

- Did the patient receive a prescription for opioid / narcotic pain medication (prescription prior to or at the time of surgical discharge from the Mohs surgeon) following Mohs micrographic surgery?
- If yes, Did the patient have one of the following reasons for prescription of opioid / narcotic pain medication?

Documented medical comorbidity(ies) which preclude the use of non-opioid analgesics and have been advised by physicians to avoid them (advanced renal dysfunction, advanced liver dysfunction, or history of bleeding peptic ulcer)

Documented allergy to non-opioid analgesics

Patient required additional pain relief despite a trial of non-opioid analgesia

None of the above

Tumor Characteristics Tab	Type of Tumor: Pre-op diag	gnosis- BCC SC	CC Melanoma	Rare tumors			
If BCC: Subtype- (Circle all that apply)							
Unspecified/missing	Superficial	Superficial Nodular		icronodular			
Infiltrative	Morpheaform	Pigment		d Sclerosing			
Adenosystic	Basosquamous	Occurring in a prior radiation field		n field			
High risk tumor	Other Specify)						
If SCC: Subtype- In situ	Well-differentiated	,					
Moderately-differentiated	Poorly-differentiate	d None spe	ecified/unknown				
If poorly-differentiated SC	C, type-						
Spindled Acantholy	rtic Desmoplastic	Adenoid/adeno	squamous (mucin-p	producing)			
If SCC high risk feature-Per	ineural/intraneural invasion	Lymphovascular	rinvasion				
Invasion to cartilage, muscl		Breslow depth >2mm Palpable lymph node					
Occurring in a prior radiation	on field	High risk tumor	(Go to additional w	vork-up)			
• If Perineural/			Dath N				
 Biopsy N Y Nerve size for biops 		Mohs N Y Mohs _>.1 mm	Both N <.1 mm	1			
If Lymphovas	cular invasion- biopsy	Mohs					
• If Invasion to	cartilage, muscle or bone-	biopsy	Mohs				
 If Breslow dej 	oth >2mm	biopsy	Mohs				
If Melanoma: Subtype- In situ	invasive	Breslow	/ depthn	nm			
Melanoma high risk feature	es- Ulceration	ceration Mitotic figures>		>1 mm2 Perineural invasion			
Lymphovascular invasion	Palpable lymph node(s)	High risk Tumo	r Non	e of above			
If Rare tumor: Subtype-	Adenocystic carcino	oma	Adnexal carcinoma				
Angiosarcoma	Apocrine/eccrine C		Atypical Fibroxanthoma				
Dermatofibrosarcoma Protuber	ans Desmoplastic tricho	pepithelioma	Extramammary P	Paget's Disease			

Leiomyosarcoma	Malignant Fibrous Histocytoma	Merkel Cell Carcinoma
Microcystic Adnexal Carcinoma	Mucinous Carcinoma	Porocarcinoma
Sebaceous Carcinoma	Undifferentiated Pleomorphic Sarcom	а

• If Leiomyosarcoma:

Primary dermal leiomyosarcoma Subcutaneous leiomyosarcoma

Surgical site main area- If tumor spans multiple areas, select the predominant area.

Cutaneous lip	Vermilion lip	Eyelid including canthus
Eyebrow	Forehead (non-eyebrow region)	Ear and external auricular canal
Nose	Temple	Cheek (including jawline)
Chin	Neck	Scalp
Hand	Upper limb (incl. shoulder, not hand)	Foot (including ankle)
Pretibial shin	Lower limb (incl. hip, not including fee	et or pretibial shin)
Nipple/areola	Trunk (excluding nipple/areola)	Anogenital

- If tumor is SCC and subtype is "in situ" including SCC and site is either "cutaneous lip, vermillion lip, eyebrow, forehead, ear and external auditory canal, nose, temple, cheek, chin, neck, or scalp: Does this tumor meet America Joint Committee on Cancer (AJCC) 8th edition staging as a tumor stage greater than or equal to T2
 - Yes No
 - If yes to meeting greater than or equal to T2, what was the tumor stage?
 T2 T3 T4a T4b
 - If T3, what is/are the defined T3 clinical characteristic(s)? (check all that apply)

Tumor >4cm in greatest diameter

Tumor > 6mm in depth from adjacent granular layer or beyond subcutaneous fat

Perineural invasion (Clinical or radiographic involvement of named nerve, Subdermal nerves, Nerve caliber >0.1mm

Minor bone erosion

Was the AJCC 8th edition tumor staging documented in the medical record Yes No

0	Side of lesion-	Right	Left	Midline	e Unknown			
0	Preop length _		_cm Preop wi	dthcm				
0	Is this tumor-	primary	Previous	ly treated				
	If Previously treated:							
	Incompletely treated (treated surgically with positive margins) Recurrent							
	Treated preoperatively to reduce tumor size using a systemic therapy							
	If recurrent how was the tumor previously treated (check all that apply)-							
	Curettage and Electrodessication Excision			Mohs Surgery				
	Radiat	ion		Superficial Brachyth	erapy			
	Cryotherapy or Cryosurgery (not including empiric) Targeted Topical Treatment (not including general field therapy for actinic keratosis)							
	Photodynamic Therapy (not including generalized field PDT for actinic keratosis)							
	Syster	nic therapy	,	Other	Unknown			

If treated pre-operatively to reduce tumor size with systemic therapy, type-Hedgehog inhibitor CTLA-4inhibitor (ipilimumab) PD-1 inhibitor EGFR-inhibitor Capecitabine Platinum-based chemo Other systemic therapy

- Has the lesion in question been confirmed to have DIFFERENT histology to the previously treated tumor? (i.e., histology confirms BCC and BCC was treated in the past)- N Y Unknown
- Is lesion in question contiguous with surgical scar after treatment of previous tumor? (i.e. inside the greatest radius of final defect measured from the center of the closure)- N Y Unclear
- Is lesion within the area of previous tumor or defect prior to reconstruction- N Y Unclear If Yes, list therapies: Hedgehog inhibitor PD-1 inhibitor EGFR-inhibitor Other systemic therapy If Other type of previous treatment, specify-_____

What is Mohs surgery Appropriate Use Criteria score-1 2 3 4 5 6 7 8 9 undefined

Mohs Surgery Tab

Mohs Surgery Tab

- Post-op length _____cm
 Post-op width ____cm
 # of Mohs stages 1
 2
 3
 4
 5
 6
- Number of CPT 17315- _____
- What features were seen on the Mohs stage-
 - SCC
 BCC
 Melanoma
 All other tumor types
 No CA seen

 •
 Were immunohistochemical stains were used on frozen sections N
 Y

 •
 What immunohistochemical stains were used Sox-10

Cytokeratins (CK-pan AL1/AL5		Del-Lp-4	Ivial (=1	201-10	
	HMB-45	MITF	MEL-5	S100	
	СК-7	CEA	CD34	CK-17	
a normanant sostions cont? (Includes frazen debulking er Mahs specimens thawed for					

 Were permanent sections sent? (Includes frozen debulking or Mohs specimens thawed for permanent sectioning, or additional margins taken for permanents.
 N Y

If sent, why- To evaluate a debulking specimen To confirm final margin To allow for special stains For tumor staging Other
 Is this a complex case? N Y

Reconstruction Tab

0

Was the tumor defect reconstructed-						Ν	Y	
•	If Yes, was the	e reconstruction	performed b	y the sa	ame Mohs s	urgeon-	Ν	Y
	or another M	ohs surgeon with	nin the same	practic	e-		Ν	Y
•	If No, what type of surgeon reconstructed the tumor-							
	A different Mo	ohs/Derm		Plastic	2		Oculo	plastic
	Otolaryngolog	gy/Head and Nec	k including	ENT Fa	cial Plastics		Gene	ral
	Other (specify)						Unknown	
•	When was the tumor reconstructed- same day delayed						Unkn	own
•	 Type of reconstruction performed- 							
	Pursestring		Linear			Flap		
	Grafts		Unknown					
•	If Linear speci	fy-						
	Simple	_cm	Intermed_		_cm	Complex_		_cm
	Cheiloplasty							

Complications Tab

NOTE: Please add complications under this tab when/if they are discovered. Thirty days post-op MohsAlQ will have a "Complications Needed" flag after every patient, if no complications have already been added. If there are no complications at the 30-day mark, click on the flag and provide the appropriate information. This is an important step as this is part of the performance measure calculation.

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