

News in LUTS pharmacotherapy

Poster Session 40

Sunday, 26 March
14:00 - 15:30

Location: Room Milan, North Hall (Level 1)

Chairs: J.C. Nickel, Kingston (CA)
P. Nyirády, Budapest (HU)
N. Thiruchelvam, Cambridge (GB)

Aims and objectives of this session

The objectives of this session is to understand how new pharmacologic research will impact on our understanding of LUTS and learn to use this knowledge to improve our management of storage and voiding LUTS associated with benign prostatic hyperplasia, overactive bladder and other causes of LUTS.

Poster viewing of 20 minutes. Presentations will take place on stage. Standard presentations are 2 minutes in length, followed by 2 minutes for discussion. Extended presentations (*) are 3 minutes in length, followed by 3 minutes for discussion.

- *531 **Impact of 5-alpha reductase inhibitors for treatment of benign prostatic hyperplasia on erectile dysfunction, treated depression, gynecomastia, and breast cancer: A real world 20 year observational study**
By: Hagberg K.W.², Divan H.A.³, Persson R.², Fang S.C.³, Jick S.S.², Nickel J.C.¹
Institutes:¹Queen's University, Dept. of Urology, Kingston, Canada, ²Boston University School of Public Health, Boston Collaborative Drug Surveillance Program, Lexington, United States of America, ³New England Research Institutes, NERI, Watertown, United States of America
- 532 **Phosphodiesterase inhibitors for BPH-LUTS: Is the benefit worth it?**
By: Pattanaik S.¹, Mayuduru R.², Panda A.³, Mathew J.⁴, Aggarwal M.⁵, Singh S.², Mandal A.²
Institutes:¹Postgraduate Institute of Medical Education and Research, Dept. of Pharmacology, Chandigarh, India, ²Postgraduate Institute of Medical Education and Research, Dept. of Urology, Chandigarh, India, ³CMC, Dept. of Urology, Vellore, India, ⁴Postgraduate Institute of Medical Education and Research, Dept. of Pediatrics, Chandigarh, India, ⁵NMC Specialty Hospital, Dept. of Urology, Abudhabi, United Arab Emirates
- *533 **Antimuscarinic use in the elderly: A poisoned apple?**
By: Meyer C.¹, Pucheril D.², Karabon P.², Gild P.¹, Von Landenberg N.¹, Atiemo H.², Menon M.², Chughtai B.³, Fisch M.⁴, Chun F.⁴, Trinh Q-D.¹
Institutes:¹Brigham and Women's Hospital, Harvard Medical School, Division of Urological Surgery and Center For, Division of Urologic Surgery and Center For Surgery and Public Health, Boston, United States of America, ²Henry Ford Health System, VUI Center for Outcomes Research, Analytics and Evaluation, Vattikuti Urology Institute, Detroit, United States of America, ³Weil Cornell Medical College/New York Presbyterian Hospital, Dept. of Urology, New York, United States of America, ⁴University Medical Center Hamburg-Eppendorf, Dept. of Urology, Hamburg, Germany
- 534 **A 52-week randomized comparative study of a triple therapy (tamsulosin, dutasteride, and imidafenacin) versus a dual therapy (tamsulosin and dutasteride) in benign prostatic hyperplasia patients with overactive bladder (DirecT Study)**
By: Yamanishi T.¹, Asakura H.², Seki N.³, Tokunaga S.⁴
Institutes:¹Dokkyo Medical University, Dept. of Urology, Tochigi, Japan, ²Saitama Medical University Hospital, Dept. of Urology, Saitama, Japan, ³Kyushu Central Hospital, Dept. of Urology, Fukuoka, Japan, ⁴Kyushu University Hospital, Medical Information Center, Fukuoka, Japan

- 535 **Comparison between tadalafil 5 mg vs. Serenoa repens/selenium/lycopene for the treatment of benign prostatic lower urinary tract symptoms secondary to benign prostatic hyperplasia. A phase IV, randomized, multicenter, non-inferiority clinical study. SPRITE study**
By: Morgia G.¹, Vespasiani G.², Reale G.¹, Di Mauro M.¹, Pareo R.³, Voce S.⁴, Madonia M.⁵, Fedelini P.⁶, Veneziano P.⁷, Carini M.⁸, Salvia G.⁹, Santaniello F.¹⁰, Ginepri A.¹¹, Bitelli M.¹², Terrone C.¹³, Gentile M.¹⁴, Giannantoni A.¹⁵, Blefari F.¹⁶, Beatrici V.¹⁷, Polledro P.¹⁸, La Rosa P.¹⁹, Arnone S.²⁰, Santelli G.²¹, Russo G.I.¹
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- 536 **A randomized, open-label, multicenter study evaluating efficacy of switch from dutasteride to tadalafil in benign prostatic hyperplasia patient with lower urinary tract symptoms (D-to-T trial)**
By: Matsumoto T.¹, Hatakeyama S.¹, Yoshikawa K.², Fukui K.³, Yanagisawa T.⁴, Kawaguchi T.⁵, Imai A.¹, Yoneyama T.¹, Hashimoto Y.¹, Koie T.¹, Saito H.⁶, Yamaya K.⁶, Funyu T.⁶, Ohyama C.¹
Institutes:¹Hirosaki University Graduate School of Medicine, Dept. of Urology, Hirosaki, Japan, ²Mutsu General Hospital, Dept. Of Urology, Mutsu, Japan, ³Fukui Urology Clinic, Dept. of Urology, Aomori, Japan, ⁴Aomori Rosai Hospital, Dept. of Urology, Hachinohe, Japan, ⁵Aomori Prefectural Central Hospital, Dept. of Urology, Aomori, Japan, ⁶Oyokyo Kidney Research Institute, Dept. of Urology, Hirosaki, Japan
- 538 **Impact of Vesomni™ on quality of life of men with lower urinary tract symptoms associated with benign prostatic hyperplasia in routine clinical practice: Interim results from the EUROPA study**
By: Rees J.¹, Arbe E.⁷, Rosa Arias J.², Skoumal R.³, Walters C.⁴, Yavuz Y.⁵, De Wachter S.⁶
Institutes:¹Tyntesfield Medical Group, Brockway Medical Centre, Nailsea, United Kingdom, ²Hospital "Santiago Apóstol", Dept. of Urology, Miranda De Ebro, Spain, ³Urocentrum Brno, Dept. of Urology, Brno, Czech Republic, ⁴Astellas Pharma Europe Ltd, Medical and Clinical Operations, Chertsey, United Kingdom, ⁵Astellas Pharma Global Development, Dept. of Data Sciences, Leiden, The Netherlands, ⁶University Hospital Antwerpen, University Antwerpen, Dept. of Urology, Edegem, Belgium, ⁷Astellas Pharma Europe Ltd, Dept. of Medical Affairs, Chertsey, United Kingdom
- 539 **Post-operative continuous use of antimuscarinics in BPH patients with storage symptoms requiring antimuscarinics before surgery – a nationwide population-based study**
By: Huang E.Y.-H., Chung H.-J., Lin C.-C., Peng R.-S., Chang Y.-H., Lin A.T.-L., Chen K.-K.
Institutes:Taipei Veterans General Hospital, Dept. of Urology, Taipei, Taiwan
- 540 **A multicenter real-life study of the efficacy of an alpha-blocker with or without anticholinergic agent (imidafenacin) treatment in patients with lower urinary tract symptoms/benign prostatic hyperplasia and storage symptoms**
By: Cho S.¹, Hoon C.¹, Park J.Y.¹, Bae J.H.¹, Lee K.W.², Yoo T.K.³, Sin D.G.⁴, Kim S.W.⁵, Kim Y.H.²
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- 541 **Testosterone therapy (TTh) improves urinary function and reduces major adverse cardiovascular events (MACE) in hypogonadal men with type 2 diabetes (T2DM) treated up to 8 years in comparison to an untreated control group**
By: Haider A.¹, Haider K.¹, Doros G.², Traish A.³

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The effect of non-steroidal anti-inflammatory drugs on risk of benign prostatic hyperplasia

By: Murtola T.¹, Nygård L.², Talala K.³, Taari K.⁴, Tammela T.¹, Auvinen A.⁵

Institutes:¹Tampere University Hospital, Dept. of Urology, Tampere, Finland, ²University of Tampere, School of Medicine, Tampere, Finland, ³Finnish Cancer Registry, Dept. of Research, Helsinki, Finland, ⁴Helsinki University Hospital, Dept. of Urology, Helsinki, Finland, ⁵University of Tampere, School of Health Sciences, Tampere, Finland

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The effect of statins on the risk of receiving transurethral resection of prostate in the outpatients of genitourinary clinic - a study by applying nation-wide population based database

By: Lin C-C.¹, Chung H.J.¹, Lin A.T.L.¹, Huang Y.H.¹, Chen T.Z.²

Institutes:¹Taipei Veterans General Hospital, Dept. of Urology, Taipei, Taiwan, ²Taipei Veterans General Hospital, Dept. of Family Medicine, Taipei, Taiwan

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The comparison in the efficacy of the two combination therapies with an anticholinergic agent and an α 1-blocker versus a α 3-adrenoceptor agonist and an α 1-blocker for patients with benign prostatic enlargement complicated by overactive bladder: A randomized, prospective trial using a urodynamic study

By: Matsukawa Y., Matsuo K., Majima T., Narita H., Kato M., Yamamoto T., Gotoh M.

Institutes:Nagoya University Graduate School of Medicine, Dept. of Urology, Nagoya, Japan