



# Coupling FAIMS and LESA: Improvements in protein analysis directly from biological substrates

Andrew Creese

School of Biosciences,  
University of Birmingham.

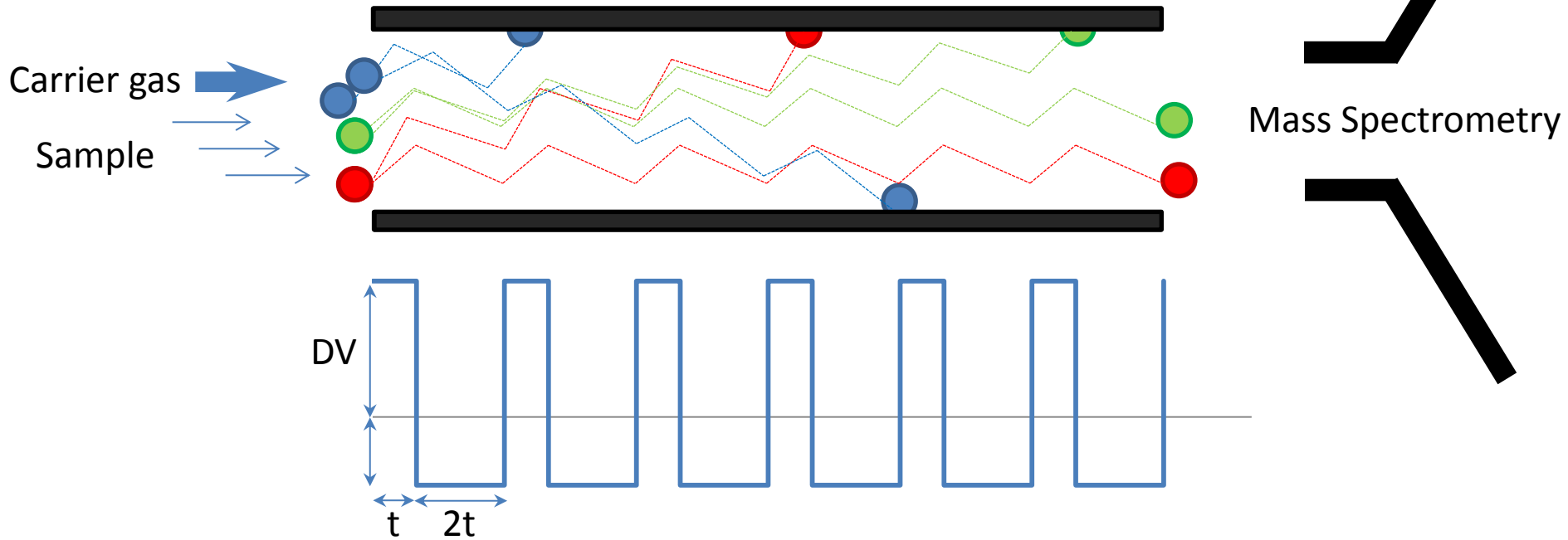
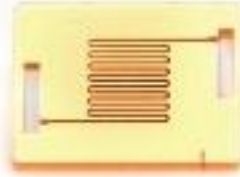
ASMS  
2015



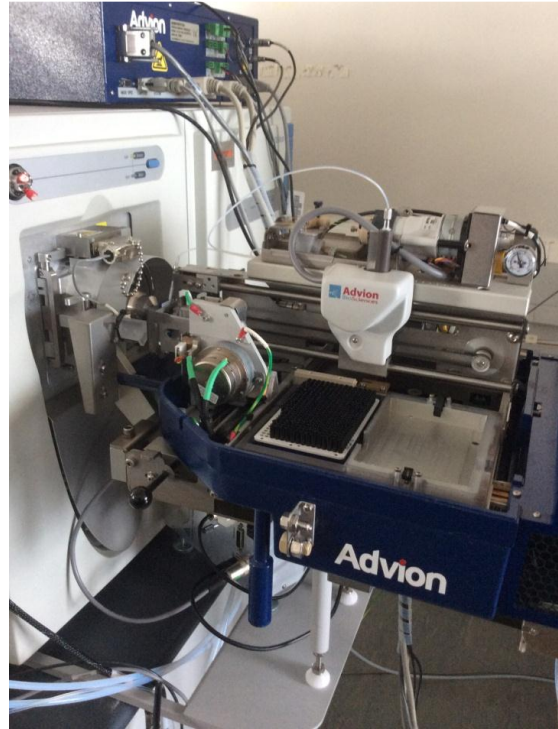
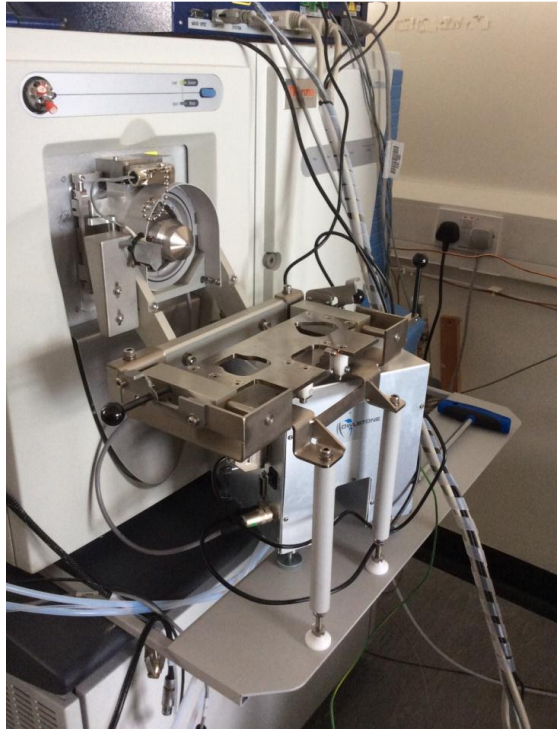
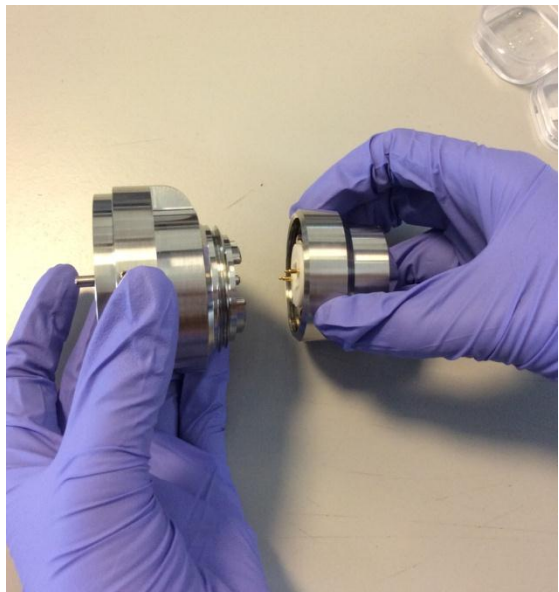
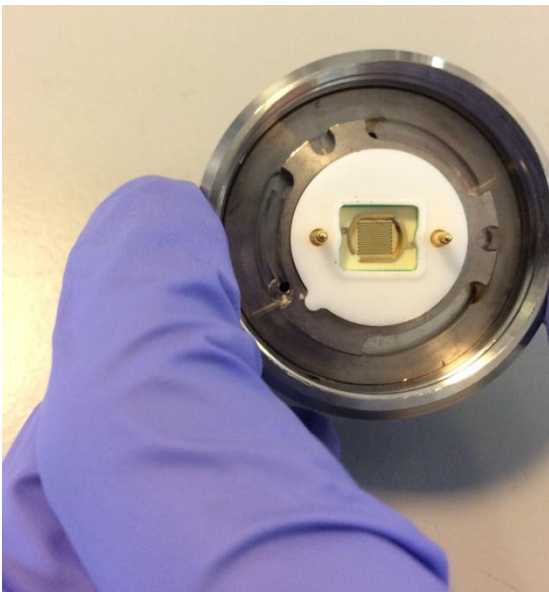
UNIVERSITY OF  
BIRMINGHAM

# FAIMS: High field asymmetric waveform ion mobility spectrometry

Gap width = 100  $\mu\text{m}$   
Thickness = 700  $\mu\text{m}$



# Coupling ultraFAIMS with TriVersa Nanomate



# Mode of operation

**Static**

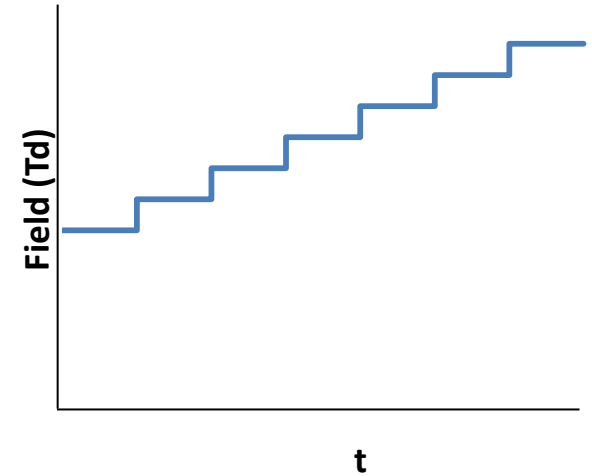
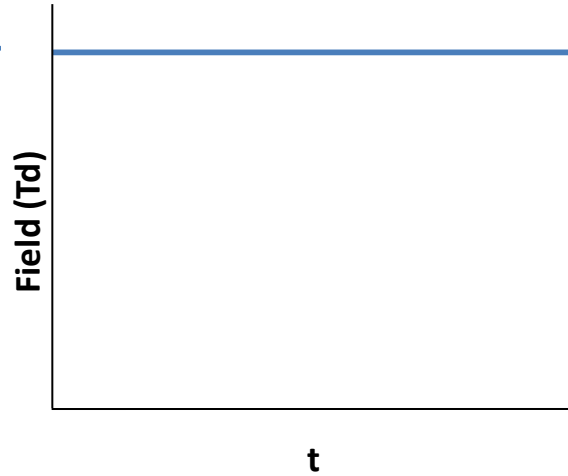
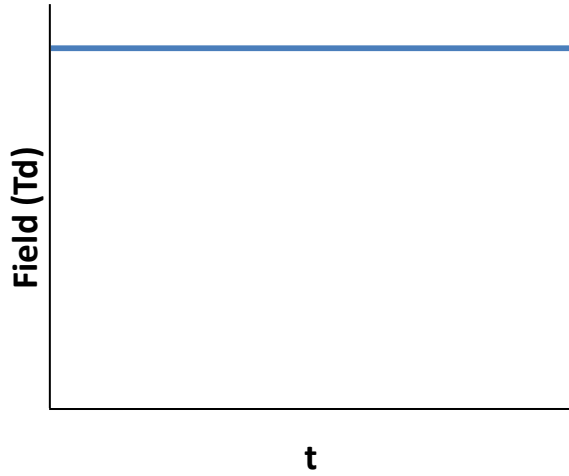
**1D**

**2D**

Dispersion Field

Dispersion Field

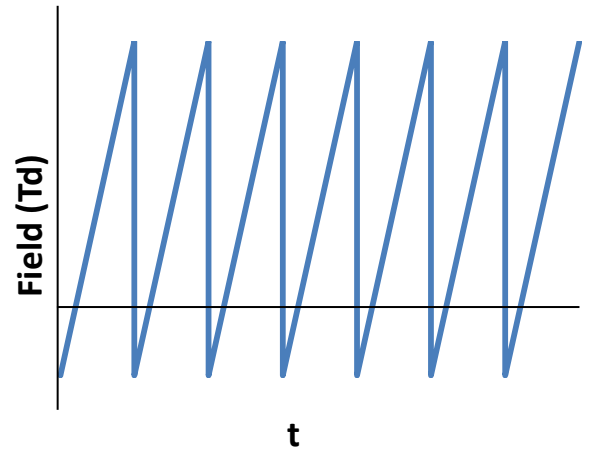
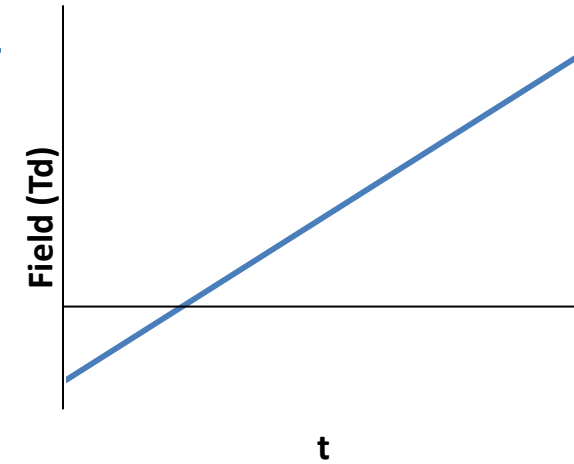
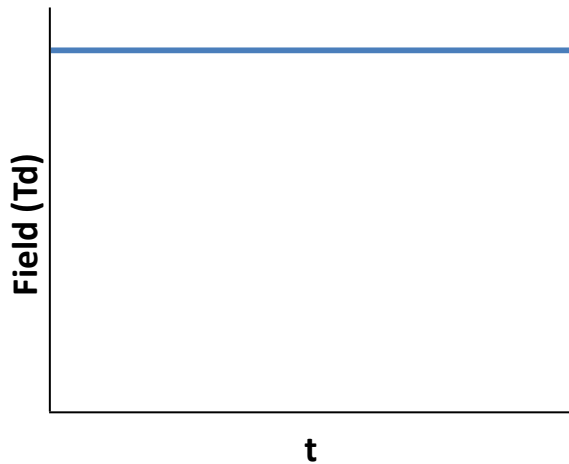
Dispersion Field

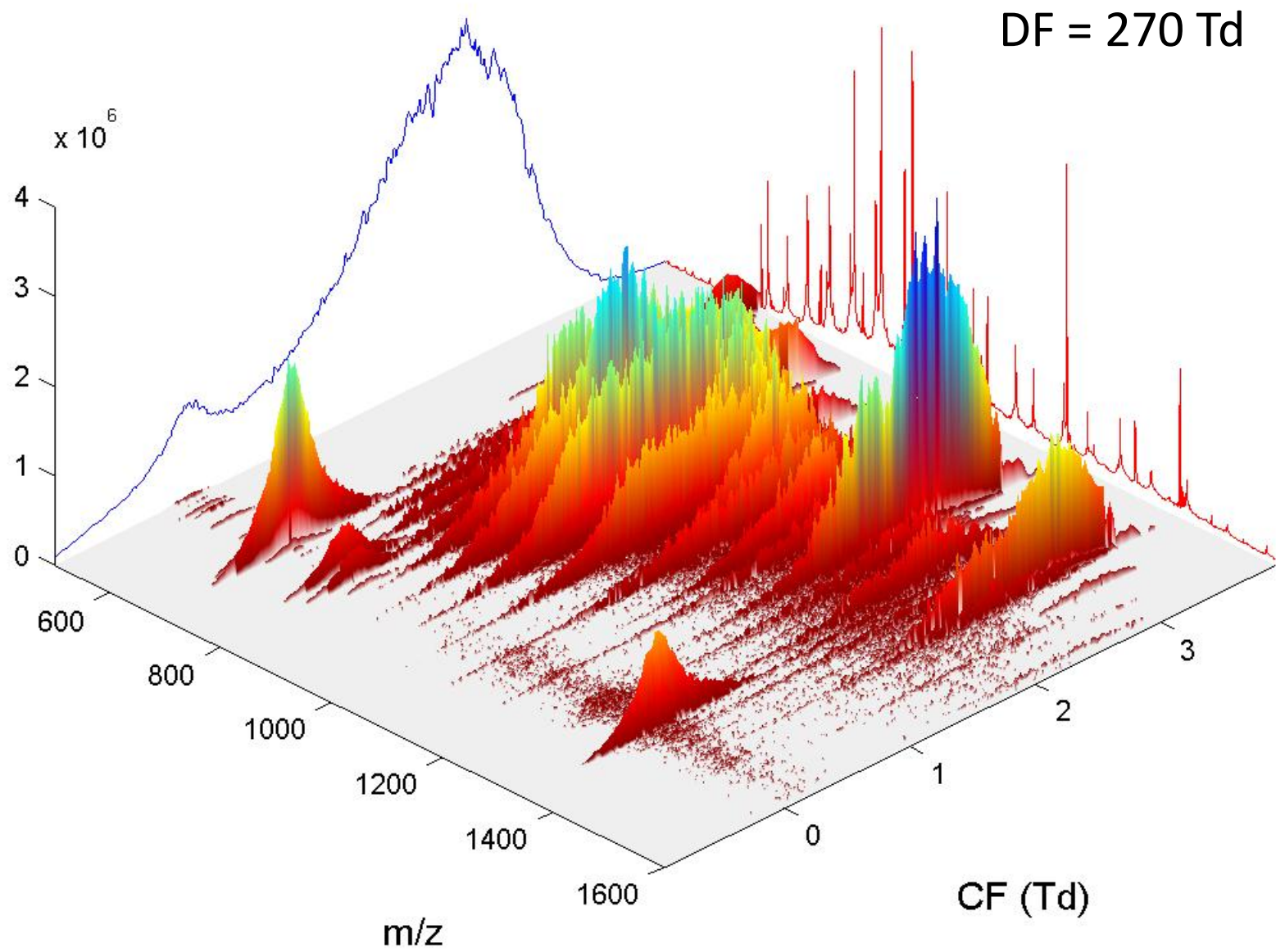


Compensation Field

Compensation Field

Compensation Field



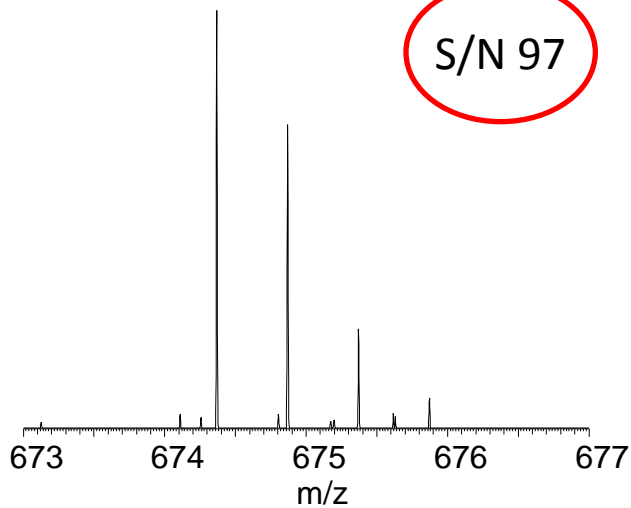


# Substance P [M+2H]<sup>2+</sup>

No FAIMS

1.27E7

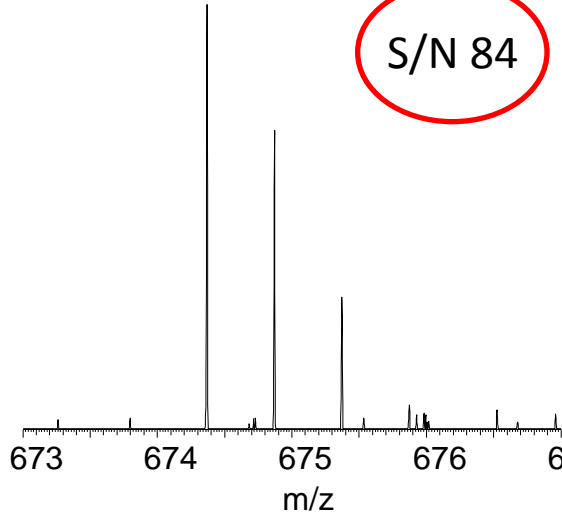
S/N 97



FAIMS no Field

8.26E5

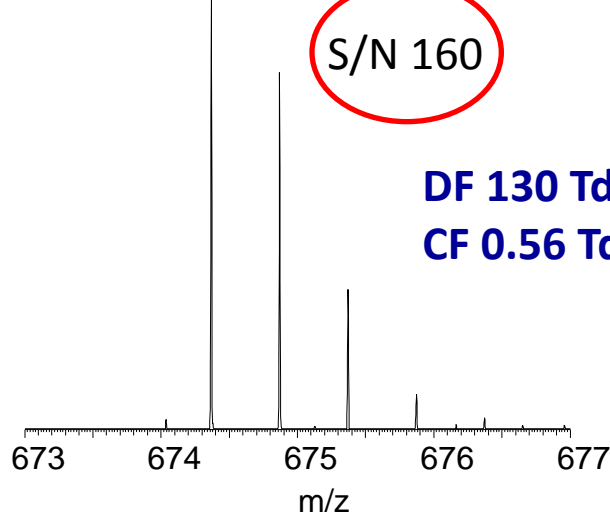
S/N 84



FAIMS

6.74E5

S/N 160



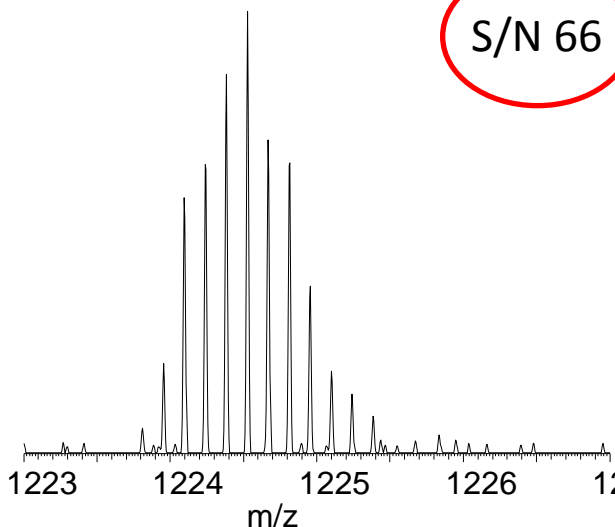
DF 130 Td  
CF 0.56 Td

# Ubiquitin[M+7H]<sup>7+</sup>

No FAIMS

1.30E7

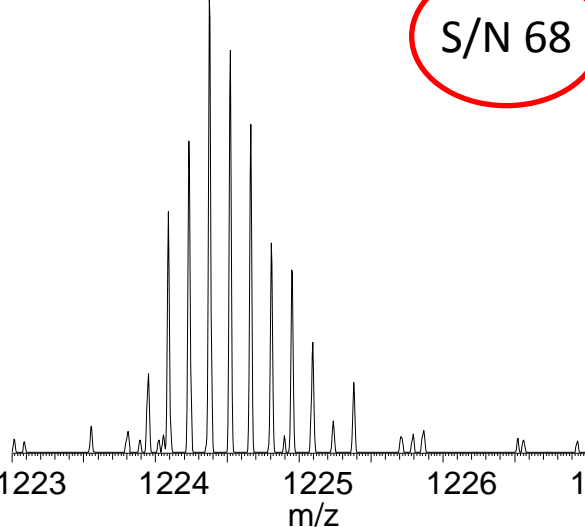
S/N 66



FAIMS no Field

1.15E6

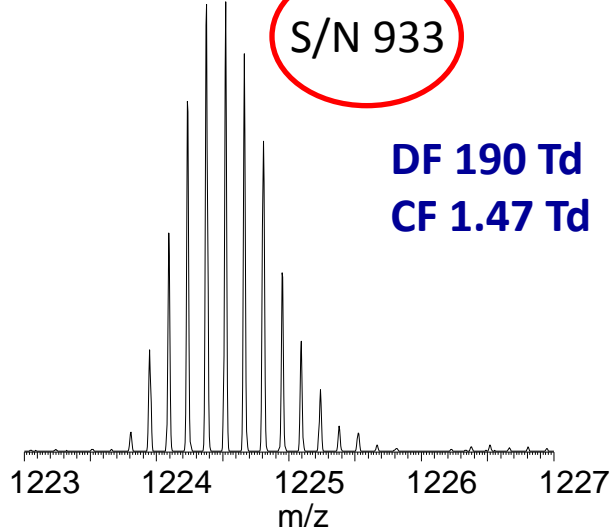
S/N 68



FAIMS

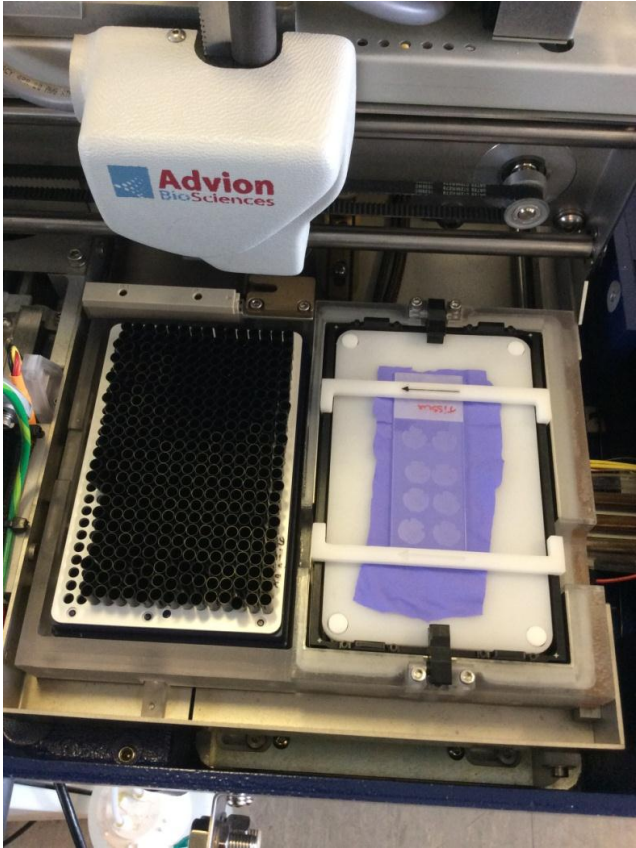
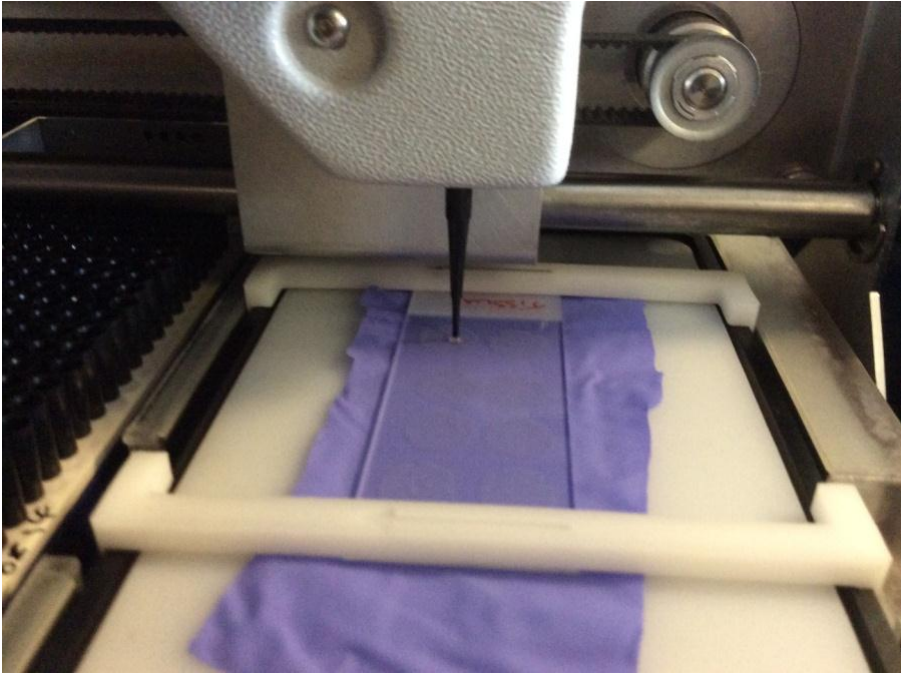
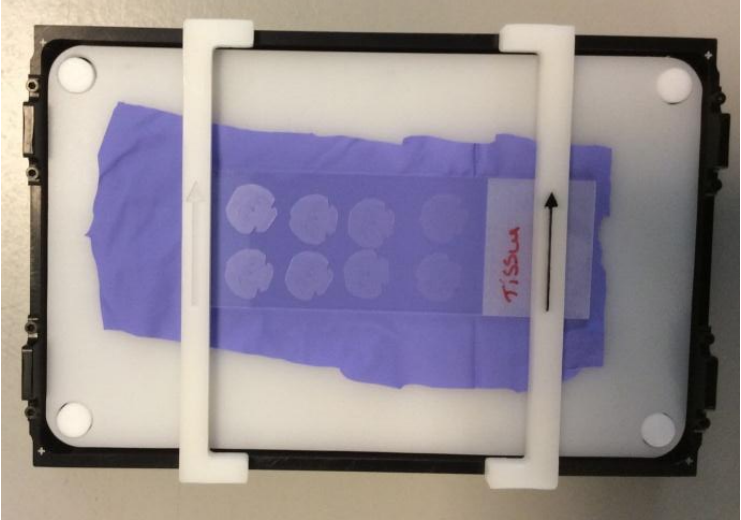
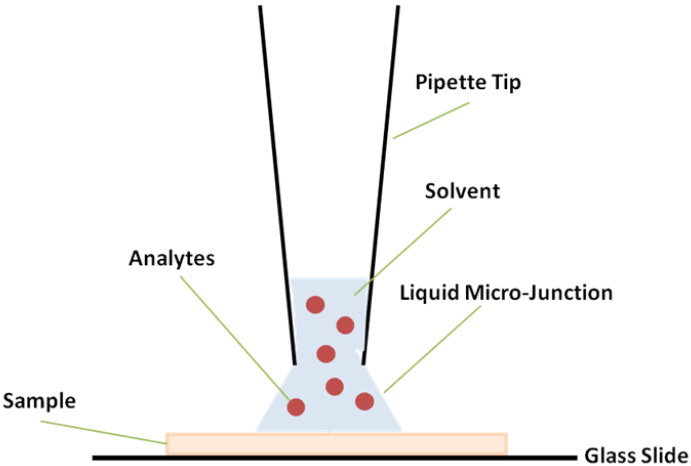
8.88E5

S/N 933



DF 190 Td  
CF 1.47 Td

# LESA: Liquid extraction surface analysis

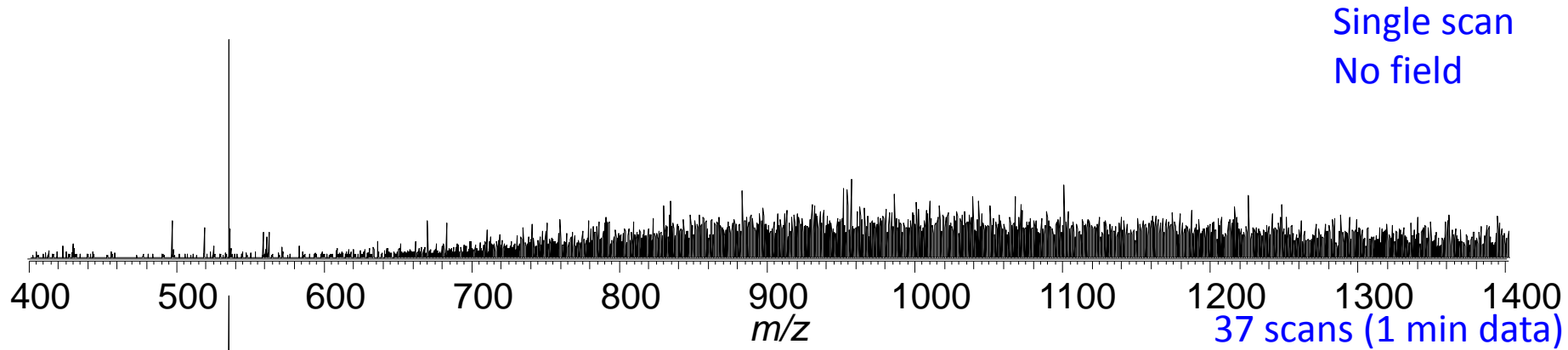


# Workflow

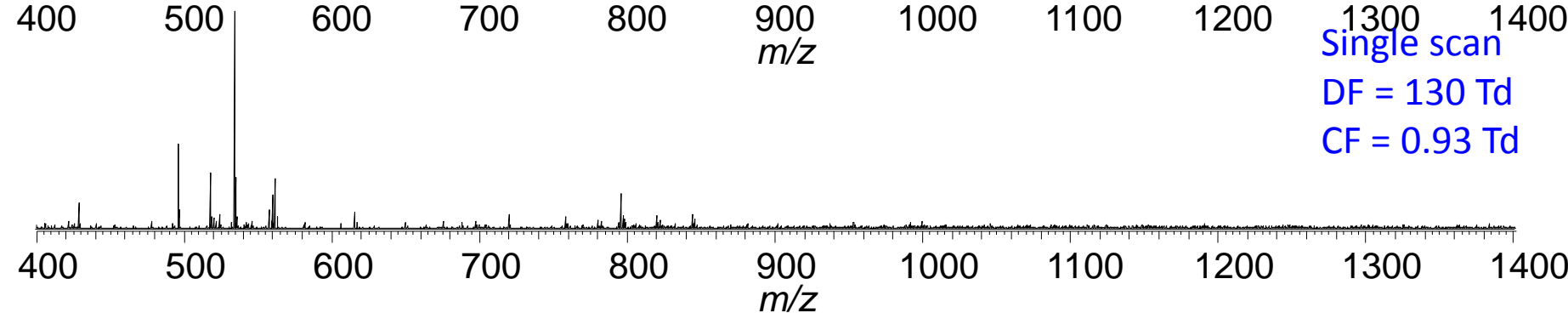
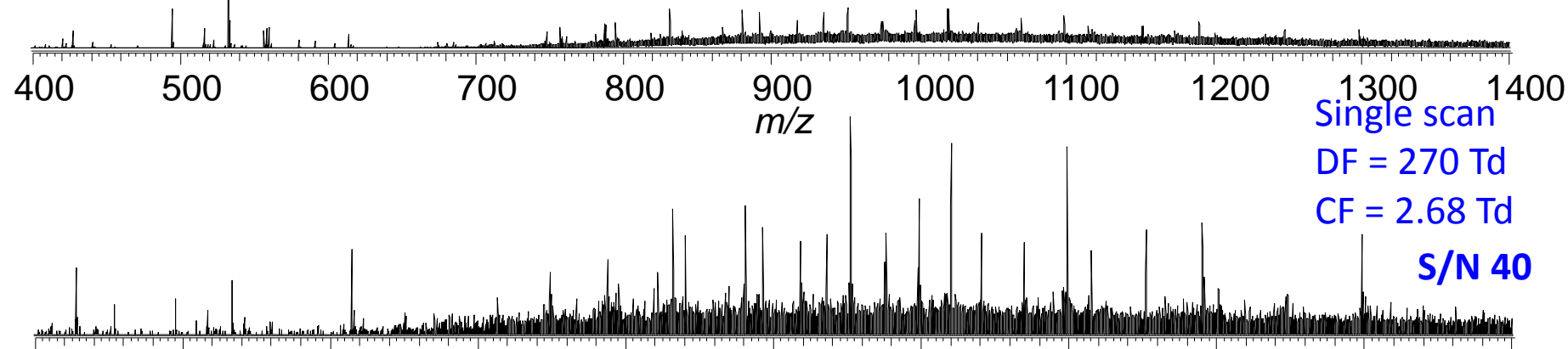




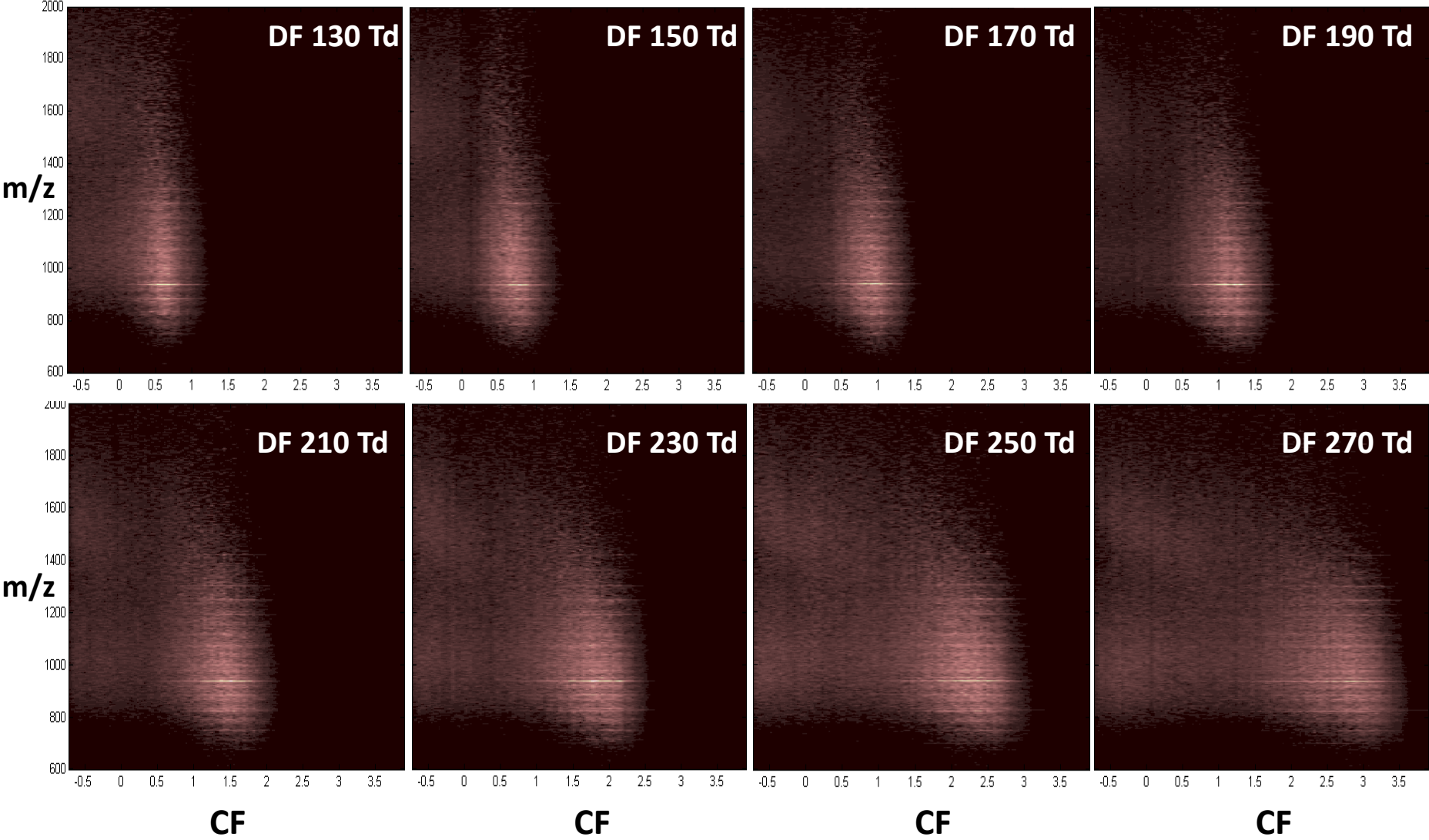
# LESA FAIMS MS of mouse liver



37 scans (1 min data)  
No field  
**S/N 11**

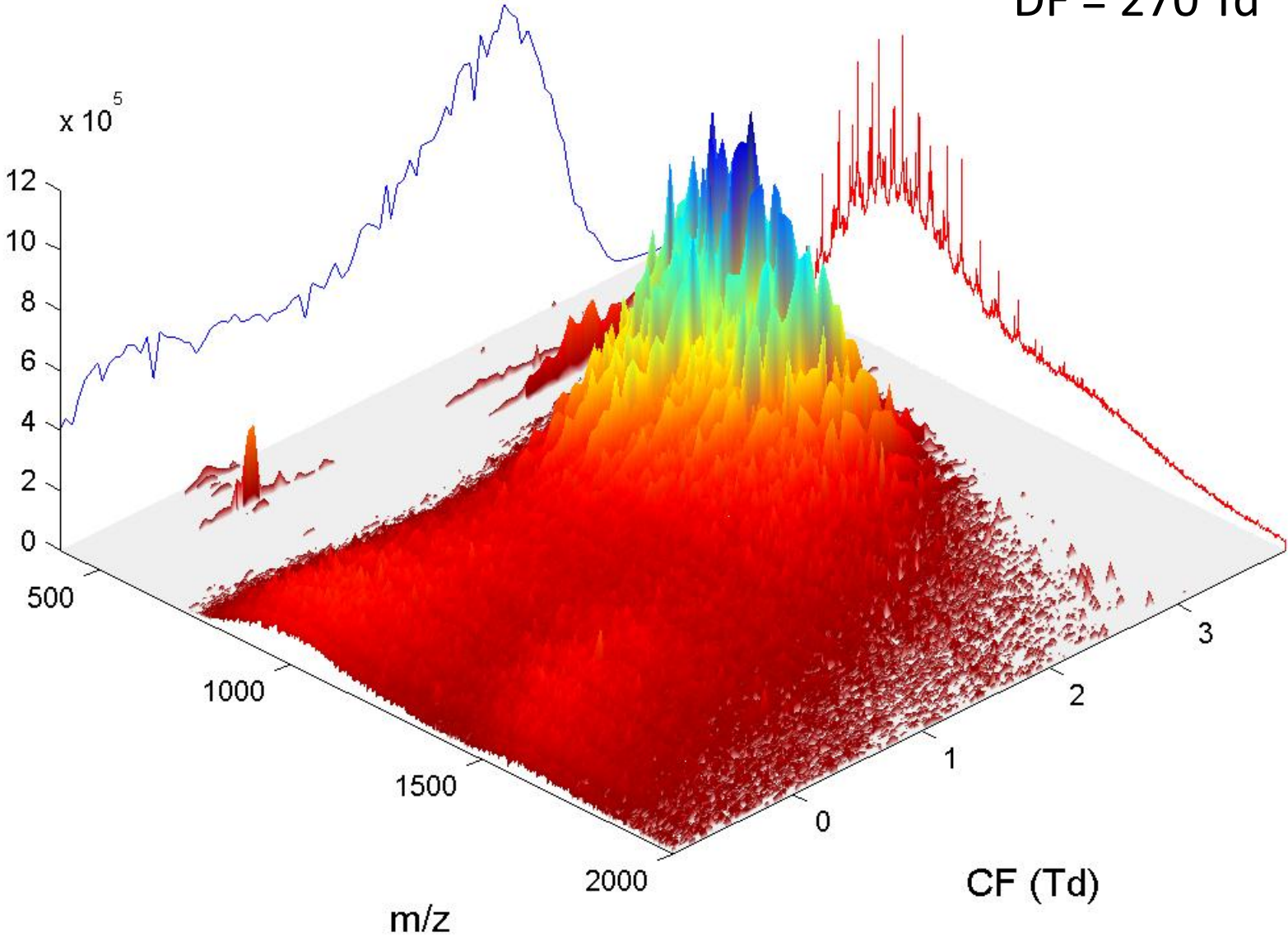


# Visualisation: Total ion transmission maps – mouse liver

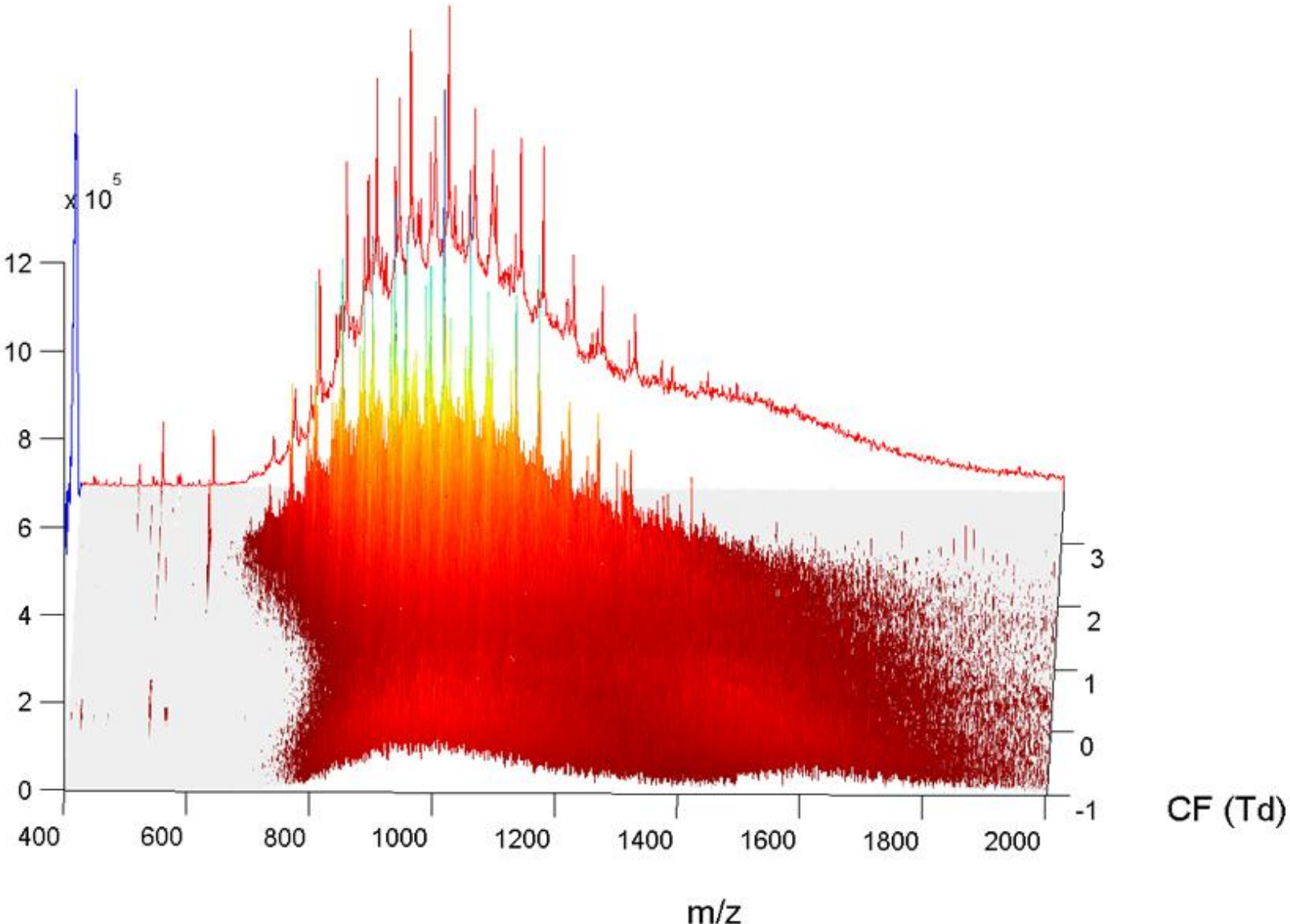


# Visualisation: Total ion transmission maps – mouse liver

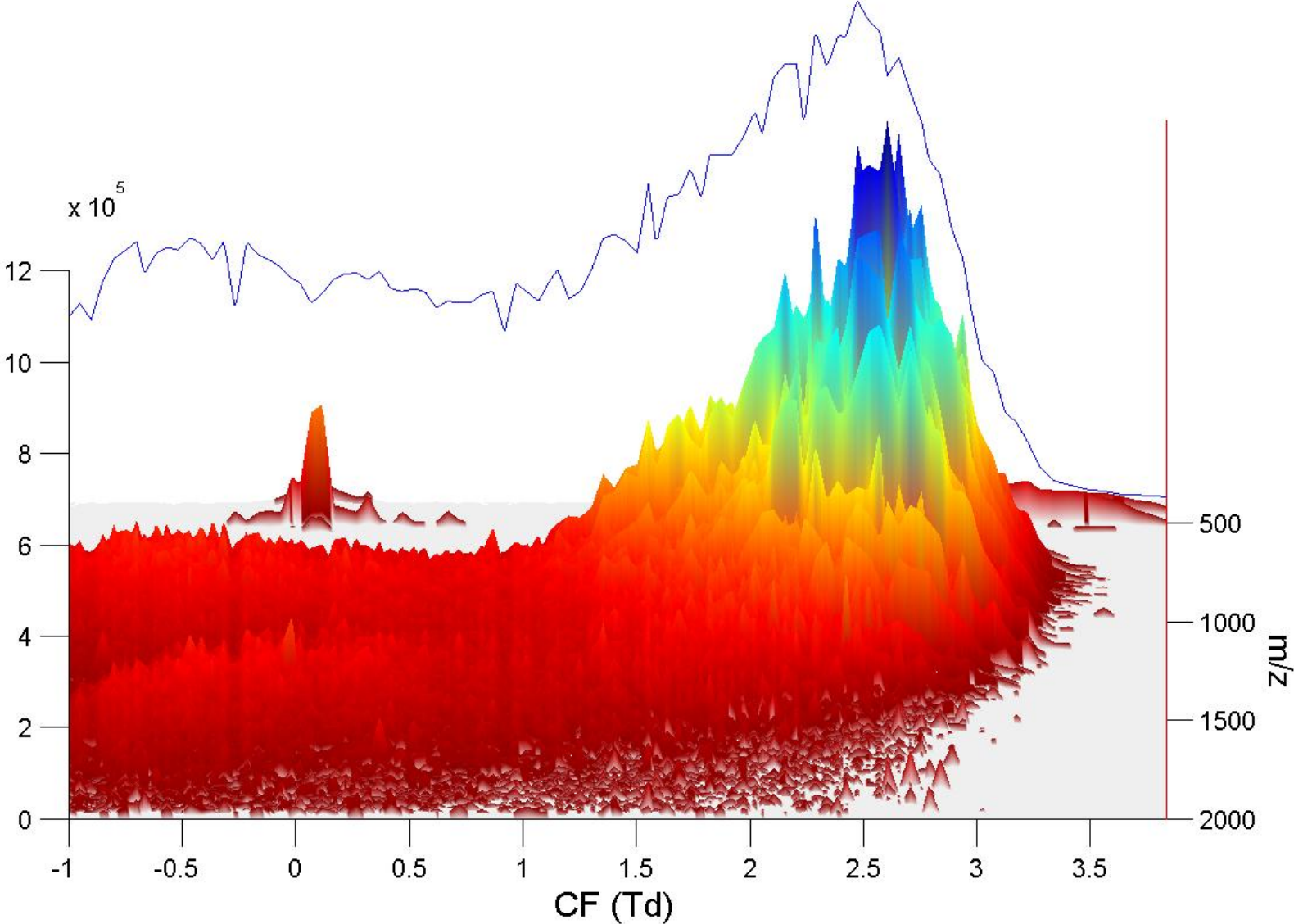
DF = 270 Td



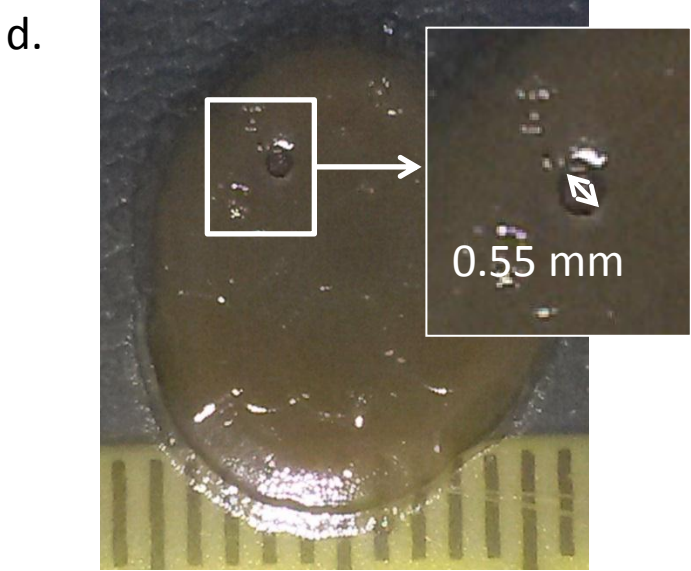
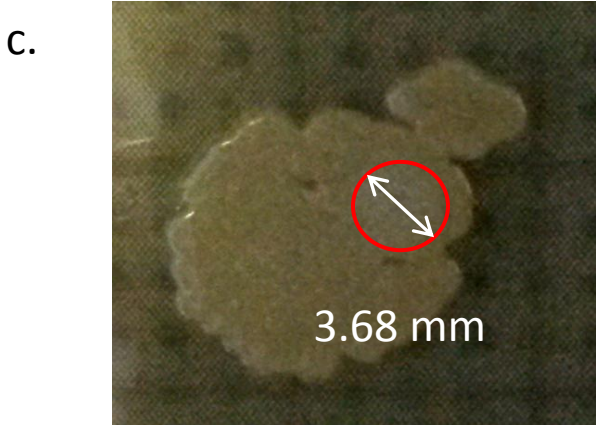
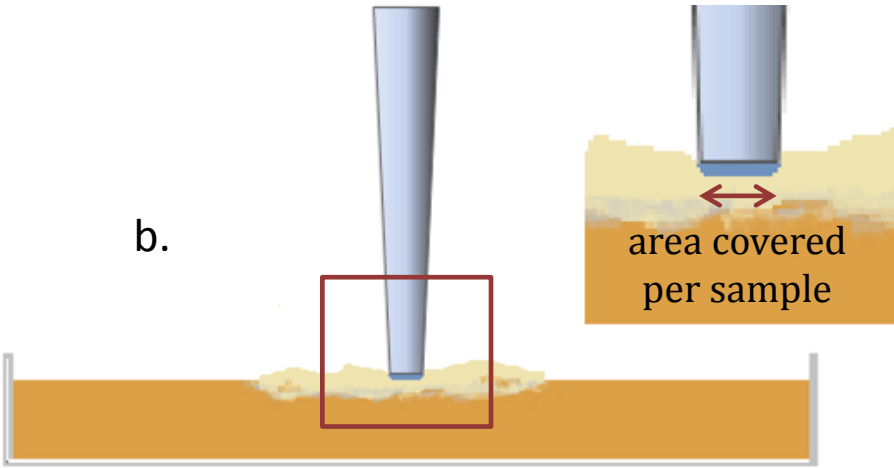
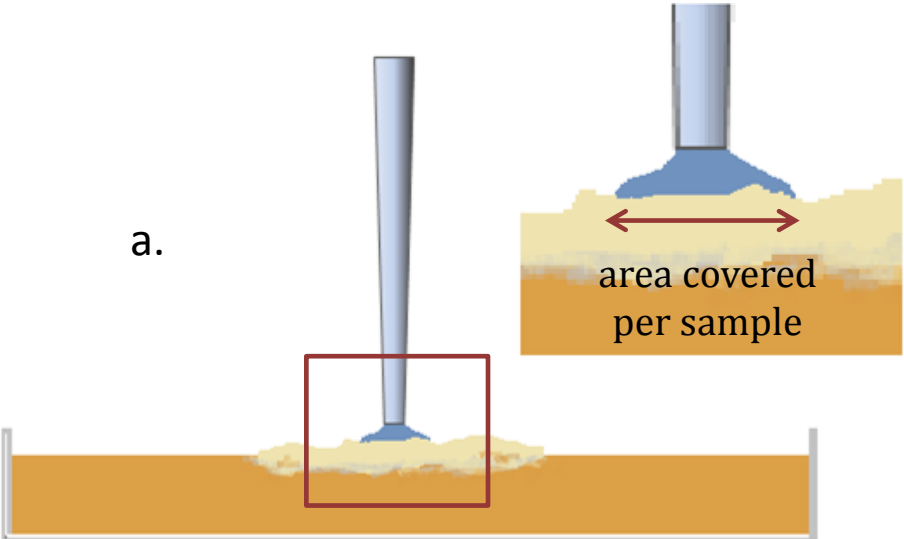
# Visualisation: Total ion transmission maps – mouse liver



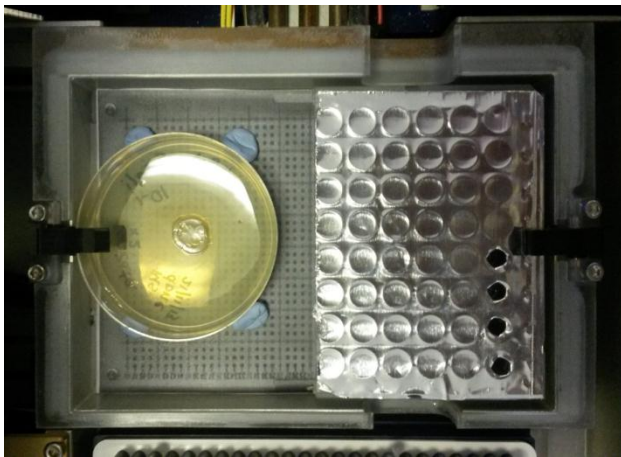
# Visualisation: Total ion transmission maps – mouse liver



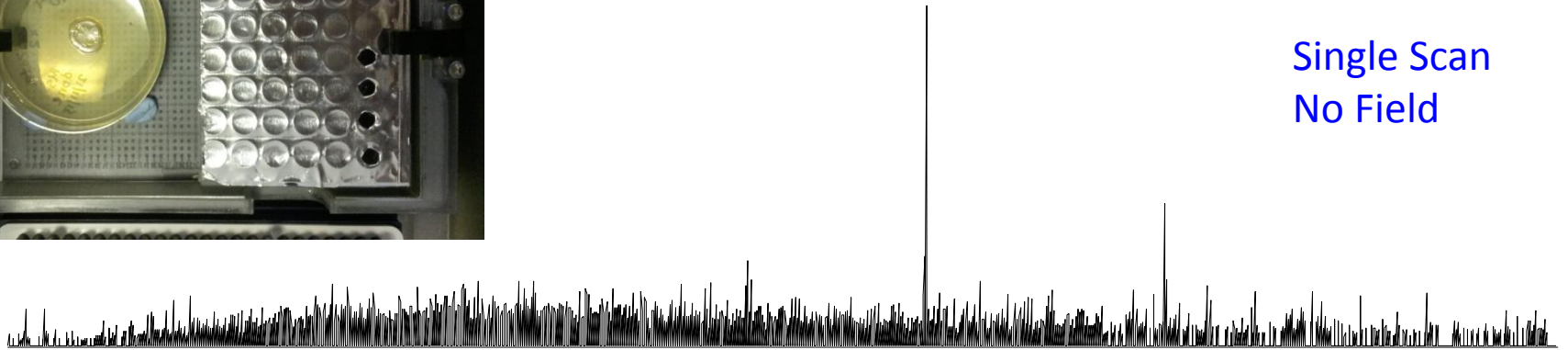
# Contact LESA: Liquid extraction surface analysis



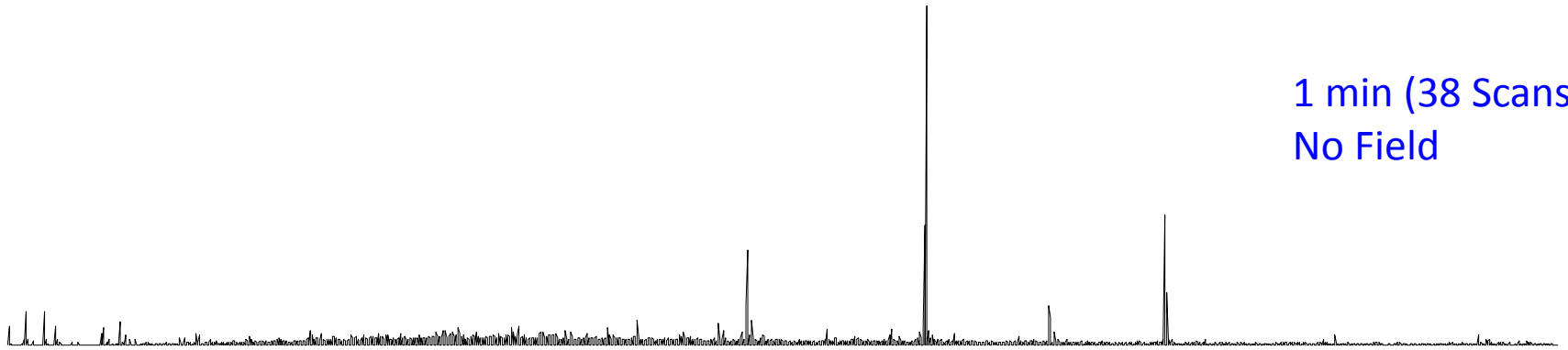
# LESA FAIMS MS of *E. coli*



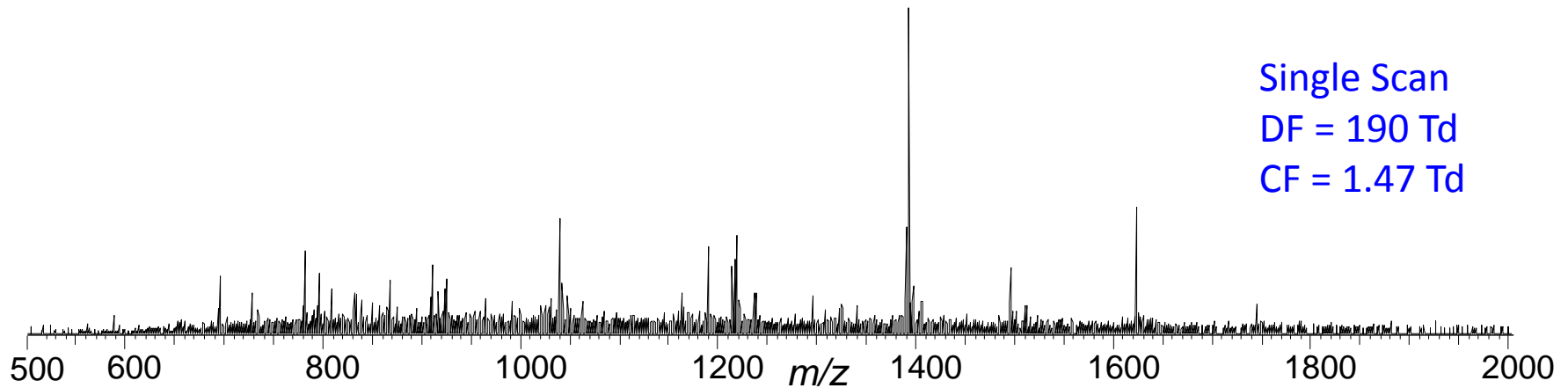
Single Scan  
No Field



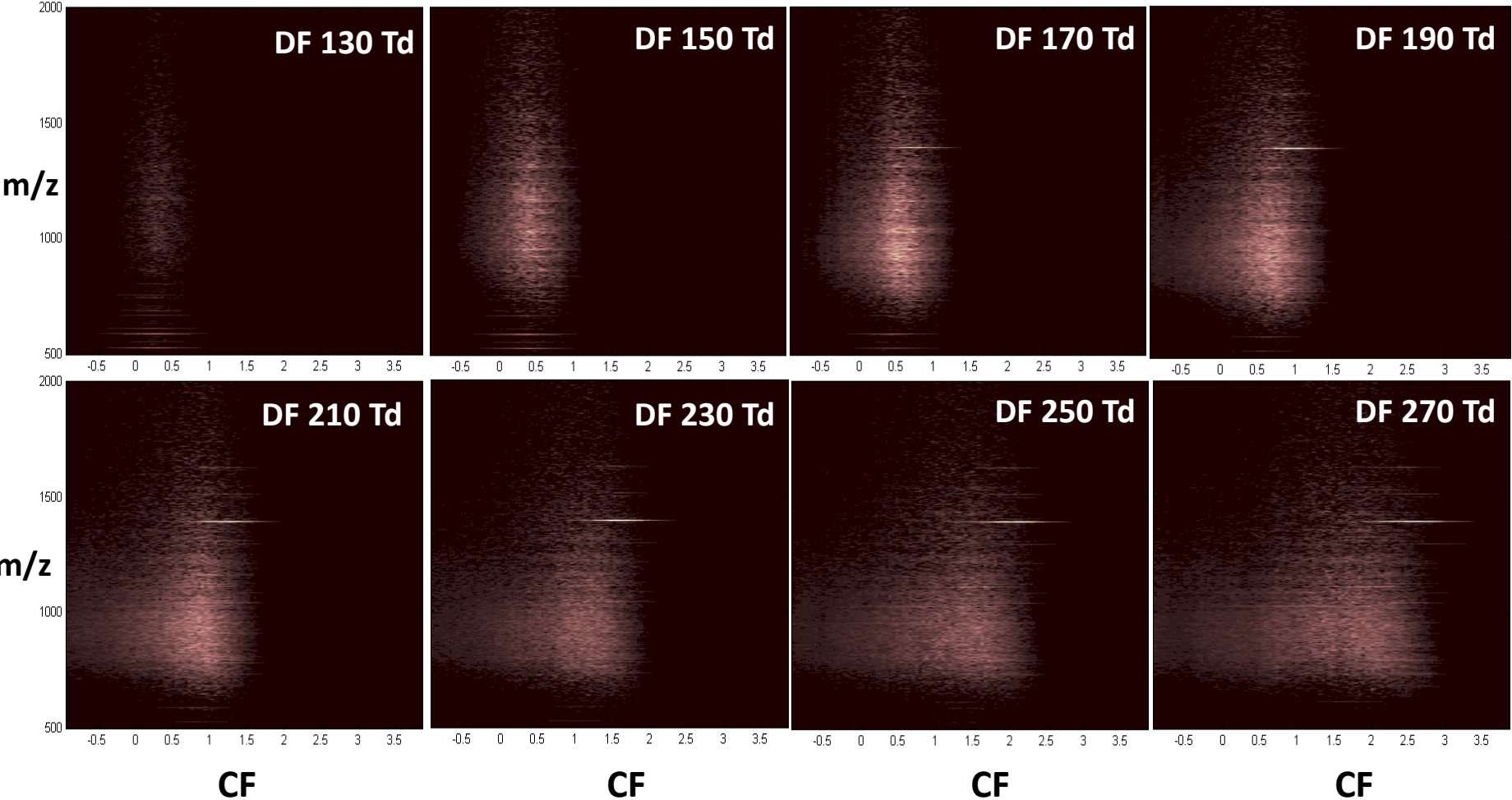
1 min (38 Scans)  
No Field



Single Scan  
DF = 190 Td  
CF = 1.47 Td



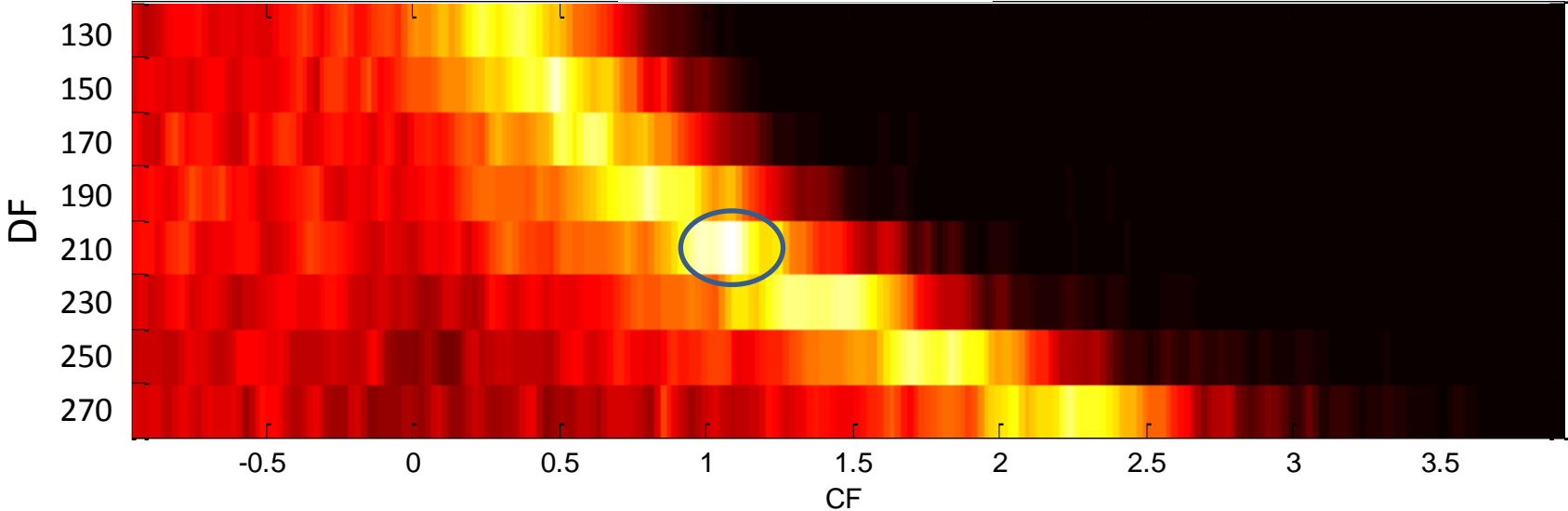
# Visualisation: Total ion transmission maps – *E. coli*



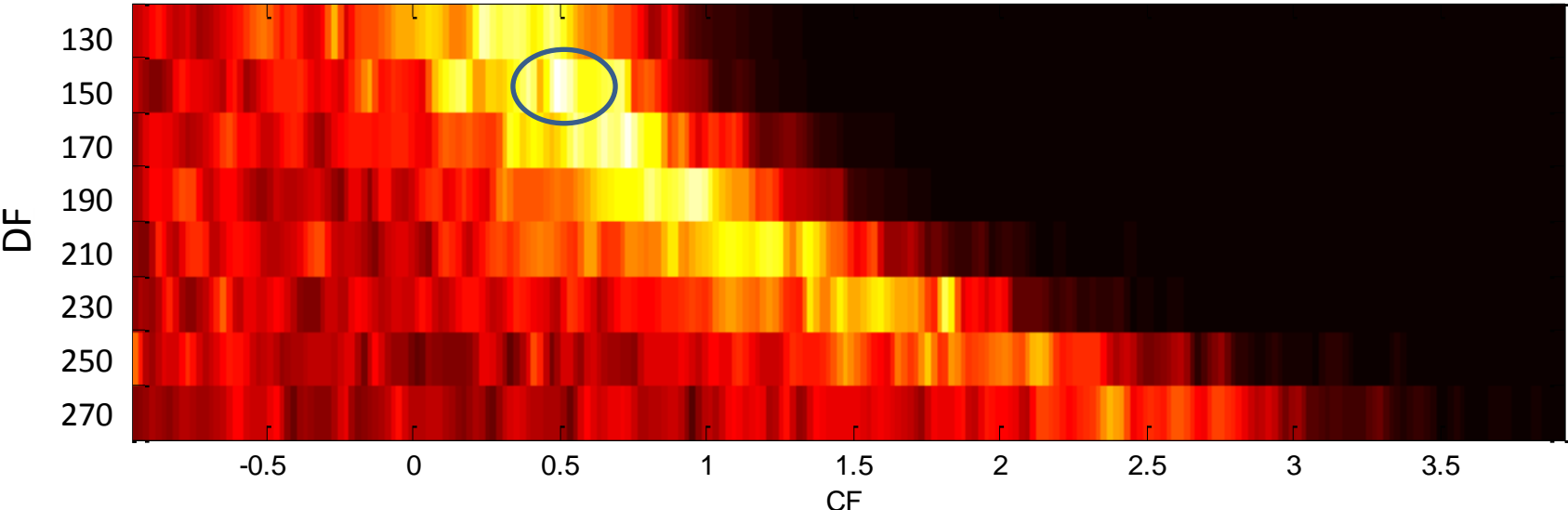


# Visualisation: Single ion transmission maps, LESA-FAIMS mouse liver.

## Beta globin 15+

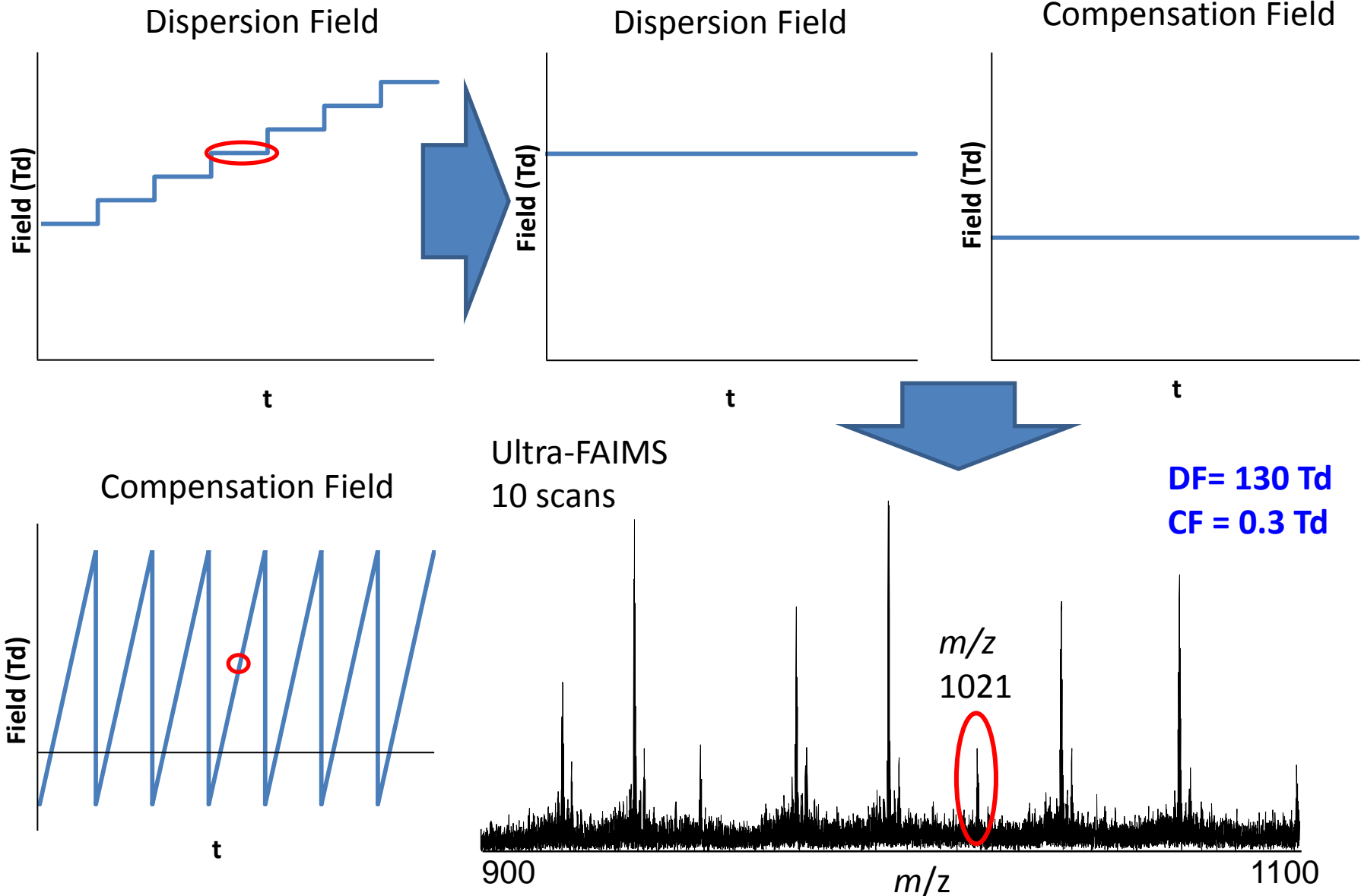


## phosphatidylcholine PC(34:2)



# Static Analysis

2D

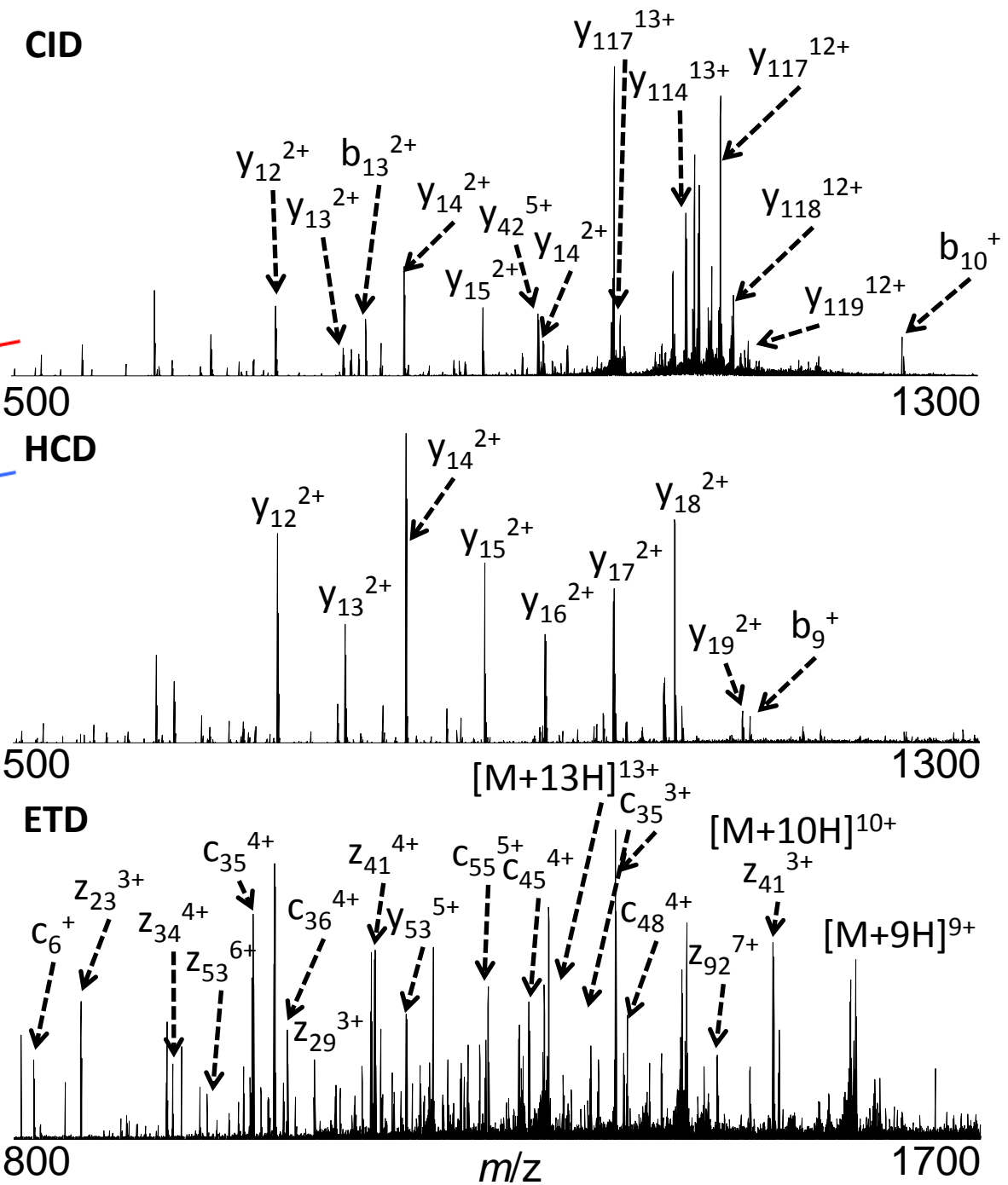


# Liver unknown protein ion m/z 1021 (14+)

MNESGKYQLQ SQENFEPFMK  
 AIGLPEDLIQ KGKDIKGVSE  
 IVHEGKKIKL TITYGPKVVR  
 NEFTLGEECE LETMTGEKVK  
 AVVKLEGDNK MVTTFKGIKS  
 VTEIINGDTIT NTMILGDIVY  
 KRVSKRI

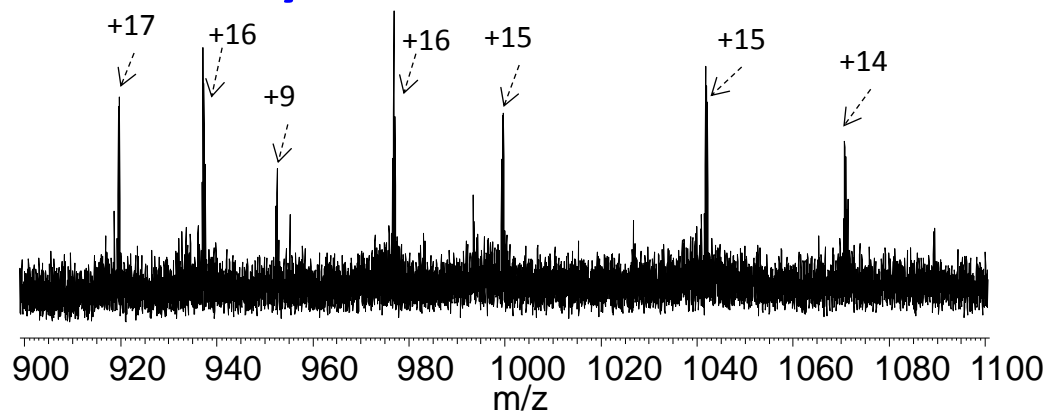
Sequence coverage:  
 CID 19%  
 HCD 15%  
 ETD 25%  
**Total: 42%**

**Liver fatty acid  
binding protein**



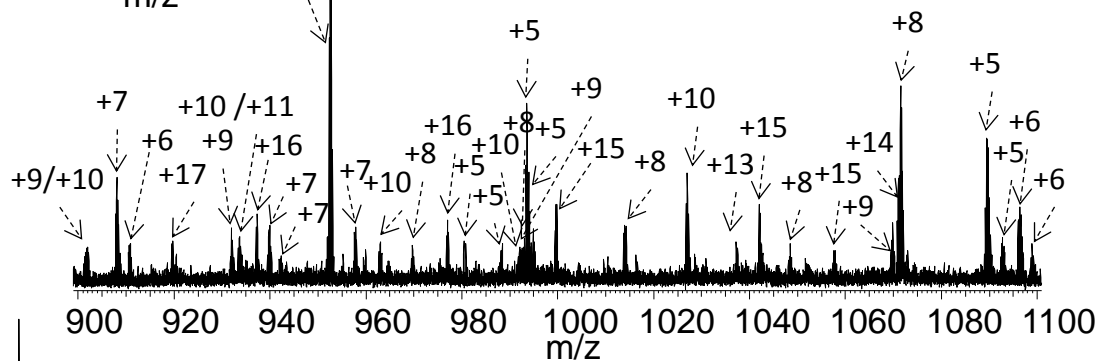
# LESA FAIMS of mouse brain: Static analysis

1 min data (37 scans)  
No Field



500 600 700 800 900 1000 1100 1200 1300 1400 1500 1600 1700 1800 1900 2000  
m/z

1 min data (23 scans)  
DF = 270 Td  
CF = 2.6 Td



500 600 700 800 900 1000 1100 1200 1300 1400 1500 1600 1700 1800 1900 2000  
m/z

# Conclusions

- Interfaced Advion Triversa Nanomate with Owlstone uFAIMS – LESA-FAIMS-MS
- Separation of different molecular species
- Reduced analysis time
- More information per spectrum
- Greater sequence coverage from MS/MS events

# Acknowledgments

## UoB

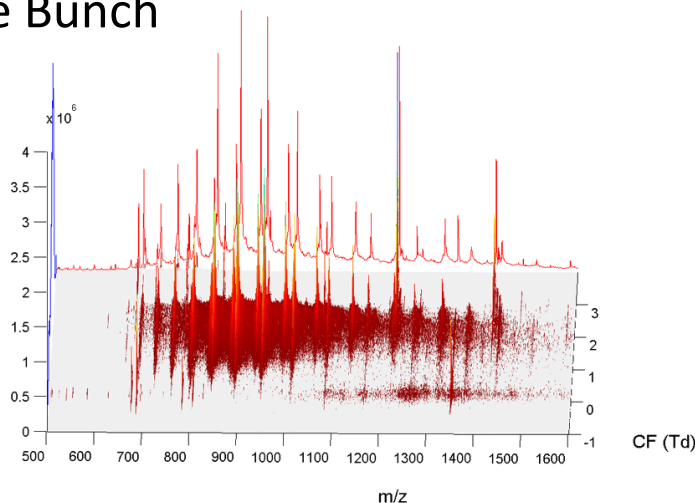
Joscelyn Sarsby  
Buffy Randall  
Alex Dexter  
Dr Rian Griffiths  
Dr Andrew Creese  
Prof Helen Cooper

## Owlstone:

Dr Danielle Toutoungi  
Dr Lauren Brown

## NPL:

Dr Josephine Bunch  
Alan Race



## Funding:

EPSRC  
Advantage West Midlands  
Birmingham Science City