

# Sequence Queries

MASCOT

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## Three ways to use mass spectrometry data for protein identification

### 1. Peptide Mass Fingerprint

A set of peptide molecular masses from an enzyme digest of a protein

### 2. Sequence Query

Mass values combined with amino acid sequence or composition data

### 3. MS/MS Ions Search

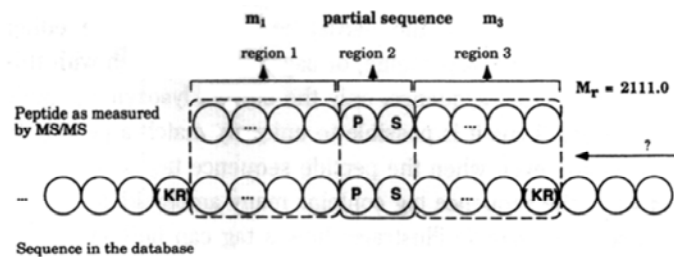
Uninterpreted MS/MS data from a single peptide or from a complete LC-MS/MS run

**MASCOT** : *Sequence Queries*

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You will remember from the introduction, that sequence queries are searches where mass information is combined with amino acid sequence or composition information



**Figure 1.** Principle of matching peptide sequence tags to a proposed sequence. The upper chain of amino acids represents the peptide sequence as measured by MS/MS (from Table 1 in this example), and the lower chain represents amino acids in the sequence database that the tag is compared to. Note that the partial sequence divides the peptide into three regions. The added mass  $m_1$  of the residues in region 1, together with the N-terminus, is a match criterion as is the added mass in region three,  $m_3$ . In region 2, the sequence is known. Furthermore, it can be required that the peptide obey the cleavage condition of the proteolytic enzyme, marked by KR for trypsin. The left pointing arrow indicates that both search directions may have to be considered.

➤Mann, M. and Wilm, M., *Error-tolerant identification of peptides in sequence databases by peptide sequence tags*. Anal. Chem. 66 4390-9 (1994).

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The best known example is a sequence tag search, where a few residues of amino acid sequence are interpreted from the MS/MS spectrum.

## Search parameters all still apply

- Enzyme
- Modifications
- Charge
- Instrument

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You can enter sequence tags, and other types of query, into the sequence query form.

Remember that all the search parameters, including enzyme specificity, modifications, and precursor charge, still apply to this type of search.

Mascot will look for a match between the tag and the ion series specified by the instrument type. Note that Mascot will only try to match the tag against ion series formed by a single backbone cleavage, and maybe a neutral loss, like y or b\* or y++. It won't try to match against side chain cleavage fragments, like d, v, w or internal fragments.

## Standard sequence tag

### Keyword is tag

### What's (probably) wrong with this tag?

1890.2 tag(1004.1, LSADTG, 1548.5)

### Very unlikely that you would be able to call L from a spectrum. Should be

1890.2 tag(1004.1, [I|L]SADTG, 1548.5)

### Ambiguity is OK as long as it is explicitly represented

877.4 tag(376.2, [I|L][Q|K][I|L], 730.2)

1869.93 tag(345.14, [I|L]A[VG|GV|R][M|F]G, 889.45)

(VG = R, F = MetOx)

Unless you have high energy fragmentation, and are able to distinguish L from I by side chain cleavage fragments, then this tag is wrong. It should be I or L.

Ambiguity in a tag is fine as long as it is recognised and spelt out. Most times, you won't know whether a residue is Q or K. F is almost identical to oxidised M. If the peaks are weak, are you sure you have a mass difference of R, or could it be VG and the intermediate peak is missing?

## Error tolerant sequence tag

**Keyword is etag**

**Peptide in database is**

GVQVETISPGDGR, MH+ = 1314.7

**b ion series tag called from TISP *should be***

1314.7 tag(513.3,T[I|L]SP,911.5)

**But, if unknown modification or SNP increases mass by 100 Da, mass values would become**

N-term side: 1414.7 etag(613.3,T[I|L]SP,1011.5)

C-term side: 1414.7 etag(513.3,T[I|L]SP,911.5)

If the sequence is in the database, it is easier and safer to perform an MS/MS search of the peak list. In this sense, the standard sequence tag is obsolete.

The error tolerant tag, which can find a match when there is an unsuspected modification or a small difference in the sequence, is very powerful and very useful.

Imagine we had an unmodified peptide of MH+ 1314.7 and we interpreted a tag of TISP in the b+ series between peaks at 513.3 and 911.5.

What happens if there is a modification or SNP that increases mass by 100 Da?

If the mod is on the N-term side of the tag, all the masses shift up by 100. However, if it is on the C-term side, only the peptide mass changes.

If the tag was in the y ion series, the reverse would be observed

## Error tolerant sequence tag

### Peptide mass is allowed to change by $\Delta m$

EITHER both fragment ion masses unchanged

OR both fragment ion masses shift by  $\Delta m$

### etags have low specificity

- Use reasonable peptide mass tolerance
- Must select an enzyme

The error tolerant tag allows for this. In effect, it allows the peptide mass to vary and allows the tag to float. However, the tag must stay attached to one end or the other. Either both fragment ion masses are unchanged or both fragment ion masses shift by the same amount as the precursor.

This causes a huge loss of specificity, so we cannot allow etag searches with very wide peptide mass tolerance ( $> 1\%$  or  $> 10$  Da) or with no enzyme specificity. The enzyme specificity in an etag search is never fully specific, in any case, because one end of the peptide can just extend until it finds a cleavage point.

## Sequence tag - general

### Tag can run either way

```
1890.2 tag(1548.5, GTDAS[I|L], 1004.1)
```

```
1890.2 tag(1004.1, [I|L]SADTG, 1548.5)
```

### Can have multiple tags per query

```
879.24 tag(1434.40, VEE, 1077.32) tag(737.22, DFW,  
289.13) tag(1644.53, [L|I]PV, 1335.36)
```

### tag and etags are scored, like ions

- the more tags that match, the higher the score
- all tags are not required to match

### If one tag in a query is etag, they are all etags

### Cannot mix ions() with tag() or etag() in same query

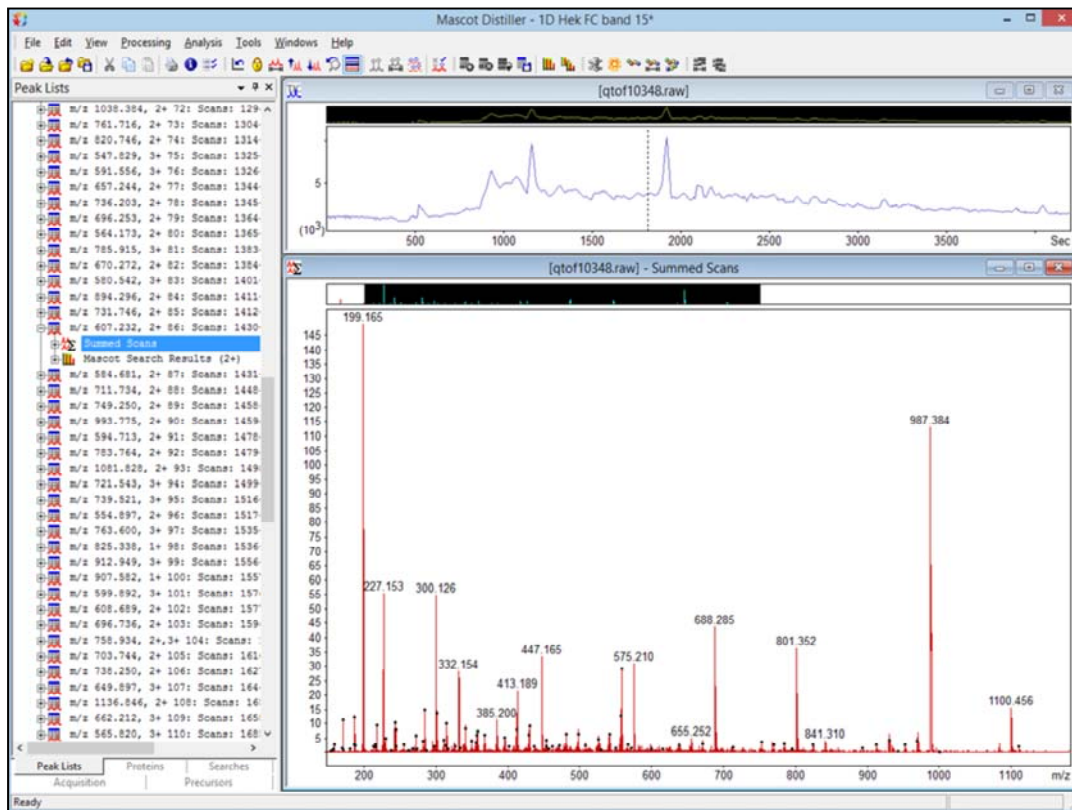
Tags can be entered with the high mass fragment on the left or the right. These two tags are identical

Mascot allows multiple tags in a single query. That is, you can call multiple tags from a single MS/MS spectrum. Tags are scored probabilistically. If one tag is wrong, you can still get a good match from the tags that are correct.

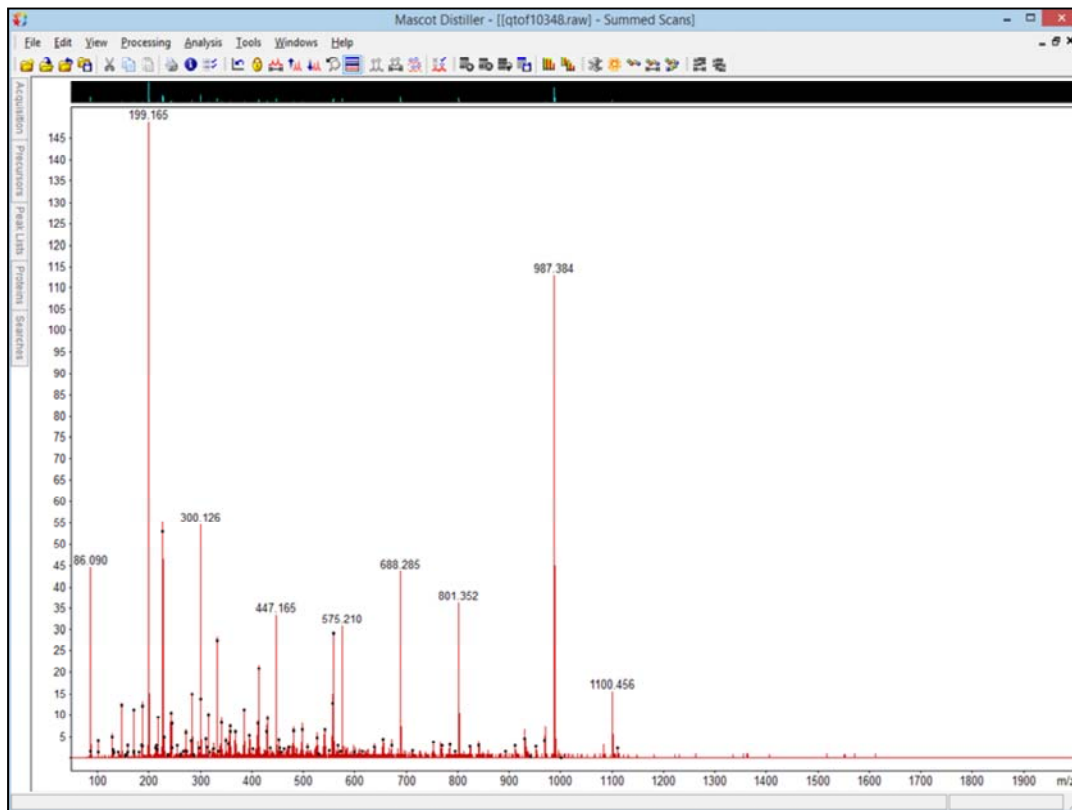
If one tag in a query is an etag then all the tags for that query are treated as etags, (not all tags in the search, just in the query)

Finally, you cannot mix ions qualifiers with tag or etag qualifiers. It would just be too complicated.



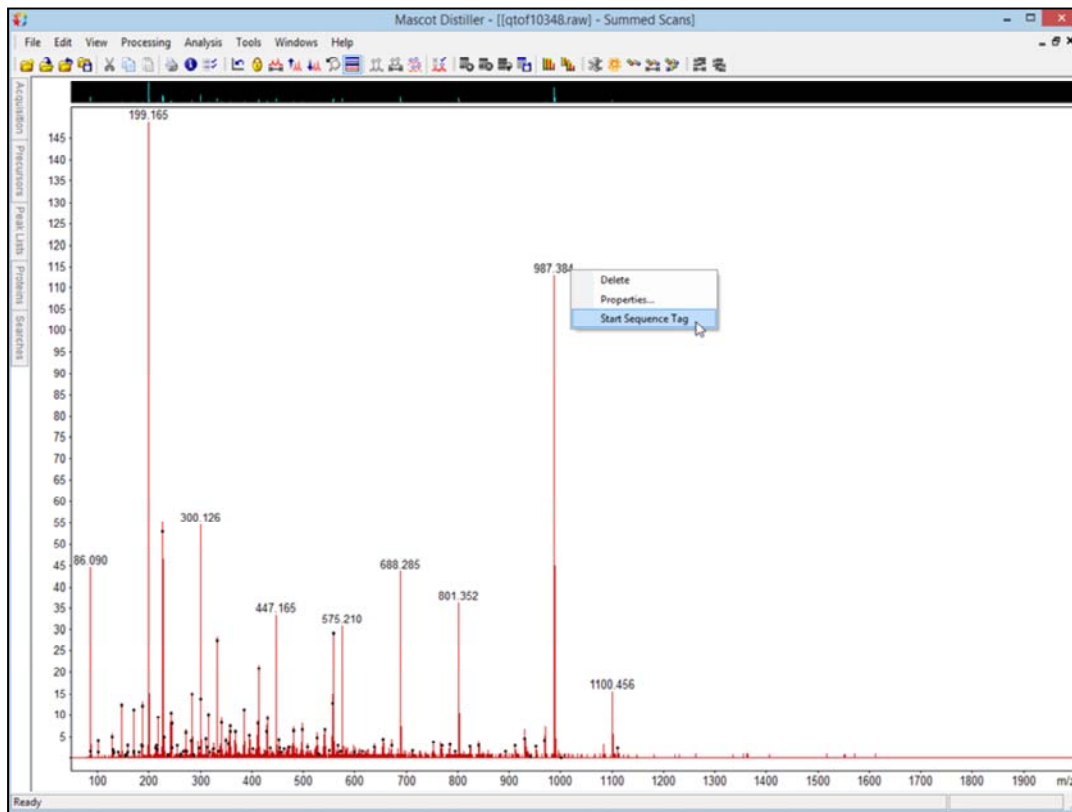


A lot of people call tags using a calculator and a table of mass values. An alternative is to use Mascot Distiller. Let's see how this works

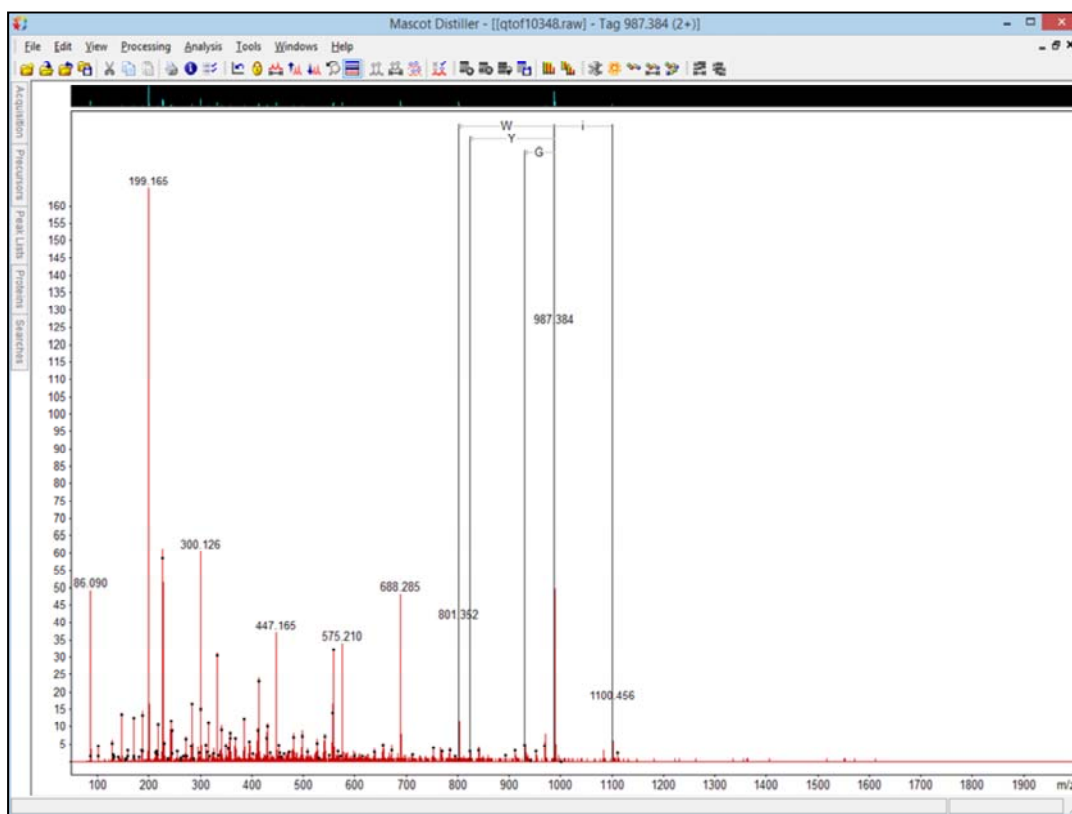


Maximise the window

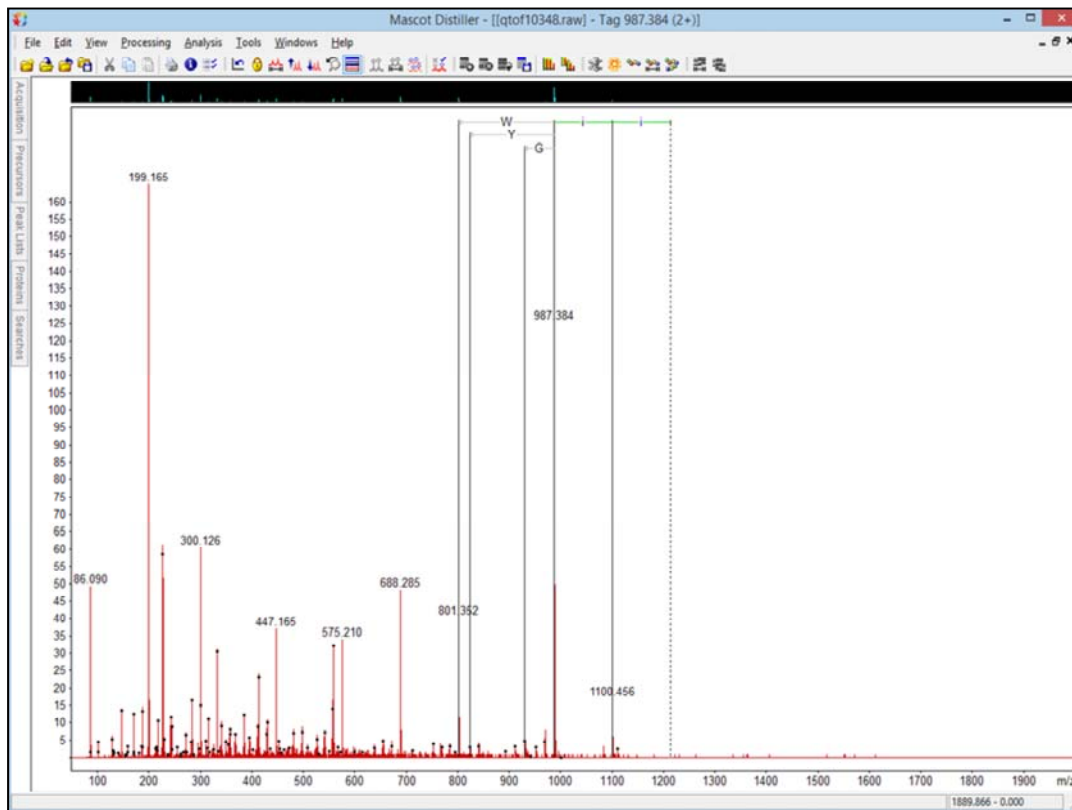
Choose a likely looking peak, such as 987.384



Right click to start a tag

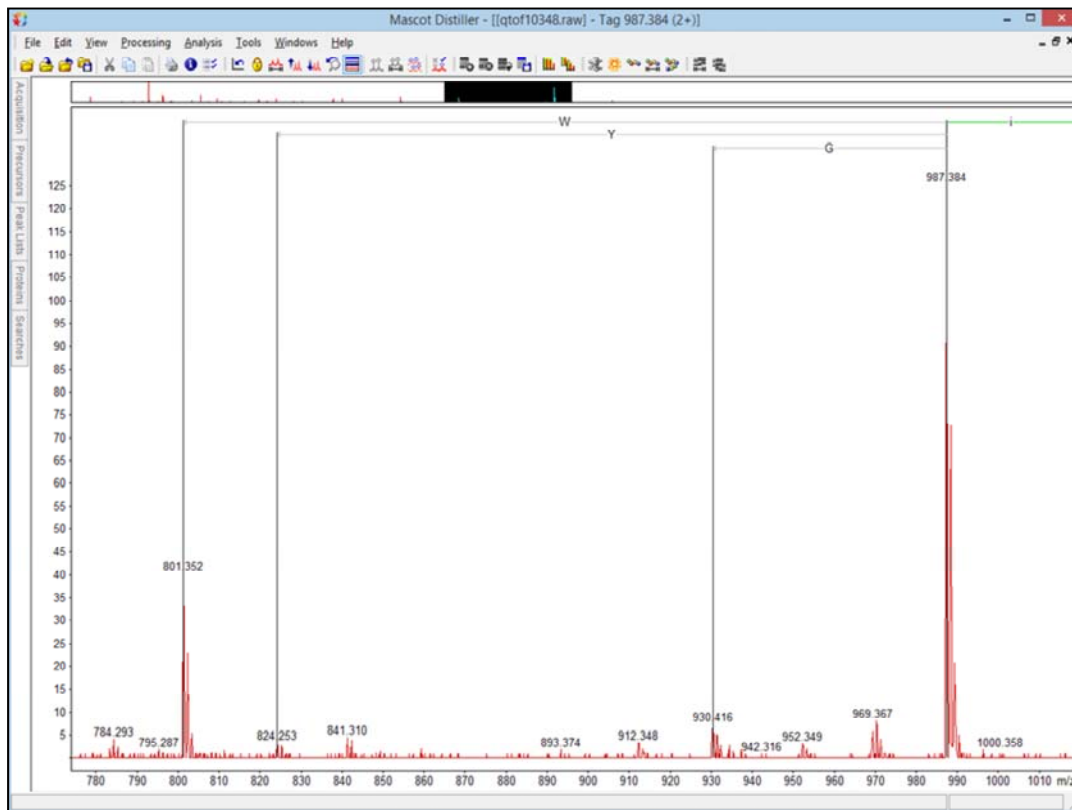


Click on any arrow to extend the tag. Where there's a choice, I'll just go for the biggest peak and stop when it starts to look tricky

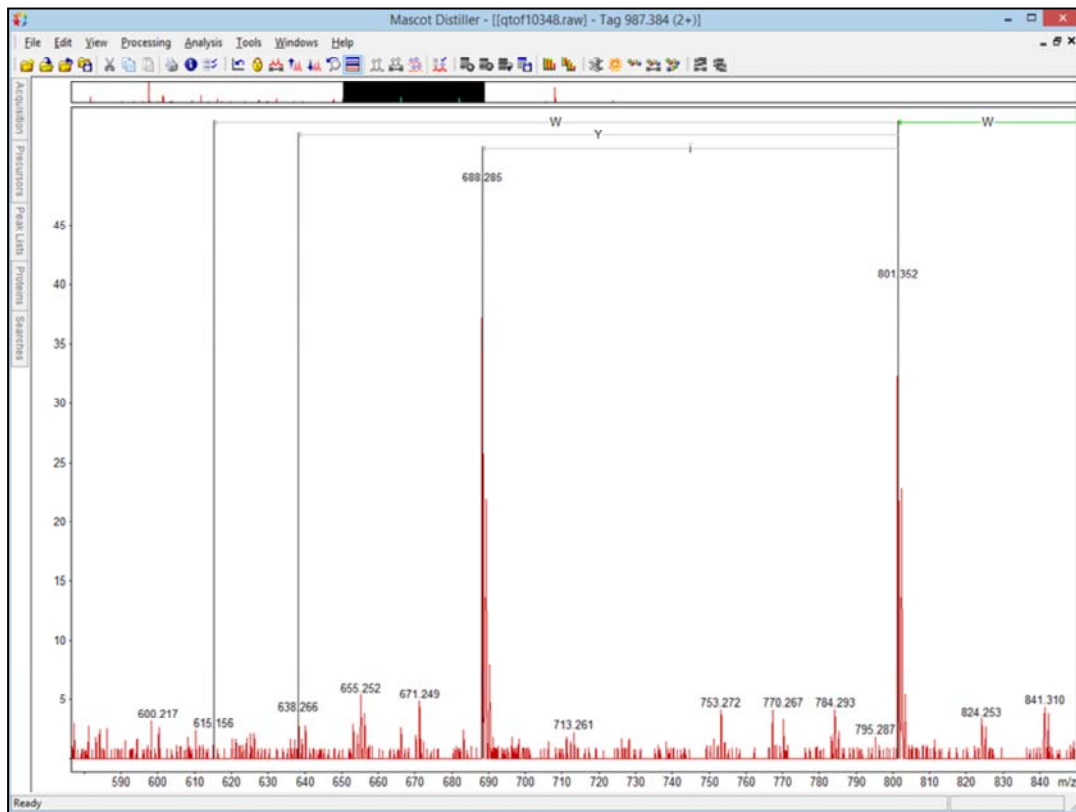


Going to the right, there's no choice and the descender from the second i (=I or L) is dotted to indicate we've reached the precursor mass, which is a bit of luck.

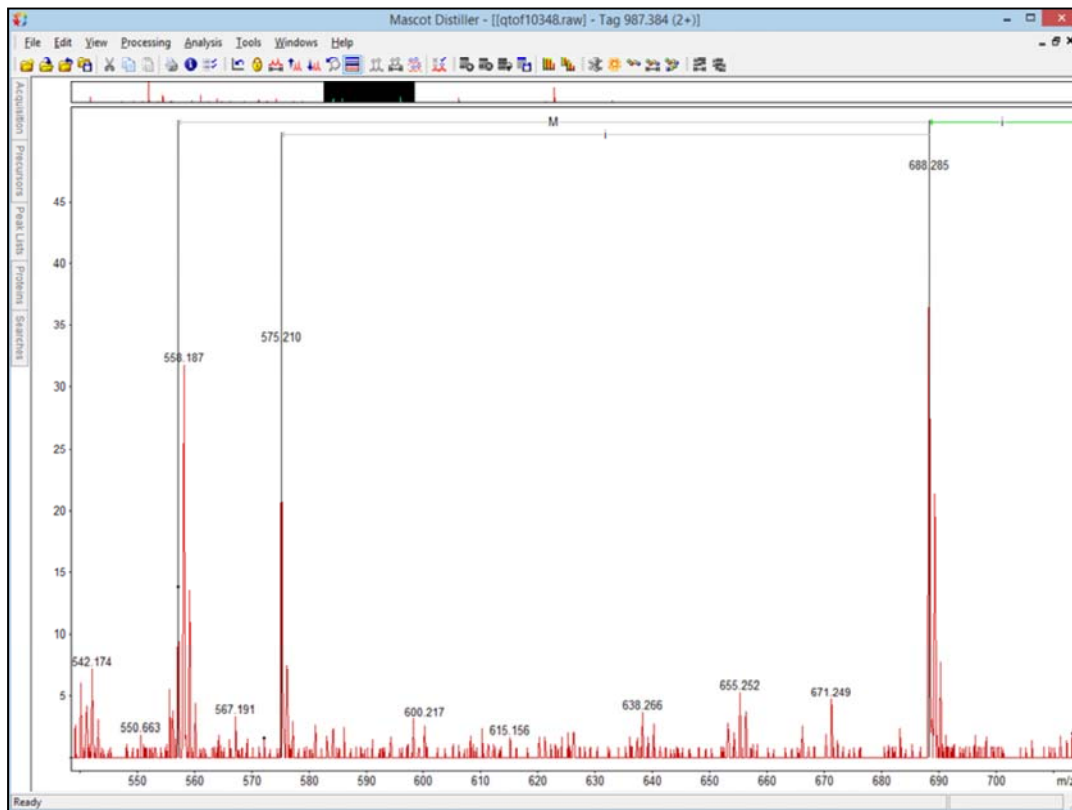
Going to the left, there is a choice of three residues. Zoom in for a closer look



No doubt that W is the most intense, so we click on the arrow head to extend it

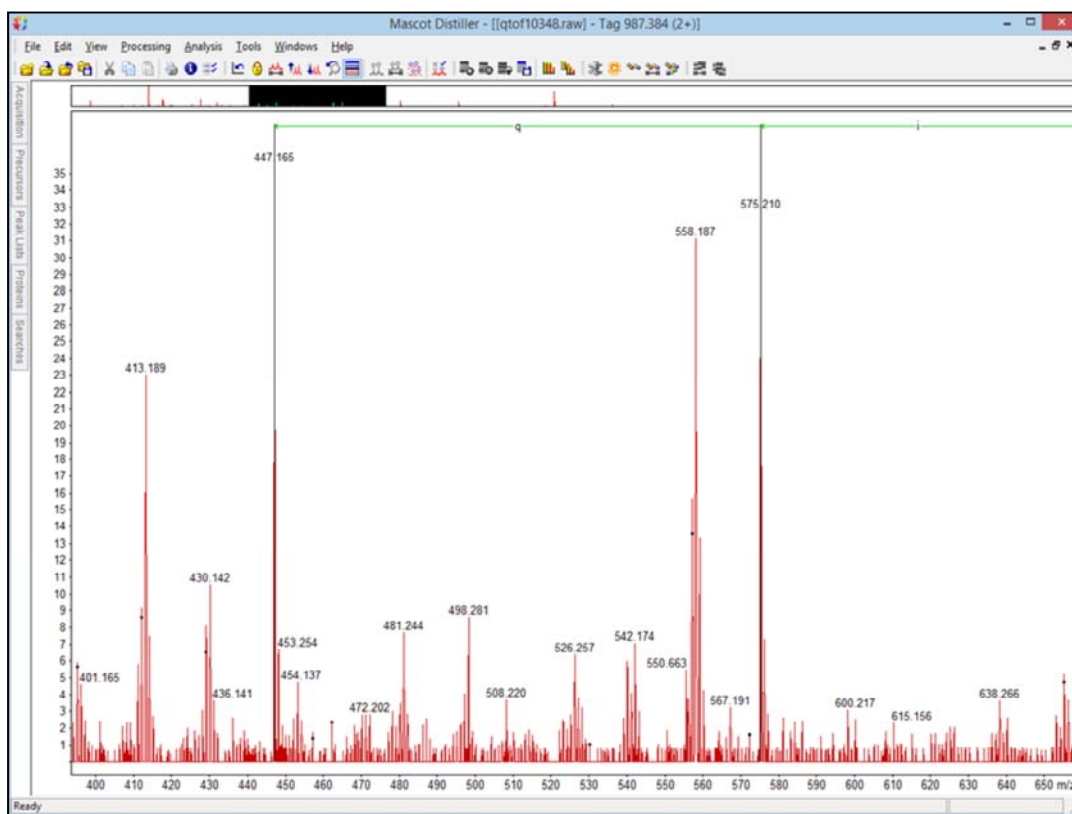


Again, a choice of three. We'll go with i

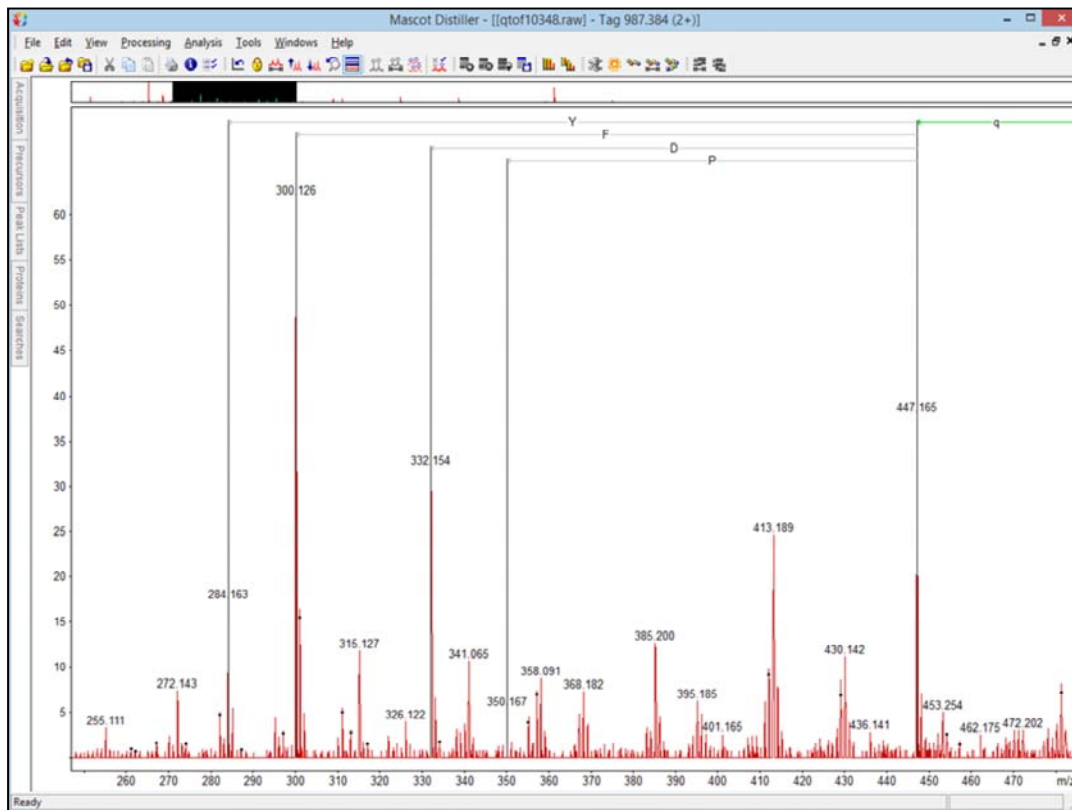


M looks a bit dodgy, so we'll go with I again

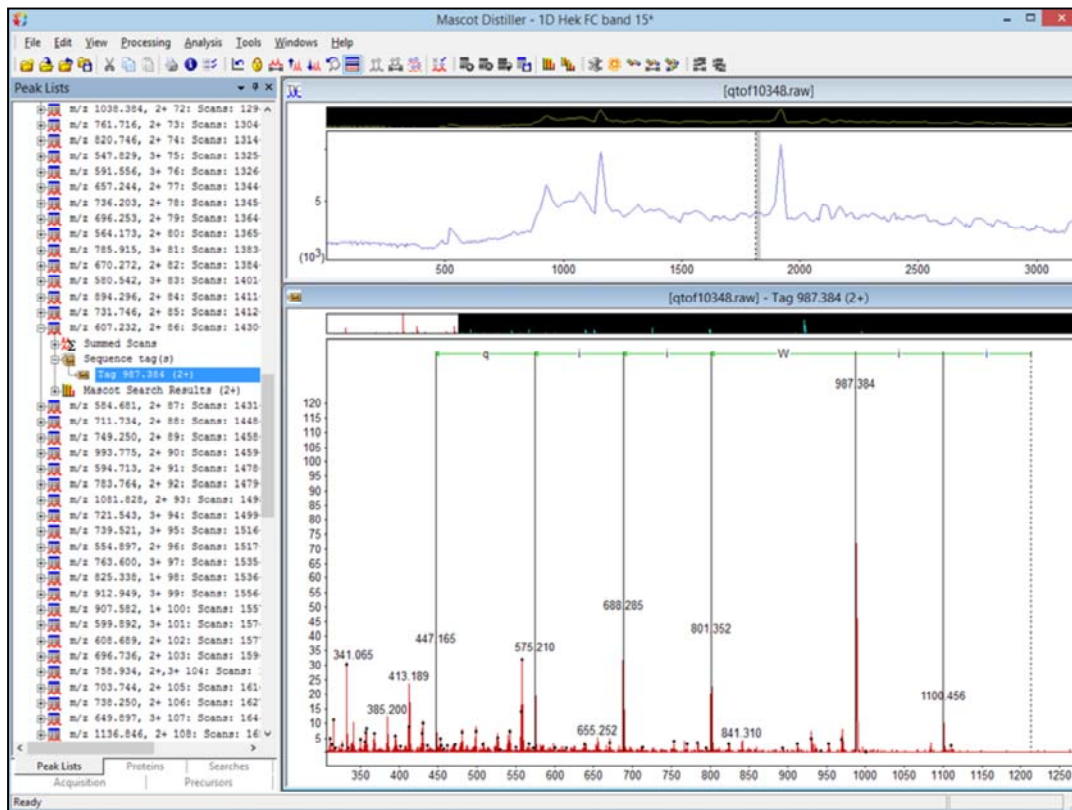




No choice ... good!

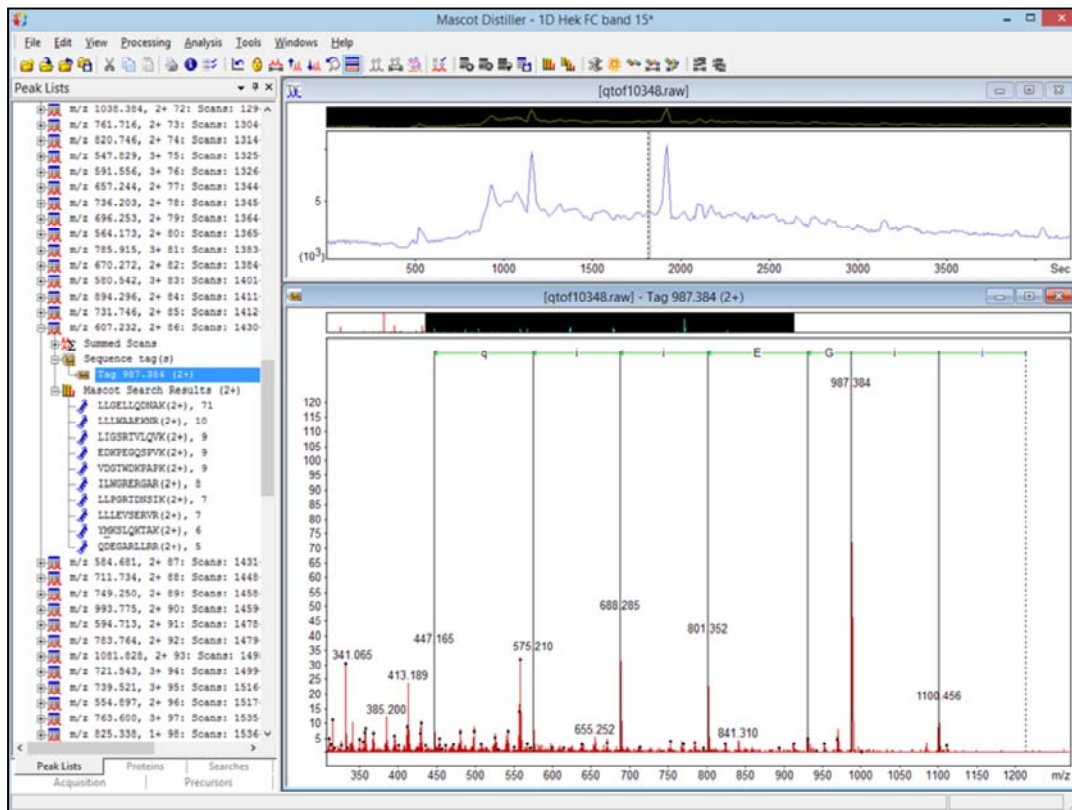


Lots of choice. Maybe time to give up



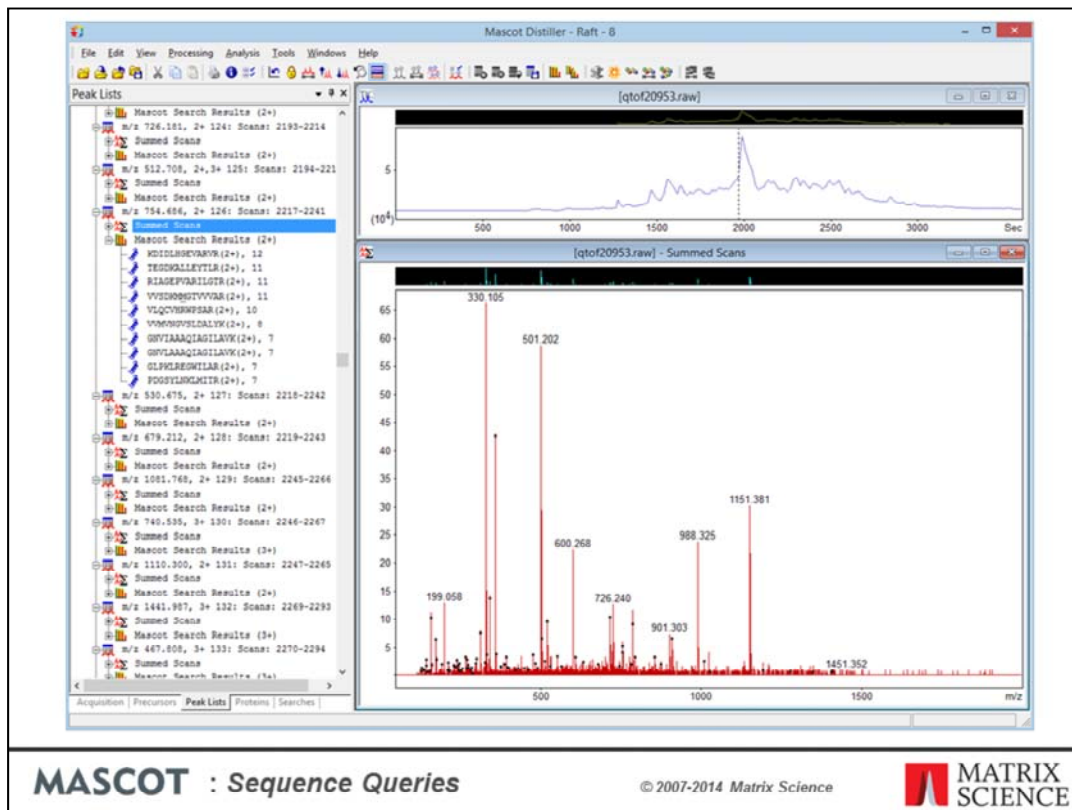
Restore the window arrangement. Here's our tag, but is it correct? Well, we just happen to have database search results from this spectrum





This is what it should have looked like. I obviously need more practice!

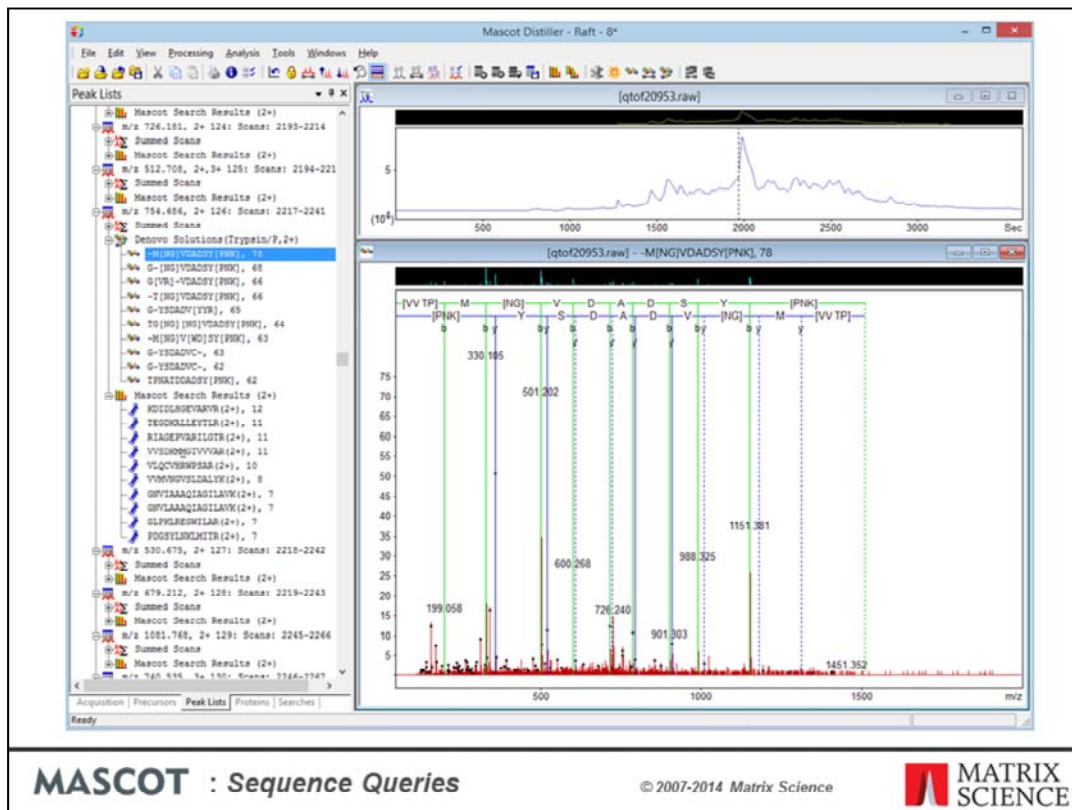
That is a rather artificial example, of course, because we have a good match from the database search, so no-one would need to call a sequence tag



Alternatively, you can automate the process entirely by using the de novo algorithm.

Here's a nice spectrum in another data set where the Mascot database search has failed to find a match

If we right click the peak list and choose de novo ...

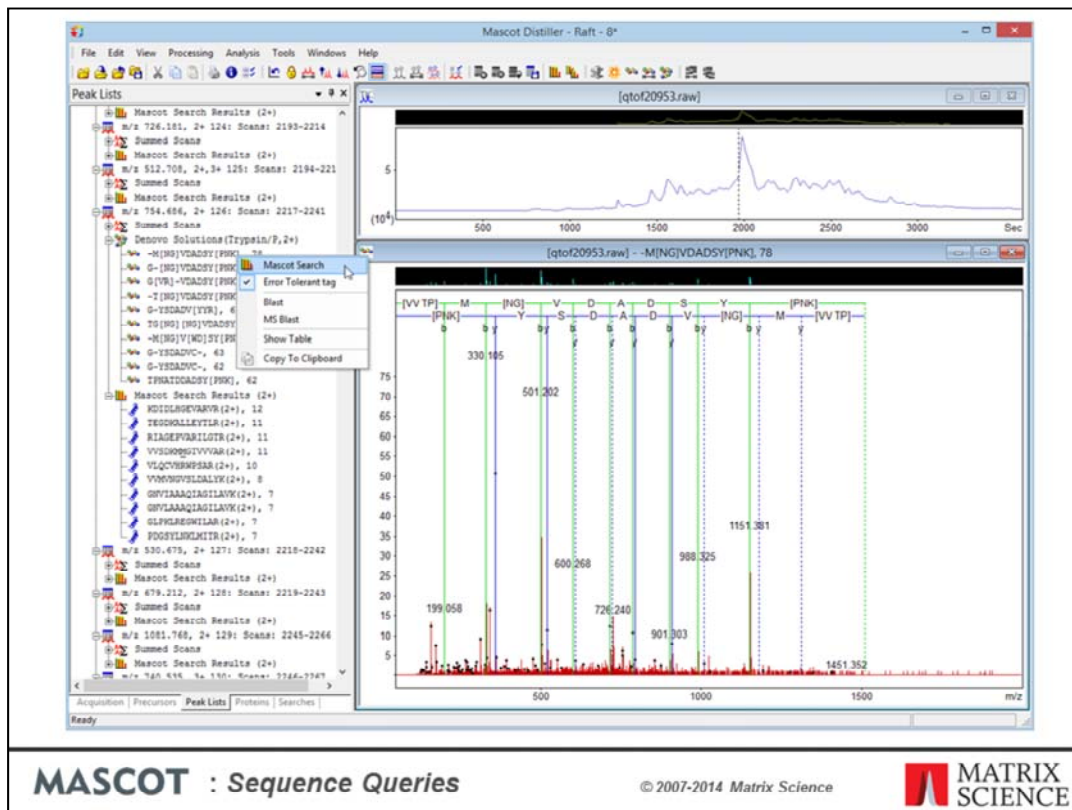


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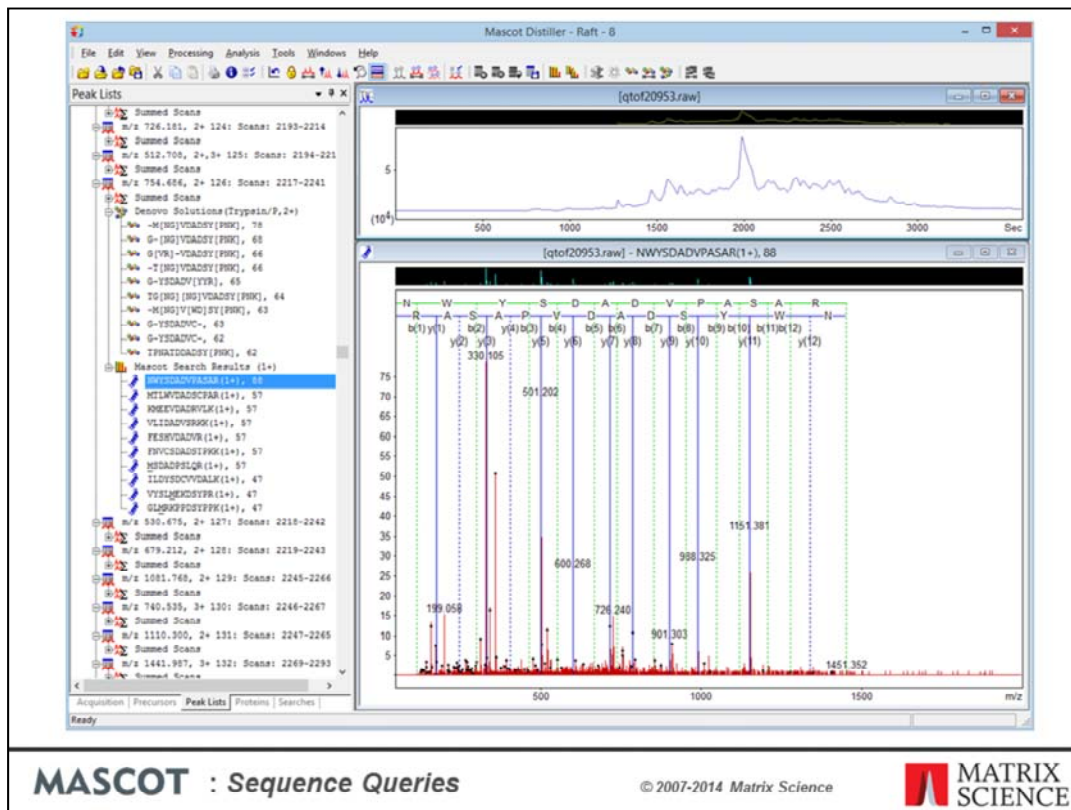
We get a reasonably high scoring solution, but with some uncertainty



Right click the solution and choose Mascot search from the context menu. Note that we have already toggled the tag type to error tolerant







And back comes the result. Note that the results from this most recent search have replaced the original database search. You can switch back to the previous results by selecting them on the searches tab.

This match looks promising. The sequence runs the opposite way from the de novo solution, but this is to be expected by chance if the de novo fails to reach one or other terminus.

If we right click the match and choose to view the full Mascot report in a browser ...

Mascot Search Results

Peptide View

MS/MS Fragmentation of **NWYSDADVPASAR**  
 Found in **PPBI\_HUMAN** in **SwissProt**. Alkaline phosphatase, placental type OS=Homo sapiens GN=ALPP PE=1 SV=2

Match to Query 1: 1507.358224 from (1508.365500.1+) index(0) etag(501.20172.VDA.786.28758) etag(786.28758.DSY.1151.38080) etag(600.26844.DAD.901.30347) etag(794.23058.ADS.521.15202)

Title: 126: Sum of 9 scans in range 2217 (m=32.8655, f=2, i=520) to 2241 (m=33.0972, f=2, i=528)

Monoisotopic mass of neutral peptide Mr(calc): 1450.6477  
 Fixed modifications: Carbamidomethyl (C) (apply to specified residues or termini only)  
 Unsuspected modification: 56.7105 Da, located in the region N-term to 102  
 Ions Score: 88 Expect: 2.1e-005 (1a1a)

#	b	b <sup>+</sup>	b <sup>0</sup>	Seq.	y	y <sup>+</sup>	y <sup>0</sup>	#
1	115.0502	98.0237		N				13
2	301.1295	284.1030		W	1337.6121	1320.5855	1319.6015	12
3	464.1928	447.1663		Y	1151.5327	1134.5062	1133.5222	11
4	551.2249	534.1983	533.2143	S	988.4694	971.4429	970.4588	10
5	666.2518	649.2253	648.2413	D	901.4374	884.4108	883.4268	9
6	737.2889	720.2624	719.2784	A	786.4104	769.3839	768.3999	8
7	852.3159	835.2893	834.3053	D	715.3733	698.3468	697.3628	7
8	951.3843	934.3577	933.3737	V	600.3464	583.3198	582.3358	6
9	1048.4371	1031.4105	1030.4265	P	501.2780	484.2514	483.2674	5
10	1119.4742	1102.4476	1101.4636	A	404.2252	387.1987	386.2146	4
11	1206.5062	1189.4796	1188.4956	S	333.1881	316.1615	315.1775	3
12	1277.5433	1260.5168	1259.5327	A	246.1561	229.1295		2
13				R	175.1190	158.0924		1

NCBI BLAST search of **NWYSDADVPASAR**  
 (Parameters: blastp, nr protein database, expect=20000, no filter, PAM30)  
 Other BLAST [web gateways](#)

All matches to this query

Score	Mr(calc)	Delta	Sequence
88.4	1450.6477	56.7105	<a href="#">NWYSDADVPASAR</a>
57.2	1520.6752	-13.3170	<a href="#">MTLWYDADSCPAR</a>

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We can see from the Peptide View that the best match was obtained by placing a modification delta of +57 Da on the N-term residue. This is almost certainly carbamidomethylation, which often derivatises amino groups. This was why the original database search failed to get a match and illustrates how the error tolerant tag can get a match when the modification is unsuspected or even unknown (not in the modifications list)

## Search strategy

1. Standard Mascot search returns the easy matches
2. Error tolerant search returns additional matches, but only for proteins already identified
3. De novo occasionally returns additional full-length peptide sequences that were not in the database
4. More often, de novo returns partial / ambiguous peptide sequences
  - No real reason to expect additional matches from a tag search
  - Use etag search to find matches to isolated peptides that have a SNP or unsuspected modification
  - Blast or MS-Blast if there is a good stretch of clean sequence

If you want to get as many identifications as possible, as efficiently as possible, you might use a strategy similar to this.

## seq()

- Like a tag, but without fragment mass information
- Most likely, from non-MS sequencing, e.g. Edman  
1234 seq(n-AC[DHK]) seq(c-HI) seq(\*-GF)
- seq() is not scored probabilistically, it is a filter

Prefix	Meaning	Example
b-	N->C sequence	seq(b-DEFG)
y-	C->N sequence	seq(y-GFED)
*-	Orientation unknown	seq(*-DEFG)
n-	N terminal sequence	seq(n-ACDE)
c-	C terminal sequence	seq(c-FGHI)

Besides tag and etag, Mascot supports a number of other sequence qualifiers. One of these is seq()

Note that seq() is a filter. It must be correct or there will be no match

## comp()

- Syntax:**

A number, followed by the corresponding amino acid between square brackets. An asterisk means “one or more”

`comp(2[H]0[M]3[DE]*[K])`

- For ICAT, you might specify**

`comp(*[C])`

- X is not allowed**

- comp() is not scored probabilistically, it is a filter**

The other important one is `comp()`. This would be useful in an ICAT search.  
Note that `comp()` is a filter. It must be correct or there will be no match

## Sequence Tag / Sequence Homology

### MultiTag

- Sunyaev, S., et. al., *MultiTag: Multiple error-tolerant sequence tag search for the sequence-similarity identification of proteins by mass spectrometry*, Anal. Chem. 75 1307-1315 (2003).

### GutenTag

- Tabb, D. L., et. al., *GutenTag: High-throughput sequence tagging via an empirically derived fragmentation model*, Anal. Chem. 75 6415-6421 (2003).

### MS-Blast

- Shevchenko, A., et al., *Charting the proteomes of organisms with unsequenced genomes by MALDI-quadrupole time of flight mass spectrometry and BLAST homology searching*, Analytical Chemistry 73 1917-1926 (2001)

### FASTS, FASTF

- Mackey, A. J., et al., *Getting More from Less - Algorithms for rapid protein identification with multiple short peptide sequences*, Molecular & Cellular Proteomics 1 139-47 (2002)

### OpenSea

- Searle, B. C., et al., *High-Throughput Identification of Proteins and Unanticipated Sequence Modifications Using a Mass-Based Alignment Algorithm for MS/MS de Novo Sequencing Results*, Anal. Chem. 76 2220-30 (2004)

### CIDentify

- Taylor, J. A. and Johnson, R. S., *Sequence database searches via de novo peptide sequencing by tandem mass spectrometry*, Rapid Commun. Mass Spectrom. 11 1067-75 (1997)

As always, there is more information in the Mascot help pages. These references are a good starting point if you are interested in learning more about the potential of combining mass and sequence information.