Genotype-Specific Changes in Vitamin B₆ Content and the PDX Family in Potato

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Vitamin B₆ is one of the most versatile cofactors in plants and an essential phytoneutrient in the human diet that benefits a variety of human health aspects. Although biosynthesis of the vitamin has been well resolved in recent years, the main research is currently based on Arabidopsis thaliana with very little work done on major crop plants. Here we provide the first report on interactions and expression profiles of PDX genes for vitamin B₆ biosynthesis in potato and how vitamin B₆ content varies in tubers of different genotypes. The results demonstrate that potato is an excellent resource for this vitamin and that strong natural variation in vitamin B₆ content among the tested cultivars indicates high potential to fortify vitamin B₆ nutrition in potato-based foods.

1. Introduction

Vitamin B₆ (vitB₆) is one of the most versatile cofactors found in living organisms. This generic term comprises three different compounds: pyridoxine (PN), pyridoxal (PL), and pyridoxamine (PN). When phosphorylated at their 5’ position (e.g., pyridoxal-5’-phosphate (PLP)) these compounds can serve as active cofactors in various biochemical reactions such as transaminations, decarboxylations, and racemizations that function primarily in amino acid metabolism [1]. However, the vitamin also plays a role in fatty acid and carbohydrate turnover [1]. In addition, B₆ vitamins function as potent antioxidants that help protect plants against abiotic stress conditions such as salt, high light, or drought, and are used by some pathogens as a neutralizing agent against photosensitizing toxins [2–4]. Due to its central role in general plant development and physiology, it is not surprising that mutants affected in vitB₆ biosynthesis display diverse defects including, for example, reduced chlorophyll, late flowering phenotypes, increased stress sensitivities, and aberrant embryogenesis [5–10]. Because the basic role of vitB₆ is conserved among humans and plants, the vitamin has also been implicated in many different health-related aspects ranging from diabetes and neurological disorders to cardiovascular diseases [11]. Plants, in contrast to humans, can synthesize vitB₆ de novo and thus represent an excellent dietary source of this essential compound. The biosynthesis requires the concerted activities of the PDX1 (pyridoxal biosynthesis 1) and PDX2 proteins that form a large multimeric complex to synthesize PLP from glutamine, a pentose-phosphate (ribose 5’-phosphate or ribulose 5’-phosphate) and a triose-phosphate (dihydroxyacetone phosphate or glyceraldehyde 3’-phosphate) [8,12].

Potato is one of the major food crops worldwide and the leading vegetable crop in the United States, with a production of 427,406 million cwt in the USA in 2011 (http://usda01.library.cornell.edu/usda/current/CropProdSu/CropProdSu-01-11-2013.pdf). Although potato is an excellent dietary source of vitamins and minerals, it is also denigrated for its high glycemic index and as a source for high fat and acrylamide containing food due to specific food processing steps [13–15]. Little is known about variations of this critical phytoneutrient in different genotypes and how the biosynthetic pathway functions in this plant.
Here we describe the PDX family in potato based on yeast-2-hybrid screens, their expression in different tissues, and how vitB6 contents vary in tubers among different potato varieties and after prolonged storage. The results indicate that the PDX family is highly conserved among different plant species, that potato is a very good resource for the vitamin, and that vitB6 content varies substantially among the tested genotypes. There is thus great potential for improving potato further through increasing the content of this specific phytonutrient, by either breeding or genetic manipulation to fortify the B6 vitamer as a healthy food resource for human nutrition.

2. Material and Methods

2.1. Plant Material. The potato cultivar Defender was grown in a culture chamber setup for long-day conditions (18 h:6 h light:dark; 20°C; 60% humidity) and used for qRT-PCR analysis, and all other samples came from a local supermarket or from field grown plants. All field grown plants were grown at the Washington State University Irrigated Agriculture Research Unit at Othello, WA, USA (46°47.277’ N Lat., 119°2.680’ W Long.) under standard commercial conditions. Baby potatoes analyzed in this work are defined as immature tubers harvested from young plants, typically 60–80 days after planting (DAP). Mature tubers were harvested from the same field at the end of the growing season, typically 160–179 DAP. Defender and Russet tubers used as samples for studies of the effects of storage time and temperature on vitB6 and reducing sugar (Glc + Fru) content, respectively, were grown as described by [16] and harvested in October 11, 2005 (181 days after planting). The harvested tubers were sorted according to weight, and tubers ranging from 170 g to 284 g were placed in storage (12°C, 95% RH) to wound heal for 11 days. Tubers of each cultivar were then stored at 6.7 and 8.9°C (95% RH) for 227 days (until May 26, 2006). Reducing sugar (Glc + Fru) and vitB6 levels were quantified in lyophilized samples of tuber tissue. Four replicates of five tubers each were sampled at 11 and 227 DAP. A central slice (approximately 1.5 mm thick, periderm attached) along the apical to basal axis from each of the five tubers was consolidated for each replicate, collectively frozen at −80°C and lyophilized. The dried tissue was ground with a mortar and pestle and sieved through a 60 mesh (0.246 mm) screen in preparation for vitB6 (see below) and reducing sugar determinations. Glucose and fructose were estimated according to a microplate modification of [17, 18] as described in [16]. The stoichiometric reduction of NADP as each hexose is converted to 6-phosphogluconate was monitored at 340 nm and quantitation was based on Glc and Fru standard curves.

2.2. Yeast-Two-Hybrid Screens. Y2H screens were performed on a mixed potato tissue cDNA library described in [19], following standard procedures as previously described in [20]. The library was in the beginning shuffled into the Gateway-compatible pACT2 (CLONTECH, Palo Alto, CA, USA; GenBank accession no. U29899) prey vector [21]. The corresponding Arabidopsis baits AtPDX1.3 and AtPDX2 were cloned into pBTM116-D9. All primary positive colonies from the initial screen were picked and first rescreened on medium containing 5 mM 3-aminol-1,2,4-triazole (3AT) to reduce the potential of analyzing autoactivators. Growing colonies under these conditions are referred to as “secondary positives.” DNA was isolated from 10 secondary positive yeast colonies, retransformed into yeast, and verified for positive interactions with either PDX1.3 or PDX2, respectively, as well as lack of auto-activation in the presence of an empty pBMI16-D9 vector. DNAs for clones that were verified to be positive were subject to sequencing. Based on gained sequence information, specific primers were designed for screening of the remaining secondary positives. All secondary positives for which no PCR product was gained underwent the same selection process of DNA as described above (isolation, retransformation, verifying interaction of auto-activation, and sequencing) to design new primer sets. This procedure was pursued till all secondary positive had been classified as summarized in Table 1. Primers used in the screen are shown in Supplementary Table 1 (see Supplementary Material available online at http://dx.doi.org/10.1155/2013/389723).

2.3. qRT-PCR Analysis. RNA isolation and first strand cDNA synthesis was done using an Isolate RNA Kit (Bioline USA Inc., MA) and a High Capacity cDNA Reverse Transcription Kit (Applied Biosystems, NY, USA). Quantitative RT-PCR reactions were performed following the protocol of 95°C for 7 mins, 40 cycles of 95°C for 15 secs, and 60°C for 1 min, on a 7500 Fast Real-Time PCR System (Applied Biosystems, NY, USA). ACTIN2 gene in potato was selected as reference gene in qRT-PCR for data modification. 50 ng of cDNA from different tissues of potato plants was used for template of each reaction. Primers used for qRT-PCR are shown in Supplementary Table 2.

2.4. Yeast Bioassay. VitB6 extraction and measurements of total vitB6 content from fresh and lyophilized tuber samples were done by a yeast bioassay, adapted from a protocol described in [9], and a vitB6 auxotrophic strain, ATCC9080. Data shown are averages of four biological replicates, each of which was in addition subject to three technical replicates. Yeast growth rates were followed at OD50 and used to determine total vitB6 content based on comparison to a standard curve.

3. Results

3.1. Screening of Y2H Library Yields Potato PDX1 Proteins and Novel Interactors. To identify expressed members of the potato PDX family, we performed two yeast-two-hybrid...
(Y2H) screens on a potato cDNA library generated from mixed tissues (source and sink leaf, root, stem, flower, petiole, and tuber; [19]). We decided to use Arabidopsis thaliana PDX proteins as baits since their function has been studied in great detail in the last years. In addition, their sequence and function appear to be conserved not only between Arabidopsis and Ginkgo biloba, but also in bacteria, making it highly likely that this would also be seen in potato [22]. One screen was done with Arabidopsis AtPDX1.3/At5g01410, and another with Arabidopsis AtPDX2/At5g60540. Through this combinatorial approach we anticipated to gain the corresponding expressed StPDX orthologs from potato, and potentially some novel interacting candidates. In total 182 and 304 primary positive clones were seen from the AtPDX1.3 and AtPDX2 screens, respectively. From those DNA was isolated and retransformed into the yeast, which resulted in 44 and 187 secondary positive clones. These remaining clones were further analyzed through sequencing and PCR to gain a better idea of the nature of these clones. Two different StPDX1 clones were identified in either screen, PGSC0003DMT400078045 and PGSC0003DMT400014535 that showed similarities to Arabidopsis AtPDX1 genes (Table 1). Phylogenetic comparison with Arabidopsis AtPDX1 proteins showed that PGSC0003DMT400078045 was most closely related to AtPDX1.1 (87.7% identity) and AtPDX1.3 (91.9% identity) (Figure 2(b)) and was named StPDX1.1. In contrast, PGSC0003DMT400014535 showed greater similarities to AtPDX1.2 (69% identity to AtPDX1.2 and 65% and 63% identity to AtPDX1.3 and AtPDX1.1, resp.), and was named accordingly StPDX1.2 (Figure 1(b)). While we found a few non-PDX encoding genes in either screen (Table I; Supplementary Table 1), no StPDX2 was present among the analyzed clones. This was unexpected since a functional PLP synthase depends on the concerted activities of PDX1 and PDX2 proteins [8]. In addition, publicly available sequence information predicts the presence of a StPDX2 gene (PGSC0003DMT400020828), which shows 75% identity on the protein level to AtPDX2.

3.2. StPDX Genes Are Expressed in All Potato Tissues. To further analyze the identified StPDX1 clones, we performed qRT-PCR analyses on different potato tissues shown in Figure 2(a). We also included the annotated StPDX2 to investigate whether and to what extent the gene is expressed in potato. As shown in Figure 2(b), all three PDX genes are expressed in all tested tissues. StPDX1.3 and StPDX2 show the highest levels of expression in leaf, while expression of StPDX1.2 is highest in sink tissues, including root and early tuber development. It was noteworthy that StPDX2 and StPDX1.3 showed high levels of expression preferentially in early tuber development (Figure 2(b)). Overall the data confirm that StPDX1.2 and StPDX1.3 are expressed in potato and demonstrate that this is also seen for the annotated StPDX2 gene.

3.3. ViB6 Content Varies Significantly among Different Genotypes and under Long-Term Storage Conditions. ViB6 is
an essential phytonutrient in the human diet, and potato is a major food crop. We were curious if there were major differences in vitB₆ content based on cultivar. Our first approach was to analyze tubers of four different potato varieties bought in a supermarket, Russet, French Fingerling, Austrian Crescent, and Purple Peruvian. The measurements showed that a Russet potato of 150 g weight contains 180 μg vitB₆ on average, while a fingerling specialty potato, like the Purple Peruvian, has 240 μg vitB₆ when consumed at equivalent amounts (Supplementary Figure 2). The data indicated that coloration of the potato may correlate with increased vitB₆ content and emphasized a remarkable difference present in the four genotypes. To verify these observations we expanded our test to 29 additional cultivars (Figure 3). Since expression of StPDX genes was highest in developing tubers we focused our analysis on young tubers. As shown in Figure 3(a), analysis of 23 different varieties of baby potatoes showed a variation in total vitB₆ content that ranged from 16.9 μg vitB₆/g dry weight (PORO07PG63-1) to 22 μg vitB₆/g dry weight (Ruby Crescent), a 30% increase from the lowest to highest level. Analysis of mature tubers from 10 different cultivars showed that vitB₆ level also varied significantly among the different genotypes with the lowest being again PORO07PG63-1 (18.6 μg vitB₆/g dry weight), with surprisingly high levels for Clearwater Russet (27 μg vitB₆/g dry weight). Samples that allowed comparison between baby and mature potatoes used in this analysis (PORO07PG63-1, La Ratte, Cecile, CO97222-1, Red Thumb, and AK3-1) showed no consistent effects of tuber maturity, indicating that vitB₆ concentrations are likely comparable throughout tuber growth. In addition, the analysis of a broader range of cultivars did not support the notion that vitB₆ levels correlate with a specific skin or flesh color. Finally, we investigated changes of vitB₆ content under long-term, low-temperature storage for two potato cultivars, Defender and Premier Russet (Figures 3(c) and 3(d)). For both cultivars, vitB₆ concentrations increased significantly over a 227-day storage period at either 6.7 or 8.9 °C, and the increases were greatest, averaging 42% (Defender) and 23% (Premier), at the highest temperature (Figures 3(c) and 3(d)).

4. Discussion

The current work identified two potato PDX1 proteins as interactors of Arabidopsis PDX1 and PDX2 proteins. In addition, the work provides first insights on tissue-specific expression profiles, and how vitB₆ content of tubers differs among various cultivars and over a 7.5-month storage period at temperatures consistent with those used in the industry.

The finding of only two different StPDX1 members in the Y2H screen with either bait is in agreement with the blast searches using Arabidopsis and potato PDX1 sequences against the available potato genome database (http://solanaceae.plantbiology.msu.edu/). This currently indicates that potato likely encodes for only two StPDX1 genes, in contrast to Arabidopsis that has three AtPDX1 family members. It was surprising that no PDX2 was found since a functional PLP synthase requires the activity of PDX2 proteins, because the potato genome encodes for a PDX2 protein, and we can demonstrate that the corresponding
gene is expressed. The reason for this absence of StPDX2 in the screen remains unclear but may be due to sequence differences among Arabidopsis and potato PDX2s that affect protein-protein interaction, since both proteins only share 69% identity. In this context it is also noteworthy that StPDX1.2 interacted with AtPDX2, which is remarkable since AtPDX1.2, to which StPDX1.2 is most closely related, is unable to assemble with AtPDX2 [6]. This again indicates that sequence variances between Arabidopsis and potato PDX proteins are responsible for this difference in interaction behavior, since AtPDX1.2 and StPDX1.2 are also only 69% identical. It will be interesting to determine in future experiments whether the potato StPDX2 is also unable to assemble with StPDX1.2 as observed for the Arabidopsis orthologs.

An interesting finding that may be worth pursuing is the novel interactions found in the two screens (Supplementary Table 1, Supplementary Figure 1). Although these interactions need to be verified for the potato StPDX proteins, they may suggest additional pathways or regulatory mechanisms by which StPDX activity is controlled in potato. These interactions may have a regulatory role on the PLP synthase or lead to a novel complex with a function distinct from vitB₆ biosynthesis (Supplementary Table 1).

VitB₆ plays a major role in various cellular reactions and confers many health benefits for humans, which may in
part be attributed to its antioxidant capabilities [11, 12, 23]. The Recommended Dietary Allowance from the American National Institute of Health (NIH) for vitB₆ is 2 mg/day for adults, and given that there is a possibility of vitB₆ malnutrition, as earlier reported for the USA [24], our data indicate that potato is a good source of this vitamin, regardless of tuber maturity or coloration. The significant variation observed among the tested genotypes indicates high potential for further increasing vitB₆ content of potato through directed breeding efforts.

Finally, starch breakdown and sweetening of tubers during long-term storage and under low-temperature conditions (cold sweetening) are undesirable for the potato processing industry due to the synthesis of dark-colored Maillard reaction products during frying. These products are a result of heat-induced nonenzymatic reactions between reducing sugars (Glc and Fru) and free amino acids to cause browning and acrylamide formation in fried potato products [15, 25]. The initial steps in starch breakdown depend in part on the activity of α-glucan phosphorylase, a PLP-dependent enzyme, which mediates the release of glucose-1-phosphate from α-1,4-linked glucan chains [26, 27].

Defender [28] is susceptible to sweetening during storage while Premier Russet is highly resistant [28–30]. Reducing sugar (Glc plus Fru) concentrations of the Defender tubers depicted in Figure 3(c) increased from 5.94±0.98 mg/g dry wt at 11 DAH to 30.6±1.8 and 26.1±1.3 mg/g dry wt when stored for 227 days at 6.7 and 8.9°C, respectively. In contrast, the Premier Russet tubers depicted in Figure 3(d) contained 1.81±0.37 mg/g dry wt reducing sugar at 11 DAH, and levels increased to only 11.9±1.4 and 4.88±0.61 mg/g dry wt when stored at 6.7 and 8.9°C, respectively, reflecting the inherently greater resistance to sweetening in this cultivar. These genotype-dependent differences in reducing sugar buildup over the 227-day storage period resulted in unacceptably dark processed fries for Defender, but fries from Premier Russet remained acceptably light (data not shown). The greater increase in vitB₆ content of Defender from 11 to 227 DAH was thus consistent with the higher levels of sweetening (and starch catabolism) in this cultivar. However, while the increases in vitB₆ may have a role in modulating the activity of α-glucan phosphorylase, vitB₆ levels were not directly correlated with the temperature-dependent buildup in reducing sugars in either cultivar. Further work is warranted to establish whether vitB₆ correlates with α-glucan phosphorylase activity and to elucidate its potential involvement in the cold-induced and genotype-dependent differences in sweetening metabolism reported here.

It is noteworthy that vitB₆ content of tubers increased postharvest, particularly when considering that vitamin C declined 62% in Defender and 51% in Premier Russet over a similar storage period [31]. Hence, with regard to vitB₆, the tubers became more nutritious with time in storage. While both cultivars had approximately the same vitB₆ content 11 DAH (ca. 17.6 μg/g dry wt), Defender tubers contained 10% higher levels than Premier tubers by 227 DAH. These data suggest that the capacity for postharvest increases in vitB₆ content of tubers may be genotype dependent, and further work is warranted to determine the heritability of this potentially important trait for breeding programs.

5. Conclusions

The current work emphasizes that potato is an excellent nutritional resource for vitB₆ with great potential for fortifying vitamin B₆ nutrition. The biosynthetic pathway is conserved among plants, and the corresponding genes are expressed in all tested tissues. The identification of new interactors is interesting and may open up new research directions with work related to PDX function in the future.

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