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Minimally invasive therapy with intralesional onabotulinum toxin A in peyronie disease

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Objective: To determine the effectiveness of intralesional application of onabotulinum toxin A in patients with Peyronie disease.

Materials & Methods: A prospective therapeutic cohort study was undertaken in patients ≥18 year with stable disease were included. Intervention: One-time intralesional application of 100 IU of onabotulinum toxin A. We included 22 patients from Urology consult from october 1st 2011 to june 30th 2012. Primary outcome measure: Grade of curvature. Secondary outcome measures: Thickness of the fibrous plaque, erectile dysfunction improvement and pain. Statistical analysis included the Pearson chi-square test for categorical variables and the student t test for quantitative variables. Any p value <0.05 was considered statistically significant.

Results: The size of the fibrous plaque was reduced from 0.34 ± 0.20 cm to 0.27 ± 0.13 cm after treatment (p=0.014). The curvature initially averaged $32.95\pm9.21^\circ$, improving to $25\pm9.38^\circ$ (p=0.025). According to the Kelami classification, the curvature was <30° in 14 cases (63.6%) and was 30–60° in 8 cases (36.4%). At 16 wk, the curvature was <30° in 19 cases (86.4%) and 30–60° in 3 cases (13.6%). The erectile dysfunction grade was 16.18 ± 4.46 before treatment and 18.22 ± 4.55 after treatment (p=0.002). Pain was reduced from 3.36 ± 3.48 before treatment to $1.14\ 1.58$ after treatment (p=0.001).

Conclusions: The application of onabotulinum toxin A can improve the clinical manifestations of Peyronie disease due to fibrosis, increasing sexual function in affected patients.

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