

# Utilizing the Pegasus® GC-HRT 4D for Improved Yeast Metabolite Characterization

Key Words: Yeast, Metabolomics, Gas Chromatography, High Resolution Mass Spectrometry

## 1. Introduction

Yeast has been used since ancient times for fermentation and production of wine.<sup>1</sup> In modern times, the *S. cerevisiae* strain is the preferred cell workhorse for the production of bioethanol.<sup>2</sup> Yeast also continues to serve as a model for systems biology and is being studied as a potential “cellular factory” for production of additional sustainable chemicals such as valuable organic acids.<sup>3,4</sup> In order for this bioengineering to become realized, the metabolism of yeast must be clearly understood. The biggest bottleneck in metabolomics research and engineering is characterization of the diverse classes of compounds found in living systems. In this study, we explored the utilization of gas chromatography—high resolution time of flight mass spectrometry (GC-HRT) and comprehensive two-dimensional gas chromatography—high resolution time of flight mass spectrometry (GCxGC-HRT) for molecular profiling of yeast. Combining the strengths of GCxGC (increased peak capacity, chromatographic resolution, and the ability to provide cleaner spectra) with high resolution and accurate mass capabilities, the Pegasus GC-HRT 4D allows researchers to take their metabolomics discovery workflows to the next level. Figure 1 clearly demonstrates the benefit of increased chromatographic resolution for complex metabolomic samples.

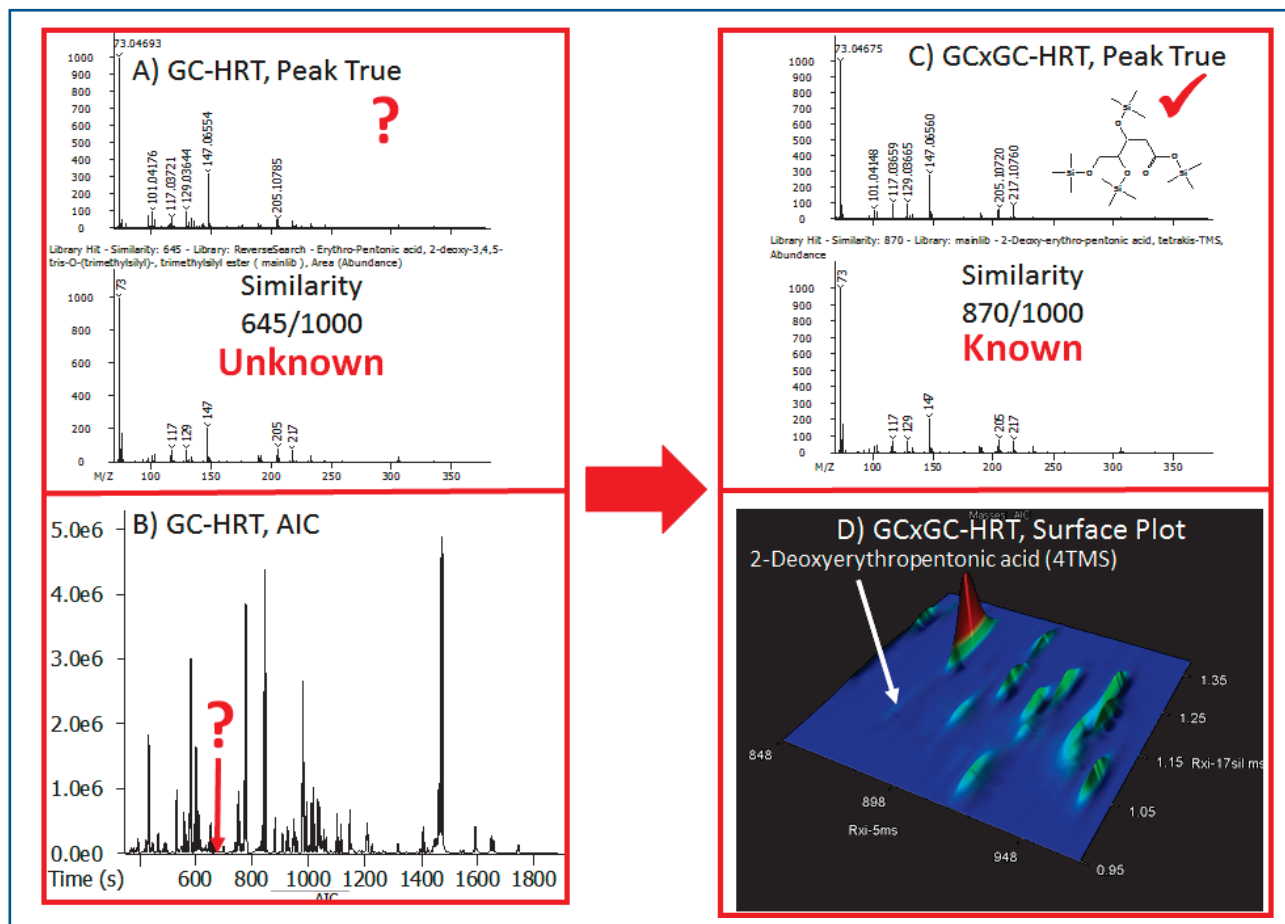


Figure 1. GC-HRT Peak True Mass Spectrum and AIC (A, B) vs. GCxGC-HRT Mass Spectrum and Surface Plot (C, D) for Characterization of 2-Deoxyerythropentonic Acid (4TMS).

## 2. Experimental

Yeast autolysate (Sigma-Aldrich) and dry Baker's yeast (local supermarket) were prepared for analysis using the following protocol. Yeast powder (15–30 mg) was extracted with 10 mL 1:1:2 H<sub>2</sub>O/CH<sub>3</sub>OH/CHCl<sub>3</sub> and filtered into a 2 mL GC vial. The solvents were removed using a Speed Vac. The residue was then derivatized using a two-step procedure: 1) Methoximation with 20 mL of MEOX (20 mg/mL methoxylamine hydrochloride in pyridine, 60 minutes at 60°C) followed by 2) treatment with 80 mL of MSTFA (60 minutes at 60°C).

Both GC and GC×GC data were collected using the instrumental parameters detailed in Table 1. Comprehensive data processing was accomplished using ChromaTOF-HRT® brand software: 1) Mass calibration, 2) Peak Find (Deconvoluted data), and 3) spectral similarity searches using NIST 2014 and Wiley 10.

Table 1. Pegasus HRT GC-High Resolution TOFMS Conditions

Gas Chromatograph	Agilent 7890 with Gerstel MPS2 Autosampler
Injection	1 $\mu$ L, Split 20:1 @ 250°C; (Splitless for CI)
Carrier Gas	He @ 1.0 ml/min, Constant Flow
Column One	Rxi-5 MS, 30 m x 0.25 mm i.d. x 0.25 $\mu$ m (Restek, Bellefonte, PA, USA)
Column Two	Rxi-17SilMS, 0.9 m x 0.25 mm x 0.25 $\mu$ m coating (Restek, Bellefonte, PA, USA)
Temperature Program	1 min at 65°C, ramped 10°C/min to 320°C, held 6 min Secondary oven maintained +5°C relative to primary oven
Modulation	1D acquisitions: 0s 2D acquisitions: 3s *Modulator temperature maintained +15°C relative to 2nd oven
Mass Spectrometer	LECO Pegasus HRT
Transfer Line	300°C
Ion Source Temperature	250°C (EI); 200°C (CI)
Acquisition Mode	High Resolution, R = 25,000 (FWHM)
Ionization Mode	El and or CI (Reagent Gas: 5% NH <sub>3</sub> in CH <sub>4</sub> )
Mass Range (m/z)	35-510 (EI); 60-1200 (CI)
Acquisition Rate	GC-HRT: 10 spectra/s; GCxGC-HRT: 200 spectra/s

## 3. Results and Discussion

A yeast autolysate sample, spiked with twenty-three D and <sup>13</sup>C isotopically labeled standards, was extracted and analyzed by gas chromatography—high resolution time-of-flight mass spectrometry (GC-HRT) operating at a resolving power of 25,000 (m/z = 219). In addition, complementary EI and CI ionization methods, spectral similarity searches of large databases (NIST, Wiley), and formulae determinations for accurate mass (<1 ppm) ions were applied for confident characterization of yeast compounds. The results of High Resolution Deconvolution® (HRD®) are characterized in the analytical ion chromatogram (AIC) for a yeast extract sample which is shown in Figure 2. Table 2 lists the names, formulas, retention times, areas, and similarity values for a representative set of yeast components. These compounds included acids, diacids, amino acids, sugars, bases, nucleosides, and nucleotides. Spectral similarity searches against the NIST and Wiley libraries resulted in an average similarity of 911/1000 for the set.

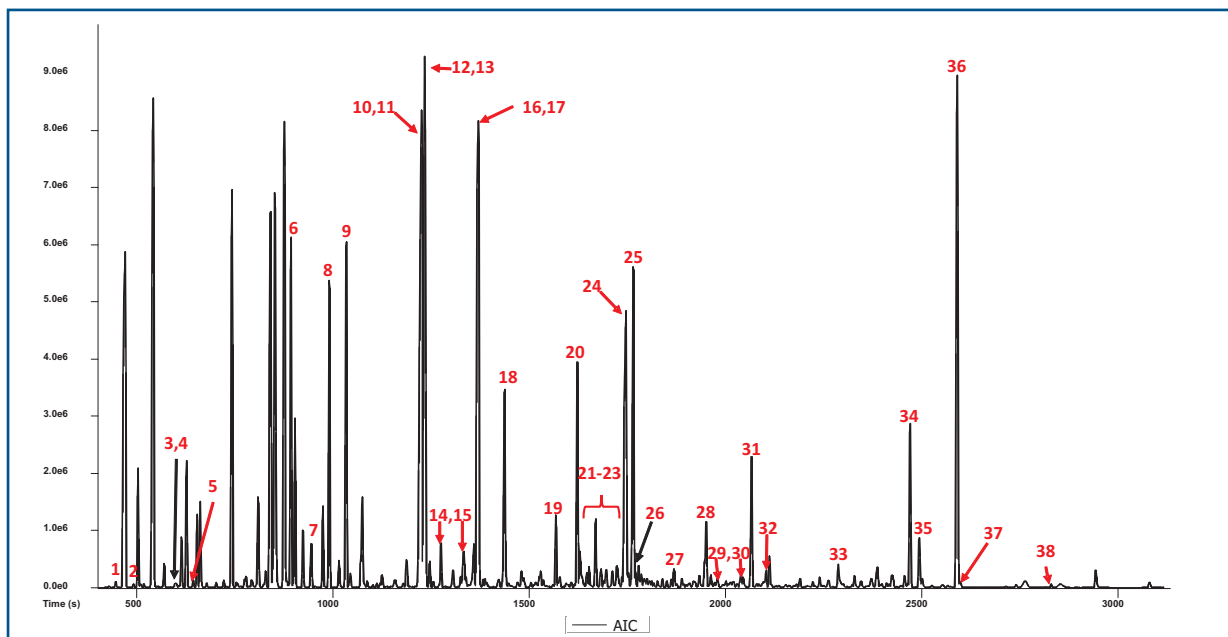


Figure 2. EI-HRT AIC of a Yeast Autolysate.

**Table 2. Representative Compounds in Yeast Autolysate.**

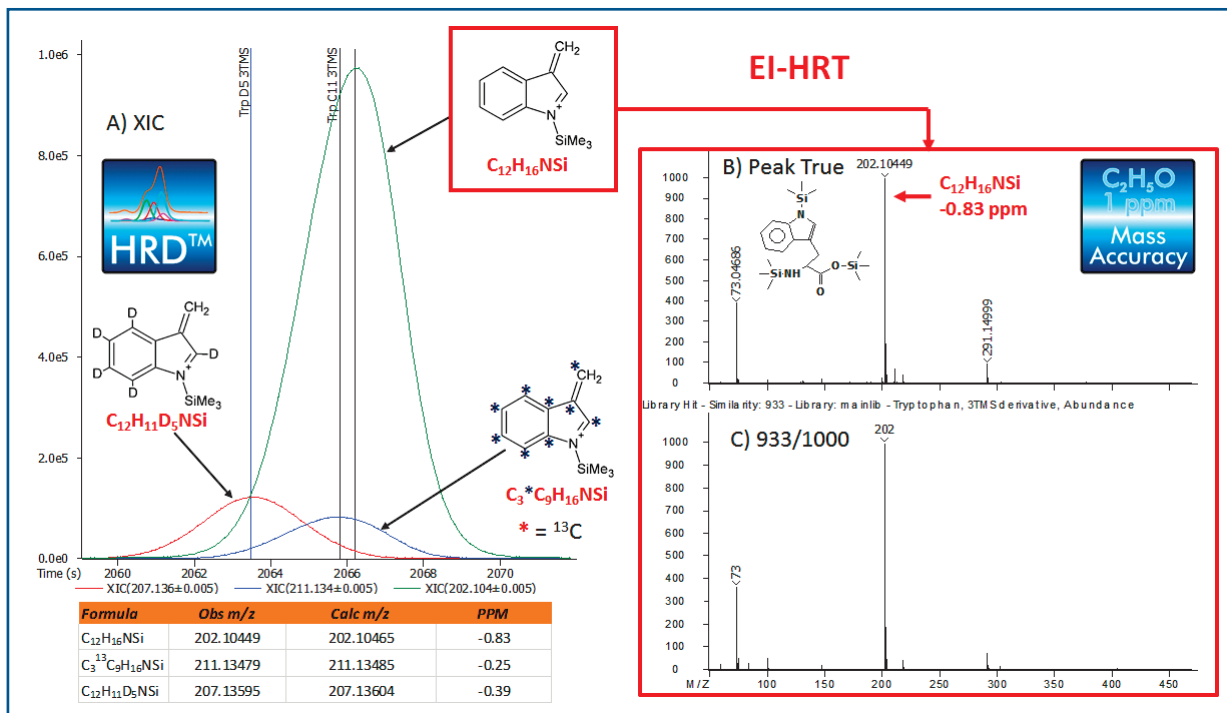
Peak #	Name	Formula	R.T. (s)	Area	Similarity	Peak #	Name	Formula	R.T. (s)	Area	Similarity
1	Pyruvic Acid (MEOX, TMS)	C <sub>7</sub> H <sub>15</sub> NO <sub>3</sub> Si	446	1137387	934	20	Ornithine (4TMS)	C <sub>17</sub> H <sub>44</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>4</sub>	1622	27497881	945
2	Glycolic acid (2TMS)	C <sub>6</sub> H <sub>20</sub> O <sub>3</sub> Si <sub>2</sub>	491	715276	920	21	Mannose (5TMS)	C <sub>21</sub> H <sub>42</sub> O <sub>5</sub> Si <sub>5</sub>	1629	3629958	840
3	2-Furancarboxylic Acid (TMS)	C <sub>8</sub> H <sub>12</sub> O <sub>3</sub> Si	588	236557	910	22	Adenine (2TMS)	C <sub>11</sub> H <sub>21</sub> N <sub>5</sub> Si <sub>2</sub>	1669	10309603	903
4	Oxalic Acid (2TMS)	C <sub>6</sub> H <sub>10</sub> O <sub>4</sub> Si <sub>2</sub>	598	982722	920	23	Cadaverine (4TMS)	C <sub>27</sub> H <sub>46</sub> N <sub>2</sub> Si <sub>4</sub>	1736	1858360	876
5	3-Hydroxybutyric Acid (2TMS)	C <sub>10</sub> H <sub>24</sub> O <sub>5</sub> Si <sub>2</sub>	643	1061101	900	24	Lysine (4TMS)	C <sub>18</sub> H <sub>46</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>4</sub>	1746	35951466	835
6	Isoleucine(2TMS)	C <sub>12</sub> H <sub>29</sub> NO <sub>2</sub> Si <sub>2</sub>	876	8087728	964	25	Tyrosine (3TMS)	C <sub>18</sub> H <sub>35</sub> NO <sub>3</sub> Si <sub>3</sub>	1765	73102265	959
7	Glyceric Acid (3TMS)	C <sub>12</sub> H <sub>30</sub> O <sub>4</sub> Si <sub>3</sub>	944	3040518	831	26	D-Sorbitol (6TMS)	C <sub>24</sub> H <sub>62</sub> O <sub>6</sub> Si <sub>6</sub>	1785	1068732	814
8	Serine (3TMS)	C <sub>12</sub> H <sub>31</sub> NO <sub>3</sub> Si <sub>3</sub>	990	37929232	925	27	Xanthine (3TMS)	C <sub>14</sub> H <sub>28</sub> N <sub>4</sub> O <sub>2</sub> Si <sub>3</sub>	1862	1355710	919
9	Threonine (3TMS)	C <sub>13</sub> H <sub>33</sub> NO <sub>3</sub> Si <sub>3</sub>	1033	42516832	948	28	myo-Inositol (6TMS)	C <sub>24</sub> H <sub>60</sub> O <sub>6</sub> Si <sub>6</sub>	1950	5944599	912
10	Methionine (2TMS)	C <sub>11</sub> H <sub>27</sub> NO <sub>2</sub> SSi <sub>2</sub>	1220	15297318	928	29	Guanine (3TMS)	C <sub>14</sub> H <sub>29</sub> N <sub>5</sub> O <sub>3</sub> Si <sub>3</sub>	1972	1391280	911
11	5-oxo-Proline (2TMS)	C <sub>12</sub> H <sub>23</sub> NO <sub>3</sub> Si <sub>2</sub>	1226	210548974	825	30	Spermine (6TMS)	C <sub>28</sub> H <sub>74</sub> N <sub>6</sub> Si <sub>6</sub>	2036	612262	907
12	Aspartic Acid (3TMS)	C <sub>13</sub> H <sub>31</sub> NO <sub>4</sub> Si <sub>3</sub>	1233	73421702	957	31	Tryptophan (3TMS)	C <sub>20</sub> H <sub>36</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>3</sub>	2066	30428930	927
13	4-Aminobutanoic acid (3TMS)	C <sub>13</sub> H <sub>33</sub> NO <sub>3</sub> Si <sub>3</sub>	1235	46969520	924	32	Spermidine (5TMS)	C <sub>22</sub> H <sub>59</sub> N <sub>5</sub> Si <sub>5</sub>	2103	1388814	940
14	Cysteine (3TMS)	C <sub>12</sub> H <sub>31</sub> NO <sub>2</sub> SSi <sub>3</sub>	1275	5484621	963	33	myo-Inositol Phosphate (7TMS)	C <sub>27</sub> H <sub>69</sub> O <sub>9</sub> PSi <sub>7</sub>	2287	2058184	943
15	Ornithine (3TMS)	C <sub>14</sub> H <sub>36</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>3</sub>	1358	5209095	908	34	Adenosine (4TMS)	C <sub>22</sub> H <sub>45</sub> N <sub>5</sub> O <sub>4</sub> Si <sub>4</sub>	2470	14863948	953
16	Phenylalanine (2TMS)	C <sub>15</sub> H <sub>27</sub> NO <sub>2</sub> Si <sub>2</sub>	1367	52692227	955	35	Maltose (8TMS)	C <sub>36</sub> H <sub>86</sub> O <sub>11</sub> Si <sub>8</sub>	2493	6188006	939
17	Glutamic Acid (3TMS)	C <sub>14</sub> H <sub>33</sub> NO <sub>4</sub> Si <sub>3</sub>	1371	97699927	866	36	Trehalose (8TMS)	C <sub>36</sub> H <sub>86</sub> O <sub>11</sub> Si <sub>8</sub>	2590	77375142	961
18	Asparagine (3TMS)	C <sub>13</sub> H <sub>32</sub> N <sub>2</sub> O <sub>3</sub> Si <sub>3</sub>	1437	24267014	952	37	Guanosine (5TMS)	C <sub>25</sub> H <sub>53</sub> N <sub>5</sub> O <sub>2</sub> Si <sub>5</sub>	2598	718577	804
19	Glycerophosphate (4TMS)	C <sub>15</sub> H <sub>41</sub> O <sub>8</sub> PSi <sub>4</sub>	1567	5723912	905	38	5'-Adenosine Monophosphate (5TMS)	C <sub>25</sub> H <sub>54</sub> N <sub>5</sub> O <sub>7</sub> PSi <sub>5</sub>	2830	454839	887

Accurate mass molecular and fragment ions were leveraged to confirm the identity of compounds. For example, observed m/z and calculated m/z values for molecular and fragment ions for a small set of amino acids are listed in Table 3. The average absolute mass accuracy in parts per million (ppm) for these amino acids was 0.72.

ChromaTOF-HRT data acquisition and data processing software combines the power of accurate mass and high resolving power for an extra dimension of separation—High Resolution Deconvolution (HRD). This is exemplified in Figure 3 where ChromaTOF-HRT processing resulted in deconvolution of coeluting native and labeled (D<sub>5</sub> and <sup>13</sup>C<sub>11</sub>) tryptophan (3TMS). The Peak True (Deconvoluted) mass spectrum was searched against the NIST 14 database and resulted in a match of 933/1000 for the native amino acid. Mass accuracy values for the D<sub>5</sub>, <sup>13</sup>C<sub>9</sub>, and native [M-C<sub>8</sub>H<sub>20</sub>NO<sub>2</sub>Si<sub>2</sub>]<sup>+</sup> fragments were -0.39, -0.25, and -0.83 ppm respectively.

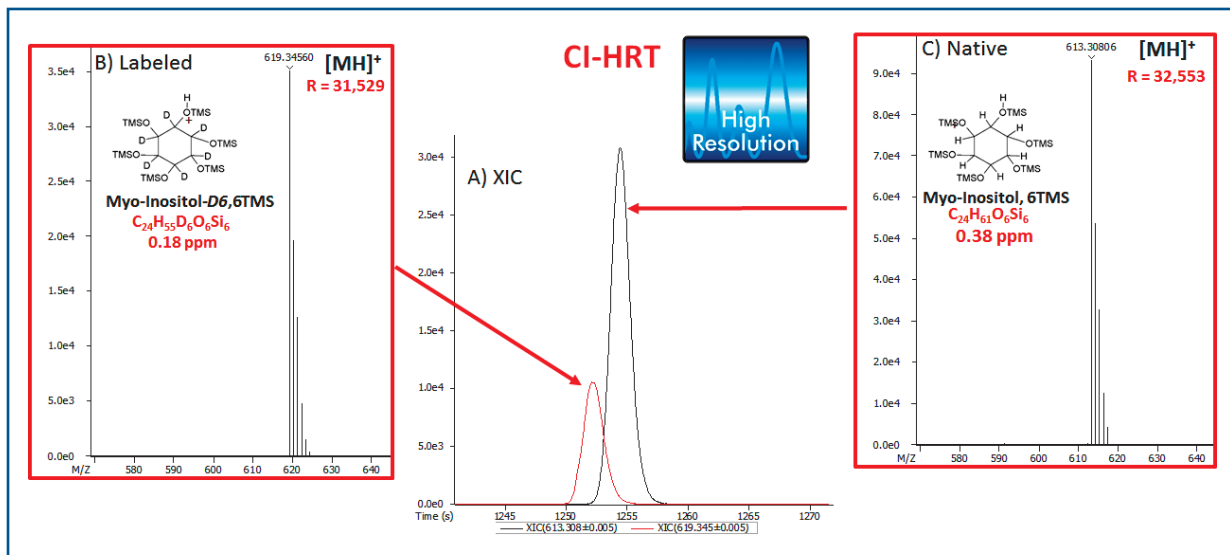
**Table 3. Accurate Mass Values for Amino Acids in Yeast Autolysate.**

Name	Formula	R.T. (s)	Ion	Obs m/z	Calc m/z	PPM
Glycine (3TMS)	C <sub>11</sub> H <sub>29</sub> NO <sub>2</sub> Si <sub>3</sub>	893	M <sup>•+</sup>	291.14974	291.15006	-1.09
Serine (3TMS)	C <sub>12</sub> H <sub>31</sub> NO <sub>3</sub> Si <sub>3</sub>	990	[M-C <sub>4</sub> H <sub>9</sub> O <sub>2</sub> ] <sup>+</sup>	204.12358	204.12344	0.68
Methionine (2TMS)	C <sub>11</sub> H <sub>27</sub> NO <sub>2</sub> SSi <sub>2</sub>	1220	M <sup>•+</sup>	293.12956	293.12955	0.00
5-oxo-Proline (2TMS)	C <sub>11</sub> H <sub>23</sub> NO <sub>3</sub> Si <sub>2</sub>	1226	M <sup>•+</sup>	273.12096	273.1211	-0.50
Aspartic Acid (3TMS)	C <sub>13</sub> H <sub>31</sub> NO <sub>4</sub> Si <sub>3</sub>	1233	M <sup>•+</sup>	349.15586	349.15554	0.92
Ornithine (3TMS)	C <sub>14</sub> H <sub>36</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>3</sub>	1358	M <sup>•+</sup>	348.20799	348.20791	0.22
Phenylalanine (2TMS)	C <sub>15</sub> H <sub>27</sub> NO <sub>2</sub> Si <sub>2</sub>	1367	[M-C <sub>7</sub> H <sub>7</sub> ] <sup>+</sup>	218.10245	218.10271	-1.18
			[M-C <sub>4</sub> H <sub>9</sub> O <sub>2</sub> ] <sup>+</sup>	192.12014	192.1203	-0.86
Glutamic Acid (3TMS)	C <sub>14</sub> H <sub>33</sub> NO <sub>4</sub> Si <sub>3</sub>	1371	M <sup>•+</sup>	363.17114	363.17119	-0.13
Asparagine (3TMS)	C <sub>13</sub> H <sub>32</sub> N <sub>2</sub> O <sub>3</sub> Si <sub>3</sub>	1437	M <sup>•+</sup>	348.17121	348.17152	-0.89
Ornithine (4TMS)	C <sub>17</sub> H <sub>44</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>4</sub>	1622	M <sup>•+</sup>	420.24718	420.24744	-0.60
Lysine (4TMS)	C <sub>18</sub> H <sub>46</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>4</sub>	1746	M <sup>•+</sup>	434.26272	434.26309	-0.84
Tyrosine (3TMS)	C <sub>18</sub> H <sub>35</sub> NO <sub>3</sub> Si <sub>3</sub>	1765	[M-C <sub>10</sub> H <sub>15</sub> O <sub>Si</sub> ] <sup>+</sup>	218.10243	218.10271	-1.30
Tryptophan (3TMS)	C <sub>20</sub> H <sub>36</sub> N <sub>2</sub> O <sub>2</sub> Si <sub>3</sub>	2066	[M-C <sub>8</sub> H <sub>20</sub> NO <sub>2</sub> Si <sub>2</sub> ] <sup>+</sup>	202.10449	202.10465	-0.83



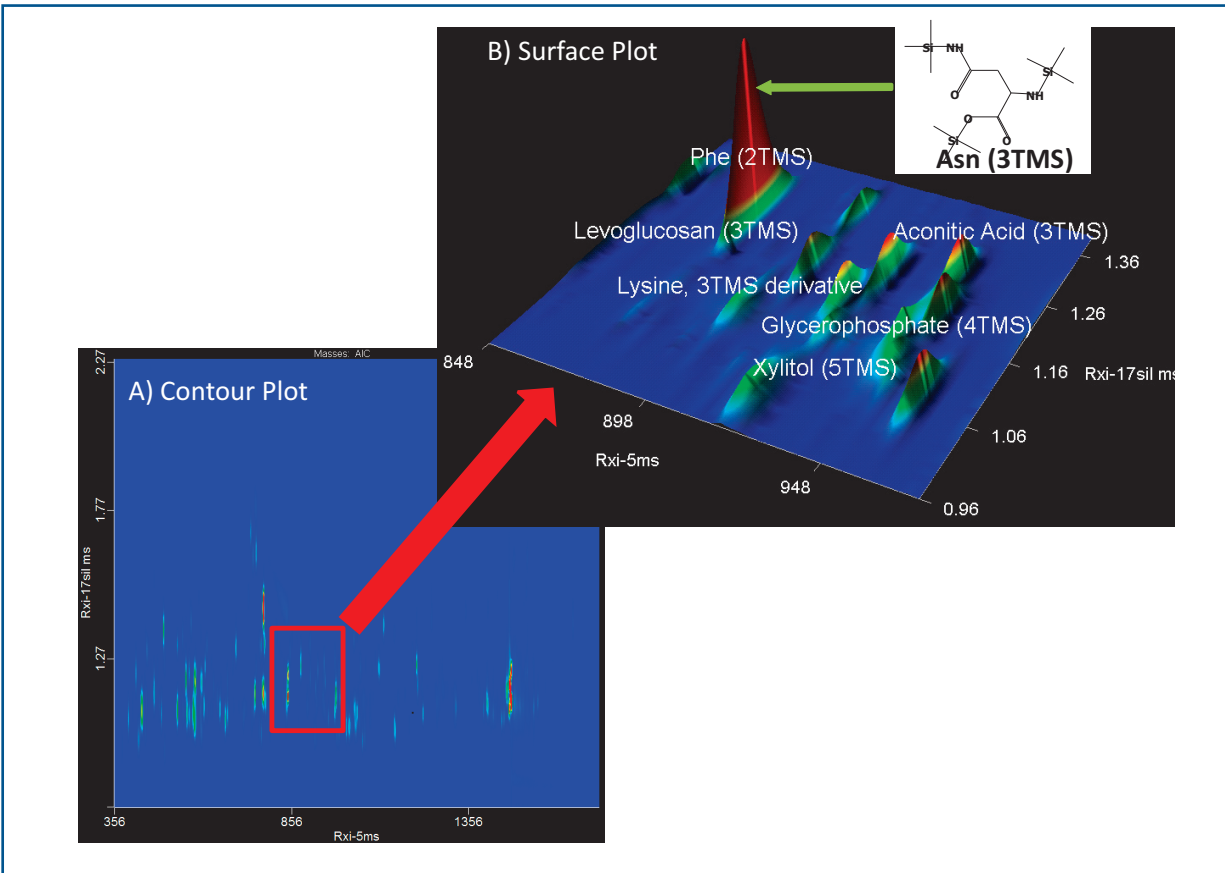
**Figure 3. A) XIC showing  $[M-C_6H_{20}NO_2Si_2]^+$  for  $D_5$ ,  $^{13}C_9$  and Labeled Tryptophan (3TMS) Fragments. B) Peak True Mass Spectrum for Native Tryptophan (3TMS). C) NIST Library Mass Spectrum—Tryptophan (3TMS).**

Complementary CI-HRT data was also used to confirm the identity of yeast compounds. For example, the XICs for the protonated, high resolution molecular ions of D6 ( $m/z = 619.34560$ ,  $R = 31,529$ ) and native ( $m/z = 613.30806$ ,  $R = 32,553$ ) myo-inositol (6TMS) are displayed in Figure 4. Formulas and mass accuracy values for these adduct ions were  $C_{24}H_{55}D_6O_6Si_6$  (0.18 ppm) and  $C_{24}H_{61}O_6Si_6$  (0.38 ppm) respectively.

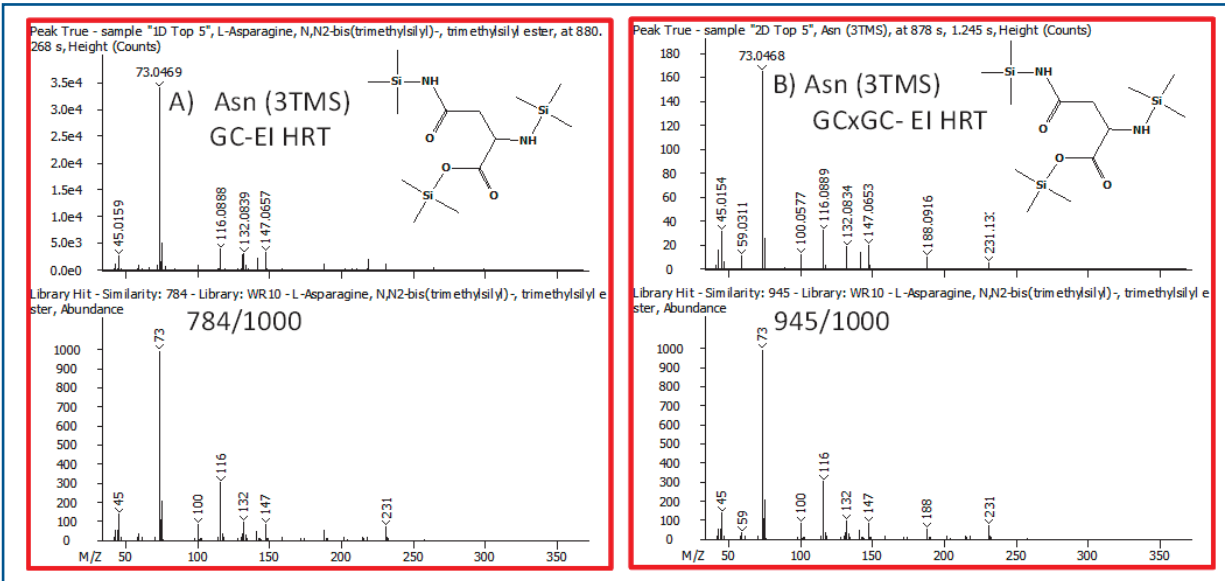


**Figure 4. A) XIC of Labeled and Native Myo-Inositol. CI-HRT Peak True Mass Spectra for Labeled (B) and Native (C) Myo-Inositol.**

The Pegasus GC-HRT 4D can be operated in 1D or 2D modes by a simple selection of operation modes in the software and specifying a modulation period. The enhanced chromatographic capability of the HRT 4D makes it an ideal discovery tool for metabolomics profiling as exemplified in the production of augmented mass spectral data with confidently identified compounds. An expansion of the contour plot for Baker's yeast extract shows the chromatographic resolution provided resulting from the GCxGC-HRT experiment (Figure 5). A comparison of the asparagine (Asn) spectral similarity results for data acquired in GC and GCxGC-HRT mode show a significant improvement in scores from 784 to 945/1000 (Figure 6). Additional examples illustrating the benefits of enhanced chromatographic resolution can be found in Table 4.



**Figure 5. A) GCxGC-HRT Contour Plot for Baker's Yeast Extract. B) Expanded Surface Plot Region Showing Enhanced Chromatographic Resolution.**



**Figure 6. Comparison of GC-HRT Spectral Similarity (A) vs. GCxGC-HRT Similarity (B) for Asn (3TMS) in Baker's Yeast. Notice Improved score due to cleaner spectra resulting from enhanced separation power of GCxGC.**

**Table 4. A Comparison of Spectral Similarity Scores Obtained From GC versus GCxGC-HRT Data Acquisition of Baker's Yeast Extract. The cleaner spectra resulting from GCxGC have a significant effect on library similarity scores.**



Name	Formula	Similarity (GC-HRT)	Similarity (GCxGC-HRT)
Phe (2TMS)	C <sub>15</sub> H <sub>27</sub> NO <sub>2</sub> Si <sub>2</sub>	<b>812</b>	<b>916</b>
2-Deoxyerythropentonic Acid (4TMS)	C <sub>17</sub> H <sub>42</sub> O <sub>5</sub> Si <sub>4</sub>	<b>645</b>	<b>870</b>
Asn (3TMS)	C <sub>13</sub> H <sub>32</sub> N <sub>2</sub> O <sub>3</sub> Si <sub>3</sub>	<b>784</b>	<b>945</b>
Levoglucosan (3TMS)	C <sub>15</sub> H <sub>34</sub> O <sub>5</sub> Si <sub>3</sub>	<b>805</b>	<b>881</b>
Methyl tetradecanoate	C <sub>15</sub> H <sub>30</sub> O <sub>2</sub>	<b>933</b>	<b>942</b>
Xylitol (5TMS)	C <sub>20</sub> H <sub>52</sub> O <sub>5</sub> Si <sub>5</sub>	<b>857</b>	<b>890</b>
(E)-Aconitic Acid (3TMS)	C <sub>15</sub> H <sub>30</sub> O <sub>6</sub> Si <sub>3</sub>	<b>768</b>	<b>910</b>
Glutamine (3TMS)	C <sub>14</sub> H <sub>34</sub> N <sub>2</sub> O <sub>3</sub> Si <sub>3</sub>	<b>835</b>	<b>923</b>
Glycerophosphate (4TMS)	C <sub>15</sub> H <sub>41</sub> O <sub>6</sub> PSi <sub>4</sub>	<b>735</b>	<b>879</b>

#### 4. Conclusion

LECO's Pegasus GC-HRT 4D and ChromaTOF-HRT brand software are powerful tools for confident analysis of metabolomics samples. The combination of EI and CI-HRT high resolution, accurate mass data facilitates rapid compound identification. This was illustrated by Peak Find processing and its ability to separate and identify coeluting deuterated, <sup>13</sup>C labeled and native compounds in complex yeast matrices. The addition of an extra dimension of chromatographic resolution dramatically enhances the discovery capabilities of the GC-HRT 4D making it an extremely valuable instrument for metabolite discovery workflows.

#### 5. References

<sup>1</sup>Chambers P.J. and Pretorius I.S. "Fermenting knowledge: the history of winemaking, science and yeast research" *EMBO reports*, **2010**, 11(12), 914-920.

<sup>2</sup>Araque E., Parra C., Rodriguez M., Freer J. and Baeza J. "Selection of thermotolerant yeast strains *Saccharomyces cerevisiae* for bioethanol production" *Enzyme and Microbial Technology*, **2008**, 43, 120-123.

<sup>3</sup>Raab, AM, Gebhardt G., Bolotina N., Weuster-Botz D. and Lang C "Metabolic engineering of *Saccharomyces cerevisiae* for the biotechnological production of succinic acid" *Metab. Eng.*, **2010**, 12(6), 518-25.

<sup>4</sup>Porro D., Bianchi M.M., Brambilla L., Menghini R., Bolzani D.E., Carrera V., Lievense J., Liu C.-L., Ranzl B.M., Grontall L. and Alberghina L. "Replacement of a metabolic pathway for large-scale production of lactic acid from engineered yeasts" *Applied and Environmental Microbiology*, **1999**, 4211-4215.



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