

#### Application Note

# MALDI micro MX Sensitivity

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This is an Application Brief and does not contain a detailed Experimental section.

## Abstract

This application brief describes the sensitivity of Waters Micromass MALDI micro MX Mass Spectrometer in reflectron and linear mode, with positive and negative polarity.

## Introduction

Matrix Assisted Laser Desorption Ionization Time-of-Flight Mass Spectrometry (MALDI-Tof MS) is one of the most sensitive mass spectrometric techniques for the analysis of biomolecules. The limit of detection of a MALDI system is determined by two factors. The absolute sensitivity of the hardware and the preparation of the sample. Of these factors the sample preparation is usually the limiting factor.

## Experimental

The sensitivity of the Waters Micromass MALDI micro MX Mass Spectrometer was tested in reflectron and linear mode, with positive and negative polarity.

#### Sensitivity reflectron mode

Sample: Glu-fibrinopeptide B at concentrations of 2 and 20 fmol/µL for positive and negative ion mode respectively made up in 0.1% trifluoroacetic acid (TFA).

Matrix: Purified a-cyano-4-hydroxy cinnamicacid matrix was obtained from Waters (186002331). The matrix was prepared at 2 mg/mLin 1:1 acetonitrile:aqueous 0.1% TFA. Samples were mixed 1:1 with matrix and 1 µL was spotted directly onto the target plate and allowed to air dry prior to analysis.

The final amount of sample per spot was 1 fmolfor positive ion mode and 10 fmolfor negative ion mode.Data acquisition: Six separate acquisitions in positive and negative mode.

#### Sensitivity linear mode

Sample: Glu-fibrinopeptide B at concentrations of 20 and 100 fmol/µL for positive and negative ion mode respectively made up in 0.1% TFA.

Matrix: The same method and matrix were used as in reflectron mode.

Data acquisition: Six separate acquisitions in positive and negative mode.

Final amount of sample per spot was 10 fmol for positive ion mode and 50 fmol for negative ion mode.

#### Data Processing

20 spectra (100 laser shots) were combined and background subtracted (polynomial order 15, subtraction 10 %). The baseline was magnified by factor 5 over the m/z range 1590–1650.

### **Results and Discussion**

The mass spectra obtained are shown in Figures 1a–1d. In all cases a section of the baseline is magnified to be compared to the monoisotopic Glu-fibrinopeptide B peak. In all experiments performed the MALDI micro MX

easily surpassed the instrument sensitivity specifications of a Signal:Noise ratio of 5:1 (Table 1).

Mode	Polarity	Average S/N ratio (six acquisitions)
Linear	negative	39:1
Linear	positive	41:1
Reflectron	negative	26:1
Reflectron	positive	16:1

Table 1. Summary of the sensitivity experiments, showing Tof analyzer mode, polarity mode and average S/N ratio.

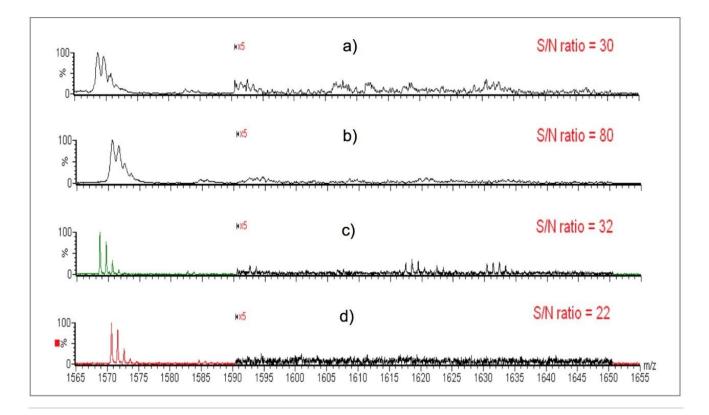


Figure 1. a) 50 fmol Glu-fibrinopeptide B on target measured in linear mode (negative), b) 10 fmol Glufibrinopeptide B on target measured in linear mode (positive), c) 10 fmol Glu-fibrinopeptide B on target measured in reflectron mode (negative), d) 1 fmol Glu-fibrinopeptide B on target measured in reflectron mode (positive).

## Conclusion

- These data easily surpass the sensitivity specifications of the instrument in both linear and reflectron mode, and in positive and negative mode
- The average S/N ratio in reflectron positive mode was 16:1 (specification is 5:1)
- The average S/N ratio in reflectron negative mode was 26:1 (specification is 5:1)
- The average S/N ratio in linear positive mode was 41:1 (specification is 5:1)
- The average S/N ratio in linear positive mode was 39:1 (specification is 5:1)

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