

Commentary

In Pursuit of Vitamin D in Plants

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Abstract: Vitamin D deficiency is a global concern. Much research has concentrated on the endogenous synthesis of vitamin D in human skin following exposure to ultraviolet-B radiation (UV-B, 280–315 nm). In many regions of the world there is insufficient UV-B radiation during winter months for adequate vitamin D production, and even when there is sufficient UV-B radiation, lifestyles and concerns about the risks of sun exposure may lead to insufficient exposure and to vitamin D deficiency. In these situations, dietary intake of vitamin D from foods or supplements is important for maintaining optimal vitamin D status. Some foods, such as fatty fish and fish liver oils, certain meats, eggs, mushrooms, dairy, and fortified foods, can provide significant amounts of vitamin D when considered cumulatively across the diet. However, little research has focussed on assessing edible plant foods for potential vitamin D content. The biosynthesis of vitamin D in animals, fungi and yeasts is well established; it is less well known that vitamin D is also biosynthesised in plants. Research dates back to the early 1900s, beginning with in vivo experiments showing the anti-rachitic activity of plants consumed by animals with induced rickets, and in vitro experiments using analytical methods with limited sensitivity. The most sensitive, specific and reliable method for measuring vitamin D and its metabolites is by liquid chromatography tandem mass spectrometry (LC-MS/MS). These assays have only recently been customised to allow measurement in foods, including plant materials. This commentary focuses on the current knowledge and research gaps around vitamin D in plants, and the potential of edible plants as an additional source of vitamin D for humans.

Keywords: vitamin D; 25-hydroxyvitamin D; 1,25-dihydroxyvitamin D; plants

1. Introduction

In most regions of the world, cutaneous synthesis following ultraviolet-B (UV-B, 280–315 nm) irradiation of 7-dehydrocholesterol in skin epidermal cells is the primary source of vitamin D for humans. When exposure to UV-B radiation is limited, because the ambient levels of UV-B radiation are low or skin is not exposed to the sun, dietary intake of vitamin D from food or supplements is required to maintain adequate vitamin D status. With the exception of mushrooms, the recognised natural dietary sources of vitamin D are animal-based (fish, meat, eggs and dairy), which raises concerns about low vitamin D intakes in populations that avoid, or consume low amounts of, animal products. For example, vitamin D deficiency is highly prevalent in India [1], where consumption of meat and dairy is low. A contribution of plant foods to dietary vitamin D intake could be important for such populations.

Liquid chromatography tandem mass spectrometry (LC-MS/MS) methods have only recently been applied to detect low concentrations of vitamin D and its metabolites in plants. These methods

are superior to the previous *in vivo* and *in vitro* bioassays, which cannot distinguish between vitamin D₂, vitamin D₃ and their hydroxylated forms. LC-MS/MS methods have been used to measure vitamin D in some animal products [2] and a small number of plants [3–5]. Although algae are not vascular plants, and so fall outside the scope of this commentary, a macroalga, *Sargassum*, has been shown to have anti-rachitic activity [6], and both vitamin D₂ and vitamin D₃ have been found in fresh water phytoplankton [7].

This commentary is an update on previous reviews of vitamin D in plants [5,8], and introduces several additional concepts: the lack of identification in plants of relevant proteins involved in vitamin D metabolism and transport; evidence suggesting that exposure to UV-B radiation may not be required for synthesis of vitamin D in plants; and the possibility of native Australian plants as a potential source of vitamin D. We briefly outline the current knowledge and research gaps around vitamin D in plants (Figure 1) and emphasise the potential role of vitamin D in plants for supplementing intake in humans.

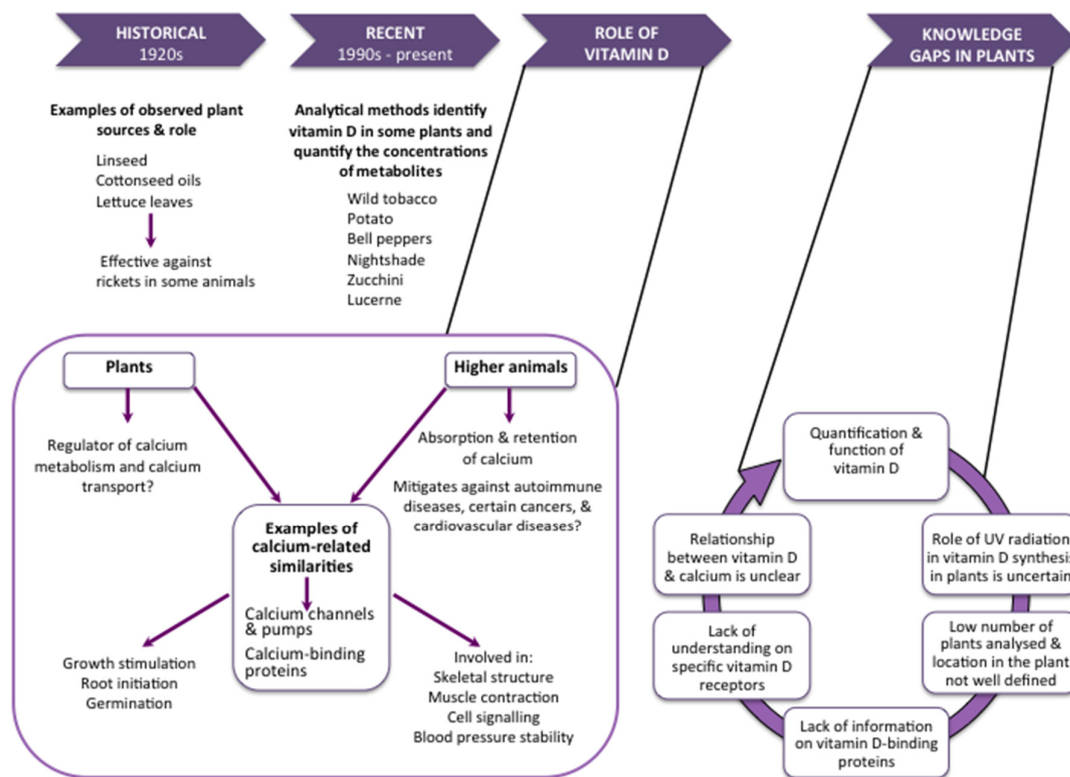


Figure 1. Vitamin D in plants: current knowledge and research gaps.

2. Metabolism of Vitamin D in Plants

UV-B irradiation of sterol precursors results in the production of vitamin D₃ in mammalian skin and vitamin D₂ in yeasts and fungi. Both forms have been used as vitamin D supplements and in food fortification. Vitamin D₄ (22-dihydroergocalciferol) has also been identified in fungi [9,10]. While mammalian and fungal cells each contain only one major sterol (7-dehydrocholesterol and ergosterol, respectively), plants have a complex sterol mixture, including sitosterol, stigmasterol, ergosterol, 7-dehydrocholesterol, campesterol and 24-methylcholesterol [11]. Cholesterol is only a minor sterol (1%–2% of total plant sterols) in most plant species, but can represent at least 10% of total sterols in some plants, such as the Solanaceae [12], a family of flowering plants including tomato, potato, eggplant, capsicum and tobacco. Plasma membranes from leaf cells of *Lycopersicon esculentum* (tomato plant) undergo changes in UV-induced absorbance with a trough at about 295 nm [11], pointing to the disappearance of a substance with an absorption peak at this wavelength. The wavelength at 295

nm is close to the action peak for previtamin D photosynthesis, which is thought to be due to the conversion of provitamin D which has an absorption maximum at this wavelength.

In humans, vitamin D₃ synthesised in the epidermis is taken up into the bloodstream tightly bound to a vitamin D-binding protein. Both vitamin D₃ (deriving from sun exposure and dietary sources) and vitamin D₂ (deriving only from dietary sources) are metabolised in the liver through an hydroxylation pathway to the intermediate compound, 25-hydroxyvitamin D (25(OH)D), the major circulating and storage form. A further hydroxylation pathway, mainly in the kidney but also in other tissues, produces the active form of vitamin D, 1,25-dihydroxyvitamin D (1,25(OH)₂D). The effects of 1,25(OH)₂D in humans are mediated through ligation with nuclear vitamin D receptors (VDR) [13] and via membrane rapid-response receptors [14,15], both of which are found in most human tissues.

Neither VDR nor vitamin D-binding protein have yet been found in plants [16], but a VDR-like binding protein for 1,25(OH)₂D₃ has been described in *Solanum glaucophyllum* (waxy leaf nightshade), suggesting that similar mechanisms of action may be present in plants [17]. The enzymes involved in the two hydroxylation reactions (25-hydroxylase and 25-hydroxyvitamin D 1 α -hydroxylase), along with 25(OH)D₃ and 1,25(OH)₂D₃, in both free and glycosidic forms, have been identified in the leaves of *Solanum malacoxylon* [18] and *Cestrum diurnum* [19], plants known to be responsible for calcinosis in animals. Vitamin D₃ is present in plants not only in free form, but also as a glycoside. Although it is not clear whether the glycoside and the free form of vitamin D₃ are equally absorbed, glycosylation does not appear to substantially reduce the activity of vitamin D₃: a study in chickens demonstrated that the vitamin D₃ glycoside has an activity of 90%–95% compared with the aglycone [20].

3. The Function of Vitamin D in Plants

There may be some similarities between plants and animals in the way in which calcium and vitamin D are associated in regulatory processes. Vitamin D has a critical role in calcium and phosphate homeostasis in animals [16]. When blood calcium concentrations fall, there is upregulation of 1 α -hydroxylase to increase 1,25(OH)₂D concentrations. This results in increased intestinal absorption of calcium, along with decreased renal excretion, to restore normal blood calcium concentrations. When this mechanism is insufficient, bone metabolism is upregulated to release calcium from skeletal stores [21]. Plants have similar calcium channels and pumps to those found in animals [22], and calcium ions are a core regulator of plant cell physiology [16]. Calcium is required for stimulation of growth, root initiation and promotion of germination in plants [22]. 1,25(OH)₂D₃ has been shown to influence growth and calcium transport in roots of *Phaseolus vulgaris* (common bean) by increasing synthesis of calmodulin [8], a calcium-binding messenger protein found in all eukaryotic cells.

4. Vitamin D₂ Content of Plants

Many plants contain endophytic fungi, which have cell membranes containing ergosterol [23–25]. Thus, vitamin D₂ has been found in plants as a result of photoconversion of ergosterol in these fungal contaminants. In 1924, Hess and Weinstock found that UV-irradiated linseed oils, cottonseed oils and lettuce leaves were effective anti-rachitic agents when fed to rats [26]. Other studies have extended this work to show benefits in cattle [27] and chickens [28] with induced rickets. These experiments provided an in vivo assay of the anti-rachitic activity of various plants. The active compound was later identified as vitamin D₂ produced from fungal contamination of the plants.

Perennial ryegrass, a common grass in permanent pastures, contains both ergosterol and vitamin D₂ as a result of fungal contamination [3], and low amounts of vitamin D₂ have been found in milk, presumably originating from fungal contamination of grass and hay [29]. However, although symbiotic fungi may improve resistance to stress and insects in the host plant [30], some fungi have been implicated in crop spoilage and/or toxicity to animals [31]. From a human nutrition perspective, there is conflicting evidence on the relative bioavailability and bioeffectiveness of vitamin D₂ versus vitamin D₃ [32–35].

5. Vitamin D₃ Content of Plants

Vitamin D₃ and 25(OH)D₃ have been found mainly in the Solanaceae family, with research focussed on the leaves, which are known to be poisonous in large amounts. It is currently not clear whether vitamin D₃ and its metabolites are present in the edible fruits of the Solanaceae. An extract from the leaves of the tomato plant was shown to significantly increase serum calcium concentrations in vitamin D-deficient rats, while the fruit was devoid of vitamin D-like activity [36]. To our knowledge, LC-MS/MS methods have not been used for analysing vitamin D₃ and its metabolites in other edible fruits.

The presence of vitamin D₃ may not be dependent on exposure to light in all plants. Previous studies in *Solanum glaucophyllum* have shown that vitamin D₃ compounds, including 7-dehydrocholesterol, vitamin D₃, 25(OH)D₃ and 1,25(OH)₂D₃, are present in cultures grown in the absence of light [37,38]. In samples grown in vitro in darkness, both 25(OH)D₃ and 1,25(OH)₂D₃ were present in the stem, leaf and (inedible) fruit [39]. Nevertheless, UV irradiation of the plants substantially increased concentrations of 1,25(OH)₂D₃ in the leaves. This suggests that treatment with UV radiation may be an effective method for increasing the content of vitamin D₃ and its metabolites in plants (at least in the leaves), similar to the effect that UV irradiation has on increasing vitamin D₂ in mushrooms [40].

Recently, Jäpelt and colleagues compared vitamin D₃ and its hydroxylated metabolites in the UV-treated and untreated leaves of the tomato plant, waxy leaf nightshade and bell pepper [41]. Using LC-MS/MS methods, vitamin D₃ was identified in the leaves of all three plants following treatment with UV radiation. Quantifiable 25(OH)D₃ was detected in the UV-treated leaves of waxy leaf nightshade, tomato plant, and bell pepper, and was also in the untreated leaves of waxy leaf nightshade. Table 1 shows the concentrations of 7-dehydrocholesterol, vitamin D₃, 25(OH)D₃ and 1,25(OH)₂D₃, measured by LC-MS/MS methods, in the leaves of various plant species with and without treatment with UV irradiation.

6. Research Gaps

There are relatively few studies quantifying vitamin D₃ and its metabolites in plants, and even fewer investigating plant parts other than leaves. Furthermore, the chemical configuration and availability of different vitamin D₃ metabolites in plants remain unknown. There is also currently little information on the occurrence and concentration of vitamin D₃ in plants, or the vitamin D₃ pathway, including binding proteins, receptors and activating enzymes. Treatment of edible plants or plant parts with UV radiation may be an approach for increasing vitamin D in the food supply, particularly for those who avoid animal products. However, with respect to the Solanaceae family, the toxicity of the leaves prohibits their use as a potential source of vitamin D. Further exploration of vitamin D₃ in native Australian plant foods may be warranted, since they grow naturally in high UV radiation environments and many have medicinal properties. Examples of native plant foods include *Tasmannia lanceolata* (Tasmanian pepper), which has a long history of use by Australian Aboriginal people as a food flavouring and is a good source of calcium; and *Solanum centrale* (bush tomato), belonging to the Solanaceae family (Figure 2). A major challenge will be to ensure that methods to analyse vitamin D₃ and its metabolites are reliable and accurate when measuring low concentrations in the complex matrices presented by different plant parts.

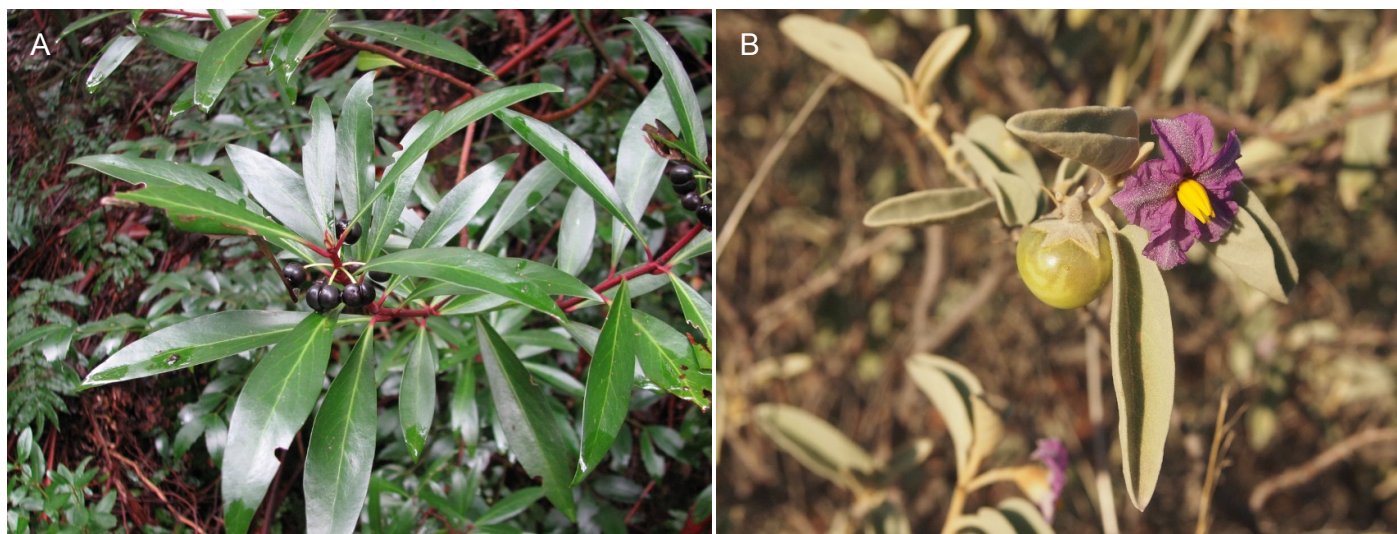


Figure 2. (A) *Tasmannia lanceolata* (Tasmanian pepper) (Credit: Mark Marathon—Own work, CC BY-SA 3.0, <https://commons.wikimedia.org/w/index.php?curid=28133459>); (B) *Solanum centrale* (bush tomato) (Credit: Melburnian—Self-photographed, CC BY 3.0, <https://commons.wikimedia.org/w/index.php?curid=10145927>).

Table 1. Concentrations of vitamin D₃ metabolites (dry weight) in plant leaves from the Solanaceae family, measured by liquid chromatography tandem mass spectrometry.

Species	7-dehydrocholesterol (µg/g)	Vitamin D ₃ (µg/g)	25(OH)D ₃ (µg/g)	1,25(OH) ₂ D ₃ (µg/g)
Non-irradiated				
<i>Lycopersicon esculentum</i> (tomato plant)	0.47 [4] ¹	Not identified [4] ¹ 0.0017 [41] ²	<0.00002 [41] ²	<0.0001 [41] ²
<i>Solanum glaucophyllum</i> (waxy leaf nightshade)	0.67 [4] ¹	Not identified [4] ¹ 0.0032 [41] ²	0.0008 [41] ²	<0.0001 [41] ²
<i>Capsicum annuum</i> (bell pepper)	0.03 [4] ¹	Not identified [4] ¹ <0.00002 [41] ²	<0.00002 [41] ²	<0.0001 [41] ²
UV-irradiated				
<i>Lycopersicon esculentum</i> (tomato plant)	0.23 [4] ¹	0.09 [4] ¹ 0.1 [41] ²	0.0043 [41] ²	<0.0001 [41] ²
<i>Solanum glaucophyllum</i> (waxy leaf nightshade)	1.26 [4] ¹	0.21 [4] ¹ 0.2 [41] ²	0.031 [41] ²	0.032 [41] ²
<i>Capsicum annuum</i> (bell pepper)	0.03 [4] ¹	Not identified [4] ¹ 0.0029 [41] ²	0.0005 [41] ²	<0.0001 [41] ²

¹ Atmospheric pressure chemical ionisation liquid chromatography tandem mass spectrometry; ² Liquid chromatography-electrospray ionisation tandem mass spectrometry.

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Conflicts of Interest: The authors declare no conflict of interest.

Abbreviations

The following abbreviations are used in this manuscript:

UV-B	Ultraviolet-B radiation
25(OH)D	25-hydroxyvitamin D
1,25(OH) ₂ D	1,25-dihydroxyvitamin D
VDR	Vitamin D receptor
LC-MS/MS	Liquid-chromatography tandem mass spectrometry

References

- Basu, S.; Gupta, R.; Mitra, M.; Ghosh, A. Prevalence of vitamin D deficiency in a pediatric hospital of Eastern India. *Indian J. Clin. Biochem.* **2015**, *30*, 167–173.
- Strobel, N.; Buddhadasa, S.; Adorno, P.; Stockham, K.; Greenfield, H. Vitamin D and 25-hydroxyvitamin D determination in meats by LC-IT-MS. *Food Chem.* **2013**, *138*, 1042–1047.
- Jäpelt, R.B.; Didion, T.; Smedsgaard, J.; Jakobsen, J. Seasonal variation of provitamin D₂ and vitamin D₂ in perennial ryegrass (*Lolium perenne* L.). *J. Agric. Food Chem.* **2011**, *59*, 10907–10912.
- Jäpelt, R.B.; Silvestro, D.; Smedsgaard, J.; Jensen, P.E.; Jakobsen, J. LC-MS/MS with atmospheric pressure chemical ionization to study the effect of UV treatment on the formation of vitamin D₃ and sterols in plants. *Food Chem.* **2011**, *129*, 217–225.
- Jäpelt, R.B.; Jakobsen, J. Vitamin D in plants: A review of occurrence, analysis, and biosynthesis. *Front. Plant Sci.* **2013**, *4*, 136.
- Darby, H.H.; Clarke, H.T. The plant origin of a vitamin D. *Science* **1937**, *85*, 318–319.
- Sunita Rao, D.; Raghuramulu, N. Food chain as origin of vitamin D in fish. *Comp. Biochem. Physiol.* **1996**, *114A*, 15–19.
- Boland, R.L. Plants as a source of vitamin D₃ metabolites. *Nutr. Rev.* **1986**, *44*, 1–8.
- Phillips, K.M.; Horst, R.L.; Koszewski, N.J.; Simon, R.R. Vitamin D₄ in mushrooms. *PLoS ONE* **2012**, *7*, e40702.
- Krings, U.; Berger, R.G. Dynamics of sterols and fatty acids during UV-B treatment of oyster mushroom. *Food Chem.* **2014**, *149*, 10–14.
- Björn, L.; Widell, S.; Wang, T. Evolution of UV-B regulation and protection in plants. *Adv. Space Res.* **2002**, *30*, 1557–1562.
- Frega, N.; Bocci, F.; Conte, L.S.; Testa, F. Chemical composition of tobacco seeds (*Nicotiana tabacum* L.). *J. Am. Oil Chem. Soc.* **1991**, *68*, 29–33.
- Haussler, M.R.; Haussler, C.A.; Bartik, L.; Whitfield, G.K.; Hsieh, J.C.; Slater, S.; Jurutka, P.W. Vitamin D receptor: Molecular signaling and actions of nutritional ligands in disease prevention. *Nutr. Rev.* **2008**, *66*, S98–112.
- Boyan, B.D.; Chen, J.; Schwartz, Z. Mechanism of vitamin D₃-dependent 1,25-dihydroxyvitamin D₃ signaling in musculoskeletal cells. *Steroids* **2012**, *77*, 892–896.
- Doroudi, M.; Plaisance, M.C.; Boyan, B.D.; Schwartz, Z. Membrane actions of 1,25(OH)₂D₃ are mediated by Ca²⁺/calmodulin-dependent protein kinase II in bone and cartilage cells. *J. Steroid Biochem. Mol. Biol.* **2015**, *145*, 65–74.
- Bikle, D. Vitamin D: An ancient hormone. *Exp. Dermatol.* **2010**, *20*, 7–13.
- Milanesi, L.; Boland, R. Presence of vitamin D₃ receptor (VDR)-like proteins in *Solanum glaucophyllum*. *Physiol. Plant* **2006**, *128*, 341–350.
- Esparza, M.S.; Vega, M.; Boland, R.L. Synthesis and composition of vitamin D₃ metabolites in *Solanum malacoxylon*. *Biochim. Biophys. Acta* **1982**, *719*, 633–640.
- Hughes, M.R.; McCain, T.A.; Chang, S.Y.; Haussler, M.R.; Villareale, M.; Wasserman, R.H. Presence of 1,25-dihydroxyvitamin D₃-glycoside in the calcinogenic plant *Cestrum diurnum*. *Nature* **1977**, *268*, 347–349.

20. Rambeck, W.A.; Weiser, H.; Zucker, H. Biological activity of glycosides of vitamin D₃ and 1 alpha-hydroxyvitamin D₃. *Int. J. Vitam Nutr. Res.* **1984**, *54*, 25–34.
21. Cranney, A.; Horsley, T.; O'Donnell, S.; Weiler, H.A.; Puil, L.; Ooi, D.S.; Atkinson, S.A.; Ward, L.M.; Moher, D.; Hanley, D.A.; et al. *Effectiveness and Safety of Vitamin D in Relation to Bone Health*; Evidence Report/Technology Assessment no. 158 (prepared by the University of Ottawa Evidence-Based Practice Center (uo-epc) under Contract No. 290-02-0021. Ahrq publication No. 07-e013; University of Ottawa Evidence-Based Practice Center: Rockville, MD, USA, 2007.
22. Dodd, A.N.; Kudla, J.; Sanders, D. The language of calcium signaling. *Annu. Rev. Plant Biol.* **2010**, *61*, 593–620.
23. De Souza Leite, T.; Cnossen-Fassoni, A.; Pereira, O.L.; Mizubuti, E.S.; de Araujo, E.F.; de Queiroz, M.V. Novel and highly diverse fungal endophytes in soybean revealed by the consortium of two different techniques. *J. Microbiol.* **2013**, *51*, 56–69.
24. Gonzaga, L.L.; Costa, L.E.; Santos, T.T.; Araujo, E.F.; Queiroz, M.V. Endophytic fungi from the genus *Colletotrichum* are abundant in the *Phaseolus vulgaris* and have high genetic diversity. *J. Appl. Microbiol.* **2015**, *118*, 485–496.
25. Schulz, B.; Boyle, C. The endophytic continuum. *Mycol. Res.* **2005**, *109*, 661–686.
26. Hess, A.F.; Weinstock, M. Antirachitic properties imparted to inert fluids and to green vegetables by ultra-violet irradiation. *J. Biol. Chem.* **1924**, *62*, 301–313.
27. Bechtel, H.E.; Huffman, C.F.; Duncan, C.W.; Hoppert, C.A. Vitamin D studies in cattle. *J. Dairy Sci.* **1936**, *19*, 359–372.
28. Wasserman, R.H.; Corradino, R.A.; Krook, L.; Hughes, M.R.; Haussler, M.R. Studies on the 1 α , 25-dihydroxycholecalciferol-like activity in a calcinogenic plant, *Cestrum diurnum*, in the chick. *J. Nutr.* **1976**, *106*, 457–465.
29. Jakobsen, J.; Saxholt, E. Vitamin D metabolites in bovine milk and butter. *J. Food Comp. Anal.* **2009**, *22*, 472–478.
30. Rodriguez, R.; Redman, R. More than 400 million years of evolution and some plants still can't make it on their own: Plant stress tolerance via fungal symbiosis. *J. Exp. Biol.* **2008**, *59*, 1109–1114.
31. Scudamore, K.A.; Livesey, C.T. Occurrence and significance of mycotoxins in forage crops and silage: A review. *J. Sci. Food Agric.* **1998**, *77*, 1–17.
32. Thacher, T.D.; Fischer, P.R.; Obadofin, M.O.; Levine, M.A.; Singh, R.J.; Pettifor, J.M. Comparison of metabolism of vitamins D₂ and D₃ in children with nutritional rickets. *J. Bone Miner. Res.* **2010**, *25*, 1988–1995.
33. Fisk, C.M.; Theobald, H.E.; Sanders, T.A. Fortified malted milk drinks containing low-dose ergocalciferol and cholecalciferol do not differ in their capacity to raise serum 25-hydroxyvitamin D concentrations in healthy men and women not exposed to UV-B. *J. Nutr.* **2012**, *142*, 1286–1290.
34. Stephensen, C.B.; Zerofsky, M.; Burnett, D.J.; Lin, Y.P.; Hammock, B.D.; Hall, L.M.; McHugh, T. Ergocalciferol from mushrooms or supplements consumed with a standard meal increases 25-hydroxyergocalciferol but decreases 25-hydroxycholecalciferol in the serum of healthy adults. *J. Nutr.* **2012**, *142*, 1246–1252.
35. Armas, L.A.; Hollis, B.W.; Heaney, R.P. Vitamin D₂ is much less effective than vitamin D₃ in humans. *J. Clin. Endocrinol. Metab.* **2004**, *89*, 5387–5391.
36. Prema, T.P.; Raghuramulu, N. Vitamin D₃ and its metabolites in the tomato plant. *Phytochemistry* **1996**, *42*, 617–620.
37. Curino, A.; Skliar, M.; Boland, R. Identification of 7-dehydrocholesterol, vitamin D₃, 25(OH)-vitamin D₃ and 1,25(OH)₂-vitamin D₃ in *Solanum glaucophyllum* cultures grown in absence of light. *Biochim. Biophys. Acta* **1998**, *1425*, 485–492.
38. Weissenberg, M.; Levy, A.; Wasserman, R.H. Distribution of calcitriol activity in *Solanum glaucophyllum* plants and cell cultures. *Phytochemistry* **1989**, *28*, 795–798.
39. Curino, A.; Milanesi, L.; Benassati, S.; Skliar, M.; Boland, R. Effect of culture conditions on the synthesis of vitamin D₃ metabolites in *Solanum glaucophyllum* grown in vitro. *Phytochemistry* **2001**, *58*, 81–89.

40. Simon, R.R.; Phillips, K.M.; Horst, R.L.; Munro, I.C. Vitamin D mushrooms: Comparison of the composition of button mushrooms (*Agaricus bisporus*) treated postharvest with UVB light or sunlight. *J. Agric. Food Chem.* **2011**, *59*, 8724–8732.
41. Jäpelt, R.B.; Silvestro, D.; Smedsgaard, J.; Jensen, P.E.; Jakobsen, J. Quantification of vitamin D₃ and its hydroxylated metabolites in waxy leaf nightshade (*Solanum glaucophyllum* Desf.), tomato (*Solanum lycopersicum* L.) and bell pepper (*Capsicum annuum* L.). *Food Chem.* **2013**, *138*, 1206–1211.



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