

Volatile organic compounds for urinary tract infection

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Policy contains: Electronic noses, mass spectroscopy, urinary tract infection, volatile organic compounds.

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Coverage policy

Volatile organic compounds for diagnosing urinary tract infection are investigational/not clinically proven and, therefore, not medically necessary.

Limitations

No limitations were identified during the writing of this policy.

Alternative covered services

- Laboratory analyzed Urine culture and sensitivity.
- Urine dipstick test.

Background

There are multiple challenges associated with the diagnosis of urinary tract infections. Current methods such as gas chromatography mass spectroscopy are expensive, require trained personnel, and are time consuming. In addition, hematuria or chronic urinary catheter use for a neurogenic or anatomically impaired bladder can complicate diagnosis (Dospinescu, 2020).

Bacteria are present in the tissues around the urethral opening and often colonize the urine. Because bacteria are more likely to ascend to the female bladder, which has a shorter urethral length, rates of urinary tract infection are higher among women. Various gram-positive and gram-negative bacteria, most commonly *Escherichia coli*, cause most urinary tract infections. Common infection diagnostic terms include cystitis, hemorrhagic cystitis, pyelonephritis, and catheter-associated urinary tract infection (Flores-Mireles, 2015; Foxman, 2014).

Risk factors for urinary tract infection include female sex; shorter urethra; prior infection; advanced age; recent sexual intercourse; use of a condom, diaphragm, or spermicide; vaginal infection; trauma/manipulation; diabetes; obesity; genetic susceptibility; or anatomic abnormalities. The estimated lifetime risk of urinary tract infection for women, based on self-reported history of diagnosis by a physician, is 60.4%. Recurrence is common. Most complicated infections are attributed to indwelling catheters (Flores-Mireles, 2015; Foxman, 2014).

Antibiotics are the standard treatment for urinary tract infections. The ever increasing rising rates of antibiotic resistance, combined with high recurrent infection rates are of great concern to health care providers, and underline the need for alternative therapies less susceptible to resistance (Flores-Mireles, 2015). An estimated 10% of urinary tract infections are resistant to antibiotics (Smart, 2019).

Improved diagnosis may enhance appropriate selection of antibiotics. Current diagnosis of urinary tract infection relies on two options. One is a dipstick test, which is rapid but often not accurate; one study places sensitivity and specificity, compared to lab culture, at 75.7% and 68.9% (Najeeb, 2015). The alternative is culturing to identify a pathogen, which takes 24 - 72 hours; during this delay, broad-spectrum antibiotics may be prescribed, leading to resistance (Dospinescu, 2020).

Volatile organic compounds are carbon-based compounds that can originate from microbial pathogens or a host response to infection and inflammation; many are associated with common urinary tract pathogens. These compounds can improve diagnosis of urinary tract infections. In addition, volatile organic compounds have been proposed as a diagnostic or screening tool for other conditions; one of the most commonly cited is cancer, although these uses are not established (Brusselmans, 2018; Catino, 2019; Farraia, 2022; Oakley-Girvan, 2017; Zhou, 2020). Several technologies have been developed, including gas chromatography, proton transfer reaction mass spectrometry, ion mobility spectrometry, selected ion flow tube mass spectrometry, field asymmetric ion mobility spectrometry, gas chromatography flame ionization detection, and odor sensors (Dospinescu, 2020).

In 2001, the U.S. Food and Drug Administration granted premarket approval for use of the Osmetech Microbial Analyser for diagnosis of urinary tract infection. The product uses "electronic nose" (odor sensor) technology to measure the presence of bacteria by semiquantitative analysis of volatile compounds (U.S. Food and Drug Administration, 2001).

Findings

The American Academy of Family Physicians practice guidelines on urinary tract infection for children/infants and adults only mentions urine microscopy and dipstick testing as diagnostic methods (Michels, 2015; Veauthier, 2020). For diagnosing recurrent urinary tract infections and asymptomatic bacteriuria, the American Urological Association and the Infectious Diseases Society of America recommend urine culture but do not mention volatile organic compound testing (Anger, 2022; Nicolle, 2019).

A systematic review of 25 studies and meta-analysis of ten studies examined the efficacy of portable electronic noses for diagnosis or monitoring various pathologies through analysis of urine samples, including four studies (n = 211) using different electronic nose systems for diagnosis of urinary tract infections. The authors stated the heterogeneity of the diagnostic measurements and findings did not permit conclusions about their use for diagnosing urinary tract infections and called for additional research and standardization of analytical methods (Afonso, 2022).

One review concludes that of the existing models, electronic noses and ion mobility spectrometry systems are still the most suitable candidates in the diagnosis of urinary tract infection, since they are easy to use, portable, relatively low-cost, and have methods which can be automated (Dospinescu, 2020).

A study of 84 urine samples that tested for 85 volatile organic compounds identified five isolates positively associated with *Escherichia coli*-resistant strains, and two with sensitive strains of urinary tract infections. The accuracy of identifying resistant and sensitive strains was 91.1% and 79.5%, respectively (Hewett, 2020).

A study used thermal desorption-gas chromatography-mass spectrometry to 'smell' antibiotic-resistant bacteria in 18 bacterial isolates (*Escherichia coli, Klebsiella pneumoniae,* and *Pseudomonas aeruginosa*). The isolates were grown with and without the presence of antibiotic. Nine and 22 compounds differed significantly between cephalexin and ciprofloxacin sensitive/resistant isolates, respectively (P < .05) (Smart, 2019).

In 2023, we added two guidelines (Anger, 2022; Nicolle, 2019) and one new systematic review (Afonso, 2022) and deleted two older individual studies and two studies that were assessed in the systematic review. No policy changes are warranted.

In 2024, no new relevant literature was found. No policy changes warranted.

References

On November 2, 2024, we searched PubMed and the databases of the Cochrane Library, the U.K. National Health Services Centre for Reviews and Dissemination, the Agency for Healthcare Research and Quality, and the Centers for Medicare & Medicaid Services. Search terms were "volatile organic compounds" (MeSH), "urinary tract infections" ([MeSH), "electronic noses," "Mass spectroscopy," "urinary tract infection," and "volatile organic compounds." We included the best available evidence according to established evidence hierarchies (typically systematic reviews, meta-analyses, and full economic analyses, where available) and professional guidelines based on such evidence and clinical expertise.

Afonso HAS, Farraia MV, Vieira MA, Cavaleiro Rufo J. Diagnosis of pathological conditions through electronic nose analysis of urine samples: A systematic review and meta-analysis. *Porto Biomed J*. 2022;7(6):e188. Doi: 10.1097/j.pbj.00000000000188.

Anger JT, Bixler BR, Holmes RS, Lee UJ, Santiago-Lastra Y, Selph SS. Updates to recurrent uncomplicated urinary tract infections in women: AUA/CUA/SUFU guideline. *J Urol*. 2022;208(3):536-541. Doi: 10.1097/JU.000000000002860.

Brusselmans L, Arnouts L, Millevert C, Vandersnickt J, van Meerbeeck JP, Lamote K. Breath analysis as a diagnostic and screening tool for malignant pleural mesothelioma: A systematic review. *Transl Lung Cancer Res.* 2018;7(5):520-536. Doi: 10.21037/tlcr.2018.04.09.

Catino A, de Gennaro G, Gilio AD, et al. Breath analysis: A systematic review of volatile organic compounds (VOCs) in diagnostic and therapeutic management of pleural mesothelioma. *Cancers (Basel)*. 2019;11(6):831. Doi: 10.3390/cancers11060831.

Dospinescu V-M, Tiele A, Covington JA, et al. Sniffing out urinary tract infection-diagnosis based on volatile organic compounds and smell profile. *Biosensors (Basel)*. 2020;10(8):83. Doi: 10.3390/bios10080083.

Farraia MV, Cavaleiro Rufo J, Paciência I, Mendes F, Delgado L, Moreira A. The electronic nose technology in clinical diagnosis: A systematic review. *Porto Biomed J*. 2019;4(4):e42. Doi: 10.1097/j.pbj.00000000000042.

Flores-Mireles AL, Walker JN, Caparon M, Hultgren SJ. Urinary tract infections: Epidemiology, mechanisms of infection and treatment options. *Nat Rev Microbiol*. 2015;13(5):269-284. Doi: 10.1038/nrmicro3432.

Foxman B. Urinary tract infection syndromes: Occurrence, recurrence, bacteriology, risk factors, and disease burden. *Infect Dis Clin North Am.* 2014;28(1):1-13. Doi: 10.1016/j.idc.2013.09.003.

Hewett K, Drabinska N, White P, et al. Towards the identification of antibiotic-resistant bacteria causing urinary tract infections using volatile organic compounds analysis – A pilot study. *Antibiotics (Basel)*. 2020;9(11):797. Doi: 10.3390/antibiotics9110797.

Michels TC, Sands JE. Dysuria: Evaluation and differential diagnosis in adults. *Am Fam Physician*. 2015;92(9):778-788. https://www.aafp.org/pubs/afp/issues/2015/1101/p778.html.

Najeeb S, Munir T, Rehman S, Hafiz A, Gilani M, Latif M. Comparison of urine dipstick test with conventional urine culture in diagnosis of urinary tract infection. *J Coll Physicians Surg Pak*. 2015;25(2):108-110. https://pubmed.ncbi.nlm.nih.gov/25703753/.

Nicolle LE, Gupta K, Bradley SF, et al. Clinical practice guideline for the management of asymptomatic bacteriuria: 2019 update by the Infectious Diseases Society of America. *Clin Infect Dis.* 2019;68(10):e83-e110. Doi: 10.1093/cid/ciy1121.

Oakley-Girvan I, Davis SW. Breath based volatile organic compounds in the detection of breast, lung, and colorectal cancers: A systematic review. *Cancer Biomark*. 2017;21(1):29-39. Doi: 10.3233/CBM-170177.

Smart A, Costello BL, White P, et al. Sniffing out resistance – rapid identification of urinary tract infectioncausing bacteria and their antibiotic susceptibility using volatile metabolite profiles. *J Pharm Biomed Anal.* 2019;167:59-65. Doi: 10.1016/j.jpba.2019.01.044.

U.S. Food and Drug Administration. 510(k) Premarket Notification database. K011043 Osmetech Microbial Analyser Urinary Tract Infection Detector.

https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpmn/pmn.cfm?ID=K011043. Decision date November 30, 2001.

Veauthier B, Miller MV. Urinary tract infections in young children and infants: Common questions and answers. *Am Fam Physician*. 2020;102(5):278-285. https://www.aafp.org/pubs/afp/issues/2020/0901/p278.html.

Zhou W, Tao J, Tao S. Volatile organic compounds analysis as a potential novel screening tool for colorectal cancer: A systematic review and meta-analysis. *Medicine (Baltimore)*. 2020;99(27):e20937. Doi: 10.1097/MD.000000000020937.

Policy updates

- 12/2021: initial review date and clinical policy effective date: 1/2022
- 12/2022: Policy references updated.
- 12/2023: Policy references updated.
- 12/2024: Policy references updated.