Test-Time Selection for Robust Skin Lesion Analysis



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Artifact-based Domain Generalization ISIC Workshop @ ECCV 2022

Artifact-based Domain Generalization of Skin Lesion Models

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Abstract. Deep Learning failure cases are abundant, particularly in the medical area. Recent studies in out-of-distribution generalization have advanced considerably on well-controlled synthetic datasets, but they do not represent medical imaging contexts. We propose a pipeline that relies







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Test-time Debiasing Literature Alignment with the Clinical Workflow

	Method	#Keypoints	AUC		
Baseline	Test-time augmentation	-	58,4		
Literature	T3A	Ι	56,7		
Literature	Tent	_	54,1		
Literature	NoiseCrop	50.176	72,7		



- Fail when only a single image is available.
- Fail when test distribution is heterogenous.



Test-time Debiasing Literature Alignment with the Clinical Workflow



- Relies on full segmentation masks, which are <u>hard to annotate</u>.
- Make modifications in the pixelspace, which <u>might introduce</u> <u>unexpected features.</u>













Test-time Selection (TTS)

- Fast to annotate.
- Avoid introducing distribution shifts by intervening on the feature space.
- Does not rely on test batch statistics.
- It's cheap as there are no model updates.

	Method	#Keypoints	AUC		
Baseline	Test-time augmentation	-	58,4		
Literature	T3A		56,7		
Literature	Tent	—	54,1		
Literature	NoiseCrop	50.176	72,7		
Ours	TTS	40	75,0		





Methodology

Methodology





Methodology

Attention Maps





PositiveNegativeKeypointsKeypoints



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Evaluation Protocol

Artifacts providing spurious correlations



Dark Corners

Hair



Ruler

Ink markings



Patches

factor	\mathbf{set}	dark	hair	ruler	ink	patches
		corner				
0	train	0.119	-0.104	0.142	0.023	-0.138
	test	0.135	-0.112	0.162	0.030	-0.149

- Mild correlations.
- Gaining robustness to artifacts will hardly impact any metric.
- We need to control/amplify the correlations.

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Spurious Features vs. Generalization



Sagawa, Shiori, et al. "Distributionally robust neural networks for group shifts: On the importance of regularization for worst-case generalization.", ICLR 2020



Spurious Features vs. Generalization



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Results

Effective throughout different training biases

Training Bias

Effective even with a single pair of keypoints

Effect of # Keypoints on Performance

Keypoints

Visualization

Keypoints

Keypoints

Before TTS

After TTS

Visualization

Keypoints

After TTS

Keypoints Before TTS ≍

After TTS

Flexible for different types of annotation

Keypoints on Artifacts

Keypoints from Segmentation Mask

Comparison of TTS Methods for Different Number of Keypoints

Number of Keypoints

Limitations

How to adapt this solution to Vision Transformers?

acquisition devices.)

• How to deal with biases uniformly spread across the image? (e.g., different

Takeaways

- Consider evaluating your models' robustness on trap sets
- TTS improves robustness across different levels of bias
- TTS is effective even with a single pair of keypoints
- TTS is flexible to different types of annotations

Code, Data & Paper: https://github.com/alceubissoto/skin-tts

Thank you

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Spurious Features vs. Generalization

<u>Uncontrolled biases (because not</u> <u>synthetic)</u>

Specific bias source not annotated:

- Image acquisition devices and protocol
- Artifacts

Benign

Malignant

Validation of artificial intelligence prediction models for cancer diagnosis using dermoscopy images: the 2019 International Skin Imaging Collaboration Grand Challe

Marc Combalia^{*}, Noel Codella^{*}, Veronica Rotemberg^{*}, Cristina Carrera, Stephen Dusza, David Gutman, Brian Helba, Harald Kittler, Nicholas R Kurtansky, Konstantinos Liopyris, Michael A Marchetti, Sebastian Podlipnik, Susana Puig, Christoph Rinner, Philipp Tsc Jochen Weber, Allan Halpern^{*}, Josep Malvehy^{*}

Summary

Background Previous studies of artificial intelligence (AI) applied to dermatology have shown AI to diagnostic classification accuracy than expert dermatologists; however, these studies did not adeque clinically realistic scenarios, such as how AI systems behave when presented with images of disease care not included in the training dataset or images drawn from statistical distributions with significant training distributions. We aimed to simulate these real-world scenarios and evaluate the effects of i institution, diagnoses outside of the training set, and other image artifacts on classification accuracy, with informing clinicians and regulatory agencies about safety and real-world accuracy.

			MEL							
		Ref	0.14	0.04	0.7	0.0079	0.039	0.037	0.0076	0.00
	ſ	No crust	0.14	0.04	0.7	0.008	0.039	0.037	0.0077	0.00
		Crust	0.016	0.019	0.83	0.0024	0.052	0.054	0	C
For skin structure of the goal	Artifacts	No hair	0.14	0.038	0.7	0.0083	0.044	0.032	0.0064	0.00
		Hair	0.15	0.052	0.67	0.0059	0.011	0.067	0.015	0.0
		No pen	0.13	0.039	0.71	0.0083	0.039	0.033	0.008	0.00
		Pen	0.23	0.048	0.53	0.0024	0.037	0.098	0.002	0.00
		No pigmentation	0.0075	0.018	0.40	0.045	0.046	0.27	0.075	0.00
		Pigmentation	0.15	0.041	0.71	0.0056	0.039	0.022	0.0032	0.00
		No ulceration	0.16	0.043	0.69	0.0053	0.042	0.024	0.005	0.00
	l	Ulceration	0.0014	0.013	0.74	0.028	0.02	0.13	0.027	0.00
	ſ	Torso	0.21	0.016	0.69	0.004	0.0076	0.037	0.012	0.00
	site	Head or neck	0.021	0.089	0.64	0.0035	0.15	0.059	0.0002	C
	lical	Lower extremity	0.17	0.022	0.73	0.011	0.0043	0.019	0.0021	0.0
	Anatom	Upper extremity	0.13	0.057	0.67	0.015	0.022	0.051	0.01	0.00
		Palms or soles	0.079	0.025	0∙8	0.017	0.012	0.016	0.018	0.00
	l (Oral or genital								

MELAN NEL-BY NEL-NEL NEL NEL-AN NEL-BC WASC NEL-DE NEL-NT

Combalia et al. "Validation of artificial intelligence prediction models for skin cancer diagnosis using dermoscopy images: the 2019 International Skin Imaging Collaboration Grand Challenge", The Lancet, 2022

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