

# Interpreting Fine-grained Dermatological Classification with Deep Learning

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**ISIC Skin Image Analysis Workshop**



LONG BEACH  
CALIFORNIA  
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# Scope

- Analyze model accuracy gap on benchmark datasets (CIFAR-10) vs. dermatological image corpus (DermAI\*)
  - SOTA on CIFAR ~98%, whereas dermoscopic ~90%
- Investigate leading label pairs by case studies
  - 3 leading pairs investigated by GradCAM/GBP
- Suggestions on better datasets of user-submitted images by our experience
  - Data Augmentation, FoV, Gamma & Illumination correction

# Dataset

User submitted Dermoscopic images across 10 most prevalent labels. 7264 images, split in 5:1 (train/test)



Acne



Alopecia



Blister



Crust



Erythema



Leukoderma



P. Macula



Tumor



Ulcer

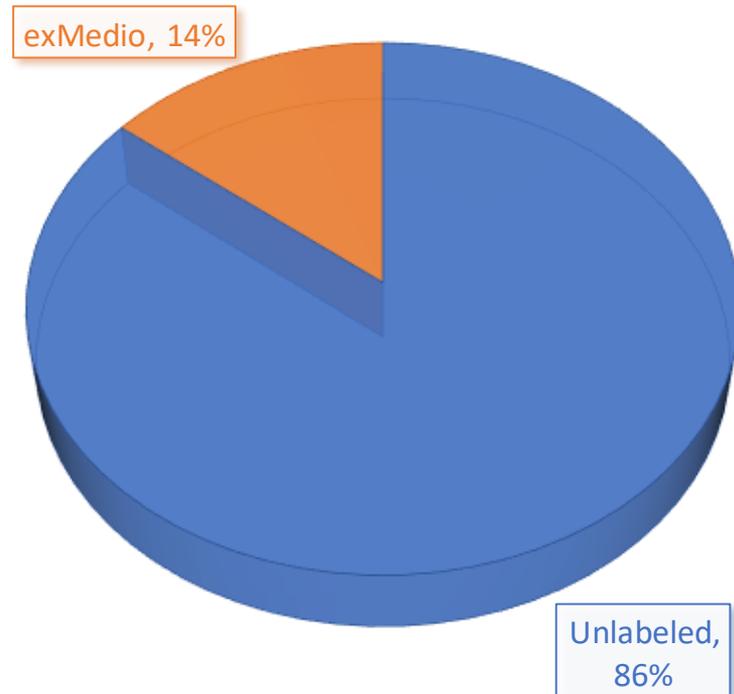


Wheal

# Dataset

- Addressing the most common dermatological complaints.
- Ultimate goal:  
To perform reliable rapid screening to reduce out-patient burden.

DERMATOLOGICAL TYPES COVERED



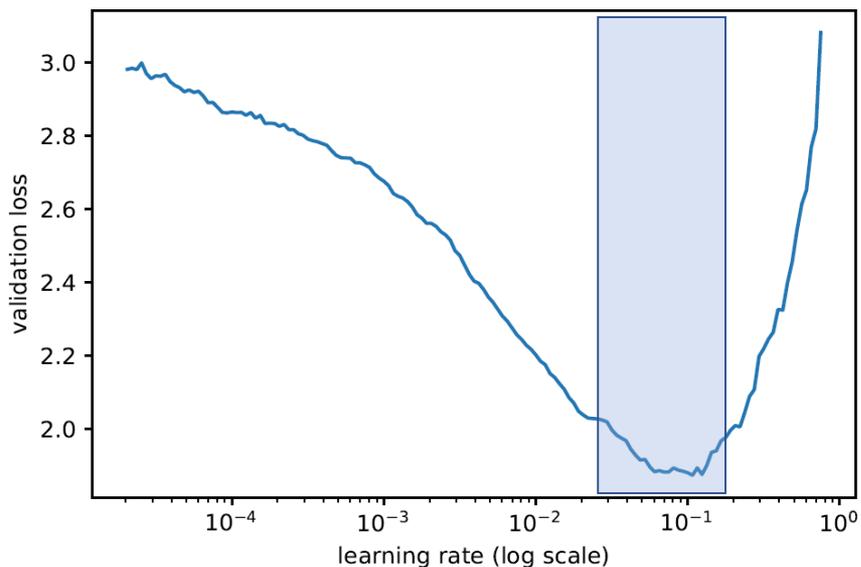
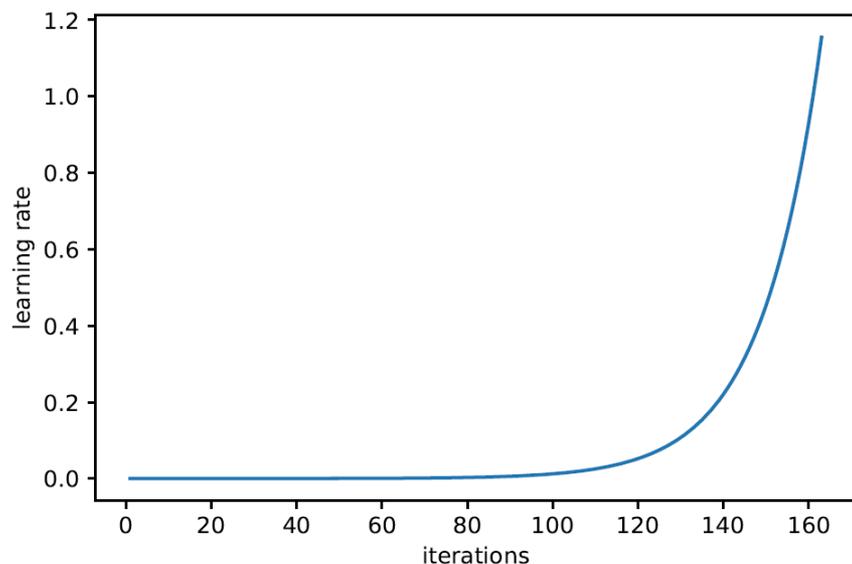
# Model Learning

- Test several architectures of increasing size/complexity

Resnet-34, ResNet-50, ResNet-101, ResNet-152

- 5:1 split, Early stopping, BCE with logits loss
  - Learning rate range test
  - SGD + Restarts (SGD-R)
  - SGD-R + Length Multiplication+ Differential Learning
- Modus operandi tested on CIFAR-10 prior\*

# Learning Rate range-test



Steadily increase the LR and observe the Cross entropy loss  
Test several mini-batches to see a point of inflexion

*Reference:*

Cyclical Learning rates for training NN, L. Smith [2017]

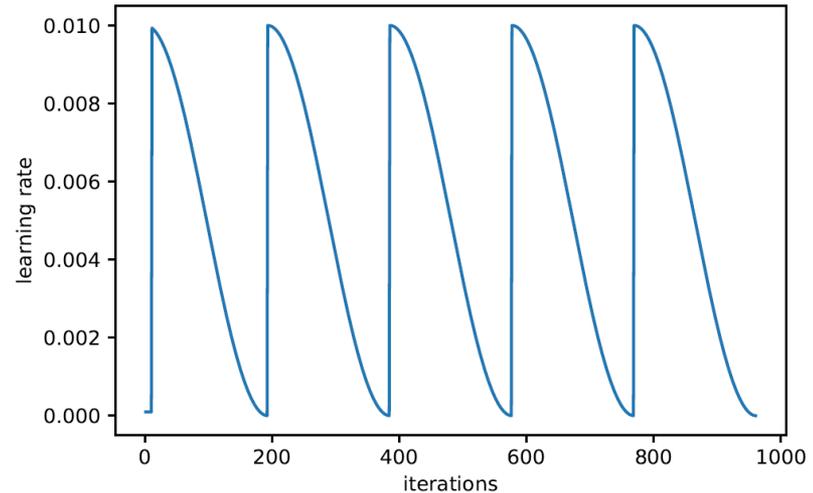
Deep Learning, S. Verma et al. 2018

# SGD-R

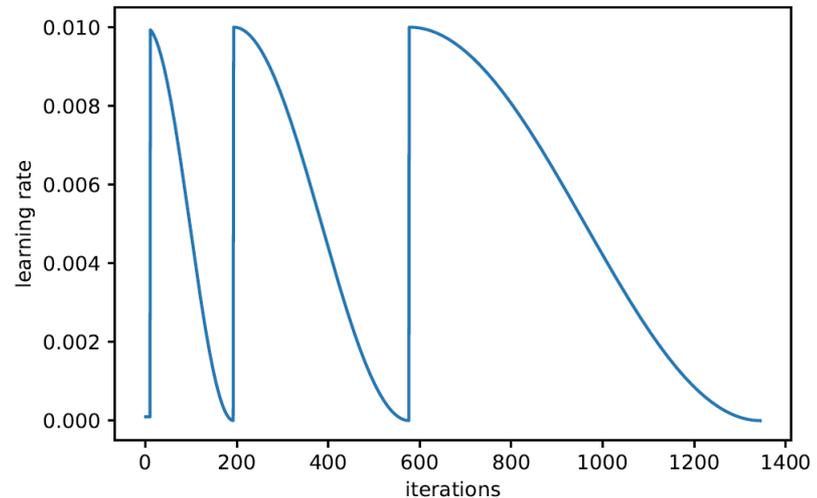
1. Avoid monotonicity by Cosine scheduling function

$$v(t) = \frac{1}{2} \left( 1 + v \cos \left( \frac{t\pi}{T} \right) \right) + \epsilon$$

2. Cycle Length Multiply by integral powers of 2 over whole architecture



Initial coarse fit by tuning the last (or last few) FC layer



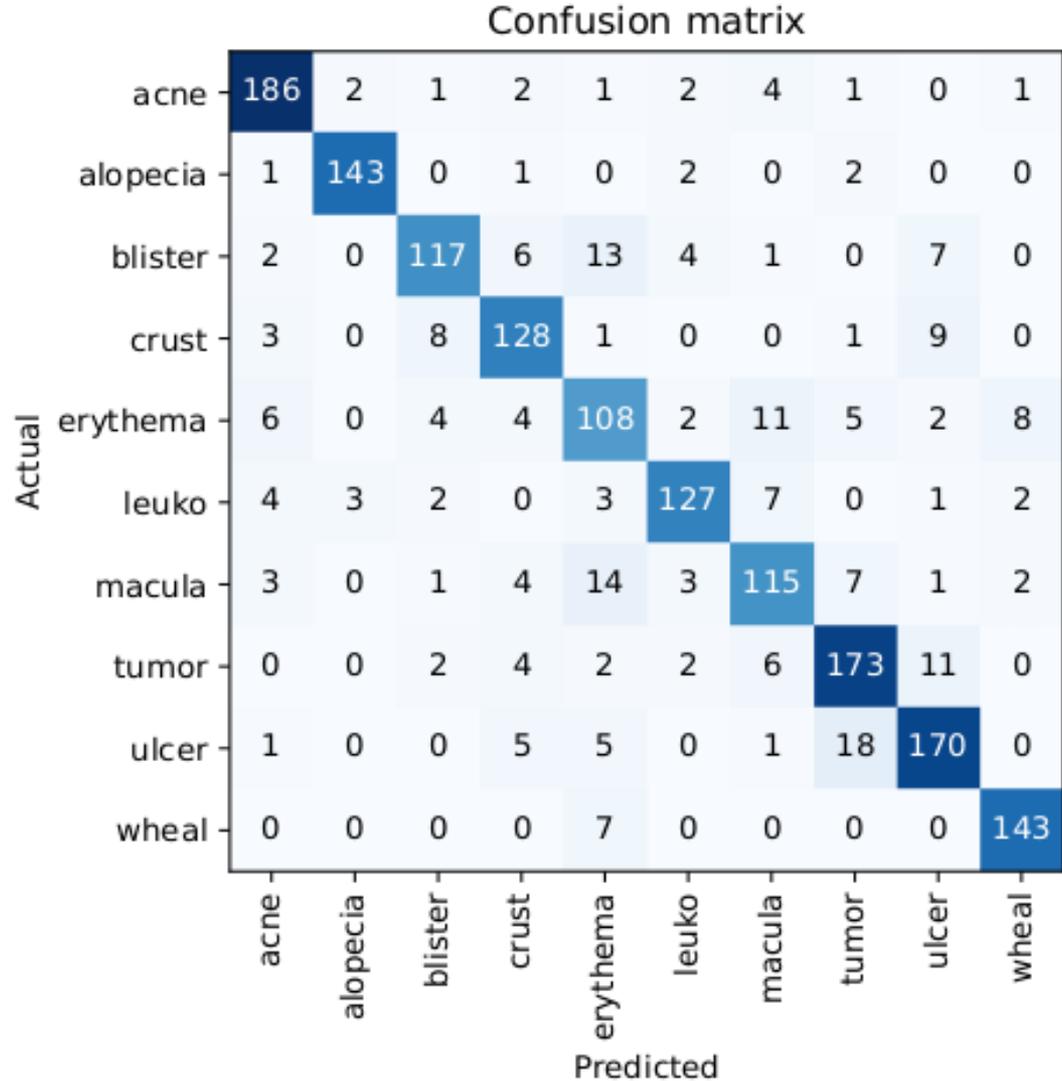
Tighter fit over all layers

Reference:

SGD with Warm restarts, Loschilov [2017]

# Application

Architecture	Acc. (Top-1)
ResNet-34	88.9%
ResNet-50	89.7%
ResNet-101	88.2%
ResNet-152	89.8%



ResNet 152 Confusion Matrix

# Analysis

- Following best practices still leaves gap.
- Focus on the label pairs which account for most errors.
- Use GradCAM and Gradient Backprop to analyze what CNNs capture in learning process.

Label 1	Label 2	Counts
Ulcer	Tumor	29
Macula	Erythema	25
Blister	Erythema	17
Erythema	Wheal	15
Crust	Ulcer	14
Blister	Crust	14
Macula	Tumor	13
Macula	Leukoderma	10
Blister	Ulcer	7
Tumor	Erythema	7
Crust	Tumor	5

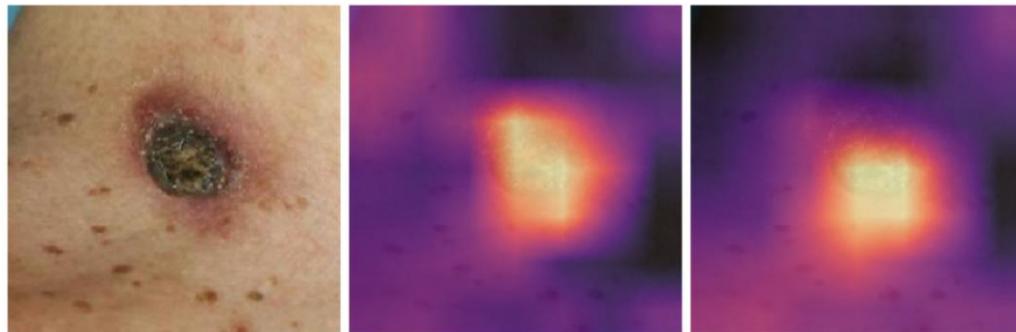
*Label pairs with at least 5 errors*

*Reference:*

GradCAM: Visual explanation from DNN, Selvaraju [2016]

Guided BP, Springenberg [2014]

# Ulcers & Tumors



**Ulcer 0.391**

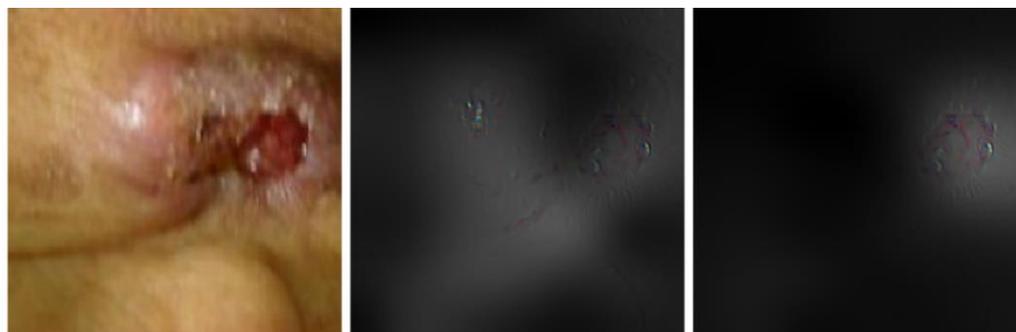
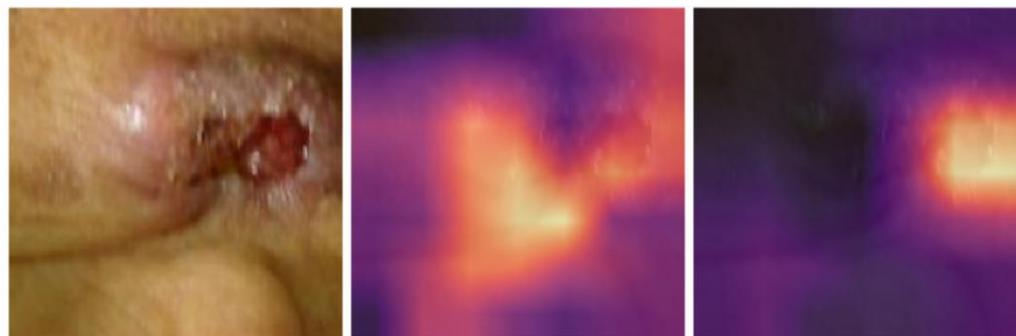
**Tumor 0.152**

High degree of geometrical (spherical) similarity is the common factor in many samples

**Tumor 0.78**

**Ulcer 0.212**

Elevations and inflammations seen in Tumors, misclassifies many ulcer samples.



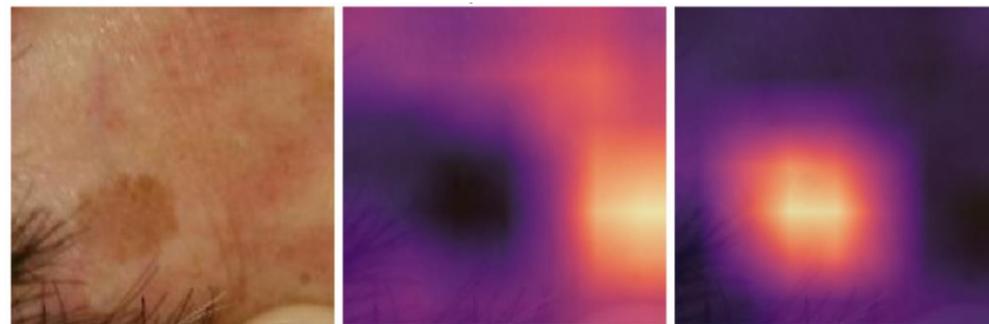
# Macula & Erythema

**Erythema 0.53**

**Macula 0.41**

Presence of pigmentation patches around the lesion can mispredict.

*FoV and ROI selection could lead to better results.*

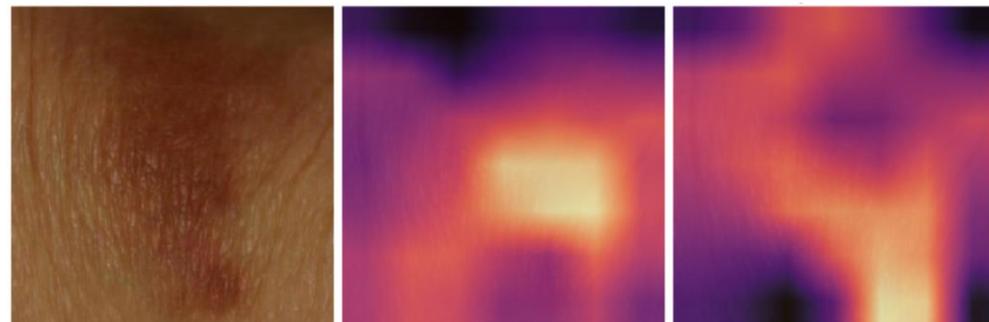


**Macula 0.69**

**Erythema 0.28**

Oval/cycloidal patches makes GBP confused with the overall shape of Macula.

FOV & Depth important factors to consider



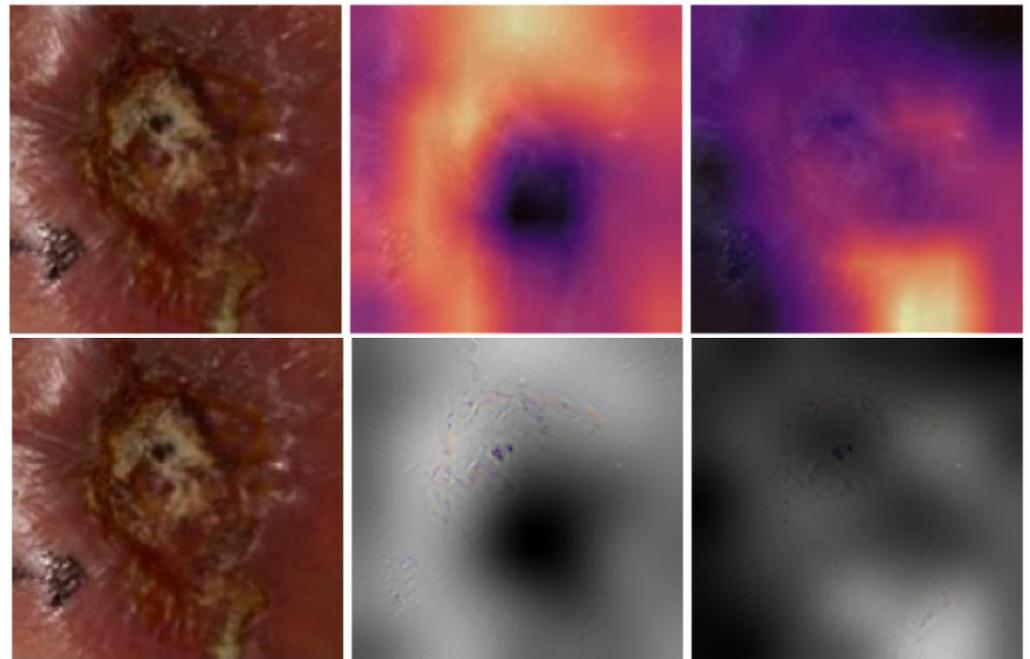
# Ulcer & Crust

**Crust 0.86**

**Ulcer 0.124**

Presence of large centroid is possible source.

*Difficult to predict as both related chronologically*

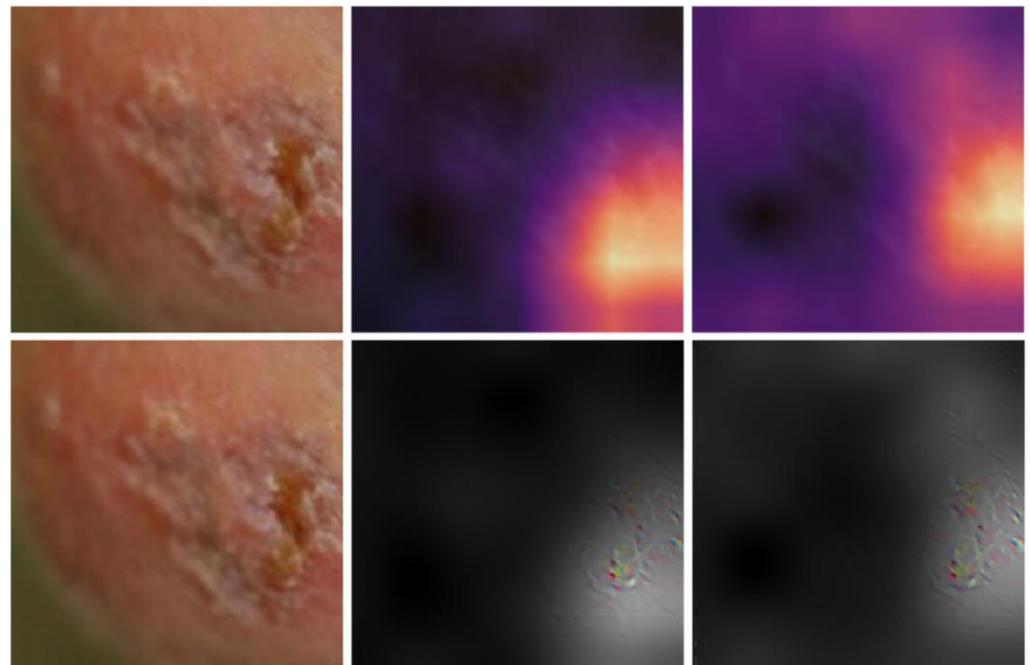


**Ulcer 0.91**

**Crust 0.06**

Oval/cycloidal patches on GBP

*Selection of right RoI, illumination could improve many cases.*



# Mitigation

Highlight some of the “hard-learned lessons” building this project from scratch.

Mitigation factors to look out:

- Balancing training sets (dynamic vs static)
- Field of View / ROI selection
- Illumination and Gamma correction

# Balancing for model learning

Confusion matrix

Actual \ Predicted	acne	alopecia	blister	crust	erythema	leuko	macula	tumor	ulcer	wheel
acne	30	0	0	0	7	0	3	0	0	0
alopecia	0	20	0	0	0	1	0	0	0	0
blister	0	0	25	1	18	0	1	0	0	0
crust	0	1	1	11	9	0	2	3	5	0
erythema	8	0	8	2	653	2	13	6	3	5
leuko	0	1	0	0	9	43	7	0	0	0
macula	0	0	0	0	39	4	201	3	3	0
tumor	0	0	0	1	10	1	6	48	14	0
ulcer	0	0	0	3	7	0	0	7	139	0
wheel	0	0	0	0	23	0	1	0	0	2

Confusion matrix

Actual \ Predicted	acne	alopecia	blister	crust	erythema	leuko	macula	tumor	ulcer	wheel
acne	186	2	1	2	1	2	4	1	0	1
alopecia	1	143	0	1	0	2	0	2	0	0
blister	2	0	117	6	13	4	1	0	7	0
crust	3	0	8	128	1	0	0	1	9	0
erythema	6	0	4	4	108	2	11	5	2	8
leuko	4	3	2	0	3	127	7	0	1	2
macula	3	0	1	4	14	3	115	7	1	2
tumor	0	0	2	4	2	2	6	173	11	0
ulcer	1	0	0	5	5	0	1	18	170	0
wheel	0	0	0	0	7	0	0	0	0	143

Custom datasets can be small, unevenly divided. Best to use dynamic in-memory augmentation during batch selection. Larger batches preferably.

# Field of View/Object Depth

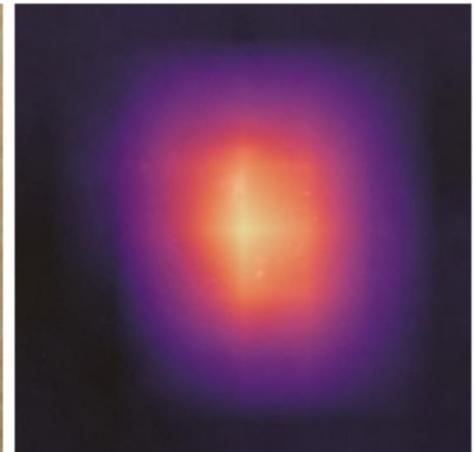
P [Blister]

0.547



P [Blister]

1.000

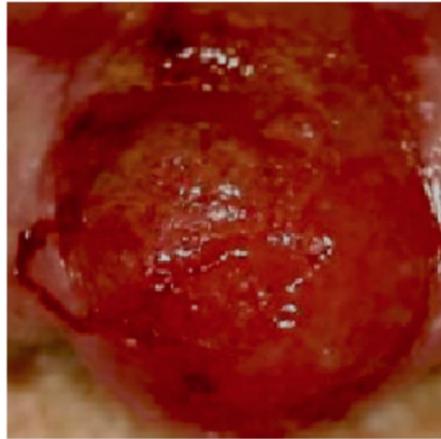


FOV selection dramatically improves performance. In user-submitted images, pre-processing needed. Bonus: if illumination stable

# Gamma & Illumination

Often illumination & shadow effects

Gamma adjustment  $\approx$   
**1.2 – 1.5**

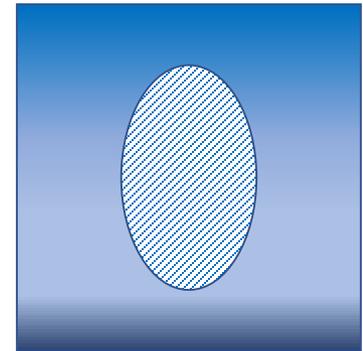
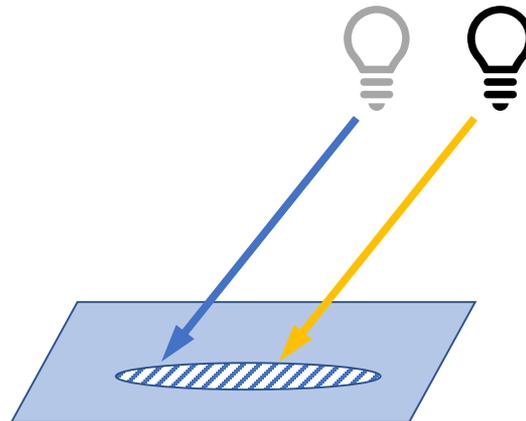


Prediction : **Ulcer (98%)**  
Actual : **Tumor (1%)**



Prediction : **Tumor 78%**

Creating illumination map & reversing imbalanced lighting by normalizing.



# Conclusion

- Gap may never be entirely removed,
- [Status Quo] Racial diversity one of the hardest problems to crack. Better to focus on single one for better performance. (But harder in developed countries).
- Not all artifacts can be fixed in user-submitted images.
- Augmentation & Photo-grammatic corrections can improve the quality of model learning/inference dramatically.
  - Balancing training data, FOV reduction, Gamma & illumination correction

<https://github.com/souravmishra/ISIC-CVPRW19>

Thank you!



# Scope

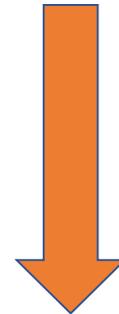
Rapid improvements in image classification tasks

- Larger better & detailed datasets
- Faster hardware resources
- Better architectures



However (the ugly truth)!

- More iterations to SOTA
- Longer train time
- Higher costs
- Small dataset reliability low



# Scope

Deployment costs can adversely impact individuals or smaller groups.

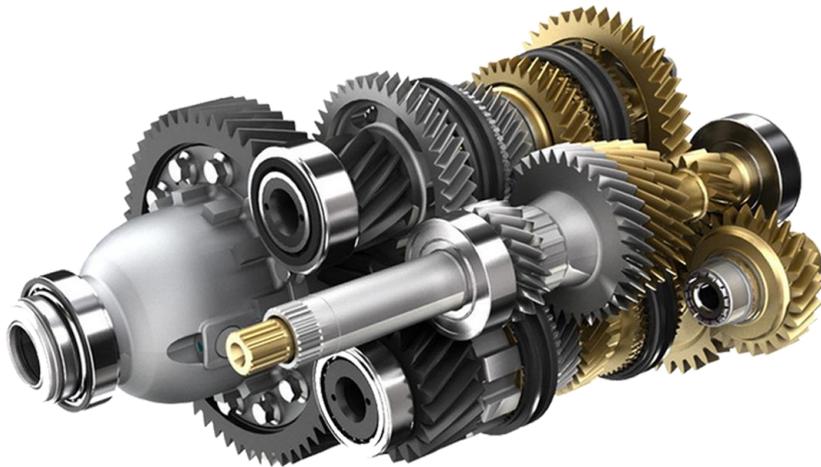
## SOLUTION?

- Organic combination of proven techniques, field tested on benchmark datasets.
- Optimization by learning rate ( $\nu$ ) adaptations.
- Transfer modus-operandi to smaller, untested data.
- Ensure repeatability.

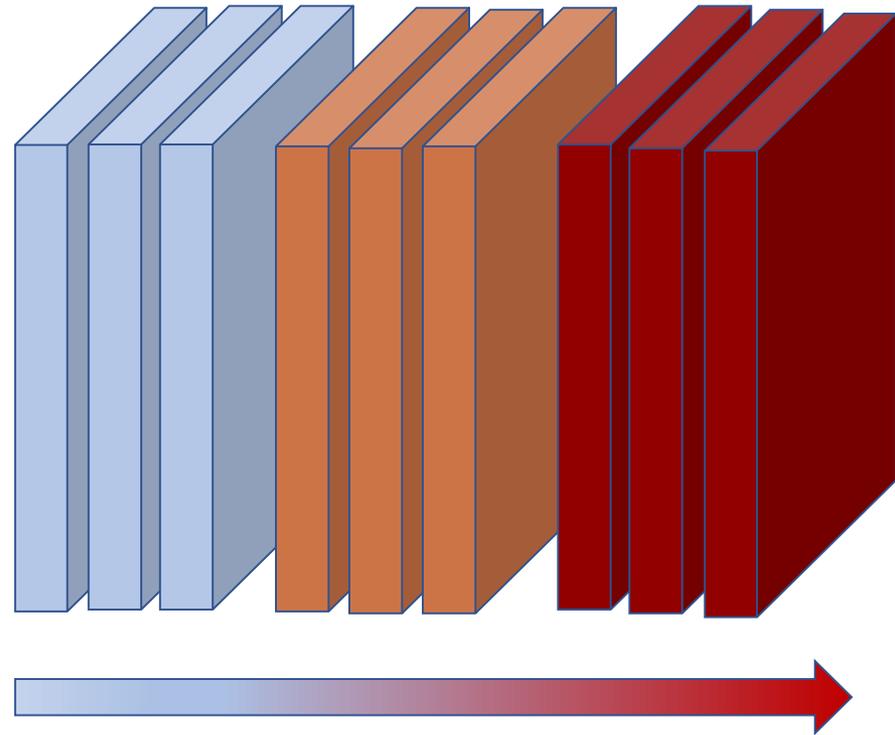
# CIFAR Baseline

- Multi-class classification on CIFAR-10
- Test candidate architectures of increasing size/complexity
  - Resnet-34, ResNet-50, ResNet-101, ResNet-152
  - DenseNet161
- Baseline Performance
  - 5:1 split, Early stopping, lower LR restarts
  - BCE with logits loss
  - Train to 90%+ validation accuracy mark

# Differential learning



Gear-box need not spin all gears equally!



Reduce computational overhead by assigning different learning rates.

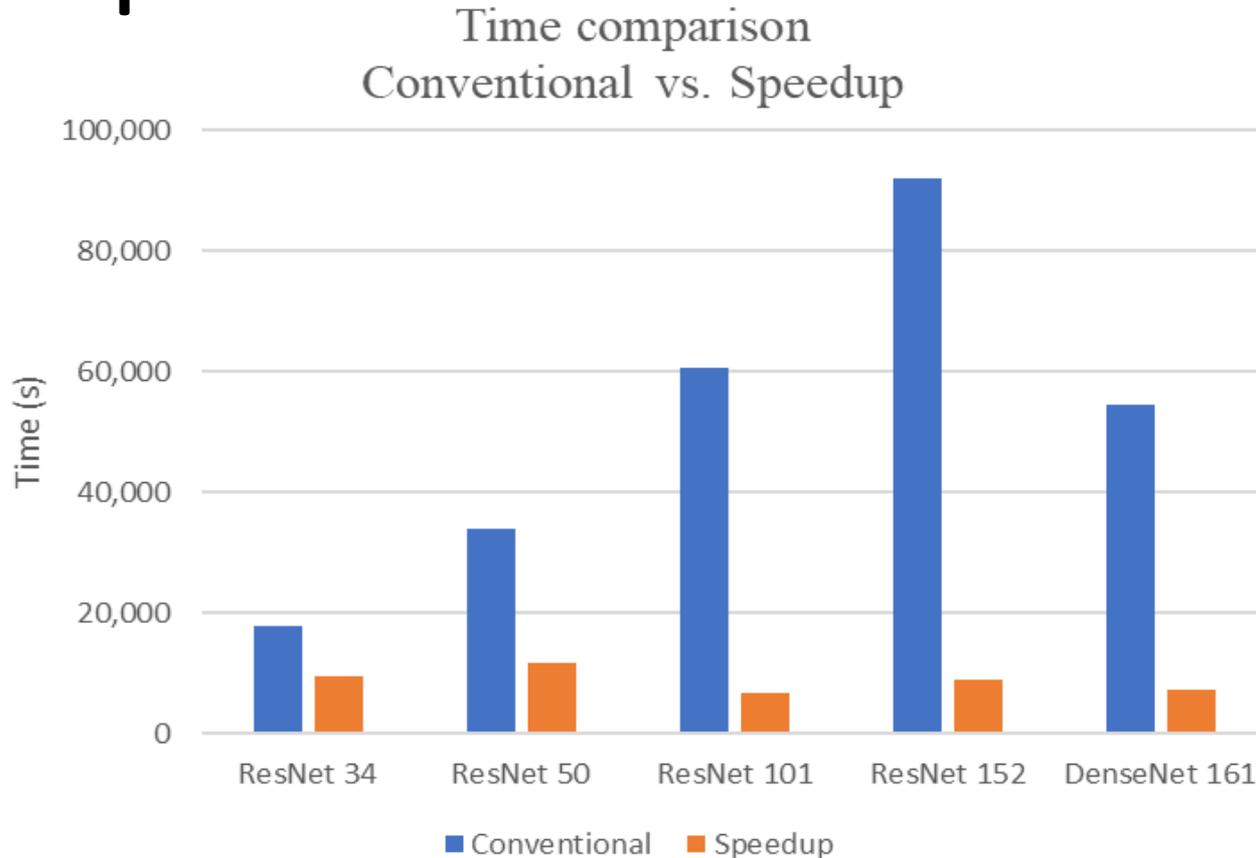
# CIFAR Baseline

Architecture	Accuracy (Top-1)	Time (s)
ResNet 34	90.36%	17,757
ResNet-50	90.54%	34,039
ResNet-101	90.71%	60,639
ResNet-152	90.68%	91,888
DenseNet-161	93.02%	54,628

# CIFAR Speedup Results

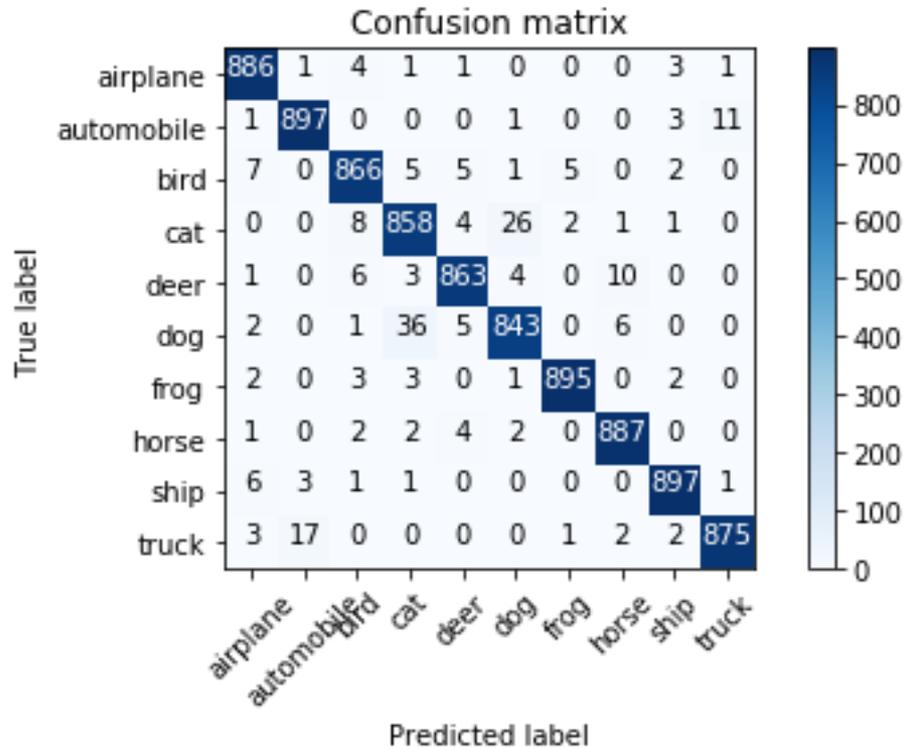
Architecture	Accuracy (Top-1)	Time (s)	$\eta$
ResNet 34	96.84%	9,565	1.84
ResNet-50	96.82%	11,817	2.88
ResNet-101	97.61%	6,673	9.09
ResNet-152	97.78%	9,012	10.2
DenseNet-161	97.15%	7,195	7.59

# Speedup Results

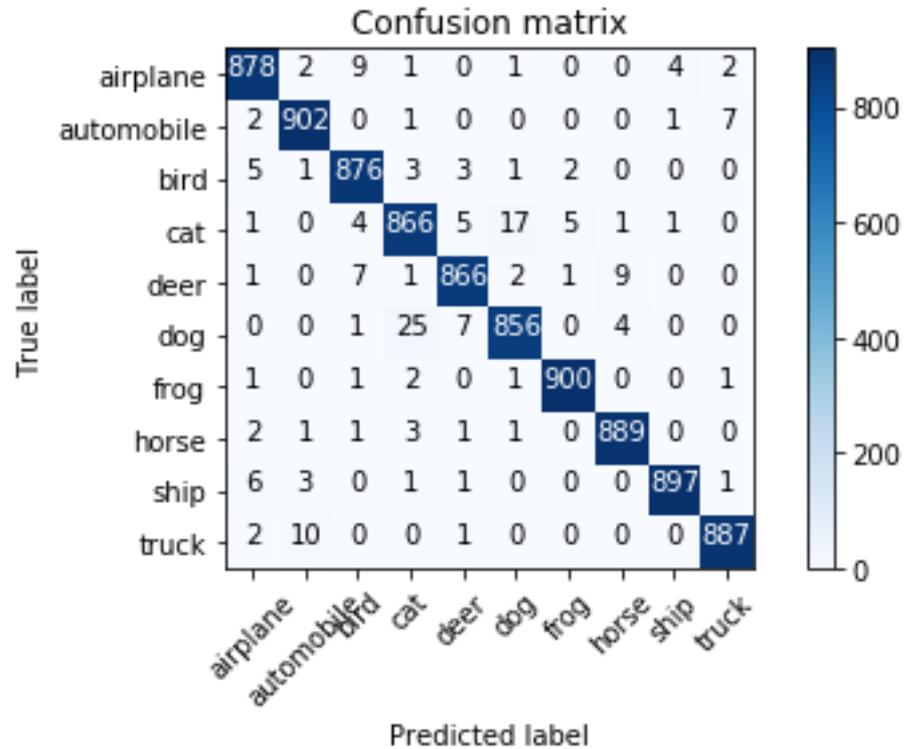


Higher dividends when architecture size grows larger.  
Possible by offsetting the computation overhead by DLR

# CIFAR Results



*DenseNet 161*



*ResNet 152*