Melanoma Identification & Biopsy

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Disclosures

- None

Goals

- 1. Melanoma Overview
- 2. Melanoma Biopsy
- 3. Review staging updates per AJCC 8th edition

Melanoma



Melanoma

- Malignant transformation of Melanocytes
 - Pigment producing cells within the basal epidermis
- Most commonly in light skinned patients
- Highly correlated with UV exposure¹
- 3% of skin cancers
 - 65% of skin cancer mortality²



Clinical Detection

| Normal Mole | mal Mole Melanoma Sign | | Characteristic | | |
|-------------|------------------------|-----------|---|--|--|
| | | Asymmetry | when half of the mole does not match the other half | | |
| | | Border | when the border (edges) of the mole are ragged or irregular | | |
| | | Color | when the color of the mole varies throughout | | |
| | | Diameter | if the mole's diameter is larger than a pencil's eraser | | |

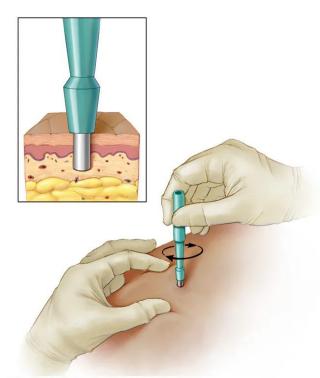
Photographs Used By Permission: National Cancer Institute

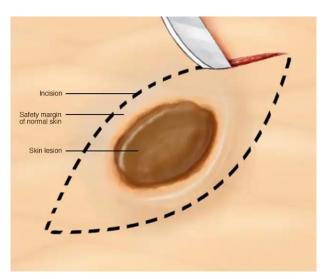
In the presence of clinical suspicion

BIOPSY

Biopsy Techniques

- 1. Excisional biopsy
- 2. Incisional biopsy
- 3. Punch Biopsy
- 4. Shave Biopsy



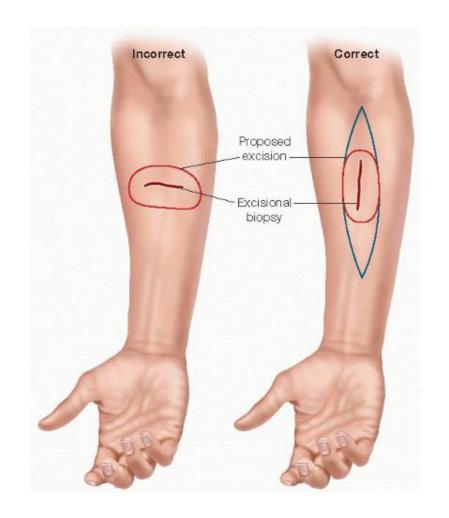


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Excisional Biopsy

- Removal of the lesion with minimal margins
- Gold standard for biopsy of pigmented lesions

- Minimize tissue disruption/undermining
- Plan resection lines for later wider resection
 - Longitudinal resection on extremities

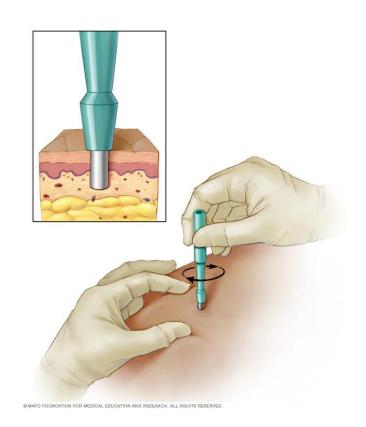


Incisional Biopsy

- A compromise
- Used if the lesion is too large for simple removal <u>OR</u>
 - Too large for your comfort level
- Select the most suspicious area for biopsy
- Still has high diagnostic value

Punch Biopsy

- A type of INCISIONAL biopsy
- Fast
- Easy
- Diagnostically valuable
- Select the thickest, most concerning portion of the lesion
- Take the biopsy into the subcutaneous fat



Punch Biopsy

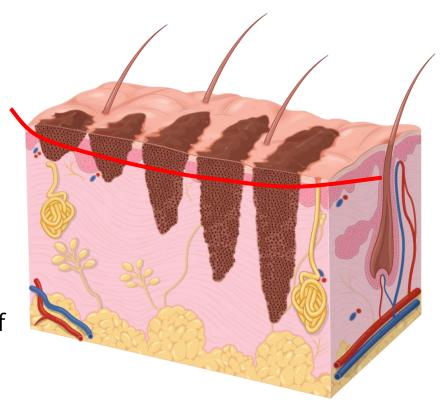
Pearls:

- 1. Use local with epinephrine. Everywhere.
- 2. Select the largest punch you can to include the whole lesion
- 3. If unable to remove lesion with punch, select at least 3-4mm



Shave Biopsy

- A tangentially excised biopsy specimen
- Usually transected in the mid dermal plane
- Irresponsible to perform on a <u>PIGMENTED</u> lesion without near certainty it is benign
- Regularly compromises staging of Melanoma **PERMANENTLY**



Biopsy Take-aways

- 1. When in doubt, take a biopsy!
 - a. Biopsy of the lesion is the fastest route to appropriate management
- 2. Excisional biopsy is best, but punch biopsy is acceptable
- 3. NEVER shave a pigmented lesion with any diagnostic uncertainty

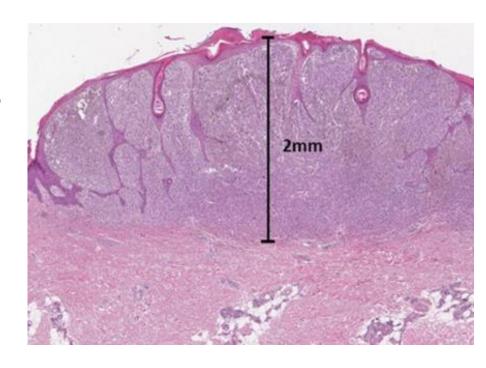
Preoperative Imaging

- Patients with Mis and clinically node negative disease do not require preoperative imaging³
- Consideration should be given to nodal assessment in equivocal Lymph node basin exams (ie/ morbidly obese patients)

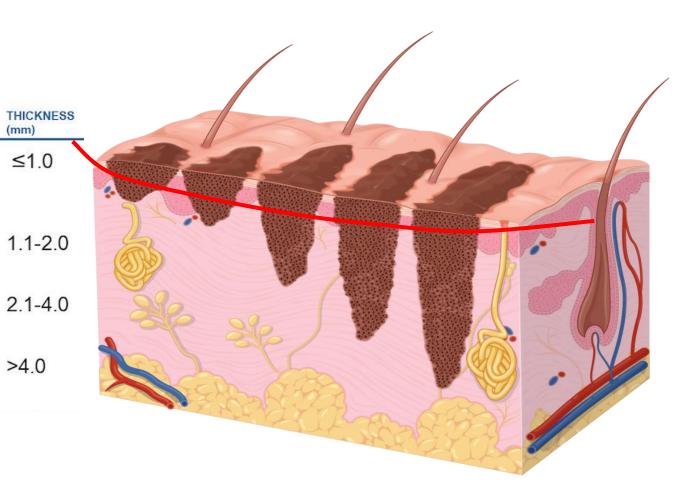
Staging - AJCC 8th Edition

Primary Tumour - T

- Based upon *Breslow Depth* and presence or absence of *Ulceration*
- Breslow Depth
 - Vertical depth of tumour extension on microscopy
 - Increases in initial depth correlate with increased risk of metastasis¹



| T CLASSIFICATION | THICKNESS (mm) | ULCERATION STATUS |
|---------------------|-------------------|--|
| T1 | ≤1.0 | a: Breslow < 0.8 mm w/o ulceration b: Breslow 0.8-1.0 mm w/o ulceration or ≤ 1.0 mm w/ ulceration. |
| T2 | 1.1-2.0 | a: w/o ulceration b: w/ ulceration |
| T3 | 2.1-4.0 | a: w/o ulceration b: w/ ulceration |
| T4 | >4.0 | a: w/o ulceration b: w/ ulceration |



CLASSIFICATION

T2

T3

T4

Regional Lymph Nodes - N

Regional Lymph Nodes (N)

NX Patients in whom the regional nodes cannot be assessed (for example previously removed for another reason)

No regional metastases detected

N1-3 Regional metastases based on the number of metastatic nodes, number of palpable metastatic nodes on clinical exam, and presence or absence of MSI²

NOTE: N1-3 and a-c subcategories assigned as shown below:

Distant Metastasis - M

Distant Metastasis (M)

M0 No detectable evidence of distant metastases

M1a Metastases to skin, sub cutaneous, or distant lymph nodes

M1b Metastases to lung

M1c Metastases to all other visceral sites

M1d Metastases to brain

Staging

| ANATOMIC STAGE/PROGNOSTIC GROUPS | | | | | | | |
|----------------------------------|-------|-------|---------------------|------|--------|--------|-------------|
| Clinical Staging ³ | | | Pathologic Staging⁴ | | | | |
| Stage 0 | Tis | N0 | MO | 0 | Tis | N0 | M0 |
| Stage IA | T1a | N0 | M0 | IA | T1a | N0 | MO |
| Stage IB | T1b | | | | T1b | | |
| | T2a | ** | •• | IB | T2a | •• | •• |
| Stage IIA | T2b | N0 | M0 | IIA | T2b | MO | M0 |
| | T3a | | | | T2a | | |
| Stage IIB | T3b | | | IIB | T3b | | |
| | T4a | | | | T4a | | |
| Stage IIC | T4b | •• | | IIC | T4b | | •• |
| Stage III | Any T | ≥N1 | M0 | IIIA | T1-2a | N1a | M0 |
| | | | <u></u> | | T1-2a | N2a | |
| | | | | IIIB | T0 | N1b-c | M0 |
| | | | | | T1-2a | N1b-c | |
| | | | | | T1-2a | N2b | ··· |
| | | | | | T2b-3a | N1a-2b | |
| | | | | IIIC | T0 | N2b-c | M0 |
| | | | | | T0 | N3b-c | |
| | | | | | T1a-3a | N2c-3c | ··· |
| | | | | | T3b-4a | Any N | |
| | | | | | T4b | N1a-2c | |
| | | | | IIID | T4b | N3a-c | M0 |
| Stage IV | Any N | Any N | M1 | IV | Any T | Any N | M1 |

Staging

- 1. Stage I Stage II disease
 - a. Based EXCLUSIVELY on the Breslow depth and ulceration

| ANATOMIC STAGE/PROGNOSTIC GROUPS | | | | | | | |
|----------------------------------|-------|--------------|-------|---------------------|---------|--------|---------------|
| Clinical Staging ³ | | | | Pathologic Staging⁴ | | | |
| Stage 0 | Tis | N0 | MO | 0 | Tis | N0 | M0 |
| Stage IA | T1a | N0 | M0 | IA | T1a | N0 | M0 |
| Stage IB | T1b | | | | T1b | | |
| | T2a | | •• | IB | T2a | •• | •• |
| Stage IIA | T2b | N0 | MO | IIA | T2b | MO | M0 |
| | T3a | | | | T2a | | |
| Stage IIB | T3b | | | IIB | T3b | | |
| | T4a | | | | T4a | | |
| Stage IIC | T4b | •• | | IIC | T4b | | •• |
| Stage III | Any T | ≥N1 | M0 | IIIA | T1-2a | N1a | M0 |
| | | | | | T1-2a | N2a | |
| | | | | IIIB | T0 | N1b-c | M0 |
| | | . . . | | | T1-2a | N1b-c | <mark></mark> |
| | | | T1-2a | N2b | | | |
| | | | | | T2b-3a | N1a-2b | |
| | | | | IIIC | T0 | N2b-c | M0 |
| | | | | | T0 | N3b-c | |
| | | | | | T1a-3a | N2c-3c | |
| | | | | | T3b-4a | Any N | |
| | | | | | T4b | N1a-2c | |
| | ** | | | IIID | T4b | N3a-c | M0 |
| Stage IV | Any N | Any N | M1 | IV | Any T | Any N | M1 |

Staging

- 1. Stage I Stage II disease
 - a. Based EXCLUSIVELY on the Breslow depth
- 2. Stage II vs. Stage III/IV
 - a. Presence of metastasis

| ANATOMIC STAGE/PROGNOSTIC GROUPS | | | | | | | |
|----------------------------------|-------|----------------|---------------------|------|--------|--------|-----|
| Clinical Staging ³ | | | Pathologic Staging⁴ | | | | |
| Stage 0 | Tis | N0 | MO | 0 | Tis | N0 | M0 |
| Stage IA | T1a | N0 | M0 | IA | T1a | N0 | MO |
| Stage IB | T1b | | | | T1b | | |
| _ | T2a | | •• | IB | T2a | •• | •• |
| Stage IIA | T2b | N0 | MO | IIA | T2b | M0 | MO |
| | T3a | | | | T2a | | |
| Stage IIB | T3b | | | IIB | T3b | | |
| | T4a | | | | T4a | | |
| Stage IIC | T4b | | | IIC | T4b | | |
| Stage III | Any T | ≥N1 | M0 | IIIA | T1-2a | N1a | MO |
| | | ·- | | | T1-2a | N2a | |
| | | . . | | IIIB | T0 | N1b-c | MO |
| | | | | | T1-2a | N1b-c | |
| | | | ··· | | T1-2a | N2b | ··· |
| | | | | | T2b-3a | N1a-2b | |
| | | | | IIIC | T0 | N2b-c | MO |
| | | | ··· | | T0 | N3b-c | |
| | | | | | T1a-3a | N2c-3c | |
| | | | ··· | | T3b-4a | Any N | |
| | | . | | | T4b | N1a-2c | |
| | ** | *** | | IIID | T4b | N3a-c | MO |
| Stage IV | Any N | Any N | M1 | IV | Any T | Any N | M1 |

Surgical Management

Resection Margins

| Breslow thickness | Additional clinical margin | | | |
|--|----------------------------|--|--|--|
| Naevus with severe cytological or architectural atypia | 5mm | | | |
| Melanoma in situ (Tis) | 5–10mm | | | |
| <1.0mm (T1) | 10mm | | | |
| 1-2mm (T2) | 10-20mm | | | |
| 2-4mm (T3) | 20mm | | | |
| >4mm (T4) | 20mm | | | |

Sentinel Lymph Node Biopsy

Sentinel Lymph Node

- The first lymph node(s) within the lymphatic basin reached by lymph draining from the primary lesion⁴
- Complete nodal staging can be obtained by focused sampling of the sentinel lymph node alone

Sentinel Lymph Node Biopsy

 Surgical technique that employs preoperative lymphoscintigraphy to facilitate surgical biopsy of the sentinel lymph nodes without complete lymphadenectomy

Sentinel Lymph Node Biopsy - Who gets one?

T1a

- <0.8mm tumour</p>
- Statistical risk of positive biopsy <5%³
- NOT typically recommended

T₁b

- 0.8mm-1.0mm tumour
- OR <1.0mm *with* ulceration

- Risk of positive biopsy 5-10%³
- Sentinel Node Biopsy recommended

Adjuvant Therapy

- Will be covered next by Dr. Mathews

Candidates:

- 1. Stage III or IV disease
 - a. Ie/ Nodal or distant metastasis
- 2. Locally advanced disease
 - a. Stage IIb/IIc
 - i. Based exclusively on breslow depth

Key Points

- 1. Melanoma is a relatively rare but Do-Not-Miss diagnosis
- 2. Early biopsy is critical
 - a. Results in most expeditious and accurate management
 - b. Surgical and adjuvant therapy decisions CANNOT be made until biopsy result is available
- 3. Full thickness biopsies are best
 - a. Excisional or punch

References

- 1. Amin MB, Edge SB, Greene FL, et al., eds. AJCC Cancer Staging Manual. 8th ed.New York: Springer, 2017
- 2. Naik, P. P. (2021). Cutaneous Malignant Melanoma: A review of Early Diagnosis and Management. *World Journal of Oncology, 12*(1), 7–19. https://doi.org/10.14740/wjon1349
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- 4. Morton, D. L., Cochran, A. J., Thompson, J. F., Elashoff, R., Essner, R., Glass, E. C., Mozzillo, N., Nieweg, O. E., Roses, D. F., Hoekstra, H. J., Karakousis, C. P., Reintgen, D. S., Coventry, B. J., & Wang, H. (2005). Sentinel node biopsy for Early-Stage melanoma. *Annals of Surgery*, *242*(3), 302–313. https://doi.org/10.1097/01.sla.0000181092.50141.fa

Questions?