Primary Sternal Osteomyelitis

EDWARD A. GILL, Jr, MD DENNIS L. STEVENS, MD, PhD Boise, Idaho

STERNAL OSTEOMYELITIS has become more common in the past two decades largely related to the increased frequency of coronary artery bypass grafting and intravenous drug abuse. Sternal osteomyelitis as a complication of sternotomy is referred to as secondary sternal osteomyelitis; primary sternal osteomyelitis has no contiguous focus of infection. Contemporary cases of primary sternal osteomyelitis occur most commonly in intravenous drug abusers.¹ Primary sternal osteomyelitis remains a rarity with only 55 cases reported in the literature.² We report two cases of primary sternal osteomyelitis in patients with no apparent risk factors.

Report of Cases

Case 1

The patient, a 72-year-old man with non-insulin-dependent diabetes mellitus, was admitted for evaluation of an enlarging anterior chest wall mass. The patient had noted fullness in the left chest wall for a year before admission. Three weeks before admission, he had had sharp anterior chest pain that was precipitated when he arose from a sitting position or coughed. He also noted night sweats and dampness over the sternum. He said he did not have hemoptysis, dyspnea, dysphagia, anorexia, fevers, chills, or weight loss. He had not recently had an operation or trauma to the chest but did recall that a splinter of wood had lodged in the area near the lesion ten years previously.

On physical examination, the patient was afebrile but had a large—12 by 8 cm—wedge-shaped, erythematous, indurated mass over the anterior chest wall that protruded to the left of the midline. A chest roentgenogram revealed an osteolytic lesion of the sternum (Figure 1). On computed tomography there was destruction of the manubrium and adjacent ribs with soft tissue swelling extending into the anterior mediastinum (Figure 2). The results of laboratory tests included a leukocyte count of 12.4×10^9 per liter (12,400 per μ l) and an erythrocyte sedimentation rate of 120 mm per hour.

An incisional biopsy of the sternum was done and a histopathologic examination showed chronic inflammation. Culture of the biopsy grew *Staphylococcus aureus* sensitive to vancomycin hydrochloride, oxacillin sodium, and cephalothin sodium. A regimen of nafcillin sodium, 8 to 10 grams per day given intravenously, was initiated but stopped after 14 days due to a skin rash. The suspicion for carcinoma or sarcoma was high, and because the initial biopsy did not show malignancy, a second biopsy was done. Further local debridement was done at the time of biopsy, but no costal cartilages were removed. The biopsy specimen again did not show a neoplasm. A Penrose drain was placed, but the wound was closed with mattress sutures. A Hickman catheter was placed, and intravenous cefazolin sodium was administered at home for a period of three months. The erythrocyte sedimentation rate dropped to 15 mm per hour after four months, and a repeat computed tomography examination eight months later revealed a decrease in the size of the chest mass (Figure 2). Currently, two years later, the patient remains asymptomatic.

Case 2

Our second patient, a 59-year-old man with peripheral vascular disease, had been placed on sodium warfarin therapy following a femoral-to-popliteal artery bypass operation. In mid-November 1983, he had chest wall pain that had developed during a hospital admission for a right lower lobe pneumonia. The patient was treated as an outpatient with ibuprofen and acetaminophen given for presumed costochondritis. A month later, itching and redness developed over the sternal area directly beneath a nickel-plated necklace recently given to him to monitor the warfarin therapy. An erythematous skin rash was noted in the sternal area, and radiographs revealed the persistence of congenital ossified centers of the sternum (Figure 3). In January of 1984, an area of fluctuance was noted beneath the erythema, and a percutaneous aspirate from this area grew S aureus. Tomograms were suggestive of an osteolytic lesion (Figure 4). Surgical exploration was done for the purpose of biopsy, culture, and debridement. Under general anesthesia, the body of the sternum underwent curettement, leaving the posterior plate intact. The wound was then packed with iodoform gauze. Three months later the patient was noted to have a sinus tract at the site of the old surgical scar, and this was explored under general anesthesia. The sternum again was curetted and packed with iodoform gauze. Because of allergic reactions to multiple antibiotics, the patient received several different drug regimens including neomycin, nafcillin, and cephapirin sodium over a period of six weeks. An eight-week course of erythromycin followed the parenteral use of antibiotics. Staphylococcus aureus was grown from the biopsy specimens of both surgical procedures. Four years later the patient remains infection free.

Discussion

Primary sternal osteomyelitis, once a rare disorder, has become more common today with the increased prevalence of intravenous drug abuse. The most common bacterial cause of primary sternal osteomyelitis in intravenous drug abusers is *Pseudomonas aeruginosa*, although the most common overall is *S aureus*.²⁻⁴ *S aureus* is also the most common pathogen in secondary sternal osteomyelitis. It is interesting that *Mycobacterium tuberculosis* has become a rare cause of sternal osteomyelitis since 1952.⁵

Vaughn's review of sternal osteomyelitis in 1918 indicated that only 13 cases had been reported before 1910,⁶ and all were in the German literature.⁷ Wilensky and Samuels (1926) reported 24 cases of primary sternal osteomyelitis,⁸ but because several of the cases described were characterized by generalized sepsis, we have excluded these in our listing

⁽Gill EA Jr, Stevens DL: Primary sternal osteomyelitis. West J Med 1989 Aug; 151:199-203)

From the Departments of Medicine, Veterans Administration Medical Center (Drs Gill and Stevens), Boise, Idaho, and University of Washington School of Medicine (Dr Stevens), Seattle.

Reprint requests to Dennis L. Stevens, MD, PhD, Department of Medicine, VA Medical Center, Boise, ID 83702.

of the total number of cases with primary sternal osteomyelitis (Table 1).¹⁻¹³ In their series, the mortality rate was 27%, which is higher than in any other series.⁸ Many of their patients died of septic shock. Since the 1926 review, an additional 30 cases have been reported in international medical journals, bringing the total to 55 cases.

Table 2 compares cases of primary and secondary osteomyelitis both in terms of treatment and complications. Note that secondary sternal osteomyelitis was not described in the preantibiotic era. Since the introduction of the coronary artery bypass procedure, however, secondary osteomyelitis has become a well-known complication. Culliford and coworkers defined predisposing risk factors for poststernotomy osteomyelitis¹⁵: a prolonged perfusion time, excessive postoperative bleeding, a depressed cardiac output, and reexploration for the control of hemorrhage. In addition, internal mammary artery bypass grafts have increased the risk of sternotomy infection as compared with vein grafts. In fact, the overall infection rate of 1.5% climbed to 2.3% with the use of one internal mammary artery bypass and to 8.5% with bilateral internal mammary artery grafts.

Primary sternal osteomyelitis presents with anterior sternal pain, redness, tenderness, and swelling. When evaluating a case of chest wall pain or a mass in the sternal area, it is most important to consider sternal osteomyelitis in the differential diagnosis. Delayed diagnosis and inappropriate treatment are common in this illness. The differential diagnosis of primary sternal osteomyelitis is broad and should include cellulitis, soft tissue abscess, benign soft tissue tumors, and soft tissue or bony sarcoma. Fibrosarcoma is the most common soft tissue sarcoma. Bone sarcomas are rare, the most common being osteosarcoma and Ewing's sarcoma,

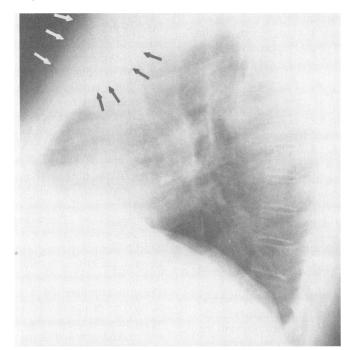


Figure 1.—A lateral chest x-ray film shows soft tissue swelling and destruction of the sternum (arrows) in patient 1.

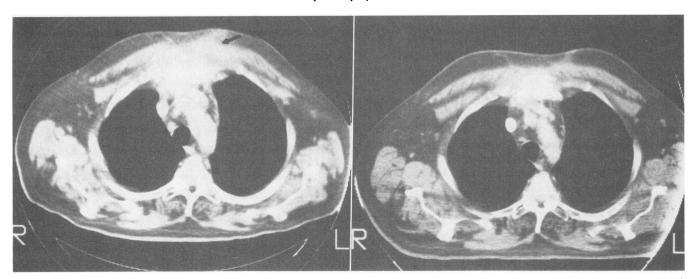


Figure 2.—Chest computed tomography in patient 1 shows, left, sternal destruction and inflammatory response extending through bone and soft tissue to the chest wall surface (arrow). Right, 9 months later, after surgical debridement and antibiotic therapy, the destructive process of the sternum is nearly absent.

Reported Case	Author, Year	Age, yr	Sex	Source of Bacteria	Organism	Surgical Therapy	Underlying Disease	Trauma	Outcome
1	Wilensky and Samuels, 19268	24	O'	Typhoid	Unknown	Debridement		No	Survived
2	•	35	ç	Sternum	Salmonella paratyphi	Debridement		No	Survived
3		24	ò	Sternum	Unknown	Radical resection*		Yes	Survived
4		8	O'	Sternum	Staph aureus	Debridement		Yes	Survived
5	Drews, 1910 ⁷ Drews, 1910 ⁷ ‡	Young†	•••	Sternum	Staph aureus	Debridement		No	Survived
	Solomon, 1926§	Youngt		Pneumonia	Unknown	Debridement		No	Died
	Sick, 1926§	40	 or	Pneumonia	Unknown	Debridement		No	Died
21	•	16	ď	General infection				No	Died
2		15	ç	Sternum	Unknown			No	Survived
23		21	ð	Sternum	Unknown			No	Survived
	Muhlein, 1926§	21		Lung	Unknown		Splenic tumor		Died
	Batut, 1926§	22	 oʻ	Sternum	Salmonella typhi	Debridement	Unknown		Survive
	Vaughn, 1918 ⁶	26	ð	Sternum	Unknown	Debridement		Yes	Survive
	Peloquin, 1926§			Sternum		4 operations		No	Died
	Bidwell, 1926§	••	•••	Liver	Unknown	Debridement	Liver abscess	No	Died
	. Koch, 1926§	30	 or	Pneumonia		Radical resection		No	Survive
	Janz, 1926§	21	ð	Pneumonia	Unknown	Debridement		No	Survive
	, Busch, 1926§ ~	18	ď	Sternum	Unknown	Radical resection		No	Survive
	. Maier, 1947 ⁹	19	ç	Pneumonia	Streptococcus sp	Debridement		No	Died
	Brown, 1952 ¹⁰	19	°	Sternum	?Mycobacterium tuberculosis	Debridement		No	Survive
34		41	ď	Sternum	M tuberculosis	Debridement		No	Survive
35		28	ď	Sternum	M tuberculosis	Several		No	Survive
	Vellacott, 195211	26	o	Sternum	Staph aureus	Debridement	Unknown	Yes	Survive
37	Biesecker, 1973 ¹²	21	ď	Sternum	Staph aureus	Debridement		No	Survive
38		21/2	œ	Sternum	Gram-negative	Debridement		No	Survive
	. Mir-Sepasi et al, 1975 ¹	22	ď	Sternum	Pseudomonas aeruginosa	Radical resection	IV drug abuse	No	Survive
10		24	ď	Sternum	P aeruginosa	Debridement	IV drug abuse	No	Survive
41		21	ď	Sternum	P aeruginosa	Debridement	IV drug abuse	No	Survive
42		24	ç	Sternum	P aeruginosa	Debridement	IV drug abuse	No	Survive
	. Enat et al, 1979 ¹³	47	ð	Sternum	Staph aureus			CPR	Survive
	. Jara et al, 1979 ⁴ ¶			•••••					
	. Sant, 1979 ³	28	ç	Sternum	Staph aureus				Survive
	. Kelly and Chetty, 1985 ²	40	ç	GI tract	Gram-negative				Survive
	. Richter, 1983 ¹⁴		•	Sternum	M tuberculosis				Survive
	. Watts et al. 1987^5	40	o	Sternum	M tuberculosis	Debridement		No	Survive
55		49	ď	Sternum	M tuberculosis			No	Survive
	pulmonary resuscitation, GI=gastrointesti				occus				
*Radical †A minim	resection was defined as complete resection al description of patients was given, with -18: Drews cited 13 cases from the Germa	on of the sterr age described	num wit I only a:	h additional recons s young.					

This may well have been a case of generalized sepsis. (Cases 44-50: 7 patients (1 woman, 6 men), all intravenous drug abusers, were treated with debridement and all survived. The causal organism in 4 was Pseudomonas aeruginosa, in 2 was Staphylococcus aureus, and in the 7th was undetermined. The ages and sources of bacteria were not given.

	Sternal Osteomyelitis						
Feature	Primary	Secondary					
Risk factors	. Trauma; ?pneumonia; ?diabetes	Internal mammary artery bypass grafting; prolonged time on heart-lun bypass; sternal massage for CPR after an operation; prolonged low cardia output state after a bypass procedure; excessive bleeding; history of reexploration for control of bleeding					
Complications	. Sepsis	Wire, saphenous vein bypass valve, and mediastinal infection; sepsis sternal erosion					
Mortality rate	. Preantibiotic era, 19% Postantibiotic era, 3%	10% all cases in postantibiotic era					
Management Organism CPR=cardiopulmonary resusc	. Antibiotics with or without surgical therapy . Staphylococcus aureus, Pseudomonas aeruginosa	Antibiotics plus reconstructive operations in most cases—left to heal open Staphylococcus aureus, Klebsiella pneumoniae, Candida albicans					

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Figure 3.—A plain lateral chest film of patient 2 shows midsternal hypertrophy and a questionable accessory joint (arrows).

both mainly pediatric tumors. Sarcoma is important to consider because a biopsy may facilitate a contiguous spread, and a complete surgical resection is necessary for cure in many cases.

The diagnosis of sternal osteomyelitis requires a bone biopsy. Helpful radiologic studies include computed tomography, a technetium Tc 99m pyrophosphate bone scan, and scans using gallium citrate Ga 67 and indium In 111. All of the nuclear medicine scans lack specificity and are perhaps best suited for follow-up examinations. Smith and associates, however, have advocated the use of gallium ⁶⁷Ga scanning as more specific than technetium ^{99m}Tc scanning.¹⁶ It is unclear from their report whether or not the ^{99m}Tc scans were multiphasic.¹⁶ We found plain and computed tomography to be the most useful in the diagnosis and follow-up of our patients. Although no studies exist supporting its use in this illness, magnetic resonance imaging has been useful in some types of osteomyelitis, especially in patients with Pott's disease.¹

The treatment of secondary sternal osteomyelitis requires more aggressive debridement than does the primary form. Hence, most cardiothoracic surgeons advocate extensive surgical intervention with some reconstruction in most cases. Johnson and colleagues have recommended total sternectomy and removal of associated cartilage, with reconstruction accomplished using bilateral pectoralis major myocutaneous flaps.18

While some debridement is often necessary in primary sternal osteomyelitis, a few cases (<10%) will resolve with antibiotic therapy alone. The extent of sternal debridement has been reviewed in a noncontrolled, nonprospective manner by two sets of authors. Mir-Sepasi and co-workers suggest removing most of the sternum but recommend leaving the posterior periosteum intact.1 Such a conservative

Figure 4.—A tomogram of the sternum shows the destructive process of the sternal osteomyelitis (arrows).

approach was curative in both of our cases, but each required a second operation.

Conclusions

We have presented two cases of primary sternal osteomyelitis, making the total of reported cases 57. Each patient had significant underlying illness: diabetes mellitus and peripheral vascular disease, respectively. Diabetes could certainly be postulated a risk factor for this disease. Case 1 also was unusual in that there was a preexisting history of a foreign body more than ten years before the patient was first seen. Our cases were more typical of the older cases in the literature in that there was no history of intravenous drug abuse. We also report what we think is the first case of this illness treated on an outpatient basis with intravenous antibiotics through a Hickman catheter. Hence, we have defined an approach to this illness combining appropriate diagnostic studies, limited surgical intervention, and appropriate antimicrobial therapy.

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Metastatic Crohn's Disease

SUSAN TERI McGILLIS, MD Cleveland ARTHUR C. HUNTLEY, MD Sacramento, California

GRANULOMATOUS LESIONS OF THE SKIN not attributable to a known foreign body reaction or infectious agents are often presumed to be sarcoidosis, a disease of exclusion. We report the case of a patient initially thought to have cutaneous sarcoidosis, who on further investigation was determined to have "metastatic" Crohn's disease.

Report of a Case

The patient, a 62-year-old man, first consulted our dermatology service for evaluation of an erythematous, scaly, nummular patch on his left lower leg that had been present for about two months.

Further history indicated that six months before he noted the lesion, he had been ill with malaise, diarrhea, vomiting, and a 9-kg (20-lb) weight loss. He was admitted to hospital and underwent an extensive evaluation, including radiographic studies that revealed tumors in the antrum of the stomach and terminal ileum (Figure 1). These were initially considered to be consistent with adenocarcinoma, but an endoscopic biopsy of the gastric lesion showed many noncaseating granulomata. No other systemic illnesses or abnormalities were identified at that time. A diagnosis of Crohn's disease was made after tuberculosis and other causes for the granulomata were ruled out. He was treated with a regimen of prednisone, 30 mg a day, with complete resolution of all gastrointestinal symptoms. The patient had been completely weaned off prednisone at the time of being seen by us.

About two months after the prednisone therapy was discontinued, he was examined in the dermatology clinic because of mild erythema and scaling that had developed on both legs. Several potassium hydroxide preparations were negative, and the patient was initially treated with midstrength topical steroids with no apparent response. The lesions progressed, and eventually thickened, indurated plaques developed with sharply marginated ulcerations and crusting (Figure 2).

Several skin biopsies revealed sarcoidal granulomas with few giant cells and little coagulation necrosis (Figure 3). Special stains and cultures were negative for deep fungi, acid-fast bacteria, and spirochetes. No foreign body material was identified by polarization. A diagnosis of sarcoidosis was initially made, and a complete evaluation for systemic involvement was embarked on. All results were normal. The patient had no adenopathy. A chest x-ray film showed no hilar

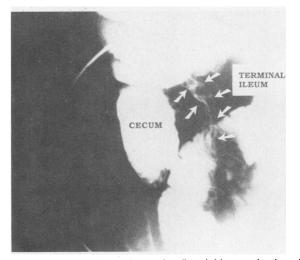


Figure 1.—A barium-enhanced radiographic examination of the intestine reveals cuffing of the terminal ileum (arrows) characteristic of Crohn's disease.



Figure 2.—The initial skin findings consisted of dermatitis and scaling of the legs within which there is ulceration and crusting.

⁽McGillis ST, Huntley AC: Metastatic Crohn's disease. West J Med 1989 Aug; 151:203-205)

From the Departments of Dermatology, Cleveland Clinic Foundation, Cleveland (Dr McGillis), and University of California, Davis, Medical Center, Sacramento (Dr Huntley).

Reprint requests to Arthur C. Huntley, MD, Department of Dermatology, University of California, Davis, Medical Center, 1605 Alhambra Blvd, Sacramento, CA 95816.