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Chemical approaches for the enhancement of 5-aminolevulinic acid-based photodynamic therapy and photodiagnosis

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The administration of 5-aminolevulinic acid (ALA) to generate enhanced intracellular levels of endogenous porphyrins is currently one of the most important approaches for photodynamic therapy (PDT) and photodiagnosis (PDD). Despite the great promise of ALA-based PDT, the physicochemical behaviour and chemical reactivity of ALA are problematic, and a variety of chemical approaches have been brought to bear to improve cellular delivery, enhance porphyrin production, and generate ALA prodrugs that have appropriate stability for convenient clinical use, as well as selectivity for cancerous tissues. While there has been considerable success, there are still a number of challenges to be addressed and opportunities to be exploited through application of chemical insight in this area.

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Introduction

In the last 40 years, 5-aminolevulinic acid (ALA) has attracted tremendous interest in the field of PDT and photobiology.^{1–5} Alongside the emergence of clinical applications and approvals for ALA-PDT for the treatment of cancerous and precancerous

conditions, as well as in photodiagnosis (PDD) and fluorescence-guided surgery, it has become apparent that there is a real need for better ways to administer ALA and also to manipulate its bioconversion to protoporphyrin IX (PpIX) *via* the heme cycle (Scheme 1).^{6,7} The aim of this review is to cast a focus on some of the chemical strategies that have been developed to improve the performance of ALA in photo-medicine, and the opportunities that exist for future exploration by medicinal chemists.

ALA is an early intermediate in the heme biosynthetic pathway and is not a photosensitiser itself (Scheme 1).

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Kunal Tewari received his PhD from University of Bath, Bath, UK in 2016 under the supervision of Dr Ian Eggleston. His research involved development of novel strategies for enhancement of delivery of 5-aminolevulinic acid (5-ALA) to specific cell types using targeted ALA dendrimeric prodrugs. He is currently working as a research associate at the University of Strathclyde and his current focus is centred around biomaterials science, especially the development of peptoid nanosheets.



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Ian Eggleston received his D.Phil. in Chemistry in 1990 from the University of Oxford, UK, and carried out postdoctoral work at the University of Wisconsin–Madison (USA), CNRS-INSERM Montpellier (France), and the University of Lausanne. In 1997 he joined the University of Dundee as Lecturer in Synthetic Organic Chemistry and moved to the University of Bath in 2006, where he is now Reader in Medicinal Chemistry. His research interests include the development of peptide-targeted agents for photodynamic therapy and light-triggered drug delivery, especially the synthesis of novel peptide-based prodrugs of 5-aminolevulinic acid.

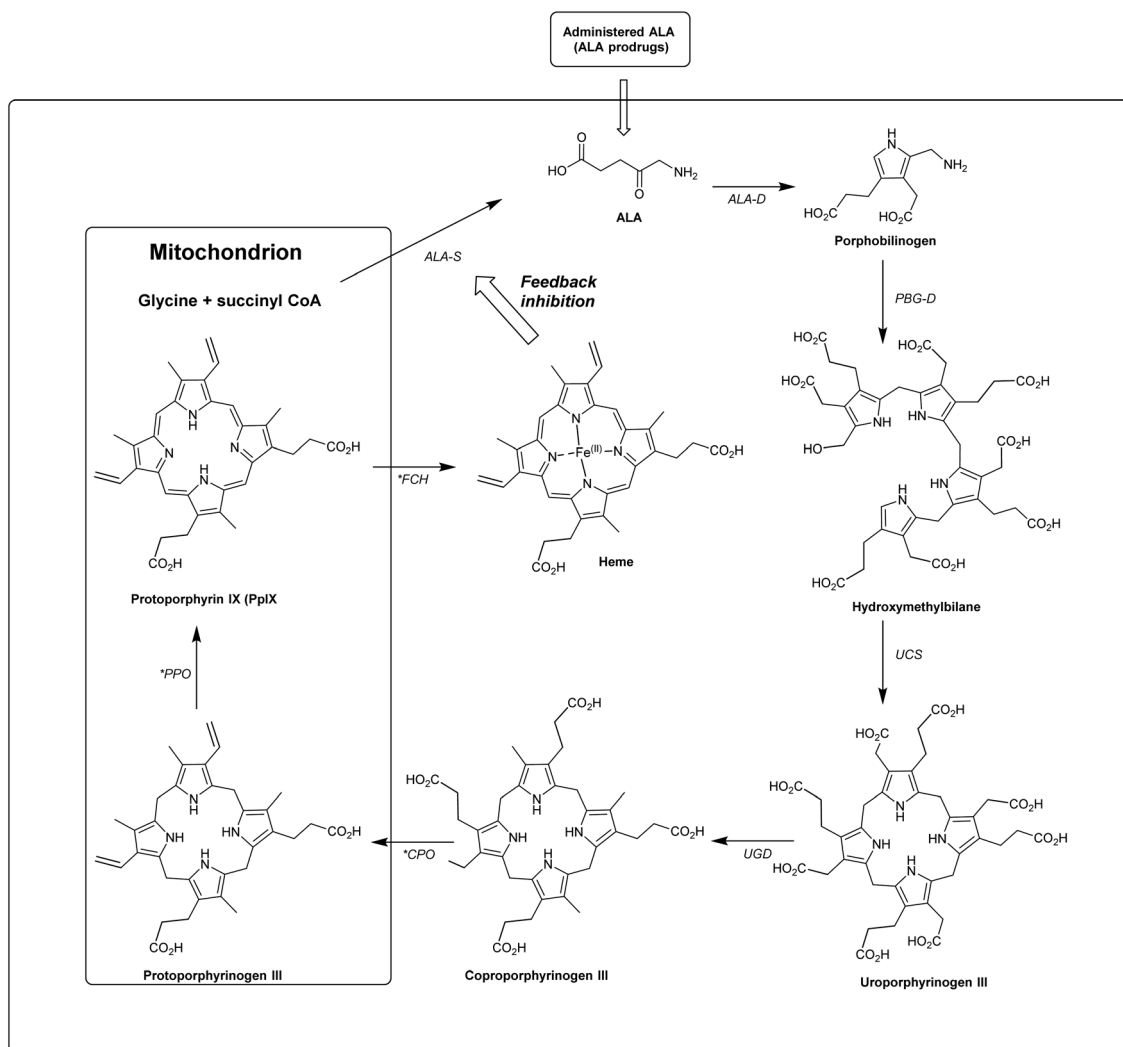




Scheme 1 5-Aminolevulinic acid (ALA) and its bioconversion to protoporphyrin IX (PpIX).

ALA-PDT and related techniques exploit instead the downstream generation of the active sensitizer protoporphyrin IX (PpIX), which is the penultimate species that is converted to heme. ALA can therefore be regarded as a prodrug for both PDT and photodiagnosis. The heme biosynthetic pathway is shown in detail in Scheme 2 and begins with a condensation

reaction between glycine and succinyl-CoA in the mitochondrion to form ALA which becomes the source of C and N atoms for the construction of the heme macrocycle. This reaction is catalysed by ALA synthase (**ALA-S**), and the synthesized ALA then enters the cytosol where a condensation reaction between two units of ALA, mediated by ALA dehydratase (**ALAD**), yields the pyrrole derivative, porphobilinogen (**PBG**). Four units of **PBG** are combined to generate a linear tetrapyrrole intermediate under the influence of porphobilinogen deaminase (**PBG-D**), followed by cyclisation by the enzyme uroporphyrinogen III cosynthase (**UCS**) to give the cyclic tetrapyrrole intermediate, uroporphyrinogen III. This intermediate now undergoes a decarboxylation of the four acetate side chains to methyl groups (catalysed by uroporphyrinogen decarboxylase, **UGD**) to give coproporphyrinogen III, which is then carried to the mitochondrion. Further oxidative steps catalysed by coproporphyrinogen oxidase (**CPO**) and protoporphyrinogen oxidase (**PPO**) then lead to formation of PpIX. Insertion of Fe^{2+}



Scheme 2 The heme biosynthetic pathway. 5-Aminolevulinic acid (ALA). Key steps that may be targeted chemically to maximise PpIX production are highlighted (*). Modified from Thunshelle *et al.*⁷



into PpIX, mediated by ferrochelatase (FC), ultimately completes the heme cycle, and under normal physiological conditions, the whole pathway and thus the production of porphyrins is tightly controlled by negative feedback by heme itself which thus controls ALA synthesis. In a clinical setting, the administration of exogenous ALA or a suitable derivative bypasses this control mechanism and the rate-limiting step in the pathway becomes the final insertion of iron into PpIX by ferrochelatase. The net result is the temporary overproduction and accumulation of excess PpIX in cells which then renders them photosensitive and susceptible to PDT. The preferential accumulation of PpIX in rapidly dividing tumour cells relative to healthy cells not only provides ALA-PDT with an inherent selectivity relative to PDT with other photosensitisers, but also presents the means to accurately image the location of tumour tissue for PDD. From a medicinal chemistry perspective, the challenge is to develop ways to optimise the amount of ALA that is available to enter into the heme cycle and its subsequent conversion into PpIX within selected tissues. This can be tackled in one of two ways: either by manipulating a specific enzyme-catalysed reaction in the heme cycle with small molecules in order to maximise PpIX production from administered ALA, or by enhancing the delivery of ALA across biological barriers to the cells of interest by incorporating ALA into derivatives with more suitable physicochemical and targeting properties and from which ALA is then released (ALA prodrugs).

Drawbacks of ALA-PDT/PDD and strategies to improve its effectiveness

ALA is a zwitterion at physiological pH, and as such its hydrophilic nature severely impairs its passage through biological barriers to ultimately reach and enter the cells of interest.³ This is a significant challenge for topical application of ALA in dermatology where penetration through the lipophilic stratum corneum^{7,8} must be achieved, and has motivated many chemical and non-chemical approaches to overcome issues of low penetration and non-homogeneous distribution in the targeted tissues. The bioavailability of ALA itself is also limited when administered parenterally, and has been associated with significant adverse effects.³

A variety of formulation-based chemical penetration enhancers have been investigated for topical ALA administration including, dimethyl sulfoxide (DMSO),⁹ 1-[2-(decylthio)-ethyl] azacyclopentan-2-one (HPE-101),¹⁰ glycerol monooleate,¹¹ 6-ketocholestanol (2% w/w)¹² and oleic acid.¹³ Recently, nanoemulsion-based formulations of ALA such as BF-200 ALA¹⁴ have begun to show very promising clinical applications, particularly for the treatment of actinic keratoses (AK), and can not only enhance penetration of ALA through the stratum corneum, but also can improve the long term stability of the prodrug.¹⁵

This highlights a further challenge for both the clinical application of ALA and also strategies for chemical modification. At physiological or alkaline pH, ALA and simple ester

derivatives (see below), are unstable with respect to self-condensation through formation of a dimeric Schiff base. Buffered solutions of ALA at physiological pH discolour even at low temperature due to decomposition of ALA, and thus stock solutions for clinical use would typically need to be freshly prepared.¹⁶ The instability of ALA under neutral and alkaline conditions has been investigated in some detail, and under alkaline conditions, it has been shown that ALA readily undergoes dimerization to form 2,5-dicarboxyethyl-3,6-dihydropyrazine (DHPY), which oxidises spontaneously to 2,5-dicarboxyethylpyrazine (PY, Scheme 3).^{17,18} Other transformations of the ALA dimer to form porphobilinogen (PBG) and pseudoporphobilinogen have also been proposed.^{18–20}

In addition to the drawbacks associated with the physical properties of ALA and its chemical stability, a further problem that needs to be addressed for both topical and systemic ALA delivery is to ensure an adequate discrimination between normal and diseased (*i.e.* tumour) cells. In this case, the challenge for the chemist is to design ALA derivatives which are either selectively targeted towards particular cell types, or which release ALA only in the right biological context in the presence of a suitable enzyme activity. Many of the chemical strategies that have been devised to enhance ALA-induced PpIX production do indeed rely on this kind of prodrug approach,⁶ and this therefore forms a major part of the rest of this review.

Alkyl ester prodrugs

The synthesis of ALA esters has so far proved to be the most effective and straightforward prodrug approach. The methyl ester of ALA (Metvix or MAL) **1a** is indeed now a mainstay for the treatment of AK by both conventional and daylight PDT, as well as being clinically approved for the treatment of basal cell carcinomas (BCC). The hexyl ester **1f** has been approved for fluorescence diagnosis in the bladder as Hexvix (now Cysview).⁷

For these clinically approved examples, and also for many other esters of ALA, it has been shown that their enhanced lipophilicity compared to ALA itself gives rise to better cellular permeability and thus quicker, more homogeneous, and more effective PpIX production in a variety of cell models. Many ALA ester prodrugs have been synthesized with linear or branched alkyl ester groups, and a selection of these are shown in Fig. 1.³ The preparation of such derivatives from the corresponding alcohols is straightforward and easily scaled-up to multi-gram quantities. Like ALA itself, simple esters have limited stability at neutral pH, although they may be stored in solution at pH 4 without significant degradation.²¹

Peng *et al.* were the first to report a comparative study on the effectiveness of a series of ALA esters against ALA. This showed that the methyl, ethyl and propyl esters **1a–c** produced more PpIX fluorescence in both murine and human tissues compared to free ALA.²² Similar observations were made by Kloek *et al.*^{23,24} and Gaullier *et al.*²⁵ with straight-chain ALA





Scheme 3 Instability of ALA at physiological pH and chemical conversion to non-heme precursors.

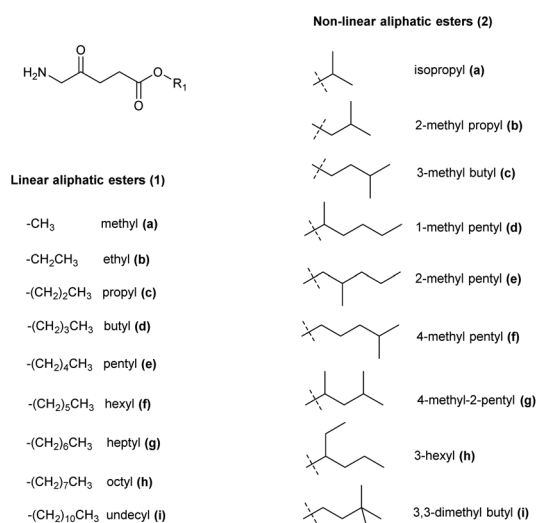


Fig. 1 Aliphatic ester prodrugs of ALA **1a–i** and **2a–i** with a linear or branched alkyl component (reviewed by Fotinos *et al.*³).

esters up to the octyl ester **1h**. Gaullier reported that for the lipophilic ALA esters **1f–h**, PpIX production was 30–150 times more efficient than for free ALA, thus suggesting faster cellular uptake of the prodrugs and ALA-induced PpIX formation, with the maximum benefit being observed for the hexyl ester **1f**. The improved performance of shorter chain esters compared to ALA upon topical administration to the skin is therefore more consistent with increased penetration through tissue towards the target cells, rather than enhanced cellular uptake as a result of a greater lipophilicity.²⁵ Several studies have confirmed that simply changing the lipophilicity of aliphatic ester prodrugs of ALA is not the only factor that must be considered

when seeking to optimise delivery and PpIX production. For example, Uehlinger *et al.*²⁶ noted that the hexyl ester **1f** produced an order of magnitude greater PpIX fluorescence than the corresponding cyclohexyl ester **3e** (see Fig. 2) even though both the conjugates had similar log *P* values. More recently, DiVenosa *et al.*²⁷ reported the evaluation of the undecyl ester **1i** of ALA which accumulates rapidly in LM3 breast cancer cells to generate PpIX, by virtue of its high lipophilicity, but is much less effective when applied topically to the skin due to slow diffusion across the stratum corneum.

The utility of ALA ester prodrugs of course depends critically upon the efficient release of ALA by cellular esterases upon internalisation. For most simple aliphatic esters, this is not problematic, however a comprehensive study by Whitaker *et al.* on the synthesis and *in vitro* biological evaluation of various linear and branched-ALA esters highlights the impact that ester structure may have on PpIX production from such prodrugs.²⁸ Here, significantly lower PpIX production was



Fig. 2 Ethylene glycol-based ester prodrugs of ALA **6a–f**.



observed in pancreatic cells for **2a**, **2d**, **2g** and **2h** compared to their linear counterparts which do not contain branching adjacent to the ester carbonyl group.

A range of ALA prodrugs have also been synthesized based on alicyclic or substituted benzyl esters which in some cases also display promising levels of PpIX compared to equimolar ALA (Fig. 3). Again, the need for a balance between lipophilicity and ease of ester hydrolysis is apparent among such compounds, and while they have not so far gained clinical approval, they provide further valuable indications to be inputted into the design of more elaborate ALA prodrug systems.

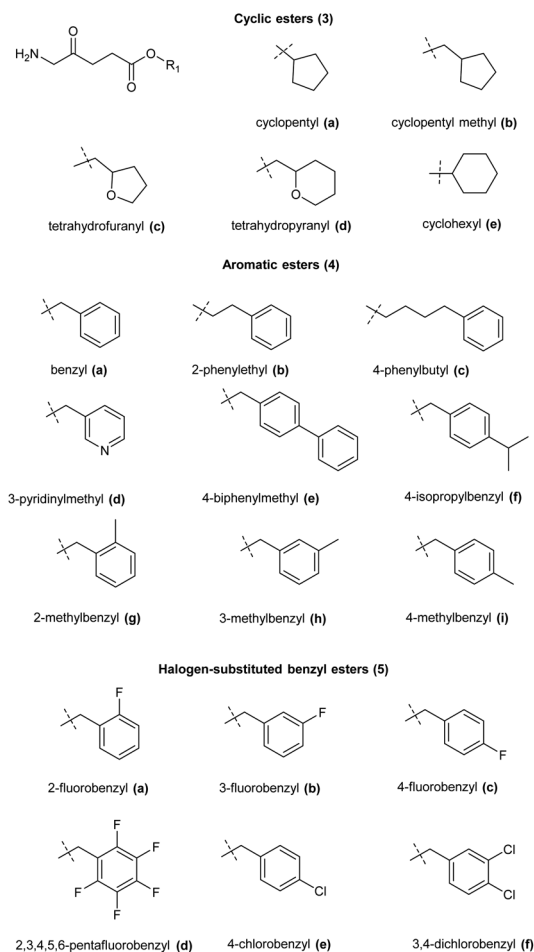


Fig. 3 Cyclic ester prodrugs of ALA **3a–e** and various substituted benzyl ester derivatives **4a–i** and **5a–f**.

For example, Kloek *et al.* reported the synthesis and biological evaluation of tetrahydrofuranyl **3c** and tetrahydropyranyl **3d** esters of ALA which both showed enhanced PpIX fluorescence compared to ALA *in vitro*.²³ Godal *et al.* reported the synthesis of the cyclohexyl ester of ALA **4e** along with a variety of substituted benzylic and arylalkyl esters **4a–f** of ALA, and biological evaluation *in vitro/in vivo* showed significantly enhanced PpIX fluorescence compared to ALA for aromatic esters, especially the benzyl and ring-alkylated benzyl esters **4a**, **4b** and **4g–i**. In contrast, the cyclohexyl ester **4e** produced the least PpIX fluorescence amongst all the prodrugs tested and was only as effective as ALA itself. Halogen-substituted benzyl esters **5a–f** showed intermediate PpIX fluorescence, and amongst them the mono-halogenated esters showed higher PpIX fluorescence than the di- or penta-substituted esters.²⁹

Although aliphatic ALA esters have been shown to induce higher PpIX production with an increase in carbon chain length, as already noted increasing ALA prodrug lipophilicity in this way can be counter-productive, bringing the limitations of slower tissue penetration as well as lower water solubility for esters which are derived from long-chain alcohols.²⁷ In contrast, poly(ethylene glycol) derivatives, which have been used for a large number of pharmaceutical applications, potentially offer the advantage of both improved water and lipid solubility, as well as giving rise to a benign promoeity upon ester hydrolysis.³⁰ To this end, Berger *et al.* reported the synthesis of ethylene glycol esters of ALA **6a–c** (Fig. 4) and tested their biological efficacy in human and rat cell lines of carcinoma and endothelial origin. It was found that a high PpIX fluorescence compared to free ALA was observed in endothelial cells in comparison to tumour cells, especially for long-chain derivatives.³¹ Godal *et al.* also reported the synthesis of ethylene glycol esters **6d** and **6e**, but biological evaluation *in vivo* showed that a lower PpIX fluorescence was produced compared to the simple hexyl ester **1f**.²⁹

As already noted, the use of ALA esters can result in increase PpIX production in various cell types, but with a greater enhancement in tumour cells. In this context, Brunner *et al.*³² observed that the simple hexyl ester **1f** and benzyl ester **4a** prodrugs which lack any specific targeting motif in their structures showed a significantly higher accumulation of PpIX in tumour cells compared to ALA, when applied at low concentrations and short incubation times. In a clinical setting, this means that appropriate PpIX levels for either PDT or PDD can be attained more rapidly and with less side effects. Preferential

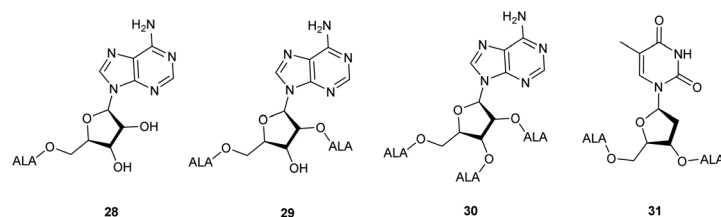


Fig. 4 ALA-nucleoside bioconjugate prodrug **28–31** bearing up to three ALA units.



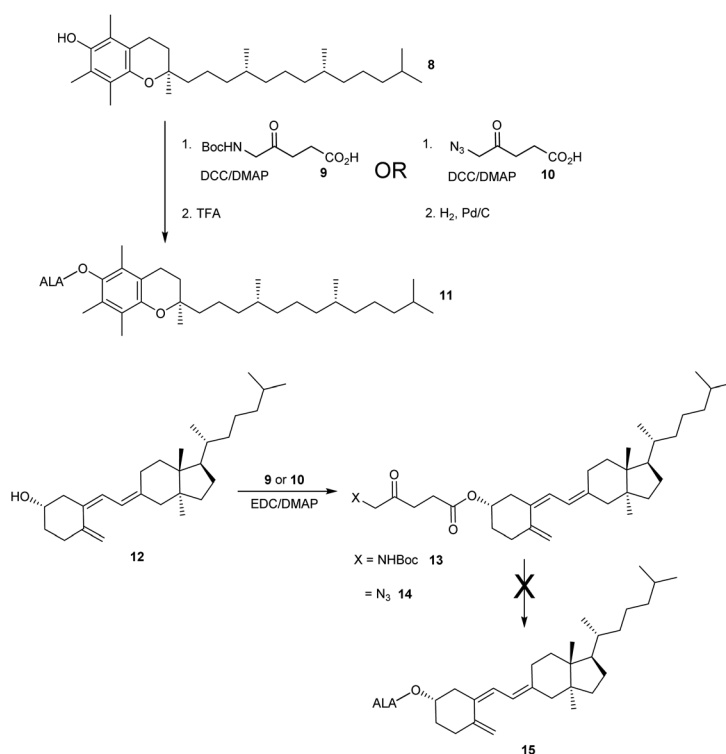
accumulation of ALA and its esters in tumour cells can also be correlated with the expression of specific uptake transporters³³ and this has provided a rationale for the development of prodrugs that contain structural features that only enhance physical properties but can provide more effective targeted delivery.

Biomolecule conjugates and polyvalent ALA ester derivatives

The esterification of ALA with various vitamins, monosaccharides and nucleoside derivatives has been explored both with a view to tailoring the physical properties of ALA and enhancing the transport of ALA into cells. For example, the attempted synthesis of novel ester bioconjugates of ALA with vitamin E (or α -tocopherol) **8** or vitamin D₃ **12** has been reported by Vallinayagam *et al.*³⁴ as shown in Scheme 4. Typically, more complex ester prodrugs of ALA such as the desired conjugates **11** and **15** are prepared using a *N*-alkoxycarbonyl-protected ALA derivative such as Boc-ALA **9** or the ALA surrogate, azidoleulinic acid **10**, with activation of the carboxylic acid function using Steglich-type esterification conditions with a carbodiimide reagent (DCC or EDC) and DMAP as catalyst. Both **9** and **10** are unable to undergo competing Schiff base chemistry during the key coupling step. For the molecules in Scheme 4, although vitamin E conjugate **11** was successfully prepared, the vitamin D conjugate **15** was unstable to final deprotection chemistry. The original rationale for the preparation of ester derivatives of ALA with these hydrophobic vitamins was to

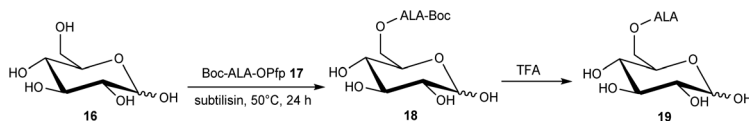
potentially facilitate the transport of ALA into cancer cells through the ease of incorporation of such bioconjugates into cell membranes.³⁵ In parallel, the enhancement of ALA-induced PpIX production by vitamin D has also been studied;³⁶ the observed enhancement of photodynamic cell death by vitamin D suggests that there may be potential for developing multifunctional ALA prodrugs of this type that can activate more than one cell-death pathway (see below).

The targeting of various photosensitisers with sugars has been very widely investigated as a means of enhancing selective accumulation in cancerous *vs.* normal cells. To this end, Vallinayagam *et al.*³⁷ reported the esterification of ALA with several simple monosaccharides. The presence of multiple hydroxyl functions in the ester component here presents an additional challenge for the chemist and required the development of distinct synthetic strategies to produce ALA conjugates selectively monoacylated on the 1-hydroxyl (anomeric) or 6-hydroxyl groups. As shown in Scheme 5, regioselective acylation of glucose **16** at the 6-position was achieved in moderate yield by enzyme-catalysed transesterification with the Boc-ALA derivative **17** using subtilisin, followed by acidic deprotection of the ALA component in **18** to give the desired conjugate **19**. Similar regioselective acylations could also be achieved with galactose **20** and mannose **24** and using porcine pancreatic lipase, again in reasonable yields. In order to effect esterification directly on the anomeric hydroxyl of these sugars, a multi-step orthogonal protection scheme was needed, using benzyl protection of the other hydroxyl groups, as outlined in Scheme 6 for the preparation of the galactose conjugate **23**. A

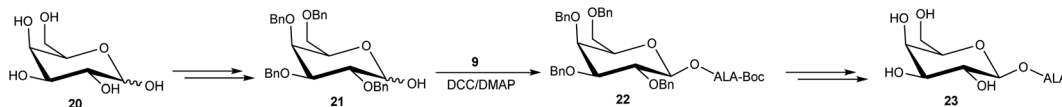


Scheme 4 Synthesis of ester bioconjugate prodrugs of ALA.





Scheme 5 Enzyme-mediated syntheses of a regioselectively 6'-O-acylated ALA-glucose bioconjugate prodrug **19**.



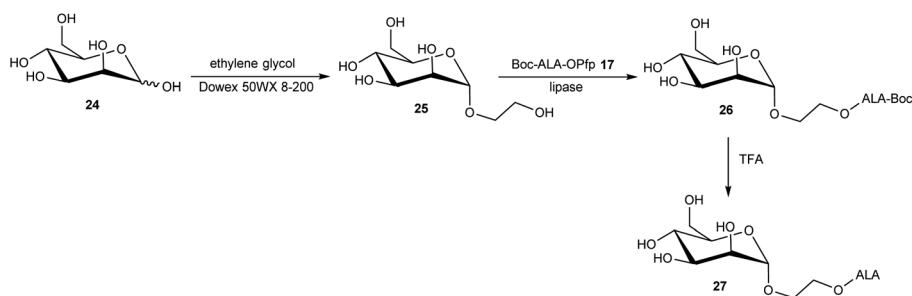
Scheme 6 Chemical synthesis of regioselectively acylated 1'-O-ALA-galactose bioconjugate prodrug **23** via an orthogonal protecting group strategy.

simpler, enzyme-mediated approach to 1-functionalised derivatives could however be achieved without additional hydroxyl protection by first derivatising the anomeric hydroxyl with an ethylene glycol spacer, as illustrated for the mannose derivative **27** (Scheme 7).

Evaluation of these prodrugs in human colon, breast and lung carcinoma cells and angiogenic endothelial cells revealed that PpIX production was similar to ALA itself, whereas in normal fibroblasts PpIX production from the prodrugs was very low. Surprisingly, there was little difference in PpIX production between the 1- or 6-acylated ALA esters such as **19** and **23**, and the insertion of a spacer as in **27** tended to decrease PpIX production. Furthermore, competition experiments with the parent sugars **16**, **20**, and **24**, or their 1-OMe derivatives had no effect on PpIX production, suggesting that such prodrugs do not appear to produce any cell-selectivity associated with sugar-mediated uptake mechanisms. It was concluded that prodrugs of this type may nonetheless be valuable for topical delivery of ALA to tumour cells with the reduced PpIX production in normal cells serving to spare them upon ALA-PDT. Moreover, the production of a harmless, alcohol component upon prodrug cleavage should again be a potential advantage for these or other prodrugs based upon natural sugars.

ALA esters with potential targeting properties have also been prepared by esterification of the hydroxyl functions of nucleoside derivatives.³⁸ A feature of these prodrugs, as with

other sugar-based derivatives, is the potential to produce derivatives with an increased payload of ALA compared to conventional esters, which is of interest since 8 copies of ALA must be delivered to the cell for bioconversion to one molecule of PpIX. Once again, prodrugs containing a single ALA unit, demand a chemical approach that allows for selective esterification of primary and secondary hydroxyl functions of the nucleoside component. To this end, an ALA prodrug of adenosine **28**, selectively monoacylated at the primary 5'-hydroxyl was prepared in modest yield directly from the parent nucleosides by Gurba *et al.* again utilising enzyme-mediated transesterification.³⁸ Attempted peracylation of the ribose component of adenosine using standard carbodiimide chemistry (DCC/DMAP) was less efficient and led to a mixture of di- and triacylated products from which di- and tri-ALA ester prodrugs **29** and **30** were subsequently obtained. In contrast, the deoxy-ribose unit of thymidine could be converted to the expected diester in excellent yield under similar conditions, providing prodrug **31** bearing two ALA units. Interestingly, evaluation of **28–31** in various cancer cell lines (human colon (SW480), breast (MCF7), lung (A549), ovarian (A2780), cervix (HeLa)) revealed that PpIX production did not clearly correlate with the number of copies of ALA present. Indeed, the adenosine conjugates **28–30** showed poor PpIX production compared to ALA in all the cell lines tested, suggesting that they may not be substrates for relevant adenosine receptors. Encouragingly, the thymidine derivative **31** did produce more PpIX (approximately



Scheme 7 Enzyme-mediated regioselective synthesis of ALA-bioconjugate prodrug **27** via introduction of a spacer unit at the anomeric position of the sugar.



1.5-fold) than equimolar ALA in most cell lines, which supports the idea that it may be possible to develop ALA prodrugs that carry multiple copies of ALA and which can be targeted to specific tumour cells.

Cyclodextrins are well-known nanocarriers of biological origin that present multiple hydroxyl functions for attachment of drug entities such as ALA. In a preliminary investigation by Aggelidou *et al.*,³⁹ beta-cyclodextrin which disposes seven

primary hydroxyl functions, could be esterified with up to three ALA units *via* the azido derivative **10**, using DCC/DMAP chemistry. The water-soluble ALA prodrug **32** that was obtained (Fig. 5) was found to effectively produce PpIX in MCF7 cells in comparison to equimolar ALA, which suggests that such compounds may have significant potential for the delivery of enhanced payloads of ALA, if effective functionalisation of all primary hydroxyls can be achieved.

The use of other polyvalent drug carriers such as dendrimers have shown significant promise for the delivery of high payloads of photosensitisers in PDT. This approach has been explored in some depth as a means of overcoming the poor systemic bioavailability of ALA in both PDT and PDD. Battah *et al.* in 2001 first reported the synthesis of a series of novel ALA-containing dendrimers bearing 6 to 18 ALA residues in which ALA residues are attached to the periphery of the dendrimer structure *via* ester linkages.⁴⁰ First-generation dendrimers **34–36** with 6 or 9 ALA units were prepared by attachment of a tris(Boc-ALA) derivative **33** to a di- or tripodal aromatic or tripodal aliphatic core *via* amide bond formation (Fig. 6). Second generation dendrimers **37** and **38** with 18 ALA units were then prepared by amplification of the building block (dendron) **33** and coupling to the tripodal aromatic cores used previously thus providing derivatives with an increased distance between the dendrimer core and the ALA units. This convergent growth approach has the advantage of ensuring the formation of prodrugs of defined size and loading, which is more challenging to achieve *via* a divergent strategy starting from a multifunctional core unit (see above), and potentially provides a route to ALA dendrimers of sufficient size to result

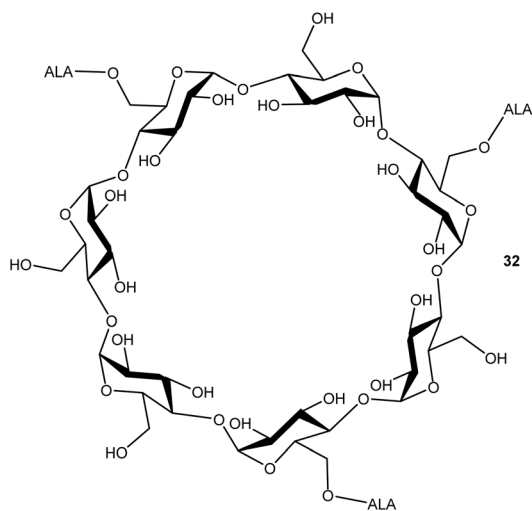


Fig. 5 Cyclodextrin-based trivalent ALA prodrug ester **32**. The conjugate obtained presents three ALA units which are presumably randomly distributed over the upper (primary hydroxyl) face of the scaffold.

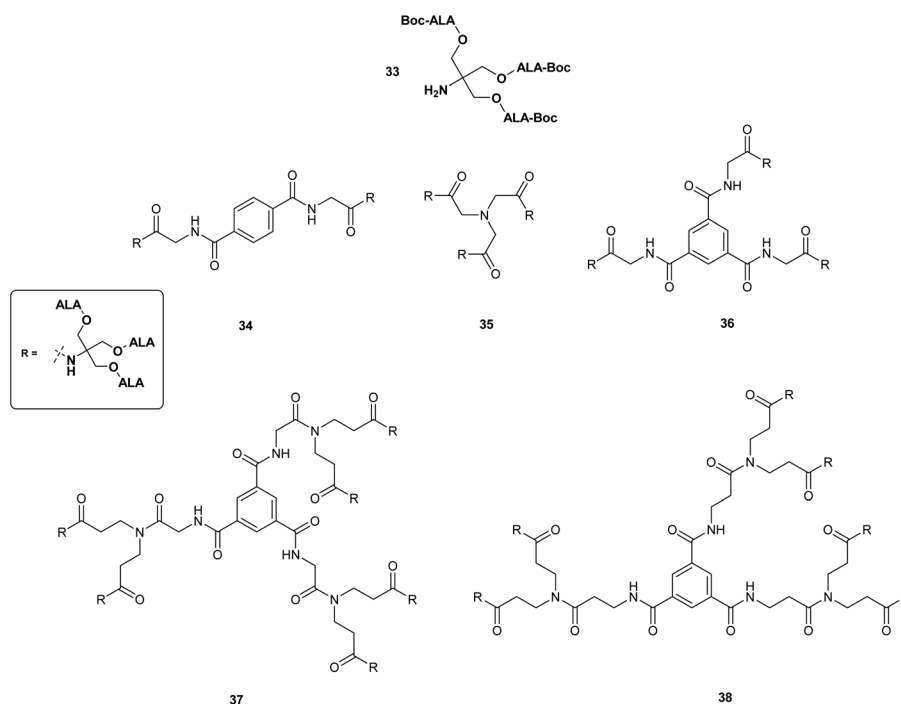


Fig. 6 First and second generation dendrimeric ALA prodrug esters **34–38**.



in enhanced permeability and retention in tumour tissue. Evaluation of first generation dendrimeric ester **35** and second generation derivative **37** in the in the keratinocyte cell line, PAM 212 at a concentration of 0.1 mM, revealed a significantly enhanced level of PpIX fluorescence after 4 h incubation compared to equimolar ALA. Although no direct correlation was observed between the number of ALA units and PpIX fluorescence, **37** appeared to have a greater than two-fold efficacy over **35**, with a correspondingly higher phototoxicity upon irradiation of cells. No dark toxicity was observed for either dendrimer at this concentration.

A further expanded second generation dendrimer **39** (Fig. 7) was subsequently prepared by MacRobert and coworkers and evaluated in detail in both PAM 212 keratinocyte and A431 human epidermoid cancer cell lines.⁴¹ This again confirmed the ability of such dendrimers to efficiently pass through cell membranes and release ALA intracellularly, with porphyrin production again being significantly higher than for equimolar ALA in both cell lines at low concentrations (0.01–0.1 mM). At higher prodrug concentrations, porphyrin production was significantly lower, possibly due to intermolecular Schiff base formation at physiological pH in culture medium as observed for most ALA ester derivatives with a free amino function.

Studies with low temperature incubation and endocytosis inhibitors suggested that cellular uptake of **39** is primarily *via* micropinocytosis and thus the improved performance of second generation derivatives **37** and **39** over first generation derivative **34** may reflect increased uptake of the larger dendrimers by endocytic processes. Uptake of **39** by endocytosis may also result in stabilisation against imine formation between ALA units due to protonation of their amino functions upon incorporation within acidic lysosomes. A further very important finding for **39** is that there is a marked difference in the kinetics of PpIX accumulation compared to ALA which ultimately may have significant importance for the application of these prodrugs, particularly for PDD. PpIX production for **39**



Fig. 7 Expanded second generation dendrimeric ALA prodrug ester **39** presenting 18 ALA units.

was found to be sustained for up to 24 h, at which point PpIX levels from ALA itself were significantly lower, consistent with the metabolism of porphyrin to heme within a few hours of incubation. The slower kinetics for **39** was proposed to result from gradual release of ALA by ester hydrolysis from the internalised prodrug which in turn is likely to be related to the steric accessibility of the multiple ALA units to intracellular esterases. It was indeed noted that the second generation dendrimer **37** which incorporates a shorter spacer structure between the ALA units and the prodrug core performed rather less effectively than **38** suggesting that the ALA ester units here are slightly less accessible. Complementary studies by the same researchers⁴² on three different ALA dendrons **40–42** each with three ALA units (Fig. 8) further support the importance of steric effects in determining the rate of ALA release from these systems and hence PpIX synthesis. Dendron derivative **41** for instance which incorporates a longer, propyl linker to the quaternary carbon, was found to be the most effective, with the ALA ester units here again being relatively more accessible for enzyme-mediated hydrolysis. **41** was also studied for *in vitro* efficacy in LM3 lung carcinoma cells and was found to produce similar porphyrin levels compared to free ALA.⁴³ It was also found to be effective *in vivo* when administered either topically or systemically to tumour-bearing mice with higher peak porphyrin levels being observed compared to the hexyl and methyl esters of ALA, although it was inferior to ALA itself. This may be due to retention of **41** within the stratum corneum at the application site when applied topically, as this triester conjugate is fairly lipophilic compared to ALA, which highlights again the fine balance that needs to be achieved in the design of ALA prodrugs with optimal physicochemical properties for both cell uptake and tissue penetration.

The sustained and enhanced PpIX production observed with the 18 ALA dendron **39** in *in vitro* studies was also confirmed by *in vivo* evaluation in a murine tumour model. Here, **39** showed a sustained release of PpIX for over 24 h, and the basal values were not reached until 48 h post-intraperitoneal administration of **39**.⁴⁴ In contrast, PpIX production for ALA was observed for only 3–4 h. While these results indicate that the kinetics of porphyrin production is a function of the rate of ester hydrolysis in **39**, it is also of significance that the integrated porphyrin accumulation for **39** and ALA at equivalent molar ratios was observed to be comparable for the 48 h time course. This demonstrates that the release of the ALA unit from **39** was essentially complete and the total porphyrin production is comparable – an essential feature to be achieved



Fig. 8 ALA dendron ester derivatives with varying structural features that may lead to altered rates of ALA release upon cleavage by esterases.



Diverse iron chelators such as EDTA, desferrioxamine (DFO, 45), and dexrazoxane all shown potential in preclinical studies for increasing PpIX levels for PDT upon ALA administration.⁷ EDTA is however, a non-selective chelator which binds other biologically relevant metal ions as well as Fe²⁺. More significantly, it has a markedly lower affinity for iron compared to DFO. DFO 45 (Fig. 10) is a hexadentate siderophore which has been shown to have antiproliferative activities against various tumour cell lines,⁵⁷ and has been extensively studied for use in the treatment of iron overload disorders.⁵⁸ However, due to its hydrophilic nature it is unsuitable for topical application, and since prolonged exposure to strong iron chelators such as DFO 45 may lead to iron starvation within healthy tissue,⁵⁹ its suitability for use in ALA-PDT or PDD is somewhat uncertain.^{60,61}

One compound that has shown particular promise for use in ALA-PDT is 1,2-diethyl-3-hydroxypyridin-4-one, more commonly known as CP94 46 (Fig. 11). The low molecular weight and greater lipophilicity of CP94 compared to both DFO and EDTA, make it particularly effective for chelating intracellular iron. CP94 has high affinity for Fe³⁺ and as such the chemical basis for its application to enhance ALA-PDT lies in its ability to promote the oxidation of Fe²⁺ to Fe³⁺ at low concentration by molecular oxygen.⁶² Following the initial observation of the efficacy of combined ALA and CP94 treatment,⁶³ many other studies have demonstrated the effectiveness of ALA and CP94 both *in vitro* and *in vivo* for PDT, as well as potential applications in fluorescence-guided surgery.^{64,65} For example, Blake *et al.*⁶⁶ investigated the combination of CP94 with ALA and MAL 1a in human glioma cells, and observed that CP94 significantly enhanced PpIX accumulation and subsequent PDT cytotoxicity. This combination may therefore be useful both in photodiagnosis and /or PpIX-induced PDT treatment of glioblastoma, where prognosis is typically poor. The combination of ALA or MAL and CP94 has also been shown to be safe and effective in clinical pilot studies for the topical PDT of nodular basal cell carcinomas.^{67,68}

The difference in pharmacological properties between ALA and a lipophilic iron chelator such as CP94, or indeed any modulator of heme synthesis, is a potential challenge that must be addressed upon coadministration. Zhou and co-

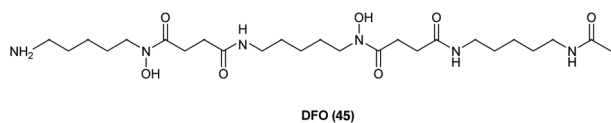


Fig. 10 Desferrioxamine (DFO) 45. A potent iron chelator that has been studied for enhancement of PpIX production in ALA-PDT.

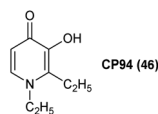


Fig. 11 Structure of CP94. A clinically validated iron chelator for ALA-PDT.



Fig. 12 General structure of ALA-HPO conjugates.

workers⁶² have sought to address this problem *via* the innovative approach of incorporating the structure of hydroxypyridone-(HPO)-type iron chelator into the ester moiety of an ALA prodrug. Conjugates of this type (Fig. 12) are more lipophilic than ALA itself and as predicted were able to efficiently deliver and release both ALA and HPO simultaneously into MCF-7 and MCF-7R cells with a corresponding enhancement in PpIX levels compared to ALA alone, or separate coadministration of ALA and HPOs. The best performing derivatives embodied a relatively long, flexible aliphatic linker in the ester unit between ALA and the bulkier HPO function, suggesting effective uptake by passive diffusion as well as the absence of any steric effects on cleavage by cellular esterases. These compounds ($n = 6-12$) were found to produce higher levels of PpIX fluorescence than the benchmark ALA hexyl ester.

A very recent report by Anayo *et al.*⁶⁹ has also revealed that a related combined ALA prodrug and iron chelator, **AP2-18**, is able to outperform the coadministration of ALA or MAL and CP94. The authors suggest that this novel patented ALA prodrug has the potential to address the challenge of delivering greater PpIX accumulation to the base of thicker nodular basal cell carcinoma (nBCC) tumours that are typically encountered clinically, and so may be a valuable new tool for dermatological PDT.

Studies by Zhou *et al.* have also revealed the potential for linking a HPO to the N-terminus of an ALA derivative,⁷⁰ while Zhu *et al.* have described ALA-HPO ester conjugates where the N-terminus of ALA is acylated (see following section).⁷¹ Once again, examples of such conjugates have been shown to be superior to ALA itself for PpIX production in a variety of tumour cell lines, with levels of intracellular porphyrin fluorescence correlating well with cellular phototoxicity upon irradiation.

Finally, Mrozek-Wilczkiewicz *et al.*⁷² have reported the concept of enhancing ALA-PDT using thiosemicarbazone (TSC) iron chelators, which have been previously developed as stand-alone anticancer agents (Fig. 13). Some of the compounds synthesised show exceptional (sub-nanomolar) dark cytotoxicity in HCT 116 colorectal carcinoma cells with a wild-type (+/+) or deleted (-/-) tumour suppressor TP53 gene. Although some synergistic effects on photocytotoxicity were observed with X and Y when used in combination with ALA-PDT, the sub-nanomolar concentrations of chelators used do not appear to be consistent with an enhancement of PpIX levels through simple sequestration of iron. A possible explanation for the effectiveness of the ALA-PDT-TSC combination may therefore be rather the result of two direct cytotoxic effects; one stemming from the PDT effect of the PpIX generated, the other due to ROS





Fig. 13 Novel thiosemicarbazones studied for synergistic iron chelation and ALA-PDT.

produced from the TSC-iron complexes *via* the Fenton reaction. This suggests an interesting new chemical approach for improved ALA-PDT that might be exploited further through the synthesis of novel combined ALA-TSC prodrugs.

N-Acylated ALA derivatives and ALA peptide prodrugs

A very obvious way to address the chemical stability issues that are associated with ALA and clinically used ester derivatives is to mask the primary amino function by the formation of an amide derivative. While this is attractive in principle, the effectiveness of such ALA prodrugs then depends upon the expression of a suitable protease activity in the targeted cells to liberate ALA. Indeed, while a number of simple *N*-acylated ALA derivatives have been reported, such as the formyl or acetyl derivatives **51** and **52** (Fig. 14), which are stable at physiological pH and also more lipophilic than ALA, *in vitro*

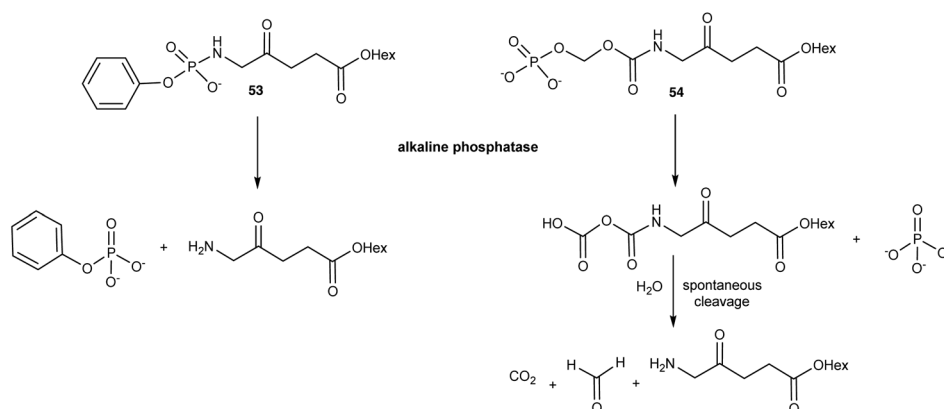


Fig. 14 Structures of formyl and acetyl ALA derivatives.

studies have in general revealed very low PpIX production, consistent with inefficient release of ALA.^{3,73}

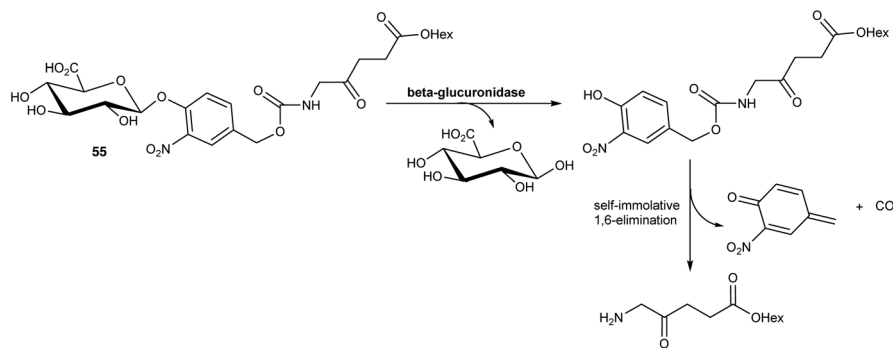
In response to the challenges posed by simple *N*-acylated ALA derivatives, Babic *et al.*⁷⁴ have developed ALA prodrugs in which the amino group of ALA is masked either by conversion to a phosphoramido group, or a novel amide where the pro-moiety contains a phosphate linkage that provides a trigger for cleavage and ALA release (Scheme 8). Both the compounds **53** and **54** were found to be chemically stable at acidic, basic, and physiological pH, however they were efficiently cleaved *in vitro* by alkaline phosphatase to liberate the hexyl ester of ALA. The activation of **53** and **54** is clearly structure-dependent, with complete conversion of **54** occurring in around 60 min, approximately ten times faster than for the phosphoramido derivative **53**. This provides the possibility to fine-tune the rate of ALA release from such compounds *in vivo*. To this end, activation by alkaline phosphatases is very attractive since they are ubiquitous enzymes that are present in all tissues and over-expressed in various tumours. This not only suggests a possible application for these prodrugs for selective PDT, but also PDD. Both compounds were shown to produce higher levels of PpIX as well as reduced toxicity compared to **1f** in U87MG glioblastoma cells over a 24 h time course, notwithstanding the slower rate of activation of **53**. Their clinical potential was further validated *in vivo* in a U87MG glioblastoma spheroid tumour model in chick embryos. Additional studies⁷⁵ with these prodrugs in a panel of different cancer cell lines revealed the level of PpIX production varied significantly, as might be expected based on known levels of phosphatase expression. As well as U87MG, the prodrugs performed best in PC3 and MCF7 cells, which points to their potential for future clinical development for applications in glioblastoma, prostate and breast cancer respectively.

In a further application of this approach, Babic and co-workers⁷⁶ have reported a prodrug of ALA **55** in which the amino function is masked by an acyl group that is susceptible to cleavage induced by β -glucuronidase activity. Such ALA prodrugs could again represent promising tools for tumour detection as elevated expression of β -glucuronidase activity has



Scheme 8 Bioconversion of phosphatase-sensitive ALA prodrugs **53** and **54** to release the hexyl ester of ALA **1f**.





Scheme 9 Cleavage of ALA prodrug **55** by β -glucuronidase activity.

been associated with various cancer types, including lung, breast and ovarian cancers. In this case, cleavage of the glucuronide unit from the *N*-acyl moiety of **55** triggers the release of the hexyl ester of ALA *via* a self-immolative 1,6-elimination process (Scheme 9). Preliminary biological evaluation showed that this prodrug produced similar levels of fluorescence as ALA-Hex in breast cancer MCF7 cell, but with much lower dark toxicity.

Peptide derivatives of ALA have also been synthesized to provide prodrugs with improved physicochemical properties relative to ALA itself. This approach also provides the possibility of achieving selective ALA release in particular cell lines which express proteases for which the prodrug is a potential substrate. Berger *et al.* investigated two types of ALA peptide prodrugs (Fig. 15) and evaluated them for their ability to produce PpIX.^{31,77} The first approach involved synthesizing acidic, basic and neutral ALA dipeptide derivatives **56a–e** as potential substrates for cellular aminopeptidases APA, APB or APN/M.³¹ Biological evaluation in endothelial and carcinoma cell lines A549, HCEC and EC219 showed that prodrugs containing a neutral amino acid sidechain such as **56c** induced the highest PpIX production, however a basic amino acid derivative (**56d**) or acidic derivative (**56e**) did not show any favourable results (minimal PpIX production). The second approach involved the synthesis and evaluation of pseudo tri- and tetrapeptide derivatives **57a–f** (Gly-Gly and Gly-Pro) of ALA as potential substrates for cellular di/tripeptidases or peptide transporters at the cell surface, to target human endothelial cells or carcinoma cells in cancer.⁷⁷ In this case, studies in human endothelial (HCEC) and human lung carcinoma cells

(A549) showed the peptide derivatives of ALA to be substrates for Gly/Pro-specific cellular transport systems and enzymatic activities. The dipeptide derivatives were also less cytotoxic in the absence of light compared to the dipeptide derivatives **56a–e**.

A significant feature in the synthetic approaches used by Berger *et al.* was that an excess of the activated amino acid components or ALA derivative was used for the formation of the critical amino acid-ALA peptide bond, presumably to compensate for the instability of ALA under the basic coupling conditions. Rogers *et al.* developed an alternative approach by employing only one equivalent each of an acylated amino acid succinimido ester and ALA-HCl.⁷⁸ The instability of ALA under basic conditions was avoided by slow neutralisation of ALA-HCl in the presence of the active esters (Scheme 6). The intermediate Boc-protected peptides were also used to synthesise the corresponding *N*-acetyl derivatives, thereby avoiding any potential epimerisation in the coupling reaction (Scheme 10).

Simple ALA peptide prodrugs of the general structure shown above embody a number of desirable features for improved topical and systemic ALA administration. As has already been noted, acylation of the amino function of ALA provides chemical stability, while esterification of the carboxyl function increases lipophilicity for enhanced membrane permeability. Compounds of this type can be termed “double prodrugs of ALA”, a term recently coined by Babic and co-workers.⁷⁶ There is also considerable scope to tailor the physicochemical properties of the prodrugs through careful choice of the amino acid residue to provide compounds that are overall more lipophilic than ALA, yet are still water soluble. Unlike other peptide prodrugs reported by Berger *et al.*,^{31,77} and also the phosphatase-sensitive prodrugs of Babic,⁷⁴ subject to the choice of a suitable amino acid residue the compounds are not charged at physiological pH.

Bourré *et al.*⁷⁹ described the biological evaluation of the dipeptide prodrugs **55** and **56** with the ALA amino group blocked with either Ac-L-Phe or Ac-D-Phe respectively, as well as the corresponding L-amino acid containing derivative **57** with a free carboxylic acid function (Fig. 16). Fluorescence spectroscopy revealed both **55** and **57** to be efficiently taken up in

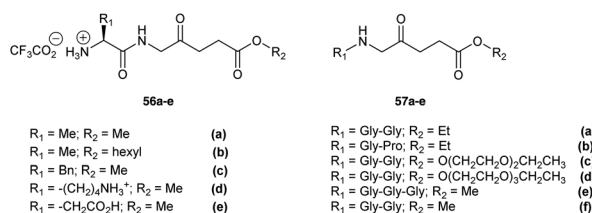
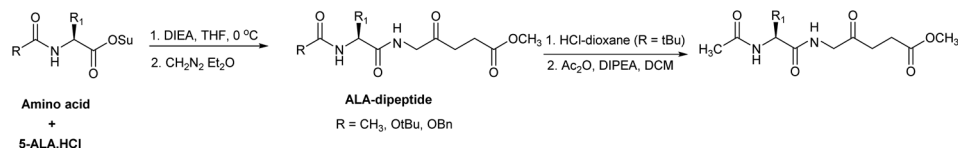


Fig. 15 Peptide prodrugs of ALA prepared by Berger *et al.*^{31,77}





Scheme 10 Synthesis of ALA peptide derivatives by Rogers *et al.* avoiding self-condensation of ALA during the coupling step.

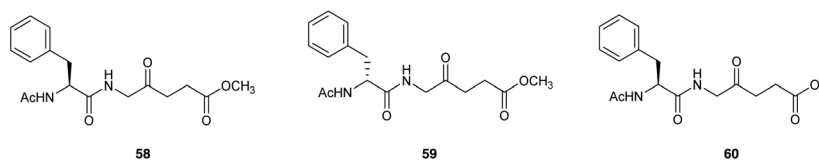


Fig. 16 Phenylalanine-derived ALA dipeptide derivatives **58–60**.

transformed PAM212 keratinocytes and generated greater levels of PpIX compared to equimolar ALA, although as anticipated better results were obtained with the neutral prodrug **58** (methyl ester) compared to **60** (free carboxy terminus). Significantly, no PpIX production was observed with the D-amino acid containing prodrug **59** consistent with the lack of a suitable intracellular enzyme with D-peptidase activity. All three dipeptide derivatives were stable in solution at physiological pH, in contrast to the corresponding dipeptides with a free N-terminus, presumably due to intermolecular Schiff base formation between the terminal amino acid amino function and the carbonyl function of an ALA residue, or possibly intramolecular cyclisation to generate a reactive six-membered Schiff base intermediate. PDT studies with both **58** and **60** in cells were in agreement with the results obtained from fluorescence pharmacokinetics, and the cell survival was significantly reduced in comparison to equimolar doses of ALA. **58** was also evaluated in a pig skin explant model in comparison to ALA and was found to produce higher porphyrin fluorescence by a factor of three demonstrating that **58** was capable of being transported into the skin and metabolised into PpIX in an *ex vivo* setting.

Eggleston and coworkers⁸⁰ subsequently reported a series of 27 dipeptide prodrugs of ALA with the same general structure Ac-Xaa-ALA-OR (*i.e.* analogues of **58** and **59**) where Xaa was an L- or D- α -amino acid providing a varying level of lipophilicity and water solubility to the prodrug. The method of Rogers *et al.*⁷⁸ was simplified by simply slowly neutralising HCl-ALA-Me or other ALA esters in the presence of the appropriate Boc or Fmoc-protected amino acid active esters, to give an intermediate dipeptide ester which could then be converted to the required acetyl derivatives in excellent yield and on a gram scale. A significant observation from this study was that there was no clear correlation as might have been expected between the lipophilicity of the prodrugs and the level of PpIX production in PAM212 keratinocytes. Quantification of the amount of ALA internalised using an improved fluorescence-based method⁸¹ revealed that in fact all the ALA prodrugs were taken up efficiently compared to ALA itself, by a combination

of passive and active transport mechanisms, but predictably only the L-amino acid derivatives produced PpIX. This clearly demonstrated not only potential for designing peptide prodrugs of ALA which are cleaved by a disease-specific/cell-line specific enzyme activity, but also the importance of being able to independently assess PpIX production and ALA uptake when attempting to rationally design ALA prodrugs. In this study, further variation of the ester component of the lead L-Phe derivative to fine-tune lipophilicity also led to the identification of a dipeptide double prodrug whose photocytotoxicity was comparable to that of the hexyl ester of ALA **1f** at concentrations where ALA itself was completely ineffective.

In a further study by the same researchers,⁸² it was shown that compounds of this general structure which contain an L-amino acid are in fact substrates for the enzyme acylpeptide hydrolase (APEH), a member of the prolyl oligopeptidase family of serine proteases, and for which D-amino acid derivatives are not substrates, as is also the case for compounds where the terminal acetyl function is replaced by a benzyloxy-carbonyl group.⁷⁹ This corroborated the earlier finding that PpIX production is also limited for these compounds in cell lines with a low expression of this enzyme such as Caco-2 and AF549.⁸⁰ Further *in vivo* studies with these compounds has provided further evidence for their potential use in topical treatment of basal cell carcinomas and their effectiveness upon systemic administration compared to ALA or its hexyl ester **1f**.^{83,84}

Some similar dipeptide-based double prodrugs of ALA have recently been reported by Chen and coworkers,⁸⁵ again exploring the variation of the ester function. This theme has also been investigated in a somewhat different way by Zhu *et al.*,⁷¹ where two of the lead dipeptide motifs identified by Eggleston and coworkers⁷⁸ were esterified with an HPO iron chelator. Zhou *et al.*⁷⁰ on the other hand, have modified the same lead dipeptides by acylation of the N-terminal amino acid residue with a suitably functionalised HPO. In both cases, codelivery of ALA and an iron chelator in the form of a dipeptide prodrug could be shown to be more effective than either ALA alone, or coadministration of ALA with CP94 as a stand-alone chelator.



Several studies have reported the synthesis and biological evaluation of more complex ALA peptide prodrugs which have been designed to provide improved cellular delivery and also highly selective ALA release. For example, Dixon *et al.* reported the synthesis of a 9-residue ALA-containing cell-penetrating peptide (CPP) conjugate **61** using Fmoc-solid phase chemistry (SPPS).⁸⁶ The synthesized peptide was based on a heptapeptide fragment of the cell penetrating peptide, penetratin (Fig. 17), and in this proof of concept study for CPP-mediated ALA delivery, it was possible to demonstrate effective uptake in PAM212 cells and intracellular conversion of the peptide to PpIX. This was the first example of the synthesis of an ALA-containing peptide by SPPS, and incorporation of the ALA residue was achieved by attachment of the preformed dipeptide, Fmoc-Leu-ALA-OH, thereby avoiding the potentially problematic acylation of ALA itself upon the solid phase. Use of a hindered 2-chlorotrityl resin support also helped prevent any potential intramolecular Schiff base formation upon acylation of the Leu-ALA dipeptide unit.

Abd-Elgaliel *et al.*⁸⁷ have adopted the same chemical strategy for the SPPS of a cathepsin E-cleavable ALA peptide prodrug **62** (Fig. 18). Biological evaluation showed that ALA was selectively liberated in cathepsin E-positive pancreatic tumour cells, but not in normal pancreatic tissue, thus allowing delivery of selective photodynamic therapy (PDT) to cancerous tissues.



Fig. 17 A cell-penetrating peptide-ALA conjugate based on the penetratin CPP sequence.⁸⁶

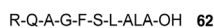


Fig. 18 A cathepsin E-activatable ALA peptide derivative.⁸⁷

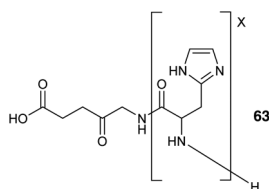


Fig. 19 General structure of ALA-oligo(L-His) prodrugs **63**.⁸⁸

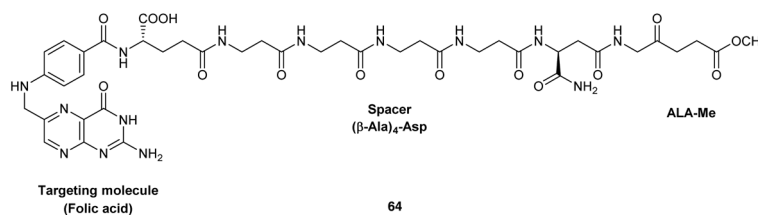


Fig. 20 A potential folic acid-targeted ALA prodrug.⁸⁹

Other peptide-based approaches for ALA delivery have also focused on the tagging of the amino function of ALA with an oligopeptide carrier. One notable example is the use of oligo(L-His) peptides which may be employed to provide tailored lipophilicity and water solubility to an attached cargo, and at physiological pH the oligo-His motif is able to act as a proton receptor and facilitate entry into tumour cells. In a study by Johnson *et al.*,⁸⁸ a series of pH-sensitive ALA-oligo(L-His) prodrugs **63** were prepared by ring opening polymerization of 1-benzyl-N-carboxy-L-histidine anhydride initiated by the amino function of ALA itself. The activity of these prodrugs was assessed in the human colon cancer HCT116 cell line revealing promising phototoxicity and selectivity compared to ALA, especially for the conjugate with 15 L-His residues (Fig. 19).

Finally, the synthesis of a novel folic acid-targeted ALA conjugate has been recently described by Guaragna *et al.*⁸⁹ Targeting of folate receptors has been frequently exploited for the delivery of anticancer drugs, and in this example the methyl ester of ALA was conjugated to folic acid *via* a peptide spacer consisting of four consecutive β -Ala units linked to the β -carboxylic acid function of the Asp residue of folic acid. The spacer was included to ensure adequate water solubility and cellular stability of the final conjugate **64**, and the whole carrier unit was prepared *via* SPPS on Rink amide resin, with an α -linked Asp residue at the C-terminus. This then left the β -acid function of the Asp residue free for acylation of the amino function of ALA (see Fig. 20). No biological and stability studies have yet been reported for this interesting targeted ALA derivative which once more illustrates the possibility of linking larger synthetic peptides to ALA.

Applications of nanotechnology and nanochemistry to the design of ALA prodrugs

The development of ALA dendrimers have already been described in detail as a promising strategy for the systemic delivery of a clinically useful payload of ALA for either PDT or PDD. In recent years, a growing number of other strategies have been explored that draw upon the knowledge gained from the synthesis and biological properties of simpler ALA esters and peptide derivatives to connect the challenge of ALA delivery to the field of nanotechnology.⁷ Notable examples



include the encapsulation of ALA into polymeric⁹⁰ or gold nanoparticles,⁹¹ and most recently the preparation of ALA-squalene nanoassemblies *via* esterification of ALA itself with squalene alcohol.⁹² Self-assembly of ALA squalene esters in water produces monodisperse systems with average diameter 70 nm, and a high ALA drug loading of 26%. These nanoassemblies show very promising PpIX production in human prostate PC3 and U87MG glioblastoma cells which is superior to that of the hexyl ester of ALA suggesting that they have great potential for PDD and tumour-selective PDT.

Several examples of peptide-targeted ALA nanosystems have also been recently reported that provide for both efficient targeting and context-dependent (*i.e.* pH-selective) ALA release.^{93,94}

Summary

Since the discovery of the clinical potential of ALA-mediated PDT considerable progress has been made in developing simple ALA derivatives that have improved bioavailability and stability compared to ALA itself, and also in identifying compounds that may be administered alongside ALA to maximise the production of PpIX in PDT and PDD applications. The methyl ester of ALA is now a mainstay for topical ALA-PDT in the clinic, while the hexyl ester is widely accepted for use in PDD. Challenges that remain to be solved are the development of effective platforms for systemic delivery of ALA, and also precise tumour targeting, particularly for applications in diagnosis and surgery. There are many opportunities for chemists to tackle these issues, drawing on the lessons from the wealth of knowledge already gathered about ALA and its derivatives and the application of modern methods of synthetic chemistry.

Conflicts of interest

There are no conflicts to declare.

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