

Requirements for Color in a Computer Aided Diagnostics Tool for Dermoscopy

Stein Olav Skrøvseth, PhD



Background

- Melanoma vs benign skin lesion based on dermoscopic imaging.
- GPs Clinical Decision Support tool (triage)
- Machine learning / pattern recognition
- Color is a feature.
- Biopsy still gold standard
- Target: reduce pathology workload

Melanoma statistics



[International Agency for Research on Cancer; 2010. http://globocan.iarc.fr]

.

The ABCD(E)-rule

- Asymmetry in shape, color or structures?
- Border

sharply delineated?

• Colors

how many distinct colors are visible?

- Differential structures which of a listed set?
- Evolution is there a change?



Machine learning





Classification of lesions

- 1. Segmentation
- 2. Feature generation
- 3. Feature selection
- 4. Classification
- 5. Validation

Sensitivity = $\frac{TP}{TP + FN}$ Specificity = $\frac{TN}{TN + FP}$



Dermoscopy

- Provides higher (OR=15.6) diagnostic accuracy compared to visual inspection.
 [Br. J. Dermatol., 2008]
- Dermoscopy is useless without training. [Lancet Oncology, 2002]
- Automated diagnosis:
 - Review (2008) found 3 systems in literature, two inferior, one on par with specialists.
 [Semin. Cutan. Med. Surg. 2008]
 - MelaFind[®], USA: Sensitivity 98.3%, Specificity: 9.9%.
 [Arch. Dermatol., 2010]
 - Many others not tested thoroughly.
- A solution could lie in multimodal imaging combined with (non-) morphological features. [Expert Rev. Anticancer Ther, 2011]



••••••

Diagnostic features : color

• The number of colors.

• A few distinctive and highly discriminate color, e.g. blue-white veil.



Blue/White veil

An irregular, structureless area of confluent blue pigmentation with an overlying white "ground-glass" haze. The pigmentation cannot occupy the entire lesion and cannot be associated with red-blue lacunes.

[dermoscopy.org]



Current approach

 53 features are extracted from each image, 11 directly related to color.

 Use perceptually uniform color space CIELab.



Color features $f = [f_1, \dots, f_{10}, f_{11}, f_{12}, f_{13}, f_{14}, f_{15}, f_{16}, f_{17}, f_{18}, f_{19}, f_{20}, \dots, f_{53}]$ Palette matching Color variability based on 3D Peripheral vs histogram. internal White, red, light brown, distribution. Mean # sample in each bin. dark brown, blue-gray, Variance # sample in each bin. • black. Pct of non-empty bins. # distinct colors % blue-gray



(a) Melanoma with all colors for training (b) Benign

(c) $f_{19} = 2, f_{20} = 0$



(d) Malignant



Results



Results, colors



Color counting

- Detect # of colors, not which.
- Color complexity.
- Very difficult to get results corresponding to dermatologist's evaluation.
- Statistically challenging.
- Will be included in the classifier.

General remarks

- In machine learning, features must be consistent.
- The exact specification must be able to extract the relevant feature.
- E.g. what is the feature relevant for the blue-white veil?

Recommendation

 Case: Image-based clinical decision support systems using machine learning methods where color is a feature.

Color specification must reflect the problem at hand.

• Colors must be consistent.



Acknowledgements

Research team:

- Thomas R. Schopf, MD
- Prof. Fred Godtliebsen
- Dr. Maciel Zortea
- Kajsa Møllersen

- Kevin Thon
- Dr. Kristian Hindberg
- Herbert Kirchesch, MD

Funding:

- Research Council Norway through Tromsø Telemedicine Laboratory (NST)
- Northern Norway Regional Health Authority

References

- Zortea, Schopf, Thon, Geilhufe, Hindberg, Kirchesch, Møllersen, Schulz, Skrøvseth, Godtliebsen, Submitted manuscipt, 2013.
- Zortea, Skrøvseth, Schopf, Kirchesch, Godtliebsen. *Automatic segmentation of dermoscopic images by iterative classification*. Int J Biomed Imaging. 2011;2011:972648.
- Zortea, Skrøvseth, Godtliebsen. Automatic learning of spatial patterns for diagnosis of skin lesions. Conf Proc IEEE Eng Med Biol Soc. 2010;2010:5601-4.
- Skrøvseth, Schopf, Thon, Zortea, Geilhufe, Møllersen, Kirchesch, Godtliebsen., "A computer aided diagnostic system for malignant melanomas," *3rd International Symposium on Applied Sciences in Biomedical and Communication Technologies (ISABEL), 2010.* pp.1,5, 7-10 Nov. 2010.