



# Common skin tumors

SOMSAK TANRATTANAKORN

28/02/2018

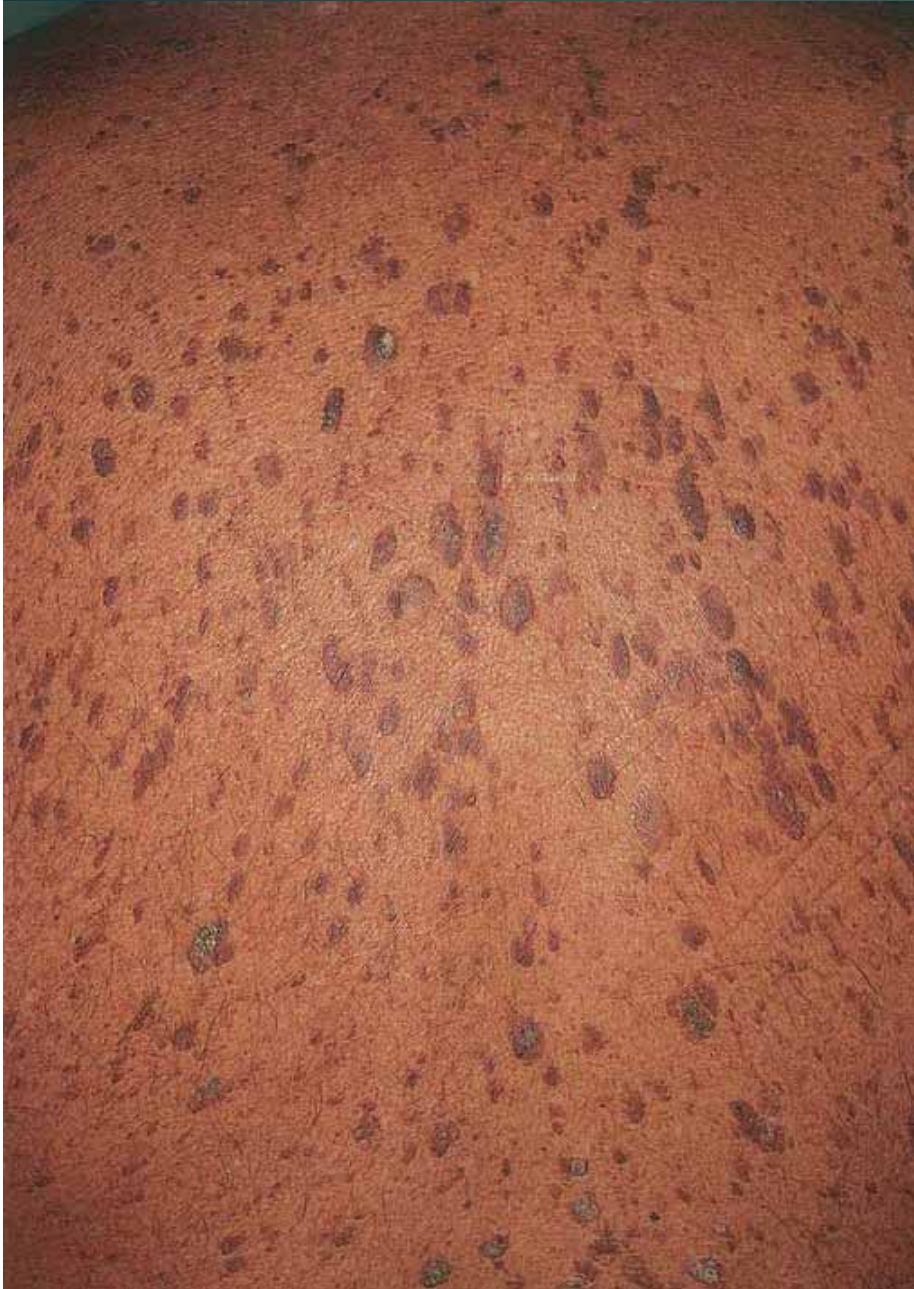
# Topic

- ▶ Benign epidermal tumors
- ▶ Skin cyst and adnexal neoplasms
- ▶ Other common skin tumor
- ▶ Common skin malignancy

# Benign Epidermal Tumors

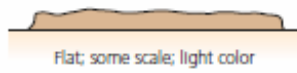
- ▶ Seborrheic keratosis
  - ▶ Dermatosi papulosa nigra
  - ▶ Stucco keratosis
- ▶ Inverted follicular keratosis
- ▶ Acrokeratosis verruciformis
- ▶ Clear cell acanthoma
- ▶ Large cell acanthoma
- ▶ Porokeratosis
- ▶ Epidermal nevus
  - ▶ Inflammatory linear verrucous epidermal nevus
- ▶ Nevus comedonicus
- ▶ Epidermolytic acanthoma
- ▶ Flegel's disease
- ▶ Cutaneous horn
- ▶ Lichenoid keratosis
- ▶ Acanthosis nigricans
- ▶ Confluent and reticulated papillomatosis
- ▶ Warty dyskeratoma

# Seborrheic keratoses



- ▶ very common brown macules, papules, plaques, or polypoid lesions
- ▶ over 40 y.
- ▶ increase number with age
- ▶ verrucous or 'stuck-on' the skin
- ▶ predilection for face, neck, and trunk
- ▶ occur anywhere except mucous membranes, palms, or soles
- ▶ sign of Leser-Trélat





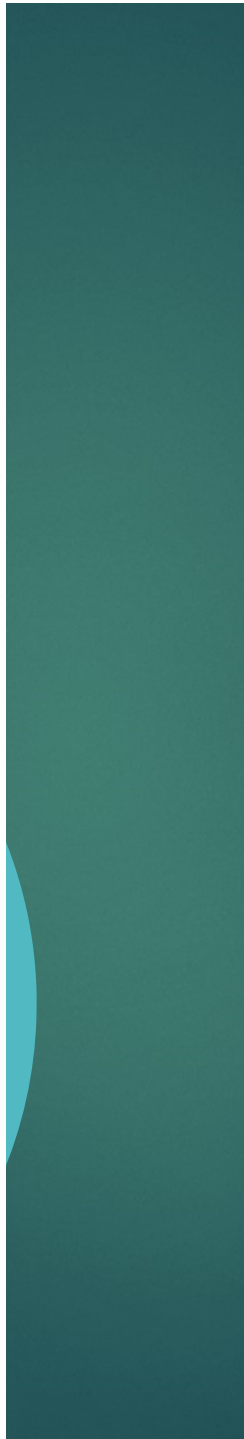
Flat; some scale; light color



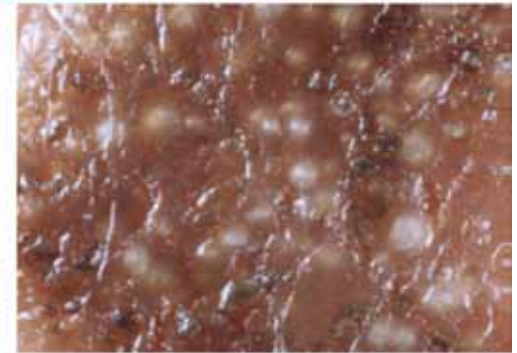
Height increase; lesion appears "stuck on" to surface; color darkens



Deep surface cracks appear; keratin can be peeled off; brown or black



Contains horn cysts, which are white or black



Height increases; horn cysts become more numerous



Smooth, dome-shaped papule; horn cysts project from surface



# Clinicopathologic Variants

- ▶ Common Seborrheic Keratosis
- ▶ Dermatosi Papulosa Nigra
- ▶ Skin Tags
- ▶ Irritated Seborrheic Keratosis
- ▶ Stucco Keratosis
- ▶ Reticulated Seborrheic Keratosis
- ▶ Clonal Seborrheic Keratoses
- ▶ Seborrheic Keratosis With Squamous Atypia
- ▶ Melanoacanthoma
- ▶ Leser-Trelat sign





Seborrheic keratosis



Dermatosis papulosa nigra









# Skin tags





# Irritated Seborrheic Keratosis





## Stucco keratoses

- ▶ Multiple gray-white keratotic papules on ankle and dorsal foot.



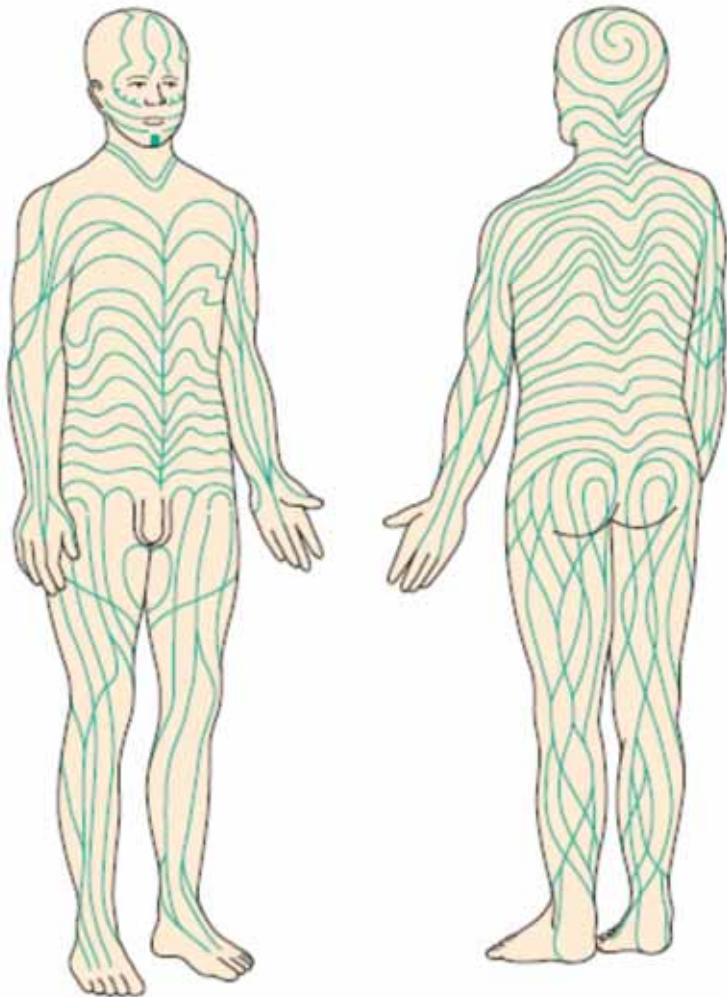
# Epidermal nevi

- ▶ benign hamartoma of epidermis and papillary dermis
- ▶ onset usually within the first year of life
- ▶ asymptomatic well-circumscribed, hyperpigmented, papillomatous papules or plaques in a linear array along Blaschko's lines
- ▶ patients with epidermal nevus syndrome have associated abnormalities, in particular musculoskeletal and neurologic





# Blaschko's lines









## Inflammatory linear verrucous epidermal nevus (ILVEN).

- Linear, psoriasiform papules or plaques
- usually on one extremity
- 75% appear before 5 years
- 4 times more common in girls
- usually persists for years despite attempts at treatment



## Nevus comedonicus

- ▶ usually present by age of 10
- ▶ closely arranged, grouped, often linear, slightly elevated papules
- ▶ center keratinous plugs



# Cutaneous horn

- ▶ firm, white to yellow, conical, markedly hyperkeratotic papule, plaque or nodule
- ▶ Most common in sun-exposed areas and arising from a hyperkeratotic actinic keratosis
- ▶ SCC is present at the base of lesion in up to 20% of patients



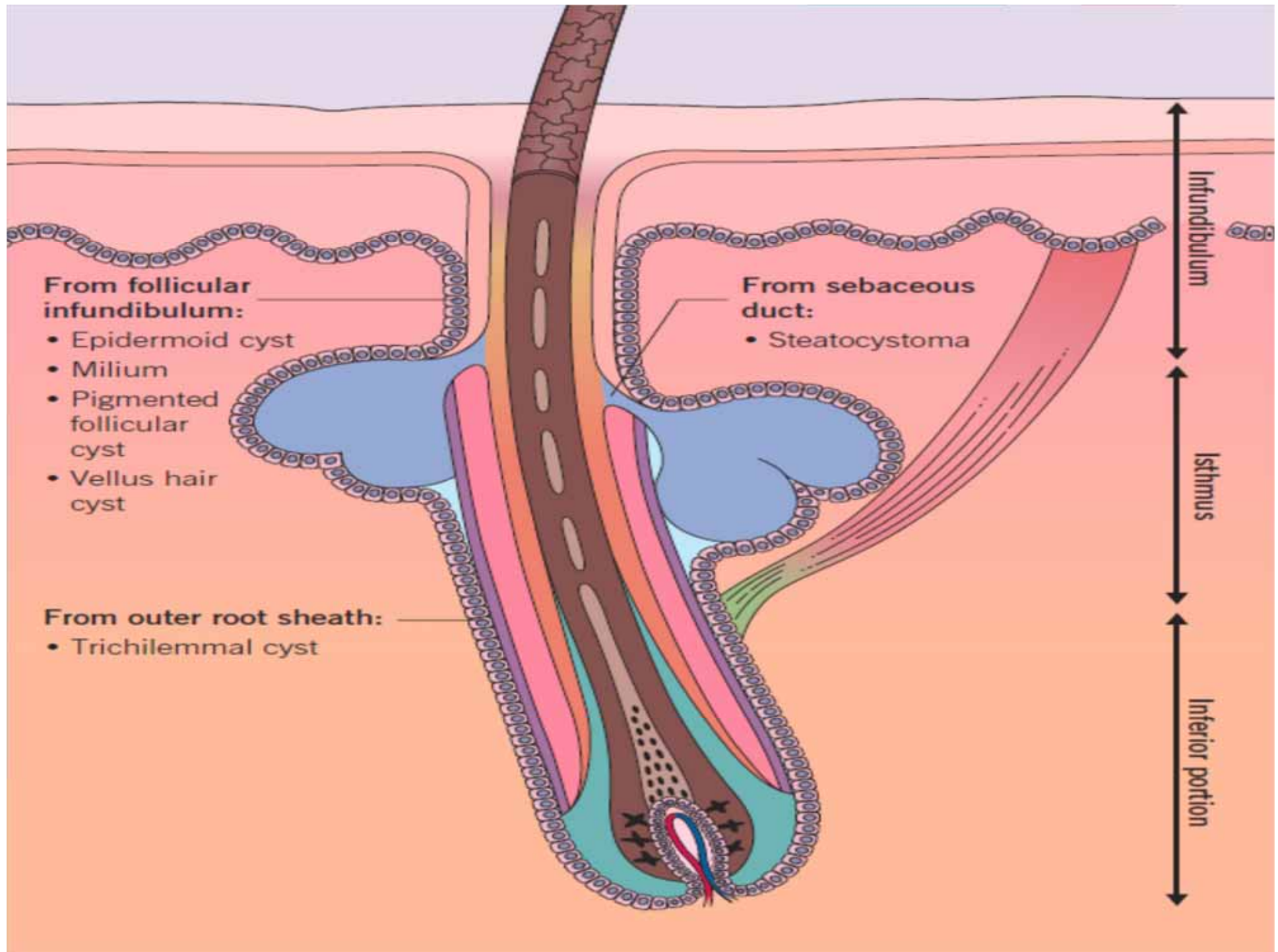
# Acanthosis nigricans





# Cutaneous cysts lining

- ▶ stratified squamous epithelium
  - ▶ Epidermoid cyst
  - ▶ Miliun
  - ▶ Trichilemmal cyst
  - ▶ Vellus hair cyst
  - ▶ Steatocystoma
  - ▶ Ear Pit / Preauricular cyst
- ▶ non-stratified squamous epithelium
  - ▶ Hidrocystoma
- ▶ no epithelium
  - ▶ Mucocele
  - ▶ Digital mucous cyst
  - ▶ Ganglion





# Epidermoid (Epithelial) cyst

- ▶ sebaceous cyst is a misnomer
- ▶ most common cutaneous cysts
- ▶ occur anywhere but common on face and upper trunk
- ▶ young and middle-aged adults
- ▶ dermal nodules with central punctum
- ▶ Multiple cysts may associated with Gardner's syndrome (familial adenomatous polyposis)





Epidermoid (Epithelial) cyst



# Inflamed epidermal cyst



# Milia



- ▶ small epidermoid cysts
- ▶ 1-2 mm white to yellow subepidermal papules
- ▶ 40-50% of infants will have milia on face, will resolve spontaneously in the first 1 month
- ▶ may secondary from blistering processes or superficial ulceration from trauma or resurfacing, topical corticosteroid-induced atrophy



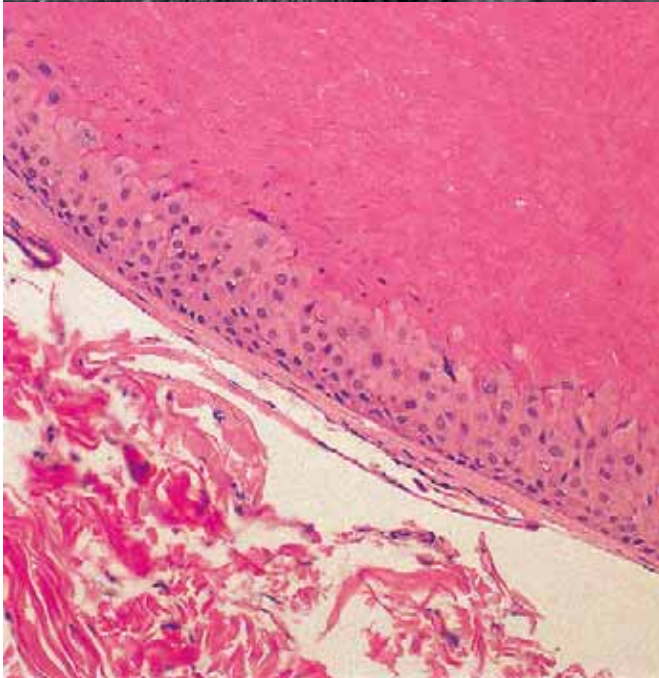
Milia





# Trichilemmal (pilar) cyst

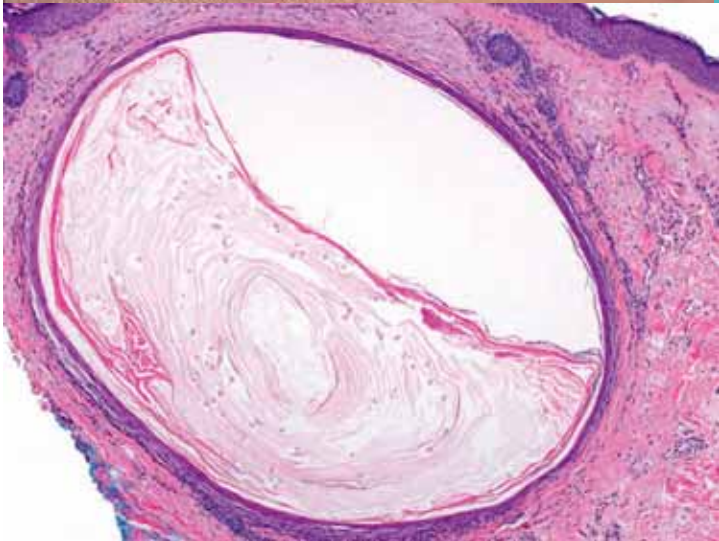
- ▶ 95% located on scalp
- ▶ Clinical = epidermal cyst





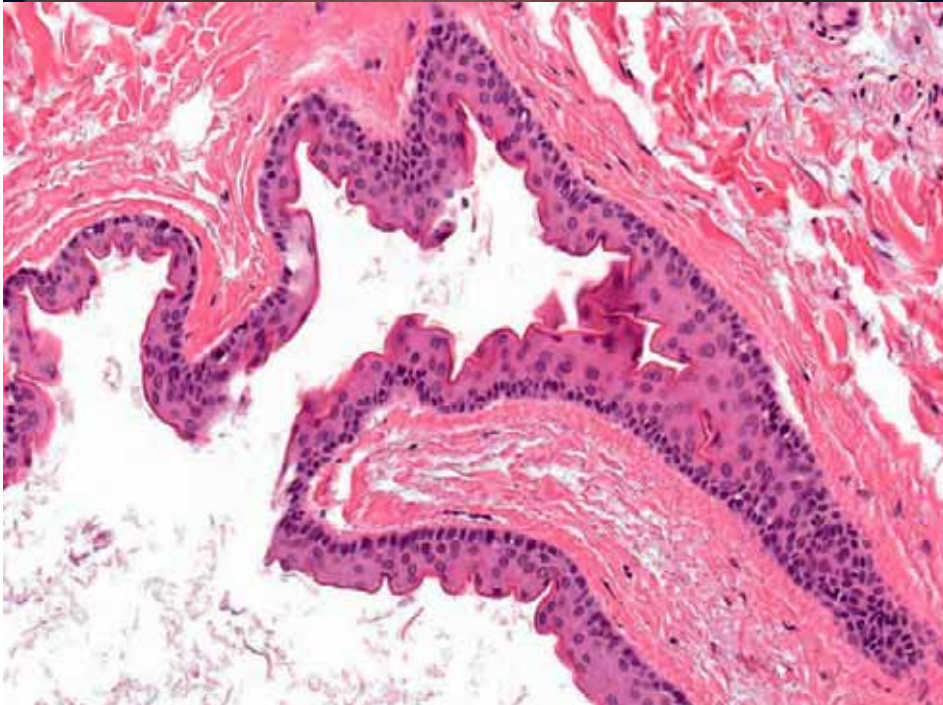
# Vellus Hair Cysts

- ▶ numerous tiny dome-shaped papules, ranging from skin-colored to darkly pigmented on trunk
- ▶ most commonly located on the trunk
- ▶ multiple may be inherited in AD



# Steatocystoma simplex/multiplex

- ▶ *sebaceous cyst*
- ▶ asymptomatic cysts in the dermis that drain oily fluid if punctured
- ▶ persist indefinitely
- ▶ chest, axillae, and groin
- ▶ Multiplex = autosomal dominant





# Steatocystoma multiplex



# Ear Pit / Preauricular cyst



- ▶ congenital defects
- ▶ 0.5–1% of normal population
- ▶ may be transmitted in AD
- ▶ usually unilateral and right-sided

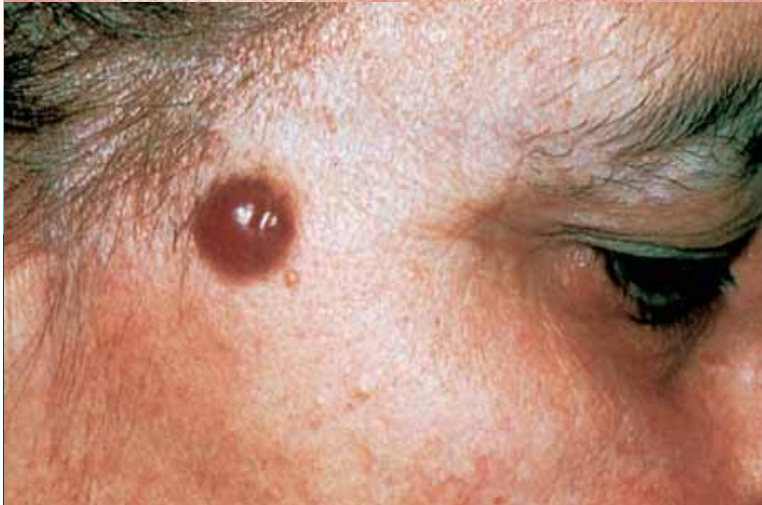


# Cutaneous cysts lining

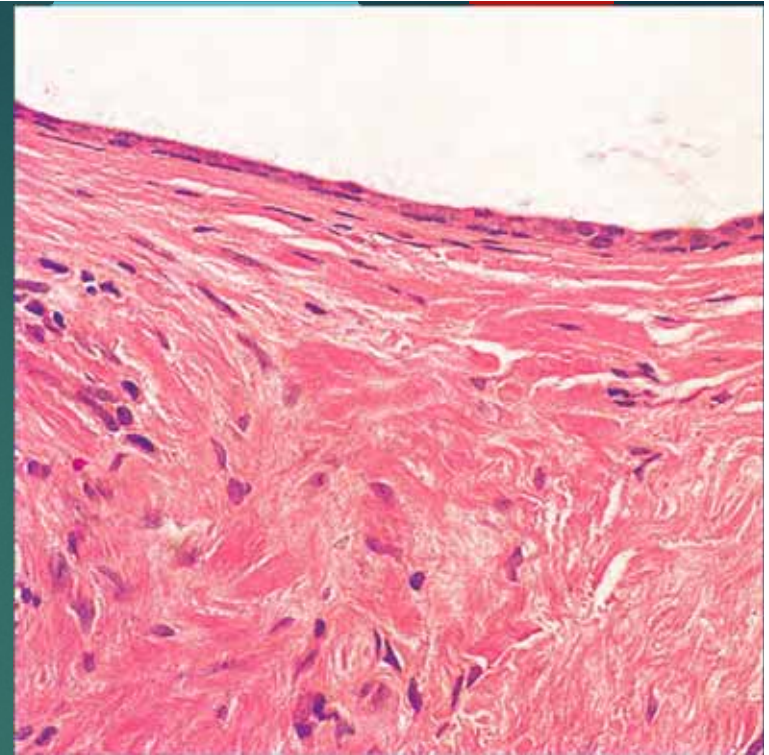
- ▶ stratified squamous epithelium
  - ▶ Epidermoid cyst
  - ▶ Milium
  - ▶ Trichilemmal cyst
  - ▶ Vellus hair cyst
  - ▶ Steatocystoma
  - ▶ Ear Pit / Preauricular cyst
- ▶ non-stratified squamous epithelium
  - ▶ Hidrocystoma
- ▶ no epithelium
  - ▶ Mucocele
  - ▶ Digital mucous cyst
  - ▶ Ganglion

# Apocrine hidrocystoma

- ▶ 1-3 mm (face, scalp)
- ▶ translucent, skin-colored to bluish cysts on face
- ▶ adenomas of apocrine sweat gland coils







## Eccrine hidrocystoma

- ▶ solitary or multiple\*
- ▶ can enlarge with heat exposure or during the summer and regress with cooler temperatures
- ▶ cystic dilation of eccrine ducts due to retention of eccrine secretions

# Cutaneous cysts lining

- ▶ stratified squamous epithelium

- ▶ Epidermoid cyst
- ▶ Miliun
- ▶ Trichilemmal cyst
- ▶ Vellus hair cyst
- ▶ Steatocystoma
- ▶ Ear Pit / Preauricular cyst

- ▶ non-stratified squamous epithelium

- ▶ Hidrocystoma (apocrine/eccrine)

- ▶ no epithelium

- ▶ Mucocele
- ▶ Digital mucous cyst
- ▶ Pseudocyst of the auricle
- ▶ Ganglion



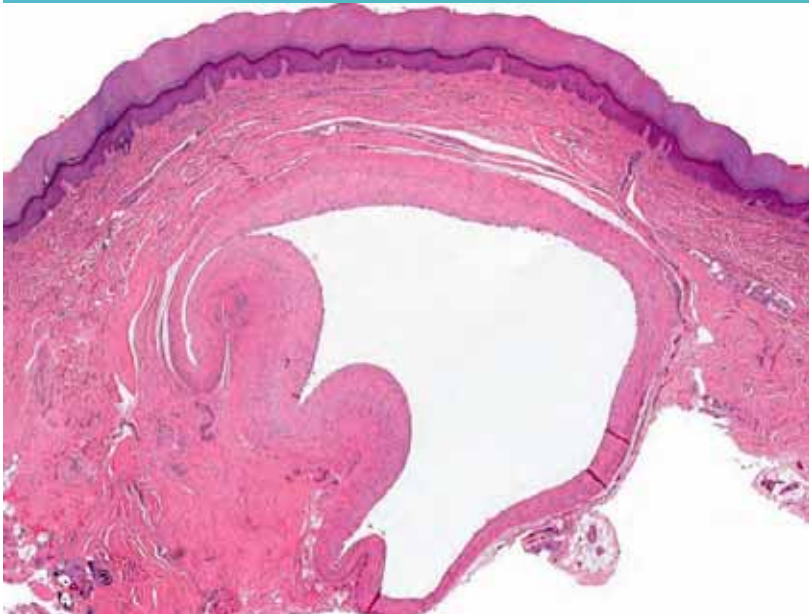
# Mucocele



- ▶ Common - lower labial mucosa
- ▶ dome-shaped, bluish, translucent papules or nodules
- ▶ disruption of ducts of minor salivary glands

# Digital mucous cyst

- ▶ dorsal surface of finger distal phalanx
- ▶ depressed nail deformity







## Pseudocyst of the auricle

- ▶ scaphoid fossa of ear in middle-aged men
- ▶ usually unilateral
- ▶ painless swelling
- ▶ ? chronic trauma

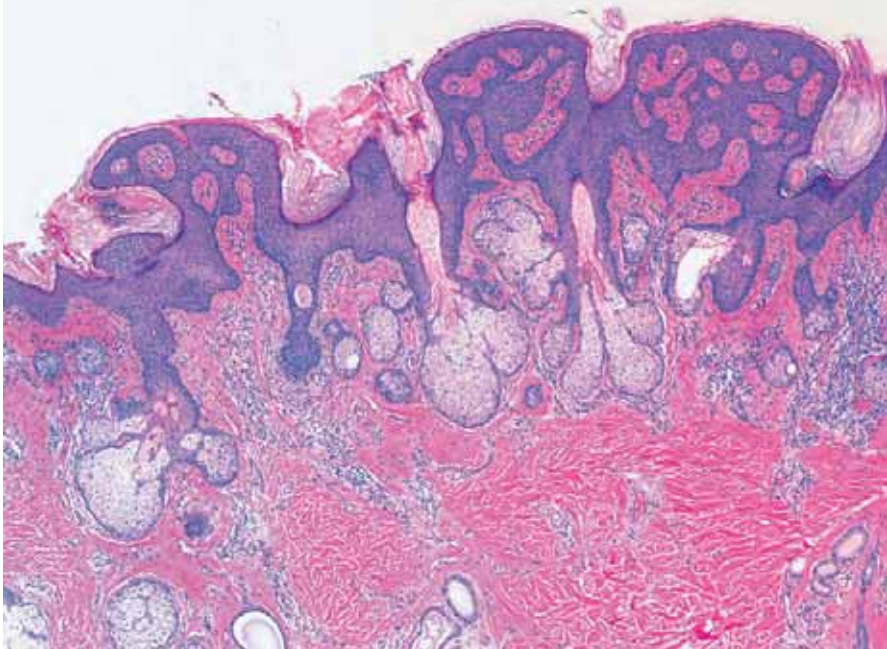
# Adnexal Neoplasms

- ▶ Follicular lineage
  - ▶ Nevus sebaceus
  - ▶ Trichoepithelioma
  - ▶ Trichofolliculoma
  - ▶ Pilomatricoma
- ▶ sebaceous differentiation
  - ▶ Sebaceous gland hyperplasia
  - ▶ Fordyce's disease
- ▶ apocrine differentiation
  - ▶ Syringoma
- ▶ eccrine differentiation
  - ▶ Eccrine nevus



# Nevus sebaceus

- ▶ papillomatous yellow-orange linear plaque on the scalp or face
- ▶ associated with alopecia
- ▶ sebaceous glands are most prominent during early infancy and post-puberty
- ▶ scalp or face



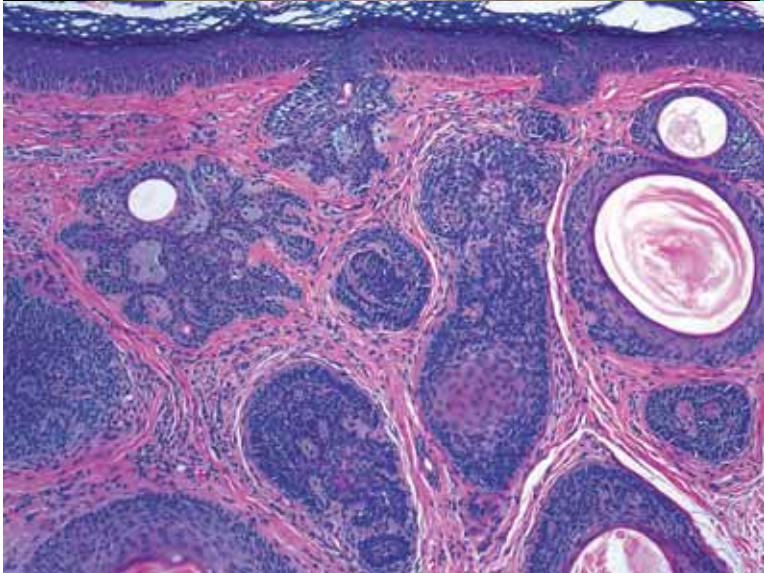
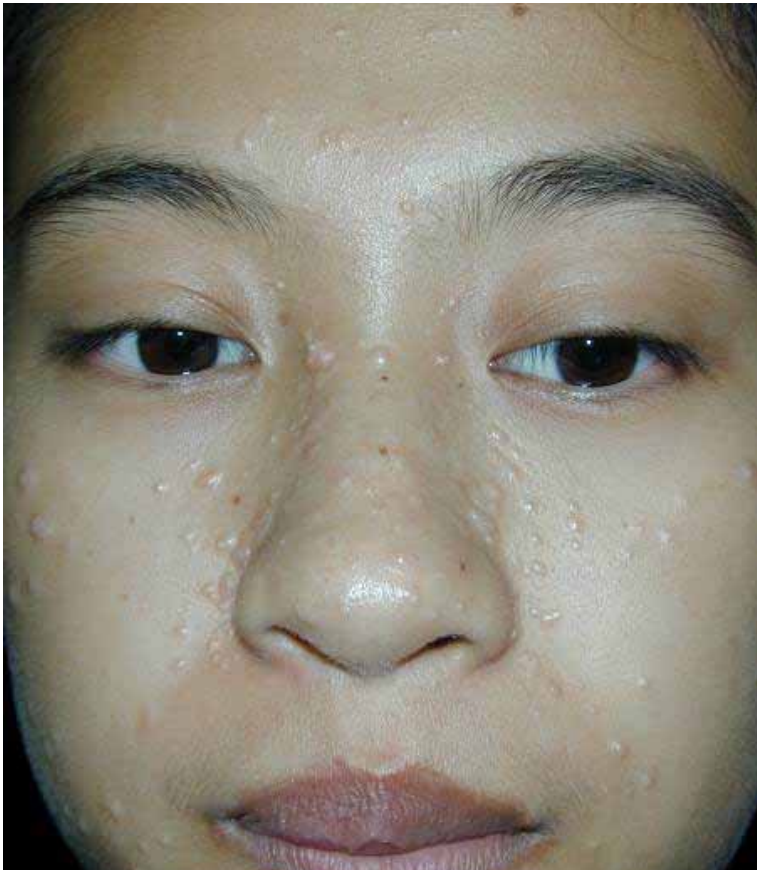






# Trichoepithelioma

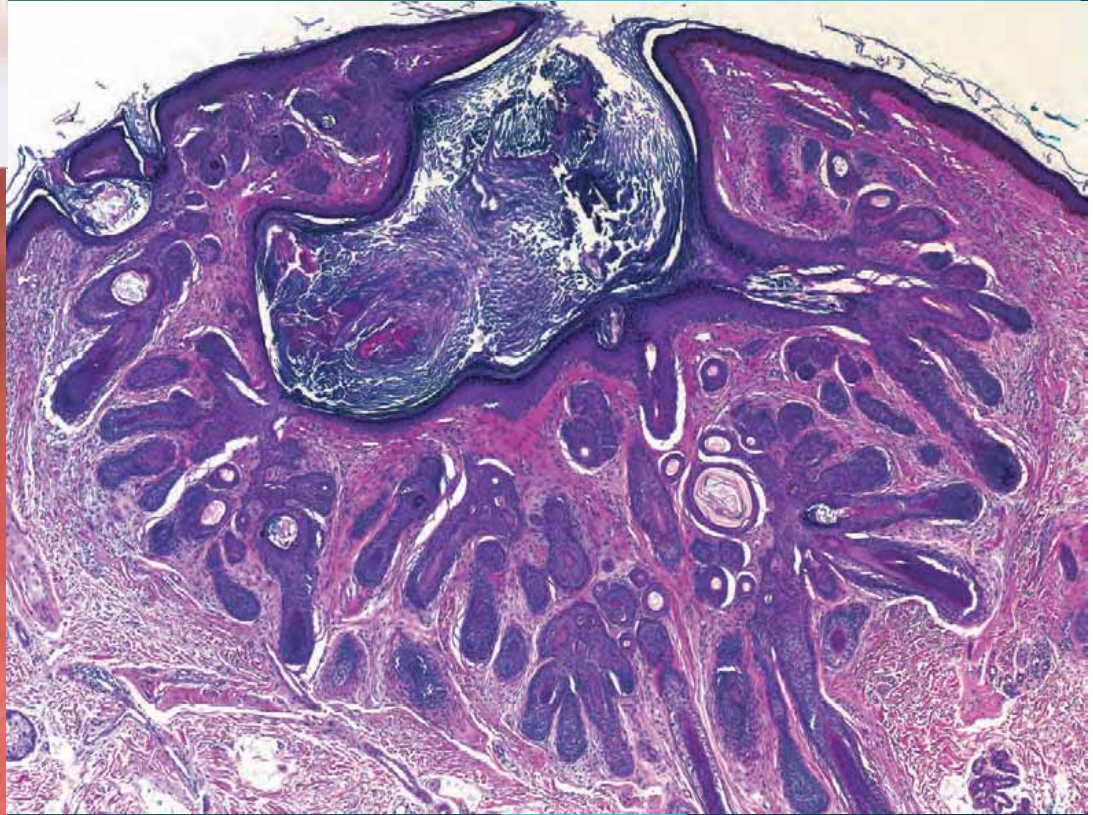
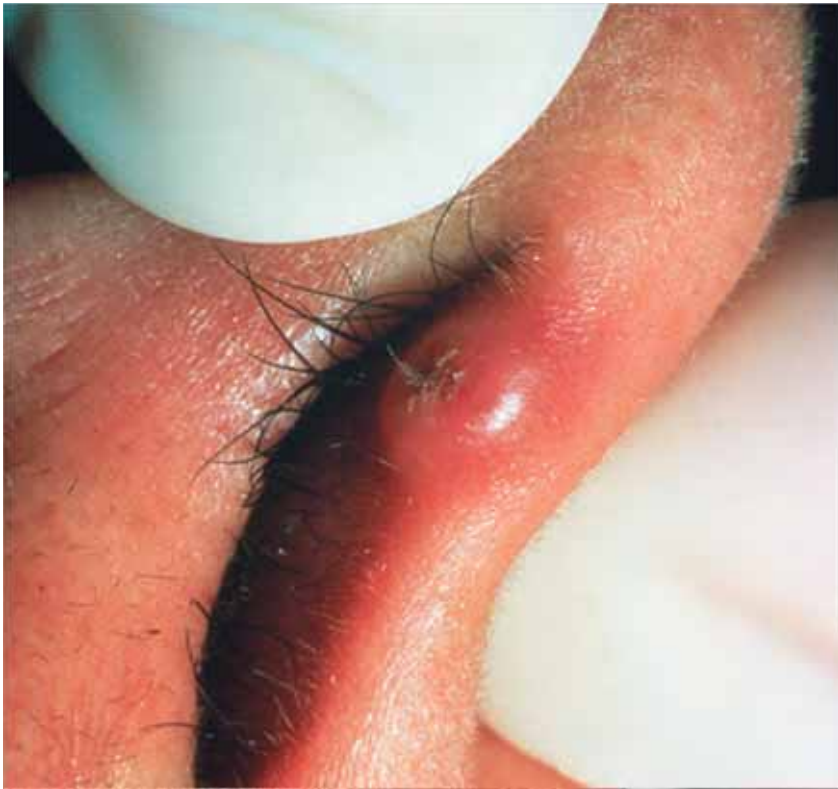
- ▶ skin-colored papule or small nodule
- ▶ face or upper trunk, lesions have a special predilection for the nose





# *Trichofolliculoma*

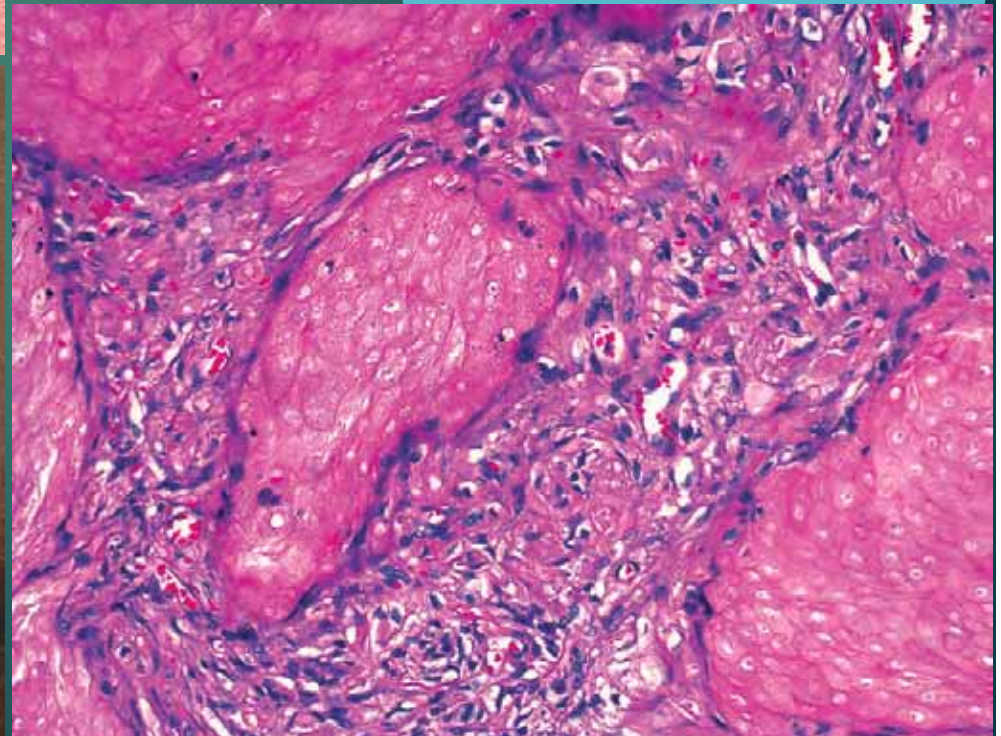
- ▶ a skin-colored papule with a dilated central pore
- ▶ no treatment is needed





# Pilomatrixoma

- ▶ solitary skin-colored or bluish nodule
- ▶ head or upper trunk
- ▶ childhood and adolescence



# Adnexal Neoplasms

- ▶ follicular lineage
  - ▶ Nevus sebaceus
  - ▶ Trichofolliculoma
  - ▶ Trichoepithelioma
  - ▶ Pilomatricoma
- ▶ Sebaceous differentiation
  - ▶ Sebaceous gland hyperplasia
  - ▶ Fordyce's disease
- ▶ apocrine differentiation
  - ▶ Syringoma
- ▶ eccrine differentiation
  - ▶ Eccrine nevus



# Sebaceous gland hyperplasia

- ▶ yellowish, occasionally telangiectatic papules
- ▶ usually on central or upper face





# Sebaceous gland hyperplasia







ectopic sebaceous glands  
(Fordyce granules/spot)

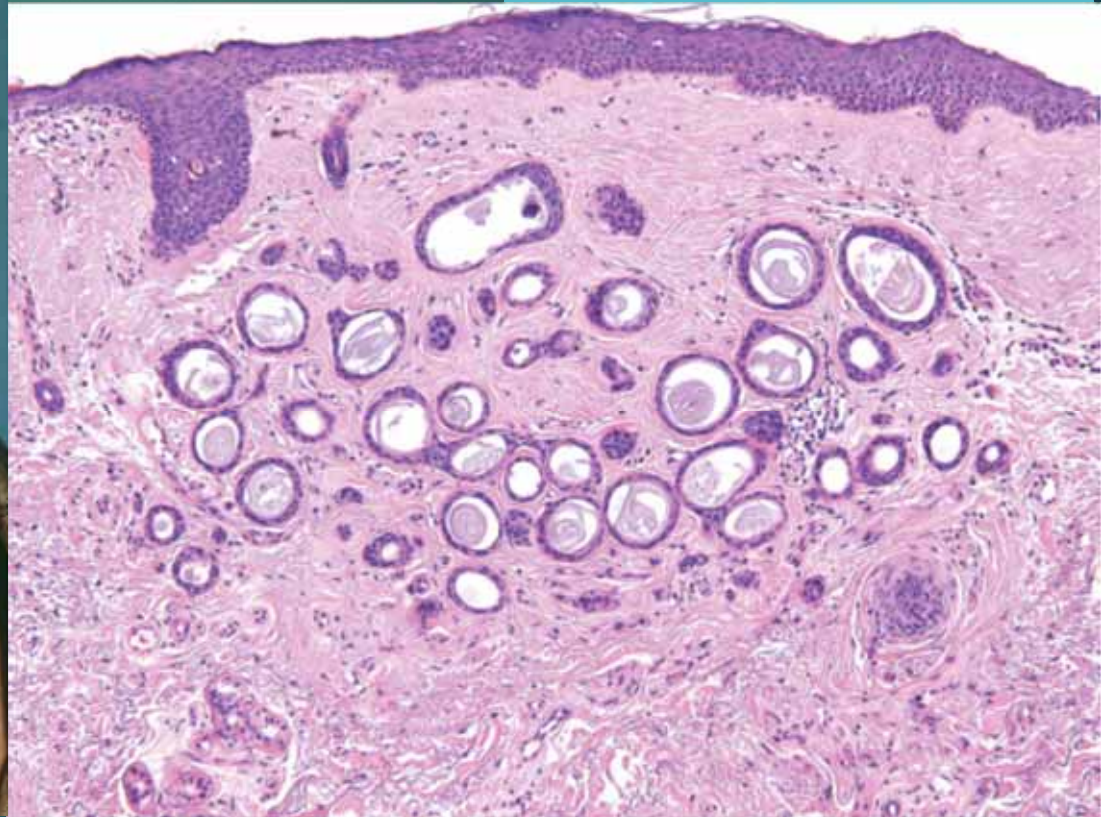
# Adnexal Neoplasms

- ▶ follicular lineage
  - ▶ Nevus sebaceus
  - ▶ Trichofolliculoma
  - ▶ Trichoepithelioma
  - ▶ Pilomatricoma
- ▶ sebaceous differentiation
  - ▶ Sebaceous gland hyperplasia
  - ▶ Fordyce's disease
- ▶ Apocrine differentiation
  - ▶ Syringoma
- ▶ eccrine differentiation
  - ▶ Eccrine nevus



# Syringoma

- ▶ small firm skin-color papules
- ▶ periorbital area (eyelids)

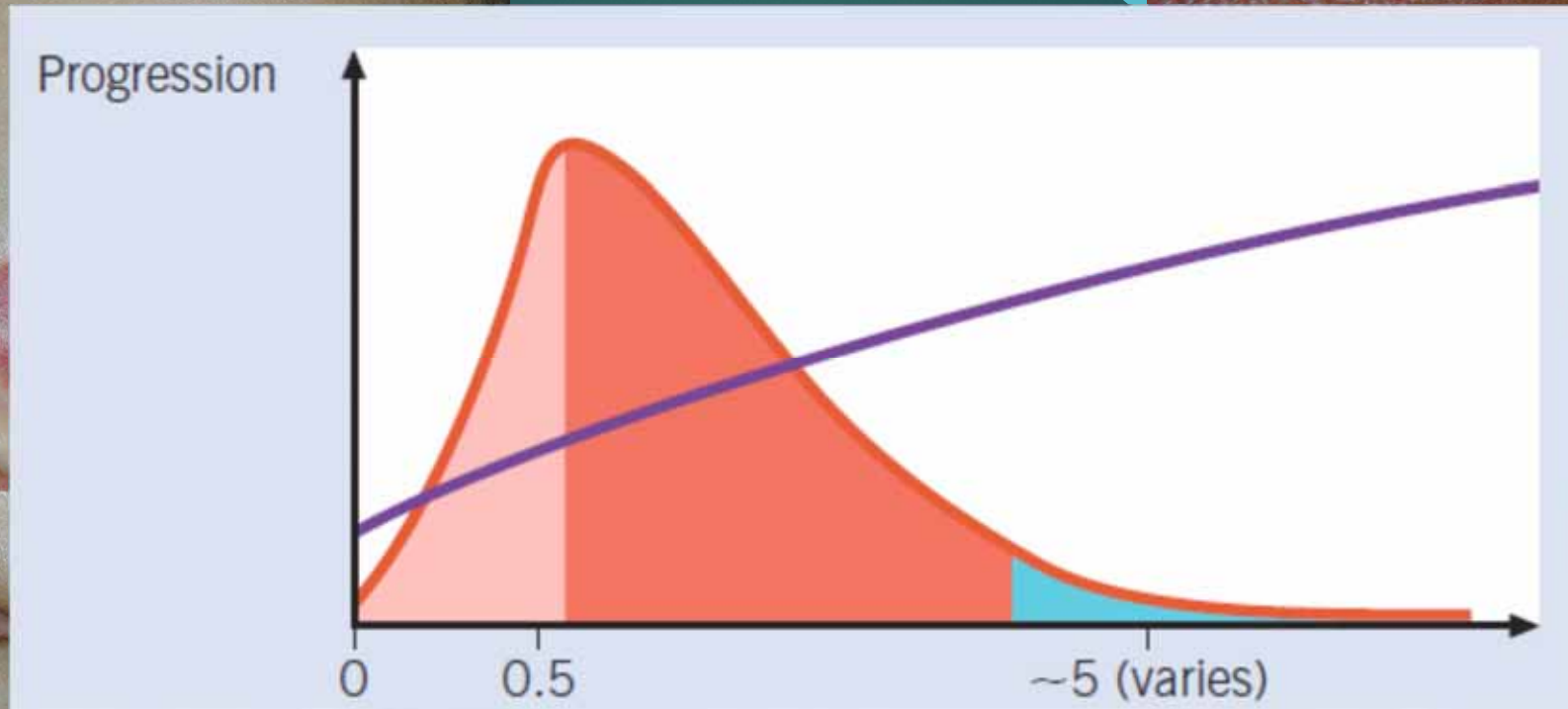


# Vascular Neoplasms

- ▶ Infantile hemangioma
- ▶ Pyogenic granuloma
- ▶ Cherry angioma



# Infantile Hemangioma



- most common benign tumors of childhood
- more common in females (2-5:1 ) and in premature
- rapid growth (proliferate phase) within first 5 months
- heal with telangiectasias, atrophy, fibro-fatty residuum, scarring



01/11/00 (2 mo.)



16/01/01 (4 mo.)





11/12/01 ( 1 y. 3 mo.)



22/04/03 (2 y. 6 mo.)

## *Cherry angiomas (senile angiomas, de Morgan spots)*



- ▶ Bright red, dome-shaped to polypoid papules 1-6 mm.
- ▶ during adult life
- ▶ on trunk and upper extremities



# Pyrogenic Granuloma (Lobular capillary hemangioma)



- Reactive vascular hyperplasia
- most common in children and young adults
- rapidly growing, friable, red papule or polyp of skin or mucosa
- frequently ulcerates
- not involute spontaneously
- Electrocautery or CO<sub>2</sub> laser

# Fibrous and Fibrohistiocytic Proliferations of the Skin

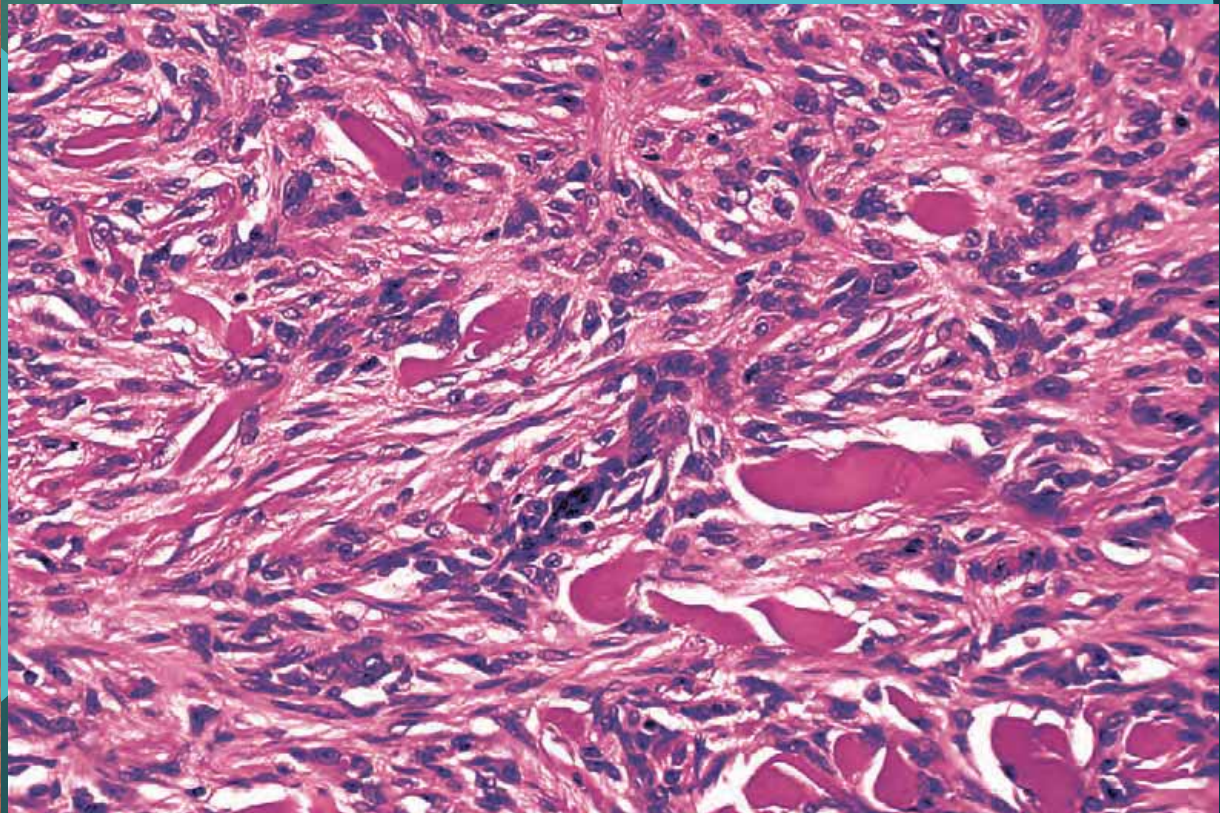
- ▶ Cutaneous angiofibroma
  - ▶ Fibrous papule
  - ▶ Pearly penile papule
- ▶ Acral fibrokeratoma
- ▶ Dermatofibroma





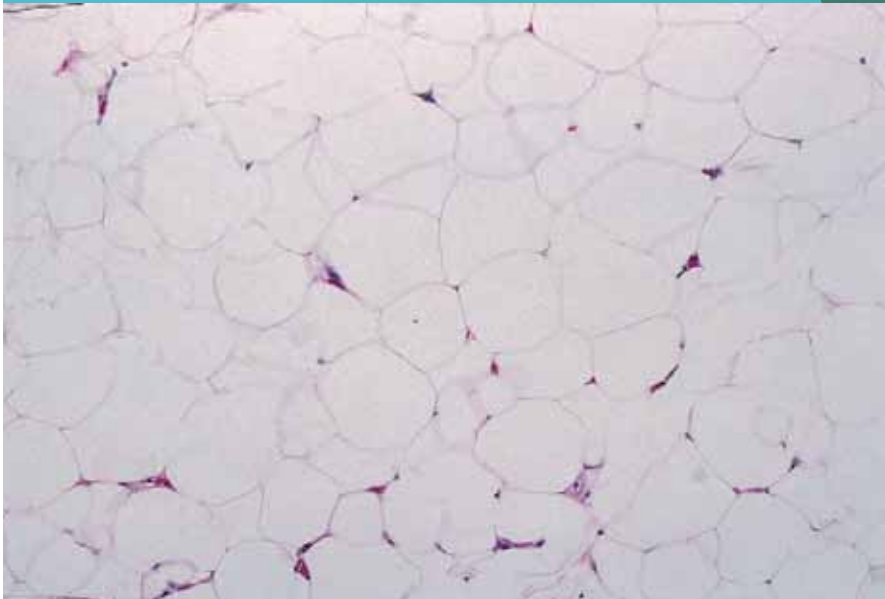
# Dermatofibroma

Hyperpigmented firm papule 0.5-2 cm.



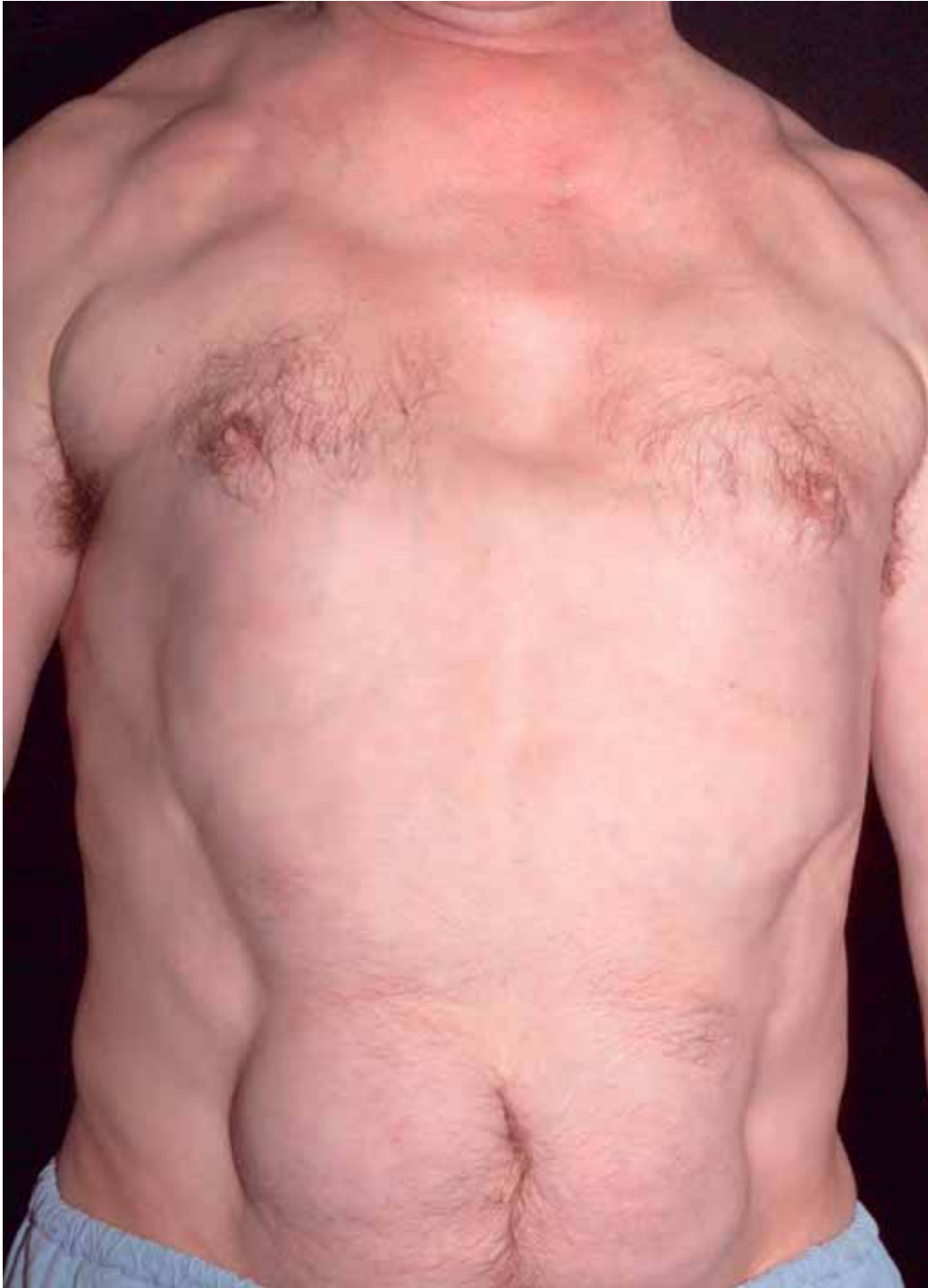


# Lipoma



- ▶ Lipoma : most common soft-tissue tumor
- ▶ Asymptomatic, soft, subcutaneous nodule arising at any site relative sparing of the head, hands, and feet
- ▶ Multiple lipomas are seen in
  - ▶ Madelung's disease
  - ▶ Gardner syndrome
  - ▶ Proteus syndrome
  - ▶ familial multiple lipomatosis
  - ▶ adiposis dolorosa
  - ▶ Bannayan–Riley–Ruvalcaba syndrome





Benign symmetric  
lipomatosis  
(Madelung's disease)



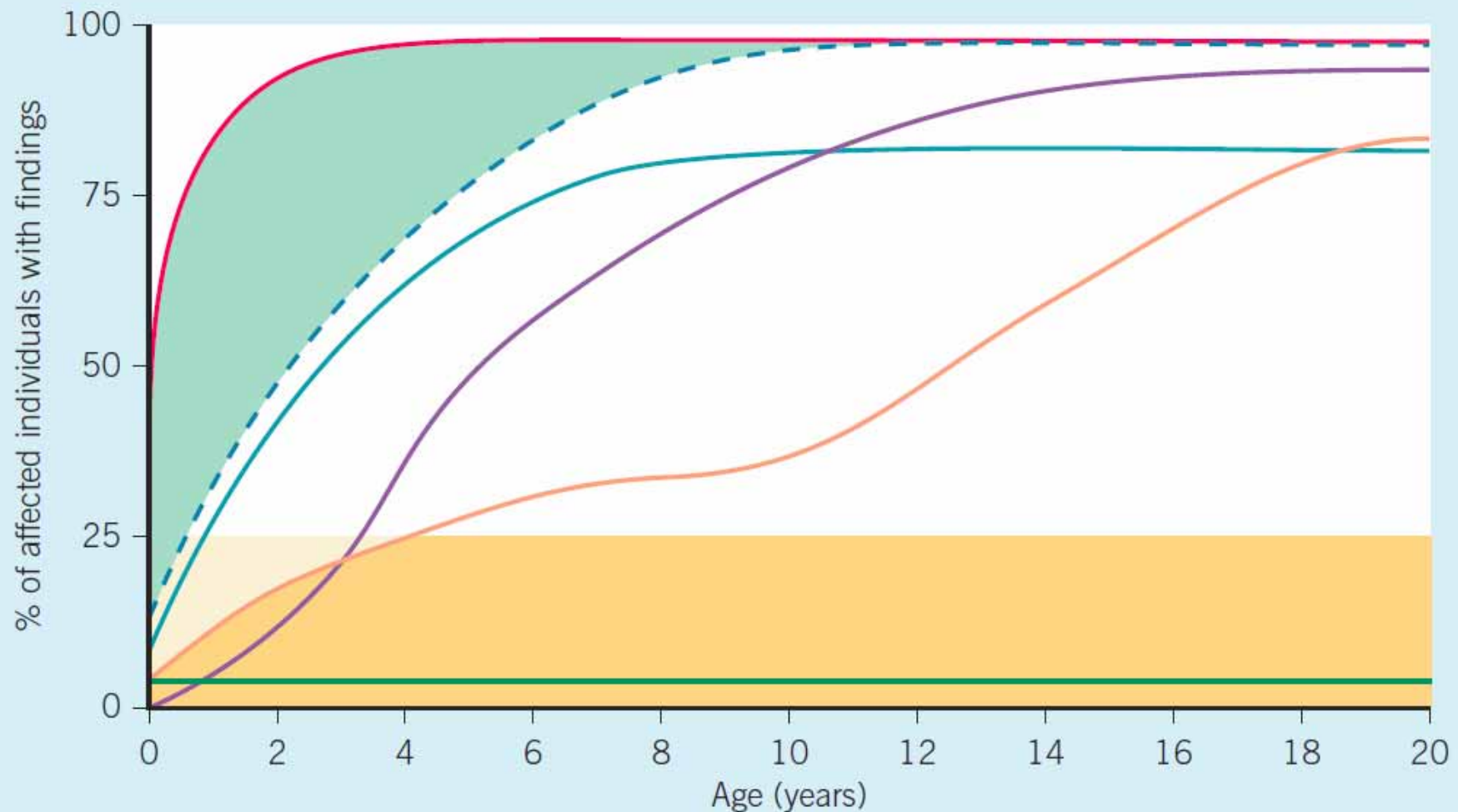
# Neurofibromatosis

- ▶ Neurofibromas (60–90%)
  - ▶ Skin-colored to tan-violet papule or nodule
  - ▶ May be pedunculated or have the “buttonhole” sign
  - ▶ Predilection for the trunk and head
- ▶ Café-au-lait macules (>90%)
- ▶ Axillary and/or inguinal freckling (~80%)
- ▶ Plexiform neurofibroma (25%)





# Development of clinical features in neurofibromatosis type 1.



- $\geq 6$  CALMS
- Intertriginous freckling
- Lisch nodules
- Neurofibromas ( $\geq 2$  of any type or 1 plexiform)
- Sphenoid wing dysplasia or pseudarthrosis
- Meets NF1 diagnostic criteria
- NF1 suspected
- Plexiform NF present but not clinically apparent
- Plexiform NF clinically apparent

# Becker's nevus

- ▶ Unilateral, hyperpigmented and often hypertrichotic patch or slightly elevated plaque
- ▶ Usually on shoulder of male patients
- ▶ Onset during adolescence





# Congenital nevocmelanocytic nevus

- ▶ present at birth
- ▶ Small <1.5 cm , medium, giant > 20cm
- ▶ Risk of melanoma ?



# Giant congenital melanocytic nevus





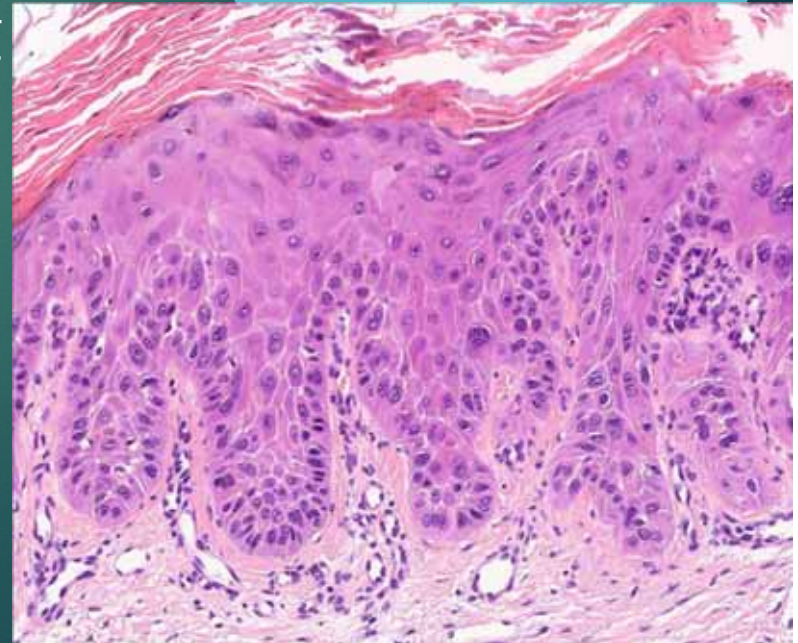
# Acquired melanocytic nevi

- ▶ Junctional
  - ▶ dark brown macule with lighter brown rim.
- ▶ Compound
  - ▶ light to medium brown papule.
- ▶ Intradermal
  - ▶ soft light pink papule.



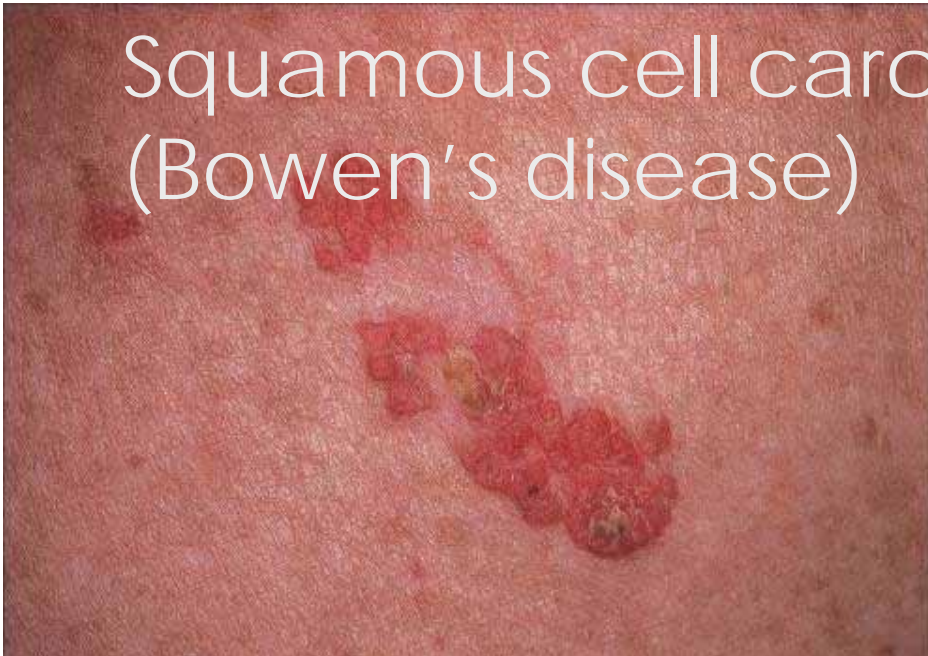
# Precancerous lesions (Squamous Cell Carcinoma in Situ)

- ▶ Bowen's disease
- ▶ Leukoplakia
- ▶ Actinic keratosis
- ▶ Erythroplasia of Queyrat



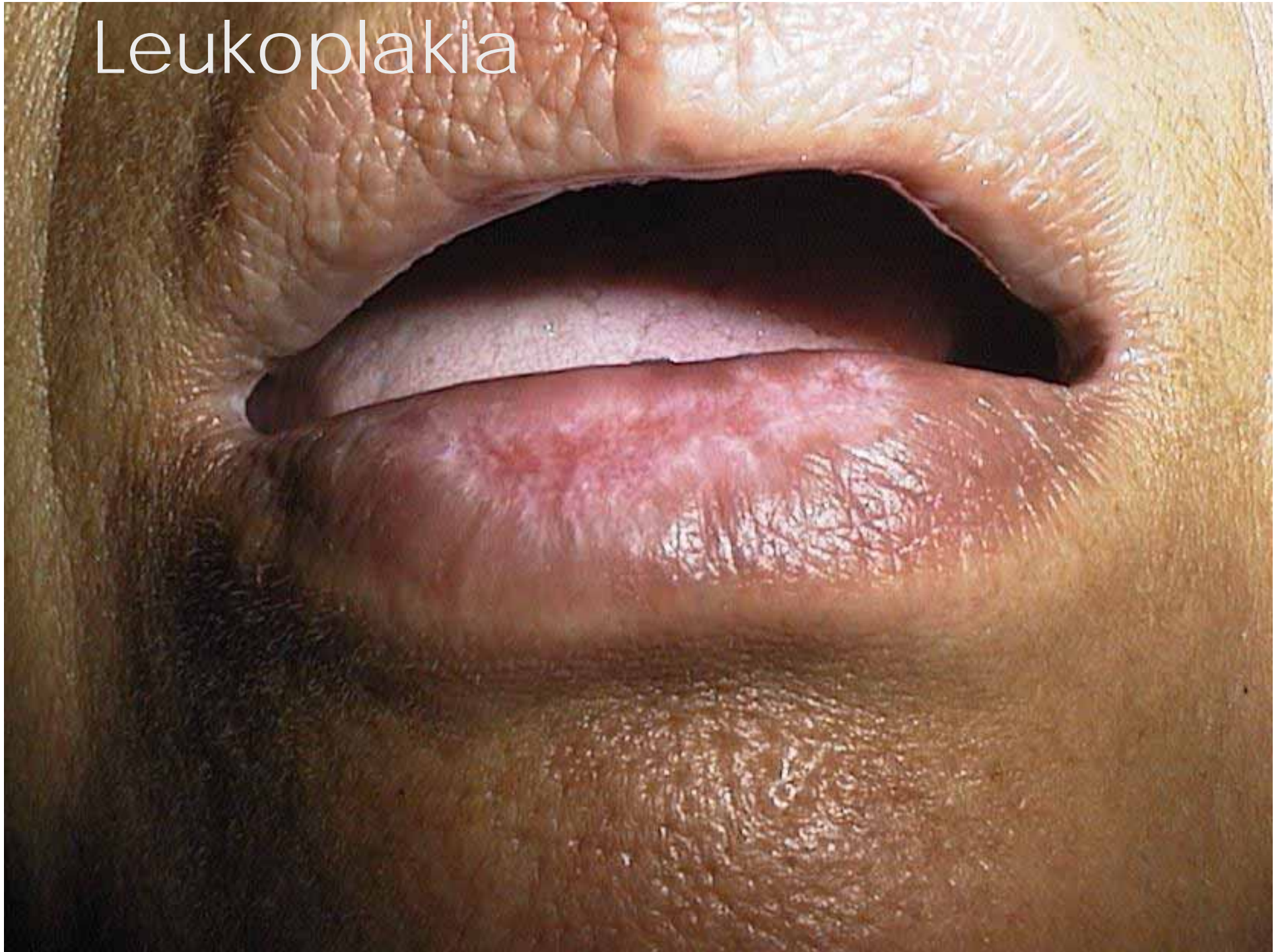


Squamous cell carcinoma in situ  
(Bowen's disease)



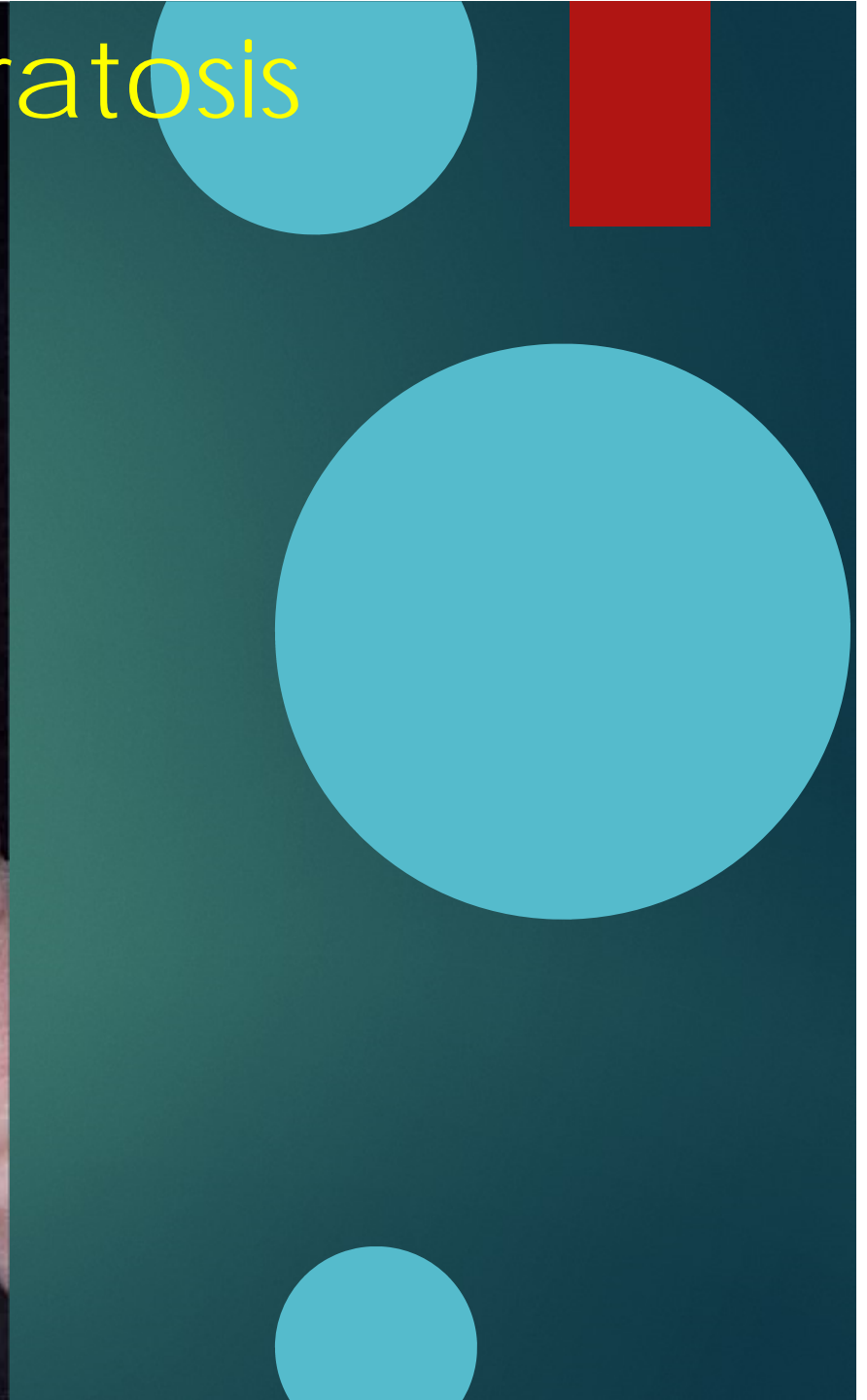


Leukoplakia





# Actinic (solar) keratosis





# Actinic (solar) keratosis





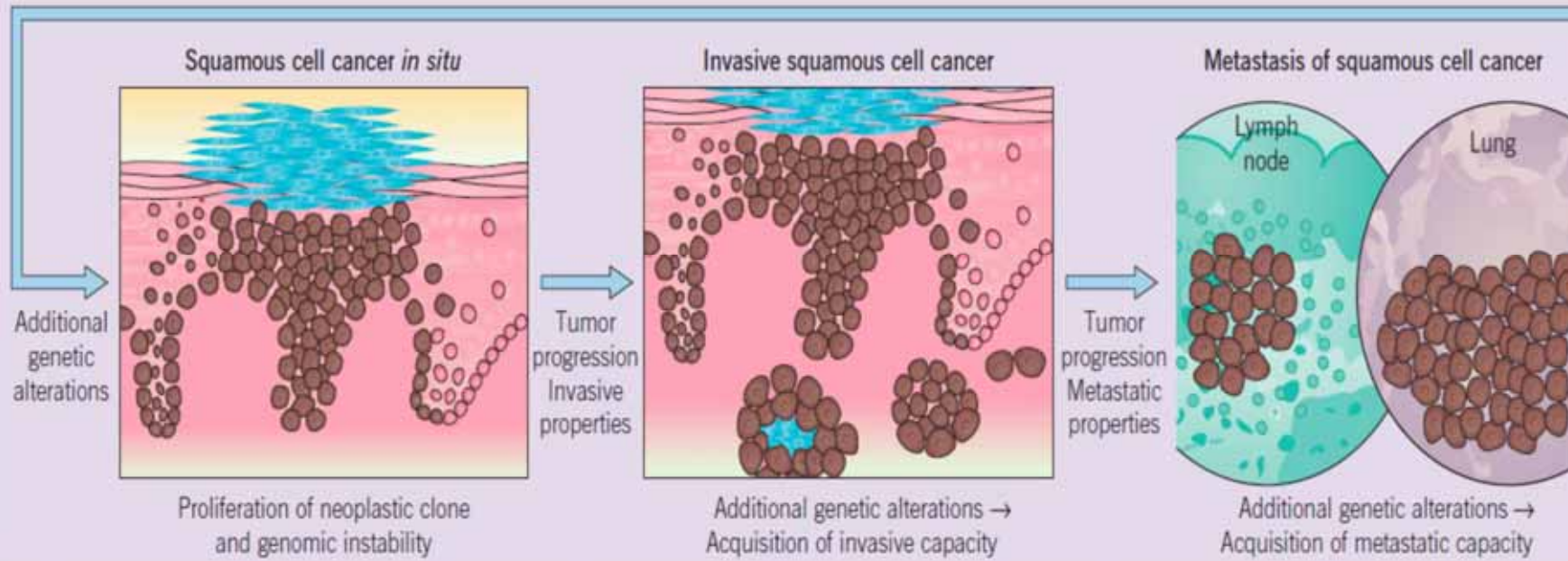
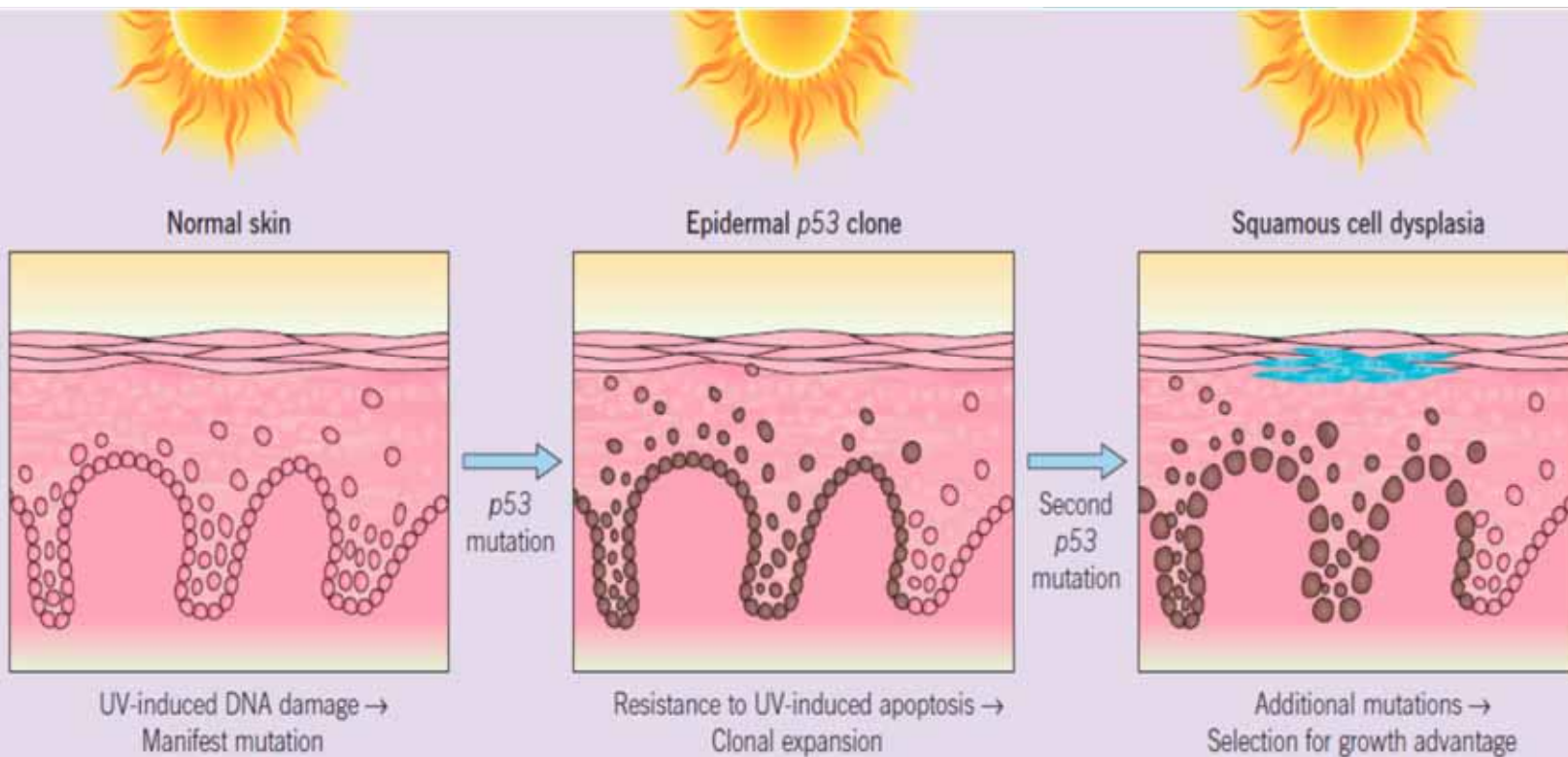
# Erythroplasia of Queyrat of the penis



# Cutaneous malignancy

- ▶ Non-melanoma skin cancers (NMSC)
  - ▶ Squamous cell carcinoma
  - ▶ Basal cell carcinoma
- ▶ Malignant melanoma





# Risk factors for development of SCC and BCC

	SCC	BCC
<b>ENVIRONMENTAL EXPOSURES</b>		
Cumulative/occupational sun exposure	+	
Intermittent/recreational sun exposure		+
Other exposures to UV light (PUVA, tanning beds)	+	+
Ionizing radiation	+	+
Chemicals (arsenic)	+	(+)
HPV	+	
Cigarette smoking	+	
<b>PIGMENTARY PHENOTYPE</b>		
Fair skin	+	+
Always burns, never tans	+	+
Freckling	+	+
Red hair	+	+



# Risk factors for development of SCC and BCC

	SCC	BCC
<b>GENETIC SYNDROMES</b>		
Xeroderma pigmentosum	+	+
Oculocutaneous albinism	+	(+)
Epidermodysplasia verruciformis	+	
Dystrophic epidermolysis bullosa (primarily recessive)	+	
Ferguson–Smith syndrome	+	
Muir–Torre syndrome	+*	(+)*
Nevoid basal cell carcinoma syndrome		+
Bazex and Rombo syndromes		+
<b>PREDISPOSING CLINICAL SETTINGS</b>		
Chronic non-healing wounds	+	
Longstanding discoid lupus erythematosus, lichen planus (erosive) or lichen sclerosus	+	
Porokeratosis (especially linear)	+	
Nevus sebaceus		+ <sup>†</sup>
<b>IMMUNOSUPPRESSION</b>		
Organ transplantation	+	(+)
Other (e.g. chronic lymphocytic leukemia treated with fludarabine, AIDS patients with HPV infection)	+	

# Basal cell carcinoma





BCC (nodular)





BCC (nodular)





BCC (pigmented)





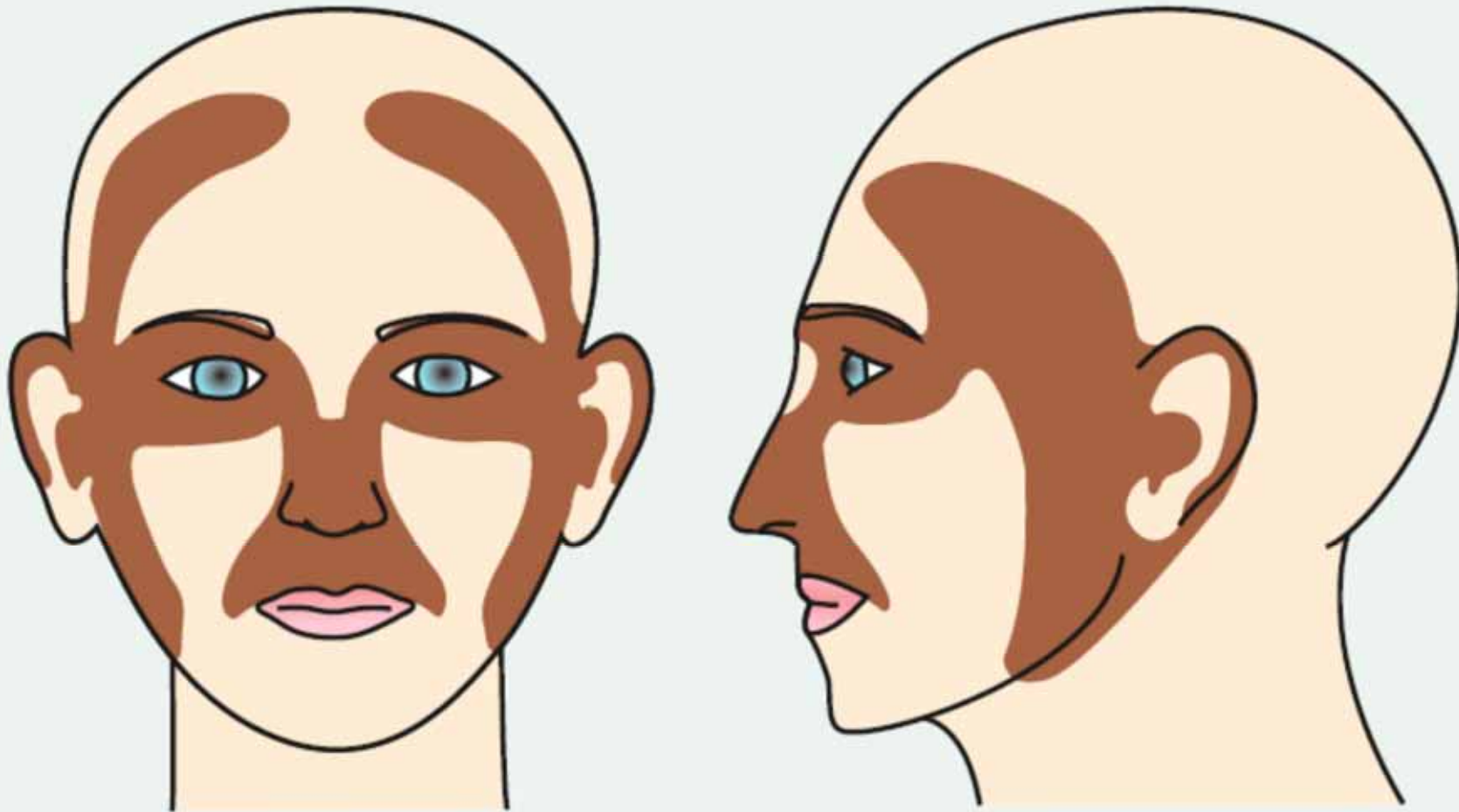




BCC (superficial)



# Indications for Mohs' Micrographic Surgery



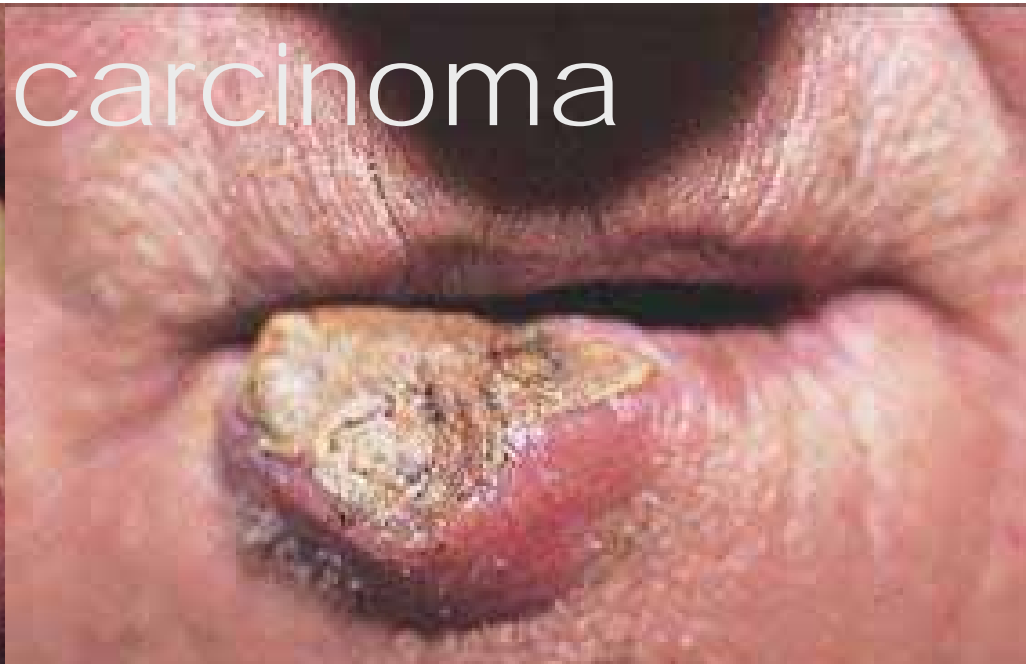


# BCC / Moh's





# Squamous cell carcinoma





# Squamous cell carcinoma





# Squamous cell carcinoma







Type of melanoma	Freq. (%)	Site	Radial growth	Special features
<b>Superficial spreading melanoma</b>	60-70	Any site, preference for lower extremities (female), trunk (male)	Yes	More pagetoid, less solar elastosis
<b>Nodular melanoma</b>	15-30	Any site, preference for trunk, head, neck	No	Nodule with vertical growth
<b>Lentigo maligna melanoma</b>	5-15	Face, especially nose and cheeks	Yes	Slower growth over years on sun-damaged skin
<b>Acral lentiginous melanoma</b>	5-10	Palms, soles, subungual	Yes	Most common melanoma in patients with darker skin types



# Superficial spreading melanoma



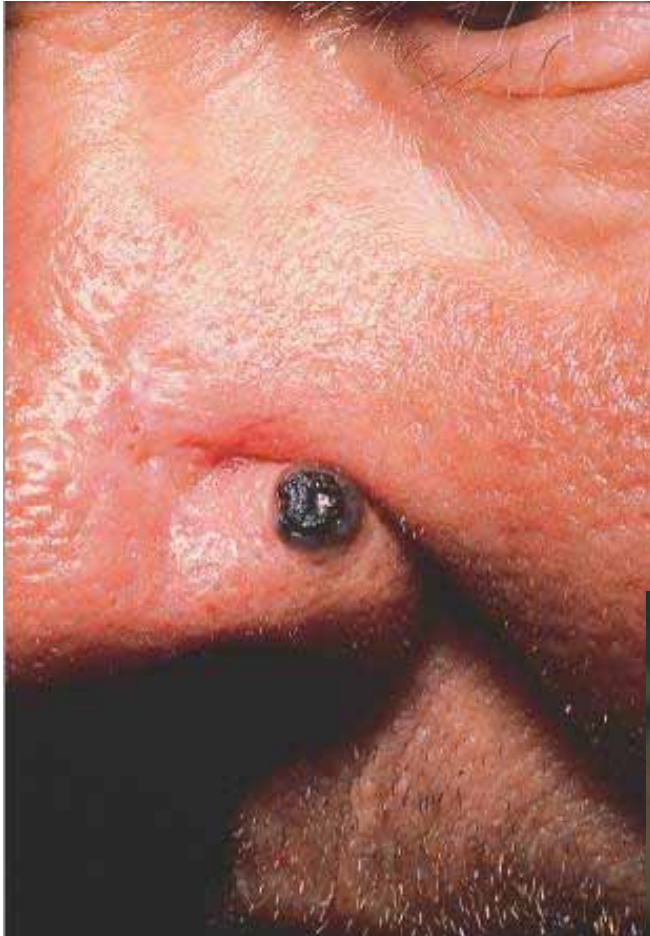


# Superficial spreading melanoma





# Nodular melanoma





# Nodular melanoma





# Lentigo maligna melanoma





# Lentigo maligna melanoma





# Acral lentiginous melanoma













END

