

1 Data Ingestion

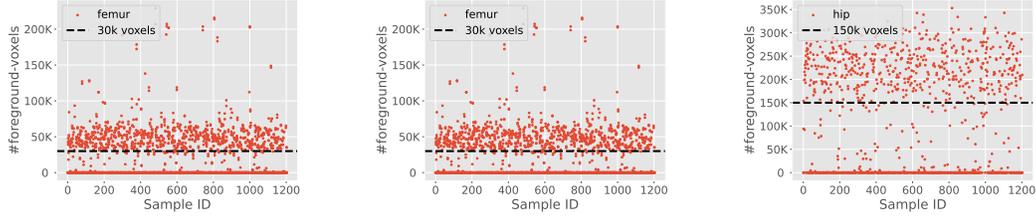


Figure 1: TotalSegmentator Dataset Ingestion: Selection of Samples was based on whether it contained a reasonable number of voxels (threshold defined individually for each anatomy) and then visually rejecting the remaining samples that contained only partial bones. Ribs were selected based on whether a full set of ribs were present.

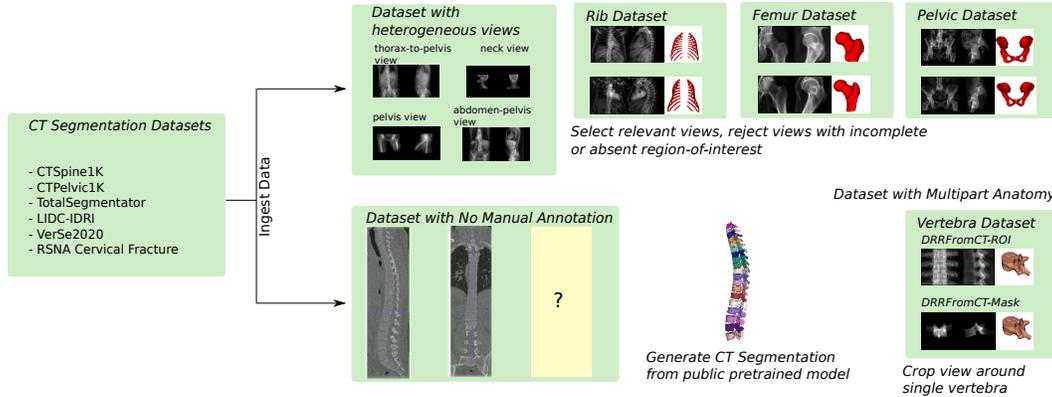


Figure 2: Data Ingestion: Various data preprocessing scenarios for Ingesting CT Segmentation Datasets for Biplanar X-ray to 3D Bone Shape Dataset

2 Benchmarking Tasks

Table 1: Benchmarking Tasks

Benchmarking Evaluation Task	Training Dataset	Testing Dataset
Architecture Comparison		
Femur	TotalSeg-Femur	TotalSeg-Femur
Hip	TotalSeg-Pelvic	TotalSeg-Pelvic
Vertebra	VerSe2019	VerSe2019
Rib	TotalSeg-Ribs	TotalSeg-Ribs
Domain Shift: Fractured Bone	TotalSeg-Pelvic	CTPelvic1k-CLINIC
	VerSe2019	RSNA Cervical Fracture
Domain Shift: X-ray with Bone Implants	TotalSeg-Pelvic	CTPelvic1k-CLINIC-Metal
Domain Shift: Cohort Shift (Population, Scanner etc.)	TotalSeg-Pelvic	CTPelvic1k-KITS19
Domain Shift: X-ray misalignment	TotalSeg-*, VerSe2019	TotalSeg-*, VerSe2019

3 Hyperparameter Tuning

- 4 We split the dataset into train-val-tests by first splitting the whole dataset into the train-test split
- 5 in the 85:15 ratio and then again splitting the train-split into the train-val split in the 85:15 ratio.
- 6 We considered model selection for each of the models using the Dice Score metric on the train-val
- 7 split. We used this validation performance to select the best hyperparameter setting and estimate the
- 8 model training epochs. We then retrain the model using both train- and val-split as training data for

9 a fixed number of epochs determined during model selection and report metrics on test-split using
 10 the last epoch checkpoint. We choose the last epoch checkpoint since choosing the epoch with the
 11 best test-split metric would result in test-split leakage. We use default model sizes for off-the-shelf
 architectures such as AttentionUNet, UNETR and UNet.

Method	Task	Encoder Channels	Kernel Size	lr
TLPredictor	femur	8,16,32 8,16,32,64 16,32,64,128,256	3,5	2e-3 2e-4

Method	Task	Encoder Channels	Decoder Channels	latent dim	kernel size	lr
AutoEncoder	rib	4,8,16,32	4,8,16,32	64	3	2e-3 2e-4
		8,16,32,64	8,16,32,64			
		8,16,32,64,128	8,16,32,64,128			
		16,32,64,128,256,128	16,32,64,128,256,128			

Method	Task	Encoder Channels	Decoder Channels	fusion channels,depth	kernel size	lr
MultiScale2DConcat	femur	4,8,16	4,8,16	32,2	3	1e-2 2e-3
		8,16,32	8,16,32	32,3		
		4,8,16,32	4,8,16,32	32,4		
		8,16,32,64	8,16,32,64	32,5		
		8,16,32,64,128	8,16,32,64,128	32,6		
		4,8,16,32,64,128	4,8,16,32,64,128			

Method	Task	Encoder Channels	Decoder channels	latent dim	kernel size	lr
1DConcat	femur	32,64,128,256	128,1024,512,8,4,4,4	128	3,5	2e-2
		32,64,128,256,512	128,1024,512,256,128,64,32 256,1024,512,256,128,64,32	256		

Method	Task	Feature Size	num heads	dropout rate	lr
SwinUNETR	vertebra	12	2,2,2,2 3,6,12,24	0.1	2e-3
		24			
		48			

Method	Task	Encoder Channels	Decoder channels	kernel size	lr
AttentionUnet/Unet		8,16, 32, 64, 128	128,64,32,16,8	3	2e-2

Table 2: Model Architecture and Hyperparameter tuning configurations

12

13 4 Replicating reimplemented architectures

14 We performed an as-close-as-possible replication of Bayat et al and obtained comparable results
 15 (95.31% DSC in the original work vs. 94.43% in our replication), which is reasonable given that
 16 there is one important step without relevant information in the original paper that precludes exact
 17 replication. The training set in the original work was not the full set of images in the dataset, but an
 18 unknown subset. This is because the ground truth for the 3D reconstruction task in their case was a
 19 silver standard mask predicted by a deep learning segmentation model, and a radiologist manually
 20 went through the masks and removed 50 data points whose masks were deemed implausible. Original
 21 work reported results on the manually cleaned silver-standard dataset, but the information regarding
 22 the exact scans from the LIDC dataset that were discarded has not been made public.

23 For all other remaining architectures, the reported results are from private datasets. Some of our
 24 key motivations for this work are because of these challenges. Lack of reproducibility and disparate
 25 dataset quality makes it difficult for new methods to be compared with existing ones, which could
 26 potentially continue for newer methods in future if a common setting is not made available.

27 5 Benchmark Framework Usage

28 Configuration File

```
1 ---
2
3 # subject-list
4 subjects:
5   subject_basepath: 2D-3D-Reconstruction-Datasets/lidc/subjectwise
6   subject_list: configs/subjects_list/lidc_subject_list.lst
7
8 # xray image properties
9 xray_pose:
10  _load: xray_pose_conf/${ROI_properties.axcode}_pose.yaml
11  res: ${ROI_properties.res}
12  size: ${ROI_properties.size}
13  drr_from_ct_mask: ${ROI_properties.drr_from_ct_mask}
14  drr_from_mask: ${ROI_properties.drr_from_mask}
15
16 # output directories
17 out_directories:
18  _load: directory_conf/dir_ct.yaml
19
20 # ROI extraction properties
21 ROI_properties:
22  axcode: PIR
23  extraction_ratio:
24    L: 0.5
25    A: 0.5
26    S: 0.5
27  ct_padding: -1024
28  seg_padding: 0
29  drr_from_ct_mask: False
30  drr_from_mask: False
31  res: 1.0
32  size: 96
33
34 # filename conventions
35 filename_convention:
36  input:
37    ct: "ct.nii.gz"
38    seg: "seg.nii.gz"
39  output:
40    vert_xray_ap: "${id}_vert-{vert}_ap.png"
41    vert_xray_lat: "${id}_vert-{vert}_lat.png"
42    vert_centroid: "${id}_vert-{vert}_centroid.nii.gz"
43    vert_centroid_xray_ap: "${id}_vert-{vert}_ap_centroid.png"
44    vert_centroid_xray_lat: "${id}_vert-{vert}_lat_centroid.png"
45    vert_ct: "${id}_vert-{vert}_ct.nii.gz" # add 'vert' for vertebra
46    vert_seg: "${id}_vert-{vert}-seg-vert_msk.nii.gz"
47    vert_overlay_ap: "${id}_vert-{vert}_ap_overlay.png"
48    vert_overlay_lat: "${id}_vert-{vert}_lat_overlay.png"
49
```

29 **6 Clinical Metrics**

30 **6.1 Vertebra Morphometry**

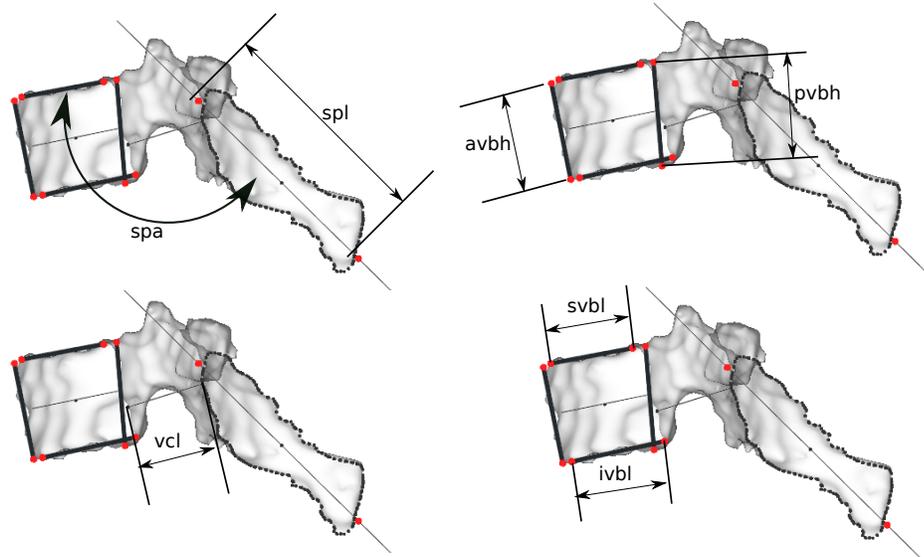


Figure 3: Vertebra Morphometry Metrics

31 **Femur Morphometry** We automatically extract Femoral Head Radius(FHR) and Neck Shaft Angle (NSA) from Femur Segmentation by adapting [?]. The following adaptations were made: i) Since
 32 full-length femur bones were not available, automatic estimation of the diaphysis axis as described in
 33 [?] was not possible. Hence, manual localization of the subtrochanteric region was performed on
 34 groundtruth segmentation and then transferred to predicted segmentation. This localization allows
 35 robust circle fitting to estimate the diaphysis axis. ii) Some of the samples do not even contain
 36 enough subtrochanteric region to reliably estimate the femur diaphysis axis. For these examples,
 37 Neck Shaft Angle(NSA) cannot be estimated. Additionally, [?] requires estimation of the diaphysis
 38 axis for robust localization of the femoral head and neck region. As an alternative, for such cases, we
 39 transfer femoral head and neck localization from the groundtruth. The manual localization of the
 40 subtrochanteric region is provided in the Benchmarking Framework Repository.
 41

42 We find that the variability due to these modifications is similar to the original method except for
 43 slightly increased variability in estimating (Femur Diaphysis Axis) FDA as shown in fig 4. We think
 44 that this ambiguity is due to not having enough subtrochanteric and diaphysis regions to accurately
 estimate FDA.

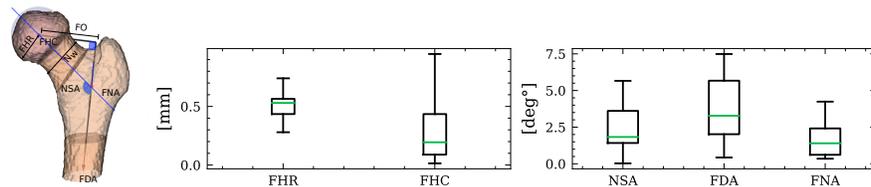


Figure 4: Repeatability of the femur morphometry extraction method as measured by error distributions for a) the landmarks/anatomical sizes and b) axis alignment identified by the adapted method.

45

Method	In-domain [TOTALSEG]	OOD [KITS19]	Δ	OOD [CLINIC]	Δ	OOD [CLINIC-METAL]	Δ	In-domain [Verse19]	OOD [RSNA]	Δ
SwinUNETR	85.78	77.68	8.09	76.04	9.74	74.71	11.06	83.59	73.42	10.18
AttentionUnet	85.03	78.64	6.39	75.22	9.81	72.14	12.89	83.66	73.23	10.43
2DConcat	84.75	79.52	5.22	75.50	9.25	69.93	14.82	83.62	72.77	10.85
UNet	84.45	77.93	6.52	73.96	10.49	72.64	11.80	82.17	69.80	12.37
MultiScale2DConcat	84.48	73.48	11.00	73.83	10.65	68.79	15.69	81.85	70.83	11.03
UNETR	82.27	75.82	6.45	72.41	9.86	69.79	12.48	81.84	71.39	10.45
TLPredictor	79.33	61.00	18.33	66.92	12.41	67.07	12.26	79.20	65.74	13.46
OneDConcat	78.85	60.39	18.46	65.52	13.33	67.36	11.49	80.92	69.35	11.57

Table 3: Reduction in Performance due to Domain Shift: The reduction in DSC (represented by column Δ) when comparing In-Domain performance with Out-of-Domain(OOD) performance shows the need for robustness against relevant shifts for clinical acceptance.

46 **7 Supplement to Quantitative Analysis of DSC vs clinical parameters**

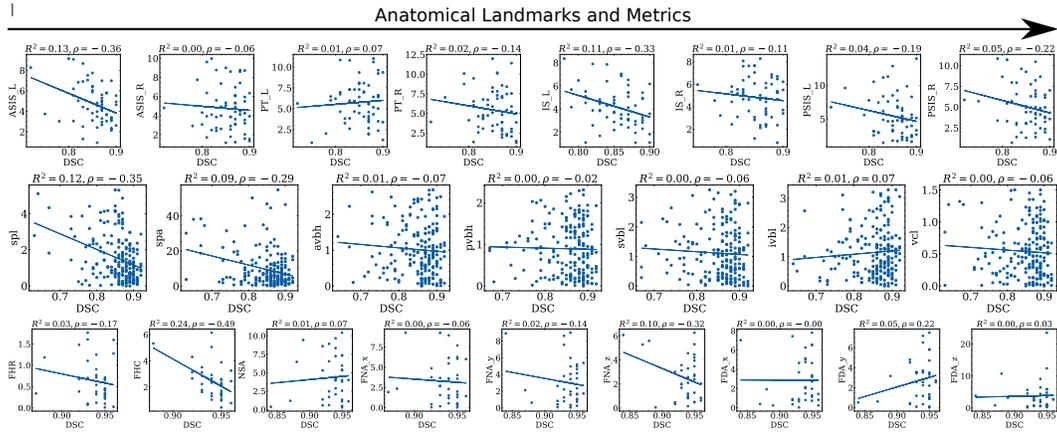


Figure 5: Relationship between Dice and Clinical Metrics across data samples on a single architecture(AttentionUnet): Top (Hip), Middle (Vertebra) and Bottom (femur)

47 **8 Qualitative Visualization**

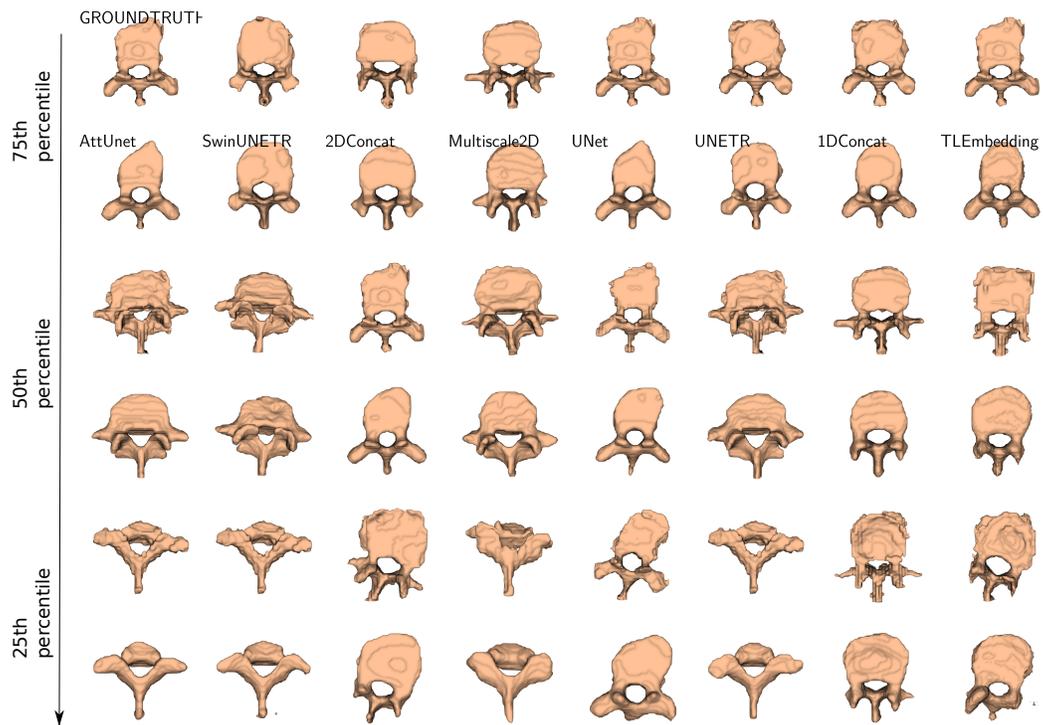


Figure 6: Vertebra Qualitative Results: 1st, 3rd and 5th row are groundtruth for corresponding architectures, whereas 2nd, 4th and 6th row are model predictions. The vertical axis represents the best(75th percentile), median and worse(25th percentile) samples for each architectures.

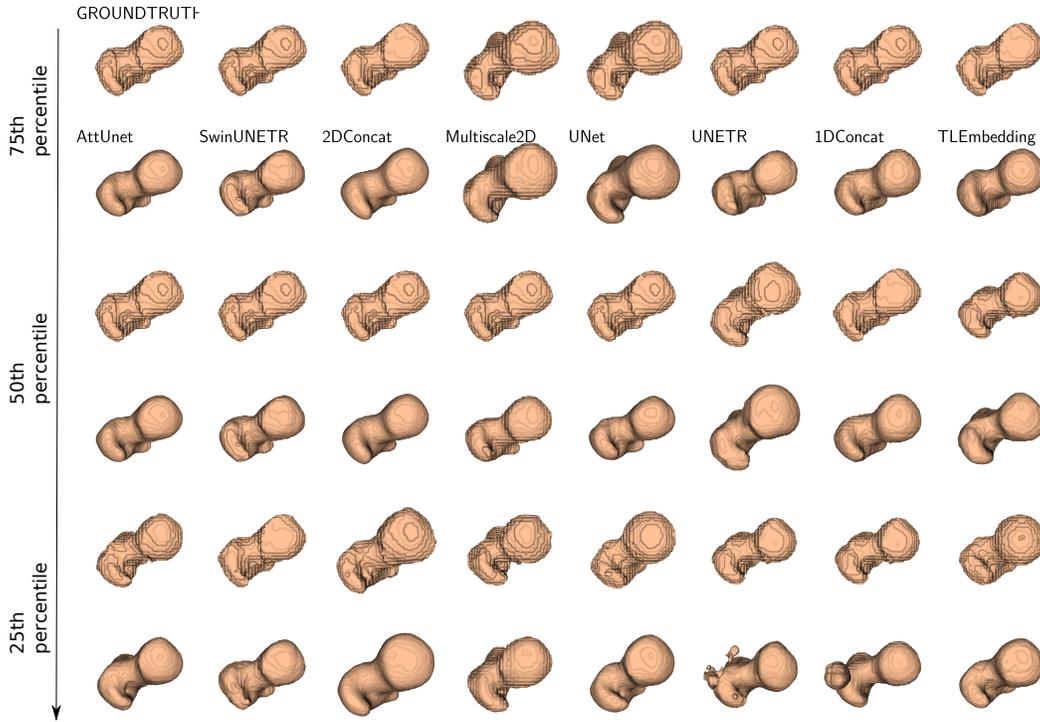


Figure 7: Femur Qualitative Results: 1st, 3rd and 5th row are groundtruth for corresponding architectures, whereas 2nd, 4th and 6th row are model predictions. The vertical axis represents the best(75th percentile), median and worse(25th percentile) samples for each architecture.



Figure 8: Rib Qualitative Results: 1st, 3rd and 5th row are groundtruth for corresponding architectures, whereas 2nd, 4th and 6th row are model predictions. The vertical axis represents the best(75th percentile), median and worse(25th percentile) samples for each architecture.



Figure 9: Hip Qualitative Results: 1st, 3rd and 5th row are groundtruth for corresponding architectures, whereas 2nd, 4th and 6th row are model predictions. The vertical axis represents the 75th percentile, median, 25th percentile and worse samples for each architecture.

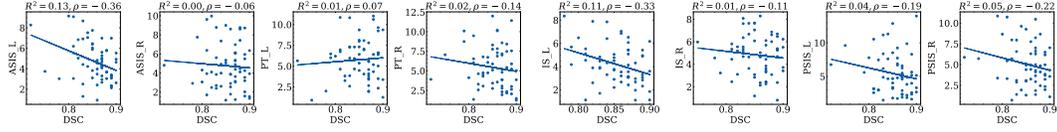


Figure 10: Hip/AttentionUnet

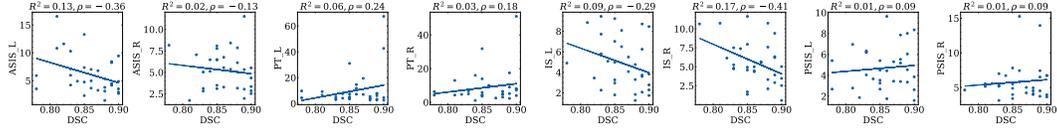


Figure 11: Hip/SwinUNETR

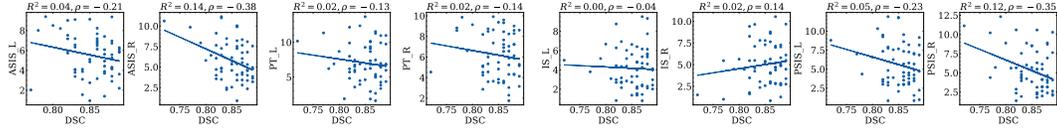


Figure 12: Hip/2DConcat

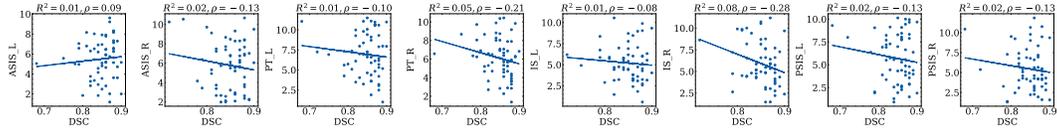


Figure 13: Hip/Multiscale2DConcat

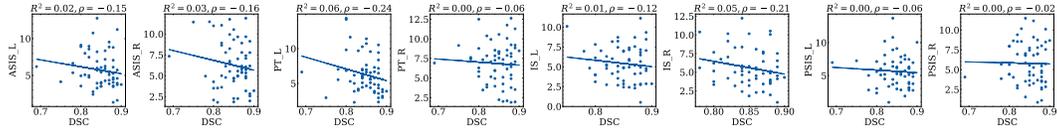


Figure 14: Hip/UNet

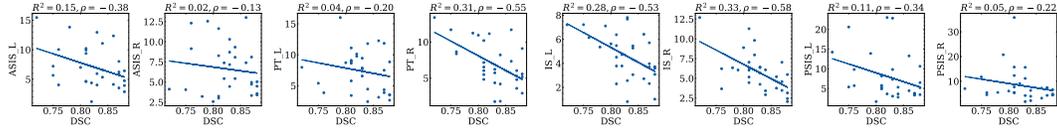


Figure 15: Hip/UNETR

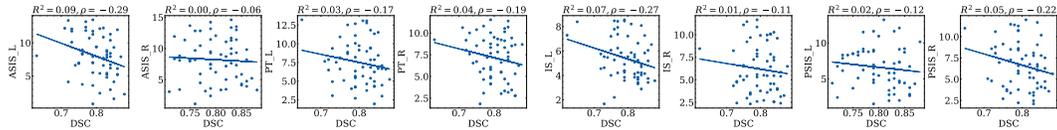


Figure 16: Hip/TLPredictor

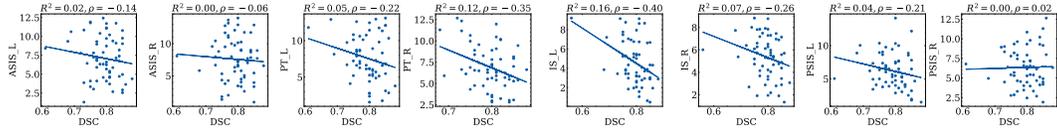


Figure 17: Hip/1DConcat

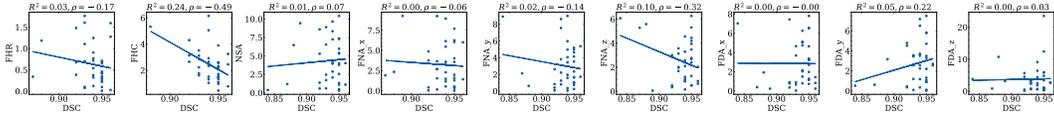


Figure 18: femur/AttentionUnet

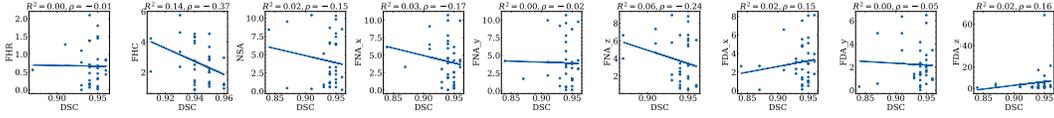


Figure 19: femur/SwinUNETR

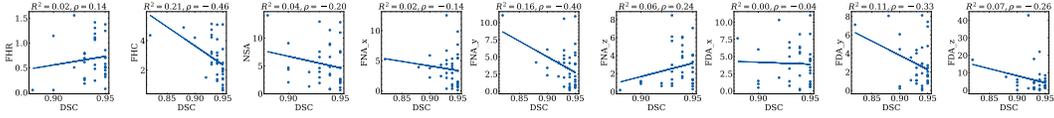


Figure 20: femur/2DConcat

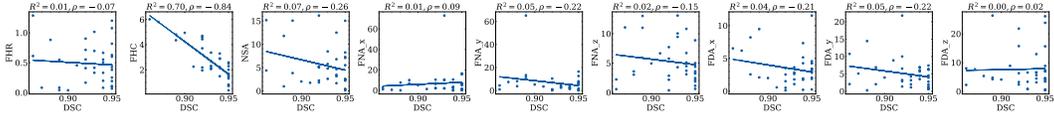


Figure 21: femur/Multiscale2DConcat

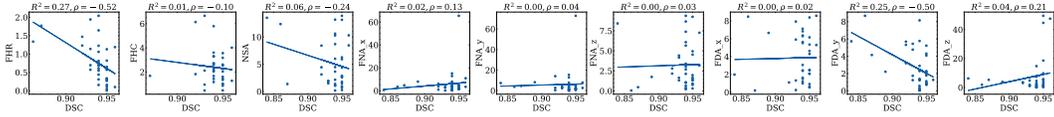


Figure 22: femur/UNet

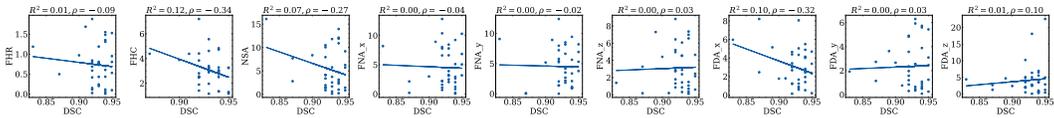


Figure 23: femur/UNETR

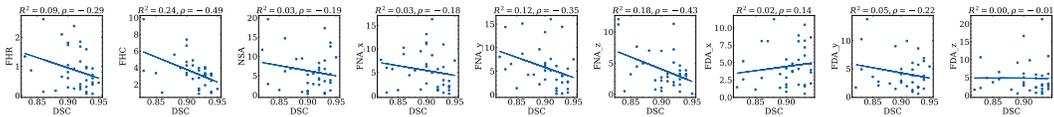


Figure 24: femur/TLPredictor

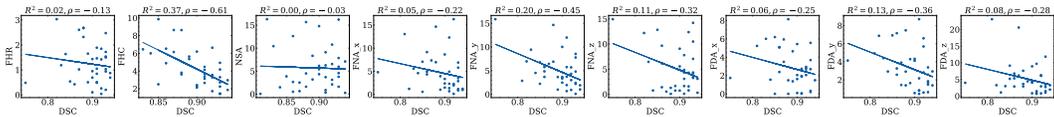


Figure 25: femur/1DConcat

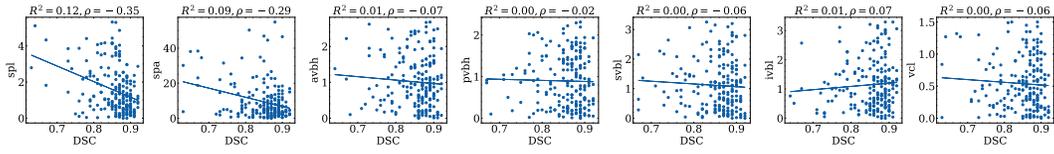


Figure 26: vertebra/AttentionUnet

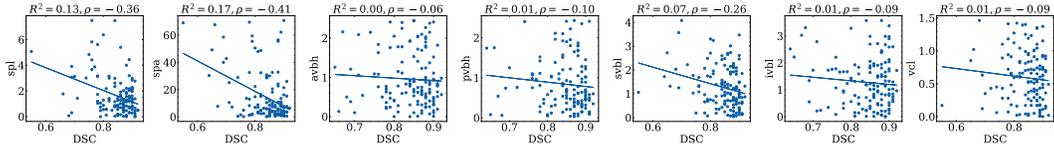


Figure 27: vertebra/SwinUNETR

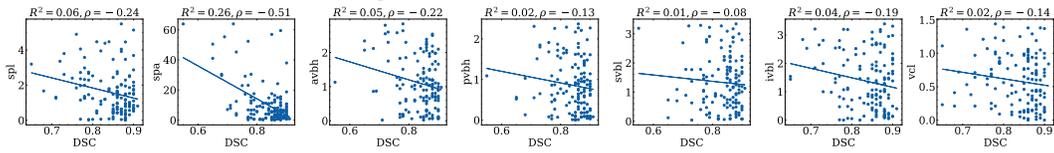


Figure 28: vertebra/2DConcat

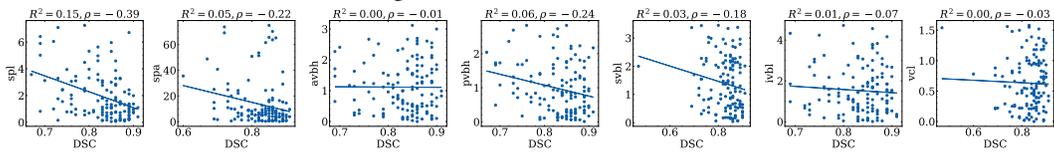


Figure 29: vertebra/Multiscale2DConcat

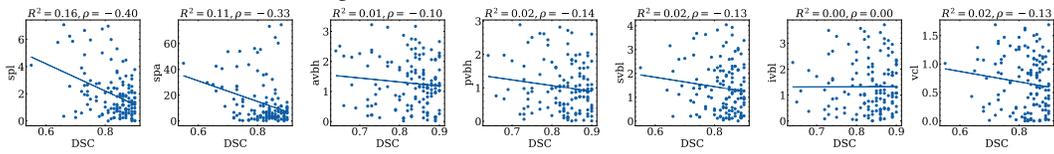


Figure 30: vertebra/UNet

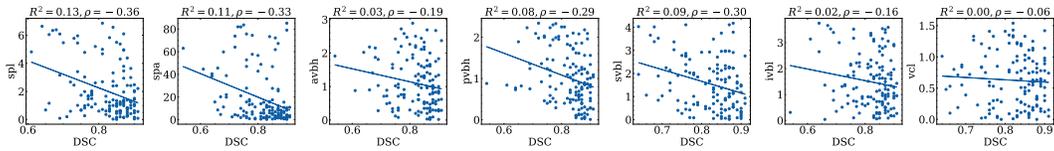


Figure 31: vertebra/UNETR

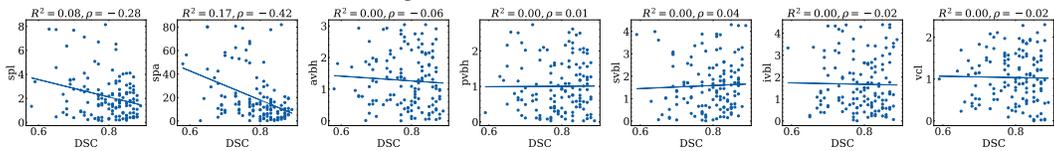


Figure 32: vertebra/TLPredictor

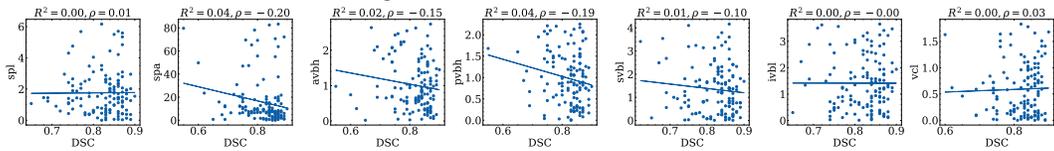


Figure 33: vertebra/1DConcat

48 **Checklist**

- 49 1. For all authors...
- 50 (a) Do the main claims made in the abstract and introduction accurately reflect the paper’s
- 51 contributions and scope? [Yes]
- 52 (b) Did you describe the limitations of your work? [Yes]
- 53 (c) Did you discuss any potential negative societal impacts of your work? [N/A]
- 54 (d) Have you read the ethics review guidelines and ensured that your paper conforms to
- 55 them? [Yes]
- 56 2. If you are including theoretical results...
- 57 (a) Did you state the full set of assumptions of all theoretical results? [N/A]
- 58 (b) Did you include complete proofs of all theoretical results? [N/A]
- 59 3. If you ran experiments (e.g. for benchmarks)...
- 60 (a) Did you include the code, data, and instructions needed to reproduce the main experi-
- 61 mental results (either in the supplemental material or as a URL)? [Yes] We will release
- 62 the GitHub repositories containing benchmarking framework and bone morphometry
- 63 extraction scripts.
- 64 (b) Did you specify all the training details (e.g., data splits, hyperparameters, how they were
- 65 chosen)? [Yes] Details for hyperparameter tuning are attached in the supplementary
- 66 section. Data splits are available in the GitHub repository.
- 67 (c) Did you report error bars (e.g., with respect to the random seed after running ex-
- 68 periments multiple times)? [No] We perform hyperparameter tuning in preliminary
- 69 experiments and perform the main experiments with one seed and one set of hyperpa-
- 70 rameters because they are computationally expensive.
- 71 (d) Did you include the total amount of compute and the type of resources used (e.g.,
- 72 type of GPUs, internal cluster, or cloud provider)? [Yes] We used a single NVIDIA
- 73 RTX3090 and NVIDIA 1080Ti GPU to train models. 50 hyperparameter tuning runs
- 74 each running 1 hour on average were performed. Full Training runs lasted from 1-2
- 75 hours.
- 76 4. If you are using existing assets (e.g., code, data, models) or curating/releasing new assets...
- 77 (a) If your work uses existing assets, did you cite the creators? [Yes]
- 78 (b) Did you mention the license of the assets? [Yes] Verse2019: CC BY-SA 4.0 License,
- 79 TotalSegmentator: Apache-2.0 license, CTPelvic1k: no explicit licence mentioned, but
- 80 made publicly available
- 81 (c) Did you include any new assets either in the supplemental material or as a URL? [Yes]
- 82 The filtered subject-id used from the original dataset and additional segmentation used
- 83 for bone morphometry will be provided in the GitHub repository.
- 84 (d) Did you discuss whether and how consent was obtained from people whose data you’re
- 85 using/curating? [N/A] All datasets we used were publicly available and hence explicit
- 86 consent was not required.
- 87 (e) Did you discuss whether the data you are using/curating contains personally identifiable
- 88 information or offensive content? [N/A] None of the Datasets we use contain personally
- 89 identifiable information or offensive content.
- 90 5. If you used crowdsourcing or conducted research with human subjects...
- 91 (a) Did you include the full text of instructions given to participants and screenshots, if
- 92 applicable? [N/A]
- 93 (b) Did you describe any potential participant risks, with links to Institutional Review
- 94 Board (IRB) approvals, if applicable? [N/A]
- 95 (c) Did you include the estimated hourly wage paid to participants and the total amount
- 96 spent on participant compensation? [N/A]