Objective: The purpose of this study is to present a review of the current understanding of glomus tumors of the hand.

Methods: Clinical cases are used to demonstrate the relevance of history and physical examination in deriving the diagnosis of this rare, but important entity. Treatment, complications, and review of the literature are presented.

Results: Glomus tumors are rare vascular lesions representing approximately 1% of all hand tumors. Derived from the glomus body, a normal intradermal arteriovenous anastomosis that arises from the normal neuromyoarterial glomus. These structures regulate skin temperature. They are present in the tips of the digits, particularly in the subungual area. The glomus body consists of an arteriole, a venule, and an anastomotic vessel without an intervening capillary bed. Histologic findings include endothelial pericytes and numerous nonmyelinated nerve fibers.

Glomus tumors account for approximately 1% of all hand tumors and are even less common in children. Colon and Upton reported a series of 349 pediatric hand tumors, 9 of which were glomus tumors. Glomus tumors are more common in women between 30 and 50 years of age, are not known to be associated with any other condition, and occur spontaneously. They are usually solitary but a multiple glomus tumor syndrome has been described that is transmitted in an autosomal dominant pattern.
Glomus Tumors of the Hand

Glomus tumors present as a classic triad of severe pain, point tenderness, and cold sensitivity. In a series of 51 patients with glomus tumors of the hand, by Van Geertruyden et al, spontaneous pain was seen in 80% of patients. Sensitivity to touch was present in 100% of patients and cold sensitivity in 63%. Bhaskaranand and Navadgi reported, in their series of 18 hand tumors, that of the 14 patients with glomus tumors all presented with severe pain. The 4 patients with other types of tumors presented with dull, aching pain. In the series, 100% of patients had point tenderness and 78% had cold sensitivity. Carroll and Berman state that excruciating, paroxysmal pain is pathognomonic for glomus tumor.

Clinical features of glomus tumors include nail deformity (Fig 1), blue discoloration (Fig 2), and palpable nodule. Van Geertruyden et al reported rates of 27%, 29%, and 6% for these findings, respectively. When only dorsal tumors are considered, the incidence of nail deformation rises to 47% and blue discoloration is seen in 43% of such lesions.

Figure 1. A 30-year-old woman with a subungual glomus tumor of the left thumb. Preoperative view of the perionychium with the tumor (left); postoperative view with minimal nail deformity (right). A 44-year-old woman with subungual tumor of the ring finger. Patient presented with 3-month history of nail discoloration and tumor. Preoperative view demonstrating tumor from the nail matrix (center, below); and postoperative view at 2 years demonstrating mild nail deformity (right).

DIAGNOSIS

Diagnosis of glomus tumors is primarily clinical. Several clinical tests are useful for diagnosing glomus tumors. Love reported that localization of the tenderness to an area and the size of a pinhead was suggestive of glomus tumor. For a positive Love’s pin test, the patient should experience severe pain and reduction in pain when the skin overlying the tumor is pressed with a pinhead, ballpoint pen, end of a paperclip, or Kirschner wire. The Love test is 100% sensitive.
and specific according to Bhaskaranand and Navadgi's series.  

The cold-sensitivity test is positive when immersing the hand in cold water elicits severe pain in and around the lesion. In addition, there should be a history of cold weather aggravating the symptoms. Hildreth's test is another reliable clinical sign for the diagnosis of glomus tumors.

This test is performed by elevating the patient's arm to exsanguinate it. A tourniquet is inflated to 250 mm-Hg and the tumor is palpated, the pain and tenderness should be reduced. A test is positive when releasing the cuff causes a sudden onset of pain and tenderness in the area of the tumor. In his series of 24 patients with hand tumors, Giele noted a positive Hildreth's test in 13 patients. Twelve of these had glomus tumors and 1 had a hemangiopericytoma with a sensitivity of 92% and specificity of 91%.

Radiography, ultrasound, magnetic resonance imaging, and angiography, all have been used with variable amounts of success for diagnosis and localization of lesions.

TREATMENT

Surgical excision is the treatment for glomus tumors; no medical therapy exists. Excision of the tumor results in resolution of symptoms in all cases.

Love's test is used to localize the tumor and it is completely excised under direct vision. Subungual tumors are approached transungually by removing the nail plate and incising the nail bed longitudinally. For lesions that are deep-seated proximally, lateral incisions through the paronychium are recommended. This approach can facilitate better access to the germinal matrix (Fig), whereas incisions through the hyponychium may cause the flap to retract and result in significant nail deformity. Meticulous nail bed repair is essential for preventing postoperative nail deformities. Pulp lesions can be approached through a lateral incision.

Figure 3. A 67-year-old woman with subungual glomus tumor of the right thumb. Preoperative...
In addition to nail deformity, recurrence is a possible complication and may occur in up to 20% of cases. Recurrence is thought to be a result of incomplete excision or, in the case of late recurrence, development of a new lesion at or near the excision site. Excision of the capsule of the tumor is required to prevent local recurrence.

**MALIGNANT VARIANT**

Glomangiosarcoma is an exceptionally rare malignant variant of the glomus tumor. It tends to appear as a painful nodule located in the subcutaneous tissue. Histology of the glomangiosarcoma tumor shows features that resemble a benign glomus tumor. Nonetheless, malignant glomus tumors arise de novo. This neoplasm is considered a low-grade malignant tumor with tendency for local recurrence, although metastasis has been reported. Treatment consists of complete local excision and close surveillance.

**REFERENCES**


