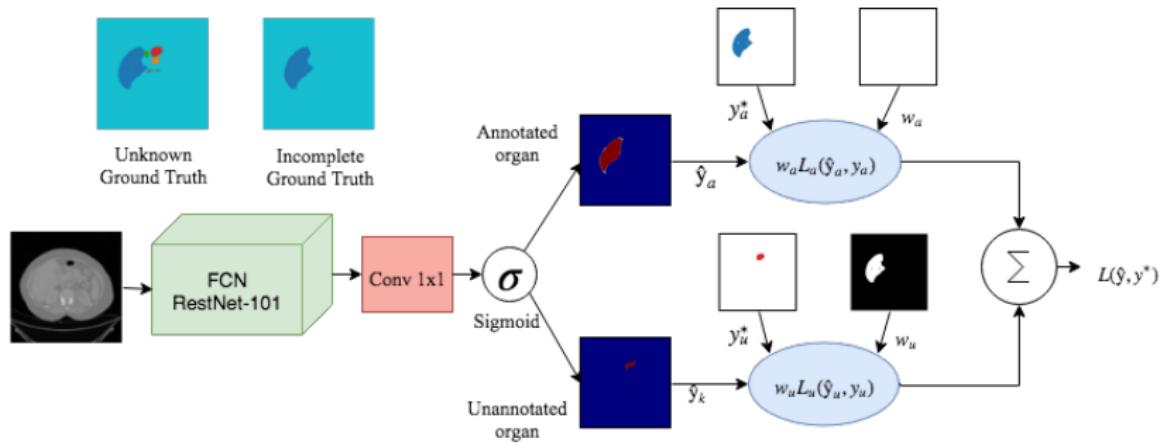
- Our approach for **Semantic segmentation with MIsing Labels and convnEts (SMILE)**
- Depart from the $(K + 1)$ multi-class classification formulation, classify each organ independently using K binary classifiers



SMILE Training

- Binary CE loss at each pixel: $L_k(\hat{y}_k, y_k^*) = - (y_k^* \log(\hat{y}_k) + (1 - y_k^*) \log(1 - \hat{y}_k))$
- Final loss: weighted sum of binary losses:

$$L(\hat{y}, y^*) = \sum_{k=1}^K w_k L_k(\hat{y}_k, y_k^*)$$

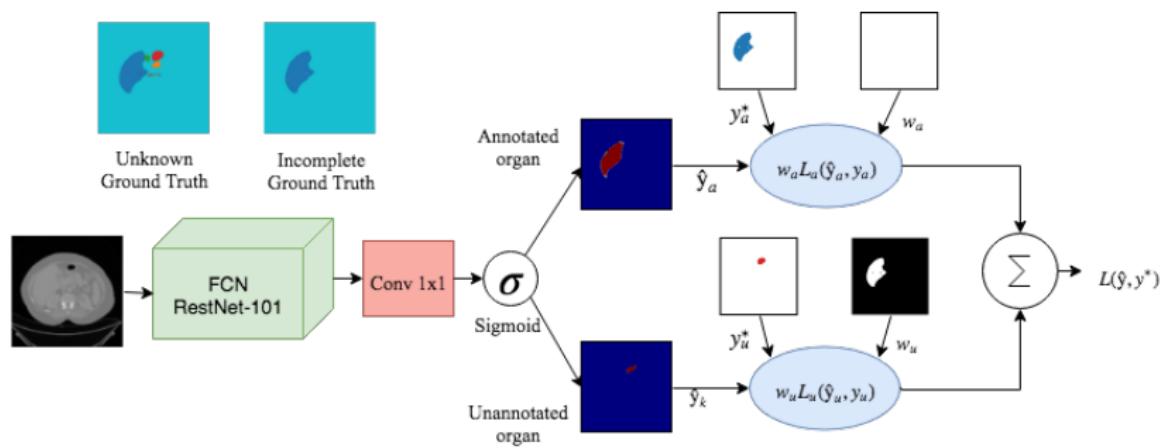


SMILE Training

Core SMILE component: binary weight maps $w_k \in \{0; 1\}$

- >Selecting or ignoring each pixel for class k
 - Class k present in volume: $w_k = 1 \forall$ pixel in volume
 - Class k absent:

$$w_k = \begin{cases} 1 & \text{if } \exists k' \neq k \text{ s.t. } w_{k'} = 1 \ (\Rightarrow y_k^* = -1), \\ 0 & \text{otherwise (pixel ignored)} \end{cases}$$



SMILE Training

- ▶ Analysis of labels used by FCN baseline and SMILE vs Ground Truth (GT)
- ▶ For class k :
 - ▶ β_k ratio of voxels in a volume
 - ▶ α the ratio of missing labels for this organ in the dataset.

Baseline FCN		SMILE	
Used	GT	Used	GT
Pos	Pos	Pos	Pos
$(1 - \alpha) \cdot \beta_k$	$\alpha \cdot \beta_k$	$(1 - \alpha) \cdot \beta_k$	0
0	$1 - \beta_k$	0	$(1 - \alpha) \cdot (1 - \beta_k) + \epsilon$

$$\epsilon = \sum_{k' \neq k} \beta_{k'}$$

- ▶ Both baseline and SMILE: only true positive
 - ▶ BUT only use $(1 - \alpha) \cdot \beta_k$ vs β_k

SMILE Training

Baseline FCN		SMILE	
GT	Used	Pos	Neg
Pos		$(1 - \alpha) \cdot \beta_k$	$\alpha \cdot \beta_k$
Neg		0	$1 - \beta_k$

GT	Used	Pos	Neg
Pos		$(1 - \alpha) \cdot \beta_k$	0
Neg		0	$(1 - \alpha) \cdot (1 - \beta_k) + \epsilon$

$$\epsilon = \sum_{k' \neq k} \beta_{k'}$$

▶ Baseline:

- ▶ **False Negatives (FN):** $\alpha \cdot \beta_k$, i.e. unannotated pixels indeed belonging to the organ

$$\boxed{\frac{TP}{FN} = \frac{1-\alpha}{\alpha}: \alpha > 0.5 \Rightarrow \frac{TP}{FN} < 1}$$

▶ SMILE:

- ▶ **Only true positives and true negatives**
- ▶ **Less true negatives than baseline:** $(1 - \alpha) \cdot (1 - \beta_k) + \epsilon$ vs $(1 - \beta_k)$
 - ▶ $\approx \alpha$ less negatives, but as $\beta \ll 1$, e.g. $\beta = 0.05^2$
 - ⇒ **in practice, largely enough negative to train**

²organs ⇔ small volume portion

Incremental self-supervision and relabeling

- ▶ SMILE True Positives (TP) labels $\propto (1 - \alpha)$
- ▶ **Motivation:** automatically increasing number of TP labels
 - ▶ Compensate for incomplete annotations
- ▶ Auto-supervision: create target positive labels
⇒ **SMILEr** (re-labeling)
- ▶ Using a curriculum strategy [Bengio et al., 2009]
 1. Train ConvNet with SMILE: certain labels only, i.e. true positives and negatives ⇒ "easy samples"
 2. Seek for new true positives with current model
 - ▶ "Harder samples", automatic labeling
 - ▶ Use this new labels as target to train a new model with more positives
 - ▶ Iterate
- ▶ $\frac{TP}{FP}$: key indicator of SMILEr success

SMILER Training

- ▶ SMILER algorithm: applied for each binary organ classifier independently^a

Algorithm 1 Algorithm for training SMILER for class k

Require: Training set $\{(\mathbf{x}_i, \mathbf{y}_i^*)\}$, γ_{max} , T , SMILE model m_0 for class k .

```
1: Initialize  $\mathbf{y}_{i,0}^* = \mathbf{y}_i^*$ ,  $N_u \leftarrow$  number of unannotated images for class  $k$ 
2: for  $t=1$  to  $T$  do
3:    $\gamma_t = \frac{t}{T}\gamma_{max}$ 
4:   for  $i=1$  to  $N_u$  do
5:      $\hat{\mathbf{y}}_i^+ \leftarrow (m_t, \mathbf{x}_i)$  // Find predicted positive pixels by  $m_t$  in image  $\mathbf{x}_i$ 
6:      $\mathbf{y}_{i,t}^{*,+} \leftarrow (m_t, \mathbf{x}_i, \gamma_t, \hat{\mathbf{y}}_i^+)$  // Assign new  $\oplus$  target labels
7:      $\mathbf{y}_{i,t}^* = \mathbf{y}_{i,t-1}^* \cup \mathbf{y}_{i,t}^{*,+}$  // Augment training set
8:   end for
9:    $m_t = train(\{(\mathbf{x}_i, \mathbf{y}_{i,t}^*)\},)$  // Re-train model with augmented training set
10: end for
```

Ensure: SMILER Model m_T

^aIgnoring the dependence on class k for the sake of clarity.

Outline

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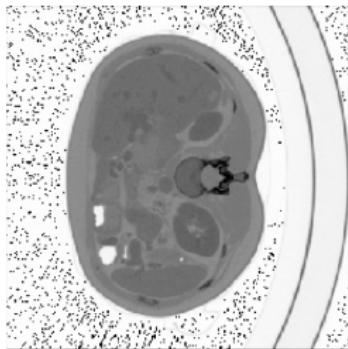
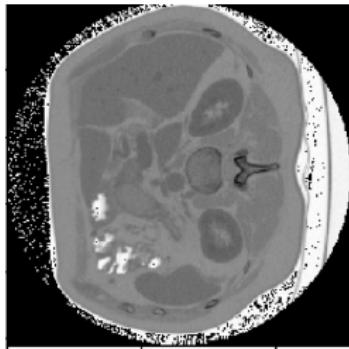
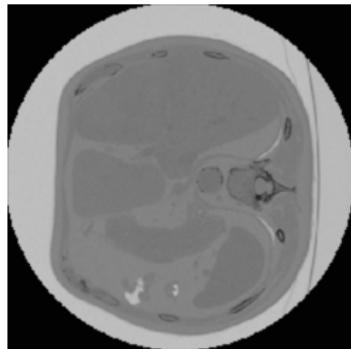
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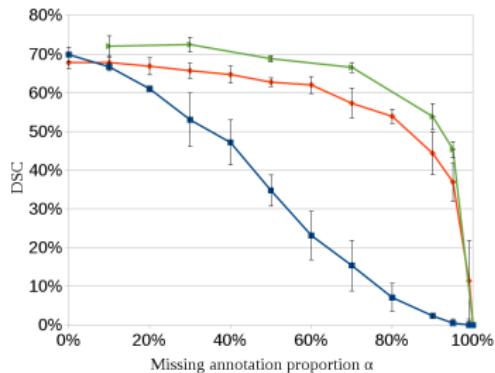
4 Ongoing Works and Perspectives

Dataset and setup

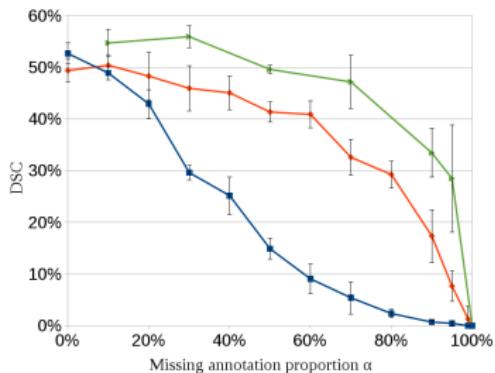
- ▶ Experiments on sub-set of our dataset with complete ground truth annotations
- ▶ 72 3D CT-scan volumes (~ 100 512×512 images) for three organs: liver, pancreas and stomach
- ▶ Partially annotated dataset generated: randomly removing $\alpha\%$ of organs in the volumes independently
- ▶ Comparison of our methods (SMILE, SMILER) wrt DeepLab baseline
 - ▶ Train 80% / Test (20%), $K = 5$ datasplits



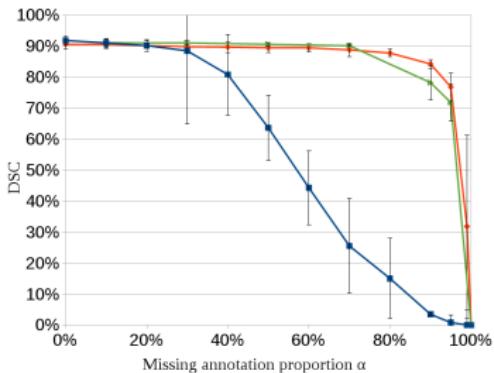
Quantitative results



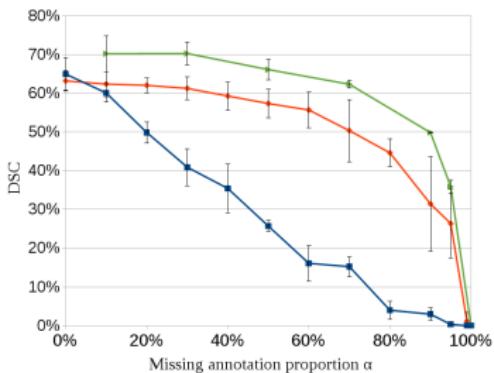
Mean



Pancreas

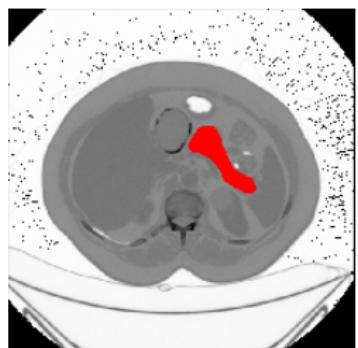
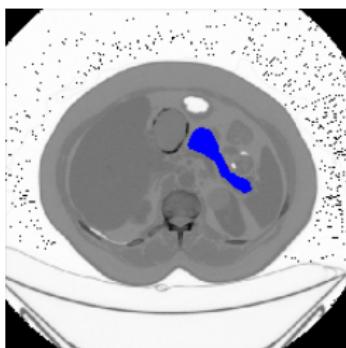
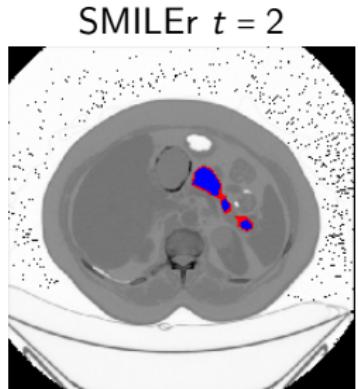
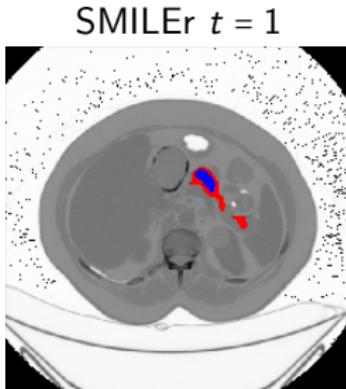
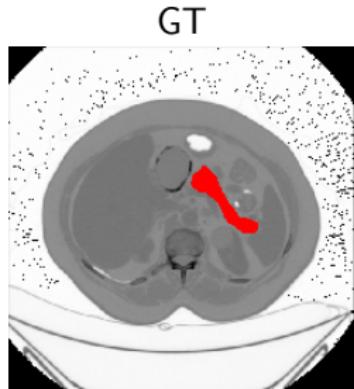


Liver

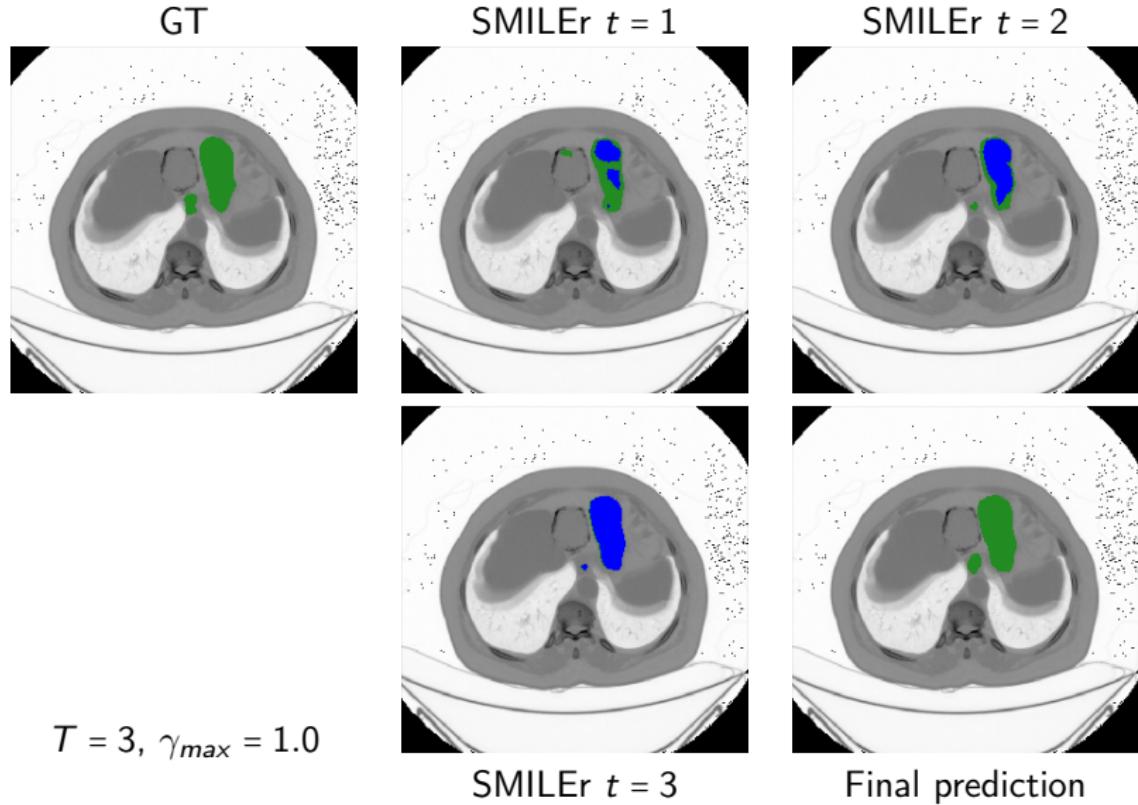


Stomach

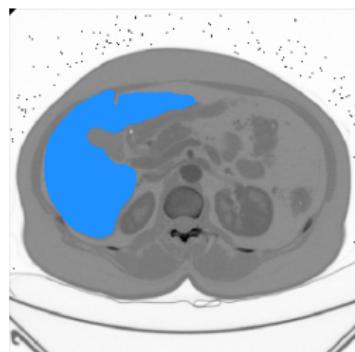
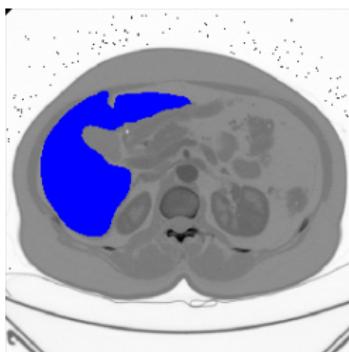
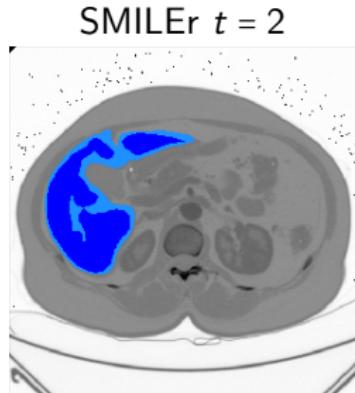
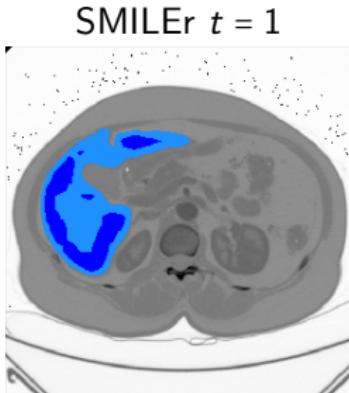
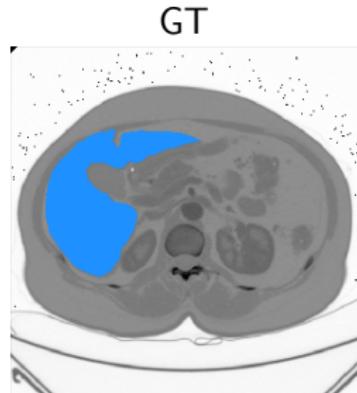
SMILEr re-labeling, $\alpha = 50\%$



SMILEr re-labeling, $\alpha = 70\%$

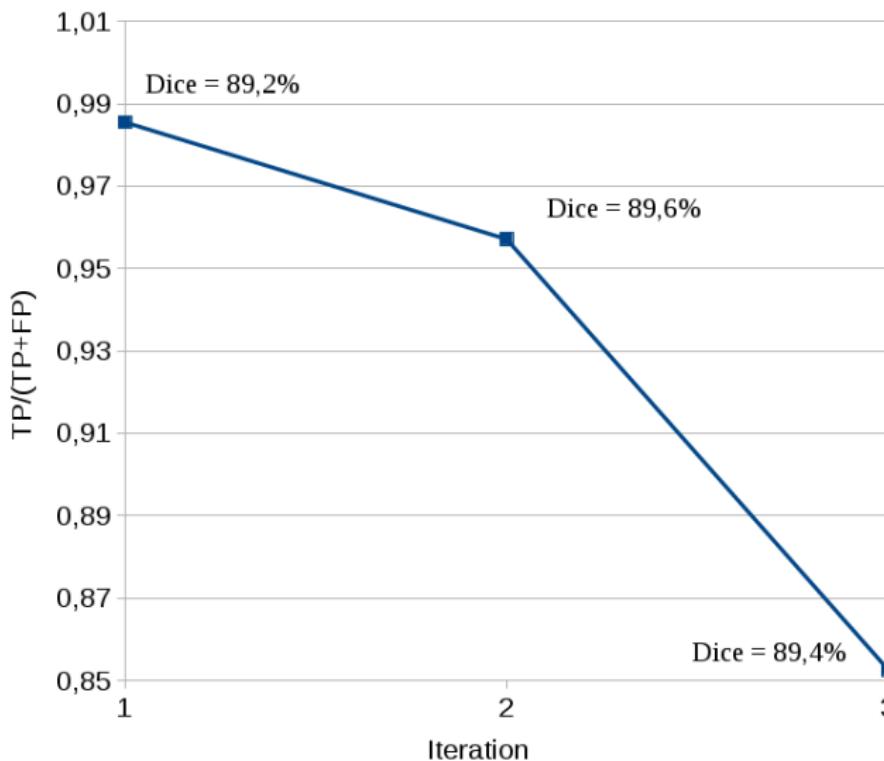


SMILEr re-labeling, $\alpha = 70\%$



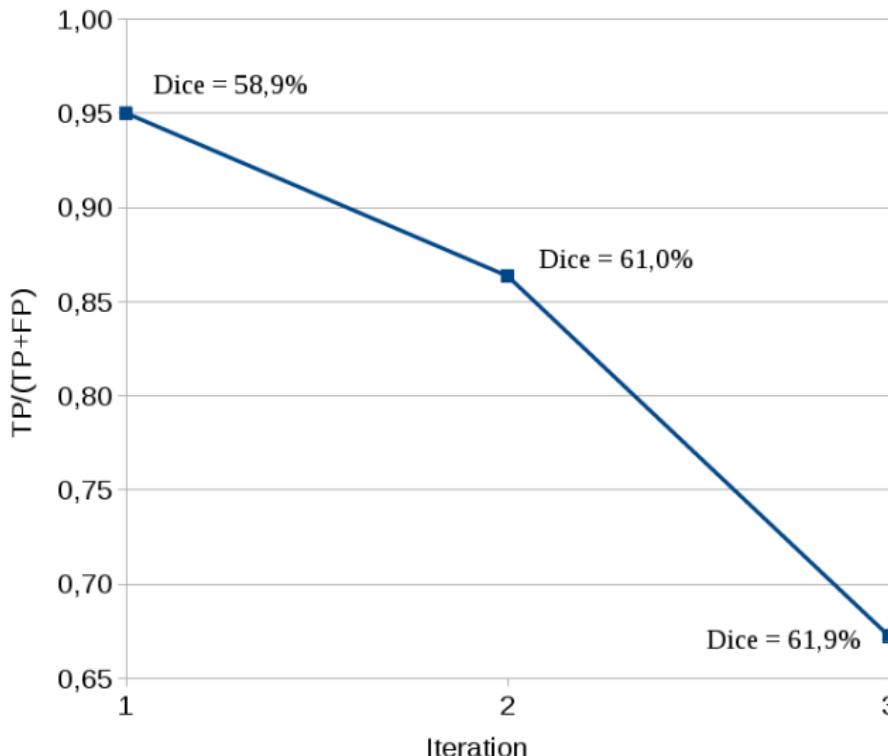
Re-labeling method

- ▶ $\frac{TP}{TP+FP}$ vs Curriculum iterations for Liver ($\alpha = 70\%$)



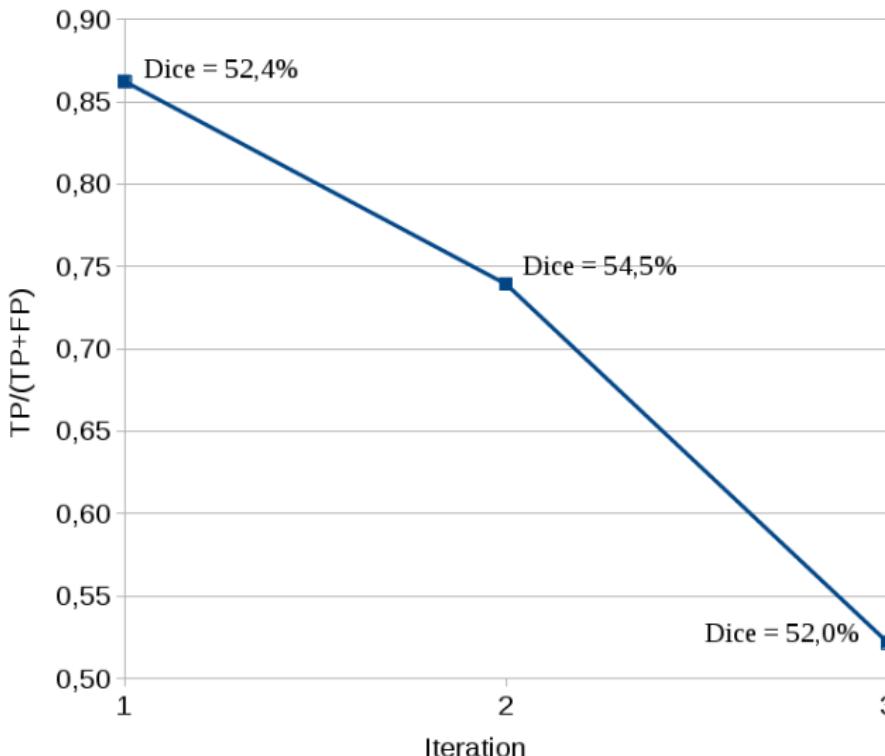
Re-labeling method

- ▶ $\frac{TP}{TP+FP}$ vs Curriculum iterations for Stomach ($\alpha = 70\%$)

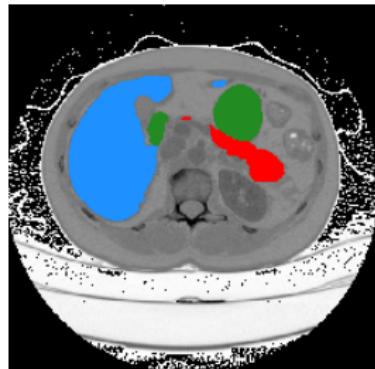


Re-labeling method

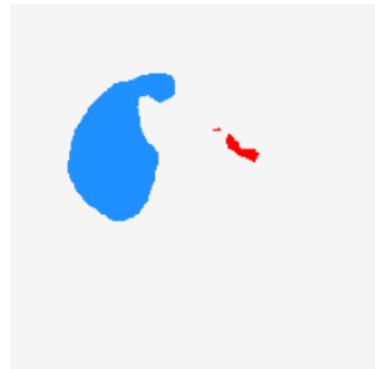
- ▶ $\frac{TP}{TP+FP}$ vs Curriculum iterations for Pancreas ($\alpha = 70\%$)



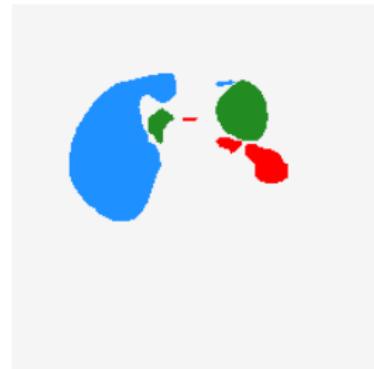
Segmentation results, $\alpha = 70\%$



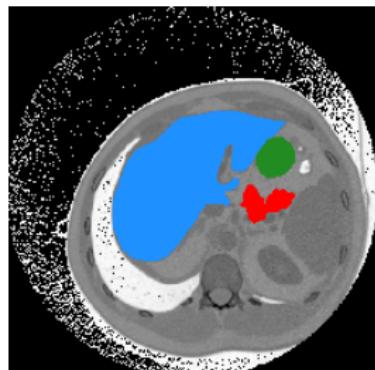
GT



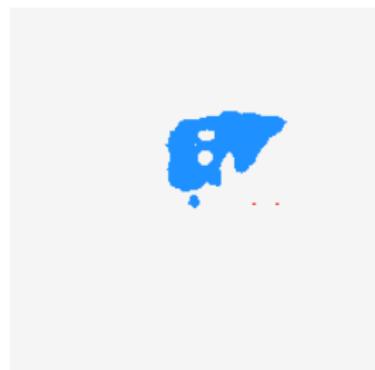
baseline



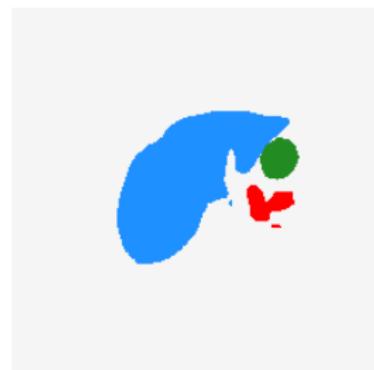
SMILEr



GT

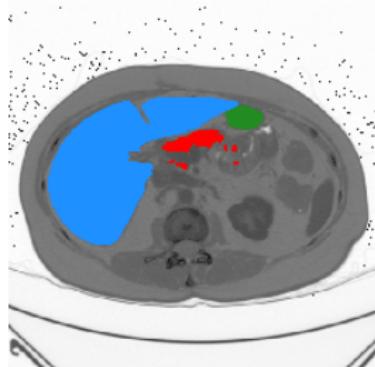


baseline

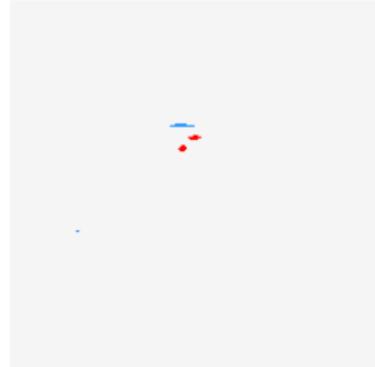


SMILEr

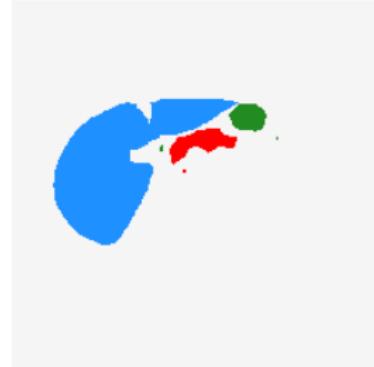
Segmentation results, $\alpha = 70\%$



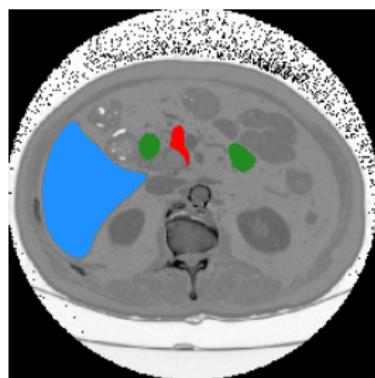
GT



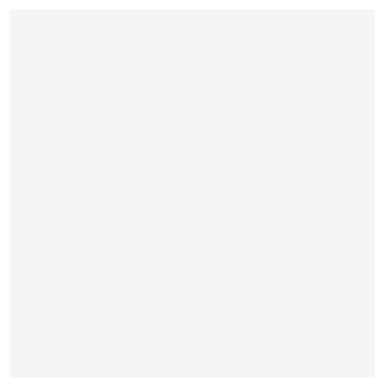
baseline



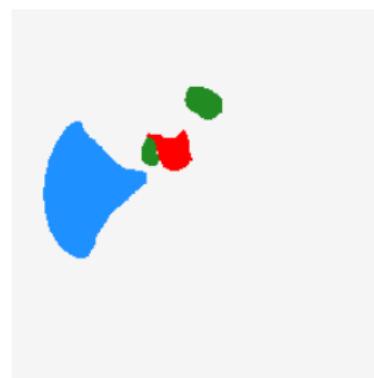
SMILEr



GT



baseline



SMILEr

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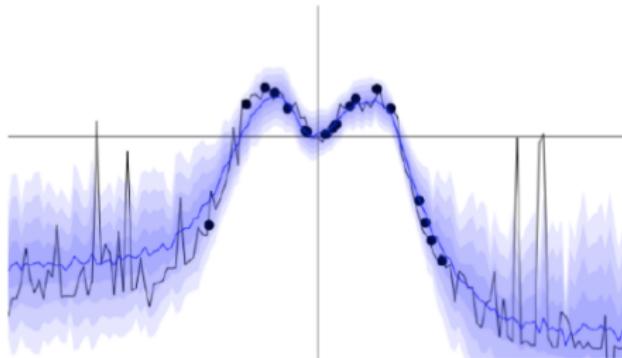
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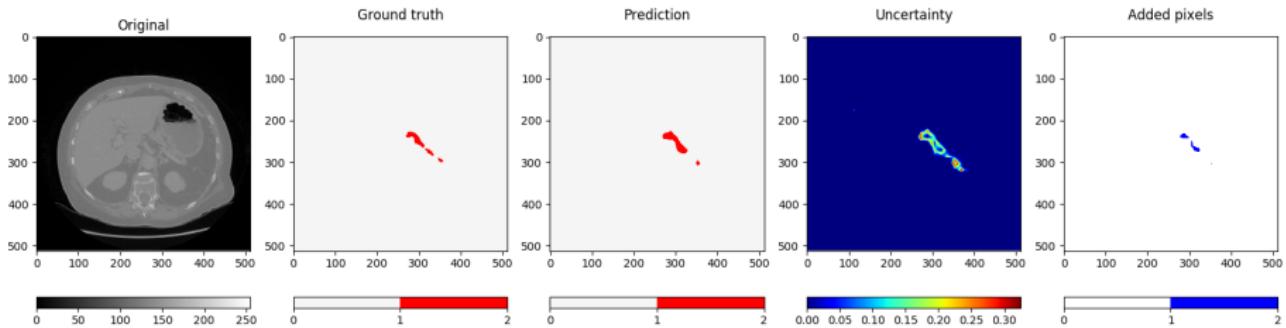
SMILER: re-labeling

- ▶ Target auto-supervision labels: top-scoring pixel
 - ▶ Last layer output in deep networks: not good confidence criterion
- ▶ Estimate uncertainty with Bayesian neural networks
 - ▶ Dropout as Bayesian approximation [Gal and Ghahramani, 2016, Kendall and Gal, 2017]
 - ▶ Simple practical implementation: variance of prediction with T dropout predictions

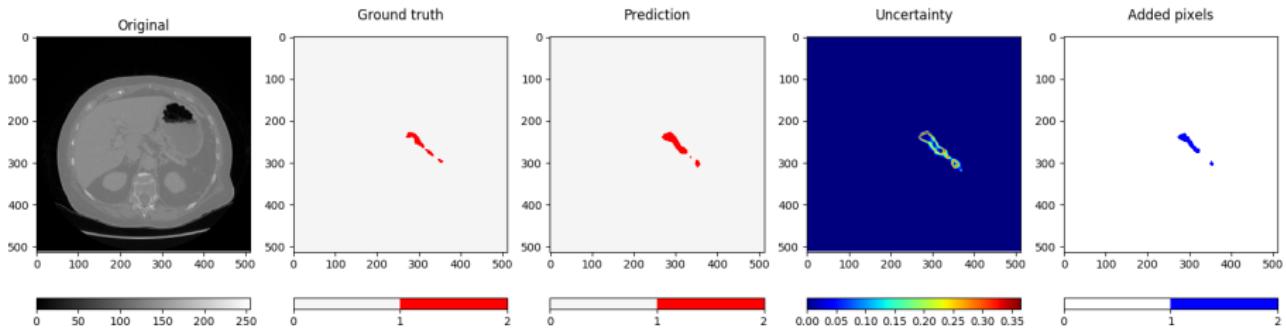


SMILER: Preliminary Results with Bayesian Dropout

$T = 1$: SMILER with lowest uncertainty (i.e. std) pixels (33%)

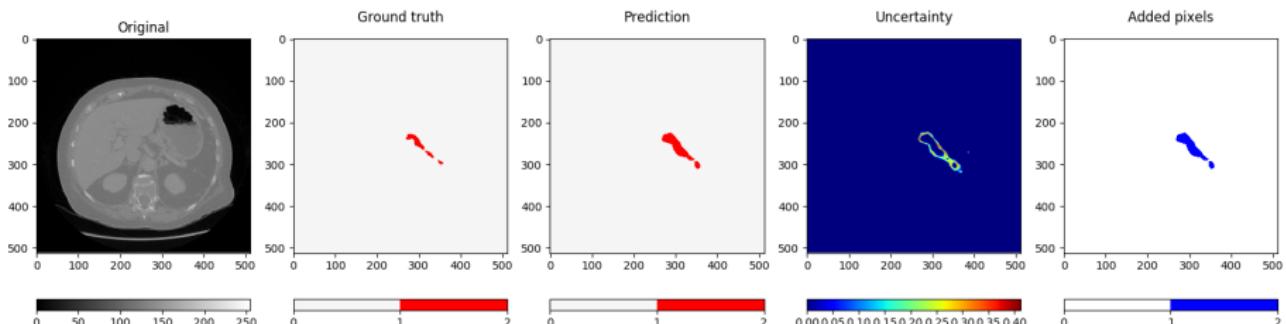


$T = 2$: SMILER with lowest uncertainty (i.e. std) pixels (66%)



SMILER: Preliminary Results with Bayesian Dropout

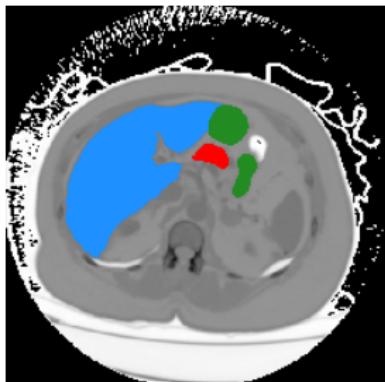
$T = 3$: SMILER with lowest uncertainty (*i.e.* std) pixels (100%)



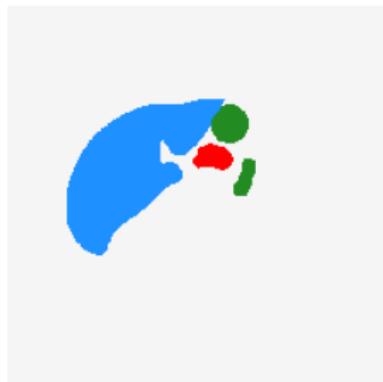
► To be continued...

Conclusion

- ▶ Method for learning with incomplete ground truth annotations
 - ▶ First stage: train only with correct label
 - ▶ Second stage: re-label positives
- ▶ Practical potential in large scale datasets with missing annotations, e.g. interactive re-labeling
- ▶ Future works (beyond uncertainty for target label selection):
 - ▶ Evaluation in larger datasets, Improving backbone architectures
 - ▶ Trained decoder, skip connections,
e.g. U-Net [Ronneberger et al., 2015], 3D ConvNets
 - ▶ Relation between medical structures, e.g. tumor (cascade)



GT



SMILEr

Thank you for your attention!

Questions?

Joint work with:

- ▶ Olivier Petit, PhD Student
- ▶ Luc Soler, Prof. at IRCAD, Visible Patient CEO

References |

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