

# **Limb Pain**

By <u>Andrea D. Thompson</u>, MD, PhD, University of Michigan; <u>Michael J. Shea</u>, MD, Michigan Medicine at the University of Michigan Reviewed/Revised Aug 2024

#### **Etiology** | **Evaluation** | **Treatment** | **Key Points**

Limb pain may affect all or part of an extremity (for joint pain, see <u>Pain in and Around a Single Joint</u> and <u>Pain in Multiple Joints</u>). Pain may be constant or intermittent, and unrelated to motion or precipitated by it. Accompanying symptoms and signs often suggest a source.

### **Etiology of Limb Pain**

The **most common causes** of pain in a limb are also readily apparent by history.

• Musculoskeletal injuries and overuse

This discussion covers extra-articular limb pain unrelated to injury or strain. Pain that is in only <u>one joint</u> or in <u>multiple joints</u> is discussed elsewhere.

There are many causes (see table <u>Some Causes of Nontraumatic Limb Pain</u>) but the most common are the following:

- <u>Deep venous thrombosis</u> (DVT)
- Cellulitis
- Radiculopathy

Uncommon but serious causes that require immediate diagnosis and treatment include

- Acute arterial occlusion
- Deep soft-tissue infection
- Acute coronary ischemia (manifesting with only referred arm pain)

TABLE					
Some Causes of Nontraumatic Limb Pain					
Cause	Suggestive Findings	Diagnostic Approach			
Musculoskeletal and soft	tissue				
<u>Cellulitis</u>	Focal erythema (or darker color on dark skin), warmth, tenderness, swelling Sometimes fever	Clinical evaluation  Sometimes blood and tissue cultures (eg, when patients are immunocompromised)			
Deep soft-tissue infection (eg, myonecrosis, necrotizing subcutaneous infection)	Deep, constant pain, typically out of proportion to other findings				
	Erythema (or darker color on dark skin), warmth, tenderness, tense swelling, fever	Blood and tissue cultures Radiography Sometimes MRI			
	Sometimes crepitation, foul discharge, bullae or necrotic areas, signs of systemic toxicity (eg, delirium, tachycardia, pallor, shock)				
<u>Osteomyelitis</u>	Deep, constant, often nocturnal pain				
	Bone tenderness, fever	Radiography, MRI, and/or CT Sometimes bone culture			
	Often risk factors (eg, immunocompromise, parenteral illicit drug use, known contiguous or remote source for infection)				
Bone tumor (primary or metastatic)	Deep, constant, often nocturnal pain	Radiography, MRI, and/or CT			
	Bone tenderness				
	Often a known cancer				
Vascular					
	Swelling, often warmth and/or redness, sometimes venous	1.116			

Doon worse	distension	UILI dSUUI IU
<u>Deep venous</u>		Sometimes D-dimer
<u>thrombosis</u>	Often risk factors (eg, hypercoagulable state, recent	testing
	surgery or immobility, cancer)	
Chronic venous stasis	Mild discomfort with swelling, erythema, and warmth of distal lower extremity	Clinical evaluation
	Sometimes shallow ulcerations	
	Sudden, severe pain	
Acute ischemia (typically due to arterial embolism, dissection, or thrombosis but sometimes due to massive iliofemoral venous thrombosis that completely obstructs blood flow in the limb)	Signs of distal limb ischemia (eg, coolness, pallor, pulse deficits, delayed capillary refill)	
	Sometimes chronic ischemic skin changes (eg, atrophy, hair loss, pale color, ulceration)	Immediate <u>arteriography</u>
	After several hours, neurologic deficits and muscle tenderness	
	Sometimes known peripheral vascular disease	
Peripheral arterial insufficiency	Intermittent leg pain triggered predictably by exertion and relieved by rest (intermittent claudication), sometimes rest pain which may worsen with leg elevation	Ultrasound Sometimes arteriography
	Low ankle-brachial blood pressure index, chronic ischemic skin changes	
Neurologic		
Plexopathy (brachial or lumbar)	Usually weakness, often decreased reflexes Sometimes numbness in a nerve plexus distribution	Usually <u>electrodiagnostic</u> <u>testing</u> (electromyography and nerve conduction velocity)
		Sometimes MRI

Thoracic outlet syndrome	Pain and paresthesias beginning in neck or shoulder and extending to medial aspect of arm and hand	Clinical evaluation
		Sometimes, electrodiagnostic testing and/or MRI
Radiculopathy (eg, caused by herniated intervertebral disk or bone spurs)	Pain and sometimes sensory deficits following a dermatomal distribution and often worsening with movement  Often neck or back pain  Usually weakness and diminished deep tendon reflexes in a nerve root distribution	Usually MRI Sometimes electromyography and nerve conduction velocity
Painful <u>polyneuropathy</u> (eg, alcoholic neuropathy)	Chronic, burning pain, typically in both hands or both feet	Clinical evaluation
	Sometimes sensory abnormalities such as hypoesthesia, hyperesthesia, and/or allodynia (pain with non-noxious stimuli)	Sometimes electromyography and nerve conduction velocity
Complex regional pain syndrome (CRPS)	Burning pain, hyperesthesia, allodynia, vasomotor abnormalities  Typically a prior injury (may be remote)	Clinical evaluation
Other		
Acute coronary ischemia (causing referred arm pain)	Absence of explanatory physical findings at the site of pain; other suggestive findings (eg, history suggesting coronary artery disease, sweating and/or dyspnea occurring simultaneously with arm pain)	ECG and serum troponin Sometimes stress testing or coronary angiography
	Chronic pain and tenderness along a taut band of muscle,	

Myotasciai pain syndrome worsening with movement and with pressure on a trigger point (focal area separate from site of pain)

Clinical evaluation

### **Evaluation of Limb Pain**

It is important to exclude <u>acute arterial occlusion</u>.

### History

**History of present illness** should address the duration, intensity, location, quality, and temporal pattern of pain. Recent injury, excessive and/or unusual use, and factors that worsen pain (eg, limb movement, walking) and relieve pain (eg, rest, certain positions) should be noted. Any associated neurologic symptoms (eg, numbness, paresthesias) should be identified.

**Review of systems** should seek symptoms of possible causes, including back or neck pain (radiculopathy), fever (infections such as <u>osteomyelitis</u>, <u>cellulitis</u>, or deep soft-tissue infection), dyspnea (<u>DVT</u> with <u>pulmonary embolism</u>, <u>myocardial infarction</u>), and chest pain or sweating (myocardial ischemia).

**Past medical history** should identify known risk factors, including cancer (<u>metastatic bone tumors</u>); immunocompromising disorders or medications (infections); hypercoagulable states (<u>DVT</u>); <u>diabetes</u>; peripheral vascular disease, hypercholesterolemia, and/or <u>hypertension</u> (acute or chronic ischemia); <u>osteoarthritis</u> or <u>rheumatoid arthritis</u> (radiculopathy); and prior injury (<u>complex regional pain syndrome</u>). Family and social history should address family history of early vascular disease and cigarette smoking (limb or myocardial ischemia) and illicit use of parenteral drugs (infections).

# Physical examination

Vital signs are reviewed for fever (suggesting infection) and tachycardia and/or tachypnea (compatible with DVT with pulmonary embolism, myocardial infarction, and infection with sepsis).

The painful limb is inspected for color, edema, and any skin or hair changes, and palpated for pulses, temperature, tenderness, and crepitation (a subtle crackling sensation indicating soft-tissue gas). Strength, sensation and deep tendon reflexes are compared between affected and unaffected sides. Systolic blood pressure (BP) is measured in the ankle of the affected extremity and compared with systolic BP of an arm; the ratio of the two is the ankle-brachial index.

# **Red flags**

- Sudden, severe pain
- Signs of acute limb ischemia (eg, coolness, pallor, pulse deficits, delayed capillary refill)
- Dyspnea, chest pain, and/or sweating
- Signs of systemic toxicity (eg, delirium, tachycardia, shock, pallor)

- Crepitation, tenseness, foul discharge, bullae, necrosis
- Risk factors for deep venous thrombosis
- Neurologic deficits

### Interpretation of findings

#### Streptococcal Cellulitis

**IMAGE** 



© SPRINGER SCIENCE+BUSINESS MEDIA

It can be helpful to categorize pain by acuity of symptom onset and then further narrow the differential diagnosis based on presence or absence of findings of

- Ischemia
- Inflammation
- Neurologic abnormalities

**Sudden, severe pain** suggests acute ischemia or acute radiculopathy (eg, due to sudden disc herniation). Acute ischemia causes generalized limb pain and manifests with weak or absent pulse, delayed capillary refill (≥ 2 seconds or, with unilateral symptoms, longer than the opposite side), coolness, and pallor; ankle-brachial index is typically < 0.3. Such vascular signs are absent with radiculopathy, in which pain instead follows a dermatomal distribution and is often accompanied by back or neck pain and diminished deep tendon reflexes. However, in both cases, weakness may be present. Acute ischemia due to massive venous thrombosis (phlegmasia cerulea dolens) usually causes edema, which is not present in ischemia due to arterial occlusion.

In **subacute pain** (ie, of 1 to a few days' duration), erythema and tenderness, often accompanied by swelling, and/or warmth, suggest an inflammatory cause. If these findings are focal or circumscribed, cellulitis is likely. Generalized, circumferential swelling is more suggestive of DVT or, much less commonly, deep tissue infection. Patients with a deep tissue infection typically appear quite ill and may have blisters, necrosis, or crepitation. Findings in DVT vary widely; swelling and warmth may be minimal

or absent. Neurologic findings of weakness, paresthesias, and/or sensory abnormalities suggest radiculopathy or plexopathy. If neurologic findings follow a dermatomal pattern, radiculopathy is more likely.

**Chronic pain** can be difficult to diagnose. If neurologic findings are present, causes include radiculopathy (dermatomal distribution), plexopathy (plexus distribution), neuropathy (stocking-glove distribution), and complex regional pain syndrome (variable distribution). Complex regional pain syndrome should be suspected if vasomotor changes (eg, pallor, mottling, coolness) are present, particularly in patients with previous injury to the affected extremity. Myofascial pain syndrome causes no neurovascular abnormalities and classically manifests with a palpably tense band of muscle in the area of pain, and pain may be reproduced by pressure on a trigger point near but not overlying the area of pain. In patients with essentially no clinical findings, cancer and osteomyelitis should be considered, particularly in those with risk factors.

#### **Cholesterol Crystal Embolization**

**IMAGE** 



© SPRINGER SCIENCE+BUSINESS MEDIA

Intermittent pain occurring consistently with a given degree of exertion (eg, whenever walking > 3 blocks) and relieved with a few minutes of rest suggests peripheral arterial disease. Such patients typically have an ankle-brachial blood pressure (BP) index of  $\leq$  0.9; an index  $\leq$  0.4 indicates severe disease. However, arterial stiffness can produce falsely negative ankle-brachial index values. Because the toe arteries are less susceptible to stiffening, the toe-brachial BP index can be measured instead in patients with suspected peripheral arterial disease and in whom the ankle arteries are likely not compressible (eg, patients with advanced diabetes or aging). Patients with exertional symptoms and normal or borderline ankle-brachial BP index (> 0.9 but < 1.40) should have repeat ankle-brachial BP index measurement after exercise on a treadmill. Patients with peripheral arterial disease may have chronic skin changes (eg, atrophy, hair loss, pale color, ulceration).

#### **Testing**

Cellulitis, myofascial pain, painful polyneuropathy, and complex regional pain syndrome can often be diagnosed clinically. Testing (see table <u>Some Causes of Nontraumatic Limb Pain</u>) is usually necessary for other suspected causes of pain.

### **Treatment of Limb Pain**

Primary treatment is directed at the cause. Analgesics can help relieve pain.

## **Key Points**

- Acute limb ischemia should be considered in patients with sudden, severe pain.
- Presence or absence of findings of ischemia, inflammation, and neurologic abnormalities plus the acuity of onset help narrow the differential diagnosis.



Copyright © 2025 Merck & Co., Inc., Rahway, NJ, USA and its affiliates. All rights reserved.