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INSTALLATION OF HYALURONAN VIA ELECTROMOTIVE DRUG ADMINISTRATION CAN IMPROVE THE EFFICACY OF THE TREATMENT IN PATIENTS WITH INTERSTITIAL CYSTITIS/PAINFUL BLADDER SYNDROME: A RANDOMIZED PROSPECTIVE STUDY

Hypothesis / aims of study

In the treatment of bladder pain syndrome/interstitial cystitis (BPS/IC) intravesical hyaluronan application is an accepted treatment. In this randomized prospective study it is aimed to identify whether installing the hyaluronan with EMDA can increase the tissue uptake and improve the efficacy.

Study design, materials and methods

The data of 31 patients who had been diagnosed with BPS/IC between 2004–2005 were examined. The patients were randomized to two groups, patients in group A received hyaluronan directly with a catheter and patients in group B received with EMDA. The patients were followed for 24 months and two groups were compared at certain time intervals.

Results

There were 6 males and 25 females. The two groups were similar for baseline parameters and efficacy of treatment with EMDA was better at month 6 and 12. The difference between the two groups was not significant at month 1 and 24. Also treatment with EMDA, positive KCl test and pretreatment voiding frequency > 17 were found to be associated with higher response rates

Concluding message

Hyaluronan installation is an effective GAG substitution therapy in patients with BPS/IC. To installate HA via EMDA can improve the efficacy of the treatment, however lack of the long term efficacy is the major problem of this GAG substitution therapy.

Parameter	Group A (n=15)	Group B (n=16)	P value
Age (mean ± SD)	43,5 ± 4,2	42,8 ± 4,0	>0.05
Voiding frequency (/day)	18.3 ± 5.3	18.0 ± 4,9	
Nocturia (/night)	4.1 ± 1.1	3.9 ± 1.1	
Mean voided volume (ml)	147± 34,2	155± 40,3	
VAS	6.7 ± 1,2	6.8± 1.4	
Symptom score (0-20)	15.8 ± 2,3	15.6 ± 2,3	
Problem score (0-16)	13.1 ± 1,9	13.8 ± 2.0	
(+) KCl test (%)	40.0	43.7	

Table 1

Disclosures

Funding: no disclosures **Clinical Trial:** Yes **Public Registry:** No **RCT:** Yes **Subjects:** HUMAN **Ethics not Req'd:** the study is established under routine investigation protocols of our department and all participants signed the informed consent form.

Helsinki: Yes **Informed Consent:** Yes