

Trace Detection of Acetamiprid on Raisins

Protecting Consumer Safety with MISA

Summary

Recent test studies conducted by the USDA [1] and the NVWA in Europe [2] have shown that raisins, the popular snack food made from dried grapes, are at the top of the list of fruits and vegetables that have been shown to contain unacceptably high levels of pesticide residues. 80% of imported raisins in the Netherlands are contaminated with an average of 11.3 different pesticides per sample, and nearly all marketed brands of raisins in the US contain at least two different pesticide residues [3]. The fundamental health concern is that the long-term, cumulative effects of consuming a variety of pesticides are unknown. Clearly, this challenges the assumption that raisins provide a child-friendly, healthy alternative to processed snack foods. To address such food safety concerns, there is a need for rapid and accurate testing to screen food samples for potentially hazardous substances. In this Application Note, MISA (Metrohm Instant SERS Analyzer) from Metrohm Raman excels in the detection of the pesticide acetamiprid on commercially sold raisins. MISA is a viable alternative to analytical laboratory testing in the quest to prevent contaminated foods from reaching and harming consumers.

Configuration



2.950.0020 - MISA Advanced

Metrohm Instant SERS Analyzer (MISA) is a high performance, portable analyzer system used for rapid, trace level detection / identification of illicit materials, food additives and food contaminants. MISA features a high-efficiency spectrograph equipped with Metrohm's unique Orbital-Raster-Scan (ORS) technology. It has a minimal footprint and extended battery life, perfect for on-site testing or mobile laboratory applications. MISA offers various Laser Class 1 attachments for flexible sampling options. Analyzer operation is available through BlueTooth or USB connectivity. The MISA Advanced package is a complete package that allows the user to perform SERS analyses using Metrohm's nanoparticle solutions and P-SERS strips. The MISA Advanced package includes a MISA Vial Attachment, a P-SERS Attachment, a ASTM Calibration Standard, a USB Mini Cable, a USB Power Supply and MISA Cal software for operating the MISA instrument. A ruggedized protective case is also provided to securely store the instrument and accessories.



2.926.0120 - MIRA XTR DS Advanced

MIRA XTR DS is an alternative for high power 1064 nm systems. Powered by advanced AI and Machine Learning, MIRA XTR DS uses a more sensitive 785 nm laser along with XTR algorithms to eXTRACT the Raman data from the sample fluorescence. MIRA XTR DS also features Orbital Raster Scanning (ORS) to provide better coverage of the sample increasing the accuracy of the results. The Advanced package includes MIRA Lib KnowItAll™ Full Raman Library Collection of over 19,500 samples, MIRA XTR DS Raman spectral library for Illicit Materials, Calibration Standard, Intelligent Universal Attachment, Right-angle Attachment, Vial Attachment, and MIRA SERS Attachment. A complete package for any type of analysis. Class 3B operation.



607506310 - Silver Colloid, 50 mL

SERS active silver colloidal solution. Potential target analytes that show favorable performance with silver colloid include illicit drugs, medicines, dyes, and amines - 50 ml package.

Introduction

Acetamiprid is a highly effective systemic neonicotinoid insecticide. Although toxicity to humans and other mammals is low, it is moderately to highly toxic for birds and aquatic life, posing a potential threat to wildlife and the food chain. This Application Note demonstrates the rapid and sensitive detection of acetamiprid extracted from raisins using the Metrohm Instant SERS (Surface-Enhanced Raman Scattering) Analyzer.

SERS detection of acetamiprid on raisins

As direct point-and-shoot Raman spectroscopy is unsuitable for trace analyte detection, SERS was used in this experiment. Dilutions of 1 mg/mL acetamiprid in methanol were pipetted onto individual 1 g portions of raisins, yielding samples containing 100, 25, 5, and 2 g/mL (ppm) and 500 ng/mL (ppb) acetamiprid. Each sample was dried and placed in a vial with 0.2 mL of dichloromethane (DCM). Each tube was vortexed for two minutes and rested for 30 minutes, and then the supernatant was transferred to a clean vial for evaporative drying. After 0.9 mL of silver colloid was added, each vial was vortexed for one minute. This was followed by the addition of 0.1 mL of 500 mmol/L NaCl and gentle agitation of the contents. Each vial was inserted into the vial holder attachment of MISA for measurement.



Results and discussion

As shown in **Figure 1**, SERS spectra for DCM extracts of acetamiprid on raisins are identical to the reference spectrum for pure acetamiprid (in dark blue). The highly resolved signature peaks tend to correlate in intensity to analyte concentration.

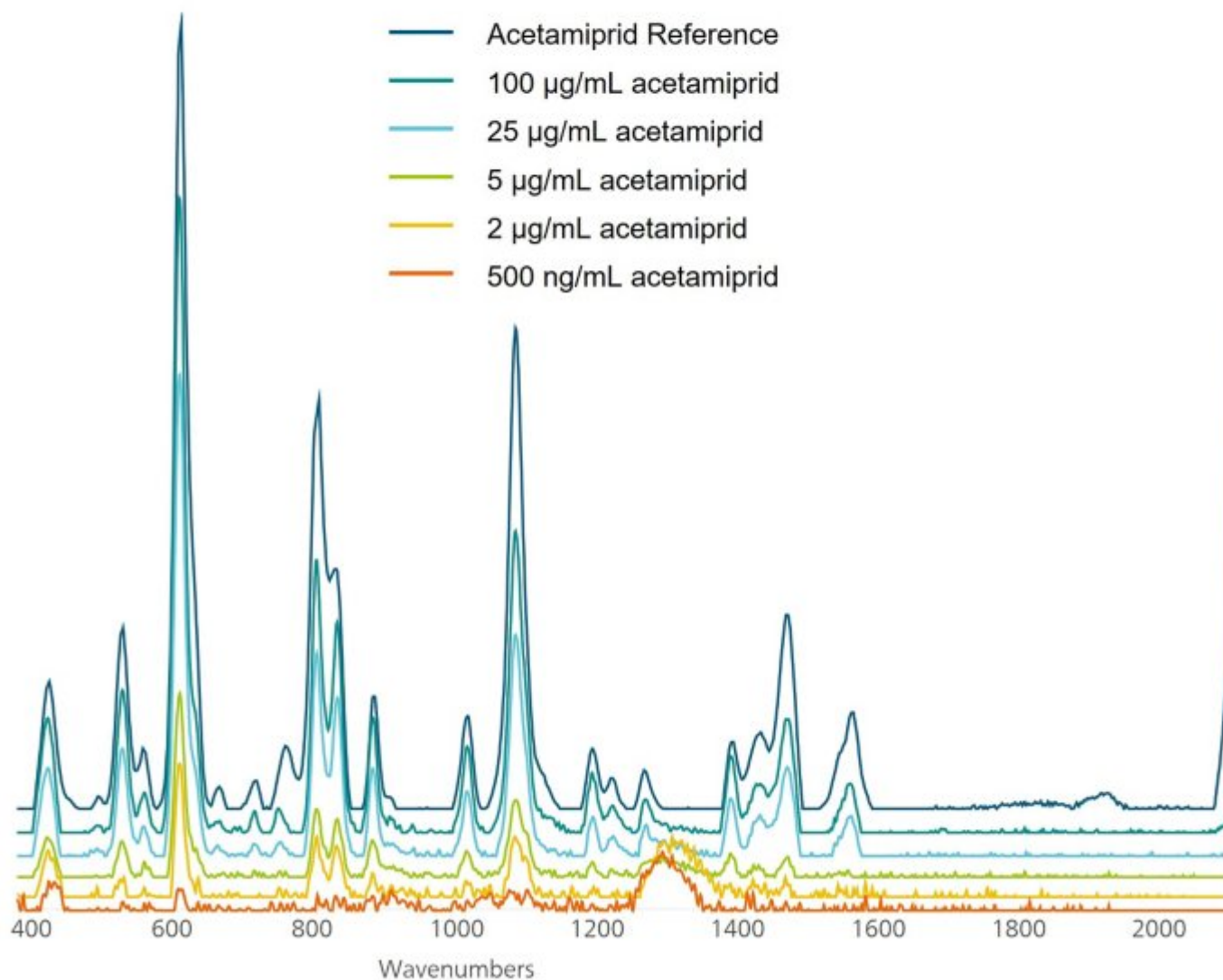


Figure 1. Raman spectra stack of acetamiprid reference and various concentrations (100 g/mL down to 500 ng/mL).

Improving SERS sensitivity

Information content from Raman spectra is greatest at higher analyte concentrations. Some poorly resolved signature peaks in **Figure 1** persist at 500 ng/mL (ppb), yet sensitivity at this level is essential because it corresponds to the maximum residue level accepted for acetamiprid in Europe.

At very low concentrations, the following two strategies may improve SERS detection:

1. Combine multiple extract aliquots into one test sample. In this case, three to four 0.2 mL DCM aliquots (from the same test batch of raisins) would be combined into one vial before evaporative drying.
2. Longer integration times on the instrument may improve sensitivity. The Auto Integration feature on MISA is adequate for higher concentrations; lower concentrations may require manual setting of integration times to four to eight seconds, for example.

Figure 2 overlays spectra for samples containing one aliquot of 5 g/mL, 2 g/mL, and 500 ng/mL acetamiprid with a sample that contains four aliquots of 500 ng/mL acetamiprid. This figure provides visual confirmation for improved signal through combined aliquots.

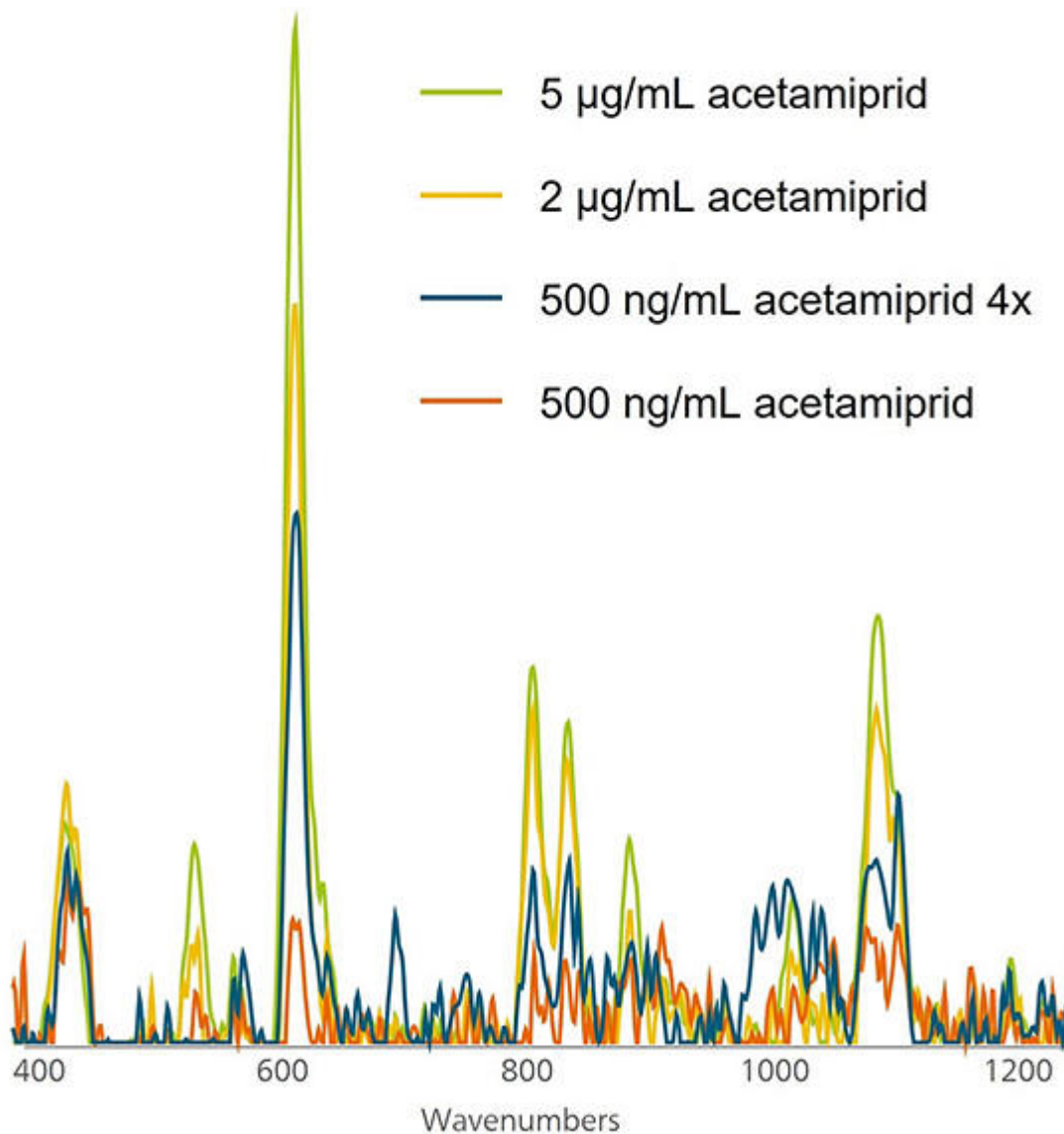


Figure 2. Very low concentration SERS spectra can be improved by combining multiple sample aliquots as seen here with a single 500 ng/mL acetaminophen aliquot (orange) compared to quadruple the 500 ng/mL acetaminophen sample evaporated to the same sample volume for analysis by MISA (blue).

Remarks

In **Figure 2**, the blue spectrum corresponding to four aliquots of 500 ng/mL acetaminophen does not reach the same levels as the yellow 2 μg/mL acetaminophen spectrum. There are several possible influences here:

1. Extraction is not always 100%. A certain amount of pesticide will bind to the sample. For example, if the raisin surface can bind 200 ng of acetamiprid, a maximum of 1.8 g can be extracted from the 2 g/g sample, whereas only 300 ng is left available from each 500 ng/g sample.
2. Extraction is never perfectly reproducible. In one instance 50% extraction may be achieved, and in another iteration, only 40%.
3. The relationship between quantity and intensity for SERS is non-linear, so small differences in analyte amount can result in a large difference in signal intensity, especially at or near detection limits.

Conclusion

MISA is a compact, user-friendly, state-of-the-art analytical tool for ensuring food safety. It facilitates informative decision making when screening food samples suspected of containing pesticide residues. Dedicated SERS substrates and a well-developed library of proven pesticides, herbicides, fungicides, and potentially harmful food additives make MISA a powerful tool for trace-detection applications.

References

1. *Raisin Grades & Standards | Agricultural Marketing Service*. <https://www.ams.usda.gov/grades-standards/raisin-grades-standards> (accessed 2022-07-26).
2. Group, E. W. *EWG's 2022 Shopper's Guide to Pesticides in ProduceTM*. <https://www.ewg.org/foodnews/> (accessed 2022-07-26).
3. Authority (EFSA), E. F. S. National Summary Reports on Pesticide Residue Analysis Performed in 2020. *EFSA Support. Publ.* **2022**, 19 (3), 7216E. <https://doi.org/10.2903/sp.efsa.2022.EN-7216>.

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