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## TREATMENT SATISFACTION WITH IMIDAFENACIN FOR LUTS ; FROM A MULTICENTER CROSS-SECTIONAL SURVEY

### Hypothesis / aims of study

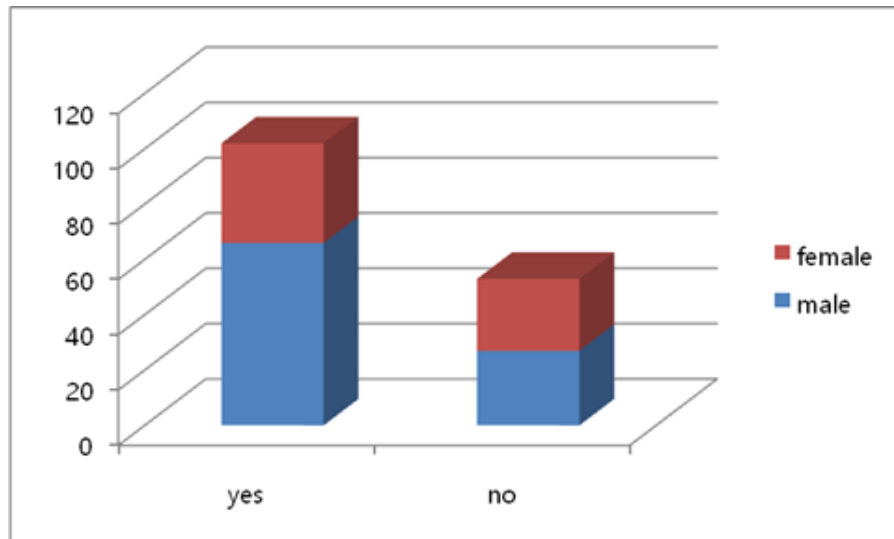
To evaluate the efficacy and treatment satisfaction of Imidafenacin in patients with lower urinary tract symptoms

### Study design, materials and methods

A prospective, cross-sectional study was conducted from July ,2016 to July ,2016 in order to evaluate the patients' satisfaction after 8-weeks treatment with Imidafenacin. The questionnaire survey has been done with 155 patients from 5 centers who were 30 - 86 years old men and women and has suffered OAB symptom for over than 3 months. The comparison survey of treatment satisfactions by the different groups has been settled by Gender, Dose(0.1mg vs >0.1mg), and Co-medication(none vs minirin, A blocker or multiple)

### Results

The average age of 155 patients are 67.5 years old (38~86 years old) and 40% of female(62 patients) and 60% of male(93 patients). 65.8% of total patients(102) were satisfied with the treatment of Imidafenacin, but 34.2%(53) were not. The male group's satisfaction ratio was higher than the female's (70.96% vs 58.06%). The main reasons of dissatisfaction were efficacy(77.35%) and side effects(13.21%). The satisfaction by the different dose was higher in the group who took 0.1mg than the other group over 0.1mg (70.7% vs 60.5%). Regarding the co-medication (minirin or a blocker) , the group without any other meditation has higher treatment satisfaction than with co-medication(70.8% vs 64.0%).



### Interpretation of results

One third of total patients has been dissatisfied with the treatment using Imidafenacin, and group, the dissatisfaction ratio was higher in female than male.

### Concluding message

The main reason was efficacy, so LUTS can be more improved if the treatment dose of Imidafesacin is increased earlier.

### References

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### Disclosures

**Funding:** none **Clinical Trial:** Yes **Public Registry:** No **RCT:** No **Subjects:** HUMAN **Ethics Committee:** institutional review board of each center **Helsinki:** Yes **Informed Consent:** Yes