

Localization of 3D Anatomical Structures Using Random Forests and Discrete Optimization

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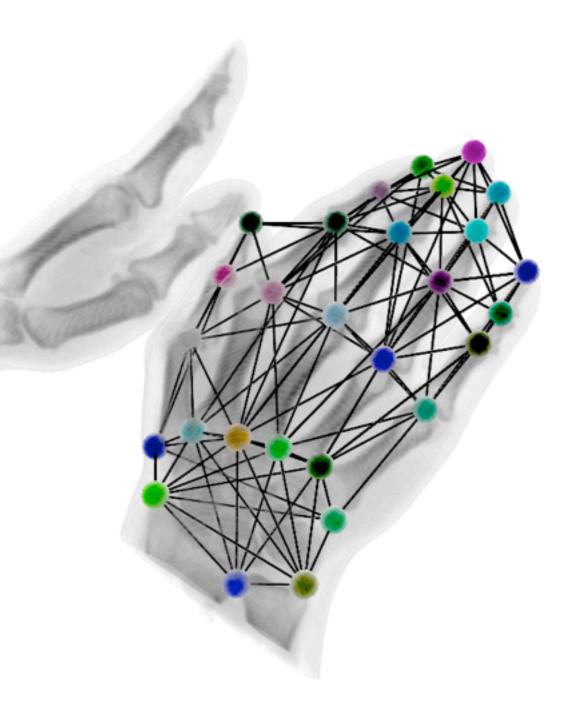


Motivation



- Most segmentation approaches require spatial initialization
 - ASM, AAM, GraphCuts, LevelSets

- Often application specific
- Localization of complex, self-similar structures

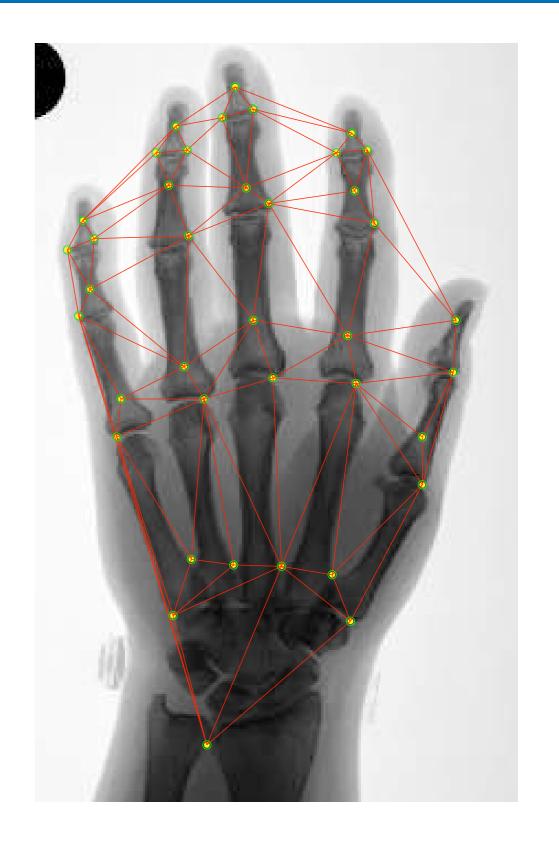


- Sparse MRF Appearance Models [Donner 2007,2010]
- Marginal Space Learning [Zheng 2009]
- Hierarchical Parsing [Seifert 2009]
- Random Forests [Criminisi 2009, Lempitsky 2009]
- A*-based optimization [Bergtholdt 2010]

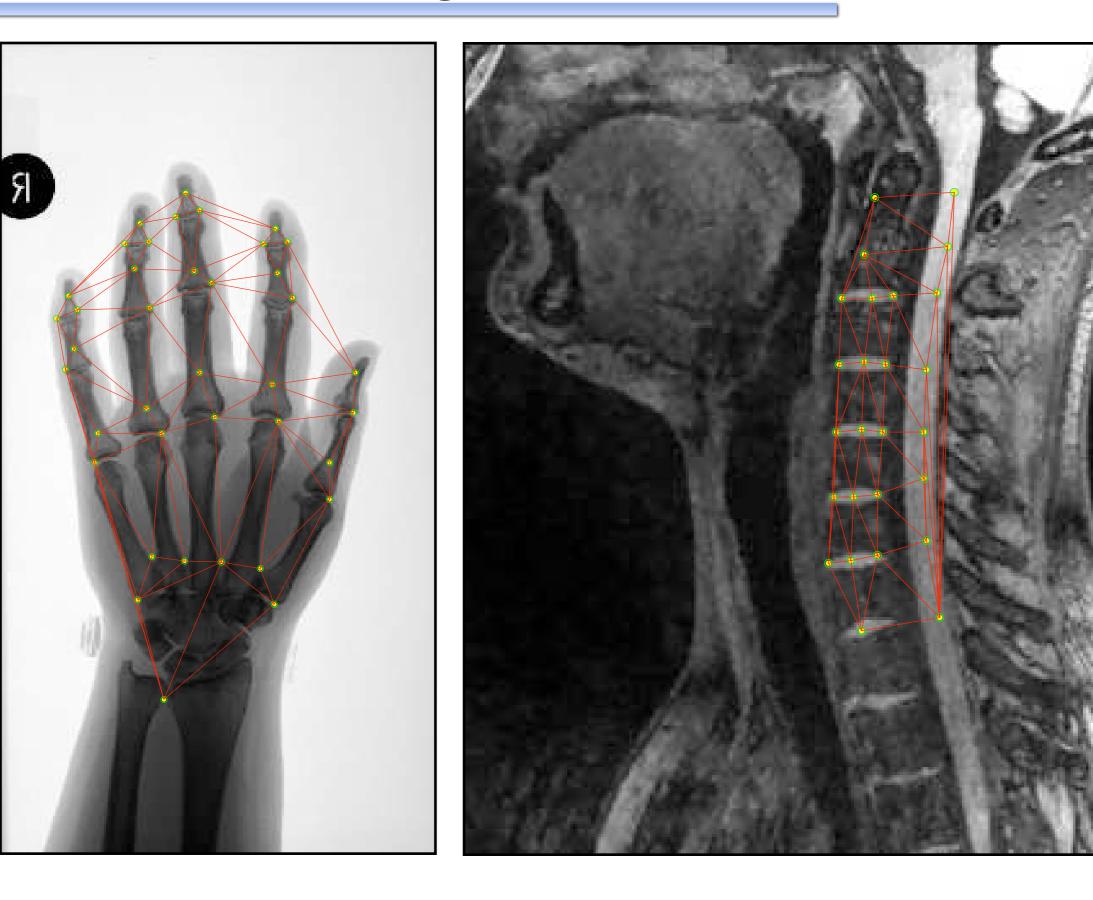


Sparse MRF Appearance Models

- Based on interest points
 - E.g. GVFpoints, Harris Corners
- Sparse Appearance Model
 - Elastic geometric model
 - Local descriptors around landmarks / edges
- Matching task formulated as MRF



Successful Matching





- Too many interest points
 - Corners or Symmetry to ambiguous
 - Multiple types of interest points?
- Local descriptors
 - How too choose?
 - Which distance metric?
- Size of MRF becomes intractable

Random Forest based Localization

- Decision Forests with Long-Range Spatial Context for Organ Localization in CT Volumes [Criminisi 2009]
 - Haar-like features with random perturbation
 - Annotation trough bounding box
 - Fast GPU implementation
 - Focused on entire organs





Finding candidate points through classification

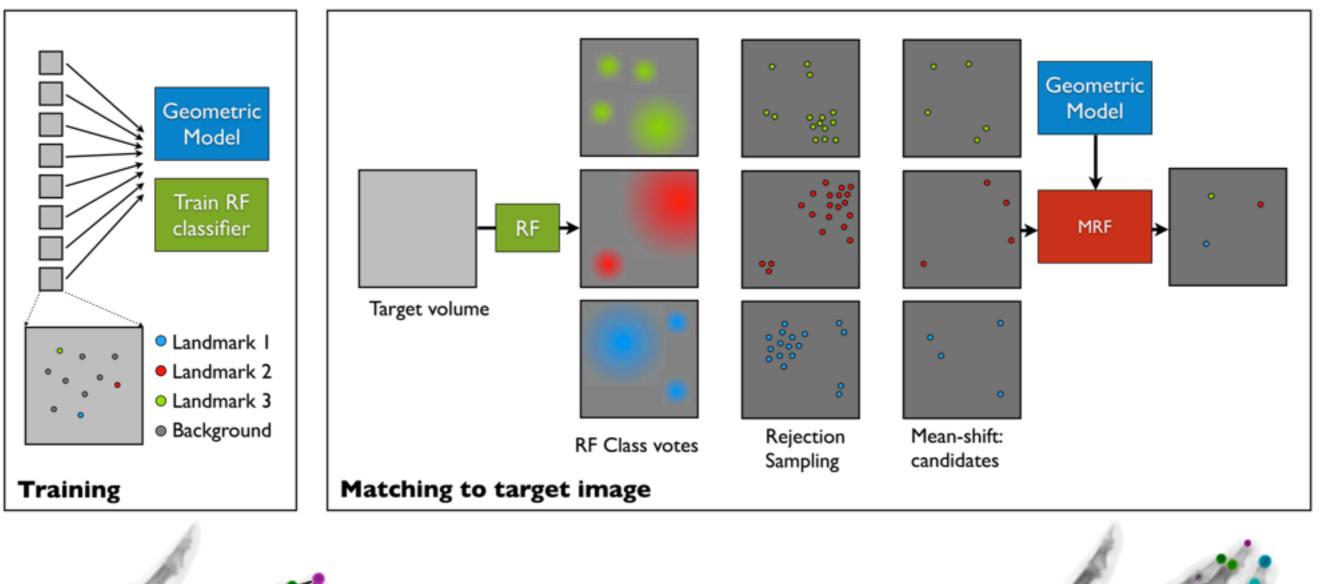
- Haar-like features
- Random forest classification
- Meanshift on sampled classification probabilities: cluster centers as candidate points

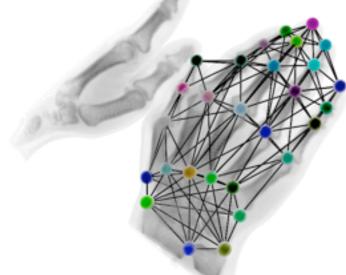
Probability estimation for MRF nodes/edges

- Local support of candidate point
- Edge probabilities from Gaussian distributions
- Simplified MRF graph through sparsity of label/edge probabilities

Flow Chart





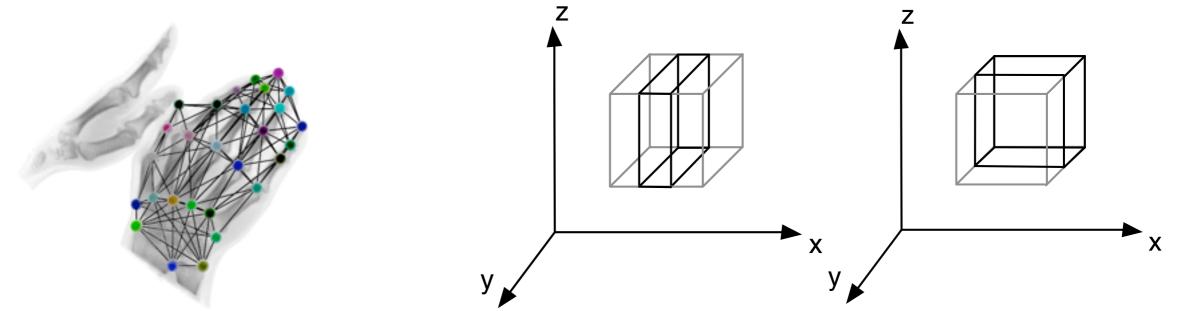


René Donner, CIR

Localization of 3D Anatomical Structures

Learning Candidate Point Detectors

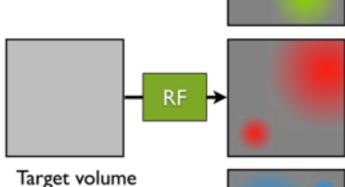
- Landmarks selected in each training sample
- Features computed for all voxels within a 3 voxel radius
- Haar-like features, using integral volumes
- 7 Features (average, 3 edge, 3 ridge) on
 3 scales (8, 16, 32 vx)



- One class per landmark
- + I background class, randomly sampled

Random Forest Classification

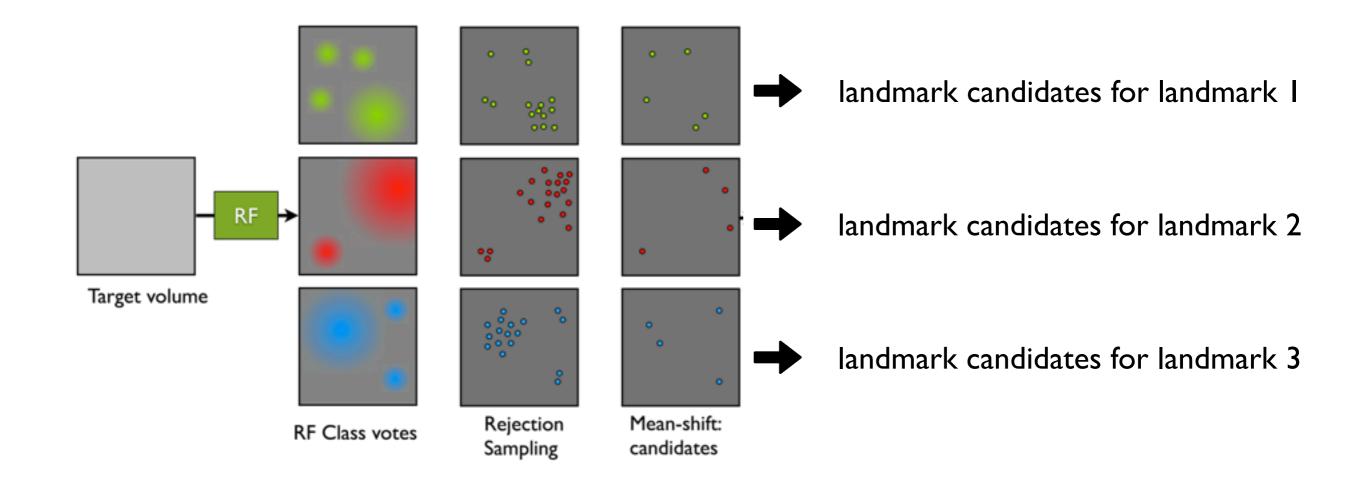
Vote count for each class = probability estimate for that landmark



RF Class votes

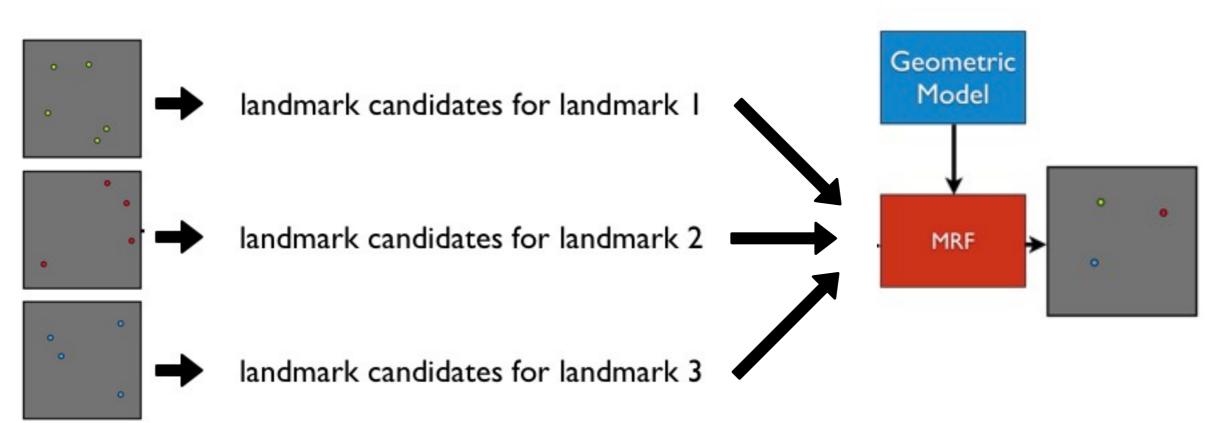
Sampling & Mean Shift

- Rejection sampling from normalized vote counts
- Sum ofvotes per cluster = local support of the candidate



Sampling & Mean Shift

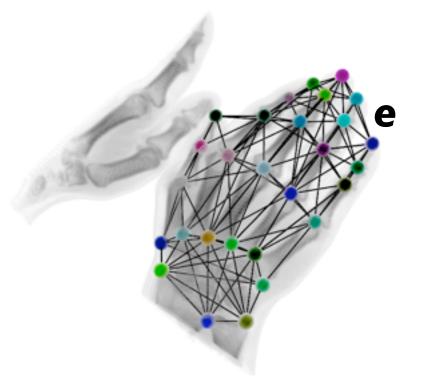
- Rejection sampling from normalized vote counts
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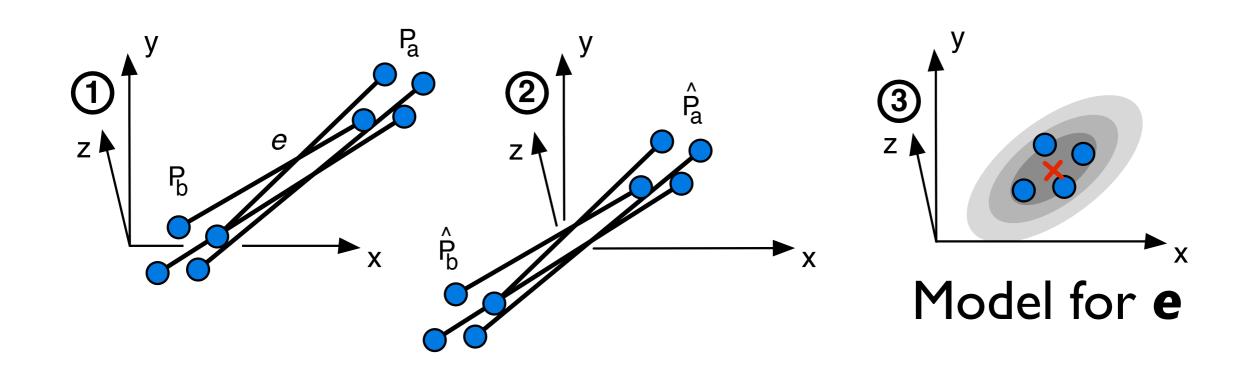


Mean-shift: candidates

Geometric Model

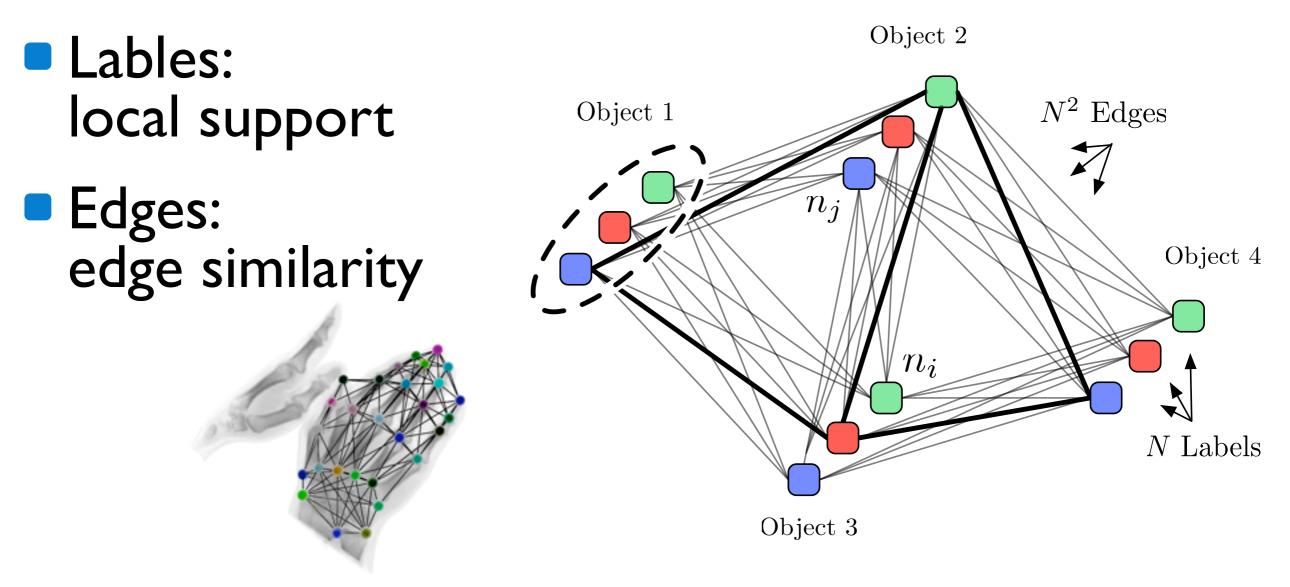
- Specified topology, following anatomical connectivity
- Estimate Gaussian distribution of length and orientation for each edge





Constuct Markov Random Field

- Same topology as geometric model
- 28 Nodes, with N~28*50 labels representing the landmark candidates



Edge similarities

- For each model edge e between model landmarks A and B:
 Object 1
 - For all pairs (Aj,Bj) of landmark candidates:

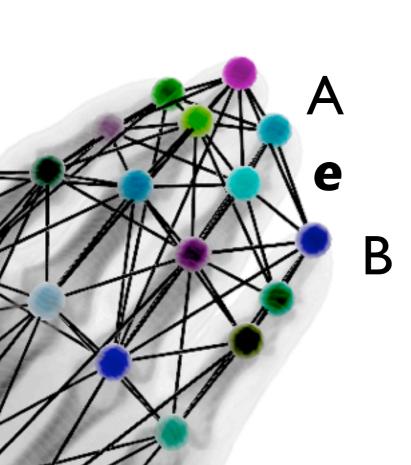
 n_i

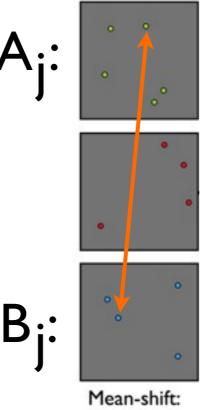
Object 3

Х

Model for **e**

Similarity = max(p_a, p_b) of the nonnormalized Gaussian model for e

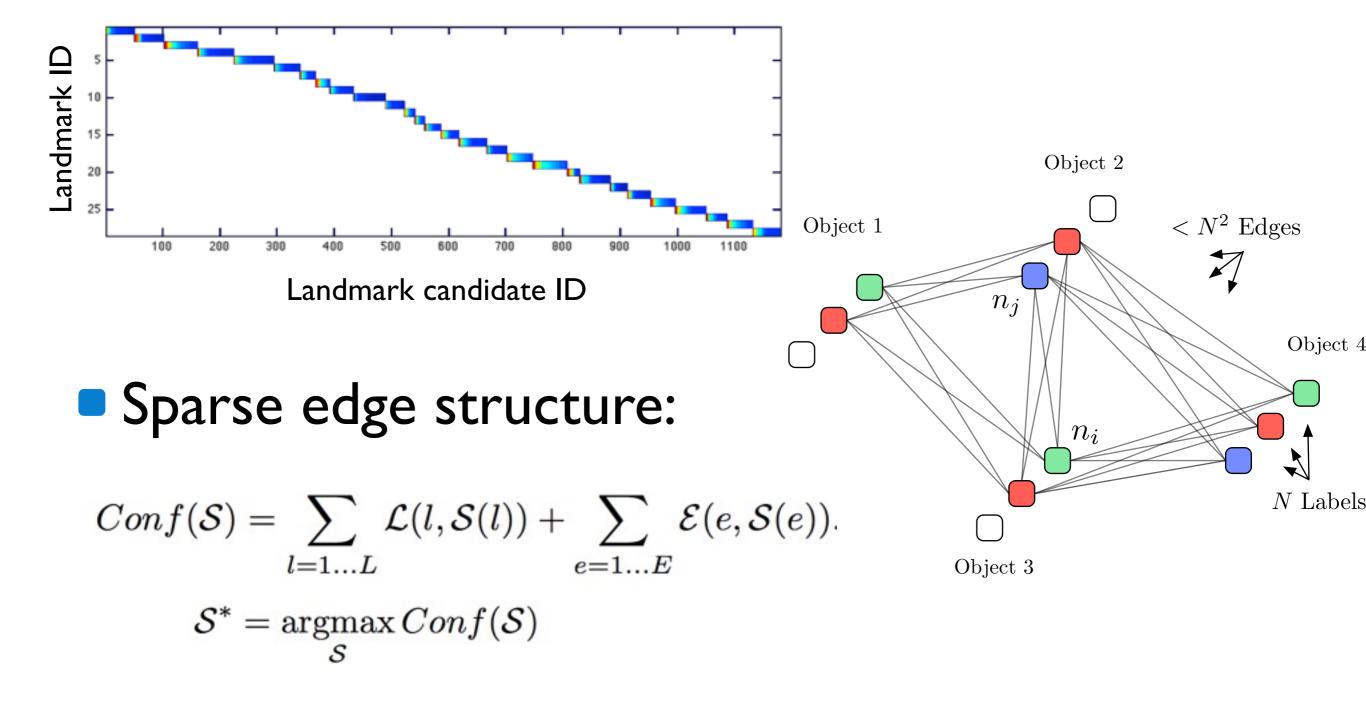




Mean-shift: candidates

Sparse MRF structure

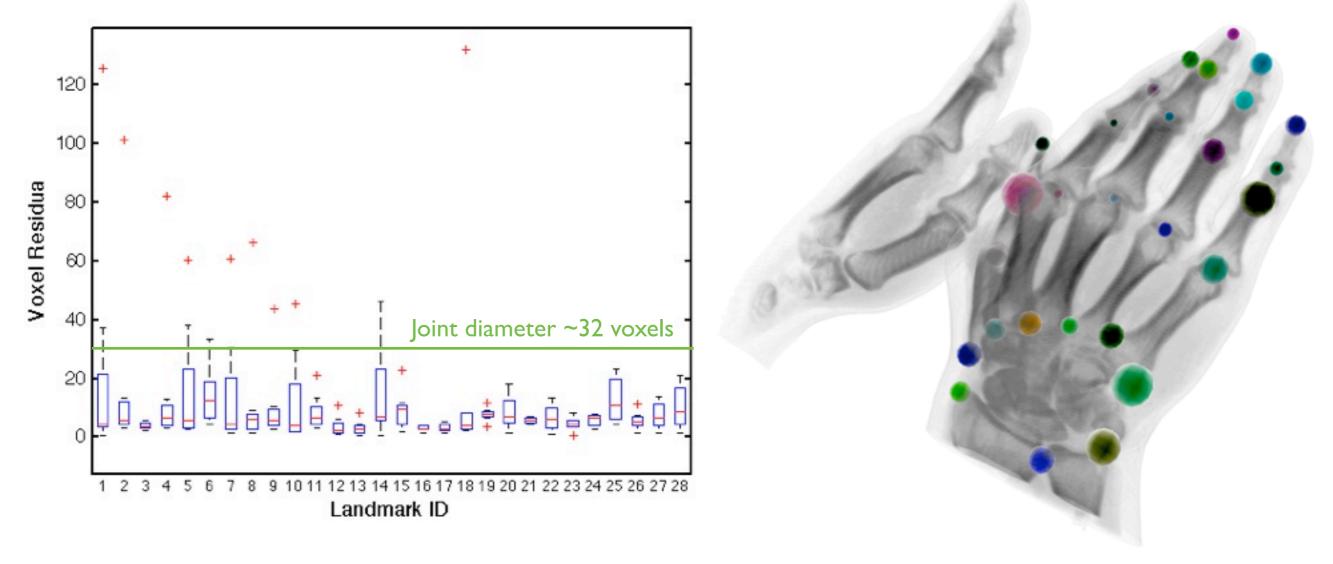
Sparse point confidences



- 8 high-resolution hand CTs
- 256 × 384 × 330 voxels
- 28 landmarks selected in each training image
- 33500 training samples for the RF
- Leave-one-out cross validation

Results

mean/median/std: 10.13 / 5.59 / 16.99 voxels



Distances of resulting localization to ground truth.

Sphere Radius = Median Error

Conlusion

- Simple method for localization of complex, self-similar anatomical structures
- Learned candidate detectors allow to cope with 3D data

Outlook

- Evaluation on other datasets (thorax MRI, whole body CT)
- Learning the graph topology learn landmarks
- Evaluation of MRF solvers



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