Treatment of Acute and Recurrent Idiopathic Pericarditis
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Case Presentation: A 56-year-old previously healthy man presented with 2 days of pleuritic left anterior chest pain, lessened by sitting forward. His examination was pertinent for low-grade fever (37.6°C), blood pressure 122/76 mm Hg without paradox, no jugular venous distension, clear lungs, and a 3-component pericardial friction rub. The ECG showed diffuse concave-upward ST-segment elevation and PR-segment depression in the inferior leads. The serum C-reactive protein level was 64 mg/L, and the cardiac troponin T was not elevated. Echocardiography showed normal left ventricular contractile function without wall motion abnormalities and no pericardial effusion. He was diagnosed with acute pericarditis, and the symptoms responded promptly to oral ibuprofen, continued for 2 weeks.

Six weeks later, he redeveloped pleuritic chest pain and clinical and ECG findings identical to the initial presentation. His primary care physician asks for advice about appropriate therapy.

Background
Pericarditis accounts for 5% of emergency department visits for chest pain in the absence of myocardial infarction. In ≈80% of cases in developed countries, the cause of pericarditis is either postviral or “idiopathic,” in that it cannot be attributed to a specific condition. Because these 2 etiologies are clinically equivalent, the term idiopathic pericarditis will refer to both in this Clinician Update. Even when managed effectively, many patients with acute pericarditis present with 1 or more repeated episodes, termed recurrent pericarditis.

Acute Pericarditis Management
Treatment of idiopathic pericarditis has long been empirical, because until recently, there have been few therapeutic trials addressing this condition. The European Society of Cardiology published the only treatment guideline for pericarditis almost a decade ago, and many of the recommendations were based on opinion because of the lack of available study evidence.

Most patients with idiopathic pericarditis experience self-limited symptoms that improve spontaneously within days to weeks. More rapid relief can be achieved with pharmacological intervention, and stable patients can be managed in the outpatient setting. Hospitalization is recommended when features suggest nonidiopathic causes or herald hemodynamic compromise, including fever >38°C (>100.4°F), the subacute development of symptoms (characteristic of tuberculosis, neoplastic disease, uremia, or collagen vascular disorders), hypotension, jugular venous distension, a large pericardial effusion, or echocardiographic features of impending tamponade (Figure). Patients who are immunocompromised or are undergoing therapy with anticoagulants should also be observed initially in the hospital.

Pharmacological Treatment
Effective agents include nonsteroidal anti-inflammatory drugs (NSAIDs), colchicine, and glucocorticoids. Concurrently, rest and avoidance of demanding physical activity help to minimize symptoms.

Nonsteroidal Anti-inflammatory Drugs
Aspirin and other NSAIDs are the first-line approach, based on clinical experience and observational reports. For example, in a 2004 study without a control group, outpatient therapy of uncomplicated pericarditis with aspirin relieved symptoms in 87% of 254 patients. Commonly used NSAID regimens are listed in the Table, with a recommended initial duration of 7 to 14 days, then treatment should be tapered until resolution of symptoms and improvement of acutely elevated serum inflammatory markers such as C-reactive protein and the erythrocyte count.
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have delayed infarct healing in animal
dial infarction because other NSAIDs
patients with pericarditis after myocar-
preferred anti-inflammatory agent for
report of 254 patients with acute pericar-
dital/idiopathic should be suspected. In the
cardial effusion develops during therapy,
persists >1 week, or a new or larger peri-
However, if chest discomfort or fever

ditions such as pericardial constriction.3
an unlikely progression to complica-
tions, and 18% had tuberculosis.5
ultimately found not to have idiopathic
not respond by 7 days of therapy were
61% of those with symptoms who did
not respond by 7 days of therapy were
ultimately found not to have idiopathic
permisodic 
pericarditis. Forty-three percent were


dation only rarely, primarily for diarrhea.12 Uncommon serious side effects,
occuring primarily in patients with chronic renal insufficiency, include hepatic toxicity and myelosuppression. It is now common practice to include colchicine, in combination with an
NSAID, as initial management of acute idiopathic pericarditis (Table).

**Glucocorticoids**
Steroid therapy has long been used to treat pericarditis because it induces prompt symptomatic relief; however, glucocorticoids should not be used as primary therapy in uncomplicated acute idiopathic pericarditis because of a high rate of relapse when the steroid is tapered or stopped.11,12,13 Glucocorticoids also appear to blunt the efficacy
of colchicine in preventing recurrences.14 As a result, and owing to the side effects associated with long-term steroid therapy, glucocorticoids should only be prescribed to patients with idiopathic pericarditis who are refractory to treatment with, or intolerant of, an NSAID plus colchicine.4

When used, the prednisone dosage recommended in the 2004 European Society of Cardiology guidelines was a relatively high 1.0 mg·kg<sup>−1</sup>·d<sup>−1</sup> for 2 weeks with rapid tapering. A lower dosage (0.25–0.50 mg·kg<sup>−1</sup>·d<sup>−1</sup>) for 2 to 4 weeks, followed by slow tapering (Table) is effective and is associated with fewer relapses than the higher dosage.15

**Recurrent Pericarditis Management**
One or more recurrences arise in 15% to
30% of patients after an initial episode of acute pericarditis.2 These attacks can

| Table. Pharmacological Treatment of Idiopathic Pericarditis |
|----------------------------------|--------|----------------|
| **Medication**                   | **Dosage** | **Duration of Therapy** |
| Nonsteroidal anti-inflammatory drugs |        |                 |
| Aspirin                          | PO: 650–975 mg 3–4 times daily | 1–2 wk (2–4 wk for recurrence) |
| Ibuprofen                        | PO: 400–800 mg 3 times daily | 1–2 wk (2–4 wk for recurrence) |
| Indomethacin                     | PO: 50 mg 3 times daily | 1–2 wk (2–4 wk for recurrence) |
| Ketorolac                        | IM: 30–60 mg once, or 15–30 mg every 6 h, or IV: 15–50 mg every 6 h | Should not exceed 5 days |
| Colchicine                       | 0.5 or 0.6 mg 2 times daily* | Up to 3 mo (Up to 6 mo for recurrence) |
| Prednisone                       | 0.25–0.5 mg/kg/d for 2 wk (2–4 wk for recurrence), then: | |
|                                  | **At dose of:** | **Taper by:** |
|                                  | 25–50 mg daily | 5–10 mg every 1–2 wk |
|                                  | 15–25 mg daily | 2.5 mg every 2–4 wk |
|                                  | <15 mg daily | 1.0–2.5 mg every 2–6 wk |

*Reduced dosage recommended for patients with advanced renal dysfunction or concurrent therapy with moderate to strong inhibitors of CYP3A4 (eg, protease inhibitors, ketoconazole, fluconazole, erythromycin, diltiazem, verapamil) or P-glycoprotein inhibitors (eg, cyclosporine).
†As recommended by Imazio et al.2
repeat over extended periods of time and may lead to substantial disability. A first recurrence typically presents within 18 months, and findings are similar to the initial episode, including pleuritic chest pain, diffuse ST-segment elevations, a pericardial friction rub, and elevated serum markers of inflammation.16

Pharmacological Treatment

Nonsteroidal Anti-inflammatory Drugs

In the absence of prospective trial evidence, aspirin or another NSAID should form the foundation of therapy for recurrences (Table).4 However, in contrast to the brief course of NSAID generally prescribed for an initial episode, a gradual tapering of the drug over 2 to 4 weeks after symptoms improve is recommended.16

Colchicine

Use of colchicine in the COPE trial was associated with fewer initial pericarditis recurrences.12 Additionally, the Colchicine for Recurrent Pericarditis (CORE) trial randomized 84 patients who had already had a first recurrence to aspirin or aspirin plus colchicine.16 Compared with aspirin alone, the combination reduced the rate of additional recurrences by 50%.

Most recently, the double-blinded Colchicine for Recurrent Pericarditis (CORP) trial randomized 120 patients with a first recurrence to colchicine or placebo, in addition to aspirin or another NSAID.17 The rate of subsequent recurrence was 24% in those randomized to colchicine compared with 55% in the placebo group. In addition, the mean number of episodes was reduced and the time to next recurrence was lengthened. The duration of colchicine therapy in the CORE and CORP trials for recurrent pericarditis was 6 months.

Glucocorticoids

Symptoms of pericarditis recurrence respond promptly to glucocorticoid therapy.1 However, when administered in this situation, slow tapering and a prolonged course may be required to prevent recrudescent symptoms, with the potential for long-term steroid-associated side effects. Furthermore, the risk of additional recurrences of pericarditis is augmented by steroid use.12,14,16 Therefore, the consensus is to initially treat recurrent episodes of pericarditis with an NSAID plus colchicine and to prescribe glucocorticoids only for refractory cases.

Patients sometimes present with chest discomfort reminiscent of prior pericarditis, which is interpreted as a recurrence, even in the absence of objective findings (no pericardial rub, ECG or echocardiographic abnormalities, or elevation of serum inflammatory markers). Although a trial of an NSAID plus colchicine may be reasonable in this situation, glucocorticoids should certainly be avoided.18

Imazio and colleagues15 compared 2 steroid dosage intensities in recurrent pericarditis: Prednisone 0.2 to 0.5 mg·kg⁻¹·d⁻¹ versus the higher commonly used dose of 1.0 mg·kg⁻¹·d⁻¹ for 4 weeks, followed by a slow taper. The lower-dose regimen was effective, whereas the higher dosage was associated with more side effects and a greater number of subsequent pericarditis recurrences and hospitalizations. Thus, a now common approach to the use of steroids in patients with recurrent pericarditis whose symptoms are refractory to an NSAID plus colchicine is the lower-dose prednisone regimen listed in the Table. With prolonged corticosteroid
use, osteoporosis prevention (eg, calcium, vitamin D, and bisphosphonates) should be considered.

A common cause of referral to specialized pericardial centers is the inability to taper glucocorticoid therapy below a certain dosage (typically ≥15 mg of prednisone daily) without reemergence of symptoms, despite concurrent NSAID plus colchicine treatment. An often effective strategy in this circumstance is to resume the lowest prior steroid dosage that had controlled symptoms, and then taper it by only 1 to 2 mg every 2 to 4 weeks.  

Other Considerations

For refractory pericarditis despite NSAID, colchicine, and glucocorticoid therapies, improved symptoms have been reported in small numbers of patients with the use of immunosuppressive agents (azathioprine or methotrexate), intravenous immunoglobulin, and the interleukin-1β receptor antagonist anakinra. Finally, pericardiectomy can be undertaken for symptomatic relief in cases of continuously relapsing pericarditis. Results have been variable, with some patients experiencing complete remission but others continuing to be plagued with ongoing symptoms after surgical intervention. The best outcomes have been reported when complete resection of the pericardium is undertaken.  

Case Presentation (Continued)

The patient’s recurrent pericarditis was treated with ibuprofen 600 mg 3 times daily plus colchicine 0.6 mg twice daily. Ibuprofen was tapered off after 3 weeks, and the colchicine was continued for 6 months. After 1 additional year of follow-up, he has had no further symptoms of pericarditis.

Summary

Appropriate therapy for acute idiopathic pericarditis is an NSAID for ≥2 weeks, and it is also reasonable to prescribe colchicine for up to 3 months (the duration used in clinical trials), especially to reduce the rate of recurrence. For initially refractory symptoms, the parenteral NSAID ketorolac may be beneficial. For recurrent episodes of pericarditis, treatment with an NSAID plus colchicine is recommended, but for a more prolonged course. During NSAID treatment, concurrent gastric protection therapy should be considered. Only for truly refractory cases should glucocorticoid therapy be used.

Disclosures

None.

References


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