Dedicated to Professor Marko D. Mihovilovic on the occasion of his 50th birthday

Abstract: This work reports on the concise total synthesis of eight natural products of the mugineic acid and avenic acid families (phytosiderophores). An innovative “east-to-west” assembly of the trimeric products resulted in a high degree of divergence enabling the formation of the final products in just 10 or 11 steps each with a minimum of overall synthetic effort. Chiral pool starting materials (L-malic acid, threonines) were employed for the outer building blocks while the middle building blocks were accessed by diastereo- and enantioselective methods. A highlight of this work consists in the straightforward preparation of epimeric hydroxyazetidine amino acids, useful building blocks on their own, enabling the first synthesis of 3''-hydroxymugineic acid and 3''-hydroxy-2'-deoxymugineic acid.

Micronutrient acquisition is an important factor in growth and survival of any living organism. Plants are stationary and need to fill all their needs from the soil they grow in. In calcareous, high pH soils, the solubility and therefore plant availability of some of these crucial metal ions, like iron, zinc and copper, is diminished, to a degree that it greatly inhibits plant growth and leads to chlorosis of the leaves. [1–2] On top of reduced yield, micronutrient deficiencies in major crops like wheat, barley, rice and maize carry over to local consumers, causing micronutrient deficiency in humans (“hidden hunger”) with severe negative effects on child growth, development and disease resistance.[3]

A unique strategy used by gramineous plants for the uptake of Fe and potentially also Zn and Cu relies on phytosiderophores (PS), which are multidentate chelators of metal ions. PS are exuded by roots of grass species into the surrounding soil (i.e. rhizosphere) where they can complex Fe\(^{III}\) ions from soil particles.[4] The PS–Fe\(^{III}\) 1:1 complex is then taken up as whole complex and the iron liberated within the cell. This complexation strategy renders grass species more efficient in Fe acquisition compared to non-grass species, particularly in high pH soil, where soluble Fe concentrations are low. Investigations into plants molecular mechanisms, and consequently efforts towards the synthesis of these natural products have been reported since the late 1970s.[5a–h]

Variation in naturally occurring PS (see Figure 1) arises from hydroxy groups present on C-2 of the western (left) and middle subunits, respectively, while the eastern hydroxyacid fragment is conserved throughout the mugineic, avenic and distichonic acid series. The general synthetic strategy is closely related to the biosynthesis[6] and specific synthetic solutions were published for avenic acid A (AVA, I), deoxymugineic acid (DMA, IV), mugineic acid (MA, VIII) and 3''-epi-hydroxy-MA (VI).

In the state-of-the-art mugineic acid synthesis by Namba et al.,[5c–d] which used a “west-to-east” approach, stereoselectiv-
ity issues arose during the introduction of the 2'-hydroxygroup by allylic oxidation requiring late stage separation of diastereomers by means of preparative HPLC, thus, limiting access to larger quantities of MA (VIII). In earlier work CS and co-workers synthesized \( \text{^{13}C}_2 \)-labelled PS in a similar fashion.\(^{[5c]}\) The activity of the synthetic community notwithstanding, to date, useful quantities of all PS in their natural and isotopically \(^{13}C\) labelled form (used as internal standards for trace analysis in soil) are currently not available, making it difficult for groups studying these plant mechanisms to efficiently carry out their research. In this light, no general approach to the more complex mugineic acids (II, III, VI and VII) has so far been reported, most likely due to lack of availability of building blocks 4 and 5. Aiming for more members of this class of compounds and due to the high diversity of „western“ fragments present in naturally occurring PS, the opposite direction of assembly was recognized as a tentatively superior approach which has thus far been widely neglected by the synthetic community.\(^{[5a]}\) Within the framework of an interdisciplinary project we set out to establish a general solution for all members of the mugineic and avenic acid family in natural form compatible with an application towards \( \text{^{13}C}_2 \)-labelled versions. These are required as standards for high performance trace quantification in complex matrices as well as for biodegradation experiments.

Results and Discussion

As mentioned above our aim was to synthesize eight different PS from „east-to-west“ from one common set of building blocks. Thus, the best transformations available to date to PS from „east-to-west“ from one common set of building blocks. Accordingly, the chiral pinanone auxiliary 13 was attached to tert-butyl protected glycine 14 under Lewis acid catalysis (See Scheme 1). The resulting imine 15 was then added in a titanium mediated aldol reaction to crotonaldehyde, giving rise to a hydroximine (not depicted), which due to limited stability (retro-Aldol) was subsequently hydrolysed under acidic conditions delivering subunit 2 in three steps in a 77% overall yield. Protected non-labelled allylglycine 3 was prepared in a straightforward fashion in one step from commercial L-allylglycine (12).

The synthesis of hydroxylated L-azetidines 4 and 5 was carried out starting from protected L- or D-threonine derivatives 18 and ent-18 in a three-step procedure. Initial attachment of a 2-picolinamide (PA) directing group (DG) was followed by cyclisation under palladium catalysed oxidative C–H activation conditions developed by Chen et al.\(^{[7]}\) The tert-butyl protecting groups in 19 and ent-19 proved to be ideal for the overall success of this route due to their excellent stability, while silyl ethers and other protecting groups were not suitable.

Next, the methyl ester and DG cleavage was evaluated, at first under basic conditions. When a hydroxide base was employed at room temperature, epimerization to the more stable\(^{[10]}\) trans-azetidines (21 and ent-21) proceeded rapidly.
and with almost full conversion. Subsequently, the amide was cleaved by further heating the reaction mixture. This turned out to be a very efficient solution to access 5. An influence in temperature and base was found, with epimerization being slower than ester cleavage at lower temperatures. Nevertheless, only a 62:38 mixture of the desired (cis-azetidine 4) and undesired product (ent-5) could be obtained with this method. The assigned stereochemistry was proven by X-ray diffraction of pure ent-5 (see Scheme 2, Deposition Number 2007994 contains the supplementary crystallographic data for this paper. These data are provided free of charge by the joint Cambridge Crystallographic Data Centre and Fachinformationszentrum Karlsruhe Access Structures service www.ccdc.cam.ac.uk/structures). While the 56% isolated yield of 4 in this step are acceptable for such an elusive material, a selective approach was also developed. Thus, the methyl ester in 19 was cleaved first with the mild reagent Me3SnOH.[11] No epimerization at C-2 was observed and the picolinamide could be subsequently cleaved with base after careful neutralization of the carboxylic acid. The amino acids 4 and 5 were separated from any minor isomers present by preparative HPLC and obtained in >60% overall yield from 18 and ent-18.

The eastern subunit aldehyde 1 was prepared in a good yield over 5 steps as reported[25] and assembly of the two central dimeric intermediates could commence. Aminoesters 2 and 3 were allylated under reductive conditions (NaCNBH3) with aldehyde 1. A smooth reaction was achieved in both instances giving 83% and 80% of dimeric amines 22 and 24 upon chromatographic purification at decent scale (Scheme 3). Attachment of a Boc protecting group was realized in high yield (Boc2O, THF) and the resulting carbamates 23 and 25 proved to be suitably stable storage compounds. The dimeric compounds 23 and 25 now enter the second stage of diversification towards two sets of four phytosiderophores each, with a 2'-hydroxy and 2'-deoxy-structure, respectively. The olefin was cleaved by ozonolysis yielding aldehydes 26 and 27 which were subsequently treated with one of the four respective western building blocks 4–7 under reductive amination conditions, as before (Scheme 4). Thanks to the universal protecting group strategy throughout the synthesis, global deprotection of the molecules proved facile and was induced by treatment with 6 m HCl. The resulting amino acids I–VIII were purified on Dowex® resin and isolated as the respective ammonium salts. For highly pure material additional trituration steps or preparative HPLC can be carried out.[26] To the best of our knowledge this represents the first total synthesis of 3’–epi-hydroxy–DMA (II), 3’–hydroxy–DMA (III), hydroxyavenic acid A (V) and 3’–hydroxy–MA (VII).

A concise and modular synthesis of phytosiderophore natural products was achieved. Starting from l-malic acid, target compounds I–VIII could be prepared in 10 or 11 steps longest linear sequence (15–25% overall yield). The required building blocks could all be efficiently accessed by C–H activation and stereoselective aldol reaction. Common key intermediates (23 and 25) allow very flexible and fast resynthesis and delivery of materials for the planned applications within our current project and beyond. As mentioned before, by employing properly protected building blocks 2 and 3 as „eastern“ fragments and compound 14 as „western“ fragment, access to the nicotianamine and distichonic acid family is well within the scope of the developed methodology.
Further work on fully assembled $^{12}$C$_2$-labelled versions of compounds I–VIII using the presented strategy as well as development towards scale up (gram-scale) of the avenic acid and mugineic acid syntheses and other PS are currently ongoing in our laboratories and will be reported in due time.

Acknowledgements

This work was supported by the European Research Council (ERC-StG-801954 “Wanted: Micronutrients! Phytosiderophore-mediated acquisition strategies in grass crops”). We thank Prof. Matthias Weil for X-ray diffraction experiments. The X-ray Centre of TU Wien is acknowledged for providing access to the X-ray diffractometer.

Conflict of interest

The authors declare no conflict of interest.

Keywords: micronutrients · mugineic acid · natural products · phytosiderophores · total synthesis

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Phytosiderophores are natural products enabling plants to absorb essential metal ions from the soil. Until now only specific solutions towards particular phytosiderophores had been reported. In this work, the entire family of these compounds is made synthetically accessible by using a divergent synthetic strategy.

Total Synthesis

N. Kratena, T. Gökler, L. Maltrovsky, E. Oburger, C. Stanetty*

A Unified Approach to Phytosiderophore Natural Products