

VERSION 2/6/2021

Case Demo	graphics entered by	Case details entered by	/ 30 day F/U er	ntered
	ACMS MohsAle	Q Registry Workshe	et Required Fields	
Surgeon plo	ease circle/complete approp	oriate elements. Information	n will be used to complete	case in MohsAIO
0		Last DOB		
Case Tab	Was this patient referred have access to the patient	Mohs Surgeon_ from a provider outside you 's electronic health record-	ır practice or by a provider	
o	a physician? If yes, Was the a	cription anticoagulant(s) an nticoagulation regiment dis or reduced perioperatively?	scontinued (including dosa	N Y
	• Disconti	nued Changed If Discontinued, Changed, discontinued, changed or i	= = = = = = = = = = = = = = = = = = =	anticoagulant
		Continuation of perioperat dangerous or because the supratherapeutic INR	-	
		Patient chose to stop thera recommendation	apy on their own or by othe	r physician
		documentation discontinuation anticoagulation No	·	orthopedic "Is there of discussion of
		Yes The patient does anticoagulant m	s not currently have a phys	ician managing
0	Were antibiotics given on If yes, Type of an	uidelines for endocarditis on the day of surgery-	or orthopedic prophylaxis- In anesthesia	N Y N Y Post-op
	Endocar		Wound infection	Other

	evidence of infection at the surgical site at time of reconstruction?					
	No Yes Antibiotics were prescribed by another physician 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? N Y 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	f the above				
Tumor Cha	aracteris	tics Tab T	ype of Tumor: Pre-op	diagnosis-	BCC SCC	Melanoma Rare tumors
If BCC: Subty Unspecified/	=	(Circle all tha	t apply) Superficial		Nodular	Micronodular
Infiltrative Adenosystic High risk tum	nor		Morpheaform Basosquamous Other Specify)		Pigmented Occurring in a p	Sclerosing orior radiation field
If SCC: Subty Moderately-0			Well-differentiat Poorly-differenti		Well-differentia	ated, keratoacanthoma /unknown
If poorly Spindled		ntiated SCC, t Acantholytic	= =	: Adeno	oid/adenosquam	ous (mucin-producing)
If SCC high risk feature-Perineural/intraneural invasion Invasion to cartilage, muscle or bone Occurring in a prior radiation field			Breslo	Lymphovascular invasion Breslow depth >2mm Palpable lymph node High risk tumor (Go to additional work-up)		
		erineural/int	N Y Nerve size for biopsy	Mohs Mohs	N Y _>.1 mm	Both N Y <.1 mm
If Lymphovascular invasion - biopsy Mohs						
		reslow depth	rtilage, muscle or bon 1 >2mm	e - biopsy biopsy	Mohs Mohs	
If Melanoma	: Subtyp	e- In situ	invasive		Breslow depth	mm
Melanor Lymphov	_	risk features- invasion	Ulceration Palpable lymph node		tic figures>1 mm risk Tumor	Perineural invasion None of above

If Post-op, Did the surgery involve breach of the oral, nasal, genitourinary or anal mucosa; area of lymphedema; exposed cartilage/bone or clinical

If Rare tumor: Subtype-Adenocystic carcinoma Adnexal carcinoma Angiosarcoma Apocrine/eccrine Carcinoma Atypical Fibroxanthoma Dermatofibrosarcoma Protuberans Desmoplastic trichoepithelioma Extramammary Paget's Disease Malignant Fibrous Histocytoma Merkel Cell Carcinoma Leiomyosarcoma Microcystic Adnexal Carcinoma Mucinous Carcinoma Porocarcinoma Sebaceous Carcinoma Undifferentiated Pleomorphic Sarcoma If Leiomyosarcoma: Primary dermal leiomyosarcoma Subcutaneous leiomyosarcoma Surgical site main area- If tumor spans multiple areas, select the predominant area. Vermilion lip Eyelid including canthus Cutaneous lip 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 feet 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? 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 Side of lesion- Right Midline Unknown Left Preop length Preop width cm cm Is this tumor- primary Previously 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** Radiation Superficial Brachytherapy Cryotherapy or Cryosurgery (not including empiric)

	Targeted Topical Treatn	nent (not including general fie	ld therapy for actinic keratosis)	
	Photodynamic Therapy	(not including generalized fiel	-	
	Systemic therapy	Other	Unknown	
		ely 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 th	e past)- N Y Unknown	
•	Is lesion in question contiguous	with surgical scar after treatme	ent of previous tumor? (i.e. inside	
	the greatest radius of final defec	t measured from the center o	f the closure)- N Y Unclear	
•	Is lesion within the area of previous	ous tumor or defect prior to re	econstruction- N Y Unclear	
	-	·	R-inhibitor Other systemic therapy	
	If Other type of previou		· · · · · · · · · · · · · · · · · · ·	
	What is Mohs surgery Appropriat	e Use Criteria score-1 2 3	4 5 6 7 8 9 undefined	
Mohs Surgery Tab				
	urgery Tab			
0	Post-op lengthcm Post-o	op widthcm # of Mohs	stages 1 2 3 4 5 6	
0	Number of CPT 17315-			
0	What features were seen on the	-		
		anoma All other tumo		
		mical stains were used on fro	zen sections- N Y	
		mical stains were used-		
	Cytokeratins (CK-pan Al	·	Mart-1 Sox-10	
	HMB-45	MITF	MEL-5 \$100	
	CK-7	CEA	CD34 CK-17	
0	Were permanent sections sent?			
	permanent sectioning, or addition			
	If sent, why- To evaluat		To confirm final margin	
	To allow for special stai	=	ing Other	
	 Is this a complex case (A 	Associate Members Only)?	N Y	
Reconstru	action Tab			
0	Was the tumor defect reconstru	cted-	N Y	
	If Yes, was the reconstruction	uction performed by the same	Mohs surgeon- N Y	
	or another Mohs surge	on within the same practice-	N Y	
If No, what type of surgeon reconstructed the tumor-				
	A different Mohs/Derm		Oculoplastic	
	Otolaryngology/Head a	nd Neck including ENT Facial		
	Other (specify)	<u> </u>	Unknown	
	When was the tumor re	constructed- same day	delayed Unknown	
	 Type of reconstruction 	•	•	
	Pursestring	Linear	Flap	
	Grafts	Unknown	•	

•	If Linear specify-						
:	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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