

# Chondromyxoid Fibroma of the Skull Base and Calvarium: Surgical Management and Literature Review

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J Neurol Surg Rep 2016;77:e23–e34.

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## Abstract

Chondromyxoid fibroma (CMF) is an exceedingly rare tumor that represents less than 1% of all primary bone neoplasms. Occurrence in the facial and cranial bones is extremely rare and frequently misdiagnosed.

**Case Reports** We report two cases of CMF, one in the sphenoclivus skull base and the other involving the parietal bone in two young female patients. Excision was performed in both cases. Presenting symptoms, treatment, and follow-up are reported.

**Methods** A retrospective review of the literature on cranial CMF was performed. The location, demographics, presenting symptoms, and treatment of all calvarial and skull base CMF cases published since 1990 are summarized.

**Discussion** In our literature review, we found 67 published cases of cranial CMF. Mean age of all calvarial and skull base CMFs at diagnosis was 38.2 years old. Of the cases affecting the cranium, the sinonasal structures were most commonly involved. To our knowledge we report only the second case of CMF involving the parietal bone published in an English-language journal. Total resection is the best treatment, and should be the goal of surgical intervention. Curettage results in high recurrence rates. Radiotherapy in the setting of subtotal resection or recurrence cannot be definitively recommended and needs further investigation.

## Keywords

- ▶ benign
- ▶ bone neoplasms
- ▶ cartilage
- ▶ calvarium
- ▶ skull base

## Introduction

Chondromyxoid fibroma (CMF) is an exceedingly rare tumor that represents less than 1% of all primary bone neoplasms.<sup>1</sup> First described by Jaffe and Lichtenstein in 1948,<sup>2</sup> CMFs need to be distinguished from other aggressive cartilaginous tumors that have significantly different treatments and prognoses. CMF is a benign tumor characterized by lobules of spindle-shaped or stellate cells with abundant myxoid or chondroid intercellular material with a varying number of multinucleated giant cells of different sizes.<sup>3</sup> Most frequently, it is found in young adults of the second and third decades of life in the lower extremity long bones, particularly arising from the metaphysis.<sup>4</sup> CMF can also arise in numerous other

anatomic sites. Its occurrence in the facial and cranial bones is extremely rare.

CMF of the cranial bones is an exceedingly challenging diagnosis to make.<sup>5,6</sup> Zilmer and Dorfman report an initial misdiagnosis rate of 22% in their series of 36 CMF cases.<sup>7</sup> Depending on its location, CMF can be difficult to distinguish from an aneurysmal bone cyst, fibrous dysplasia, giant cell tumor, osteoblastoma, osteosarcoma, Ewing sarcoma, mucocele, Langerhans histiocytosis, or even a schwannoma.<sup>5,8,9</sup> More often, CMF is mistaken for three other myxoid tumors: chordoma, chondroid chordoma, and chondrosarcoma that have a greater frequency of occurrence in the craniofacial skeleton.<sup>10,11</sup>

received  
August 15, 2015  
accepted  
October 21, 2015  
published online  
January 4, 2016

DOI <http://dx.doi.org/10.1055/s-0035-1570033>.  
ISSN 2193-6358.

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Here we update the total English literature count to 59 cases of CMF arising from the skull base, including our case of sphenoclivical CMF. Additionally, to our knowledge we report the second case of CMF involving the parietal bone published in an English-language journal.

### Case 1

A 38-year-old right-handed woman presented with a 1.5-year history of nasal obstruction and serous rhinorrhea. Approximately 1 month prior to her visit, she also noticed diplopia. A computed tomographic (CT) scan and magnetic resonance imaging (MRI) of the paranasal sinuses were obtained, which showed an expansile lesion of the central skull base, in the sphenoclivical region, with extension into the left infratemporal fossa (→**Fig. 1**). An endoscopic biopsy of the mass was obtained and an initial diagnosis of a malignant fibrohistiocytoma was made. On further review, the pathology was interpreted as showing a “Fibro-mixoid tumor, locally aggressive, and with unknown metastatic potential.” This patient’s case was discussed at our Joint Planning Conference and the recommended treatment was surgery.

The patient underwent a transmaxillary approach to the anterior cranial fossa with resection of the extradurally

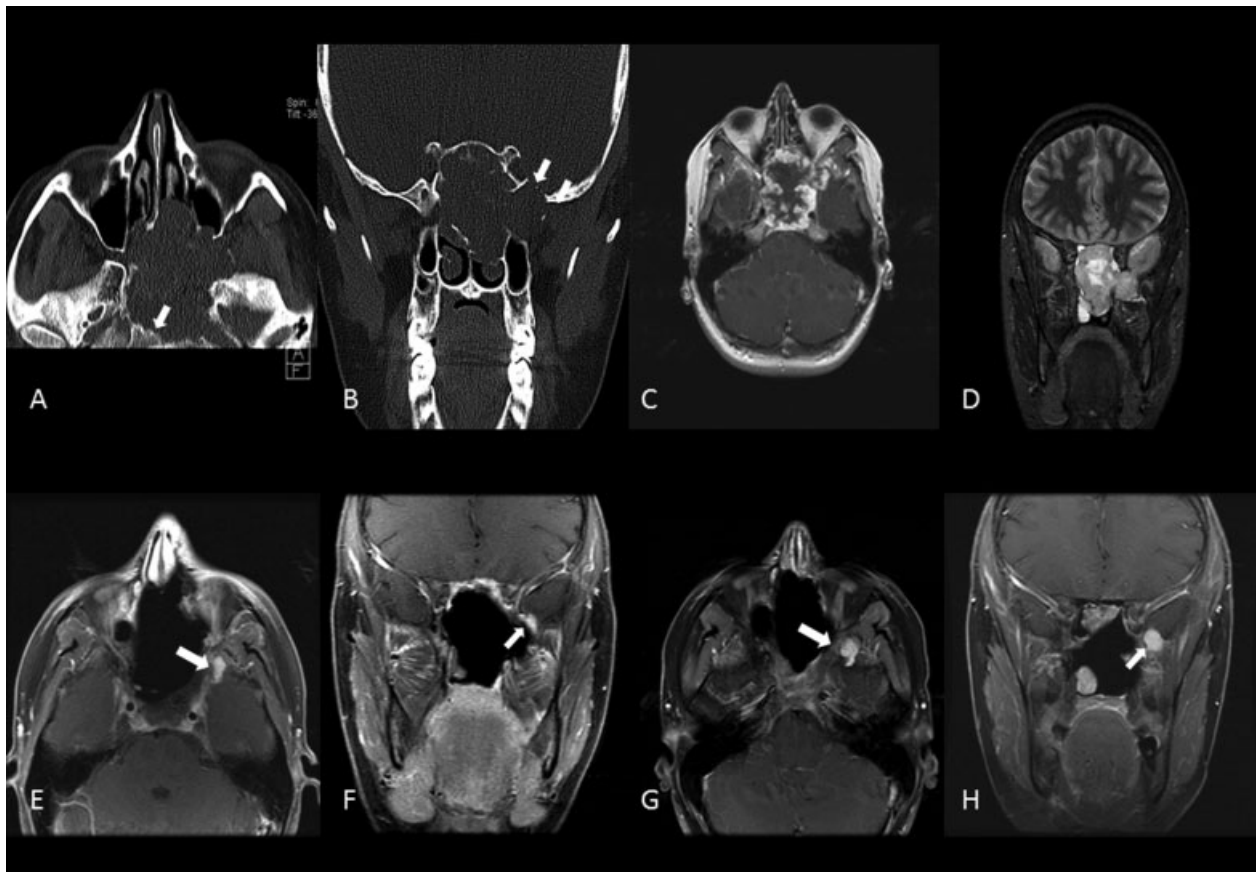
located tumor. Postoperative course was uneventful and the patient was discharged on postoperative day 4.

On follow-up, the patient reported left facial numbness in the V2 distribution and postoperative MRI showed residual tumor at the left lateral portion of the pterygopalatine fissure. The final pathology report was CMF. The minimal residual disease was followed with serial MR imaging.

Over the ensuing year, the patient’s tumor showed slow but progressive growth. Surgical resection of the progressive residual sphenoclivical CMF was performed. She underwent a left orbitocranial approach to middle cranial fossa with resection of the tumor. Postoperative course was uneventful and the patient was discharged on postoperative day 2. She is disease free at 4.5 years.

### Case 2

A 31-year-old right-handed woman presented with a 1-year history of brief, mild, left-sided headache. Approximately 3 months prior to her visit, she noticed a lump and tenderness over the left parietal area. As the tenderness increased, she ultimately underwent MR imaging. This revealed the presence of a large lesion centered within the diploe of the parietal bone. There was no intradural



**Fig. 1** Case 1. Preoperative axial (B) and coronal (B) CT scans identify this destructive central skull base mass. There is erosion of the clivus (arrow in A) and of the middle fossa floor (arrow in B). Preoperative pos-contrast axial T1-weighted MRI (C) and coronal T2-weighted MRI (D) reveal findings typical for most “chondroid” tumors. Low signal intensity with homogeneous enhancement on T1-weighted postcontrast imaging and high signal intensity on T2-weighted imaging. Images (E) and (F) identify residual disease following the patient first surgery (arrows). Follow-up MRI imaging (G) and (H) reveal progressive enlargement of the residual tumor (arrows) prompting further surgical management.

extension. At the time of presentation, the patient was in her third trimester of pregnancy, so treatment was postponed to follow her delivery. Subsequent high-resolution CT and MRI images showed a stable tumor, and the patient elected for surgery (►Fig. 2).

She underwent resection of the left parietal calvarial lesion and reconstruction with a custom cranial implant. Postoperative course was uneventful and the patient was discharged on postoperative day 3. Her pathology report indicated left parietal calvarial CMF in bone and 3 month follow-up CT showed no evidence of recurrent or residual tumor. She is disease free at 5.5 years.

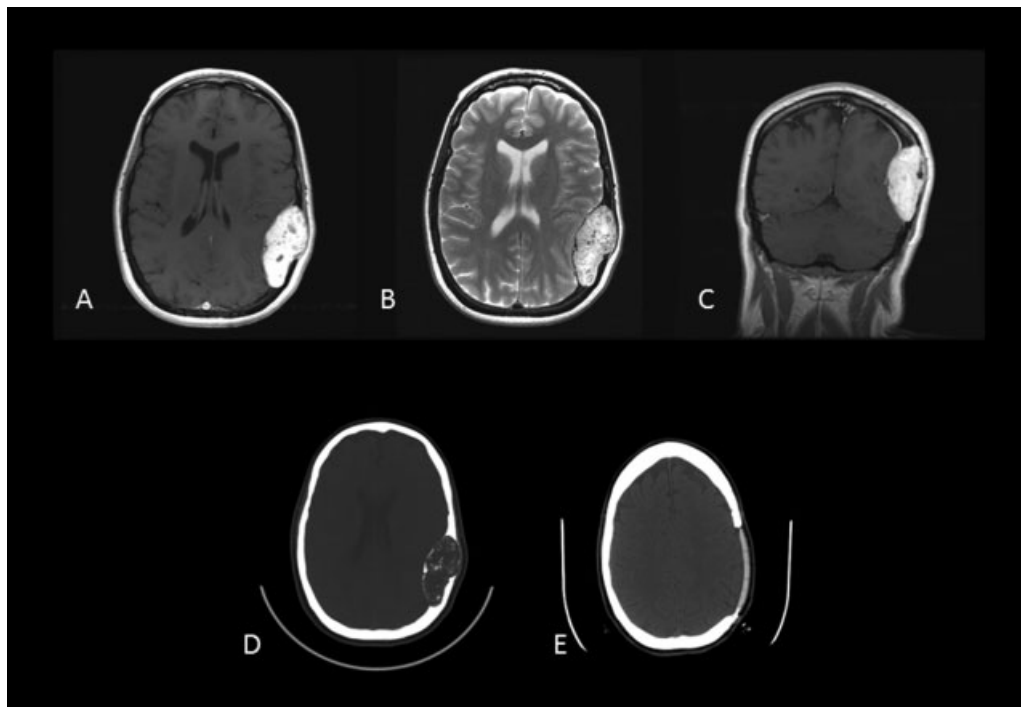
## Methods

A retrospective literature review was performed. Pubmed search terms included “chondromyxoid fibroma” with English-language filter. Publication dates were limited to January 1, 1990 to April 21, 2013. Search was secondarily refined by article type, case report, and tumor location, calvarial and skull base. Skull base tumors were further categorized into sinonasal, clival/sellar, sphenoid/parasellar, orbit/zygoma, or temporal bone/occiput. All included articles were analyzed and relevant information extracted for the table. Tumor locations and clinical symptoms were standardized for ►Table 1. This study was conducted with approval from The University of Texas M. D. Anderson Cancer Center Institutional Review Board (IRB) under protocol PA14-0505.

## Discussion

CMF is an exceptionally rare tumor, primarily affecting the long bones and accounting for less than 1% of all osseous neoplasms.<sup>1</sup> Its occurrence in the craniofacial skeleton is even less frequent. Wu et al, in their review of 278 cases, reported only 15 tumors involving the skull and facial bones.<sup>4</sup> The location, demographics, presenting symptoms, and treatment of all of calvarial and skull base CMF cases published since 1990 are summarized in ►Table 1.

In our literature review, we have found 67 published cases of cranial CMF since 1990. The mean age of all calvarial and skull base CMFs is 38.2 years, which is consistent with the literature.<sup>4</sup> Patients with clival/sellar and sphenoid/parasellar sites of origin were, on average, a decade older than those with other sites of origin. Ages ranged from newborn to 73 years. Additionally, we found a slightly greater predilection for CMF in females, with 35 females and 24 males. This is conflicting in the literature with some studies reporting a slight male predilection and others a 2:1 female-to-male occurrence. We found a slight male predominance in the temporal bone/occipital site subgroup. Of the cases affecting the cranium, the sinonasal structures were affected most commonly, with the second most common tumor location being the temporal bone and occiput. Most tumors rarely affected a single bone, and therefore appeared to grow without respect to the bony anatomy by involving multiple surrounding bones. Most patients were symptomatic at the time of presentation. The insidious onset of symptoms can



**Fig. 2** Case 2. Preoperative axial postcontrast T1-weighted MRI (A), axial T2-weighted MRI (B), and coronal postcontrast T1-weighted MRI reveal a large tumor of the parietal bone with its epicenter in the diploe of the skull. The typical pattern of inhomogeneous enhancement on T1-weighted imaging and hyperintensity on T2-weighted imaging is again seen. Preoperative (D) and postoperative (E) axial CT scans reveal complete tumor removal with custom cranial implant replacing the removed bone.

**Table 1** Reported cases of CMF in the calvarium and skull base

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
1	Wilson et al	1991	26	F	Parietal bone Squamous temporal bone	—	Excision	—	None	0	—	—
2	Morimura et al	1992	41	M	Frontal bone	Hand hypesthesia Painless swelling	Excision	Gross total	None	0	—	—
3	Wu et al	1998	—	—	Frontal bone	—	—	—	None	0	—	—
4	Wu et al	1998	—	—	Frontal bone	—	—	—	None	0	—	—
5	Wu et al	1998	—	—	Frontal bone	—	—	—	None	0	—	—
6	Karkuzhali et al	2005	11	F	Parietal bone	Convulsions HA	Excision	Gross total	None	0	15	No
7	Hakan et al	2008	45	F	Frontal bone	Painless swelling	Excision	Gross total	None	0	20	No
8	Our study	2013	31	F	Parietal bone	HA Tender swelling	Excision	Gross total	None	0	3	No
SKULL BASE (59)												
Sinonasal (17)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
1	Koay et al	1995	57	F	Ethmoid sinus Frontal sinus Nasal bone Orbit	Painless swelling	Excision	Subtotal	None	0	6	No
2	Isenberg	1995	34	F	Ethmoid sinus Nasal bone	Nasal obstruction Painless swelling	Excision	(1) Subtotal (2) Gross total	None	0	(1) 36 (2) 8	Yes
3	Nazeer et al	1996	New born	M	Ethmoid sinus	Apnea Nasal obstruction Cardiorespiratory arrest	Excision	Gross total	None	0	12	No

Table 1 (Continued)

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
4	Mendoza et al	1998	New born	M	Ethmoid bone Nasal bone	Nasal obstruction	Excision	Gross total	None	0	24	No
5	Wu et al	1998	—	—	Ethmoid bone	—	—	—	None	0	—	—
6	Shek et al	1999	6	F	Ethmoid sinus Frontal bone Maxillary bone Nasal cavity Sella turcica	Blindness, Epistaxis Nasal obstruction	Curettage x4	Subtotal x4	+	50 Gy	120	Yes x3
7	Wang et al	2000	60	F	Ethmoid bone Nasal bone	Asymptomatic	Excision	Gross total	None	0	6	No
8	Baujaj et al	2001	50	F	Ethmoid sinus Frontal sinus Nasal bone	HA Nasal obstruction Nasal pain	Excision	Gross total	None	0	18	No
9	Azorin et al	2003	46	M	Frontal sinus	Painless swelling	Excision	Gross total	None	0	22	No
10	Smith et al	2006	49	F	Nasal cavity Palate	—	Biopsy	Gross total	None	0	30	No
11	Veras et al	2009	60	F	Nasal cavity	Asymptomatic	Excision	Gross total	None	0	12	No
12	Chen et al	2009	—	—	Maxillary bone	—	Excision	—	None	0	—	—
13	Kadom et al	2009	14	M	Ethmoid sinus Frontal bone Frontal sinus	Diplopia Exophthalmos HA Nasal obstruction	—	—	None	0	—	—
14	Castle et al	2011	43	F	Ethmoid sinus	Sinus pressure	—	—	None	0	—	—
15	Thomas et al	2011	49	M	Nasal cavity Palate	Face pain Nasal obstruction Sinus pressure	Excision	Gross total	None	0	24	No
16	Yoo et al	2012	New born	M	Nasal cavity Nasal bone	Nasal obstruction	Excision	Gross total	None	0	24	No

(Continued)

**Table 1 (Continued)**

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
17	McClurg et al	2012	49	F	Nasal bone Hard palate	Nasal obstruction Maxillary teeth numbness	Excision	Gross total	None	0	16	No
Clival/Sellar (7)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
1	Linskey et al	1990	73	M	Clivus	Ataxia CN 7, 10 impairment Deafness, Diplopia Dysphagia HA	Excision	Gross total	None	0	3	No
2	Keel et al	1997	34	F	Clivus Nasopharynx Prepontine Cistern	Blindness Diplopia HA	Curettage	Subtotal	+	68 Gy	11	No
3	Keel et al	1997	65	F	Clivus Ethmoid sinus Sphenoid sinus	HA	Curettage	Gross total	None	0	26	No
4	Keel et al	1997	42	M	Clivus Sphenoid sinus	-	-	-	-	-	-	-
5	Patino-Corredoba et al	1998	41	F	Clivus Ethmoid sinus Nasopharynx Sella turcica, Sphenoid sinus	Ataxia, Deafness, HA	Excision x3	(1) Subtotal (2) Gross total (3) Gross total	(1) + (2) - (3) -	-	(1) 72 (2) 12 (3) 48	Yes x2
6	Bloom et al	2004	38	F	Clivus Jugular foramen Occipital bone	CN 12 impairment Dysarthria Dysphagia HA Neck pain	Excision	Gross total	None	0	5	No

Table 1 (Continued)

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
7	Xu et al	2011	55	M	Sella turcica	Blindness Decreased ACTH HA	Excision	Gross total	None	0	6	No
Sphenoid/Parasellar (12)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
1	Nazeer et al	1996	66	F	Nasopharynx Sella turcica Sphenoid sinus	Nasal obstruction	Curettage	Subtotal	None	0	12	Yes
2	Keel et al	1997	51	F	Ethmoid sinus Sphenoid sinus	—	—	—	—	—	—	—
3	Keel et al	1997	66	F	Ethmoid sinus Nasopharynx Sphenoid sinus	Nasal obstruction	(1) Curettage (2) Excision	Subtotal	(1) — (2) +	61 Gy	(1) 6 (2) 20	Yes
4	Wu et al	1998	—	—	Sphenoid bone	—	—	—	None	0	—	—
5	Wu et al	1998	—	—	Sphenoid bone	—	—	—	None	0	—	—
6	Wu et al	1998	—	—	Sphenoid bone	—	—	—	None	0	—	—
7	Feuvret et al	2005	19	M	Petrous temporal bone Sphenoid bone	Diplopia	Excision	Subtotal	+	59 CGE, 45 Gy	(1) 14 (2) 12	Yes
8	Feuvret et al	2005	28	F	Sphenoid bone	CN 6 impairment Deafness Diplopia	Excision x2	Subtotal	+	59 CGE, 45 Gy	48	Yes
9	Vernon and Casiano	2006	44	M	Sphenoid sinus	Retro-orbital pain	Curettage	Gross total	None	0	—	No
10	Morris et al	2009	52	F	Sphenoid sinus	Vertigo	Curettage	—	None	0	24	No
11	Yu et al	2009	39	M	Cavernous sinus Mandibular ramus Petrous temporal bone sphenoid sinus	CN 5, 6 impairment Diplopia Face numbness	Excision	Subtotal	None	0	6	No

(Continued)

**Table 1** (Continued)

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
12	Our study	2013	38	F	Clivus Sphenoid bone	Diplopia Nasal obstruction Rhinorrhea	Excision	(1) Subtotal (2) Gross total	None	0	12	(1) Yes (2) No
Orbit/Zygoma (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
1	Carr et al	1992	41	F	Orbit Zygoma	Painless swelling	Excision	Gross total	None	0	18	No
2	Wolf et al	1997	35	F	Frontal bone Orbit Sphenoid bone	HA	Excision	Gross total	None	0	—	—
3	Hashimoto et al	1998	32	M	Ethmoid sinus Frontal sinus Nasal cavity Orbit	Exophthalmos Painless swelling	Excision	—	None	0	24	No
4	Bucci et al	2006	51	M	Orbit Zygoma	Painless swelling	Excision	Gross total	None	0	24	No
5	Cruz et al	2007	10	F	Ethmoid sinus Orbit	Exophthalmos	Excision	Gross total	None	0	12	No
6	Heindl et al	2009	37	F	Frontal bone Orbit	Exophthalmos HA Painless swelling	Excision	Gross total	None	0	24	No
7	Khalatbari et al	2012	14	M	Orbit Sphenoid bone	Exophthalmos Face pain Swelling	Excision	Gross total	None	0	96	No
8	Ditta et al	2012	51	F	Orbit Sphenoid bone Zygoma	Exophthalmos Face pain HA	Both	Gross total	None	0	5	Yes



Table 1 (Continued)

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
Temporal bone/Occiput (15)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
1	Dumas et al	1994	43	F	Jugular foramen Petrous temporal bone Occipital bone	Painless swelling Intermittent LOC tongue atrophy	—	—	None	0	—	—
2	Maruyama et al	1994	67	M	Jugular foramen Petrous temporal bone	CN 5–10 impairment Deafness HA Nausea	Excision	Subtotal	None	0	—	—
3	LeMay et al	1997	22	M	Mastoid temporal bone	Deafness HA Otalgia	Excision	Gross total	None	0	3	No
4	Patino-Corredoba et al	1998	20	M	Mastoid temporal bone	Deafness	Excision	Gross total	None	0	—	—
5	Wu et al	1998	—	—	Occipital bone	—	—	—	None	0	—	—
6	Wu et al	1998	—	—	Occipital bone	—	—	—	None	0	—	—
7	Suzuki et al	1999	49	M	Squamous temporal bone	Blindness Painless swelling	Excision	Gross total	None	0	—	—
8	Tarhan et al	2000	44	F	Tympanic temporal bone	Facial pain	Excision	Gross total	None	0	—	—
9	Otto et al	2007	58	F	Mastoid temporal bone	Syncope Vertigo	Excision	Gross total	None	0	6	No
10	Thompson et al	2009	33	F	Mastoid temporal bone Stylomastoid foramen	CN 7 impairment	Excision	—	None	0	—	—
11	Crocker et al	2009	22	M	C1 Clivus Occipital bone	Deafness	Excision	Gross total	None	0	24	No

(Continued)

**Table 1** (Continued)

Calvarial (8)												
Case no.	Author	Year	Age	Sex	Location	Clinical symptoms	Treatment	Gross total/subtotal resection	Radiation	Radiation dose	Follow-up (mo)	Recurrence
12	Ozek et al	2011	17	M	Cerebropontine angle Petrous temporal bone	CN 6, 7 impairment Deafness, Diplopia HA	Excision	Subtotal	None	0	—	No
13	Wang et al	2011	31	M	Petrous temporal bone	HA	Excision	—	None	0	—	—
14	Gupta et al	2012	42	M	Mastoid temporal bone Sigmoid sinus	Deafness Otalgia	Biopsy	—	None	0	—	—
15	Sharma et al	2012	12	F	Squamous temporal bone Zygoma	HA Otalgia Painful swelling	Excision	Gross total	None	0	—	No

Abbreviations: ACTH, adrenocorticotrophic hormone; CGE, cobalt gray equivalents; CMF, chondromyxoid fibroma; CN, cranial nerve; F, female; Gy, gray; HA, headache; LOC, loss of consciousness; M, male; Mo, months.

Reported cases of CMF in the calvarium and skull base. A literature review was performed identifying all cases of CMF. Eight calvarial tumor cases were reported. Skull base tumors were further categorized into sinonasal (17 cases), clival/sellar (7 cases), sphenoid/parasellar (12 cases), orbit/zygoma (8 cases), and temporal bone/occiput (15). Relevant data were extracted including, presenting clinical symptoms, treatment, gross total versus subtotal resection, radiation therapy if used, follow-up time, and recurrence.

result in a potentially delayed diagnosis for skull base lesions.<sup>12</sup> Symptoms included painless or tender swelling, headache, nasal obstruction, exophthalmos, diplopia, deafness, and otalgia. The most common presenting symptom for calvarial CMF was swelling. For the skull base lesions, the most common presenting symptoms appeared to be related to their respective locations. For sinonasal CMF, the most common presentation was nasal obstruction, whereas clival/sellar lesions presented most commonly with headache. Sphenoid/Parasellar lesions presented with diplopia and orbital/zygomatic CMF presented most commonly with exophthalmos. Last, temporal bone/occipital CMF was most commonly associated with deafness at presentation.

Given this tumor's rare propensity for the calvarium and skull base, we also reviewed our own institution's experience with CMF. Since 1957 to present, MDACC has seen 36 patients with CMF. Of these patients, there were two cases affecting the spine: one at C2 and the other at S1/lilium. There were only one calvarial case and one skull base case,<sup>13</sup> both reported here.

Radiologic findings are not diagnostic, but they can offer insight into the diagnosis before intervention. Classically CMF is described as a "radiolucent, lobulated, circumscribed lesion with a sclerotic rim and cortical expansion or erosion"<sup>14</sup> and calcification is rare. Because of the low occurrence rate of this tumor, MRI findings have not been clearly established. However, similar to other bone tumors, CMF has low signal on T1-weighted and high signal on T2-weighted images owing to its cartilaginous nature.<sup>14</sup> The most challenging aspect to radiologic diagnosis of CMF is the high variability of involved sites. Therefore suspicion for CMF should be maintained when evaluating solitary bone lesions.

Histopathologic analysis of CMF reveals a myxoid lesion containing a paucicellular center, bland stroma cells, reactive bony spicules, and hyaline cartilage with up to 75% of lesions in the skull and facial bones also containing matrix calcifications.<sup>8,15</sup> Though not always present, the characteristic features of CMF include lobular appearance, chondromyxoid stroma, and fibrous tissue with multinucleated giant cells.<sup>16</sup> Nielsen et al performed ultrastructural examination on six tumor samples finding populations of cells with features of three different cell types: chondrocytes, myofibroblasts, and a mixture of both chondrocytes and myofibroblasts.<sup>17</sup> It should be noted that if the lesion shows significant atypia or mitotic activity, the diagnosis of CMF should be reconsidered. CMF is most commonly found to be positive for vimentin, smooth muscle actin, desmin, S-100 (variably), and CD34.<sup>18</sup> Generally CMF is negative for pancytokeratin, carcinoembryonic antigen (CEA), and GFAP, and also has a low proliferation rate visualized by Ki-67 staining.<sup>19</sup> Veras et al describe SOX9 staining in a case of sinonasal CMF. SOX9 has been previously described as a chondrogenesis "master regulator" and plays a role in early-phase chondrocyte differentiation.<sup>15</sup>

Chondrosarcoma, chondroblastic osteosarcoma, fibrous dysplasia, chondroblastoma, and chordoma are included in the differential diagnosis.<sup>20,21</sup> This is an important distinction to make given the relatively benign nature of CMF. Low-grade chondrosarcomas and CMF share stellate cells, S-100 protein stain positivity, and negative keratin stain. However,

chondrosarcomas are more infiltrative than CMFs, which tend to "push" adjacent bony trabeculae.<sup>10</sup> Additionally, chondrosarcomas lack the fibrous component seen in CMF.<sup>22</sup> Desai et al, in their study of 10 CMF cases, reviewed the characteristics of the histologic differential diagnosis.<sup>20</sup> Chondroblastic osteosarcoma diagnostically has osteoid production, of which CMF does not, but it may be difficult to differentiate when there are large patches of CMF-like tissue present. Fibrous dysplasia has a myxoid appearance, but it is differentiated from CMF by its irregular osteoid seams. Chondroblastoma has prominent calcifications and an eosinophilic polygonal cytoplasm that is differentiated from the stellate cytoplasm of CMF. Last, chordoma tends to have large tumor cells, epithelioid with eosinophilic or vacuolated cytoplasm arranged in nests or cords.<sup>22</sup> Additionally, chordoma stains positive for S-100, epithelial membrane antigen (EMA), and cytokeratins whereas CMF is only positive for S-100.<sup>22</sup>

Though it had previously been thought that CMF is an acquired lesion, there is evidence suggesting a possible genetic link. Though cytogenetic studies are limited, Smith et al report there are 14 cases of CMF in the literature with known karyotypes. Notably 11 of the 14 cases had nonrandom clonal abnormalities of chromosome 6. In particular, rearrangements of the chromosome 6 long arm were most frequent, with four of the cases having pericentric inversion *inv(6)(p25q13)*.<sup>23</sup> Importantly, chromosome 6 has been implicated in normal cartilaginous development, carrying genes BMP6 (bone morphogenetic protein 6), COL9A1 (collagen type 9  $\alpha$  1), COL10A1 (collagen type 10  $\alpha$  1), and IGF2 (insulin-like growth factor 2). Additionally, supporting the possible genetic propensity for CMF, our literature review revealed three cases of congenital CMF presenting in newborns.

Recommended first-line treatment is surgical resection.<sup>6</sup> Given the benign nature of CMF, surgery can provide a cure. In our literature review, most patients received excision. Of the 67 cases reported here, 43 had excision, 6 curettage, and 2 received both. Two patients received only biopsy, and 14 cases did not report treatment. Of all the cases included here, there were nine recurrences. All but one case of recurrence was associated with subtotal resection, five with excision and three with curettage. Only one recurrence occurred in the setting of gross total resection in which the patient had the extraosseous component excised, while the bone itself was curetted until all visible tumor removed. Twenty-five cases had no data on recurrence.

Efficacy and need for radiotherapy is controversial. Some authors have argued in support of postoperative radiation to prevent local recurrence,<sup>24</sup> whereas others have reported concern for secondary risks of irradiation induced malignant transformation.<sup>25</sup> It remains to be clear if radiotherapy is of benefit in patients with subtotal resection or curettage. In our literature review, five patients received radiotherapy. Dosing ranged from 50 to 68 Gy. In two cases, the patient also received proton therapy, 59 CGE with 45 Gy. No dosing recommendations are available for CMF, but Feuvret et al proposed 55 to 60 Gy, consistent with other benign tumors.<sup>26</sup> Feuvret et al believe that radiotherapy should be part of standard treatment and proposed a treatment flow diagram

for skull CMFs. Recommendations for radiation include patients who are unresectable or received curettage or partial resection in vital neurologic areas, or have a recurrent tumor.<sup>26</sup> In our literature review, four of the five patients who received radiation had recurrence postirradiation. All of the patients who received radiotherapy had an initial subtotal resection. Though efficacy of radiotherapy in the setting of subtotal cranial CMF resection cannot be dismissed, our review did not show a benefit.

One of our patients and three patients identified in our literature review had symptom onset and subsequent diagnosis during pregnancy. This raises the possibility that these tumors are hormonally sensitive. To our knowledge, however, there is only one reported case where a hormonal influence was potentially associated with CMF.<sup>27</sup> Cytogenetic analysis of a scapular CMF revealed mutation in the parathyroid hormone/parathyroid hormone-related peptide receptor gene (PTH/PTHrP). Halbert et al indicate that this suggests the existence of autocrine/paracrine regulatory loops thought to be essential for normal chondrocyte maturation and/or endochondral bone formation.

## Conclusion

Cranial CMF is rare, and consequently frequently misdiagnosed. Key features of diagnosis are radiographic and histologic findings. It is important to keep CMF in the differential diagnosis when evaluating solitary cranial bone lesions, as CMF is curable by total excision. Total resection is the best treatment, and should be the goal of any surgical intervention. Curettage results in high recurrence rates. Radiotherapy in the setting of subtotal resection or recurrence cannot be definitively recommended and needs further investigation.

### Acknowledgments

Thanks to Dima Suki and Lamonne Crutcher for their help obtaining Institutional Review Board approval for this manuscript.

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