Large-scale metabolic profiling is expected to develop into an integral part of functional genomics and systems biology. The metabolome of a cell or an organism is chemically highly complex. Therefore, comprehensive biochemical phenotyping requires a multitude of analytical techniques. Here, we describe a profiling approach that combines separation by capillary liquid chromatography with the high resolution, high sensitivity, and high mass accuracy of quadrupole time-of-flight mass spectrometry. About 2,000 different mass signals can be detected in extracts of Arabidopsis roots and leaves. Many of these originate from Arabidopsis secondary metabolites. Detection based on retention times and exact masses is robust and reproducible. The dynamic range is sufficient for the quantification of metabolites. Assessment of the reproducibility of the analysis showed that biological variability exceeds technical variability. Tools were optimized or established for the automatic data deconvolution and data processing. Subtle differences between samples can be detected as tested with the chalcone synthase deficient tt4 mutant. The accuracy of time-of-flight mass analysis allows to calculate elemental compositions and to tentatively identify metabolites. In-source fragmentation and tandem mass spectrometry can be used to gain structural information. This approach has the potential to significantly contribute to establishing the metabolome of Arabidopsis and other model systems. The principles of separation and mass analysis of this technique, together with its sensitivity and resolving power, greatly expand the range of metabolic profiling.

Metabolomics, the comprehensive analysis of metabolites present in a biological sample, has emerged as the third major path of functional genomics beside mRNA profiling (transcriptomics) and proteomics (Fiehn, 2002; Sumner et al., 2003). Metabolomic approaches seek to profile metabolites in a nontargeted way, i.e. to reliably separate and detect as many metabolites as possible in a single analysis. Combined with information on transcript and protein abundance, this would ideally lead to a nearly complete molecular picture of the state of a particular biological system at a given time.

A characteristic of plant life is the production of a vast number of natural compounds, often called secondary metabolites. Secondary metabolites have crucial roles in plant development as well as in the interaction of a plant with its biotic and abiotic environment (Kutchan, 2001). They are "secondary" only in the sense that in different plant species, different sets of metabolites occur (Pichersky and Gang, 2000).

Ample evidence has been obtained in the past three decades for a wide range of functions of secondary metabolites. Arrays of antimicrobial compounds help defend against pathogens (Dixon, 2001; Hahlbrock et al., 2003) and deter herbivores. Secondary metabolites can function as signals internally or in communication with a symbiont (Peters et al., 1986). They provide protection against abiotic stresses such as UV light, drought, or high salt concentrations (Jin et al., 2000).

The majority of steps in the biosynthesis of secondary metabolites is assumed to be catalyzed by specific enzymes (Pichersky and Gang, 2000). Appropriately, completion of the Arabidopsis genome sequence revealed that a significant proportion of the approximately 25,000 predicted genes encode proteins assumed to function in secondary metabolism. The Arabidopsis genome contains >250 cytochrome P450 genes (The Arabidopsis Genome Initiative, 2000), >100 acyl transferase genes, >300 glycosyl transferase genes, >300 glycoside hydrolase genes (http://www.Arabidopsis.org/info/genefamily/genefamily.html), and a large number of genes encoding enzymes such as dioxygenases, O-methyl transferases, terpene synthases, or polyketide synthases. There is a huge discrepancy between the number of these genes and the number of known reactions catalyzed by these types of enzymes in Arabidopsis...
Arabidopsis, leading to the conclusion that a large number of metabolites have yet to be identified. Some of them might even belong to compound classes that so far have not been known to occur in Arabidopsis at all (The Arabidopsis Genome Initiative, 2000). Thus, understanding a significant part of Arabidopsis biology requires methods allowing the sensitive detection and quantification as well as the identification of secondary metabolites. Applying such techniques to various genetic backgrounds, and to different environmental and developmental conditions then would help elucidate the function of such compounds and of the genes involved in their biosynthesis.

Profiling schemes for Arabidopsis and other plants have been developed in recent years (Roessner et al., 2000, 2001; Fiehn et al., 2000; Wagner et al., 2003). The main focus of these mostly gas chromatography (GC)-mass spectrometry (MS)-based approaches have been metabolites of the primary metabolism such as sugars, amino acids, organic acids, or sugar alcohols. Several hundred compounds can be robustly and reliably detected. However, these first pioneering reports already emphasize the need for complementary liquid chromatography (LC)-MS-based approaches to allow a more comprehensive profiling of metabolites (Roessner et al., 2000). The coupling of electrospray ionization (ESI) MS with capillary (Cap) electrophoresis (Soga et al., 2002) and hydrophilic interaction chromatography (Tolstikov and Fiehn, 2002) has been successfully applied to metabolomics problems. Every analytical procedure is necessarily limited as to what type of compounds can be separated and detected. GC-MS is predominantly applied to very polar or unpolar substances, and the main application range of LC-MS is more related to compounds of medium polarity. Furthermore, LC coupled to an MS technique providing soft ionization and high mass accuracy has the potential to generate information useful for the identification of unknown compounds because molecular ions and characteristic fragments can be detected (De Hoffmann, 1996; Niessen, 1999).

We initiated a project aiming at exploiting the potential of hybrid mass spectrometers developed in the past few years for the profiling of Arabidopsis metabolites. Here, we introduce an experimental system based on Cap LC coupled to ESI quadrupole time-of-flight MS (CapLC-ESI-QqTOF-MS). Using this approach, we were able to detect more than 800 mass signals (m/z values) in root extracts and more than 1,400 mass signals (m/z values) in leaf extracts of Arabidopsis. Electrospray as a soft ionization method combined with the enhanced resolution and mass accuracy of the TOF instrument and tandem MS make the identification of unknown compounds feasible, and examples are presented.

RESULTS

The major objective of the work described here was to develop a metabolite profiling scheme using the robust and well-established separation of extracts by LC on reversed-phase material in combination with state-of-the-art MS. The basic assumption was that such an approach would very well complement the pioneering GC-MS-based schemes by extending the range of metabolome analysis to those compound classes not amenable to GC analysis. These would include a significant fraction of the plant secondary metabolism.

A few years ago, QqTOF mass spectrometers were introduced. They combine TOF mass analysis with the established technique of ESI, resulting in high sensitivity, high mass resolution and high mass accuracy (Chernushevich et al., 2001), and are therefore principally well suited for comprehensive metabolite profiling. However, to our knowledge, no reports of such an application of hybrid mass spectrometers have yet been published.

Plant Growth, Extraction, and Cap LC

To be able to grow Arabidopsis with high reproducibility and to have easy access to leaf and root material, we established an aseptic hydroponic growth system. This more laborious approach limits throughput to some extent. The major benefit in addition to a more complete analysis of plant biology is that this system allows us to also biochemically profile the responses in a plant organ not directly exposed to a certain change in the environment.

Plant tissues were routinely harvested after 6 weeks and were frozen in liquid nitrogen. Freeze-drying was omitted to minimize loss of potentially susceptible compounds, such as volatiles, glucosinolates, and glycosides. Frozen plant material was homogenized in liquid nitrogen and was extracted twice with 80% (v/v) aqueous methanol. A third extraction did not produce any considerable gain in yield. Extracts were separated by Cap LC using C18 material. Flow rate and composition of the gradient had to be optimized to obtain good separation and maximum stability of the electrospray. A flow rate of 5.5 μL min⁻¹ and a gradient starting from 5% (v/v) acetonitrile were found to produce the best results.

Manual Data Analysis and Reproducibility

Cap LC was coupled to ESI-QqTOF-MS analysis operated in positive ion mode. Instrument parameters were set in such a way that the mass range from 106 to 1,000 D was monitored simultaneously. The signals generated by ions arriving at the detector were routinely summarized in a spectrum every 2 s for a total of 1,590 spectra during a 53-min run. The resulting total ion chromatogram did not resolve distinct peaks (Fig. 1A). Extensive data deconvolution
was required to extract defined mass spectra. This was done manually by dividing the mass range of 106 to 1,000 D into mass windows of 25 D (Fig. 1B). Mass spectra were then extracted from peaks apparent within a particular mass window (Fig. 1C). Several hundred mass signals are extractable in this way from a single run.

At this stage of manual data deconvolution, quantifiability of mass signals, as well as the reproducibility of the mass spectrometric analysis, of the extraction and of the biological material were assessed. The isoflavone biochanin A was selected as an internal standard. A calibration curve was recorded, and 5 pmol was determined to lie within the range of linearity between amount and signal intensity. When added to a plant extract, the recovery rate of biochanin A was practically 100%. Twenty-one and 24 mass signals in root and leaf extracts, respectively, across the Cap LC run and representing a wide variety of intensities and masses were then randomly chosen (Tables I and II). Analysis of different dilutions of extracts and inspection of resulting calibration curves showed that, for most signals, the linear range covered almost two orders of magnitude. Up to 50-fold dilution resulted in a linear decrease in intensity. Untreated wild-type Arabidopsis Columbia (Col-0) plants were analyzed in four independent experiments. To obtain maximum mass accuracy, each run was recalibrated against five characteristic fragment or adduct ions of sinapoylmalate (Fig. 1C, left) in the case of leaf samples, and five characteristic fragment or adduct ions of hirsutin in the case of root samples. The selected signals were then quantified by integrating the peak areas using AnalystQS and by normalizing them to the fresh weight of the plant material extracted.

The reproducibility of the CapLC-ESI-QqTOF analysis was determined by extracting the same material three times and by running three times the same extract. The cumulative sd for repeated analysis of the same extract was 11.1% (Fig. 2, A and C). When different extracts of the same material were compared, an average variability of 25% ± 17.9% was found for leaves, and 21.6% ± 13.4% for roots. The numbers indicate that the technical reproducibility varies considerably, indicating a dependence on the mass signal in question. When four independent experiments were analyzed, an average biological variation was measured of 35.5% ± 14.0% for leaf signals and 55.9% ± 26.0% for root signals (Fig. 2, B and D). Again, the degree of variation was dependent on the mass signal.

Signals were identified based on tR and mass. The stability of tR was determined for six mass peaks eluting between 25 and 51 min. Seventeen analyses of six different extracts were performed over a period of several days. The maximum sd was 0.3 min and the average sd was 0.25 min, indicating the stability of the Cap LC separation. The mass accuracy was eval-
uated by averaging the mass difference for 11 signals from known compounds. Signal intensities covered a range of about two orders of magnitude. The average mass difference through 18 runs was 9.0 ppm.

Thus, we concluded at this stage that CapLC-ESI-QqTOF-MS can be a valuable tool for the robust and reproducible simultaneous detection and quantification of several hundred mass signals.

### Identification of Known Compounds

As mentioned above, it has to be assumed that most of the Arabidopsis metabolites are as yet unknown. Moreover, unlike in the analysis of the major primary metabolites, standards are available for only very few compounds monitored by LC-MS and, in contrast to GC-MS, extensive mass spectral libraries do not exist for LC-MS. Therefore, being able to directly gain structural information on metabolites detected in profiling experiments is a key requirement for metabolomics. The QqTOF-MS technology has the potential to provide this information because the HR mass of a single ion can be used for calculation of a number of elemental compositions corresponding to those within a window of less than 30 ppm, usually 5 to 15 ppm. As shown, this mass accuracy is routinely achievable in profiling experiments with complex samples. A first evaluation of the potential of QqTOF-MS was carried out with the list of mass signals selected for the assessment of reproducibility. Elemental compositions were calculated. To further restrict the number of possible elemental compositions, the isotopic patterns of the ions of interest were checked for the presence of characteristic heteroatoms, such as sulfur, by comparing the calculated with the measured isotopic distribution. The possible presence of nitrogen can be deduced from even-mass numbered $[M + H]^+$ and/or $[M + Na]^+$ ions, respectively. With this information, a literature search for known natural products could now be performed using commercially available sources such as SciFinder, the Chapman & Hall Dictionary of Natural Products on CD-ROM, the NIST database, etc. Tables I and II list the information obtained on these peaks based on the exact mass and literature data. In leaf extracts, nine out of 24 signals could tentatively be identified, and in root extracts,
10 out of 21 signals could tentatively be identified. For six additional peaks in leaf extracts, it was found that they represent glycosides.

**Automatic Data Deconvolution and Analysis**

Using manual data deconvolution, we were able to demonstrate the potential of CapLC-QqTOF-MS for metabolomics with respect to sensitivity and robustness, as well as the ability to quantify and identify compounds. However, manual data deconvolution is time consuming and therefore obviously not acceptable for any type of metabolomic analysis that is aiming at a considerable throughput. To overcome this limitation, we tested and optimized MetaboliteID, a metabolite processing software (Applied Biosystems, Foster City, CA). MetaboliteID allows us to automatically extract mass spectra from the total ion chromatogram and to generate peak lists displaying $t_R$, and mass and intensity of a peak. We first evaluated MetaboliteID by comparing its output with the results of manual data deconvolution. Following the optimization of data extraction parameters such as minimum signal strength (50 counts s$^{-1}$, 2.5-fold higher than background) and XIC window width (0.35), we analyzed the same set of mass signals as before in the reproducibility experiments. The average variability calculated from the same data sets with the unsupervised method was only slightly higher than that determined manually. The technical variability changed from 25.0% to 34.1% for leaf extracts and from 21.6% to 26.9% for root extracts. Values for the biological variability changed from 35.5% to 40.7% and from 55.9% to 67.5% for leaves and roots, respectively. Thus, the cumbersome manual data deconvolution could be replaced by unsupervised automatic deconvolution without significant losses in reproducibility.

Next, the number of detectable mass signals was determined using the optimized MetaboliteID parameter settings. Through 72 runs of 24 extracts from four independent experiments, on average, 1,415 sig-

### Table II. Mass signals from leaf extracts used for assessing the variance of the CapLC-ESI-QqTOF-MS analysis

A total of 24 signals and the internal standard biochanin A were chosen. Listed here are their HR masses and $t_R$ as identifiers. Based on the mass, elemental compositions were calculated and the corresponding compounds tentatively were identified.

<table>
<thead>
<tr>
<th>MR</th>
<th>$t_R$</th>
<th>Elemental Composition (if available)</th>
<th>Compound</th>
<th>Substance Class</th>
</tr>
</thead>
<tbody>
<tr>
<td>114.0371</td>
<td>23.8</td>
<td>C$_5$H$_8$NS</td>
<td>4-Methylsulphinylbutyl-ITC</td>
<td>Isothiocyanate</td>
</tr>
<tr>
<td>116.0706</td>
<td>4.9</td>
<td>C$<em>5$H$</em>{10}$NO$_2$</td>
<td>Proline [M + H]$^+$</td>
<td>Amino acid</td>
</tr>
<tr>
<td>170.0997</td>
<td>35.9</td>
<td>C$<em>8$H$</em>{12}$NS</td>
<td>8-Methylsulphinyloctyl ITC</td>
<td>Isothiocyanate</td>
</tr>
<tr>
<td>200.0180</td>
<td>23.8</td>
<td>C$<em>6$H$</em>{12}$NO$_2$Na</td>
<td>4-Methylsulphinylbutyl-ITC</td>
<td>Isothiocyanate</td>
</tr>
<tr>
<td>207.0652</td>
<td>25.4</td>
<td>C$<em>{11}$H$</em>{11}$O$_4$</td>
<td>Sinapoyl malate [CVF: sinapic acid + H-H$_2$O]$^+$</td>
<td>Phenylpropanoid</td>
</tr>
<tr>
<td>285.0756</td>
<td>40.7</td>
<td>C$<em>{18}$H$</em>{27}$O$_5$</td>
<td>Biochanin A (internal standard) [M + H]$^+$ or [M + H-H$_2$O]$^+$</td>
<td>Isoflavone</td>
</tr>
<tr>
<td>321.1960</td>
<td>30.3</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>349.2379</td>
<td>31.2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>363.0687</td>
<td>25.4</td>
<td>C$<em>{15}$H$</em>{16}$O$_4$Na</td>
<td>Sinapoyl malate [M + Na]$^+$</td>
<td>Phenylpropanoid</td>
</tr>
<tr>
<td>365.1536</td>
<td>31.2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>423.2206</td>
<td>49.9</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>431.2107</td>
<td>50.4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>494.2567</td>
<td>45.0</td>
<td></td>
<td></td>
<td></td>
</tr>
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<td>502.2501</td>
<td>45.0</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>510.2358</td>
<td>45.0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>579.1708</td>
<td>23.7</td>
<td>C$<em>{22}$H$</em>{31}$O$_{14}$</td>
<td>Kaempferol-3-O-a-L-rhamnopyranoside-7-O-a-L-rhamnopyranoside [M + H]$^+$</td>
<td>Flavone glycoside</td>
</tr>
<tr>
<td>675.2524</td>
<td>31.4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>703.1481</td>
<td>25.4</td>
<td>C$<em>{18}$H$</em>{27}$O$_{16}$Na</td>
<td>[2 Sinapoyl malate + Na]$^+$</td>
<td>Phenylpropanoid</td>
</tr>
<tr>
<td>719.1220</td>
<td>25.4</td>
<td>C$<em>{19}$H$</em>{27}$O$_{18}$K</td>
<td>[2 Sinapoyl malate + K]$^+$</td>
<td>Phenylpropanoid</td>
</tr>
<tr>
<td>735.0755</td>
<td>25.3</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>802.4232</td>
<td>47.6</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>815.4580</td>
<td>46.6</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>829.4661</td>
<td>50.3</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>965.5394</td>
<td>45.0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>984.5300</td>
<td>45.0</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
nals were detected in leaf extracts and 827 in root extracts. The standard deviation was 204 for the number of leaf signals and 180 for the number of root signals. When the same extract was analyzed three times, the numbers varied on average by only 5.8% for leaves and 5.5% for roots.

Tools are required for the processing of such large data sets. We decided to use Excel macros and programming in Visual Basic to establish automatic analysis of peak lists generated by MetaboliteID. As a first step, a procedure for establishing master lists for each analysis was developed. This was necessary to summarize ions that were detected more than once; for instance, because of problematic peak shape. The correct assignment of corresponding signals is essential for comparing different CapLC-ESI-QqTOF-MS analyses.

Figure 2. Technical and biological variance of the CapLC-ESI-QqTOF-MS analysis assessed by analyzing 25 mass signals from leaf extracts and 22 mass signals from root extracts. Signals are identified by mass and $t_r$. Technical reproducibility of the LC-MS analysis of methanolic root (A) and leaf extracts (C) was determined by extracting the same material three times as well as performing three repetitions of the LC-MS measurements. Quantification of 22 selective ion traces in root extracts and 25 ion traces in leaf extracts was carried out. The SD of the three independent LC-MS analyses of a single extract is represented in the white, gray, and black bars of the charts. The cumulative SD for repeated analysis of the same extract showed a value of 11.1%. Comparison of the three different extracts of the same material (striped bar) resulted in an average variability of 25% ± 17.9% for leaves and 21.6% ± 13.4% for roots. When four independent experiments were analyzed, an average biological variation was measured of 35.5% ± 14.0% for leaf signals (D) and 55.9% ± 26.0% for root signals (B).

Figure 3. Mass spectra of two metabolites differentially accumulating in Arabidopsis wild-type (Ler WT) and mutant plants (Ler tt4 mutant). The flavonoid glycoside Rha-Rha-kaempferide ([M+H]$^+$ at $m/z$ 579.1731, in-source fragments at $m/z$ 433.1107 and 287.0500) was not detected in leaves of the tt4 mutant line (A), whereas the glucosinolate gluconasturtiin ([M+Na]$^+$ at $m/z$ 446.0611) accumulated to much higher amounts in the roots of tt4 plants (B).
Table III. List of the most abundant mass signals from root (top) and leaf (bottom) extracts incremented in 100-Da steps

From left to right and indicated by the shading, the five strongest signals are listed. HR mass (m/z) and tR (min) are used as identifiers. Intensities (Int.) are given as area per milligram of fresh weight. Doubly charged ions are marked with the charge state in brackets (2).

<table>
<thead>
<tr>
<th>HR mass [m/z]</th>
<th>tR</th>
<th>Int.</th>
<th>HR mass [m/z]</th>
<th>tR</th>
<th>Int.</th>
<th>HR mass [m/z]</th>
<th>tR</th>
<th>Int.</th>
<th>HR mass [m/z]</th>
<th>tR</th>
<th>Int.</th>
<th>HR mass [m/z]</th>
<th>tR</th>
<th>Int.</th>
</tr>
</thead>
<tbody>
<tr>
<td>100-200</td>
<td>160.08</td>
<td>40</td>
<td>1853.145.05</td>
<td>40</td>
<td>1343.117.06</td>
<td>40</td>
<td>1193.129.06</td>
<td>40</td>
<td>924.128.05</td>
<td>40</td>
<td>445.06</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>200-300</td>
<td>234.10</td>
<td>37</td>
<td>683.228.23</td>
<td>52</td>
<td>572.202.13</td>
<td>27</td>
<td>517.288.29</td>
<td>32</td>
<td>328.281.17</td>
<td>51</td>
<td>257.01</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>300-400</td>
<td>317.12</td>
<td>5</td>
<td>477.315.16</td>
<td>5</td>
<td>384.381.08</td>
<td>5</td>
<td>358.301.14</td>
<td>52</td>
<td>255.353.21</td>
<td>5</td>
<td>144.01</td>
<td></td>
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<tr>
<td>400-500</td>
<td>429.24</td>
<td>5</td>
<td>276.411.14</td>
<td>24</td>
<td>238.415.12</td>
<td>29</td>
<td>113.438.13</td>
<td>24</td>
<td>81.437.37</td>
<td>47</td>
<td>65.01</td>
<td></td>
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</tr>
<tr>
<td>500-600</td>
<td>535.18</td>
<td>29</td>
<td>625.592.16</td>
<td>22</td>
<td>605.577.10</td>
<td>5</td>
<td>482.532.06</td>
<td>22</td>
<td>27.525.18</td>
<td>32</td>
<td>27.01</td>
<td></td>
<td></td>
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<tr>
<td>600-700</td>
<td>623.06</td>
<td>41</td>
<td>360.607.41</td>
<td>52</td>
<td>226.699.36</td>
<td>39</td>
<td>463.444.25</td>
<td>51</td>
<td>364.147.25</td>
<td>39</td>
<td>31.01</td>
<td></td>
<td></td>
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<tr>
<td>700-800</td>
<td>764.50</td>
<td>54</td>
<td>1970.13</td>
<td>5</td>
<td>379.33</td>
<td>41</td>
<td>1781.36</td>
<td>41</td>
<td>1781.36</td>
<td>41</td>
<td>1781.36</td>
<td></td>
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<tr>
<td>800-900</td>
<td>815.25</td>
<td>29</td>
<td>3080.89</td>
<td>28</td>
<td>7810.20</td>
<td>28</td>
<td>6829.23</td>
<td>28</td>
<td>4844.20</td>
<td>28</td>
<td>4844.20</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>900-1000</td>
<td>938.71</td>
<td>54</td>
<td>1970.13</td>
<td>5</td>
<td>379.33</td>
<td>41</td>
<td>1781.36</td>
<td>41</td>
<td>1781.36</td>
<td>41</td>
<td>1781.36</td>
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</tr>
</tbody>
</table>

* Mass signals detected in <75% of the runs.  Signal manually integrated because it was below the 50 cps threshold normally used for MetaboliteID analysis.

Example for Detection of Metabolic Differences between Samples

Having established rapid unsupervised automatic data extraction and processing, we assessed the applicability of our profiling scheme by analyzing a well-characterized mutant with a metabolic defect. Landsberg erecta (Ler) tt4 plants lack a functional chalcone synthase (Shirley et al., 1995) and are therefore deficient in flavonoid biosynthesis. Kaempferol glycosides are the main flavonoids biosynthesized by Arabidopsis plants under normal laboratory conditions (Veit and Pauli, 1999). When Ler and tt4 mutant plants were compared by CapLC-ESI-QqTOF-MS, the expected lack of kaempferol and its glycosides rha-rha-kaempferide and rha-kaempferide in leaves of tt4 plants was detected (Fig. 3A; Table IV). Addi-
tionally, two methoxy-substituted indole carboxalde- 
hydes, probably the 1- and 4-methoxy isomers, 
8-methylthiooctyl amine (m/z 176.1472; Kawabata et 
al., 1989), and probably a methoxy-substituted 
indole-glutathione conjugate, such as L-γ-glutamy-
S-[(1-methoxy-1H-indol-3-yl) methyl]-L-cysteinyl-
Gly (m/z 467.1600; Bjergegaard et al., 2000), were 
found to be reduced in amount. However, both sub-
stances have not been obtained from natural sources.
To also test the feasibility of handling very diverse 
data sets, we performed a comparison of metabolite 
composition of root and leaf tissue. Through nine 
samples each, the average overlap was only 8.7% 
(±1.1%).

Identification of Compounds by Tandem MS

Frequently, specific metabolites display changes 
correlated to a particular genetic background or elic-
ited by an environmental or developmental stimulus.
After detection of such changes by nontargeted 
metabolomic analysis, the identification of the re-
spective metabolites is of key importance for bio-
chemical phenotyping and gene function analysis.

Additionally, identification is desirable to extend the 
catalog of known metabolites for Arabidopsis or any 
other plant species under study.

In-source fragmentation of compounds can already 
provide valuable structural information sufficient for 
identification. This was demonstrated above for the 
mass signals used in the reproducibility experiments (see Tables I and II). Tandem MS represents an ad-
ditional powerful option. The first quadrupole of the 
mass spectrometer can be used as a mass filter and 
the second quadrupole can be used as a collision cell. 
Collision-induced dissociation (CID) yields frag-
ments that can be used for elucidation of structures. 
In the following three examples from the method 
evaluation experiments are documented.

Roots of tt4 mutants were found to accumulate 
high amounts of gluconasturtiin, whereas only minor 
amounts were found in the corresponding root ex-
tracts of the Ler plants (Fig. 3B). The isotopic pattern 
of the observed ion at m/z 446.0623 ([M+Na]+) 
clearly indicates the presence of two sulfur atoms 
(see Fig. 4A). The CID spectrum is dominated by 
fragment ions at m/z 284 ([M+Na-162(glucosyl)]+), 
m/z 266 ([m/z 284-H2O]+), whereas m/z 186

![Figure 4](https://example.com/figure4.png)

**Figure 4.** Identification of significant metabo-
lites of Arabidopsis by HR-ESI-MS/MS (product ion scan). A, Positive ion CID-MS of gluconas-
turtiin (in the box: calculated and measured iso-
topic pattern of the [M+Na]+ ion at m/z 446). B, 
Positive ion ESI-CID-MS of hirsutin.
Table IV. Mass signals differentially expressed between Ler wild-type and tt4 mutant plants Tentative identification is listed whenever possible.

<table>
<thead>
<tr>
<th>HR Mass</th>
<th>(t_r)</th>
<th>Average Fold Change</th>
<th>Molecular Formula (if available)</th>
<th>Substance Class</th>
<th>Compound</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>m/z min</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leaf, reduced</td>
<td>287.0550  23.7</td>
<td>17 (+/−7)</td>
<td>C_{13}H_{11}O_{6}</td>
<td>Flavone Glycoside</td>
<td>Kaempferol [M + H]^+</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>([m/z 579−2x146(Rha)]</td>
</tr>
<tr>
<td></td>
<td>579.1708 23.8</td>
<td>88 (+/−12)</td>
<td>C_{25}H_{20}O_{12}</td>
<td>Flavone Glycoside</td>
<td>Rha-rha-kaempferide [M + H]^+</td>
</tr>
<tr>
<td></td>
<td>433.1129 24.0</td>
<td>11 (+/−7)</td>
<td>C_{25}H_{20}O_{10}</td>
<td>Flavone Glycoside</td>
<td>Rha-kaempferide [M + H]^+</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>([m/z 579−146(Rha)]</td>
</tr>
<tr>
<td>Leaf, higher</td>
<td>661.3178 38.8</td>
<td>4.2 (+/−1.2)</td>
<td></td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>789.4299 49.4</td>
<td>4.2 (+/−1.5)</td>
<td></td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td>Root, reduced</td>
<td>116.0500 33.6</td>
<td>3 (+/−0.1)</td>
<td>C_8H_6N</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>133.0521 33.6</td>
<td>3.4 (+/−0.4)</td>
<td>C_{9}H_{8}NO</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>144.0443 33.6</td>
<td>4.2 (+/−0.8)</td>
<td>C_{9}H_{9}NO</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>176.0711 33.6</td>
<td>3.2 (+/−0.1)</td>
<td>C_{9}H_{9}NS</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>176.1472 22.9</td>
<td>3.5 (+/−0.9)</td>
<td>C_{9}H_{9}NS</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>136.0579 5.3</td>
<td>4.7 (+/−0.1)</td>
<td>C_{9}H_{9}NS</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>467.1600 23.7</td>
<td>5.1 (+/−0.3)</td>
<td>C_{26}H_{33}N_{5}O_{7}S</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td>Root, higher</td>
<td>419.2765 52.1</td>
<td>11.8 (+/−3.2)</td>
<td>C_{26}H_{33}N_{5}O_{7}S</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>423.0106 6.1</td>
<td>7.8 (+/−3.2)</td>
<td>C_{26}H_{33}N_{5}O_{7}S</td>
<td>Methoxy-subst. indole</td>
<td>Methoxyindolecarboxaldehyde</td>
</tr>
<tr>
<td></td>
<td>446.0555 26.6</td>
<td>581.5 (+/−418)</td>
<td>C_{26}H_{35}NO_{5}S_{2}Na</td>
<td>S-Containing compound</td>
<td>2-Phenylethyl glucosinate</td>
</tr>
</tbody>
</table>

(phenylethylisothiocyanate + Na)^+ represents the phenylethyl moiety. Hirsutin was used for our root analysis as a stably occurring reference compound. It is known to arise as a myrosinase-catalyzed degradation product from its parent compound, 8-methylsulphinyloctyl glucosinolate (glucohirsutin). Both metabolites were originally obtained from the seeds of rock cress (Arabis hirsuta) by Kjær and Christensen (1958). EI-MS spectra were given by Bichler (1980). The ESI-TOF mass spectrum displays a prominent key ion at m/z 234 (see Fig. 4B). The CID spectrum displays a prominent key ion at m/z 170 ([M+HCH_4SO]^-), also appearing as an intense in-source fragment, which is a group-characteristic ion of methylsulphinyl isothiocyanates (ITCs). Additional typical ions are m/z 161 ([M+HCH_3-NCs]^-), which is a key fragment of mustard oils, and the stable cyclic immonium ion at m/z 114.

We could confirm the expected reduction of kaempferol-3-O-α-L-rhamnopyranoside-7-O-α-L-rhamnopyranoside in tt4 mutants compared with Ler control plants. Its ESI-CID mass spectrum showed a subsequent loss of the two rhamnosyl moieties from the [M+H]^+ ion at m/z 579 generating fragments at m/z 433 and 287, respectively, and was found to be identical with that of a standard sample (data not shown). The structure of the aglycon was established by comparing the CID-MS of m/z 287 with the [M+H]^+ ion of authentic kaempferol. Thus, characteristic aglycon fragments are formed by subsequent losses of water and carbon monoxide, respectively, or from the retro-Diels-Alder reaction.

**DISCUSSION**

The current status of metabolomics can be viewed as being in some ways equivalent to the situation of sequencing programs such as the Human Genome Project around 1990 (Sumner et al., 2003). The enormous potential of comprehensive biochemical phenotyping for the functional analysis of biological systems is realized and numerous projects have been initiated. However, major technological limitations need to be overcome. For instance, the chemical diversity of the metabolome necessitates the use of different analytical techniques to cover the wide range of polarities found among the metabolites occurring in a cell.

Metabolic approaches aim at monitoring the biochemical status of an organism by simultaneously measuring as many metabolites as possible. A robust and reproducible analysis that provides qualitative and quantitative data and allows high sample throughput is desired. Maybe equally important at this stage is to contribute to cataloging the metabolome of an organism. No metabolome is completely known as of yet. Systematic identification of the metabolites occurring in a species is particularly relevant for plants, given the wealth of natural products they produce. Sequence data indicate that Arabidopsis expresses a large number of enzymes for which substrates and products are unknown (The Arabidopsis Genome Initiative, 2000).

LC coupled to HR-MS has great potential to play an important role in metabolomics as a complement

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to GC-MS. However, reports of such a use are few, and thus far, most concern targeted profiling approaches (Lange et al., 2001; Huhman and Sumner, 2002). Our objective was to explore Cap LC coupled to state-of-the-art ESI-QqTOF-MS for the nontargeted profiling of Arabidopsis extracts. Reversed-phase LC on C18 was chosen because this was found to be suitable and highly reproducible for secondary metabolite profiling in a number of plant species including Arabidopsis (Graham, 1991). In principle, other techniques such as Cap electrophoresis could also be coupled to ESI-QqTOF-MS to further expand the range of metabolite profiling. Hallmarks of TOF-MS are the HR and high mass accuracy that make it an invaluable tool for the identification of compounds and its use for the sensitive simultaneous detection of large numbers of metabolites separable by LC on reversed-phase material. Major question marks concerning this approach were the robustness of the analysis, its reproducibility, and its suitability for high-throughput and quantitative analysis. Our results demonstrate that CapLC-ESI-QqTOF-MS principally meets all the necessary criteria for powerful metabolic analysis.

$t_\text{Rs}$ of compounds eluting from the Cap LC column were found to show only little variation. Optimized flow rate and gradient composition produced a stable ion spray. Evaluation of the mass accuracy showed that the target value in the range of 5 ppm error (Chernushevich et al., 2001) can be reached for many signals even in a complex mixture of compounds, allowing, as discussed below, the calculation of elemental compositions and thereby the tentative identification of compounds. Obviously, the accuracy is correlated with signal strength and the error is higher for signals with intensities closer to background. Robustness of the analysis is illustrated by the fact that the average deviation in intensity for a range of mass signals was no higher than 11% when the same extract was run several times. Reproducibility of $t_\text{R}$ and mass accuracy provided a sufficient basis for the unequivocal detection and identification of mass signals.

Quantification of metabolites is a serious hurdle for metabolomics (Sumner et al., 2003). There are inherent limitations to the dynamic range of MS and, as emphasized by Trethewey et al. (1999), for nontargeted approaches, compromises have to be made concerning the quantification of metabolites. Dilution series of Arabidopsis extracts indicated a satisfactory linear range of the CapLC-ESI-QqTOF-MS analysis for many of the signals tested. However, the linear range is dependent on the metabolite in question, and there is currently no feasible way of exactly assessing it for every signal. As predicted by Fiehn (2002), we therefore see the immediate application of CapLC-ESI-QqTOF-MS predominantly in the highly sensitive and rapid detection of qualitative and also pronounced quantitative metabolic differences.

The potential for such a use is very high because of the large number of mass signals that can be resolved. On average, about 1,400 signals were detected in leaf extracts and about 800 signals were detected in root extracts. Given the limited overlap between root and leaf metabolites, a total of about 2,000 different signals can be detected in extracts from Arabidopsis plants grown under control conditions. Most likely, the numbers will be significantly higher once the analysis is extended to other tissues such as flowers (Chen et al., 2003), to different developmental stages, and to plants exposed to environmental stimuli. Increases in the number of detectable mass signals can be achieved by extending the analysis to the negative mode, which is less effective yet allows the measuring of metabolites not seen in positive mode, such as the intact glucosinolates or salicylic acid (data not shown).

The robustness of the analysis is again indicated by the stability of mass signal numbers when the same extract was analyzed repeatedly. The average variation was no greater than 6%. Because of in-source fragmentation and the formation of adduct ions, the number of mass signals detected is definitely higher than the number of metabolites. It is impossible at this stage to reliably estimate the actual number of metabolites detected.

Technical variance (as the sum of variation introduced by the extraction and by chromatography and mass spectrometric analysis) was below 30% for all but two of the chosen mass signals. This provides a solid basis for the detection of differences between samples. The biological variability of around 40% is similar to what has been reported for other large-scale profiling techniques. Technical and biological variance are dependent on the mass signal in question. Possible causes are, for instance, a coelution of ions that affects quantification through ion suppression and high metabolic fluctuation. Processing of a large number of profiling experiments will be required to eventually define the range of variability for individual mass signals.

The efficient use of CapLC-ESI-QqTOF-MS for gene function analysis implies a considerable throughput of samples. Automatic unsupervised deconvolution of MS data and processing of the extracted information is a must. After extensive optimization, we found the MetaboliteID software able to function sufficiently well for deconvolution. A comparison with the results from manual stepwise extraction and integration of a range of mass signals showed only small differences. Peak lists generated by MetaboliteID are further processed using Excel macros and Visual Basic programs. These allow us to assign peaks correctly based on $t_\text{R}$ and mass. The tools are tailored to discover qualitative and quantitative differences between samples and sets of samples. We tested the procedures by analyzing Ler
wild-type and tt4 mutant plants. The expected difference in kaempferol glycosides due to the lack of a functional chalcone synthase was detected. In addition, we found several as yet unknown differences. Comparison between root and shoot samples demonstrated that very diverse data sets can also be handled. The limited overlap we found is further proof for the fundamental difference between root and leaf metabolism.

CapLC-QqTOF-MS is to be used for the detection and elucidation of metabolic changes elicited by environmental or developmental stimuli, as well as for the determination of metabolic differences attributable to a particular genetic background. Furthermore, substantial progress in establishing the metabolome of Arabidopsis and other model species is envisioned. All of these uses will rely on the potential to identify metabolites. We explored a non-targeted and a targeted way of gaining structural information. The highly accurate mass of the molecular ion and ion-source fragments obtained during routine analysis can be used to tentatively identify metabolites with a good success rate. By systematically analyzing the mass data generated, we predict the ability to identify a large number of compounds that to date have not been known to occur in Arabidopsis. In selected cases of mass signals showing a change of immediate biological interest, CID can be used to generate fragments that, in many cases, provide a basis for an assignment of structure.

As is the case for other profiling schemes, aqueous methanol is used here for the extraction of Arabidopsis metabolites. Obviously, the choice of solvent limits the scope of the analysis, and the use of additional extraction methods is likely to result in the detection of further substance classes. A survey of the compounds tentatively identified in this study shows that the relevant known classes of Arabidopsis secondary metabolites such as indole-derived compounds (e.g. indole acetic acid derivatives), degradations products of glucosinolates (sulfinylthioamides), phenylpropanoids (sinapylmalate), and flavonoids as well as their glycosides (e.g. kaempferol-3-O-α-L-rhamnopyranosid-7-O-α-L-rhamnopyranosid) can be detected by the method described. Indole derivatives, for instance, can be clearly classified by LC-ESI-MS/MS measurements according to their substitution pattern at the ring skeleton (e.g. methoxyindoles). The same is true for the ITCs (e.g. hirsutin). The example of glucostaurin, which contains two sulfur atoms, demonstrates the usefulness of the isotopic pattern as an additional feature to support the proposed elemental composition. Arabidopsis secondary metabolites not covered by this method are mono- and sesquiterpenoids, triterpenoid alcohols, phytoestrogens, waxes, and carotenoids.

**MATERIALS AND METHODS**

**Plant Growth**

Surface-sterilized seeds of the Arabidopsis ecotypes Col-O and Ler as well as of the tt4 mutant line were sown on agarose plugs (0.5%, w/v) and were grown hydroponically in one-tenth Hoagland nutrient solution No. 2 (pH 5.3–5.5; Sigma, St. Louis). The medium was supplemented with iron, chelated by N,N’-di-(2-hydroxybenzoyl)-ethylenediamine-N,N’-diacetic acid to a final concentration of 5 μM Fe-N,N’-di-(2-hydroxybenzoyl)-ethylenediamine-N,N’-diacetic acid (Chaney, 1988). Seedlings were grown for 6 weeks in hydroponic greenhouse boxes containing 10 plantlets each. The nutrient solution in the boxes was changed on a weekly basis and was aerated through 0.2-μm filters. Light conditions in the growth cabinet were fixed to 230 to 240 μE m⁻² s⁻¹ and a photoperiod of 8 h of light/16 h of dark at 23°C, day and night. Leaves and roots of plants were harvested separately approximately 1 h into the light period, pooled, and stored at −80°C.

**Extraction and CapLC-ESI-QqTOF-MS**

For the metabolite LC-MS analysis, freshly ground plant material (about 100 mg) was subjected twice to the following extraction procedure: mixing of the plant material with 200 μL of 80% (v/v) MeOH, sonication for 15 min (20°C–22°C), and centrifugation at 19,000g for 10 min. The extracts were combined, filtered (polytetrafluoroethylene filter, pore size of 0.2 μm), and analyzed by LC-MS. Positive LC-ESI-TOF mass spectra were recorded on an API QSTAR Pulsar Hybrid Quadrupole TOF instrument (Applied Biosystems) coupled to a capillary HPLC system (Ultimate; Dionex, Sunnyvale, CA). Typical MS instrument settings were an electrospray voltage of 5.5 kV with nebulizer gas being N₂ and collision gas being N₂ as well. The LC separation was performed on a Fusa C18 column (3 μm, 0.3 × 150 mm, PepMap; Dionex) applying a gradient system starting from 95% eluent A (0.1% [v/v] HCOOH/water) and 5% eluent B (0.1% [v/v] HCOOH/acetonitrile) to 95% eluent B in 45 min at a flow rate of 5 μL min⁻¹ (sample injection volume: 1 μL). Formic acid was used because of its compatibility with MS analysis.

**Data Analysis**

Peak finding and quantification of selective ion traces was accomplished using the instrument’s AnalystQS software. Automatic raw mass data deconvolution was performed using the MetaboliteID software (Applied Biosystems). Peak lists generated by MetaboliteID were further analyzed using Excel and Visual Basic.

**CID-MS**

2-Phenylethyl glucosinolate (glucostaurin): tᵣ, 26.69 min; collision energy (CE), 20 eV; declustering potential (DP), 50 V; m/z (relative intensity, in percentages), 446.0556 (calculated for 446.0555: C₂₅H₂₃NO₈S₄Na⁺, [M + Na⁺]⁺, 10); 284.0124 (calculated for 284.0021: C₁₂H₁₀NO₅S₂Na⁺, [M + Na⁺-glucosyl]⁺, 100); 266.0101 (calculated for 265.9922: C₁₂H₁₀NO₅S⁺, [m/z 284-H₂O]⁻, 45); 186.0300 (calculated for 186.0353: C₈H₁₂NSNa⁺ [2-phenethylthio-ITC+Na⁺]⁺, 8); 8-Methylsulphinyl/loctyl isothiocyanate (hirsutin, 8-MSOOC-ITC): tᵣ, 26.54 min; CE, 30 V; DP, 50 V; m/z, 234.0993 (calculated for 234.0986: C₁₀H₁₅NO₅SNa⁺, [M – H⁺]⁻, 80); 170.0444 (calculated for 170.0397: C₉H₁₄NS⁺, [M+H⁻]⁻, 100); 161.1029 (calculated for 161.0994: C₈H₁₄OS⁺ [M+CH₂CO¹⁻], 18); 137.1201 (calculated for 137.1198: C₈H₁₄N⁺, [m/z 170]⁺, 20); 114.0407 (calculated for 114.0371: C₇H₁₄NS⁻, 18); 69.0711 (calculated for 69.0698: C₅H₁₀N⁺, 19). Kaempferol-3-O-α-L-rhamnopyranosid-7-O-α-L-rhamnopyranosid can be detected by the method described. Indole derivatives, for instance, can be clearly classified by LC-ESI-MS/MS measurements according to their substitution pattern at the ring skeleton (e.g. methoxyindoles). The same is true for the ITCs (e.g. hirsutin). The example of glucostaurin, which contains two sulfur atoms, demonstrates the usefulness of the isotopic pattern as an additional feature to support the proposed elemental composition. Arabidopsis secondary metabolites not covered by this method are mono- and sesquiterpenoids, triterpenoid alcohols, phytoestrogens, waxes, and carotenoids.
controls production of UV-protecting sunscreens in Arabidopsis. EMBO J 19: 6150-6161