

A summary of "Isomer interferences observed during the development of a 47-analyte HRAM LC-MS/MS method for urine drug testing"

Overview:

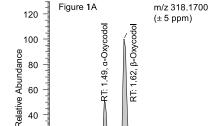
Results:

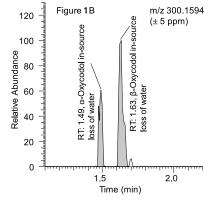
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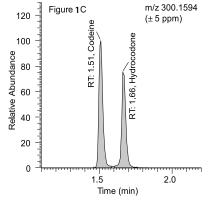
The rise in the use of opioids for pain management and illicit use has increased the need for urine drug testing laboratories to monitor a wide range of opioid analytes. Detection and confirmation of drug analytes in urine is typically done with a preliminary immunoassay and subsequent analysis using GC/MS or LC/MS. Liquid chromatography coupled with quadrupole mass spectrometers and high resolution accurate mass (HRAM) instrumentation is becoming more prevalent for confirmation of drug analytes due to its improved specificity, shorter run times, simpler sample preparation, and lower detection limits. However, laboratories must be aware that this instrumentation may not be able to separate interferences arising from isomeric metabolites that are not typically monitored. The use of a non-selective β -glucuronidase may increase the detection levels of these metabolites. The presence of these interferences was discovered during the development of a 47-analyte HRAM LC-MS/MS UDT method and these interferences are reported here.

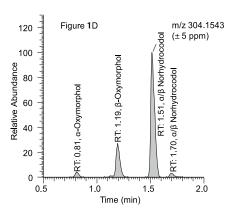
Materials and Methods:

75 μ L of patient urine sample was combined with 300 μ L of master mix containing internal standards and IMCSzyme®. Samples were incubated at 65°C for 60 minutes and centrifuged at 4000 rpm for 7 minutes before analysis on a Waters ACQUITY UPLC® I-Class coupled with a Thermo Scientific Q-Exactive OrbitrapTM.



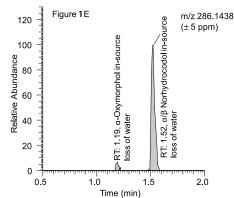






Time (min)

2.0



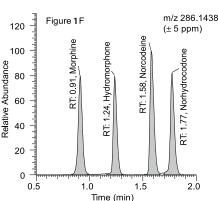


Figure 1. LC-MS/MS Chromatograms of (A) α / β -Oxycodol in a patient specimen positive for Oxycodone; (B) α -Oxycodol and β -Oxycodol loss of water due to in-source fragmentation; (C) Codeine and Hydrocodone in a standard sample (D) α / β -Oxymorphol and α / β -Noroxycodol in a patient specimen positive for Oxycodone; (E) β -Oxymorphol and α or β -Noroxycodol loss of water due to in-source fragmentation; (F)Morphine, Hydromorphone, Norcodeine and Norhydrocodone in a standard sample.

Results Continued:

Table 1. Potential Analyte Interferences

| Analyte | Precursor m/z (H+) | Formula (H+) | Interference | Interference m/z (H+) | Formula (H+) | Source of Interference | Interference loss of water (H+) | Formula loss of water (H+) | Reporting Outcome |
|----------------|-----------------------|-----------------|---|--------------------------|-----------------|---|---------------------------------------|----------------------------------|--|
| Codeine | 300.1594 | C18H22NO3 | Oxycodone metabolites, α/β-oxycodol | 318.1700 | C18H24NO4 | In-source loss of water produces isomeric precursor and product ions | 300.1594 | C18H22NO3 | Over-reporting or false positive for codeine |
| Hydrocodone | 300.1594 | C18H22NO3 | Oxycodone metabolites, α/β-oxycodol | 318.1700 | C18H24NO4 | In-source loss of water pro- duces isomeric precursor and product ions | 300.1594 | C18H22NO3 | Over-reporting or false positive for hydrocodone |
| Norcodeine | 286.1438 | C17H20NO3 | Oxycodone metabolites, α/β-noroxycodol | 304.1543 | C17H22NO4 | In-source loss of water produces isomeric precursor and product ions | 286.1438 | C18H22NO3 | Over-reporting or false positive for norcodeine |
| Norhydrocodone | 286.1438 | C17H20NO3 | Oxycodone metabolites, α/β-noroxycodol | 304.1543 | C17H22NO4 | In-source loss of water produces isomeric precursor and product ions | 286.1438 | C18H22NO3 | Over-reporting or false positive for norhydrocodone |
| Hydromorphone | 286.1438 | C17H20NO3 | Oxycodone metabolites, α/β-oxymorphol | 304.1543 | C17H22NO4 | In-source loss of water pro- duces isomeric precursor and product ions | 286.1438 | C18H22NO3 | Over-reporting or false positive for hydromorphone |
| Morphine | 286.1438 | C17H20NO3 | Oxycodone metabolites, α/β-oxymorphol | 304.1543 | C17H22NO4 | In-source loss of water produces isomeric precursor and product ions | 286.1438 | C18H22NO3 | Over-reporting or false positive for morphine |
| Oxycodone | 316.1543 | C18H22NO4 | Hydrocodone metabolite, hydrocodone N-oxide | 316.1543 | C18H22NO4 | Presumptive hydrocodone metabolite | N/A | | Over-reporting or false positive for oxycodone |
| Oxymorphone | 302.1387 | C18H22NO4 | morphine metabolite/ impurity, morphine N-oxide | 302.1387 | C17H20NO4 | Presumptive hydrocodone metabolite | N/A | | Over-reporting or false positive for oxymorphone |
| Noroxycodone | 302.1387 | C18H22NO4 | Hydrocodone metabolite, N-hydroxynor- hydrocodone | 302.1387 | C17H20NO4 | Presumptive hydrocodone metabolite | N/A | | Over-reporting or false positive for noroxycodone |

Conclusions:

The presence of isomeric metabolites not typically monitored in patient urine samples is something urine drug testing labs must be aware of to avoid over-reporting or obtaining false positives. Because these metabolites are present only in patient samples, external controls are not a good indicator that a method is free of interference. This work provides a summary of some of the potential opioid interferences in an effort to raise awareness of the importance of running patient samples early on in the process of method development.

his information was summarized by IMCS from the poster "Isomer interferences observed during the development of a 47-analyte HRAM LC-MS/MS method for urine drug testing" presented by Ana Grenier Dominion Diagnostics at MSACL 2017