

DNA analysis by MALDI-TOF mass spectrometry

Matrix

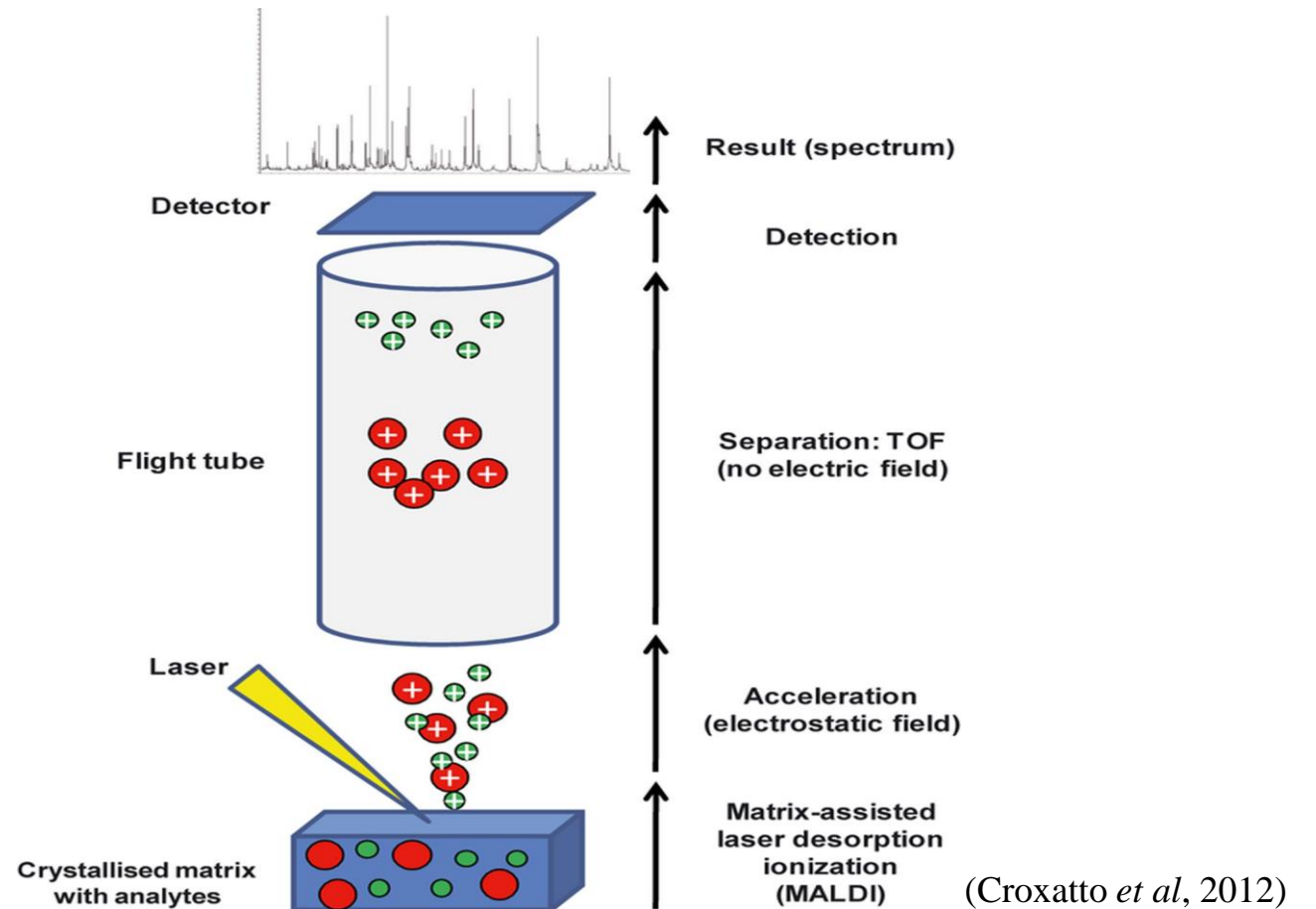
Assisted

Laser

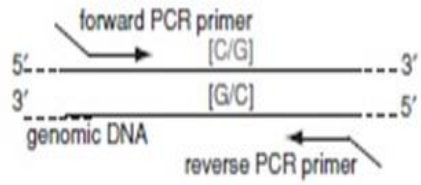
Desorption

Ionization

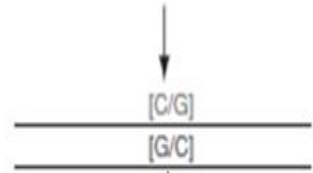
Time **O**f **F**light



amplification

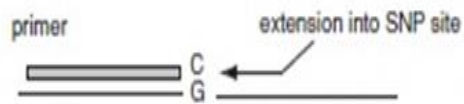


PCR product



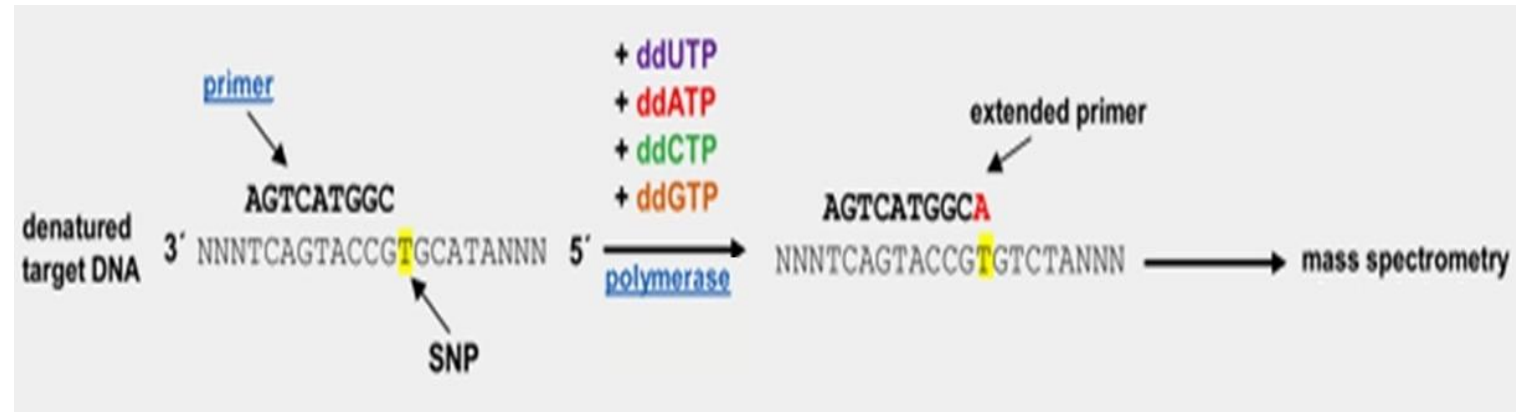
Purification reaction

Primer extension

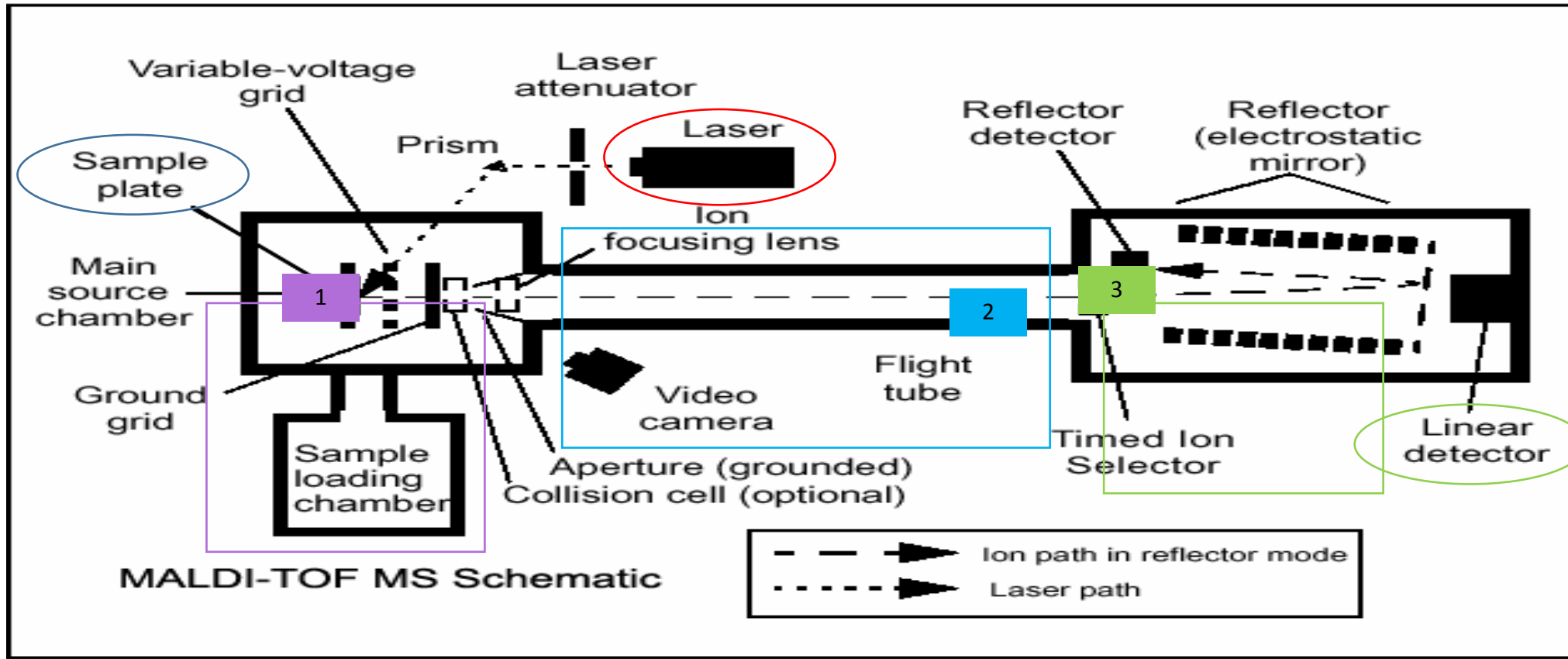


sample conditioning, dispensing, and MALDI-TOF MS

MALDI-TOF mass spectrometry analysis



(Modified from Gabriel *et al*, 2009)



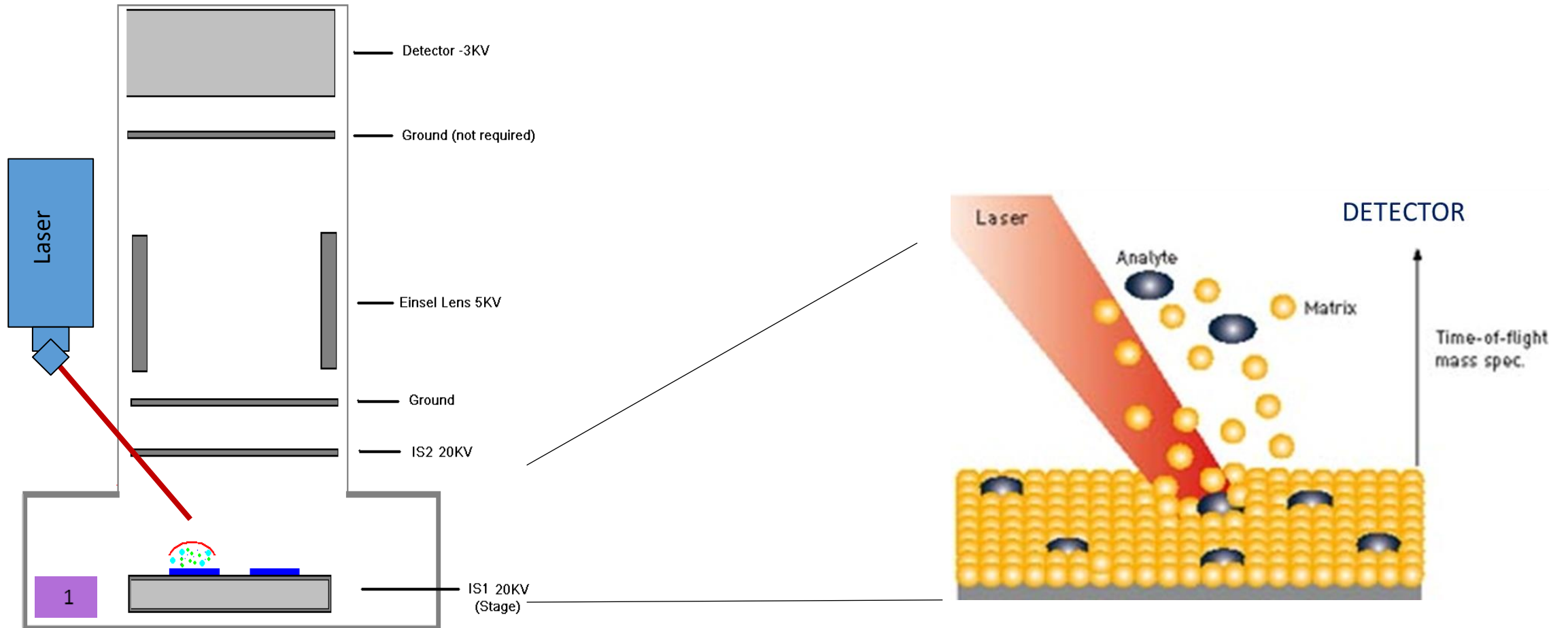
1 IONIZATION CHAMBER

2 ANALYZER

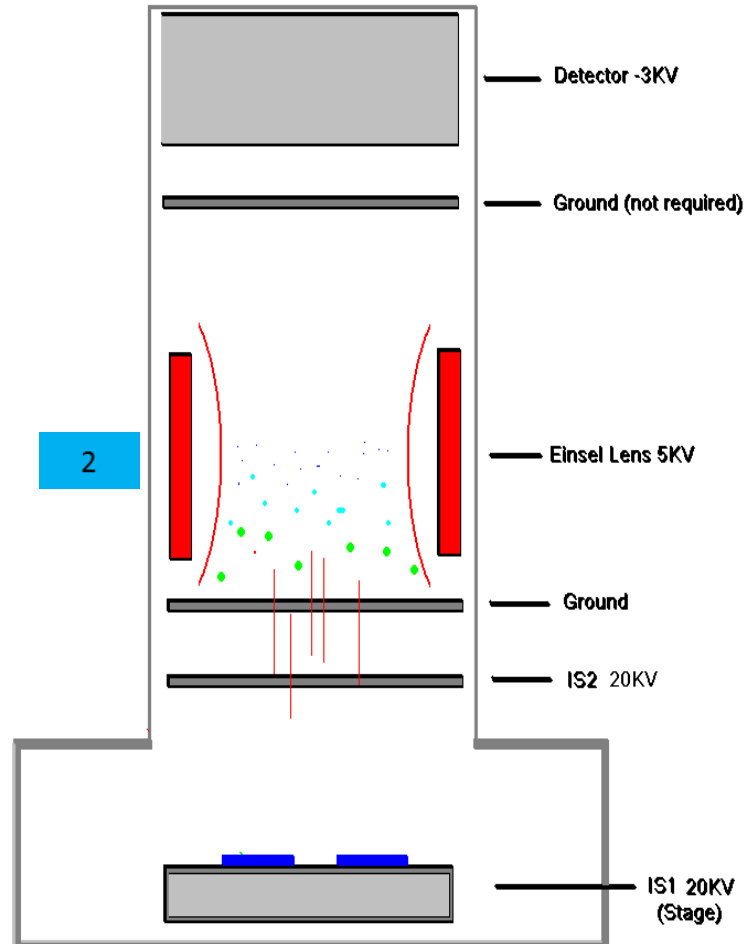
3 DETECTOR

IONIZATION CHAMBER:

Ions are generated by a **desorption/ionization process matrix** assisted by a laser (MALDI)



ANALYZER: Based on time of flight (ToF)

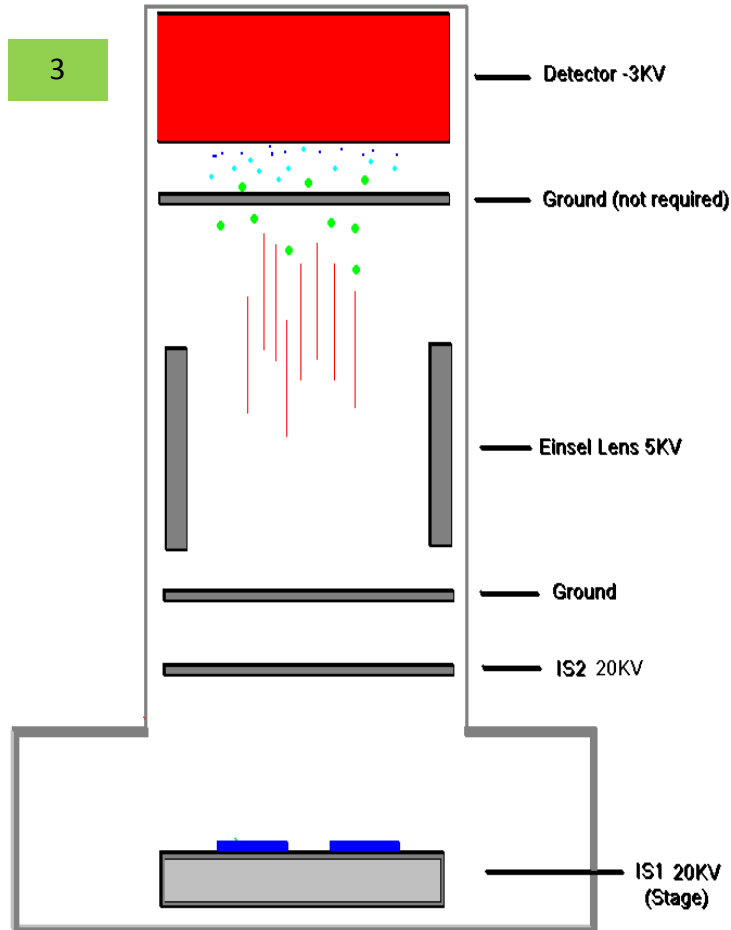


Time of flight of a ion is related to its mass/charge ration m/z

Lighter ions hit the detector first, while the slower traveling heavier ions hit the detector later

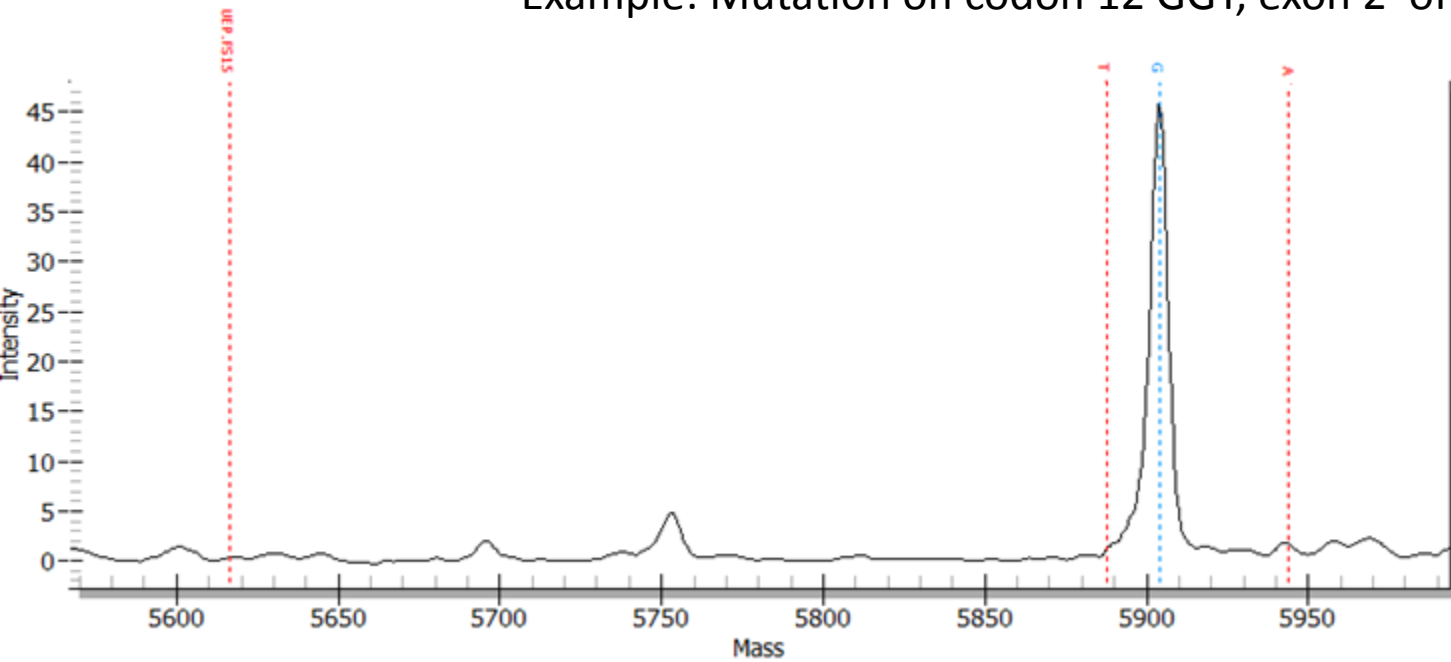
DETECTOR:

Ions are multiplied in electrons, further used to obtain an electric signal

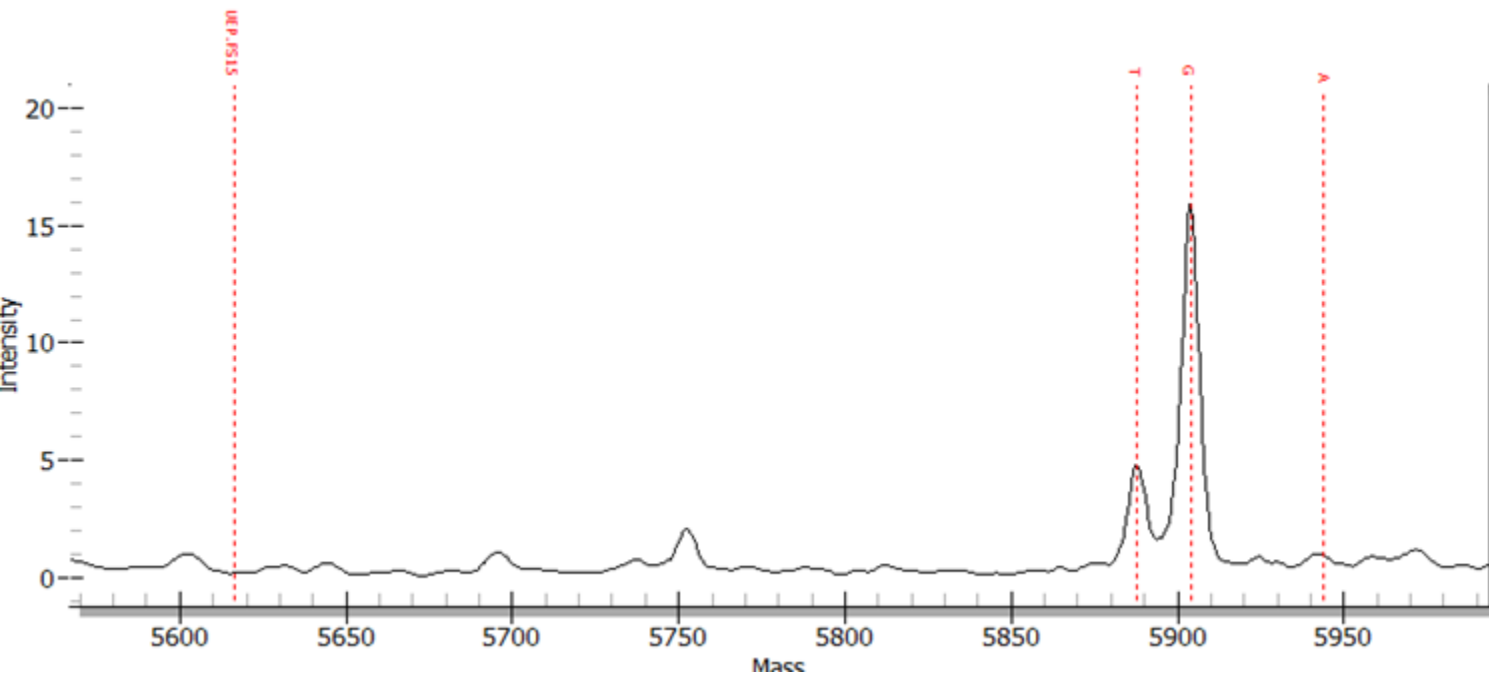


1. Each ion reaches the detector and 1-3 electrons will be released
2. Electrons move into another plate and 1-3 new electrons will be released (secondary emission)
1. This process will be repeated many times, leading to a signal amplification → digital signal visible in real time on the computer.

Example: Mutation on codon 12 GGT, exon 2 of *KRAS* gene



Mass spectrometer analysis graph, with an expected peak (G), as in wild-type sequence **GGT**.



Mass spectrometer analysis graph, with an expected peak (G) and an unexpected peak (T) representing c.34G>T transversion mutation **GGT>TGT**