

# Top Dermatologic Issues in Primary Care

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**E~~X~~PLORE**  
HEALTHCARE SUMMIT



## Disclosure

- I have served as a consultant for Castle Biosciences
- I have served as a consultant for Aegle Therapeutics

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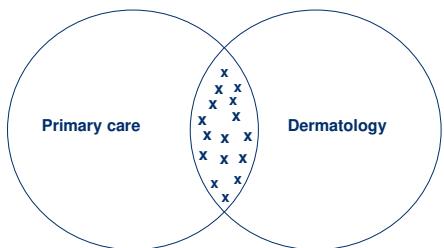
## Goals of today's talk

- Review the morphologic range of dermatologic disease
- Emphasize dermatologic diagnoses in primary care setting
- NOT to review the entirety of relevant dermatology
- Emphasize the essential role of a biopsy in making a diagnosis

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## Scope of this talk



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## Our why:

- “Skin conditions are the most common reason for a new presentation to a primary care physician”\*



\* Roux E Le, Edwards PJ, Sanderson E, Barnes RK, Ridd MJ. The content and conduct of GP consultations for dermatology problems: A cross-sectional study. *Br J Gen Pract*. 2020;70(699):e723–30.

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Grada et al. *J Clin Aesthet Dermatol*. 2022 May; 15(5): E82–E86.

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## A quick tour through the world of dermatologic morphology

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**Describing Lesions**

- Size
- Color
- Primary Lesion Type
- Secondary Lesion Type (if present)
- Configuration
- Location

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**Lesion Types****Primary**

Changes in the skin directly caused by the disease process.

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**Secondary**

Changes in the skin caused by external forces (scratching, trauma, infection, or the healing process).

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**Primary Lesions**

Macule	Bulla
Patch	Pustule
Papule	Wheal
Plaque	Telangiectasia
Nodule	Cyst
Tumor	Comedones (open & closed)
Vesicle	

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**Macule** < 1cm flat, non-palpable, change of skin color.

Examples



Freckles  
(Ephelides)



Solar  
Lentigines



Junctional  
Nevus

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**Patch** > 1cm flat, non-palpable, change of skin color.



Vitiligo



Port Wine  
Stain

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**Papule** < 1cm superficial, raised, palpable lesion with distinct borders



Skin Tags  
(Acrochordons)



Molluscum  
Contagiosum



Seborrheic  
keratoses



Intradermal nevus

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**Plaque** – >1 cm raised, flat-topped, palpable lesion greater than 1 cm in diameter.



Psoriasis



Atopic Dermatitis

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**Nodule** – Firm lesion less than 1 cm in diameter. It can be located in epidermis, dermis, or subcutaneous tissue. Increased depth differentiates nodules from papules.



Rheumatoid Nodules



Nodular Acne



Lipoma

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**Tumor** – Solid mass in skin or subcutaneous tissue > 2 cm.




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**Fluid filled sacs:**

< 1 cm → Vesicle  
> 1 cm → Bulla



Herpes Simplex (vesicle)



Contact Dermatitis



Bullous Pemphigoid (Bulla)

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**Pustule** – vesicle containing “puss” which is neutrophil-rich.  
Can be sterile or infectious.



Folliculitis



Acne



Pustular psoriasis

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**Wheal** – Edema in upper dermis.



Urticaria

**Telangiectasia** – Dilated superficial blood vessel.



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**Cyst** – Cavity containing fluid, solid or semi-solid material



Epidermal Inclusion Cyst

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**Comedones** – A plug of keratin or sebum within the dilated orifice of a hair follicle (non-inflammatory)



Closed "whitehead"

Open "blackhead"

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### Secondary Lesions

Scale  
Excoriation  
Lichenification  
Fissure  
Erosion  
Ulcer

Crust  
Atrophy  
Purpura  
Hyper/Hypo-pigmentation

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**Scale** – Flakes or plates of desquamated stratum corneum

Seborrheic Dermatitis



Xerosis

**Crust** Dried plasma or exudates.

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Impetigo

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**Atrophy** – Thinning or absence of epidermis, dermis, or subcutaneous fat.



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**Lichenification** – Thickening of epidermis with exaggerated skin lines. Usually from chronic scratching/rubbing.



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**Erosion** — Loss of part or all of the epidermis.



(Pemphigus Vulgaris)

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**Ulcer** — Loss of epidermis and dermis due to necrosis.



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**Excoriation** — Loss of superficial epidermis due to trauma.



(ie: scratching, picking)

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**Fissure** — Crack in skin due to dryness.



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**Petechiae, Purpura, & Ecchymosis** — Non-blanchable bleeding in skin.

Size: petechiae < 3 mm  
purpura 3 mm – 1 cm  
ecchymosis > 1 cm



Petechiae



Palpable Purpura



Ecchymosis

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## Hypo/ Hyper-pigmentation

Secondary lightening or darkening of the skin.



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## Skin Configurations

Annular  
Linear  
Grouped  
Serpiginous  
Arcuate

Disseminated/Generalized  
Confluent  
Reticulated

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### Annular: Ring shaped



Tinea  
Corporis

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**Linear:** In a line.



Koebner's Phenomenon

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**Grouped:** Lesions that are clustered together.



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**Serpiginous:** wavy or “snake-like” in appearance.



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**Arcuate:** crescent or “half-moon” shaped



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**Reticular:** lesions with a “net-like” arrangement.



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**Disseminated/Generalized:** Describes a lesion that is usually localized that has spread



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**Confluent:** running together



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**Location**

- Intertriginous
- Photodistributed
- Palmar/Plantar
- Dermatomal
- Symmetrical
- Blaschko's Lines

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**Intertriginous:** Area where two skin surfaces touch or rub together.



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**Photodistributed:** in areas exposed to sunlight.



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**Palmar/Plantar:** relating to the palm of the hand or sole of the foot.



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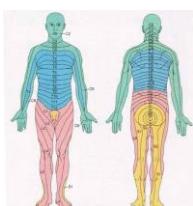
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**Dermatomal:** corresponding to a dermatome of the body.



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**Symmetrical:** Made up of exactly similar parts facing each other or around an axis



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**Blaschko's Lines:** skin lines that trace the migration of embryonic cells.



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### What are the most common dermat diagnoses in primary care?

- Study in 2022: on the National Ambulatory Medical Care Survey (NAMCS) between 2007 and 2016, the most recent years available:
- The NAMCS is an ongoing survey which provides objective information about the use of ambulatory medical services in the United States.
  - The survey is conducted annually by the National Center for Health Statistics (NCHS) at the Centers for Disease Control and Prevention (CDC).
  - The NAMCS surveys a large, generalizable sample of physicians and non-physician providers and has achieved high response rates of up to 77%.

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Ahn CS, Allen MM, Davis SA, Huang KE, Fleischer AB, Feldman SR. The National Ambulatory Medical Care Survey: A resource for understanding the outpatient dermatology treatment. *J Dermatolog Treat.* 2014;25(6):453–458.

Arafa AE, Anzengruber F, Mostafa AM, Navarini AA. Perspectives of online surveys in dermatology. *J Eur Acad Dermatol Venereol.* 2019;33:511–520.

※ Grada et al. *J Clin Aesthet Dermatol.* 2022 May; 15(5): E82–E86.

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## The most common skin diagnoses in primary care

- In the population-based, cross-sectional analysis using the National Ambulatory Medical Care Survey between 2007 and 2016:
- The five most common skin diagnoses among all medical specialties were
  - contact dermatitis
  - acne vulgaris
  - actinic keratosis
  - "benign neoplasm" of the skin
  - epidermoid cyst



✉ Grada et al. J Clin Aesthet Dermatol. 2022 May; 15(5): E82–E86.

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## Other "Top" Dermatologic Issues for Primary Care

- Identify a skin malignancy
- Identify eczematous, psoriasiform, lichenoid, and drug-induced conditions
- Identify potential autoimmune connective tissue diseases
- Identify autoimmune bullous dermatoses
- Barriers to sampling the skin in primary care
  - Requires proper set up, equipment for procedures, photography/ triangulation of lesions, proper sample containers (ex. Michel's media for direct immunofluorescence).
- Delay in referral / wait times for patients to be seen by dermatology
- Delay in diagnosis and treatment

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## A bit of a deeper dive into

### The most common issues

- Acne vulgaris
- Epidermoid cyst
- "Benign" neoplasms of the skin
- Actinic keratosis
- Contact dermatitis

### Other top issues

- Cutaneous malignancy
  - Basal cell carcinoma
  - Squamous cell carcinoma
  - Melanoma
- Refractory inflammatory dermatoses
  - Eczematous
  - Psoriasiform
  - Lichenoid
- Autoimmune connective tissue diseases
  - Ex. cutaneous lupus
- Autoimmune bullous diseases
  - Ex. bullous pemphigoid



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The top most common

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**Acne vulgaris  
vs rosacea – diagnosis**



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**Acne  
vs  
Rosacea**

OTHER FEATURES: Acne Vulgaris	OTHER FEATURES: Rosacea
<ul style="list-style-type: none"> <li>Most prevalent in adolescents and young adults</li> <li>Variable distribution on face</li> <li>Frequent shoulder, chest, and/or back involvement</li> <li>Sequela of postinflammatory hyperpigmentation, postinflammatory erythema, and scarring</li> <li>Association with hyperandrogenic disorders (e.g., polycystic ovarian syndrome)</li> </ul>	<ul style="list-style-type: none"> <li>Most prevalent in adults &gt;30 years old</li> <li>Centrifugal distribution (cheeks, nose, chin)</li> <li>Ocular involvement (e.g., symptoms of eye irritation, eyelid erythema, conjunctival injection, crusting, recurrent hordeolum and chalazion)</li> <li>Sensitive skin</li> <li>Flushing</li> </ul>
<b>KEY CONCEPTS</b>	
<p><b>Acne vulgaris and rosacea are common causes of inflamed papules or pustules on the face.</b> Recognition of other characteristic features is helpful for distinguishing these conditions. Patients may exhibit some or all of the displayed features.</p> <p>Distinguishing between acne vulgaris and rosacea is important because of differences in the approach to patient evaluation and treatment. For example, an assessment for signs of ocular involvement (e.g., eyelid erythema, conjunctival injection, crusting) is an important component of the initial evaluation of female patients with acne vulgaris, particularly in the presence of severe, sudden onset, or recalcitrant acne. In patients with rosacea, an assessment for signs or symptoms of ocular involvement is important for identifying patients who may benefit from ophthalmologic examination.</p>	

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**Acne vulgaris vs rosacea – treatment****Acne**

- Daily wash with benzoyl peroxide-containing wash (Ex. CeraVe with benzoyl peroxide) or salicylic acid wash
- Topical clindamycin solution, gel, or lotion
- Daily retinoid (ex. OTC adapalene gel, or tretinoin creams) – a pea-sized amount only across entire face at night
- Oral medications: doxycycline 100 mg BID (or minocycline) for up to 1 month, can consider refills for flares
- Hormonal driven: start with spironolactone 50 mg daily, increase to 100 mg daily as tolerated (consider checking potassium; warn of side effects; not for use in woman trying to get pregnant)
- Also consider topical Winlevi (clascoterone) – androgen receptor inhibitor

**Rosacea**

- Start topical metronidazole gel
  - If fails, consider topical ivermectin (Soolantra)
- Dermatologist: can perform lasers (example PDL to target hemoglobin in telangiectasias)
- Wash with sensitive skin cleaners (Cetaphil, CeraVe, Vanicream, etc).
- Can consider long-term, low dose doxycycline 50 mg daily, or 40 mg Oracea (slow-release)
- Can consider vasodilators (topical brimonidine – α2 adrenergic receptor agonist)
- Identify and reduce triggers as much as possible (alcohol, spicy foods, heat, stress, etc)
- Refer to ophthalmology if ocular involvement

⌘ Isotretinoin for severe cases

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**Epidermoid inclusion cysts - diagnosis**

© 2015 VisualDx.com

© 2015 VisualDx.com

Beware of the "cyst" – if deeper with no punctum, it may not be a "cyst"

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**Epidermoid inclusion cysts - differential****Lipoma****Ganglion cyst****Pilar cyst****Dermoid cyst**

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Cysts? Unfortunately not.



✉ Pajaziti, L., Hacıçı, S.R., Dobruna, S. et al. Skin metastases from lung cancer: a case report. BMC Res Notes 8, 139 (2015). <https://doi.org/10.1186/s13104-015-1105-0>

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#### Benign neoplasms of the skin (examples)



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#### "Pyogenic granuloma" (lobular capillary hemangioma) vs other?



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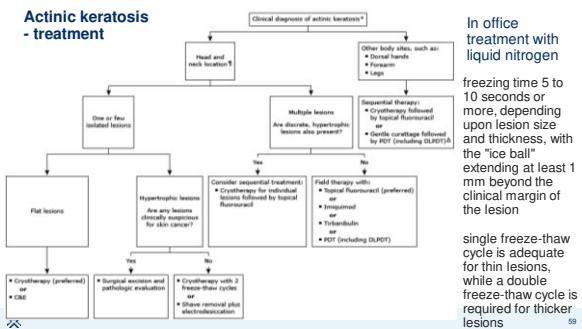
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## Actinic keratoses



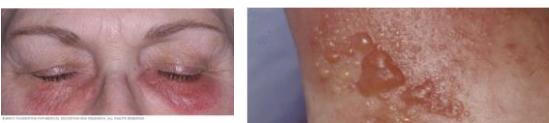
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## Contact dermatitis - diagnosis



- Common contact allergens include plant allergens, metals, fragrances, acrylics, medicaments, and preservatives.

History and geometric distribution are important

Useful resource: Contact Dermatitis Institute ([www.contactdermatitisinstitute.com](http://www.contactdermatitisinstitute.com))

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### Contact dermatitis – treatment/ avoidance



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### The other “Top” issues

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### Skin cancer – The “big 3” – diagnosis - clinical



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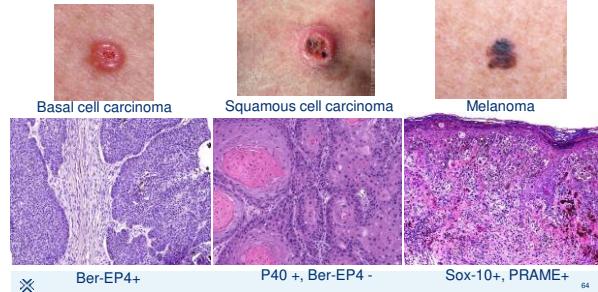
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### Skin cancer – The “big 3” – diagnosis - dermatopathology




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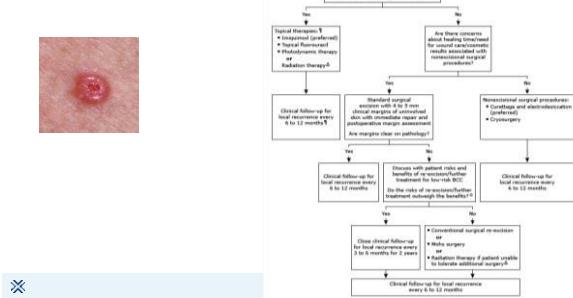
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### Skin cancer/BCC - treatment




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### Appropriate Use Criteria for Mohs

Mohs Surgery Appropriate Use Criteria  
American Academy of Dermatologists

OPEN

Details    Reviews    Related

iPhone    Cancer Type    Score    Restart

Area H    MEDIAN SCORE IS 9

APPROPRIATE

The use of Mohs is appropriate for the specific clinical presentation.

SELECTED CRITERIA

- Basal Cell Carcinoma
- Area H
- Aggressive

View decision tree

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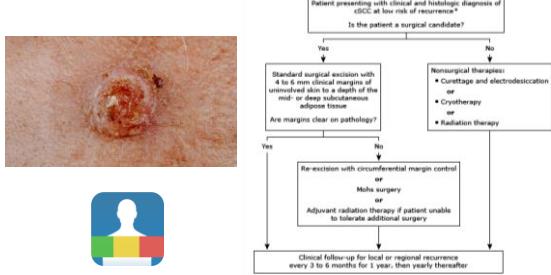
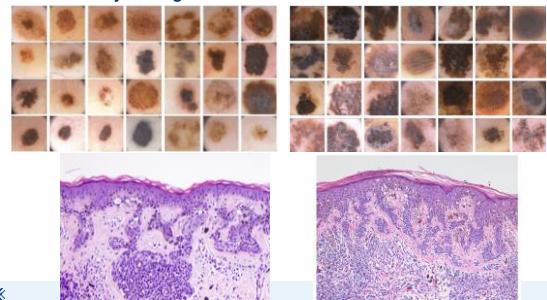
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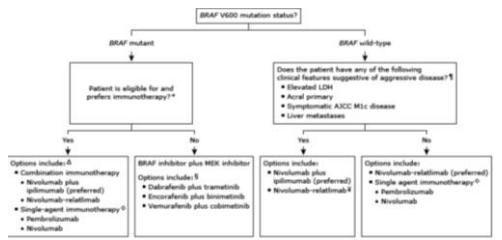
**Skin cancer/SCC - treatment****The melanocytic diagnostic dilemma****Melanoma- staging****Definition of Primary Tumor (T) - AJCC 8<sup>th</sup> Edition**

T Category	Thickness	Ulceration status
Tis (melanoma <i>in situ</i> )	Not applicable	Not applicable
T1	$\leq 1.0$ mm	Unknown or unspecified
T1a	<0.8 mm	Without ulceration
T1b	>0.8 mm 0.8–1.0 mm	With ulceration With or without ulceration
T2	>1.0–2.0 mm	Unknown or unspecified
T2a	>1.0–2.0 mm	Without ulceration
T2b	>1.0–2.0 mm	With ulceration
T3	>2.0–4.0 mm	Unknown or unspecified
T3a	>2.0–4.0 mm	Without ulceration
T3b	>2.0–4.0 mm	With ulceration
T4	>4.0 mm	Unknown or unspecified
T4a	>4.0 mm	Without ulceration
T4b	>4.0 mm	With ulceration

Gersbach, Sclimenti, et al. Melanoma. In Arns, M.B., Edge, S.B., Greene, F.L., et al. (Eds.) AJCC Cancer Staging Manual. 8th Ed. New York: Springer; 2017.

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### Melanoma- treatment



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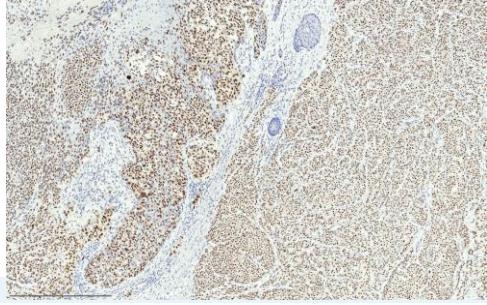
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### PRAME (PReferentially-expressed Antigen in MElanoma)



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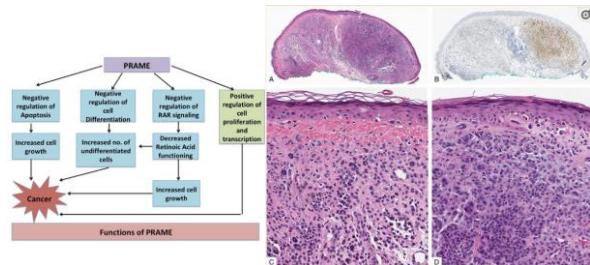
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### PRAME in melanoma



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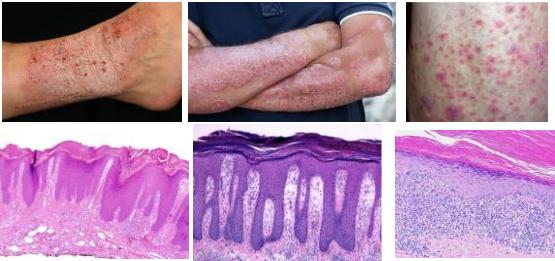
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### Eczematous vs psoriasiform vs lichenoid - diagnosis



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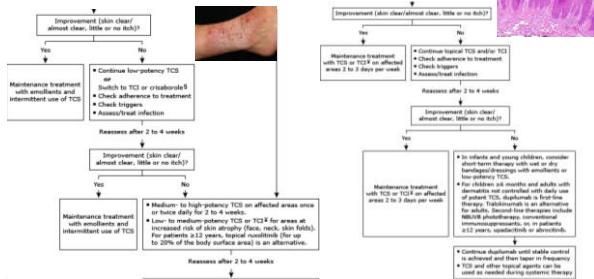
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### Eczema / atopic dermatitis - treatment



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### Psoriasis – treatment - biologics



Biologic	Other Compatible Conditions	Contraindications	Dosing	Approx. Cost (first Year)	Common Adverse Reactions (>10%)	Efficacy - Primary Outcome and Long term Outcome
Adalimumab (Humira) TNF	Psoriasis Rheumatoid Arthritis Gerd/ECZ*	Acute ( $\geq$ 70) or other severe infections; Malignancy; Reproductive disorder; Developing/treated Tuberculosis	Every 2-wk (SC)*	\$21,000*	Infection at site; Headache; Nausea; Antibody development CEST/Other infections*	PASI 75 at Week 16: 71.7%; PASI 50 at Week 16: 71.7%; PASI 30 at Week 16: 71.7%*
Cetuximab (Regen-Cet) TNF	Acute ( $\geq$ 70) or other severe infections; Heart failure*		Every 2-wk (SC)*	\$18,275*	Headache; Nausea; Antibody development CEST/Other infections*	PASI 75 at Week 16: 71.4%; PASI 50 at Week 16: 69.3%; PASI 30 at Week 16: 68.6%*
Tacrolimus (Protopic) TNF	PAH Dyslipidemia*	Hypersensitivity to tacrolimus; Pain or changes systemic*	Topical twice for 3 mos, then once weekly (SC)*	\$21,000*	Infection at site; Headache; Nausea; CEST/Other infections*	PASI 75 at Week 12: 47.4%; PASI 75 at Week 36: 73.9%*
Infliximab (Remicade) TNF	Crohn's PAH Dyslipidemia*	Sinus infection*; Heart failure*	IV infusion q 8-12 wks, then every 4-8 wks*	\$30,000*	Infection at site; Antibody development Antibiotic resistance CEST/Other infections*	PASI 75 at Week 16: 70.0%; PASI 75 at Week 30: 70.0%*
Secukinumab (Cosentyx) TNF		Sinus infection*; Heart failure; Dyslipidemia*	SC, IM, or IV infusion every 4-8 wks*	\$21,000*	Infection at site; Antibody development Contractile synapse CEST/Other infections*	Not reported (refer to infliximab)*

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<https://www.skintherapyletter.com/psoriasis-tool-biologics/>

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### Psoriasis – treatment - biologics

Biologic	Other Considerable Conditions	Contraindications	Dosing	Approx. Cost (First Year)	Common Adverse Reactions (>30%)	Efficacy – Primary Outcome and Long-term Outcome
Biologics (IL-1)	PtA Pregnancy/Breastfeeding*	Crohn's Disease Hyperthyroiditis by levothyroxine*	Weeks 0-16: Once every 1 week (QW) Week 17-24: Once every 4 weeks (Q4W)	\$14,000*	ER375mg subcutaneous*	PtA A 61% vs Week 12 PtA B 51% vs Week 12 *of PtA C responds maintained until Week 48 *of PtA D
Biologics (Tnf)	PtA Pregnancy/Breastfeeding*	Hyperthyroiditis by levothyroxine Disease*	Every two weeks until week 11, then every four weeks (Q2W)	\$70,000*	ER375mg subcutaneous*	PtA A 61% vs Week 12 PtA B 51% vs Week 12 *of PtA C responds maintained until Week 48 *of PtA D
Biologics (Cytokines)	PtA Pregnancy/Breastfeeding*	Hyperthyroiditis by levothyroxine HIV Chronic Infectious*	Loading dose weekly for 4 weeks, then every 4 weeks (Q4W)	\$24,000*	ER375mg subcutaneous*	PtA E 75% vs Week 12 PtA F 75% vs Week 12 *of PtA G responds maintained until Week 48 *of PtA H
Biologics (Tumor)	PtA (plus IL-17C)* Neurofibromatosis*	Hyperthyroiditis by levothyroxine HIV Chronic hepatitis C HIV or lymphocytic maligancy HIV	Once or twice a week, then every 8 weeks after (Q8W)	\$31,400*	ER375mg subcutaneous*	PtA M 62% vs Week 16 PtA N 62% vs Week 16 *of PtA O responds maintained until Week 48
Biologics (Infect)	PtA Cushing Disease Hyperthyroiditis by levothyroxine Disease*	After infection Untreated leg B HIV Hyperthyroiditis by levothyroxine Disease*	Once or twice a week, then every 12 weeks (Q12W)*	\$22,000*	Anti-tumor necrosis factor ER375mg subcutaneous*	PtA P 75% vs Week 12 PtA Q 75% vs Week 12 *of PtA R responds maintained until Week 48
Biologics (Oral)	Crohn's Disease (Plus IL-17C)*	Pregnancy/Breastfeeding*	Once or twice a week, then every 12 weeks (Q12W)*	\$24,000*	Anti-tumor necrosis factor ER375mg subcutaneous*	PtA S 61% vs Week 12 (PtA T)* PtA U 61% vs Week 12 (PtA V)*



<https://www.skintherapyletter.com/psoriasis-education-tool-biologics/>

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### Drug-induced lichenoid dermatitis – treatment

1. Eliminate potential drug causes



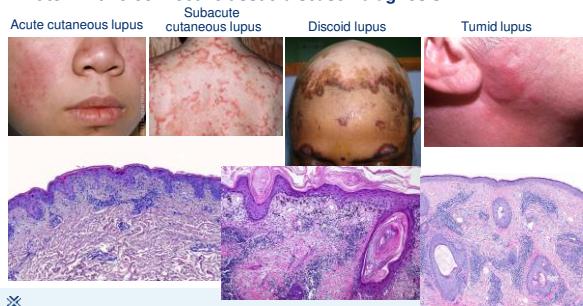
2. Topical steroids
3. Wide range of immunosuppressives



Group of drug
Antimicrobial substances
Antihistamines (H <sub>1</sub> -blocker)
Antihypertensives/antimigraines
Antimalarial drugs
Antipsychotics
Anticoagulants/antithrombotic medications
Antidiabetics
Metals
Nonsteroidal anti-inflammatory drugs
Oral contraceptives/inhalers
Lipid lowering drugs
Tumor necrosis factor alpha antagonists
Checkpoint inhibitors
Miscellanea

The bolded drugs are the ones most frequently implicated.

### Autoimmune connective tissue disease - diagnosis



### Autoimmune connective tissue disease - treatment

Management of discoid lupus erythematosus and subacute cutaneous lupus erythematosus in adults



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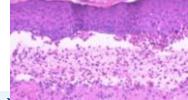
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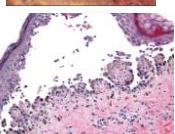
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### Autoimmune bullous dermatoses, examples - diagnosis

Bullous pemphigoid



Pemphigus vulgaris



Bullous lupus




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### Autoimmune bullous dermatoses, examples - diagnosis

Bullous pemphigoid



IgG, C3

Pemphigus vulgaris



IgG

Bullous lupus



IgG, IgA, IgM, C3 (full house)

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Likely association*	Probable association†	Uncertain association‡
• Angiotensin	• Acitretin (0)	• Atorvastatin (1-2)
• Angiotensin	• Adalimumab	• Aztreonam
• Aspirin	• Aripiprazole	• Aztreonam
• Biotin	• Aspirin (ibuprofen)	• Aztreonam
• Cetuximab	• Atazanavir	• Aztreonam
• Chagapil	• Aztreonam	• Captopril
• Cisplatin	• Azaconazole	• Celecoxib
• Everolimus	• Cefazolin	• COVID-19 mRNA vaccines
• Fenvoxade	• Chloroquine	• Dabrafenib
• Fingolimod	• Cholestyramine	• Dasatinib
• Levofloxacin	• Ciclosporine	• Desacetylpilocarpine
• Linagliptin	• Clofibrate	• Disulfiram
• Lomustine	• Clofazimine	• Doxycycline
• Pembrolizumab	• Clofazimine	• Enzalutamide
• Poxvirus	• Clofazimine	• Favipiravir
• Poxvirus with ultraviolet A	• Colchicine	• Favipiravir
• Rituximab	• Cyclosporine	• Gefitinib
• Sirolimus	• Cyclosporine	• IgM antibody
• Sitagliptin	• Cyclosporine	• Interferon
• Tacrolimus	• Cyclosporine	• Interleukin
• Tenaya tozell	• Cyclosporine	• Ivermectin
• Thalidomide	• Cyclosporine	• Irinotecan
• Vincristine	• Cyclosporine	• Ketotifen
• Zoledronic acid	• Cyclosporine	• Levetiracetam
• Zoledronic acid	• Cyclosporine	• Meloxicam
• Zoledronic acid	• Cyclosporine	• Methotrexate
• Zoledronic acid	• Cyclosporine	• Nitrofurantoin
• Zoledronic acid	• Cyclosporine	• Neostigmine (benzyl benzoate)
• Zoledronic acid	• Cyclosporine	• Ondansetron
• Zoledronic acid	• Cyclosporine	• Placebo extracts
• Zoledronic acid	• Cyclosporine	• Plasmapheresis
• Zoledronic acid	• Cyclosporine	• Raspberries
• Zoledronic acid	• Cyclosporine	• Rotavirus vaccine
• Zoledronic acid	• Cyclosporine	• Sulfasalazine
• Zoledronic acid	• Cyclosporine	• Sulfite fu vaccine
• Zoledronic acid	• Cyclosporine	• Tamoxifen
• Zoledronic acid	• Cyclosporine	• Tenofovir
• Zoledronic acid	• Cyclosporine	• Tinzaparin
• Zoledronic acid	• Cyclosporine	• Tocopherol
• Zoledronic acid	• Cyclosporine	• Tramadol
• Zoledronic acid	• Cyclosporine	• Ursodeoxycholic acid
• Zoledronic acid	• Cyclosporine	• Ustekinumab

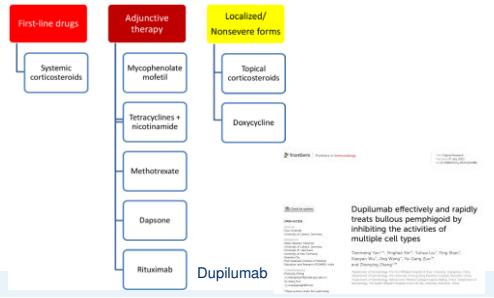
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## Treatment – autoimmune bullous disease – BP as an example

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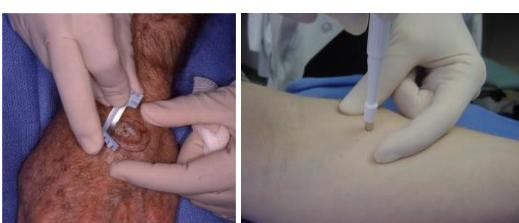
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## Basic dermatologic procedures

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### Types of Biopsies and Indications




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### Biopsy Site Selection

BIOPSY SITE SELECTION	
Lesion/disorder	Appropriate site
Tumor	Thickest portion; avoid necrotic tissue
Blister	Edge of lesion, including perilesional skin (see Fig. 0.11)
Ulcerated/necrotic lesion	Edge of ulcer or necrosis plus adjacent skin
Generalized polymorphous eruption	Characteristic lesion of recent onset ( $\pm$ more developed lesion)
Small vessel vasculitis	Characteristic lesion of recent onset

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### Patient Preparation

- Determine the type of biopsy
- Informed consent: bleeding, discomfort, infection, and scarring
- Site preparation:
  - Identification and marking
  - Time Out
  - Photograph
  - Close up for lesional details
  - Distant for identification of landmarks

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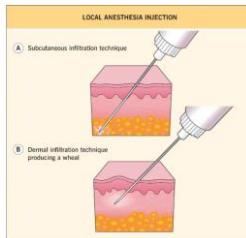
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### Anesthesia Techniques

- Lidocaine 1% with or without epinephrine
- Small lesions: direct infiltration of anesthetic into lesion
- Larger lesions: a field block by placing a ring of anesthesia around surgical site
- Bevel up
- Use small gauge needle (30), insert quickly at a 45° angle
- Slow injection to create an intradermal wheal, then may proceed to subcutaneous injection depending on shave vs. punch
- Additional sticks should be done through areas that are already numb
- Use smaller syringes – require lower pressure for injection
- Warm anesthetic to body temperature
- Slow injection
- Verbal and tactile distraction



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Bologna et al. Dermatology

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### Patient Preparation Continued

- Prep
  - ETOH swab
  - Iodine
  - Chlorhexidine
- Anesthesia
  - Plane of injection
- Procedure
  - Hemostasis: Aluminum chloride, hemostatic sponge, compression, cautery, suture, ferric subsulfate
  - Label specimen bottle with formalin



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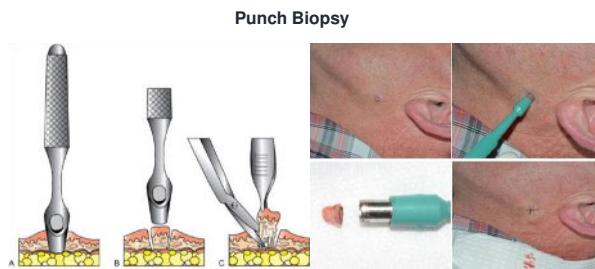
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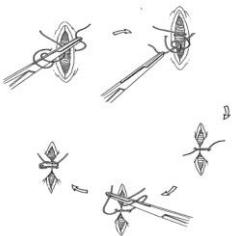
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**Instrument tie**

- Needle holder is held parallel to the wound incision
- Needle end of suture is looped twice around the holder before grasping the free end of suture
- The free and needle end of the suture exchange sides across the wound
- Additional throws are done in a similar manner, except with one loop



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**Biopsy for direct immunofluorescence**

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## CONTINUING MEDICAL EDUCATION

**Skin biopsy****Biopsy issues in specific diseases**Dale W. Braverman, MD; Mark J. Rosenzweig, MD; and Barbara J. Wilkes, MD<sup>1</sup>  
<sup>1</sup>Charlotte, North Carolina; Allergenics, Milwaukee, Wisconsin; and Bellvitge, Barcelona, Spain

Elston DM, Stratman EJ, Miller SJ.  
Skin biopsy: Biopsy issues in specific diseases.

J Am Acad Dermatol. 2016 Jan;74(1):1-16; quiz 17-8. doi: 10.1016/j.jaad.2015.06.033.  
Erratum in: J Am Acad Dermatol. 2016 Oct;75(4):854. PMID: 26702794.

**Table 1. Suspected disease entities with recommended biopsy type, site, and required laboratory tests**

Condition	Biopsy Type	Site	Test
Acne vulgaris	Punch or shave biopsy	Lesion or comedone	Direct immunofluorescence for IgA3 and IgG4 in comedones
Atopic dermatitis	Punch or shave biopsy	Lesion or normal skin	Direct immunofluorescence for IgG4 in lesional skin
Candidiasis	Punch or shave biopsy	Lesion	Microscopic examination for yeast and/or hyphae
Chronic graft-versus-host disease	Punch or shave biopsy	Lesion	Immunohistochemistry for CD30 and CD20
Collagen vascular diseases	Punch or shave biopsy	Lesion	Immunofluorescence for IgA, IgG, IgM, C3, and C4 in lesional skin
Epidermolysis bullosa	Punch or shave biopsy	Lesion	Immunofluorescence for IgG and IgA in epidermal basement membrane zone
Folliculitis decolorans trichophytinis	Punch or shave biopsy	Lesion	Microscopic examination for fungi
Hypothyroidism	Punch or shave biopsy	Lesion	Immunofluorescence for IgG4 in lesional skin
Impetigo herpetiformis	Punch or shave biopsy	Lesion	Microscopic examination for fungi
Infectious mononucleosis	Punch or shave biopsy	Lesion	Immunofluorescence for IgG and IgM in lesional skin
Linear IgA bullous dermatosis	Punch or shave biopsy	Lesion	Immunofluorescence for IgA in epidermal basement membrane zone
Lupus erythematosus	Punch or shave biopsy	Lesion	Immunofluorescence for IgG and IgM in epidermal basement membrane zone
Molluscum contagiosum	Punch or shave biopsy	Lesion	Microscopic examination for virus
Necrotizing fasciitis	Punch or shave biopsy	Lesion	Microscopic examination for bacteria
Onychomycosis	Punch or shave biopsy	Lesion	Microscopic examination for fungi
Periorificial dermatitis	Punch or shave biopsy	Lesion	Immunofluorescence for IgG4 in lesional skin
Psoriasis	Punch or shave biopsy	Lesion	Immunofluorescence for IgG4 in lesional skin
Rosacea	Punch or shave biopsy	Lesion	Microscopic examination for bacteria
Sarcoidosis	Punch or shave biopsy	Lesion	Immunofluorescence for IgG4 in lesional skin
Syphilis	Punch or shave biopsy	Lesion	Microscopic examination for spirochetes
Tinea versicolor	Punch or shave biopsy	Lesion	Microscopic examination for fungi
Vitiligo	Punch or shave biopsy	Lesion	Immunofluorescence for IgG4 in lesional skin
Wegener's granulomatosis	Punch or shave biopsy	Lesion	Immunofluorescence for IgG4 in lesional skin
Yaws	Punch or shave biopsy	Lesion	Microscopic examination for spirochetes

**Billing/coding**

Code	Description
11102	Tangential biopsy of skin (e.g., shave, scoop, saucerize, curette) single lesion
+11103	each separate/additional lesion (List separately in addition to code for primary procedure)
11104	Punch biopsy of skin (including simple closure, when performed) single lesion
+11105	each separate/additional lesion (List separately in addition to code for primary procedure)
11106	Incisional biopsy of skin (e.g., wedge) (including simple closure, when performed) single lesion
+11107	each separate/additional lesion (List separately in addition to code for primary procedure)



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**Dermatology in the Primary Care Setting**

Primary care providers are in a prime position to take care of dermatologic issues.

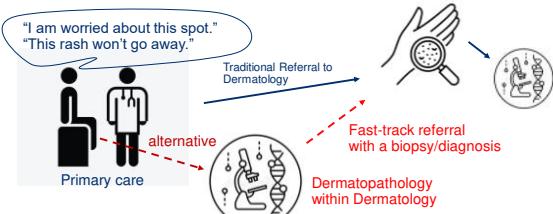


Jeffrey D. McBride, OU Dermatology, OU Dermatopathology

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## Dermatology in the Primary Care Setting

Primary care providers are in a prime position to take care of dermatologic issues.



✉ Jeffrey D. McBride, OU Dermatology, OU Dermatopathology

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Thank you for your attention.



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[Jeffrey.McBride@ouhealth.com](mailto:Jeffrey.McBride@ouhealth.com)

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