Antisense oligonucleotides are now entering the clinic for hard-to-treat diseases. New chemical modifications are urgently required to enhance their drug-like properties. We combine amide coupling with standard oligonucleotide synthesis to assemble backbone chimera gapmers that trigger an efficient RNase H response while improving serum life time and cellular uptake.

Antisense oligonucleotides (ASOs) are short (~20mer) chemically modified oligomers that bind to their complementary RNA targets to modulate gene expression at the mRNA level. Thus, ASOs can target proteins that are considered undruggable through conventional approaches. As such, they hold enormous promise for hard-to-treat diseases as evidenced by a number of recently approved oligonucleotide (ON)-based drugs.

Chemical modifications are essential to improve the serum stability and pharmacodynamic properties of ASOs, as unmodified ONs are rapidly digested by nucleases in vivo and suffer from poor cellular uptake and tissue distribution. Whilst there have been considerable advances to modify ONs either at the nucleobase, sugar or backbone, a set of distinct chemical modifications to confer ideal drug-like properties has not yet been achieved. Commonly used ribose modifications include 2′-F, 2′-OMe, 2′-O- (2-methoxethyl) and locked nucleic acids, all of which have been shown to improve target affinity and serum stability. The most commonly used phosphodiester (PO) mimic is the phosphorothioate (PS) linkage which is compatible with ribonuclease H (RNase H) activation, a mechanism resulting in degradation of an mRNA upon formation of an ASO:mRNA heteroduplex. PS linkages also induce RNA target degradation and we evaluate serum stability and pharmacodynamic properties through interactions with plasma proteins. However, unspecific protein binding can contribute to the toxic potential of PS-ASOs and the PS linkage is P-chiral resulting in a mixture of diastereomers (more than half a million isomers in Mipomersen). Moreover, inefficient cellular uptake remains a major challenge for ASO therapeutics. Therefore, the investigation of other artificial backbone linkages is urgently needed.

Charge-neutral backbone modifications represent an interesting class of PO mimics. Among those, (thio)phosphonooacetate esters, phosphotriesters and alkyl phosphonates can enhance cellular uptake by eliminating the PO negative charges. Moreover, incorporation of a single methylphosphonate can eliminate hepatotoxicity of PS-ASOs. Phosphorodiamidate morpholino oligomers (PMOs) combine backbone and sugar surrogates. Within DNA, isolated amides can slightly increase duplex stability with target RNA while consecutive amides induce RNA target degradation and we evaluate serum stability and cellular uptake of a partially uncharged AM-gapmer ON.

Here we report the introduction of isolated and consecutive amide linkages into ONs for RNase H-based antisense applications. We discuss possible ASO designs that contain the AM backbone to induce RNA target degradation and we evaluate serum stability and cellular uptake of a partially charged AM-gapmer ON.

Isolated AM internucleoside linkages were introduced via dinucleotide phosphoramidite 6 (Fig. 1A) with an internal amide as the building block for standard ON synthesis. Preparation of phosphoramidite 6 required 5′-tritylation of monomer 1 followed by ester hydrolysis to obtain acid 3. Amide coupling with amine 4 resulted in dimer 5 and subsequent phosphitylation gave 6 which can be used in standard ON synthesis. Phosphoramidite 6 was then used to synthesise ASOs (ON2, ON4, ON5, Fig. 1B)
commercially available phosphoramidite PO-cycle to the AM-cycle can be achieved by PO-coupling (iii) of build up sections with PO linkages (PO-cycle). Transition from the ON synthesis using standard phosphoramidite monomers builds up sections of nucleosides consecutively linked through the PO bond (PO-cycle). Efficient RNase H response mediated by ON2 would confirm that the AM linkage could be accommodated inside the PO binding pocket of RNase H. Efficient RNase H response mediated by ON2 would confirm that the AM linkage could be accommodated inside the PO binding pocket of RNase H. The same rational was applied to ON4 confirming that the AM linkage could be accommodated inside the PO binding pocket of RNase H with the ASO.12,29 ON2 contains three isolated AM linkages at various positions. The design of the tested ASOs was based on three consecutive interactions inside the PO binding pocket of RNase H with the ASO.12,29 ON2 contains three isolated AM linkages which are interspaced by two consecutive POs. Efficient RNase H response mediated by ON2 would confirm that the AM linkage could be accommodated inside the PO binding pocket of RNase H. The same rational was applied to ON4 and ON5 whose gapmer design further narrows the window for RNase H activity while utilising the properties of otherwise RNase H-incompatible modifications, including the 2′-OMe modification31 (ON3, Fig. 1B). However, this strategy has not yet been reported for the AM-cycle as described before. Using PyBOP and NMM as the coupling agent and base gave minimal side products and the combined PO- and AM-cycles gave chimeric gapmer ON6 with charge-neutral wings in an overall isolated yield of 16% (Fig. 3 and Fig. S1, S2 ES1†). The crude AM-modified ONs gave clean chromatographic traces (Fig. 3A and D) with the main peak corresponding to the desired product which was confirmed following purification (Fig. 3B and E) and mass analysis (Fig. 3C and F). We rationalise that the absence of a 2′-fuctionality can increase coupling efficiencies compared to consecutive amide couplings of more challenging RNA-type monomers.31,32

The consecutive AM backbones in ON6 constitute the flanking wings of the ASO, a commonly used strategy to retain RNase H activity while utilising the properties of otherwise RNase H-incompatible modifications, including the 2′-OMe modification33 (ON3, Fig. 1B). However, this strategy has not yet been reported for the AM-cycle as described before. Using PyBOP and NMM as the coupling agent and base gave minimal side products and the combined PO- and AM-cycles gave chimeric gapmer ON6 with charge-neutral wings in an overall isolated yield of 16% (Fig. 3 and Fig. S1, S2 ES1†). The crude AM-modified ONs gave clean chromatographic traces (Fig. 3A and D) with the main peak corresponding to the desired product which was confirmed following purification (Fig. 3B and E) and mass analysis (Fig. 3C and F). We rationalise that the absence of a 2′-fuctionality can increase coupling efficiencies compared to consecutive amide couplings of more challenging RNA-type monomers.31,32

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AM backbones. Thus, ON6 was tested in the RNase H assay while ON7 and ON8 (Fig. 4 and Table 1) were used as positive controls to induce degradation of a 5'-FL-labelled complementary target RNA (FL-RNA2). Aliquots of the reaction buffers containing E. coli RNase H, FL-RNA2 and a catalytic amount of ASOs (ON6, ON7 or ON8) were quenched at different time points and analysed by gel electrophoresis (Fig. 4 and Fig. S3, ESI†). The gels show that all tested ASOs induce complete target degradation within 30 min at 37 °C. Quantification confirmed that AM-gapmer ON6 activates RNase H as efficiently as the controls ON7 and ON8. This is a clear improvement on previously reported gapmers with charge-neutral wings using peptide nucleic acids (PNAs) which only induce target degradation in a non-catalytic way.34 Watts et al. reported that improved gapmer designs and optimisation of linkers between PNA and DNA sections can lead to catalytic activity but the charge-neutral section was limited to only one wing.35 In comparison, AM-gapmer ON6 shows efficient catalytic degradation of the target RNA.

No correlation was observed between the cleavage rate of an RNA target by RNase H with ON6-8 and the measured melting temperatures (TmS, Table 1 and Fig. S4, ESI†). The 2'-OMe modifications in gapmer ON7 increased target affinity to RNA by +2.5 °C (+0.25 °C per modification (mod)) whereas the amide backbone in ON6 slightly decreased duplex stability with RNA by −1.5 °C (−0.19 °C per mod) compared to unmodified ON8. This is consistent with the literature for both modifications.11,25

A flexible 4'-endo sugar pucker, a rigid sugar phosphate backbone, and a duplex conformation between A and B are essential for an efficient RNase H response.36 In this context, we investigated the structural changes in duplexes induced by the AM backbone. We did not observe significant perturbation of the duplex structures formed by ON6 with DNA or RNA targets when compared to ON8 (an unmodified DNA ON) in circular dichroism experiments (Fig. S3, ESI†), which is consistent with the high efficiency of ON6 in inducing an RNase H response.

Enzymatic stability of ASOs is important to ensure optimal biological half-life and therapeutic efficacy. The terminal amides in ON6 result in enhanced nuclease resistance and longer serum lifetime in foetal bovine serum (FBS) compared to ON7 and ON8 which were both rapidly degraded (Fig. S6, ESI†). As previously reported,27 the 2'-OMe modifications in ON7 slightly extend serum lifetime but are not as effective as the amide bonds in ON6.

Potential enhanced cellular uptake of the AM-gapmer design was evaluated by confocal laser scanning microscopy (CLSM). For this experiment, ON6 was fluorescently labelled at the 5'-end to give ON9 (Fig. 3D-F) in which eight PO linkages were replaced by charge-neutral AM bonds and the ON has a charge-to-linkage ratio of 0.56 (including the PO linkage between FL and the 5'-end). No solubility issues were encountered with this oligomer. ON10 and ON11 represent the fluorescently labelled derivatives of ON7 and ON8 respectively in which all 17 internucleoside linkages consist of negatively charged PO bonds, with an additional PO bond for the attachment of FL (Fig. 5). HeLa cells were incubated with 5 μM ON9-ON11 in serum-free medium and analysed by CLSM after fixation. AM-gapmer ON9 showed increased intracellular accumulation compared to ON10 and ON11 for which a distinct fluorescent signal inside the cell was absent (Fig. 5 and Fig. S7 for a general view, ESI†). The localised fluorescent signals observed for ON9 (Fig. 5A) suggest that the uptake mechanism leads to its partial entrapment within subcellular compartments. Co-incubation of ON9 with fluorescently tagged epidermal growth factor (EGF) shows that only a low number of FL signals were co-localised with stained endosomes (Fig. S8, ESI†). Moreover, a clear increase of punctuated fluorescence for ON9 was only detected after incubation for 16 h while receptor-mediated endocytosis is known to happen within <30 min. Together, our preliminary results suggest that ON uptake may be occurring through fluid phase endocytosis which results in unspecific uptake from the extracellular fluid into vacuoles, while their fate upon maturation can be highly variable.38 These observations are also in agreement with reports that fully charge-neutral methyl phosphonate-modified ASOs enter the cell via fluid phase endocytosis after 10–12 hours of incubation.21 Similar results showed enhanced cellular uptake of ASOs in which half the charge was neutralised by phosphotriesters.19

Table 1 TmS of AM-gapmer ON6 and 2'-OMe-gapmer ON7 compared to unmodified DNA ON8 against a complementary RNA and DNA target. Values were obtained from the maxima dΔα/ΔT vs. T for 3 mM of each ON in 10 mM phosphate buffer, 200 mM NaCl, pH 7.0, n = 2 ± SD. ΔTm is relative to the unmodified control ON8

| ON     | (5' → 3')                                                                 | RNA target | DNA target |
|--------|---------------------------------------------------------------------------|------------|------------|
|        |                                                                           | Tm         | ΔTm        | Tm         | ΔTm        |
| ON6    | TpUpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpT               | 54.6       | −1.5       | 52.3       | −3.2       |
| ON7    | UpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUpUp      | 58.6       | +2.5       | 47.8       | −7.7       |
| ON8    | TpUpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpTpT               | 56.1       | −         | 55.5       | −         |

a = AM, p = PO, U = 2'-OMe uridine.
In conclusion, different ON chemistries and their effect on pharmacodynamics, pharmacokinetics, cellular delivery and toxicity are poorly understood and backbone modifications have been mainly focused on the PS linkage, leaving other chemistries under-explored. The recently reported applications of charge-neutral backbones to enhance cellular uptake\textsuperscript{29–31} and mediate toxicity\textsuperscript{32} emphasise the importance of exploring other artificial backbone structures. Here we report the partial replacement of the natural ON PO backbone by an amide internucleoside linkage and its effects on antisense activity, target engagement, serum stability and cellular uptake. We show that the AM linkage is well tolerated in the wings of an ASO gampner design while retaining an efficient RNase H response. The stability of the formed duplex is only slightly decreased (–0.19 °C per mod) compared to unmodified DNA, and this does not compromise target engagement and RNA cleavage efficiency. The ease of chimeric gampner design with synthesis with reduced steric complexity, retained catalytic efficiency, high serum stability and potentially enhanced cellular uptake all add desirable drug-like properties. Thus, the AM backbone mimic represents a valuable candidate for further development.

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Conflicts of interest

There are no conflicts to declare.

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