

The role of DICOM in

# Digital imaging in dermatology

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# Overview

- Drivers for skin imaging
- Modalities skin imaging
  - Digital photography
  - Dermoscopy
  - Total body photography
  - Reflectance confocal microscopy
  - Optical coherence tomography

# Digital imaging in dermatology

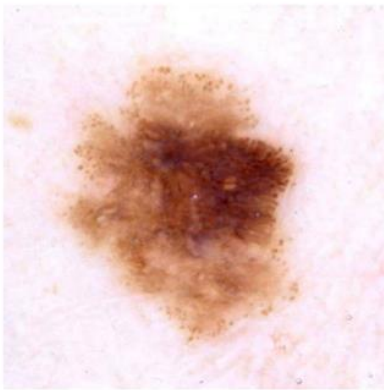
*“Imaging is increasingly being used in dermatology for documentation, diagnosis and management of cutaneous disease.”*

# Digital imaging in dermatology

- Ubiquity of digital cameras, mobile devices and EMR
- Diagnostic imaging
  - Sequential examination for changes in mole (size, shape, colour) marker for melanoma
  - Teledermatology
  - Advance modalities e.g. dermoscopy, TBP, RCM
- Diagnostic aids
  - Consensus diagnosis
  - Technology assisted full skin examination
  - Artificial intelligence

# Melanoma

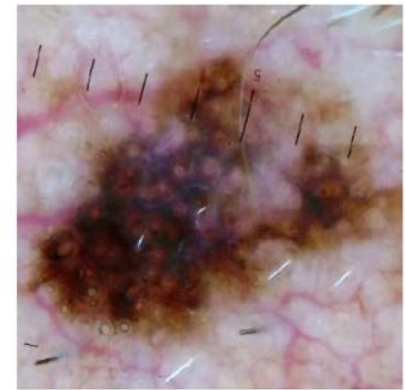
- The accuracy of the clinical diagnosis of cutaneous melanoma with the unaided eye is only about 60% (Kittler et al. 2002)
  - One of every three melanomas an error in clinical diagnosis was made



Superficial spreading  
melanoma



Nodular melanoma



Lentigo maligna melanoma

# Melanoma

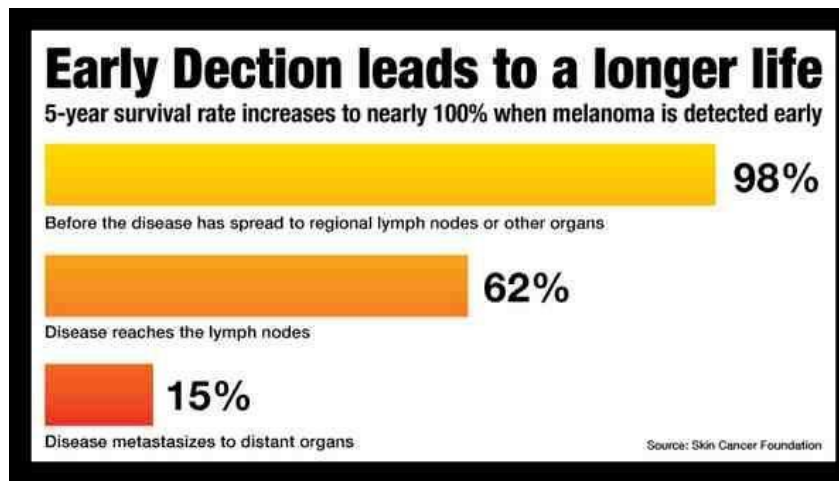
- Currently, 28.4 moles are biopsied or excised for every melanoma detected [1]
- In Australia, reducing unnecessary biopsy or excision can potentially save \$70 million in direct health system costs [2]

1. Argenziano G, Albertini G, Castagnetti F, De Pace B, Di Lernia V, Longo C, Pellacani G, Piana S, Ricci C, Zalaudek I. Early diagnosis of melanoma: what is the impact of dermoscopy? *Dermatol Ther*. 2012 Sep-Oct;25(5):403-9. doi: 10.1111/j.1529-8019.2012.01482.x.

2. Elliott, T.M., Whiteman, D.C., Olsen, C.M. et al. Estimated Healthcare Costs of Melanoma in Australia Over 3 Years Post-Diagnosis *Appl Health Econ Health Policy* (2017) 15: 805. <https://doi.org/10.1007/s40258-017-0341-y>

# Melanoma

- Earlier detection
  - 98% curative rate at 5 years for early melanoma
  - estimated average annual cost for an early stage versus late stage melanoma is \$AUD1,681 versus \$AUD115,109



# Dermatologist-level classification of skin cancer with deep neural networks

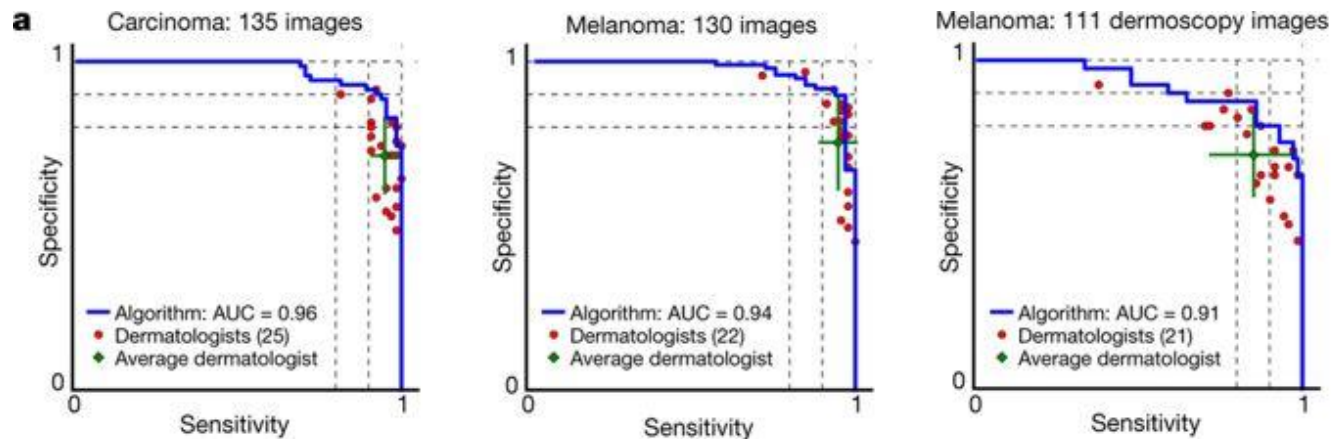
Andre Esteva<sup>1\*</sup>, Brett Kuperl<sup>1\*</sup>, Roberto A. Nova<sup>2,3</sup>, Justin Ko<sup>2</sup>, Susan M. Swetter<sup>2,4</sup>, Helen M. Blau<sup>5</sup> & Sebastian Thrun<sup>6</sup>

Skin cancer, the most common human malignancy<sup>1-3</sup>, is primarily diagnosed visually, beginning with an initial clinical screening and followed potentially by dermoscopic analysis, a biopsy and histopathological examination. Automated classification of skin lesions using images is a challenging task owing to the fine-grained variability in the appearance of skin lesions. Deep convolutional neural networks (CNNs)<sup>4,5</sup> show potential for general and highly variable tasks across many fine-grained object categories<sup>6-11</sup>. Here we demonstrate classification of skin lesions using a single CNN, trained end-to-end from images directly, using only pixels and disease labels as inputs. We train a CNN using a dataset of

images (for example, smartphone images) exhibit variability in factors such as zoom, angle and lighting, making classification substantially more challenging<sup>23,24</sup>. We overcome this challenge by using a data-driven approach—1.41 million pre-training and training images make classification robust to photographic variability. Many previous techniques require extensive preprocessing, lesion segmentation and extraction of domain-specific visual features before classification. By contrast, our system requires no hand-crafted features; it is trained end-to-end directly from image labels and raw pixels, with a single network for both photographic and dermoscopic images. The existing body of work uses small datasets of typically less than a thousand

## Summary

- Dataset of 129,450 clinical images
- Malignant melanoma *versus* benign nevi with an accuracy equivalent to 21 board-certified dermatologist





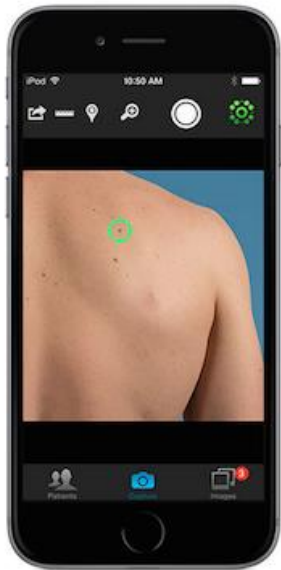
# Man against machine: diagnostic performance of a deep learning convolutional neural network for dermoscopic melanoma recognition in comparison to 58 dermatologists

H. A. Haenssle<sup>1\*,†</sup>, C. Fink<sup>1†</sup>, R. Schneiderbauer<sup>1</sup>, F. Toberer<sup>1</sup>, T. Buhl<sup>2</sup>, A. Blum<sup>3</sup>, A. Kalloo<sup>4</sup>,  
A. Ben Hadj Hassen<sup>5</sup>, L. Thomas<sup>6</sup>, A. Enk<sup>1</sup> & L. Uhlmann<sup>7</sup>

## Summary

- Dataset 100 images
- 58 dermatologists
- Algorithm outperformed most dermatologists irrespective of experience

# Digital photography



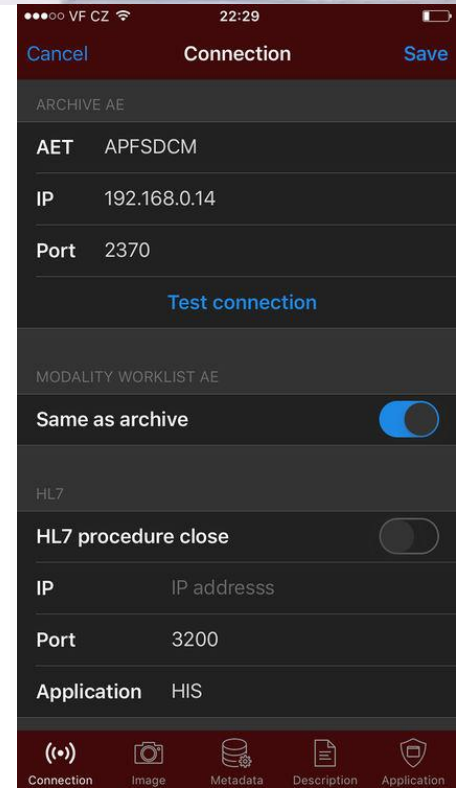
Mark image location



Image source  
<https://www.dermnetnz.org/cme/dermatoscopy-photography/dermatoscopes-and-cameras/>  
<https://vector.childrenshospital.org/2012/10/rash-conclusions-teledermatology-as-a-shared-care-model/>

Image source  
<https://www.meso.biz/products/dicom-camera/>  
<http://www.dicompass.cz/dicom-camera/>

# DICOM cameras

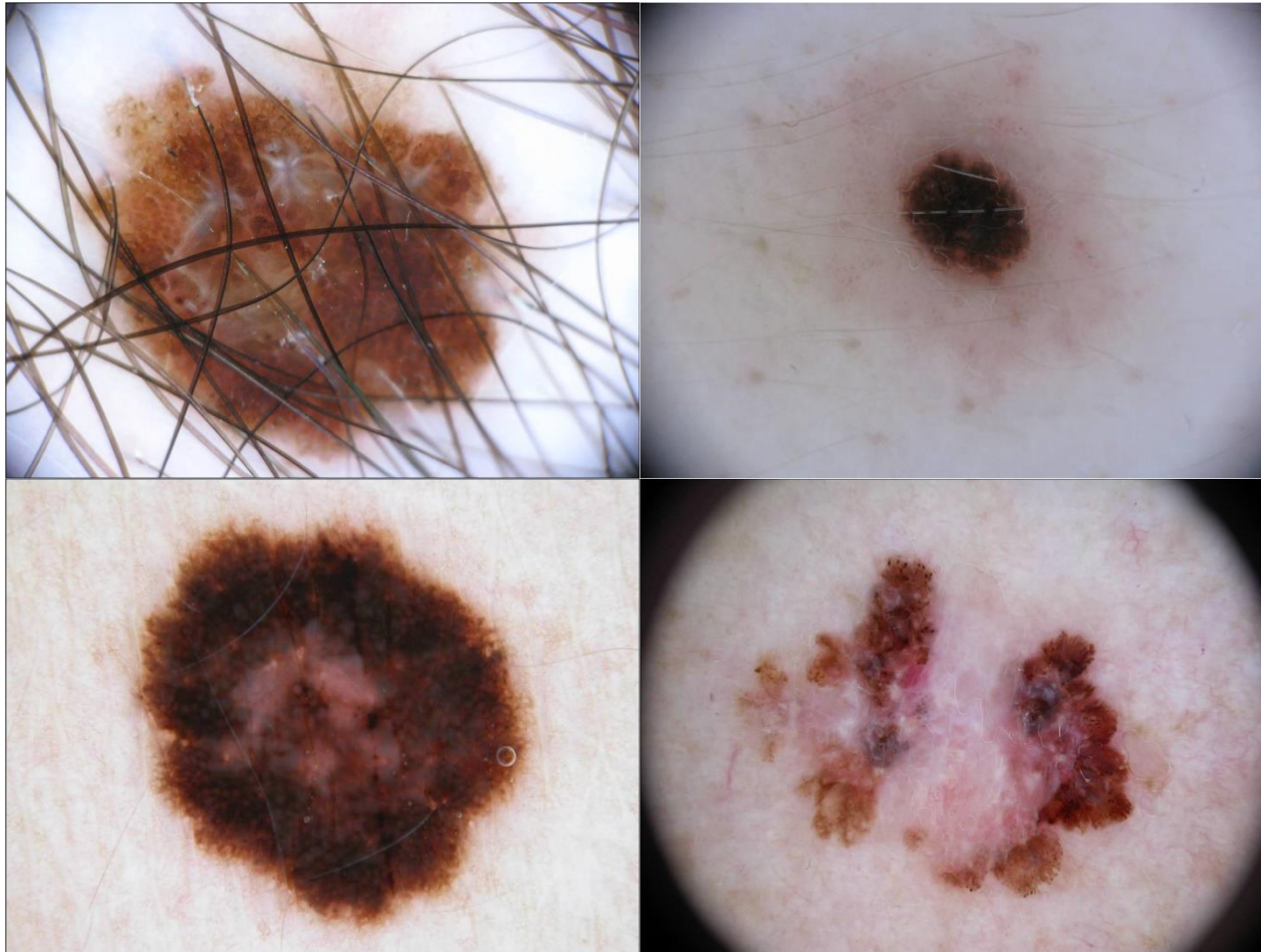


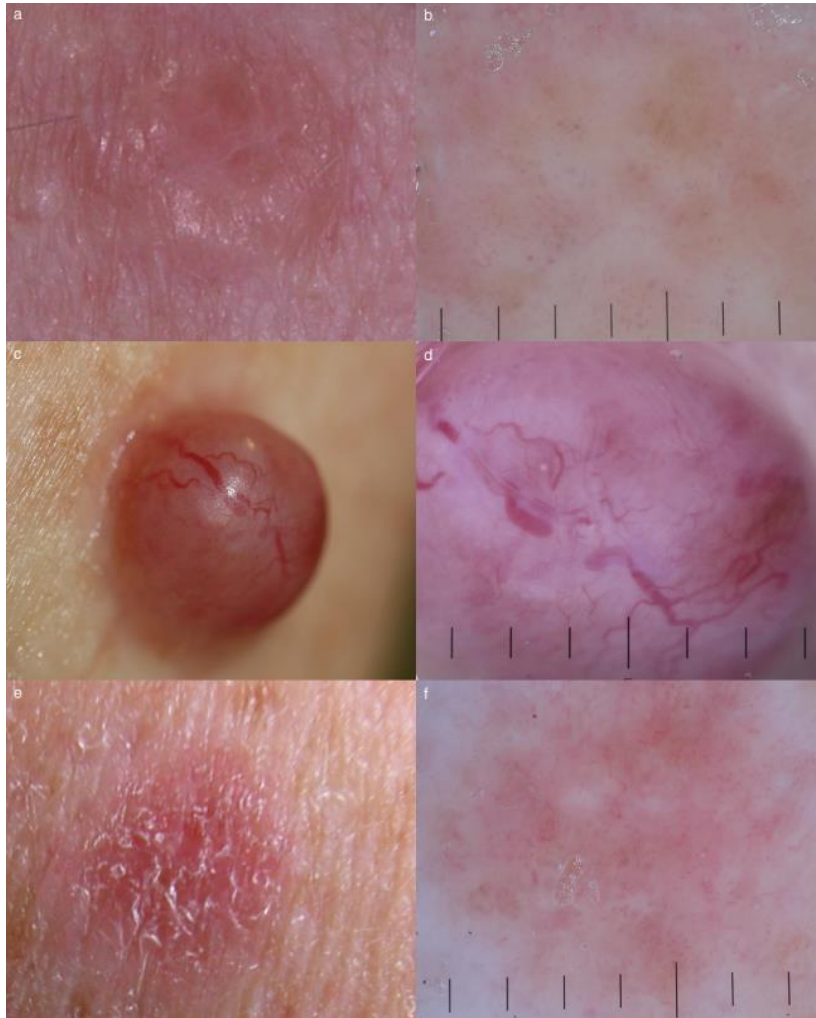
# Dermoscopy

- Dermoscopy is a non-invasive diagnostic technique that enable the visualisation of sub-surface morphological (structural) features not seen by the naked eye.



Argenziano G1, Soyer HP. Dermoscopy of pigmented skin lesions--a valuable tool for early diagnosis of melanoma. *Lancet Oncol.* 2001 Jul;2(7):443-9.



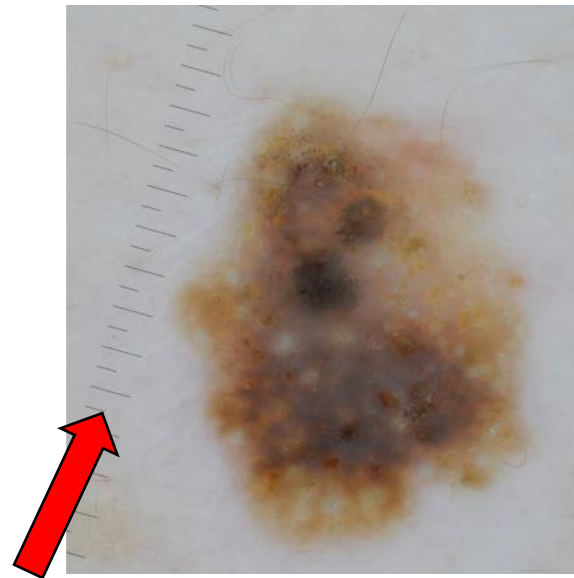


# Dermoscopy – diagnostic accuracy

- Improve diagnostic accuracy for melanoma in comparison with inspection by the unaided eye
- Dermoscopy by untrained or less experienced examiners was no better than clinical inspection without dermoscopy.
- A consensus diagnosis involving two or more experts is recommended to yield the highest possible diagnostic accuracy.

# Dermoscope - equipment

- Magnifying lens
- Light source (LED +/- polarized filters)
- Can switch between non-polarized and polarized light source
- Glass plate + scale

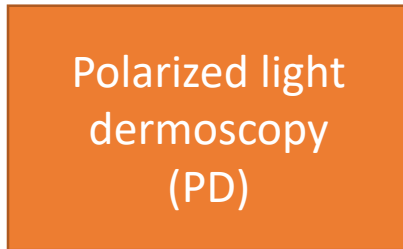




# Clinical examination

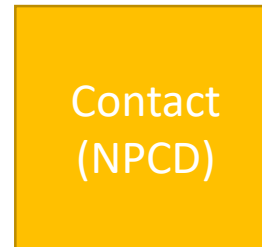
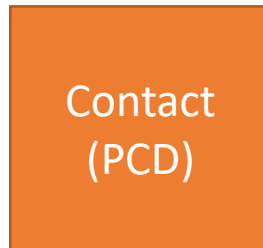


Light: LED +  
polarized filters  
to achieve **cross-  
polarization**



Light: LED

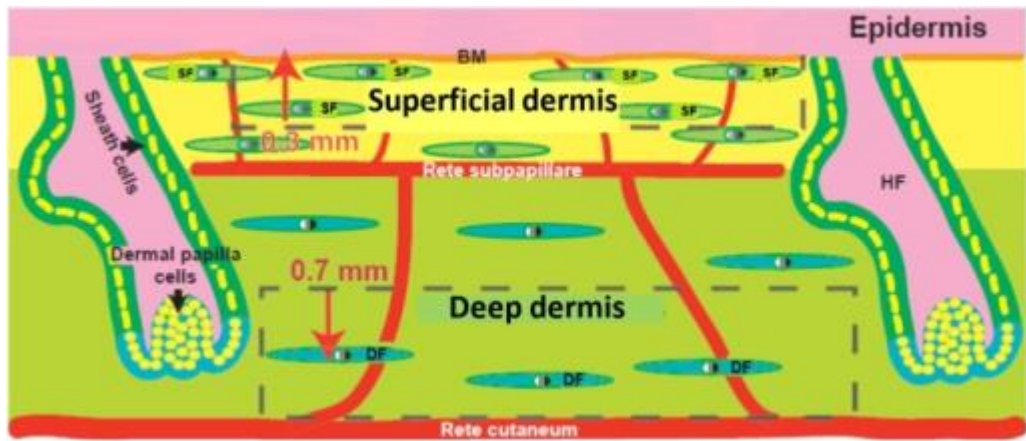
**Technique**



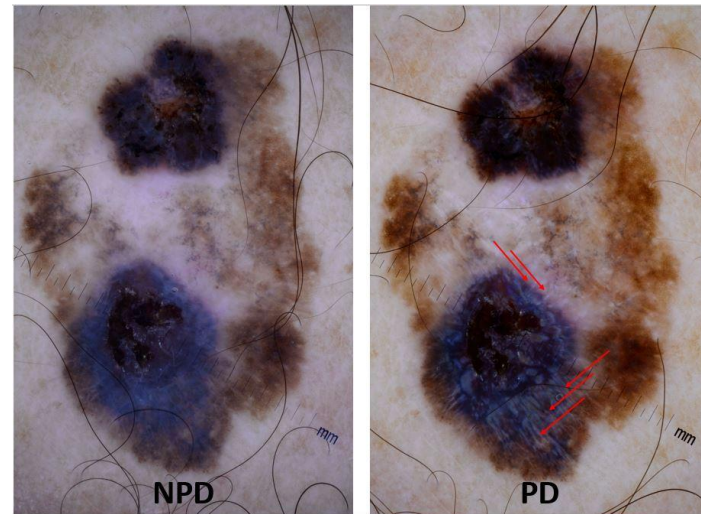
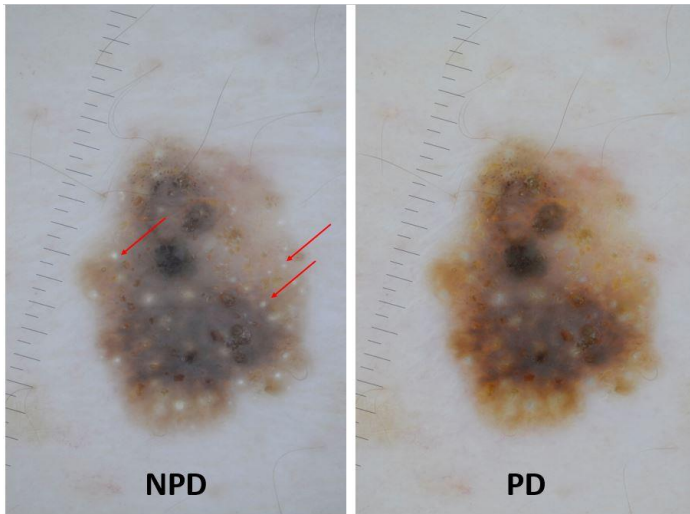
**Non-contact** does not require  
direct contact with the skin

**Direct contact** of the glass plate  
with the skin surface  
+ **liquid interface**

# Difference PD and NPD

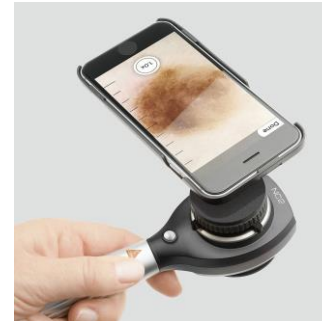


NPD – superficial epidermis to dermo-epidermal junction  
 PD – dermo-epidermal junction to superficial dermis



# Video dermoscopes

- Camera attached
- Smart-phone attached
- Handheld
- System

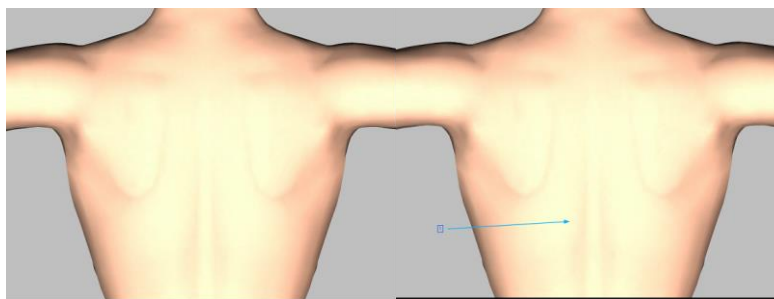


# DICOM support

Vendor	Type	IOD	Service class
3Gen	CA, SPA	*	*
Canfield / Visiomed	CA,SPA,S	US, SC	Storage SCU
Derma	H,S	SC	
Fotofinder	SPA, H, S	Multi-frame True Color SC	
Heine	CA	*	*
Pixience	S	SC	

CA = camera attached; SPA=smart phone attached, H=handheld, S=System

\* DICOM support if coupled with a DICOM camera software



# Total Body Photography



**3D**

Canfield Scientific  
<https://www.canfieldsci.com/imaging-systems/vectra-wb360-imaging-system>



**2D**

Fotofinder  
<https://www.fotofinder.de/en/>

# Total body photography?

## Advantages

- Unimaged melanoma
- “Ugly duckling”

## Disadvantages

- Genital, acral, scalp and body folds
- Maturity
- Cost
- Resolution +/- dermoscopy

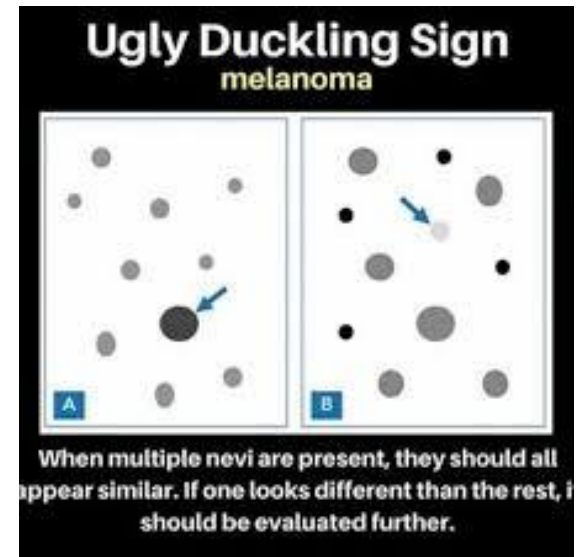
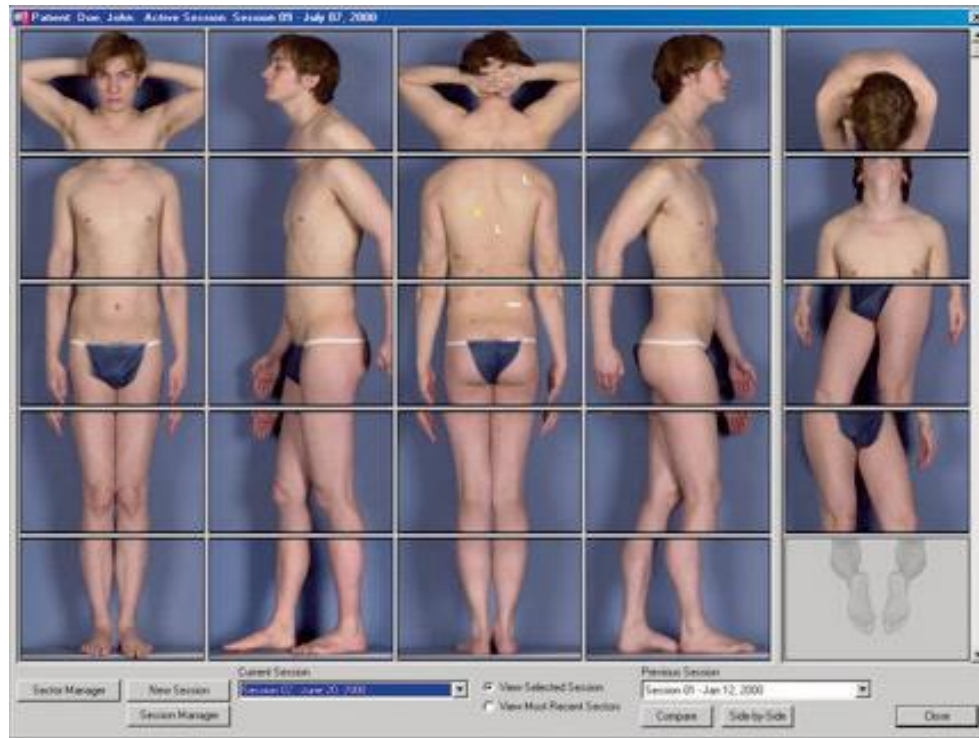


Image source  
<https://www.skincancer.org/skin-cancer-information/melanoma/melanoma-warning-signs-and-images/the-ugly-duckling-sign>

# 2D versus 3D

- Examination time
  - 25 exposures 2D *versus* 1 (simultaneous) exposure 3D
- Orthogonal imaging
  - Skin lesions on curved surfaces

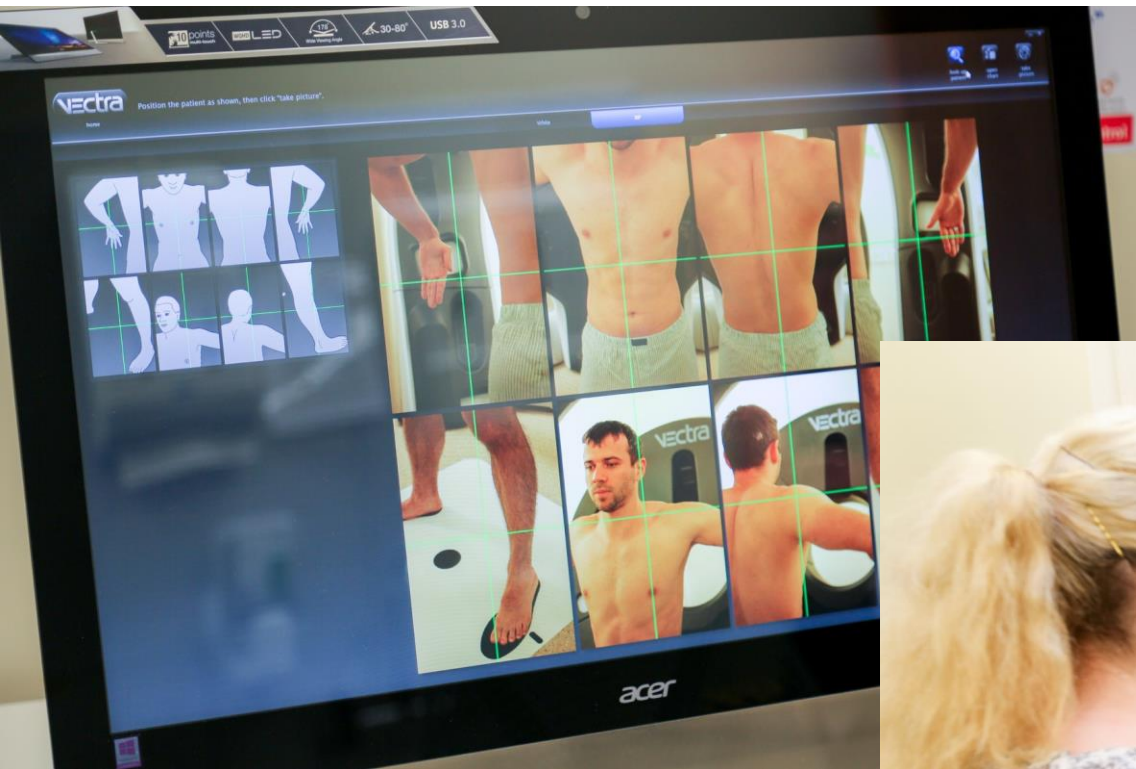
# 2D Total Body Photography

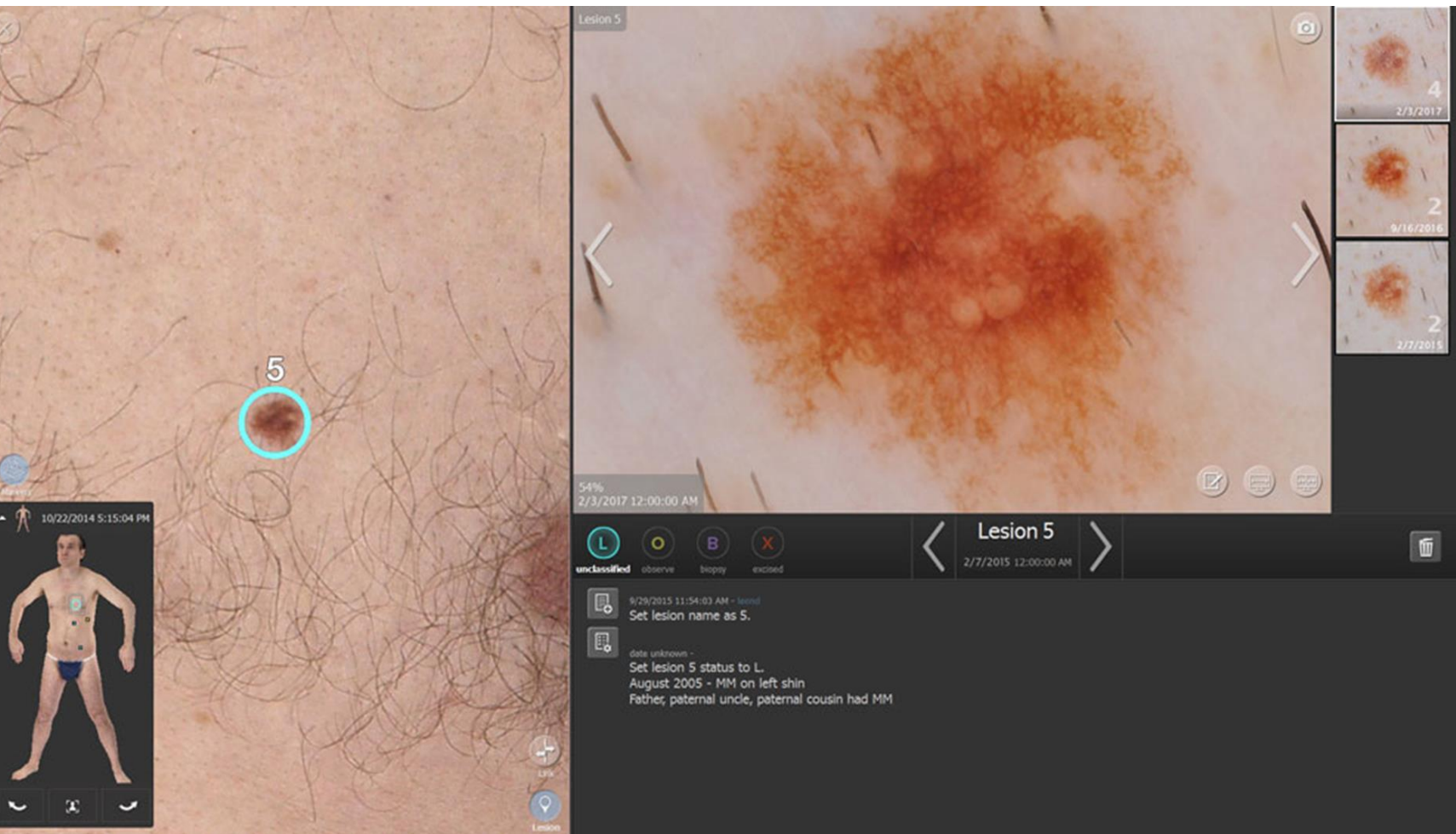




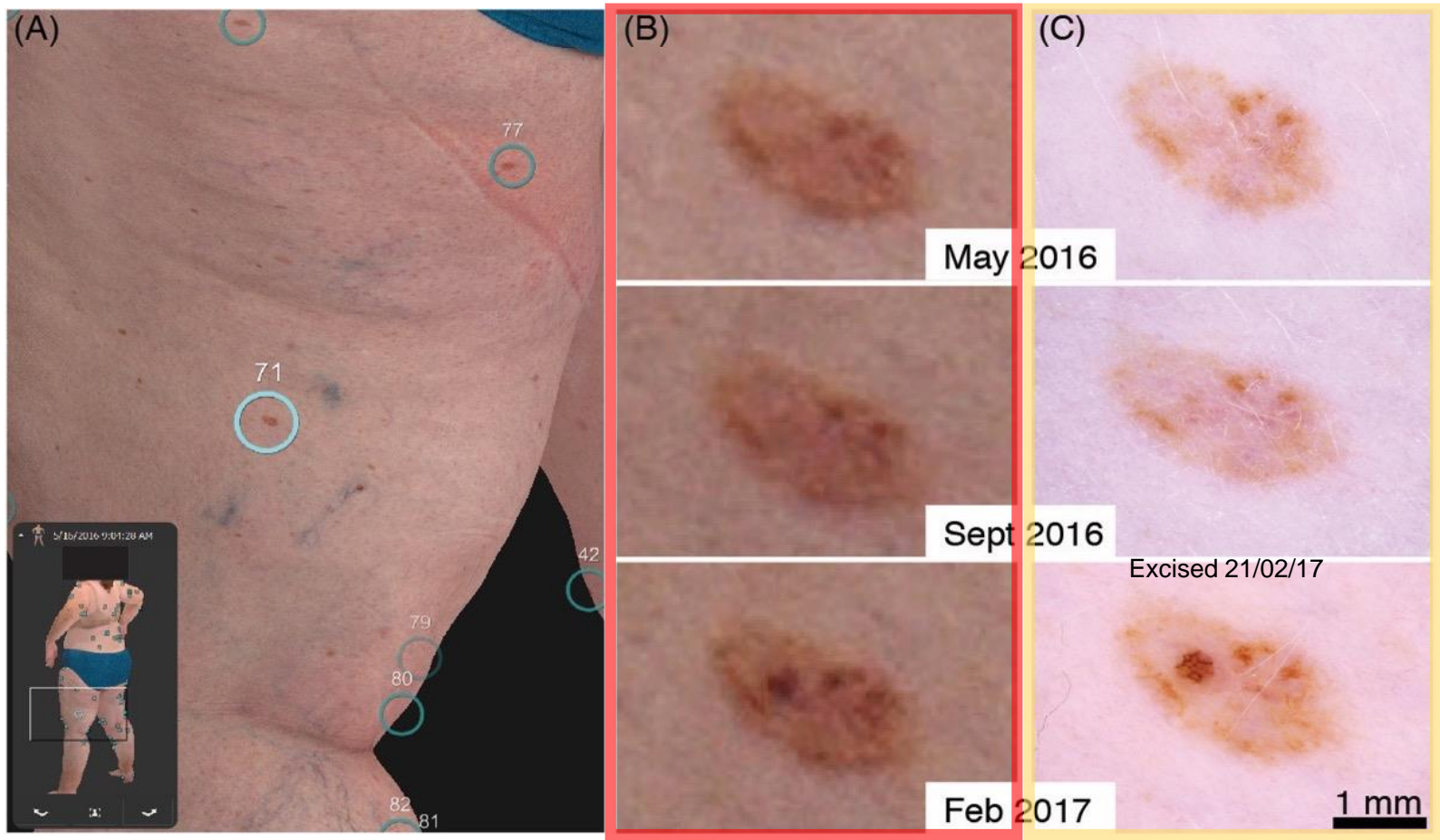


















Canfield Scientific  
<https://www.canfieldsci.com/imaging-systems/vectra-wb360-imaging-system>



TBP

Dermoscopy

	Benign		Malignant	
Symmetrical		<b>A</b> Asymmetry		Assymetrical (the two sides do no match)
Borders are even		<b>B</b> Border		Borders are uneven
One color		<b>C</b> Color		Two or more colors
Smaller than 6 mm (1/4 inch)		<b>D</b> Diameter		Larger than 6 mm (1/4 inch)
Ordinary mole		<b>E</b> Evolution		Changing in size, shape, color, or another trait



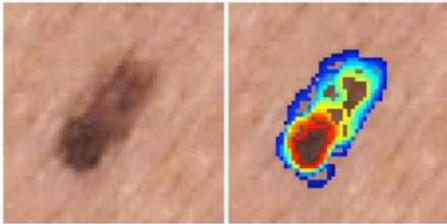
Toggle Details

## Lesion Data

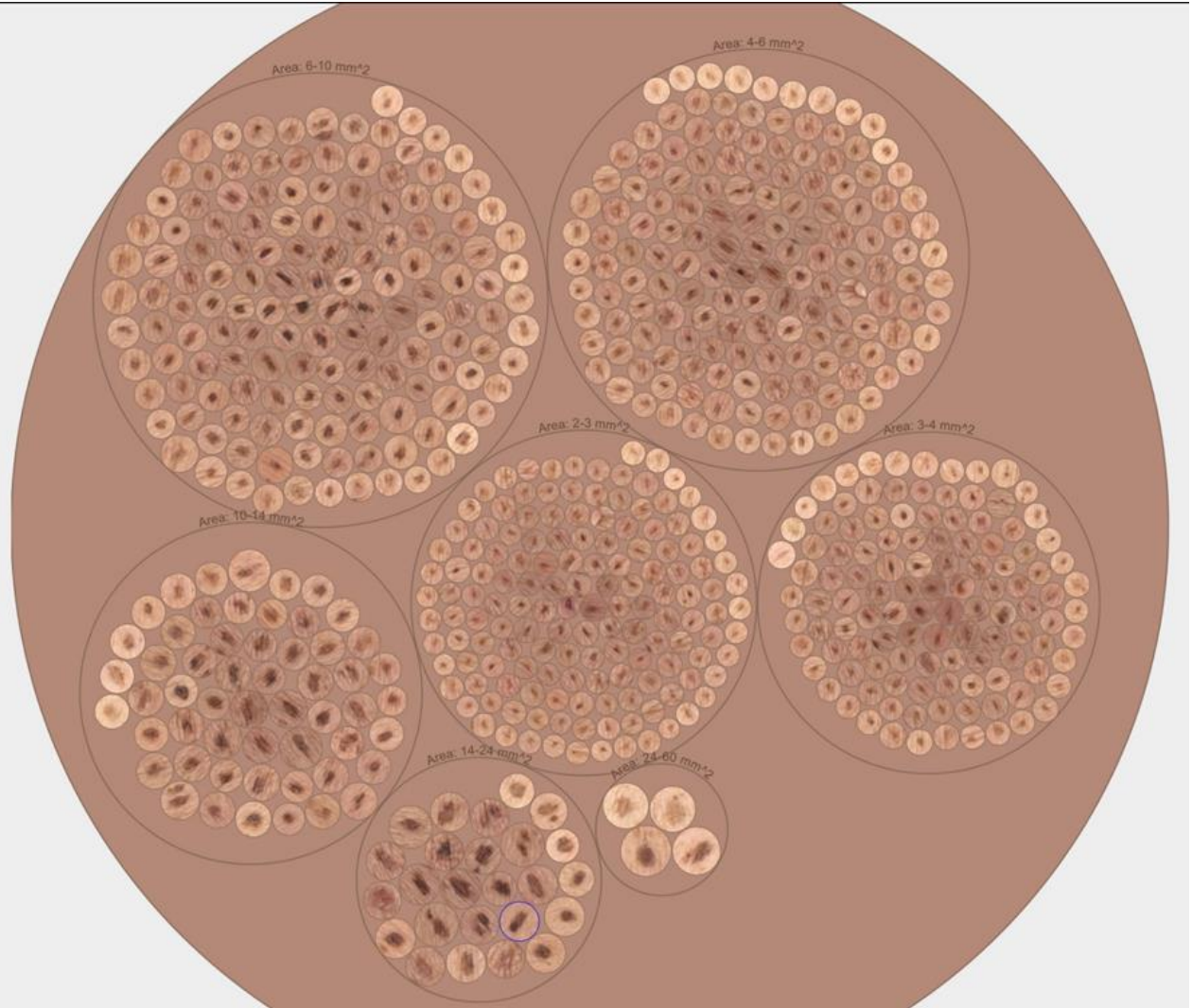
Group By:  ▼  
Sort By:  ▼  
Location:  ▼

Lesion #24-126

< Prev Next >



Lesion ID #	24-126
Location	torso-back
Area	16.02 mm <sup>2</sup>
Major Axis Length	6.41 mm
Minor Axis Length	3.30 mm
Perimeter	17.93 mm
Eccentricity	0.86
Levels	18.00
L*	46.07
a*	19.61
b*	25.72
Delta L*	21.98
Delta E*	24.48





# Reflectance Confocal Microscopy

- Non-invasive, in vivo imaging of the cellular and tissue architecture (resolution equivalent to histology)
- Images up to 200  $\mu\text{m}$  (papillary dermis)

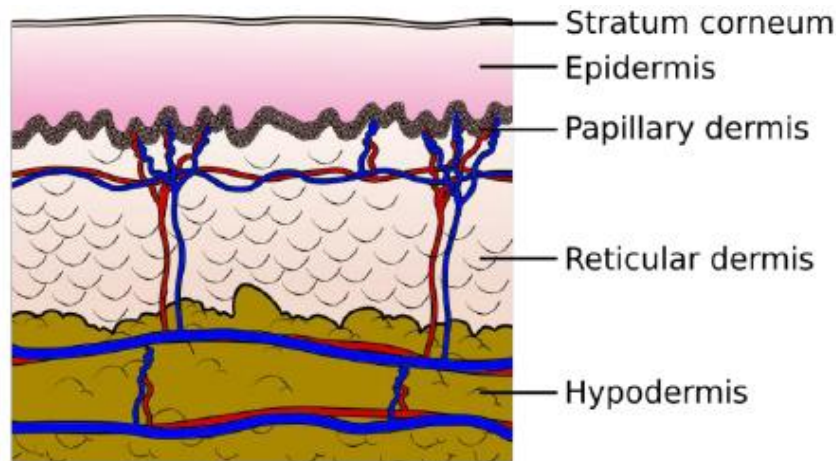
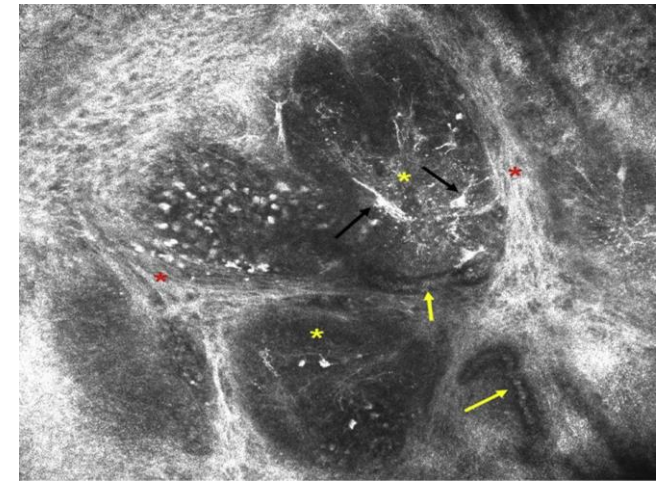


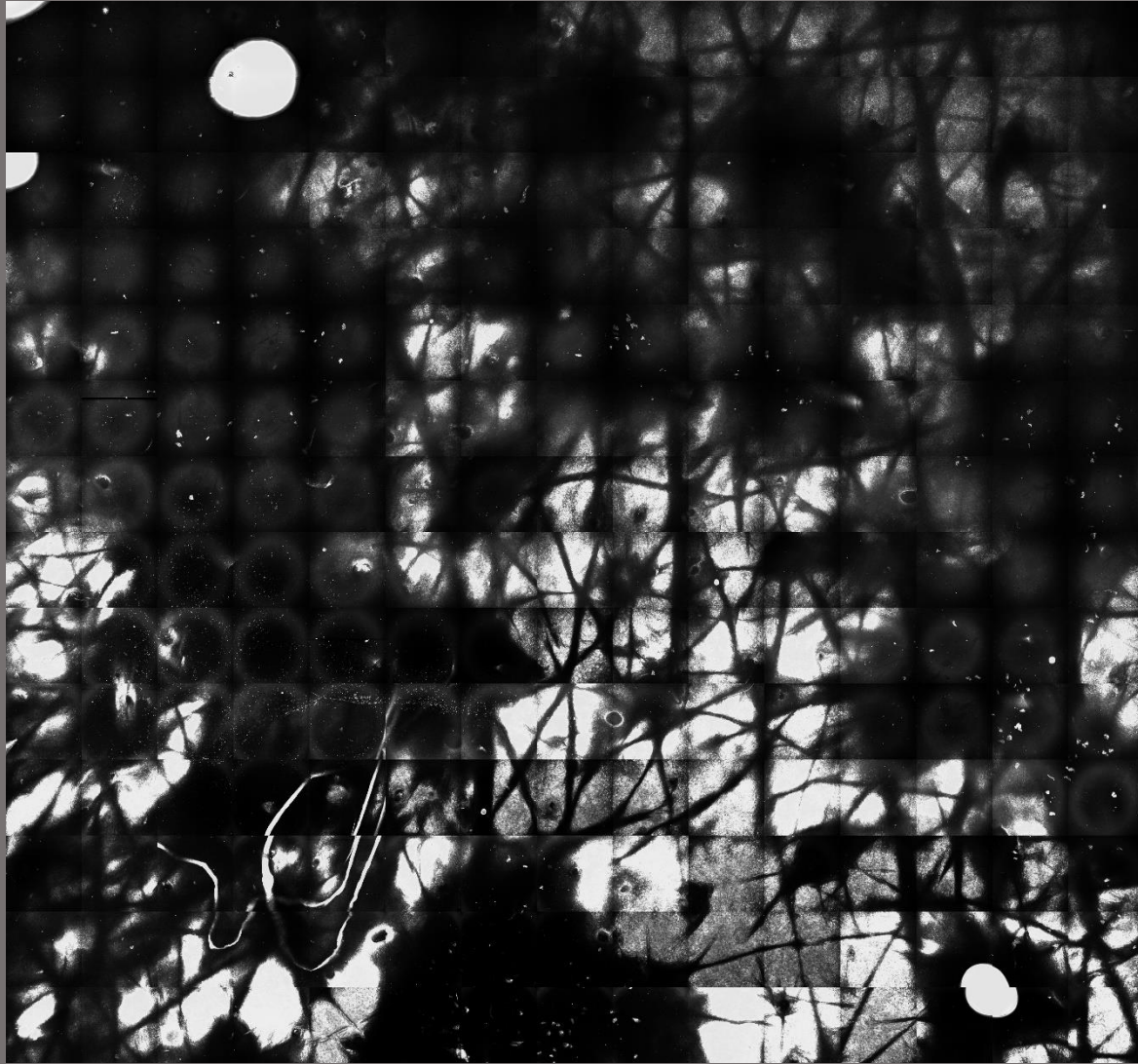


Image source <http://www.caliberid.com/>

# Reflectance Confocal Microscopy

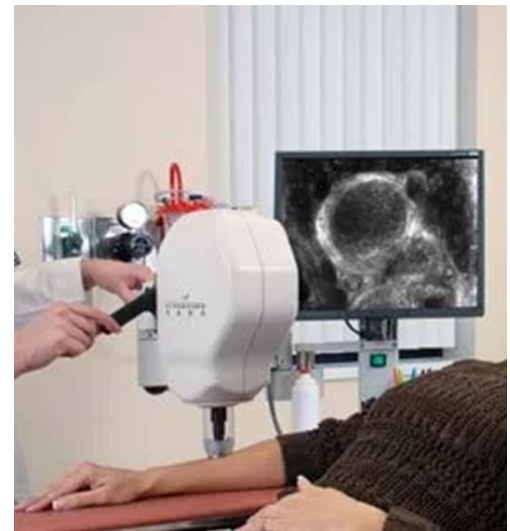


<http://www.caliberid.com/vivascope1500-Overview.html>



# Reflectance Confocal Microscopy

- Dramatically increase the accuracy of skin cancer diagnosis [9]
- “Virtual biopsy”
  - Reduce the number of biopsy
- Delineation of surgical margins



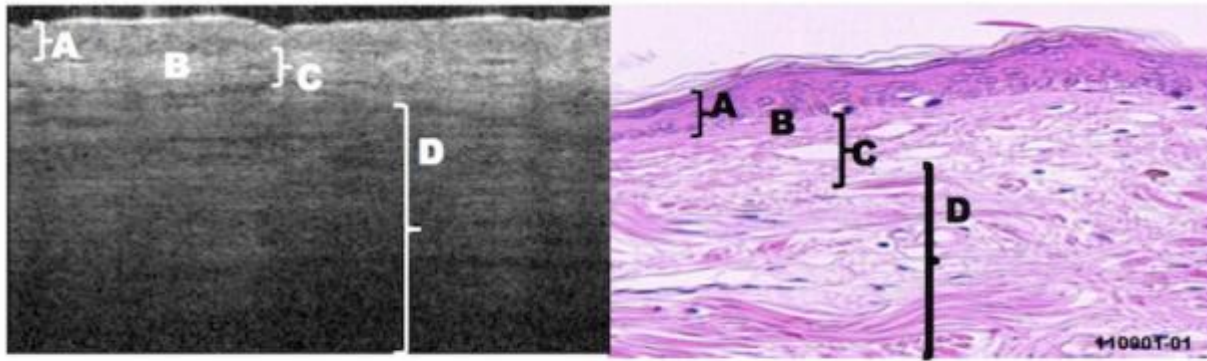
# Vivanet <sup>TM</sup> and Vivastore <sup>TM</sup>

- DICOM cloud storage + scheduling system
- Facilitates teledermatology
  - Imaging expertise
  - Second opinion
- Facilitates international community of practice

# Optical Coherence Tomography

- Mainly used in ophthalmology
- Non-invasive, in vivo imaging skin architecture
- Depth up 1.5mm
  - Greater depth but lower resolution than RCM
- Inflammatory disease + sometimes skin cancer

# Optical Coherence Tomography

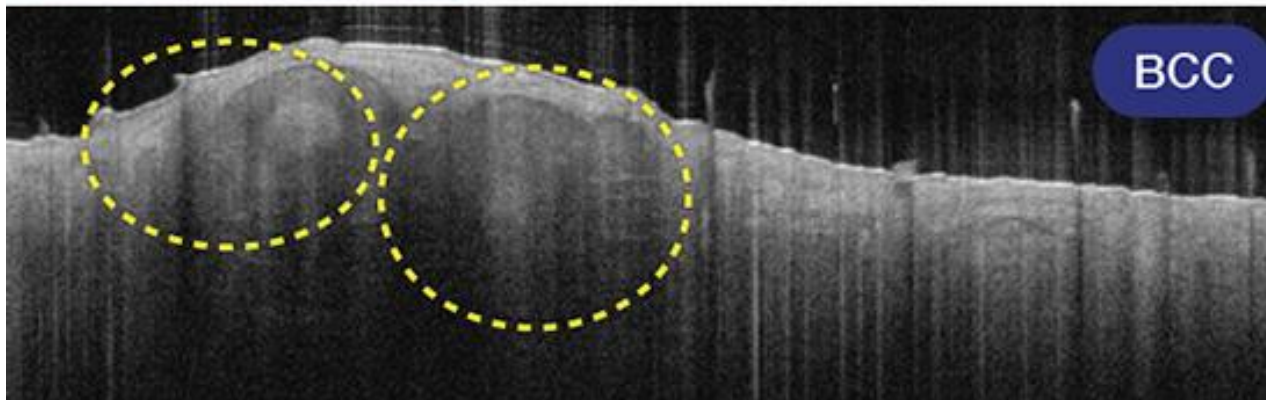
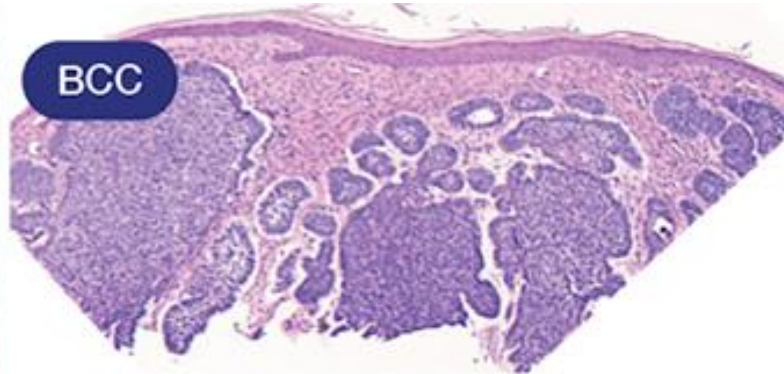


- A: Epidermis
- B: Dermal Epidermal Junction (DEJ)
- C: Papillary Dermis
- D: Reticular Dermis



Image source  
<https://www.dermnetnz.org/>





The resolution enables the visualization of architectural changes, but not of single cells.

# Summary

- Increasing evidentiary skin imaging
- Increasing diagnostic skin imaging
- DICOM WG 19 dormant
- No dermatology specific DICOM IODs
  - Visible Light, Ultrasound, OCT
- Many vendors have some level of DICOM (inconsistent)
- Majority of dermatologists do not use DICOM

# Thank you

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