

KORPATH

Urine-ID Case Review

• Patient History:

97-year-old male resident in Assisted Living.

• Disease State:

Pain while urinating, cloudy urine, and confusion. Suspected UTI, specifically from E. coli because most urine cultures previously sent in were only positive for E. coli.

• Why This Test was Ordered:

Male cystitis is considered complicated, so a Urine-ID was ordered to look for possibility of coinfections.

• Outcome:

Vikor's full Urine ID results were positive for E coli. Plus, Klebsiella oxytoca at 10⁶, Pseudomonas aeruginosa at 10⁶, Klebsiella pneumoniae at 10⁶, Enterococcus faecalis at 10⁶, Enterobacter cloacae at 10⁵ and 6 Antibiotic Resistant Genes. We were able to show that Urine-ID was able to identify coinfections including a Pseudomonas infection, which due to the potential seriousness, of this type of infection, treatment should be aggressive and monitored closely for treatment effectiveness.

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Electronically approved on 07-09-2021 by: John Ekstrand •Email: pharmconsult@vikorscientific.com • Phone: 1-855-742-7635, 1-855-PharmD5

| | VIKORSCIENTIFIC | | Urine-ID [™] Molecular Pathogen Report |
|----------------------|--|--|--|
| • | 22 WestEdge Street 8th Floor Charleston, SC 29403 Ph# (854) 429-1069 Fx# (833) 247-4091 www.vikorscientific.com | ACCREDITED COLLEGE of AMERICAN PATHOLOGISTS #8359749 | Clinical Euboratory Imendments #42D2150400 |
| | Patient Name | Date of Birth | Gender Race |
| | | XX-XX-1927 | M UNDISCLOSED |
| | Drug Allergies: | NKDA | |
| | Notes from Ordering Physician: | SWAB | |
| MEDICATION REVIEW | Notes from Pharmacist: | The treatment guidance liste treatment references, the org contribute to medication resis such as comorbidities, renal appropriateness of therapy. allergies into account when t take the entire clinical preser treatment decisions. Not all o antimicrobial therapy as som non-pathogenic colonizers. | ed is based on infectious disease ganisms detected, and genes known to istance. Important clinical information function, etc. may influence the overall The provided guidance only takes drug they are provided. The provider should intation into account when making detected microbes will require the are part of the normal flora or can be |
| | | Due to the potential seriousn treatment should be aggress effectiveness. Treatment rec fosfomycin, cefepime, and m | ness of a Pseudomonas infection, sive and monitored closely for treatment commendations include quinalones, neropenem. |
| | | Male cystitis is considered contract treatment recommendations cystitis. | omplicated. The dosing regimens in the below contain dosing for complicated |
| | | Prostatitis due to Enterococcus is more difficult to treat. Acute, mild cases with prompt response may require only 14 days of antibiotics. Chronic infection should be treated longer. Courses shorter than 4 weeks are associated with higher relapse rates. Preferred regimens are ceftriaxone, ciprofloxacin, levofloxacin, and fosfomycin | |
| | | ***NOTE*** Please add CEF recommendations below due | TRIAXONE 1g x 1 dose to the treatment e to prostate involvement. |
| | | Due to patient allergies and/or recommendations are very li | or resistance genes, treatment imited. |

| | Medication | Route | Dose |
|------------|---------------|-------|--|
| FIRST LINE | ciprofloxacin | oral | 500 mg BID x 10-14 days(complicated) |
| | | | Considerations: (Pseudomonas, Klebsiella, Enterococcus, E. Coli, Enterobacter) |
| | | | Fluoroquinolones have been associated with serious and possible irreversible reactions; tendonitis/tendon rupture, peripheral neuropathy, CNS effects. These may occur all together or months after tx. Increased risk in patients over 60 and pt on corticosteroids. Avoid in Myasthenia Gravis. Reserve for pts with no alternative tx options for acute bacterial sinusitis, acute bacterial exacerbation of chronic bronchitis, or uncomplicated UTI. Adjust dose for CrCl <50ml/min. Not recommended in children under 18 y/o. Not first line agent in children with complicated UTI or pyelonephritis. |

| | VIK 22 West Charlest Ph# (854 www.vik | OR SCIENTIFIC Edge Street 8th Floor on, SC 29403 4) 429-1069• Fx# (833) 247-4091 orscientific.com | | COLLEGE # AMERICAN PATHOLOGISTS #8359749 | Urine-ID™ Molecular Pathogen Report |
|------------|---|--|-------------|--|---|
| | Patient N | Name | | Date of Birth Gende | |
| | | OR fosfomycin | oral | 3g q72h x 3 doses(complicated) Considerations: (Pseudomonas, Enterobacter) Do not use for pyelonephritis, C Empty stomach preferred. Now av prostatitis resistant E.coli. | Klebsiella, Enterococcus, E. Coli, an be used for chronic prostatitis. ailable as a generic. can be used in |
| SECOND LIN | JE | meropenem / vaborbactam | intravenous | 4g q8h x 7-10 days Considerations: (Pseudomonas, Enterobacter) Treatment option takes into accour Adjust dose per renal function | Klebsiella, Enterococcus, E. Coli, It the potential resistance |

| Methodology | The infectious disease and antibiotic resistance detection panels are tested utilizing Real-time PCR technology to detect the presence of genes associated with pathogens and antibiotic resistance via amplification of genomic DNA. Amplification and detection are performed using the Applied Biosystems™ QuantStudio™ 12K Flex Real-time PCR system, which includes the QuantStudio™ 12k Software v1.3 and Thermo Fisher Scientific TaqMan™ assays. The assays are preloaded onto TaqMan™ OpenArray plates. |
|-------------|---|
| Limitations | This test only detects microorganisms and antibiotic resistance (ABR) genes specified in the panel. ABR genes are detected in the specimen and are not specific to a detected pathogen. ABR genes may be detected in bacterial strains not tested for in the panel. |
| | The resistance genes for Ampicillin, selected Extended-Spectrum-Betalactamases, Vancomycin, Carbapenems, Sulfonamide, Trimethoprim, Aminoglycosides and the Quinolone gyrase groupings are assays customized by pooling the individual genes listed in the associated group. If listed as positive, this indicates that at least one of the genes in the group was detected and the class of medication could have potential resistance. |
| Disclaimer | This test was developed and its performance characteristics determined by Vikor Scientific™. It has not been cleared or approved by the FDA. The laboratory is regulated under CLIA as qualified to perform high complexity testing. This test is used for clinical purposes. It should not be regarded as investigational or for research. Pharmacy guidance and recommendations therein are not under the purview of the laboratory or agencies which accredit the laboratory. |
| | The treatment guidance listed in the report is based on infectious disease treatment references, the organisms detected, and genes known to contribute to medication resistance. Important clinical information such as comorbidities, renal function, patient weight, platelet count, microbiology results, etc. may influence the overall appropriateness of therapy. The provided guidance only takes drug allergies into account when they are provided and available to the pharmacist making the recommendation. The overall appropriateness of therapy must be determined by the physician treating the patient. The provider has all the patient information necessary to make that determination and should take the entire clinical presentation into account when making treatment decisions. Should the treating physician wish to discuss the provided guidance, the pharmacist is available for consult at the email and phone number provided. |



VanB, VanA1, VanA2

Proteus vulgaris Providencia stuartii Serratia marcescens Staphylococcus aureus

Staphylococcus saprophyticus Streptococcus agalactiae Uncultured Megasphera 1