# Effectiveness of Solifenacin Succinate in the Treatment of Mixed Urinary Incontinence: A Randomized Controlled Trial







## Affiliations to disclose:

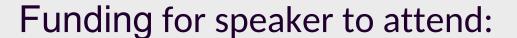
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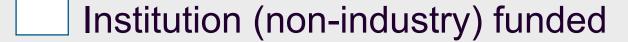
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No conflict of interest to disclose







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# THE TYPES OF URINARY INCONTINENCE













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## **Double-blind randomized controlled trial:**

□Taleghani Hospital in Tehran □2022\_2024

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# A total of 200 women diagnosed with mixed urinary incontinence

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### **Inclusion Criteria**

- ☐ Be 18 years or older
- ☐ No treatment with any other medication
- ☐ Given a written informed consent to participate in the study

## **Exclusion Criteria**

- 1. Use of other antimuscarinic medications besides solifenacin
- 2. Failure to complete the full treatment course or irregular solifenacin use
- 3. Diagnosis of high Post-Void Residual (PVR) volume via ultrasound
- 4. Presence of Stage 3 or 4 pelvic organ prolapse
- 5. Urinary tract infection
- 6. Diabetes
- 7. History of urinary tract surgery

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## **Self-Report Questionnaires**

1. Demographic and clinical data collection: Includes variables such as age, BMI classification, Delivery history, physical activity classification, history of medication use (especially hormonal therapy), and history of any diseases.

2. Incontinence diagnosis: Used self-report Questionnaire for Urinary Incontinence Diagnosis( QUID ) as standardized diagnostic tool.

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# 200 patients were randomly divided equally into two groups as follows:

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# **Experimental Group:**

Received oral
administration of
Solifenacin Succinate
(VSOL5) at a dose of 5
milligrams. The medication
was taken in tablet form
after breakfast for a period
of three months
in Tehran,
Iran.



# **Control Group:**

Received a placebo tablet that was visually identical to Solifenacin Succinate but contained no active ingredient. This was also administered orally after breakfast for a period of three months.

The severity of urinary incontinence was assessed using the validated QUID (Questionnaire for Urinary Incontinence Diagnosis) at baseline, 1.5 months, and 3 months post-treatment.

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# What Is QUID?

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#### The Questionnaire for female Urinary Incontinence Diagnosis (QUID)

Do you leak urine (even small drops), wet yourself, or wet your pads or undergarments	None of the time	Rarely	Once in a while	Often	Most of the time	All of the time
2. when you bend down or lift something up?	0		0	0	0	
3. when you walk quickly, jog or exercise?	п		0		п	п
4. while you are <b>undressing</b> in order to use the <b>toilet</b> ?	0	п	0	0	0	П
5. Do you get such a strong and uncomfortable need to urinate that you leak urine (even small drops) or wet yourself before reaching the toilet?			П		0	
6. Do you have to <b>rush to the bathroom</b> because you get a <b>sudden, strong need</b> to urinate?			П	а	п	П

#### Scoring:

Each item scores 0 (None of the time), 1 (Rarely), 2 (Once in a while), 3 (Often), 4 (Most of the time) or 5 (All of the time). Responses to items 1, 2 and 3 are summed for the Stress score; and responses to items 4, 5, and 6 are summed for the Urge score.

- □ Physical activity level based on the questionnaire was 3.59 ± 1.96 in the intervention group and 3.56 ± 2.00 in the control group (p = 0.880), showing no significant difference.
- Age: In the intervention group, the mean ± standard deviation of age was 52.80 ± 13.78, while in the control group it was 51.65 ± 11.90 (p= 0.653), indicating no significant difference.
- □ Body Mass Index (BMI) in the intervention group was 26.89 ± 3.10, and in the control group it was 26.95 ± 2.99 (p = 0.806), again showing no significant difference.
- Number of pregnancies in the intervention group was 3.01 ± 1.79 and in the control group it was 2.98 ± 1.74 (p = 0.964), which was not statistically significant.

# Results

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- □ Participants in the solifenacin group demonstrated a statistically significant reduction in QUID scores at both follow-up points compared to the placebo
- ☐ At 1.5 months, the mean QUID score was 6.65 ± 1.08 in the intervention group versus 14.80 ± 2.52 in the control group (p < 0.001)
- $\Box$  At 3 months, scores were 3.25 ± 1.11 vs. 9.98 ± 4.34 (p < 0.001)



The magnitude of reduction from baseline was significantly greater in the solifenacin group (-8.47 and -11.87 at 1.5 and 3 months, respectively) than in the control group (-0.88 and -5.70, respectively) with p < 0.001 for all comparisons.

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# **Discussion**

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www.icseus.org/2025/abstract/ DOI: 10.1111/luts.12476

#### ORIGINAL ARTICLE

WILEY

TOT in combination with solifenacin or intravaginal prasterone in postmenopausal women with mixed urinary incontinence: A retrospective analysis in 112 patients

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Federica Sala<sup>1,2</sup> | Melania Loggia<sup>1,2</sup> | Giorgia Cardella<sup>1,2</sup> | Claudia Morgani<sup>1,2</sup> | Giovanni Grossi<sup>2</sup> | Marzio Angelo Zullo<sup>3</sup> | Herbert Carmelo Carlo Valensise<sup>1</sup> | Pier Luigi Palazzetti<sup>2</sup> | Michele Carlo Schiavi<sup>2</sup>
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The results of these study are consistent

Original Article

MEDICINE

DOI: 10.32322/jhsm.1106031

11 levith 5ct Med 2022, 707, 1207, 1211.

Comparison of tolterodine, trospium chloride, solifenacin treatments and its side effects on patients with pure urinary and mixed incontinence

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# Explanation of the mechanism behind this effectiveness





▶ Pharmaceuticals (Basel). 2024 Jan 16;17(1):116. doi: 10.3390/ph17010116

## Regular and Irregular Use and Reasons for Discontinuation of Solifenacin Therapy in Patients with Overactive Bladder Managed by Urologists

Mateusz Małkowski <sup>1</sup>, Agnieszka Almgren-Rachtan <sup>2,\*</sup>, Magdalena Olszanecka-Glinianowicz <sup>3</sup>, Jerzy Chudek <sup>4,\*</sup>, Piotr Chłosta <sup>5</sup>

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▶ Braz J Med Biol Res. 2022 Jan 25;55:e11721. doi: 10.1590/1414-431X2021e11721 🗷

Treatment of bladder dysfunction with solifenacin: is there a risk of dementia or cognitive impairment?

<u>LP Dantas</u> <sup>1,2</sup>, <u>ARCC Forte</u> <sup>1</sup>, <u>BC Lima</u> <sup>1</sup>, <u>CNS Sousa</u> <sup>1</sup>, <u>EC Vasconcelos</u> <sup>1</sup>, <u>PHC Lessa</u> <sup>3</sup>, <u>RF Vieira</u> <sup>3</sup>, <u>MCA Patrocínio</u> <sup>4,5</sup>, SMM Vasconcelos <sup>1</sup>

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Studies have shown that prolonged use of certain muscarinic antagonists—especially those with high lipophilicity—can increase the risk of cognitive decline. These substances have different pharmacological properties, including the ability to cross the blood-brain barrier, and affect central nervous system regions such as the hippocampus and cerebral cortex. As a result, they may cause adverse effects on memory and cognition.

# Due to the limited sample size

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☐ Cannot be generalized to all patients with the same condition.

☐ In addition to this, the presence of various confounding factors and the complexity of the situation make it difficult to draw definitive conclusions from these results.

☐ It is clearly evident that the observed effect was present in this particular study, and the findings are acceptable and justifiable within the context of the research. Broader and more extensive studies across larger populations are needed to confirm these results.

## Conclusion

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- ☐ Solifenacin succinate represents a promising pharmacologic option for treating mixed urinary incontinence in women.
- ☐ Its efficacy and tolerability suggest it could be an effective first-line or adjunctive treatment, particularly in cases where behavioral therapies alone are insufficient.

