Matrix-Assisted Laser Desorption/Ionization-Mass Spectrometry Imaging of Metabolites during Sorghum Germination

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biosynthesis started immediately after water uptake. Dhurrin accumulation increased rapidly within the metabolites in two of the lines. Little or no dhurrin or recycling products were present in the dry grain, but their de novo during 72 h of grain imbibition, germination and early seedling development, as well as the spatial distribution of these investigate accumulation of dhurrin, its recycling products and key general metabolites in four different sorghum lines embryo, con

laser desorption/ionization-mass spectrometry imaging demonstrated that dhurrin primarily accumulated in the germinating were mainly located in the scutellum and/or pericarp/seed coat region, suggesting unknown key functions in germination.

Dhurrin is the most abundant cyanogenic glucoside found in sorghum (Sorghum bicolor) where it plays a key role in chemical defense by releasing toxic hydrogen cyanide upon tissue disruption. Besides this well-established function, there is strong evidence that dhurrin plays additional roles, e.g. as a transport and storage form of nitrogen, released via endogenous recycling pathways. However, knowledge about how, when and why dhurrin is endogenously metabolized is limited. We combined targeted metabolite profiling with matrix-assisted laser desorption/ionization-mass spectrometry imaging to investigate accumulation of dhurrin, its recycling products and key general metabolites in four different sorghum lines during 72 h of grain imbibition, germination and early seedling development, as well as the spatial distribution of these metabolites in two of the lines. Little or no dhurrin or recycling products were present in the dry grain, but their de novo biosynthesis started immediately after water uptake. Dhurrin accumulation increased rapidly within the first 24 h in parallel with an increase in free amino acids, a key event in seed germination. The trajectories and final concentrations of dhurrin, the recycling products and free amino acids reached within the experimental period were dependent on genotype. Matrix-assisted laser desorption/ionization-mass spectrometry imaging demonstrated that dhurrin primarily accumulated in the germinating embryo, confirming its function in protecting the emerging tissue against herbivory. The dhurrin recycling products, however, were mainly located in the scutellum and/or pericarp/seed coat region, suggesting unknown key functions in germination.

Cyanogenic glucosides (CNglcs) are nitrogenous defense compounds that are widely distributed in the plant kingdom (Gleadow and Møller, 2014). They are of particular interest due to their presence in several crops, including the economically important cereals barley (Hordeum vulgare; Nielsen et al., 2002), wheat (Triticum aestivum; Erb et al., 1981; Jones, 1998), and sorghum (Sorghum bicolor; Kojima et al., 1979; Poulton, 1990), and because they often accumulate in high amounts in a wide range of plants (Kojima et al., 1979; Swain and Poulton, 1994; Crush and Caradus, 1995; Gleadow and Woodrow, 2002; Busk and Møller, 2002; Forslund et al., 2003; Jørgensen et al., 2005; Gleadow et al., 2008; Sánchez-Pérez et al., 2008). The CNglcs exert their function as defense compounds by releasing hydrogen cyanide (HCN), which is toxic to many living organisms and potentially auto-toxic to the plant (Lechtenberg, 2011). The HCN release is caused by β-glycosidase catalyzed hydrolysis of the CNglcs, initiated by tissue disruption, as for example caused by chewing herbivores (Morant et al., 2008). Besides this well-established function, CNglcs serve roles as storage and transport forms of reduced nitrogen to be reincorporated in general metabolism upon demand (Selmar, 1993; Swain and Poulton, 1994; Pičmanová et al., 2015; Nielsen et al., 2016; Del Cueto et al., 2017; Bjarnholt et al., 2018; Blomstedt et al., 2018; Schmidt et al., 2018). Although all plants possess a detoxification and nitrogen recovery pathway for HCN produced as a by-product of ethylene biosynthesis (Matilla, 2000; Piotrowski, 2008; Iqbal et al., 2013), nitrogen recovery from CNglcs proceeds via pathways that avoid release of the auto-toxic HCN. One such pathway has been elucidated (Bjarnholt et al., 2018), whereas the existence of others is documented by the coherent series of CNglc-derived metabolites identified in several cyanogenic plants. These include CNglcs decorated with extra sugars or acyl groups, or amide and carboxylic acid derivatives (Li et al., 2013; Pičmanová et al., 2015; Blomstedt et al., 2016; Del Cueto et al., 2017; Bjarnholt
Sorghum contains the Tyr derived CNGlc dhurrin and the dhurrin derived metabolites shown in Figure 1 (Pičmanová et al., 2015; Blomstedt et al., 2016; Bjarnholt et al., 2018). The biosynthetic pathway and the enzymes catalyzing dhurrin biosynthesis are well known (Gleadow and Møller, 2014), and the genes and enzymes forming a dhurrin recycling pathway have also been identified (Bjarnholt et al., 2018). This pathway proceeds via an intermediate glutathione conjugate (Fig. 1, 5) to form the end products ammonia and p-hydroxyphenylacetic acid. The latter compound exhibits hormonal activity in algae (Fries, 1976, 1977), and it is stored as the glucoside p-glucosylxyphenylacetic acid in sorghum (Fig. 1, 6; Pičmanová et al., 2015; Blomstedt et al., 2016; Bjarnholt et al., 2018). The dhurrin derived carboxylic acid p-hydroxymandelic acid has antifungal and allelopathic activity (Kope et al., 1991; Yamamoto et al., 1999; Gu et al., 2008) and is likewise stored as a glucoside (dhurrin acid; Fig. 1, 5) in high concentrations in sorghum (Pičmanová et al., 2015; Blomstedt et al., 2016). The sorghum grain contains little or no dhurrin (Tipples and Norris, 1983; Panasiuk and Bills, 1984; Velickovic et al., 2014; Nielsen et al., 2016), but dhurrin is rapidly biosynthesized at the early stages of plant development where it accumulates in high amounts to protect the newly formed tissues (Halkier and Møller, 1989; Busk and Møller, 2002). Interestingly, dhurrin recycling occurs concurrently with dhurrin de novo biosynthesis at the early stage of sorghum seedling development (Adewusi, 1990; Busk and Møller 2002; Bjarnholt et al., 2018). At that stage, nitrogen recovery may not be the main purpose of dhurrin recycling. Previous studies did not investigate if dhurrin biosynthesis and recycling takes place already during seed germination.

The seeds, formally called caryopsis, of Poacea are characterized by a typical tissue compartmentalization, which depends on their taxonomical clade. Grains of cereals such as sorghum are constituted mainly by a starch and protein filled endosperm and a dormant embryo composed of the embryonic axis and scutellum (Fig. 2; Dante et al., 2014). Seed germination and initiation of central metabolism is induced by imbibition and hormone-controlled mobilization of resources from the endosperm storage tissue (Yamaguchi, 2008; Sponsel, 2016). Hydrolytic enzymes produced in the aleurone layer surrounding the endosperm are released to degrade the storage polymers. The released general metabolites are then transported via the scutellum into the developing embryo as a source of energy and building blocks (Dante et al., 2014; Rosental et al., 2014) ensuring rapid growth of the vulnerable young seedling. Channeling of resources toward formation of specialized metabolites would thus entail that these play a role in germination or early growth of the seedling. In germinating barley seeds, hordatines have been found to accumulate in the embryonic axis, presumably for the purpose of defending the tissue against pests and diseases at this vulnerable growth stage (Gorzolka et al., 2016). Plant hormones and specialized metabolites are biosynthetically and structurally closely related, and several specialized metabolites are suspected of playing a role in hormonal signaling. The production of hydrolytic enzymes excreted by the aleurone and scutellum layers is triggered by gibberellins, hormones produced in the scutellum and transported to the aleurone cells (reviewed by Yamaguchi, 2008; Sponsel, 2016). In wheat for example, the phenolic specialized metabolite coumarin has been shown to induce production of hydrolytic enzymes in aleurone cells (Saleh and Abu El-Soud, 2015). Tissue-wise localization of compounds during early germination and seedling development may, therefore, lend clues to their significance and biological activity.

In this study, we investigate the spatial distribution of dhurrin and its derivatives in the germinating sorghum grain using mass spectrometry imaging (MSI). This technique enables determination of the in situ localization of metabolites based on their molecular masses. A number of different methods are available, but the most popular is matrix assisted laser desorption ionization (MALDI)-MSI (Bjarnholt et al., 2014; Boughton et al., 2016). This method allows the detection of a wide variety of metabolites and offers the possibility of very high spatial resolution (so far demonstrated down to 5 μm in plant tissue analyses; Bjarnholt et al., 2014; Korte et al., 2015; Boughton et al., 2016). Coupling the MALDI ion source with mass analyzers that provide high mass accuracy (<2 ppm), such as the Orbitrap, provides...
identification of compounds based on their accurate mass. Here, we applied highly sensitive liquid chromatography (LC)-triple quadrupole mass spectrometry (MS) analyses in combination with MALDI-Orbitrap-MSI to study the emergence and spatial distribution of dhurrin and its derivatives compared with selected general metabolites. To gain more insight into the significance of dhurrin biosynthesis and its endogenous recycling pathways, we monitored the changes in accumulation of dhurrin and its derived metabolites during the early stages of the germination process, from the dry seed until the protrusion and early growth of the coleoptile where dhurrin biosynthesis is at its highest rate (Busk and Møller, 2002), in four varieties of sorghum, one of which lacked the capacity to synthesize dhurrin (Blomstedt et al., 2012).

RESULTS

To explore the formation and recycling of dhurrin in the course of grain imbibition and germination, four different genotypes of sorghum were analyzed. The wild types used were a commercial breeding line (wild-type parent; Blomstedt et al., 2012) and the cultivar BTx623 for which the genome sequence has been reported (www.phytozome.org). The mutant line tcd1 (totally cyanide deficient) obtained by ethyl methanesulfonate mutagenesis does not produce dhurrin because it carries a mutation in CYP79A1 rendering it inactive (Blomstedt et al., 2012). The tcd1 line was generated in the commercial breeding line (Blomstedt et al., 2012) and has now been backcrossed twice into the cv BTx623. This was included as a unique reference system to confirm the identities of dhurrin and its derivatives detected in the wild type, and to explore their metabolic relationship and possible significance in the germination process. A sibling from the last tcd1 × BTx623 cross named TCD1, which does not contain the mutation, was analyzed as an additional control. Grains and seedlings from all four genotypes were collected at specific time points (Fig. 2A) following imbibition and germination in the dark. All samples were subjected to quantitative metabolite analysis by
LC-MS. Selected cv BTx623 and tcd1 samples were analyzed by MALDI-MSI to monitor the spatio-temporal developments in accumulation of interesting analytes at four stages of germination (Fig. 2, B and C). The grains were imbibed for 12 h, and the levels of dhurrin and its recycling products were quantified in the whole-grain extracts at 0, 12, 24, 36, 48, 60, and 72 h after imbibition (AI), as well as in samples from the mature, unimbibed grain.

Figure 2. Sorghum germination process; embryo and seedling development and sampling time points. A, Timeline of imbibition, germination, and collection of samples. Sorghum grain were imbibed in water for 12 h and germinated for 72 h. Samples were collected every 12 h at the desired time points, as indicated with arrows. All eight time points were subjected to metabolite profiling. From the eight time points, four developmental stages (S) were selected for analysis by MSI and named: S0, dry seed; S1, emerged radicle and coleoptile; and S2, developed shoot and primary root. B, Graphical representation of the stages collected for MSI analyses. For the cvBTx623 line these stages matched the indicated time points (for tcd1 see Fig. 3). C, Schematic of longitudinal sections of the four developmental stages selected for MSI analysis, showing detailed morphological features and main compartments.

Accumulation of Dhurrin and its Recycling Products during Germination and Seedling Formation

Analyses of the dry unimbibed grain of the three dhurrin producing sorghum lines confirmed the previous observation from other genotypes that dhurrin is absent in the dry grain (Tipples and Norris, 1983; Panasiuk and Bills, 1984; Velickovic et al., 2014; Nielsen et al., 2016). Previously, the earliest reported presence of dhurrin was in shoots after a total of 44 h of imbibition and germination in the dark (Halkier and Møller, 1989). In this study of germinating grain and the germinated grain including emerging shoots and roots, dhurrin reached detectable levels already at 0 h AI in cv BTx623 and at 12 and 24 h AI, respectively, in the TCD1 and wild-type lines (Fig. 3, A–C). Dhurrin levels were generally found to be higher in cv BTx623 compared with other dhurrin producing lines and continued to increase in cv BTx623 at the end of the experiment, whereas it was more stagnant in the two other lines. At the end of the experiment, at 72 h AI, the dhurrin concentration in cv BTx623 and TCD1 was ~1.3 nmol mg⁻¹ (SE = 0.08) and 1.1 nmol mg⁻¹ (SE = 0.04), respectively. In the tcd1 parent line dhurrin content peaked at 48 h AI (0.9 nmol mg⁻¹, SE = 0.08).

The targeted analysis also documented the presence of previously reported dhurrin derivatives, namely the dhurrin glucosides (Fig. 1, 4), dhurrin acid (Fig. 1, 3), p-hydroxyphenyl(S-glutathione)acetonitrile (GS-pOH-PACN; Fig. 1, 5), and p-glucosyloxyphenylacetic acid (pGPAAc; Fig. 1, 6). Other products and intermediates of the proposed recycling pathways were not detected (Fig. 1, 2, 7, and 8). With a few specific exceptions discussed below, the accumulation patterns of the detected metabolites matched their expected formation from dhurrin, i.e. in most cases they were present in detectable levels at one or two time points after the appearance of dhurrin, and the concentrations were either
increasing or leveling out toward the end of the experiment, similar to those of dhurrin.

An overall trend of proportionality between the level of dhurrin and the recycling products was observed, i.e. the \( tcd1 \) parent samples exhibited the lowest levels of dhurrin and also of recycling products compared with the other wild-type lines. Average values and SEs for all compounds/time points and lines are presented in the Supplemental Table S1. The dhurrin derivative present at the highest concentration was \( \text{p} \text{GPAAc} \). The timing of the appearance of \( \text{p} \text{GPAAc} \) was slightly shifted between the lines, and in the \( tcd1 \) parent line the concentration of \( \text{p} \text{GPAAc} \) was already at its maximum level at the first time point where dhurrin was detected (Fig. 3C). After the initial increase in concentration all dhurrin-producing lines maintained a stable level at 0.3 nmol mg\(^{-1}\) throughout the experimental period. The same accumulation pattern was observed for GS-\( \text{p} \text{OHPACN} \), although the concentration of this compound was much lower reaching 0.006, 0.002, and 0.0011 nmol mg\(^{-1}\) in cv BTx623, TCD1, and \( tcd1 \) parent line, respectively (Supplemental Fig. S1). Such a pattern is in accordance with the position of GS-\( \text{p} \text{OHPACN} \) as an intermediate in the recycling pathway of dhurrin to \( \text{p} \text{GPAAc} \) (Bjarnholt et al., 2018). In all three wild types, dhurrin acid was present as the second highest accumulated recycling product, reaching \( \approx 0.1, 0.06, \) and 0.03 nmol mg\(^{-1}\) in cv BTx623, TCD1, and parent, respectively. Although at barely detectable levels (between 0.005 and 0.01 nmol mg\(^{-1}\)), dhurrin acid was also detected in dry mature seeds. Strikingly, the mutant line \( tcd1 \) accumulated \( \text{p} \text{GPAAc} \) to roughly the same level as the wild-type and sibling lines (Fig. 3D), but with a distinct time delay of approximately 24 h. This was unexpected, as \( tcd1 \) in the original parent background was previously reported to not accumulate \( \text{p} \text{GPAAc} \) during plant development (from 7 to 46 d after sowing; Blomstedt et al., 2016; Bjarnholt et al., 2018). In this study, we did detect trace amounts of dhurrin in one replicate from each of two growth stages, indicating that it may in fact be produced in the mutant line at levels below the detection limit. However, at this stage it cannot be concluded whether dhurrin was actually present and could be a precursor for the detected \( \text{p} \text{GPAAc} \).

**Accumulation of Free Amino Acids**

To investigate the timing of the onset of dhurrin biosynthesis in more detail, we compared it with one of
the very early events in general metabolism during seed germination: the increase in concentration of free amino acids, caused by hydrolysis of storage proteins or de novo biosynthesis, to feed the need of the developing embryo. Dhurrin is biosynthesized from the amino acid Tyr, and furthermore, the recycling of dhurrin into embryo. Dhurrin is biosynthesized from the amino acid novo biosynthesis, to feed the need of the developing acids, caused by hydrolysis of storage proteins or de germination: the increase in concentration of free amino acids. Interestingly, the free amino acid levels were different in tcd1 compared with its sibling TCD1, as illustrated for the total sum in Figure 4. The tcd1 line seemed to be delayed with respect to amino acids accumulation until ~48 h AI, after which the levels were often higher in tcd1 compared with TCD1 and consequently also the parent line. With only three biological replicates to measure changes in such a dynamic pool of metabolites undergoing rapid change, the SEs were relatively high. However, Gln, Pro, and Tyr stood out, as the apparent levels of these amino acids reached higher or similar levels in tcd1 than in cv BTx623 at one or more of the time points 48 to 72 h AI (Supplemental Fig. S3). Almost all other amino acids were higher in cv BTx623 than in all other lines at most time points, with the most pronounced differences being found for Asp, Asn, His, Ile, Met, Ser, Thr, and Val. Levels of Ala, Arg, and Lys appeared somewhat unaffected or differently affected by the genetic background, and the remaining amino acids followed the trends of the total sum, but with moderate differences between the lines (Supplemental Fig. S3).

### Spatial Distribution of General and Specialized Metabolites

The stages selected for MSI analyses were chosen by a combination of the metabolite analyses and development of particular anatomical features. The four stages were (1) S0, the dry grain representing a base line; (2) S1, the imbibed grain, which was the first time point where dhurrin was detected in the wild type; (3) S2, where the radicle and early shoot emerged and dhurrin recycling products were first detected; and (4) S3, the stage where the primary root was fully developed and the coleoptile, containing the primary leaf, completely protruded from the embryonic axis of the embryo. For cv BTx623 these stages corresponded to the time points shown in Figure 2. The development of tcd1 was slightly delayed compared with cv BTx623, and therefore the seeds selected for MSI analysis from these lines were chosen similar to cv BTx623 until 48 h AI. After 48 h, the concentrations of free amino acids in the parent and TCD1 lines leveled out or even decreased, whereas they were still increasing in the cv BTx623 line. These trajectories and line-dependent differences completely matched the dhurrin trajectories (Fig. 3). Differences in concentrations of free amino acids in germinating seeds were previously shown to depend on the genetic background (Feenstra et al., 2017). As previously mentioned, the tcd1 line was originally generated in the genetic background of a near isogenic commercial breeding line, and the tcd1 and TCD1 lines used in the current study are the F2 generation of the tcd1×BTx623 cross. Consequently, these two lines have a mixed parent×BTx623 genetic background, which is presumably what is reflected by the somewhat intermediate position of TCD1 between cv BTx623 and the mutant parent line with regard to concentrations of dhurrin and free amino acids.
based on the developmental stage rather than time (time points shown in Fig. 3D). After the first 36 h AI the mutant line was delayed by ~12 h compared to the wild type. At this stage the radicle and the coleoptile containing the primary leaves were fully emerged in cv BTx623, whereas in tcd1 that was delayed until 48 h AI. For each stage and line, three seeds (including roots and seedlings when present) were sectioned longitudinally at 10 μm thickness, placed on precooled glass slides, and immediately freeze-dried, using the method reported by Kawamoto and Kawamoto (2014), with the few modifications reported by Gorzolka et al. (2016), Boughton and Thinagaran (2018), Sarabia et al. (2018), and Schmidt et al. (2018). Three consecutive sections were selected and subjected to respectively fluorescence microscopy and MSI analyses in both positive and negative mode to allow detection of an extensive range of metabolites.

Dhurrin and its recycling products and a number of general metabolites were detected in positive mode using 2,5-dihydroxyxybenzoic acid (DHB) as matrix. Amino acids were detected in negative mode using 1,5-diaminonaphthalene (DAN) as matrix on a different section. To minimize the crystal size and obtain uniform matrix deposition and thereby minimize delocalization of analytes, sublimation was used for matrix deposition (Hankin et al., 2007). The sections were analyzed with 30 μm pixel size, which provided sufficient spatial resolution for matching the morphological features observed in the fluorescence microscopy images while maintaining a reasonable acquisition time (~4–6 h per sample). During data processing to produce the images, each ion was normalized to the total ion count (TIC) in each pixel, making up for some of the variation between analyses. Many factors, including sample handling, matrix deposition, and plant-tissue background can cause differences in ionization efficiencies for individual metabolites across the sample. Potentially this may result in images of metabolite distributions that do not provide a true representation of the sample constituents. To validate our results, we therefore compared with previous data obtained using different approaches.

During the germination process, breakdown of the starchy endosperm into mono- and disaccharides provides an important source of energy for the developing embryo (Lopes and Larkins, 1993; Aoki et al., 2006; Leonova et al., 2010; Sánchez-Linares et al., 2012; Yan et al., 2014; Feenstra et al., 2017). Quantification of mono- and disaccharides of microdissected germinating cereal seeds from maize (Zea mays) and oat (Avena sativa) showed that monohexose (Hex) and dihexose (Hex2) accumulate in the embryo and scutellum (Leonova et al., 2010; Feenstra et al., 2017). The same distribution was observed in the MALDI-MSI analyses of the germinating sorghum seeds (Fig. 5A). Hex2 was less abundant in unimbibed samples, but its signal greatly increased in the embryo and scutellum already after imbibition (S1). As the germination proceeded and the axis protruded, Hex2 was concentrated more extensively in the emerging radicle and coleoptile (S2; Fig. 5A). Accumulation of Hex occurred after the protrusion of the radicle from the embryo (S1), where it was mainly found in the differentiation/elongation zone of the emerging radicle. At the last stage analyzed (S2), the Hex signal was highly concentrated in the coleoptile and slightly less abundant in the radicle. At the S3 stage, Hex and Hex2 were detected in the pericarp/aleurone layers (the two layers could not be distinguished with the lateral resolution applied in the MSI analyses). This is consistent with observation of MALDI-MSI in germinating maize kernels (Feenstra et al., 2017).

It is important to note that the compounds are detected based on their accurate mass and corresponding sum formulae. In the scutellum and developing embryo Hex and Hex2 are the major general metabolites Glc and Suc, as demonstrated in the microdissection experiments (Leonova et al., 2010; Feenstra et al., 2017); however, it is possible that Hex and Hex2 in the layer(s) encasing the seeds are different saccharides. For Hex and Hex2, the potassium adducts \([\text{C}_6\text{H}_{12}\text{O}_6\text{K}]^+\) and \([\text{C}_{12}\text{H}_{22}\text{O}_{11}\text{K}]^+\) gave the strongest signals in positive mode and were therefore used to map the distribution of these compounds. This was in accordance with the maize kernel study using a similar MALDI-MSI method (Feenstra et al., 2017), which has also been shown to provide a good signal quality from Hex2 in negative mode (Korte et al., 2015).

Based on this knowledge, the efficiency and consistency of our ionization methods were compared by analyzing the distribution of Hex and Hex2 as deprotonated species \([\text{C}_6\text{H}_{12}\text{O}_6\text{H}]^-\) and \([\text{C}_{12}\text{H}_{22}\text{O}_{11}\text{H}]^-\) detected in negative mode (Fig. 5A). At all stages analyzed, the distribution of \([\text{Hex}^+\text{K}]^+\) and \([\text{Hex}^2\text{K}]^+\) versus \([\text{Hex}^-\text{H}]^-\) and \([\text{Hex}^2\text{H}]^-\) were as identical as can be expected for different sections, demonstrating the efficiency of both ionization methods.

Choline is an important plant constituent as a precursor in the formation of many metabolites including acetylcholine and phosphatidylcholine (PC; Miura and Shin, 1984). In chick pea (Cicer arietinum) seedlings, choline is highly abundant in the cotyledon and growing part of the embryo including the radicle (Ahmad and Karim, 1951), whereas PC is preferentially stored in the endosperm together with many other seed oil reserves (Fincher, 1989; Leonova et al., 2010). The same tissue-specific distributions were demonstrated for germinating barley seeds using MALDI analysis, showing strong accumulation of PC in the endosperm, whereas choline was detected in the scutellum + embryo and the aleurone layer (Gorzolka et al., 2016). Similar to Gorzolka et al. (2016), we detected choline as protonated species and PC (16:0) as a potassium adduct \([\text{C}_6\text{H}_{13}\text{NO}_3\text{K}]^+\) and \([\text{C}_{23}\text{H}_{43}\text{NO}_7\text{K}]^+\) in positive mode. These analyses demonstrated that our MALDI-MSI method afforded efficient ionization across all examined germination stages and tissues, including the endosperm (Fig. 5B). Nearly identical results were obtained using the tcd1 mutant line (Supplemental Fig. S5), demonstrating similar ionization
efficiencies and basic metabolite distributions in the two genotypes and corroborating the robustness of the methods.

Localization of Dhurrin and its Recycling Products during Germination and Seedling Formation

Dhurrin and its recycling products were detected as potassium or sodium adducts in positive mode. The ions providing the most intense signal were selected for generation of the images. Dhurrin was most consistently detected as a potassium adduct ([C$_{14}$H$_{17}$NO$_7$+K]$^+$), and only in a few individual replicates also detected as its sodium adduct with much lower signal intensity. Therefore, the potassium adduct was used to generate the images. Dhurrin glucosides were the only recycling products detected as a potassium adduct [M+K]$^+$, whereas dhurrin acid, GS-pOHPACN, and pGPAAc were exclusively detected as their sodium adducts ([M+Na]$^+$). Supplemental Table S3 provides a list of accurate masses applied to generate the images. The intensities varied between replicates, but the distributions were the same. The most clear results were selected for presentation in Figure 6, and the replicate results can be viewed in Supplemental Figure S7. In the metabolic extract analysis, dhurrin was first detected after imbibition (S$_0$) in the BTx623 line, and the recycling products were first detected after axis protrusion (S$_1$). In the MSI analyses, these compounds were all detected in the dry seeds as well as after imbibition (Fig. 3; Supplemental Fig. S7, S$_0$ + S$_1$), but the signal intensities were minute at these early stages, and the compounds not detected in all replicates. This explains why the very same constituents

Figure 5. Distribution and compartmentalization of general metabolites during germination and early seedling development of the cv BTx623 wild-type line. Three consecutive cryo-sections were imaged: The optical images on the far left (cv BTx623) were acquired with fluorescence microscopy, and the remaining images using MALDI-MSI in both negative and positive ion mode, using respectively DAN and DHB as matrix at 30 μm spatial resolution. Developmental stages are as in Figure 2. The dashed lines define the outlines of the recorded MS images. A, Comparative distribution of hexose and dihexose using the two ionization modes. B, Specific tissue compartmentalization of choline [M+H]$^+$ and phosphocholine (PC 0:16) [M+K]$^+$. Scale bar = 1 mm. Signals for all MSI images are normalized to TIC on each pixel, and maximum values for generating images are listed in Supplemental Figure S4.
were not detected in the total extracts that were analyzed by LC-MS, due to the dilution effect. The weak signals make it difficult to assess the exact localization of the compounds, but it appears that dhurrin and its glucosides were present in both scutellum and embryo of the dry seeds. The three dhurrin glucosides have the same sum formula and thus identical exact mass and could therefore not be differentiated in the MSI analysis. The distribution of the ions corresponding to the dhurrin glucosides consistently followed the same distribution pattern as dhurrin until the end of the experiment. On the other hand, the dhurrin recycling products dhurrin acid and pGPAAc mainly accumulated in the scutellum and other tissues enclosing the embryo. After water uptake, this difference was much more pronounced than at S0, as the signal for dhurrin and the dhurrin glucosides were clearly much more intense in the embryo than in the scutellum, whereas the recycling products seemed to be almost exclusively located in the scutellum (Fig. 6, S). As the germination proceeds, the radicle is the first part of the seedling to visibly emerge from the seed coat. Subsequently, the coleoptile containing the plumule (primordial leaf) protrudes from the embryonic axis, and this marks the end of germination. In cv BTx623, this
developmental stage, S1, was reached at 24 h AI. At this stage the dhurrin and dhurrin glucosides appeared to be more concentrated in the differentiation/elongation zone of the emerging radicle and in the area protecting the plumule. The remaining compounds were still mainly found in the scutellum.

At the last time point, S2, all compounds were exclusively detected in the newly formed root and coleoptile tissue. In particular, dhurrin was detected at very high intensity in the tip of the fully emerged coleoptile, consistent with a previous study (Halkier and Møller, 1989). The dhurrin derivatives were weakly detected at this stage and there were variations between replicates (Supplemental Fig. S7), making it difficult to determine whether the compounds were uniformly distributed or not. The strongest signal was found for the dhurrin glucosides, whereas dhurrin acid, pGPAAc and GS-pOHPACN were barely detectable. The distribution of GS-pOHPACN diverged from the remaining compounds, as it was generally more broadly distributed. At S0 it was detected at highest intensity in the embryo, whereas at S1 it was equally intense in embryo and scutellum. In addition, in some replicates it was detected in the pericarp/aleurone layer at S0, S1, and possibly S2. As such, GS-pOHPACN and pGPAAc, which are respectively intermediate and product of the same metabolic pathway, displayed only partially overlapping distributions.

At all imaged stages, MSI signals corresponding to dhurrin and its recycling products were absent in the tcd1 mutant, at the applied mass tolerances (Supplemental Table S3) and default abundance threshold of MSIReader (Robichaud et al., 2013). This is visualized for dhurrin in Supplemental Figure S6A, showing average mass spectra of cv BTx623 and tcd1 at S1. This is difficult to visualize for earlier stages, where dhurrin and its derivatives were only detected in scattered pixels in cv BTx623 samples, making the average spectra uninformative. For dhurrin the most abundant isotope of the [M+K]+ ion is that containing a single 13C, which has a theoretical abundance of 15.1%. In the three pixels with highest dhurrin abundance in the S0 sample shown in Supplemental Figure S7, we were able to detect this isotope (mass-to-charge ratio [m/z] 351.0670) at, respectively, 8%, 9%, and 15% of the monoisotopic [dhurrin+K]+ peak (Supplemental Fig. S6C). Given the very low absolute peak intensities, which affects the accuracy of the measured ratios, and that this m/z value was absent from other pixels and tcd1 samples, this further supports the presence of low quantities of dhurrin and its derivatives in the BTx623 line at S0 and S1. The second most abundant isotope is [dhurrin+41K]++; however, at a theoretical abundance of 7.2%, this m/z value was not detectable at the S0 and S1 stages. At S2 and S3, the MSI findings were corroborated by the LC-MS based metabolite analyses of the tcd1 mutant (Fig. 3D). Taken together, the results support that the detected ions in BTx623 do in fact correspond to the proposed compounds. pGPAAc was detected at later time points in the LC-MS analyses of tcd1, but unfortunately the size of the protruded axis (root and coleoptile) at those stages of seedling growth caused embedding and sectioning to be impossible and therefore the distribution of this compound could not be explored by MSI.

Distribution of Amino Acids

In the germinating maize kernels, only three proteinaceous amino acids were reliably detected by MALDI-MSI (Feenstra et al., 2017). In our study, all amino acids except Ala and Gly were readily detected as deprotonated species in negative mode [M-H]-. However, the relative signal intensities in the MS images did not completely reflect the concentrations determined in the LC-MS analyses. This serves to emphasize the well-known fact that the ionization efficiency in MS imaging depends on the type of applied matrix, the tissue matrix background, and in particular the physicochemical properties of the analytes. Figure 7 shows results for a selected subset of amino acids with a full data set provided in the supplementary data (Supplemental Figs. S8 and S9). Although MSI is not a quantitative technique, the overall trends in intensity of amino acid ions in images from samples of increasing age did match the overall trends found in the analysis of free amino acids in extracts. In cv BTx623 as well as in tcd1, there was a sharp increase in signal intensity from S0 to S1 for most amino acid ions, reflecting the sharp increase in amino acid concentration detected in extracts of whole samples at 24 to 36 h AI. Asp, Glu, and Asn were present in high concentrations at all stages, and genotypes analyzed by LC-MS and these amino acids were also detected at all stages in the MSI analyses of cv BTx623 and tcd1 (Fig. 7; Supplemental Figs. S8 and S9). In general, small, neutral amino acids such as Val, Met, Leu/Ile (which have identical sum formulae and cannot be differentiated in the MSI analyses), Ser, Pro, and Thr were only clearly detectable by MSI at the later growth stages, regardless of their concentration in the extract analysis. Conversely, Lys and Trp were found in low concentrations at all growth stages, but were nevertheless in some cases detectable in the MSI analysis already at stage S0.

No free amino acids were detected in the endosperm, indicating that the rate of hydrolytic release of free amino acids from the storage proteins was lower than the transport of the amino acids toward the developing tissues. At the active growth stages from S0 onward, where the amino acids are also synthesized de novo, the presence of amino acids was detected both in the scutellum and embryo. At these early stages of germination, the free amino acids are required for biosynthesis of protein and metabolites, presumably in both tissues, but eventually only the embryo will be active. This was reflected by the MSI results with a tendency toward the scutellar amino acid signals decreasing and the embryonic axis localized amino acid signals increasing from S0 to S2. Asp, Glu, Gln, and Trp were in some cases also detected in the...
aleurone/pericarp layer at stage $S_0$ and/or $S_1$. The inconsistency in detection between stages (Supplemental Fig. S8) and replicates (Supplemental Fig. S9) may be exacerbated by the low concentrations present. At the last imaged growth stage, some amino acids displayed distinct localizations in the newly formed tissues. For example, the highly abundant Glu was almost detected everywhere, whereas Lys was mainly detected in the elongation zones of both root and shoot where Trp, on the other hand, was specifically not detected (Fig. 7). We observed no striking differences for $tcd1$ compared with cv BTx623, as illustrated for a subset of amino acids in Figure 7B and Supplemental Figure S9.

DISCUSSION

The spatio-temporal investigation of the accumulation of dhurrin and its recycling products in the germinating sorghum grain demonstrates that initiation of dhurrin biosynthesis and even the endogenous metabolism of dhurrin are very early events in seed germination and seedling development, initiated already during imbibition. The in situ localization of dhurrin metabolites showed that dhurrin and related cyanogenic di-glucosides accumulate in the newly formed tissues, protecting the vulnerable developing embryo. On the other hand, dhurrin derivatives accumulate mainly in the scutellum or are distributed across embryo, scutellum, and aleurone/pericarp layer, indicating key functions in the germination process.

Imbibition of the dry seeds is the starting point of germination. In the model plant Arabidopsis ($Arabidopsis thaliana$) metabolite abundances undergo substantial changes after the first few hours of imbibition (Rajjou and Debeaujon, 2008; Howell et al., 2009). The metabolic activity generates the essential building blocks for plant growth, energy production, and
protein biosynthesis. In accordance with previous studies, we found only low concentrations of free amino acids in the dry sorghum grain (Bewley et al., 2013; Feenstra et al., 2017). The concentration of total and most of the individual free amino acids increased slowly at the first few time points after imbibition, and then increased dramatically at 24 h AI in cv BTx623 and slightly later or less steeply in the other control lines (Fig. 4; Supplemental Fig. S3). The trajectories of dhurrin concentration were exactly the same (Fig. 3), showing that dhurrin biosynthesis was switched on as early as the key event of amino acids release from the endosperm storage proteins. This suggests that dhurrin is of primary importance in the early phase of germination. Dhurrin localization at the tissue level as determined by MALDI-MSI furthermore points toward the presumed primary functionality of the metabolite. The presence of dhurrin in the differentiation/elongation zone of the emerging radicle and in the area protecting the plumule confirms the position of dhurrin as a key player in the sorghum defense system. Specialized metabolites that provide defense against general biotic stresses are commonly found in peripheral or key tissues where they can provide most benefit to the plant as a first line of chemical defense (Bellotti and Riis, 1994; Shroff et al., 2008; Ha et al., 2012; Bøgeskov Schmidt et al., 2018). In germinating barley, the antifungal hordatines were found to accumulate predominantly in the outer layer of the shoot (Gorzolka et al., 2016), similar to dhurrin accumulation in the outer layer of the cotyledon. However, the concurrent onset of the dhurrin recycling processes early in the development is indicative of other roles in the germinating grain.

The dhurrin glucosides were detected in the metabolite extracts as soon as dhurrin levels increased at 24 h AI. In the MS images of cv BTx623, they were colocalized with dhurrin. The biological functions of these di-glucosides are unknown, but as their aglycone is the dhurrin cyanohydrin, they will release HCN following possible hydrolysis resulting in the removal of both Glc residues. The cyanogenic di-glucosides found in cassava (Manihot esculenta) and the rubber plant (Hevea brasiliensis) have been proposed to be transport forms protected against cleavage by the monoglucosidase that hydrolyses the cyanogenic monoglucosides (Selmar, 1993). By contrast, in almonds (Prunus dulcis, syn. Prunus amygdalus), the cyanogenic di-glucoside amygdalin accumulates in the kernels and is responsible for
the toxic cyanide release in the bitter seeds following the serial action of two glucosidases (Sánchez-Pérez et al., 2008, 2009, 2012; Del Cueto et al., 2017).

Dhurrin-6-glucoside was originally isolated from guttation droplets of sorghum leaves (Selmar et al., 1996), indicating a role in transport for this compound. The dhurrin glucosides may serve different functions in different tissues, e.g. transport forms in aerial tissues and defense purposes in roots where dhurrin is lowly abundant or absent. Determination of the distribution of the individual di-glucosides would require dissection and extract analysis where the isobaric compounds can be differentiated by retention time and their linkage types be determined by ion mobility MS (Pičmanová et al., 2015) or by NMR spectrometry of the isolated compounds.

The remaining dhurrin recycling products detected in the extracts of germinating seeds from the dhurrin producing lines either emerged at the same time as the dhurrin concentration increased (pGPAAc) or shortly after (GS-\textit{pOHPCN}), or were detected at all time points, but found to increase shortly after the increase in dhurrin concentration (dhurrin acid). This early emergence or increase likewise indicate important roles of these compounds at the early stages of plant life. In fact, all derivatives were detected already at S1 in the MS images, supporting their importance for the germinating seed or early seedling formation. Despite their apparent metabolic relation to dhurrin, the two major derivatives, dhurrin acid and pGPAAc, were apparently absent from the growing tissues until after germination at S2. This would suggest that before this stage, supply of reduced nitrogen to the growing tissues is not the main purpose of the pathways for conversion of dhurrin into these acids. The presence of dhurrin acid in the dry seed and its localization around the embryonic axis after imbibition at S1 supports that it functions as a storage form of an antifungal agent (Kope et al., 1991), protecting the seed and the early embryo stage. In cassava, the main cyanogenic glucoside is linamarin, and its dhurrin acid analog, linamarin acid, was found to be localized to the stem and in the vascular tissue of the cassava tuber (Bøgeskov Schmidt et al., 2018). The presence in the vascular tissue proposes that the glucosylated CNGlc-derived carboxylic acids may be transport forms of their potentially bioactive aglucones.

\textit{p}-Hydroxyphenylacetic acid, the aglucone of pGPAAc, has been shown to exert growth promoting effects and auxin-like activities in algae (Fries, 1976, 1977). The compound has been isolated from plants, algae, and associated microbes, and its nonhydroxylated analog phenylacetic acid is well-established as an auxin-like plant hormone (Koepfl et al., 1938; Ferro et al., 2010; Simon and Petrásék, 2011). Glucosylation is a well-known mechanism used to control auxin homeostasis (Korasick et al., 2013) and the activity of gibberellins, both key hormones controlling seed germination (Yamaguchi, 2008). The observed localization of pGPAAc in the scutellum, the tissue that is considered the main source of gibberellin production in germinating seeds, renders it tempting to suggest that \textit{p}-hydroxyphenylacetic acid has an equally important role in sorghum seed germination. Assignment of an important function to \textit{p}-hydroxyphenylacetic acid is supported by the fact that the \textit{tcd1} mutant was shown to produce pGPAAc, which after germination accumulated at wild-type levels (Fig. 3D). The presence of pGPAAc in \textit{tcd1} was surprising in the light of previous reports where pGPAAc was found to be absent or present in trace amounts in vegetative tissues of \textit{tcd1} (Bjarnholt et al., 2018) and a second mutated line, \textit{tcd2}, which is mutated in the last enzyme of dhurrin biosynthesis (Blomstedt et al., 2016). In the present study, we did detect traces of dhurrin in a few replicates of \textit{tcd1} extracts, and this may be what fuels the production of pGPAAc in the germinating seeds and/or young seedling tissues of this line, with the low dhurrin supply explaining the delayed accumulation. On the other hand, the eventual accumulation of wild-type levels of pGPAAc suggests that this compound is produced from a completely different pathway, which remains to be elucidated.

GS-\textit{pOHPCN} is an intermediate in the production of pGPAAc from dhurrin (Bjarnholt et al., 2018). As such, it was convincing to find it localized to both embryo and scutellum, where its precursor, dhurrin, and its downstream product, pGPAAc, accumulated. At S1, the accumulation of precursor and product appeared almost completely separated between the two tissues. The presence of GS-\textit{pOHPCN} in both tissues could therefore suggest that it is produced from dhurrin in the embryo and subsequently transported to the scutellum for further processing into pGPAAc. Compartmentalization between the different tissues of germinating seeds has also been suggested for the gibberellin biosynthetic pathway in Arabidopsis (Urbanova and Leubner-Metzger, 2016). However, this does not explain the presence of GS-\textit{pOHPCN} in the aleurone/pericarp layer. In the aleurone tissue, the small molecules present can be expected to be mainly signaling compounds or products resulting from the cell wall degradation that happens during germination. It can be hypothesized that GS-\textit{pOHPCN} is a transport molecule of the bio-active nonglucosylated \textit{p}-hydroxyphenylacetic acid or even of free \textit{pOHPCN} (\textit{p}-hydroxyphenylacetonitrile), which may have an undescribed bio-activity. \textit{p}-Hydroxyphenylacetic acid was not detected in the extract analyses, maybe because the locally produced and aleurone localized \textit{p}-hydroxyphenylacetic acid were diluted too much in the total extracts. Conversion of GS-\textit{pOHPCN} to \textit{p}-hydroxyphenylacetic acid requires the action of one of the two lambda GSTs (glutathione S-transferases) \textit{SbGSTL1} or \textit{SbGSTL2}, followed by the nitrilase hetero-dimer \textit{SbNIT4A/B2}. The transcripts of these enzymes colocalized at the cellular level in sorghum seedling leaves (Bjarnholt et al., 2018) and were coexpressed in developing sorghum seeds (Nielsen et al., 2016). Transcriptome analyses of the individual tissues in the
germinating sorghum seed, such as recently carried out for barley (Betts et al., 2017), could help identify source and sink tissues of GS-pOHPACN and lend further clues as to the nature of its biological function.

The overall pattern of free amino acid accumulation showed that the siblings tcd1 and TCD1 were somewhat intermediate between their parent wild-type lines. The tcd1 mutant displayed a delay in accumulation of free amino acids compared with its sibling. This delay was mirrored by the delay in germination in tcd1, as germination of TCD1 and the parent line were not significantly different from cv BTx623. Amino acids have been shown to influence the germination process. In Arabidopsis, de novo biosynthesis of Met was described to be of primary importance for seed germination and seedling growth (Gallardo et al., 2002). In other plants, such as barley and sugar beet (Beta vulgaris var. altissima), enzymes related to Met metabolic pathways dramatically increase in the first hours after imbibition before radical emergence (Soeda et al., 2005; Catusse et al., 2008). In cv BTx623 and TCD1, the concentration of Met increased much more steeply than in tcd1, in which comparable levels were reached with a delay of 24 to 36 h. It is possible that an imbalance in the free amino acids caused by possible overaccumulation of Tyr in the absence of the dhurrin biosynthesis could influence the germination rate. Indeed, Tyr as well as Pro and Gln accumulated to higher levels in tcd1 compared with its sibling and even cv BTx623 after germination. Like other aromatic amino acids, Tyr is produced in the plastids and exported into the cytosol as precursor for synthesis of other metabolites. Degradation of Tyr via a pathway initiated by Tyr amino transferase (Tyr-AT) and transamination of α-ketoglutarate into Glu (Hildebrandt et al., 2015; Schenck and Maeda, 2018) avoids the risk of overaccumulation. Glu did not accumulate in the tcd1 mutant relative to the wild type, but as Glu is a precursor for Pro and Gln, it is likely that the observed increase in levels of the latter two in tcd1 is a consequence of Tyr overaccumulation and degradation. Tyr may also be directed into synthesis of other metabolites such as tyramine, tocopherols, Vitamin E, and numerous phenolic compounds such as the flavonoids. We did not target these compounds in our analyses, but it is also possible that an imbalance in accumulation of such metabolites may have caused the perturbation of the germination rate.

Our investigations of the MALDI-MSI method showed that the applied matrices worked similarly and well across tissue types, genotypes, and growth stages for a range of general metabolites with different physicochemical properties (Fig. 5). Our MSI results for hexoses and amino acids were in accordance with other studies using both MALDI-MSI and LC-MS analyses of microdissected tissues (Figs. 5 and 7; Leonova et al., 2010; Gorzolka et al., 2016; Feenstra et al., 2017). Although MSI is not a quantitative technique, the overall trends in intensity of the amino acid ions in samples taken at different stages of the germination process (Fig. 4) matched the overall trends found in the analyses of extracts (Supplemental Fig. S3). Amino acids are poorly ionizable compounds, and because of their amphiprotic nature they have been reported to be hardly detectable by MALDI-MSI (Toue et al., 2014; Feenstra et al., 2017). Despite this, we visualized the distribution of 18 out of the 20 free proteinaceous amino acids, many with highly intense signals. The very high selectivity of the Orbitrap operating at 140,000 mass resolving power (applying a bin width of 0.01 Th in image generation) has likely improved detection compared with experiments on low-resolution MALDI-time of flight systems (Toue et al., 2014), but this does not explain the difference between the current study and the work of Feenstra et al. (2017). Factors such as sample preparation, mounting technique (carboxymethylcellulose, or CMC, versus gelatin), matrix (DAN vs. DHB), and sample drying (freeze drying versus desiccation) as well as differences in plant background matrices might have a significant impact on signal intensity and therefore on the successful detection of metabolites of interest. Some discrepancies were indeed observed between the LC-MS and the MALDI-MSI analyses of dhurrin and dhurrin-derived metabolites. At the early stages of germination, the discrepancies might be explained by the more localized accumulation, allowing compounds to be more easily detected by MSI than in the total seed extracts. However, at the last point of seedling development investigated, where the radicle has evolved into the primary root and the coleoptile was fully emerged from the seed coat, the concentrations of dhurrin and its recycling products in the dhurrin producing wild-type lines were just as high as at the previous growth stage, but only weakly or barely detected by MALDI-MSI. This may be explained by the substantial change in the plant background during the rapid transition from the seed to a seedling, which may have affected the ionization efficiency of the compounds. Ion suppression caused by matrix background is a common problem in MS-based studies. This study demonstrates the power of combining LC/MS and MALDI-MSI to gain insight into important biological processes. The results have demonstrated an apparent importance of dhurrin and its recycling processes at the earliest stages of sorghum grain germination and seedling formation and unveiled the high complexity of this metabolic grid and compartmentalizations. These insights point toward specific physiological functions of dhurrin and its recycling products.

MATERIALS AND METHODS

Plant Material

Seeds of four different sorghum (Sorghum bicolor) lines, namely ‘BTx623’, TCD1, tcd1 parent, and tcd1 mutant, were imbibed for 12 h in the dark at room temperature and germinated on moist filter paper in petri dishes at 28°C for 3 d. Samples were collected every 12 h at eight time points, dry grain and 0, 12, 24, 36, 48, 60, and 72 h, with 0 h corresponding to those taken immediately after 12 h of imbibition. At each time point, three seeds combined as one biological
Sample Preparation for Metabolite Profiling by LC-MS

Frozen plant material was ground in liquid nitrogen using mortar and pestle and extracted in 85% (v/v) aqueous methanol with 0.5% (v/v) formic acid (80°C, 5 min) in screw-cap microtubes (Sarstedt) and immediately cooled on ice. The extracts were transferred to screw cap glass vials and stored at −20°C until further pretreatment. For analyses, samples were diluted 5 times (1:5 [v/v]) in water and filtered through a membrane filter (0.45 μm; Merck Millipore) by centrifugation (2000 g, 5 min). After filtration, samples were stored at 4°C until analysis.

Quantification of Dhurrin and Related Metabolites by LC-3Q-MS

The diluted extracts were directly analyzed by LC-MS. Chromatography was performed on an Advance UHPLC system (Bruker). Separation was achieved on a Zorbax Eclipse XDB-C18 column (100 × 3.0 mm, 1.8 μm, 100 Å; Agilent Technologies), using a gradient of A = MilliQ grade water with 0.05% (v/v) formic acid and B = 100% methanol. The elution profile was 0 to 0.03 min, 2% B; 0.03 to 0.9 min, 2% to 15%; 0.9 to 1.4 min, 15% to 60%; 1.4 to 3.3 min, 60% to 100%; 3.3 to 3.9 min, 100%; 3.9 to 4.0 min, 100% to 2% B; and 4.0 to 5.0 min 2% B. The mobile phase flow rate was 500 μl min−1, and column temperature was maintained at 40°C. The liquid chromatography was coupled to an EVOQ Elite TripleQuad mass spectrometer (Bruker) equipped with an electrospray ion source (ESI) operated in combined positive and negative ionization mode. Instrument parameters were optimized by infusion experiments with pure standards. The ion spray voltage was maintained at 5000 and −4000 V in positive and negative ionization mode, respectively. Cone temperature was set to 300°C and cone gas to 20 psi. Heated probe temperature was set to 200°C and probe gas flow to 50 psi. Nebulizing gas was set to 60 psi and collision gas to 1.6 mTorr. Nitrogen was used as probe and nebulizing gas and argon as collision gas. Active exhaust was constantly on. Multiple reaction monitoring (MRM) was used to monitor analyte molecular ion to fragment ion transitions: MRMs were chosen based on direct infusion experiments. Detailed values for mass transitions can be found in Supplemental Table S2A. Both Q1 and Q3 quadrupoles were maintained at unit resolution. Alternatively, a modified MRM setting was used to select and filter the parent ion in both Q1 and Q3 while not applying any collision energy in Q2. This variation of selected ion monitoring (SIM) was called SIM2 and increased the signal strength by ~3.5- to 15-fold and improved signal-to-noise ratio up to 10-fold compared with normal SIM where only Q1 is used for ion selection. Also for SIM2, both Q1 and Q3 were kept at unit resolution. Bruker MS Workstation software (Version 8.2, Bruker) was used for data acquisition and processing. Linearity in ionization efficiencies was verified by analyzing dilution series of standard mixtures. Reference compounds for dhurrin, dhurrin amide, dhurrin acid, pGPAAc, p-hydroxyphenylacetic acid, p-hydroxyacetone, and GS-pOHPACN were used for absolute quantification by using 12 points external calibration curves in a range of concentrations from 0.003 to 90 μM.

Detection of Dhurrin Glucosides by LC-Quadrupole Time-Of-Flight-MS

Full scan analysis of sorghum extracts was performed using a Thermo Scientific Dionex Ultimate 3000 RS (ThermoFisher Scientific) coupled with a Bruker compact quadrupole-time-of-flight MS (Bruker Daltonics) equipped with an ESI source. For data acquisition and processing, Compass DataAnalysis software (version 4.3, Bruker Daltonics) was used. Then 8 μl of samples were injected on a LC Kinetics XB-C18 column (150 × 2.1 mm, 1.7 μm, Phenomenex), and compound separation was achieved using a gradient of A = MilliQ grade water with 0.05% (v/v) formic acid and B = acetonitrile with 0.05% (v/v) formic acid. The elution profile was 0 to 5 min, 2%; 4 to 9 min, 30%; B to 14 min, 100%; 14 to 21 min, 100% to 2%; 21 to 28 min, 2%. The mobile phase flow rate was 300 μl min−1, and column temperature was maintained at 40°C. The full-scan mass spectra were obtained within a range of m/z 50 to 1300 with the instrument operated in the positive ion mode. The spray voltage was set at 4000 V with the maximum dry temperature set at 250°C at the pressure of 1.2 Bar. The up-front collision-induced dissociation was set to off, and the optimized collision energy was set at 7eV in the collision cell. The quadrupole mass filter was set with low mass and with the ion energy of 4 eV. Extracted ion chromatograms for specific [M+H]+ adduct ions were used to detect dhurrin glucosides. Their fragmentation patterns, accurate masses, and retention times were used to identify peaks according to Pičmanová et al. (2015).

MALDI-MSI Sample Preparation and Fluorescence Microscopy

Samples were prepared according to the Kawamoto method (Kawamoto, 2003; Kawamoto and Kawamoto, 2014) with slight modifications. Frozen samples were embedded into a 2% (v/v) aqueous medium of CMC into a cryomold. The mold containing the embedded sample was submerged into a coolant solution of hexane/dry ice for ~5 min. When completely frozen, CMC blocks were removed from the mold and stored ~80°C until cryo-sectioning. Frozen samples were transferred to a Leica CM3050S cryostat (Leica Microsystems) at ~30°C to thermally equilibrate for 1 h. The embedded samples were fixed on the sample holder of the cryo-microtome using optimum cutting temperature gel (Tissue-Tek), sectioned on 10 μm thickness, collected with cryolipim II C9 (Section Lab), which was attached to prechilled glass slides with double-sided adhesive carbon tape (Agar scientific), and finally freeze-dried under vacuum at 1.0 mbar overnight. Three consecutive sections were selected for each sample; one was used for fluorescence microscopy analysis inspection and the others were coated with, respectively, DHB and DAN by sublimation, as described by Hankin et al. (2007). All the fluorescence images were captured on a Leica DMR HC (Leica Microsystems) wide-field fluorescence microscope, using a filter cube with a 340 to 380 nm excitation filter, 400 nm beam splitter, and 420 nm long-pass emission filter. Autofluorescence exposure times varied were based on the stage of the grain and the tissue composition (i.e. dry grain or germinated grain with protruded axes) but generally were set around between 600 and 800 ms. FIJI Image (Schindelin et al., 2012)
was used for any additional image preprocessing, such as cropping, adding the scale bar, or other relevant operations.

MALDI-MSI Analyses

MALDI-MSI analysis was performed on a Thermo QExactive Orbitrap mass spectrometer (Thermo Scientific) equipped with the AP-SMALDI10 ion source (TransMIT). The AP-SMALDI10 ion source was equipped with a MNL<sub>3</sub>00 nitrogen laser (LTB Lasertechnik) with 337 μm wavelength and 60 Hz frequency, operated using 30 laser pulses per pixel. Spectra were acquired in positive and negative ion mode (scan range m/z 70–700) at mass resolving power 140,000 at m/z 200. A matrix peak was used as lock-mass for internal mass calibration, ensuring a mass accuracy of 2 ppm or better. The acquired raw files were converted to mzML files using an mzML converter (Schramm and al., 2012), and images were generated using MSIReader v.1.01 (MATLAB; Robichaud et al., 2013) using a bin width of 0.01 Th and the following default parameters: MaxColorScale: 100,000, NormalizeOption: TIC, NormScale: 1 NormCutoff: 1, Minimum Abundance Threshold: 0.001. All MS images were normalized to the TIC in each pixel, and the maximum values for generating images are listed in Supplemental Figure S4. The same color scale was used regardless of genotype or germination stage. Metabolites were identified by accurate mass, and the complete list of m/z values and the mass tolerance applied for each is available in supplementary information (Supplemental Table S3).

Supplemental Data

The following supplemental materials are available.

Supplemental Table S1. The concentrations of dhurrin and dhurrin recycling products in four different lines.

Supplemental Table S2. Transitions and response factors used for MRM for analysis by LC-3Q-MS.

Supplemental Table S3. Compounds visualized by MALDI-MSI.

Supplemental Figure S1. Concentration of GS-PHPACN during germination and early seedling development in sorghum wild types.

Supplemental Figure S2. Dhurrin glucosides content during germination and early seedling development in sorghum wild types.

Supplemental Figure S3. Concentrations of individual amino during germination and early seedling development in four sorghum lines.

Supplemental Figure S4. Maximum values for generating images.

Supplemental Figure S5. Distribution and compartmentalization of general metabolites in tcd1.

Supplemental Figure S6. Comparison of average raw mass spectra of two selected samples of cv BTx623 and tcd1 at stage S1.

Supplemental Figure S7. All replicates of images in Figure 6.

Supplemental Figure S8. Distribution of all detected amino acids during germination in cv BTx623 by MALDI-MSI.

Supplemental Figure S9. Distribution of a subset of amino acids across biological replicates.

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