Abstract:

Medical Management of Peyronie’s Disease

Peyronie’s disease (PD) is a wound-healing disorder in which a fibrotic plaque forms in the tunica albuginea layer of the penis. It clinically presents as any combination of penile pain, angulation, and erectile dysfunction. Recent studies indicate that PD has a prevalence of 3%–9% in adult men. Although the exact etiology has not been established, PD likely results from a predisposing genetic susceptibility combined with an inciting event such as microtrauma during intercourse. During the initial acute phase (6–18 months), the condition may progress, stabilize, or regress. For this reason authorities recommend a more conservative treatment approach, with a trial of oral and/or intralesional pharmacotherapy, before surgical reconstruction is considered. Oral therapies most commonly employed include tocopherol (vitamin E) and paraaminobenzoate (Potaba), with colchicine, tamoxifen, propoleum, and acetyl-L-carnitine being used less often. There are a limited number of long-term placebo-controlled studies with these oral agents, and for the most part, studies have failed to show a consistent beneficial effect. Intralesional injection therapy for PD is more commonly used as a first-line therapy. The current approach includes injection with interferon-a-2b, verapamil, or collagenase (Xiaflex, Auxilium, Philadelphia, PA). Other available therapies that have not consistently shown efficacy in placebo-controlled studies include corticosteroids, orgotein, radiation, and extracorporeal shockwave therapy. Surgery is considered when men with PD do not respond to conservative or medical therapy for approximately 1 year and cannot perform satisfactory sexual intercourse. Ongoing basic research in PD will likely identify future targets for medical exploitation.