Author Year; Country Score Research Design Total Sample Size	Methods	Outcome
Kim et al. 1995; USA Prospective controlled trial Level 2 N=20	Population: 13 men with SCI, 7 non-SCI Age: range 19-73 yrs, Duration of erectile dysfunction: range 0.6-27 yrs.  Treatment: Papaverine gel or placebo gel, dose range: 133-500mg.  Outcome Measures: Safety and efficacyof topical papaverine gel.	For SCI patients (n=12)  1. 3 patients with papaverine gel had full erections, but full erections also occurred with placebo gel.
Kim et al.1995; USA Pre-post Level 4 N=10	Population: Men with SCI (n=9), 1 arterial insufficiency; Age: mean 33 yrs, range 19-50; Injury level: cervical (n=4), thoracic (n=5)  Treatment: Topical prostaglandin E1 to penis, scrotum, and perineum.  Outcome measures: Color flow Doppler Ultrasound for cavernous artery diameter and peak systolic flow velocity, vital signs: systolic blood pressure (SBP), diastolic blood pressure (DBP), heart rate (HR), clinical erection, adverse events.	<ol> <li>Mean cavernous artery diameter increased from 0.09 to 0.11 cm.</li> <li>Mean peak systolic flow velocity increased from 15.4 to 22.8 cm/sec.</li> <li>Clinical erections were observed in 2 men.</li> <li>Vital signs were unaffected by PGE1.</li> <li>No adverse events.</li> </ol>
Chancellor et al. 1994; USA Post-test Level 4 N=18	Population: 18 men with SCI, Age: range 19-65 yrs, Level of injury: C7-L3 (15 thoracic).  Treatment: Minoxidil spray, papaverine injection, or vacuum constriction device (VCD).  Outcome Measures: Erectile response.	<ol> <li>Papaverine injections increased median rigidity 77% (range 30-100%). Rigidity was significantly less with minoxidil.</li> <li>Vacuum constriction device changed rigidity a median of 57% (range 30-80%).</li> <li>No difference between vacuum constriction device and papaverine.</li> <li>Patient subjective rating scale was significantly lower for minoxidil than vacuum constriction device or papaverine.</li> <li>Physicians' subjective ratings were significantly lower for minoxidil than other treatments.</li> </ol>
Beretta et al. 1993; Italy Post-test Level 4 N=15	Population: 15 men, age range: 20-38 yrs, Level: T2-L5. Treatment: Prostaglandin E1 and 2% Minoxidil solution. Outcome Measures: Erectile response.	<ol> <li>4 patients had complete responses, 5 had partial, 6 had no response.</li> <li>9 patients with complete/partial response continued to use minoxidil at home for 1 month.</li> <li>26.6% obtained an erectile response sufficient for vaginal penetration.</li> </ol>
Sonksen et al. 1992; Denmark Post-test Level 4 N=17	Population: 17 men, age range: 19-51 yrs, level: C2-L4, 13 complete, 4 incomplete.  Treatment: Transiderm-Nitro plaster (10mg/24hrs), which contains 50mg glyceryl trinitrate.  Outcome Measures: Erectile response.	<ol> <li>5 patients had complete responses (full rigidity), 7 had partial responses (some rigidity and/or increase in penile circumference), and 5 had no response (no noticeable erection).</li> <li>Erection duration (complete response): 20-45 min.</li> <li>5 (29%) had erections sufficient for vaginal penetration.</li> </ol>