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Hydrophobic alkyl-linked base-modified pyrimidine and 7-deazapurine 2'-deoxyribonucleoside phosphoramidites: synthesis and application in solid-phase synthesis of modified and hypermodified oligonucleotides

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A series of 2'-deoxyribonucleoside 3'-phosphoramidites bearing hydrophobic alkyl-linked modifications at nucleobases was synthesized, namely 5-phenylethyluracil, 5-pentylcytosine, 7-(indol-3-yl)ethyl-7-deazaadenine, and 7-isopentyl-7-deazaguanine derivatives. These nucleoside phosphoramidites were used for solid-phase synthesis of modified and hypermodified oligonucleotides containing up to fifteen modified nucleotides in a row. Their hybridization with complementary non-modified or modified oligonucleotides and thermal stability of the resulting DNA duplexes was studied using UV-vis denaturing experiments and CD spectroscopy. The results indicate that the partially modified hydrophobic DNA can still retain B-conformation, although with lower thermal stability. On the other hand, the hypermodified ONs containing all four modified nucleotides did not hybridize to duplexes likely due to formation of aggregates as indicated by dynamic light scattering measurement. This work expands the toolkit of chemically modified nucleotides for applications in functional nucleic acids or nucleic acid therapeutics, but also shows the scope and limitations of the use of hydrophobic nucleotides in hypermodified oligonucleotides and DNA.

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Introduction

Chemically modified nucleic acids have become indispensable tools in chemical biology, diagnostics, therapeutic development, and material sciences.¹ Synthetic oligonucleotides, bearing site-specific modifications on nucleobases, offer a powerful approach for tuning hybridization behaviour, enhancing binding specificity and affinity, and introducing new functionalities. This is particularly relevant in the context of aptamer selection and engineering,² where introduction of one^{3–10} or two^{11,12} aromatic or hydrophobic moieties at nucleobases has led to significantly improved affinity and selectivity for protein targets, when compared to their unmodified counterparts.

The substitution of nucleobases with rigid alkynyl-linked substituents has been studied extensively. These modifications can be easily introduced through Pd-catalyzed Sobnoga-shira reactions of halogenated nucleosides with terminal alkynes and the resulting 5-alkynylpyrimidine^{13–18} and 7-alkynyl-

7-deazapurine^{19–21} nucleotides were reported to stabilize (but in some cases destabilize²²) DNA and RNA duplexes *via* enhanced π - π stacking interactions and increased hydrophobicity the and decrease the toxicity of antisense oligonucleotides (ASOs).²³ On the other hand, flexible alkyl-linked modifications at the nucleobase have been studied less frequently mostly focusing on derivatives of 5-methylpyrimidines.^{24–26}

In most of the above mentioned works, the studies have been limited to nucleic acids containing just one or two modified bases, whereas just a handful of scattered papers reported substitution on all four canonical nucleobases.^{27,28} In our lab, we have recently developed the general concept of hypermodified DNA and RNA where each and every nucleobase is bearing a different modification and their enzymatic synthesis using sets of all four modified (d)NTPs. In hypermodified DNA we reported examples containing four different hydrophobic modifications,^{29,30} anionic³¹ or cationic³² groups, sugars,³³ amino acid-like functional groups³⁴ and combinations of substituents from these classes. These hypermodified DNA that can contain up to 150 modified nucleotides in a row can still be sequenced and thus have potential in selection of aptamers and other functional nucleic acids. However, for these applications a phosphoramidite-based chemical synthesis³⁵ of the hypermodified oligonucleotides will be mostly needed to provide

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these polymers in larger quantities. We have previously reported³⁶ the design and synthesis of a complete set of base-modified 2'-deoxyribonucleoside 3'-phosphoramidites, each bearing alkyne-linked hydrophobic arylethyl or alkynyl substituents (analogous to the hydrophobic alkyne-linked dNTPs used for enzymatic synthesis of hypermodified DNA) and found an interesting effect that one or several alkynyl modifications slightly destabilize the DNA duplexes while the full modification at every nucleotide leads to stabilization of duplexes. To complement the previous study and extend the portfolio of hydrophobic modifications also to flexible alkyl substituents, we report here the synthesis of full set of all four 2'-deoxyribonucleoside 3'-phosphoramidites bearing four different hydrophobic alkyl or arylalkyl substituents and their use on the solid-phase synthesis of ONs.

Results and discussion

Synthesis of modified nucleoside phosphoramidites

The set of modified nucleoside phosphoramidite building blocks was designed based on previously reported alkynyl-linked phosphoramidites (**1a–1d**)³⁶ and on the set of modified dNTPs²⁹ used for the enzymatic synthesis of hypermodified hydrophobic DNA (Fig. 1A). Hence the new target nucleoside phosphoramidites were derived from 5-phenylethyl-2'-deoxyuridine (**2a**), 5-pent-1-yl-2'-deoxycytidine (**2b**), 7-(indol-3-yl)ethyl-7-deaza-2'-deoxyadenosine (**2c**) and 7-isopentyl-7-deaza-2'-deoxyguanosine (**2d**) (Fig. 1B), where the substituents resemble and mimic side chains of amino acid residues phenylalanine (Phe), methionine (Met), tryptophan (Trp), and valine (Val), respectively.

The synthesis of modified pyrimidine nucleosides consisted of attaching the modifications through Sonogashira cross-coupling to the iodo-nucleoside according to the previous work,³⁶ followed by hydrogenation of the ethynyl to the ethyl linker, introduction of protecting groups and attachment of the phosphoramidite function (Scheme 1, see details in SI). The synthesis of functionalized pyrimidine nucleosides commenced with the reduction²⁹ of the ethynyl linker to ethyl in compound

3a and **3b** (prepared according to ref. 36) by H₂ with 10% Pd/C (0.1 equiv for **3a** and 0.3 equiv. for **3b**) in MeOH. The reaction afforded the alkyl-linked nucleosides **4a** and **4b** in good yields of 90% and 84%, respectively. The 5'-hydroxyl group was then selectively protected using dimethoxytrityl chloride (DMTrCl) in the presence of *N,N*-dimethylaminopyridine (DMAP),³⁷ resulting in DMTr-protected nucleosides **5a** (77%) and **5b** (69%). Subsequently, the exocyclic amino group of modified 2'-deoxycytidine **5b** as a dimethylformamide derivative by treatment with *N,N*-dimethylformamide dimethylacetal (DMF-DMA) in DMF³⁸ at 40 °C resulting in compound **6b** in a good yield (92%). Finally, coupling of **5a** and **6b** with 2-cyanoethyl-*N,N*-diisopropylchlorophosphoramidite³⁵ in the presence of *N,N*-diisopropylethylamine (DIPEA) led to fully protected nucleoside phosphoramidites **2a** (65%) and **2b** (54%).

For the 7-substituted 2'-deoxy-7-deazapurine nucleoside phosphoramidites, a different strategy was employed. First, the protecting groups were first installed, followed by attachment of the hydrophobic alkyne moiety, reduction of the alkyne linker, and conversion of the modified nucleosides to the corresponding phosphoramidites. Previously,³⁶ for the introduction of indolylethyl moiety, we used 1-acetyl-3-(trimethylsilyl) ethynylindole, which required TEA·3HF for *in situ* cleavage of the TMS during the coupling reaction. Since we were not able to scale up the reaction under such conditions, we deprotected the trimethylsilyl from the indole moiety beforehand and used it as such in the Sonogashira cross-coupling. Therefore, we synthesized known 1-acetyl-3-ethynylindole³⁹ which was then used in the Sonogashira reaction with protected 7-iodo-2'-deoxy-7-deazaadenosine **7c** (see details in SI).

Hydrogenation of protected and functionalized nucleosides **3c** and **3d** by H₂ with 10% Pd/C in MeOH resulted in compound **4c** (77%) and **4d** (89%), however, we needed to use large amount (1 equiv.) of the palladium catalyst to be able to perform the reduction in case of **3c** (0.2 equiv. in case of **3d**). Final reactions with 2-cyanoethyl-*N,N*-diisopropylchlorophosphoramidite, under the conditions mentioned above, resulted in target fully protected nucleoside phosphoramidites **2c** (41%) and **2d** (57%), respectively.

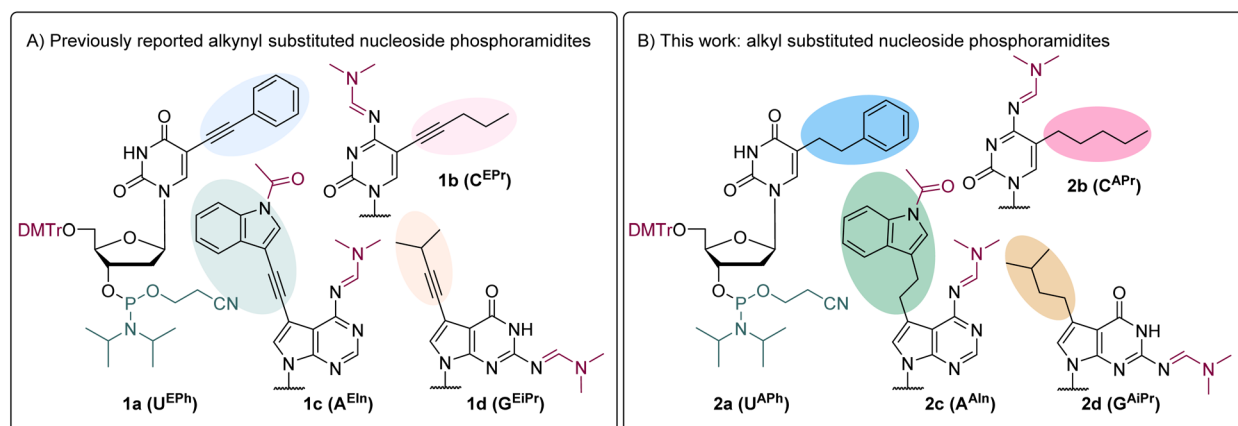
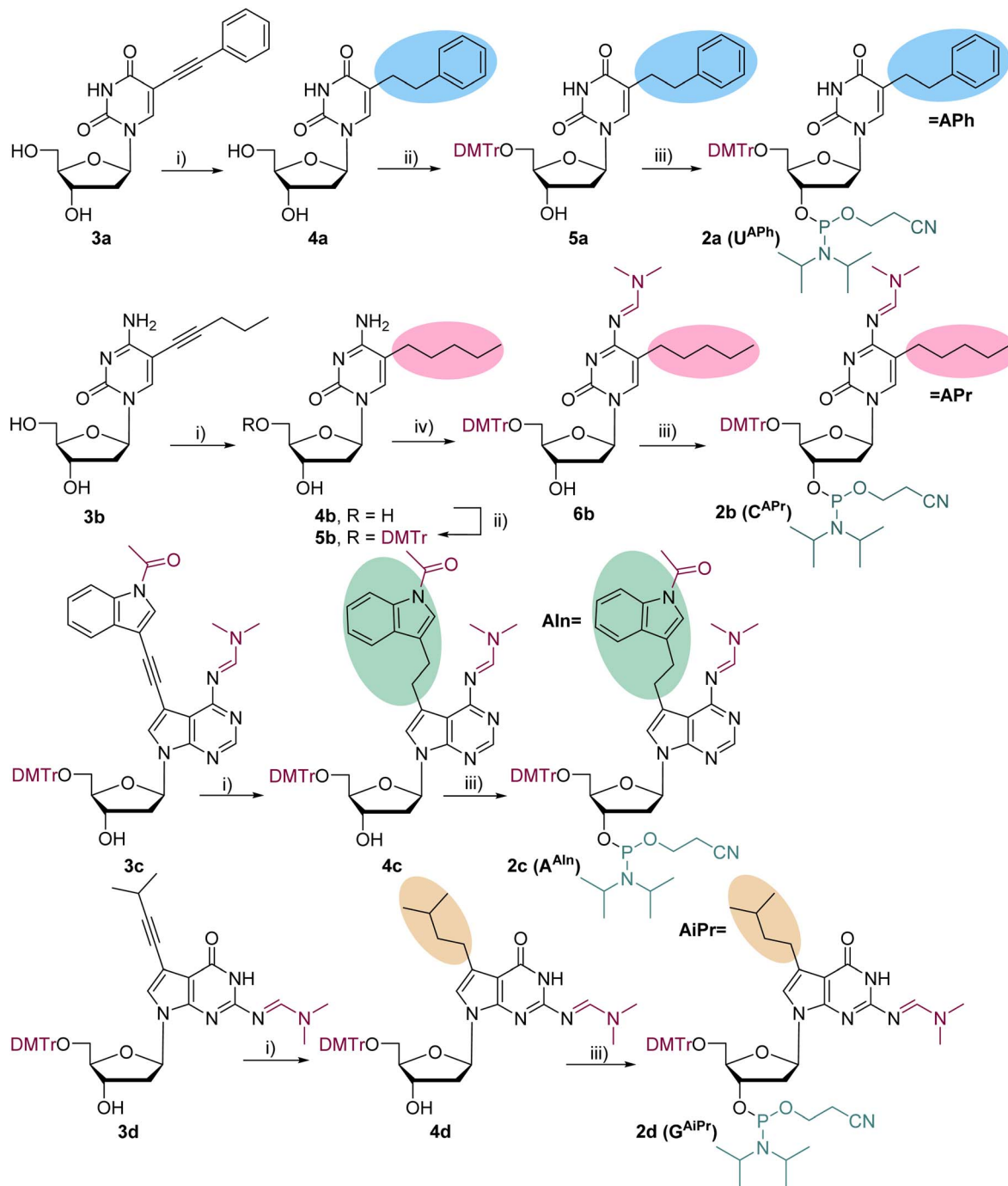


Fig. 1 (A): Previously reported series with modifications linked to the nucleobase via ethynyl linker; (B): new series with ethyl linker between the modification and the nucleobase.



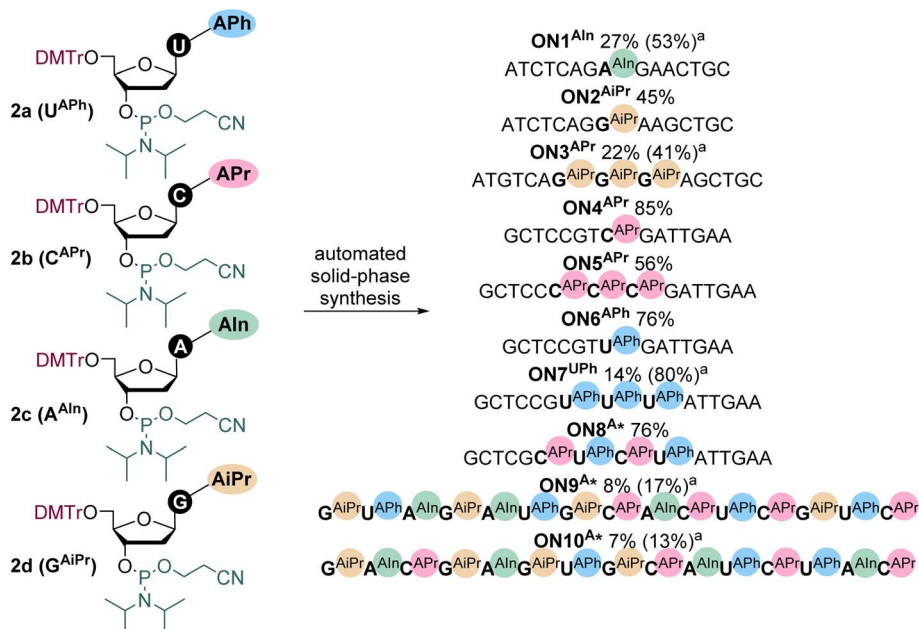
Scheme 1 Reagents and conditions: (i) H_2 , 10% Pd/C, MeOH, iPrOH (in case of **3c** and **3d**), RT, 3 h (**3a** 90%, **3b** 84%, **4c** 77%, **4d** 89%); (ii) DMTrCl (1.2 equiv.), DMAP (0.1 equiv.), pyridine (dry), RT, overnight (**5a** 77%, **5b** 69%); (iii) 2-cyanoethyl-*N,N*-diisopropylchlorophosphoramidite (1.2 equiv.), DIPEA (2.5 equiv.), DCM (dry), 0 °C to RT, 1.5 h (**2a** 65%, **2b** 54%, **2c** 41%, **2d** 57%); (iv) DMF-DMA (14 equiv.), DMF (dry), 40 °C, under Ar, 4 h (**6b** 92%).

Solid-phase synthesis of modified oligonucleotides

The modified nucleoside phosphoramidites **2a–2d** were used for the automated synthesis of partially and fully modified oligonucleotides (ONs). We designed ten target sequences containing one or three modified nucleotides in the central part of the chain, combination of two different modified nucleotides and then fully modified complementary sequences containing

combinations of all four modified nucleotides, corresponding to those we previously published³⁶ in the alkyne-linked series (for sequences see Scheme 2 or Table S2 in SI). Since the final amount of the nucleoside phosphoramidite **2c** was not sufficient for all types of ONs to be synthesized, oligonucleotides containing three **2c** were neglected (Scheme 2), but we still synthesized the hyper-modified sequences (ON9^{A*} and





Scheme 2 Solid-phase synthesis of the partially and hyper-modified oligonucleotides with their yields (a) HPLC yields given in parenthesis in case of lower isolated yield).

ON10^{A*}). The ONs were synthesized using standard synthesis protocol for solid-phase synthesis employing the phosphoramidite method. The partially modified **ON1^{Aln}**–**ON8^{A*}** were synthesized on standard solid-phase columns with one nucleoside attached at the 3' end, whereas the hyper-modified **ON9^{A*}** and **ON10^{A*}** were synthesized on the universal solid-phase columns, with no nucleoside on the solid phase support. The

synthetic scale was set to 1 μ mol using the 0.1 M concentration of all the phosphoramidites. The coupling duration for the natural phosphoramidites was 1 minute 30 seconds while for the modified phosphoramidites the coupling time was increased to 6 minutes to elevate the probability of successful incorporation. After the synthesis, the cleavage of the ONs from the solid phase was performed by aqueous ammonia, followed

Table 1 List of synthesized modified oligonucleotides (ON^{A*}) with purchased non-modified complementary strands (cON) forming modified duplexes DNA^{A*} (15 bp) with modifications attached to the nucleobase via ethyl linker (DNA^{A*}; A = alkyl/ethyl linker, * = modification)

Code		DNA ^{A*}
DNA1^{Aln}	ON1 ^{Aln} cON1	5'-ATCTCAGA ^{Aln} GAAGTGC-3' 3'-TAGAGTC T CTTGACG-5'
DNA2^{AiPr}	ON2 ^{AiPr} cON2	5'-ATCTCAGG ^{AiPr} AAGCTGC-3' 3'-TAGAGTC C TTCGACG-5'
DNA3^{AiPr}	ON3 ^{AiPr} cON3	5'-ATGTCAG ^{AiPr} G ^{AiPr} G ^{AiPr} AGCTGC-3' 3'-TACAGT C C C TCGACG-5'
DNA4^{APr}	ON4 ^{APr} cON4	5'-GCTCCGTC ^{APr} GATTGAA-3' 3'-CGAGGCA G CTAAGT-5'
DNA5^{APr}	ON5 ^{APr} cON5	5'-GCTCCC ^{APr} C ^{APr} C ^{APr} GATTGAA-3' 3'-CGAGG G G G CTAAGT-5'
DNA6^{Aph}	ON6 ^{Aph} cON6	5'-GCTCCGTU ^{Aph} GATTGAA-3' 3'-CGAGGCA A CTAAGT-5'
DNA7^{Aph}	ON7 ^{Aph} cON7	5'-GCTCCGU ^{Aph} U ^{Aph} U ^{Aph} ATTGAA-3' 3'-CGAGGC A A A TAAGT-5'
DNA8^{A*}	ON8 ^{A*} cON8	5'-GCTCGC ^{APr} U ^{Aph} C ^{APr} U ^{Aph} ATTGAA-3' 3'-CGAGC G A G A TAAGT-5'
DNA9^{A*}	ON9 ^{A*} cON9	5'-G ^{AiPr} U ^{Aph} A ^{Aln} G ^{AiPr} A ^{Aln} U ^{Aph} G ^{AiPr} C ^{APr} A ^{Aln} C ^{APr} U ^{Aph} C ^{APr} G ^{AiPr} U ^{Aph} C ^{APr} -3' 3'-C A T C T A C G T G A G C A G-5'
DNA10^{A*}	ON10 ^{A*} ON9	5'-G ^{AiPr} A ^{Aln} C ^{APr} G ^{AiPr} A ^{Aln} G ^{AiPr} U ^{Aph} G ^{AiPr} C ^{APr} A ^{Aln} U ^{Aph} C ^{APr} U ^{Aph} A ^{Aln} C ^{APr} -3' 3'-C T G C T C A C G T A G A T G-5'
DNA11^{A*}	ON9 ^{A*} ON10 ^{A*}	5'-G ^{AiPr} U ^{Aph} A ^{Aln} G ^{AiPr} A ^{Aln} U ^{Aph} G ^{AiPr} C ^{APr} A ^{Aln} C ^{APr} U ^{Aph} C ^{APr} G ^{AiPr} U ^{Aph} C ^{APr} -3' 3'-C ^{APr} A ^{Aln} U ^{Aph} C ^{APr} U ^{Aph} A ^{Aln} C ^{APr} G ^{AiPr} U ^{Aph} G ^{AiPr} A ^{Aln} G ^{AiPr} C ^{APr} A ^{Aln} G ^{AiPr} -5'



by deprotection of the protecting groups by heating the solution to 65 °C for 6 h. Subsequently, the HPLC purification of the ONs was performed, followed by the mass spectrometry characterization and purity control by UHPLC-MS (see details in SI). Concentration was approximately measured by nanodrop and precise concentration and yield determination was done by the elemental analysis of phosphorus content.

All the partially modified (**ON1^{Aln}**–**ON8^{A*}**) and fully modified ONs (**ON9^{A*}**, **ON10^{A*}**) were successfully synthesized, purified and characterized (see details in SI). According to the retention times obtained during HPLC purification on a C18 column (Table S1 in SI), the relative hydrophobicity of the resulting partially modified ONs can be estimated. Singly modified ONs show the lowest hydrophobicity from the set (**ON4^{APr}** < **ON6^{Aph}** < **ON2^{AiPr}** < **ON1^{Alm}**), indicating pentyl-C modification is the least hydrophobic and indole-modified A the highest. Generally, increasing the number of incorporated modifications further enhances hydrophobicity, as is reflected in the trend **ON5^{APr}** (3 mod.) < **ON7^{Aph}** (3 mod.) < **ON3^{AiPr}** (3 mod.) < **ON8^{A*}** (4 mod.). However, the isolation of the hyper-modified strands (**ON9^{A*}**, **ON10^{A*}**) was more difficult as it was needed to prolong the HPLC gradient (of MeCN, phase B, in 0.1 M TEAB aqueous solution, phase A) from 1 h to 2 h to obtain pure **ON9^{A*}** and **ON10^{A*}**. Isolated yields (Scheme 2 and Table S2) in some cases were good: 45–85% for ONs containing one modification (45% for **ON2^{AiPr}**, 85% for **ON4^{APr}**, and 76% for **ON6^{Aph}**), three pentylC modifications (**ON5^{APr}**, 56%) and combination of pentylC with phenylethylU modifications (**ON8^{A*}**, 76%). On the other hand, only 14% isolated yield was obtained when three phenylethylU modifications were incorporated in a row (**ON7^{Aph}**). Lower yields (7–27%) were also obtained for other ONs containing multiple modifications (22% for **ON3^{AiPr}**, 8% for **ON9^{A*}**, and 7% for **ON10^{A*}**) or containing single modification of indolyethylA (**ON1^{Alm}**, 27%).

The purity of synthesized ONs* was determined by UHPLC-MS and in most cases was above 96% except **ON6^{Aph}** and **ON10^{A*}** with purities 91% and 88%, respectively (see details in SI, Table S2). Pure modified ONs were successfully isolated in sufficient amounts for following studies (avg. 2 mM in 200 μl). It should be mentioned, that during the course of our work, a preprint⁴⁰ has been released reporting the use of related unprotected indol-linked phosphoramidites in the oligonucleotide synthesis, but this approach did not work in our hands.

The hydrophobic modifications most likely decrease the solubility of the resulting ONs in water, since the individual modifiers have lower solubilities in water (benzene 1.8 g L⁻¹, indole 3.6 g L⁻¹, propane 0.07 g L⁻¹ at 25 °C) than the nucleosides (uridine 50 g L⁻¹, adenosine 25 g L⁻¹, cytidine 36 g L⁻¹, guanosine 10 g L⁻¹).⁴¹ However, we did not observe any precipitation in water or in buffer solutions, therefore, we imply all of the studied ONs are soluble in water and commonly used buffers.

The yields of the modified ONs (Scheme 2 and Table S2 in SI) synthesized from the hydrophobically modified phosphoramidites **2a–2d** are lower compared to usual yields for non-modified ONs, especially for the hypermodified sequences, but still mostly somewhat better than for the previously reported alkynyl-linked phosphoramidites (**1a–1b**).

Hybridization of modified oligonucleotides

All partially modified ONs (**ON1^{Aln}**–**ON8^{A*}**) were annealed with non-modified complementary strands (**cON1**–**cON8**) from commercial supplier (to form **DNA1^{Aln}**–**DNA8^{A*}**, Table 1) and compared by gel electrophoresis with matching non-modified DNA (see sequences in SI, Table S3). As expected, the size of unmodified and partially modified DNA was very similar, as shown by mobility on the agarose gel (Fig. 2).

Nonetheless, in case of the hypermodified strands combined with non-modified complementary strand (forming **DNA9^{A*}**, **DNA10^{A*}**) and in case of both hypermodified complementary strands (forming **DNA11^{A*}**), we were unable to confirm the formation of double-stranded DNA on the gel (Fig. 2), even though we tried different buffer compositions and various ionic strengths to promote the annealing (see details and gels in SI).

Circular dichroism spectroscopy

The structure of synthetic DNA duplexes (**DNA1^{Aln}**–**DNA8^{A*}**) was studied by circular dichroism (CD) spectroscopy. CD spectra for non-modified DNAs had characteristic conservative pattern for B-form DNA thus showing positive band at ~280 nm and a negative band at around ~255 nm (Fig. 3). Eventual difference in spectral shape and/or their intensity is due to difference in primary sequence of nucleotides. CD spectra of the partially

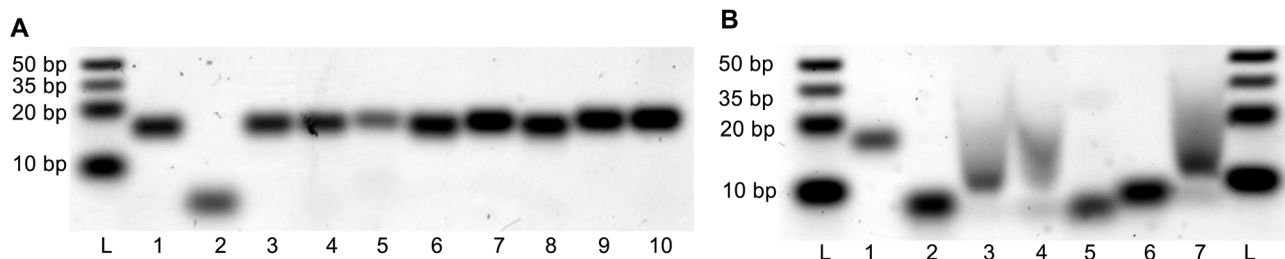


Fig. 2 (A): Agarose gel analysis of (1) non-modified **DNA1**, (2) non-modified **ON1** (3–10) modified **DNA1^{Aln}**–**DNA8^{A*}**; (L): DNA ladder; (B): agarose gel analysis of (1) non-modified **DNA9**, (2) non-modified **ON9**, (3) fully modified **ON9^{A*}**, (4) fully modified **ON10^{A*}**, (5) fully modified **ON9^{A*}** with non-modified **cON9** (**DNA9^{A*}**), (6) fully modified **ON10^{A*}** with non-modified **ON9** (**DNA10^{A*}**), (7) fully modified **ON9^{A*}** with fully modified **ON10^{A*}** (**DNA11^{A*}**); (L): DNA ladder.



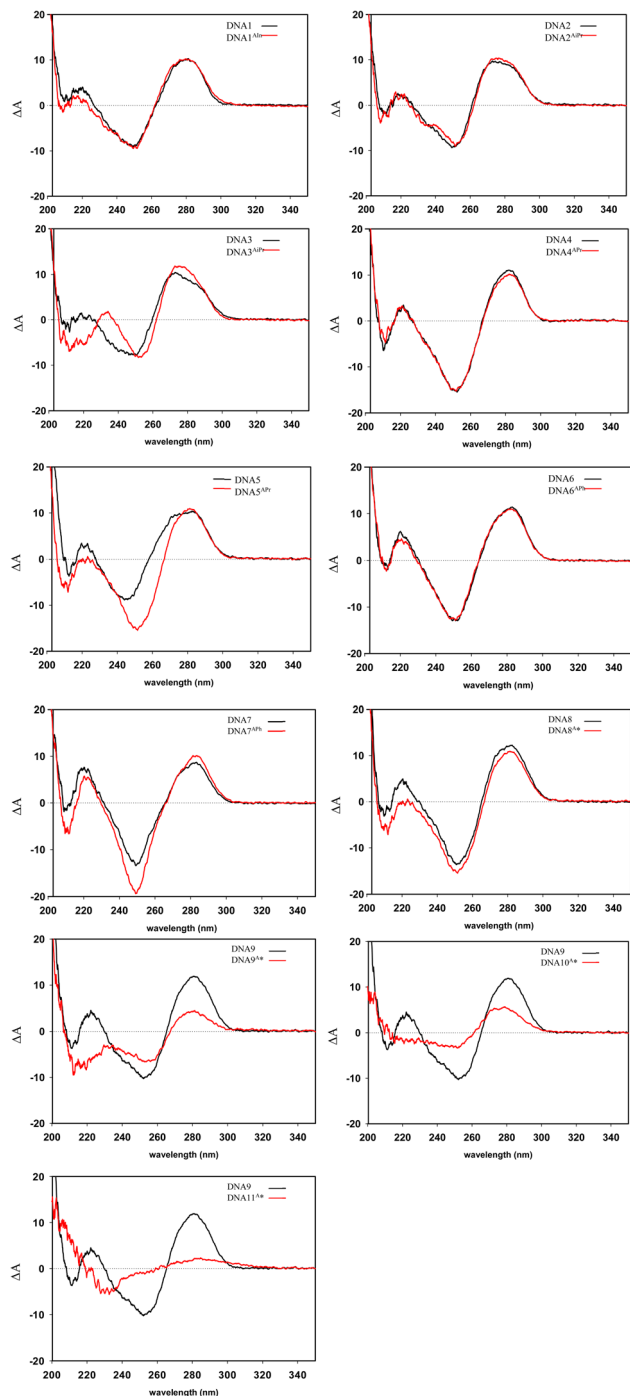


Fig. 3 CD Spectra of modified DNA1^{AIn}-DNA11^{A*} (red curves) compared with their non-modified equivalent DNA1-DNA9 (black curves).

modified DNA had rather similar pattern indicating that the modified DNA duplexes probably adopt B-form conformation. The observed variations in spectral patterns and intensities, relative to the corresponding unmodified DNA, can be attributed to the introduced oligonucleotide modifications (Fig. 3). While a single modification exerts minimal influence on the CD spectrum, the incorporation of three or more modifications results in pronounced spectral alterations.

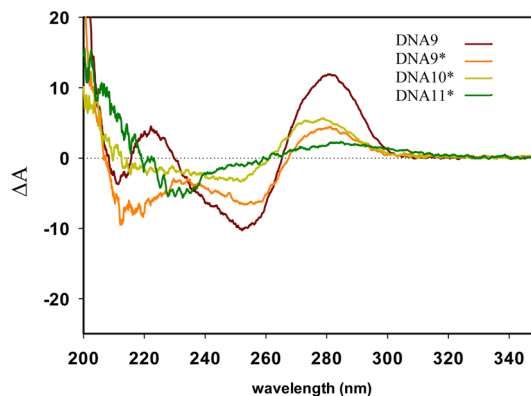


Fig. 4 Comparison of CD spectra at 5 °C of non-modified DNA9 and corresponding modified sequences of DNA9^{A*}, DNA10^{A*} and hyper-modified DNA11^{A*}.

Though the pattern of CD spectrum obtained for DNA9^{A*} (Fig. 4), with one strand modified and the complementary non-modified, seems rather similar as those for non-modified DNA9, however, its low intensity reflects mostly the presence of unannealed oligonucleotides in agreement with agarose gel analysis (Fig. 2, lane 5).

For DNA10^{A*}, similarly as DNA9^{A*} with one strand modified and the complementary strand non-modified, or for DNA11^{A*}, the hyper-modified duplex with two fully modified strands, the duplex formation is not detected, neither by the agarose gel analysis, nor by the CD spectroscopy (Fig. 4). Presumably, the multiple adjacent highly hydrophobic moieties tend to form aggregates or micelle-like structures that prevent the hyper-modified ONs from hybridization into DNA duplex. This is in contrast to previously reported hypermodified rigid alkynyl-linked ONs that were able to form the duplexes. However, amphiphilic DNA is generally known to aggregate, as is in the case of cholesterol-modified DNA,⁴²⁻⁴⁴ where aggregation causes a challenge while studying membrane proteins.⁴⁵ Example of base-modified ONs with 5-(dodec-1-ynyl)uracil exhibited micelle formation, whose size, and stability depended strongly on the number of hydrophobic units, rather than their position in the sequence.⁴⁶

To further examine the effect of the linker on the duplex behavior, CD spectra of DNA6^{EPh} and DNA7^{EPh} were additionally measured (Fig. 5). One modification in the middle for both types of modification linkers cause only minor changes in the CD spectra compared to non-modified DNA duplex (DNA6^{E/Ph} in comparison with non-modified DNA6, Fig. 5A). The overall shape of obtained CD spectra confirms that DNA6^{E/Ph} form B-type DNA conformation.

Modified DNA7^{E/Ph} still retain B-form DNA, according to measured CD spectra (Fig. 5B). Introducing three modifications cause spectral changes especially in case of ethynyl-phenyl modifier. Observed effects are influenced by the presence of additional chromophores in conjugation with the nucleobase and their eventual exciton coupling. Specifically, the spectral changes including the emergence of the negative spectral band at ~305 nm accompanied by the blue shift of positive band from



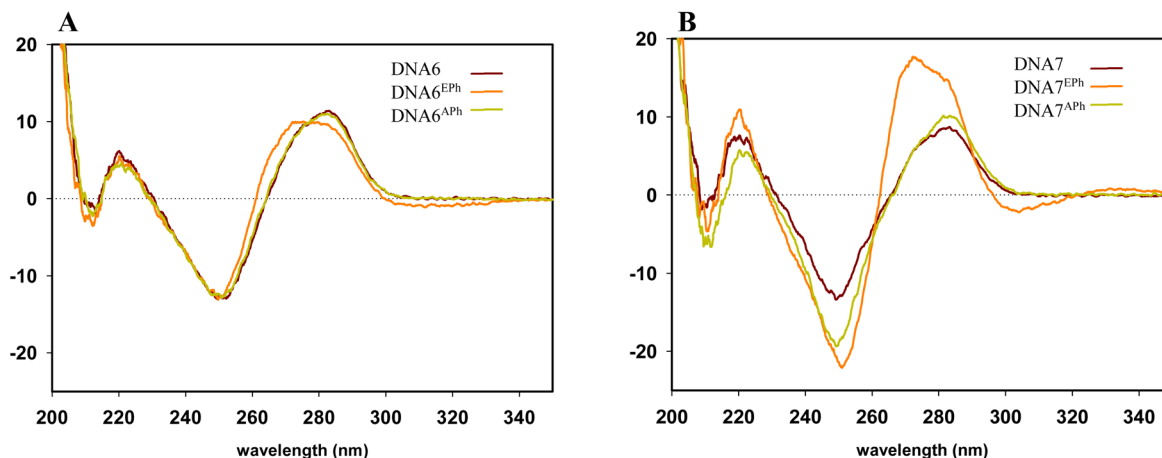


Fig. 5 (A): CD spectra of modified DNA6^{EPh} and DNA6^{Aph} of the same sequence compared with the corresponding non-modified sequence (DNA6). (B): CD spectra of modified DNA7^{EPh} and DNA7^{Aph} compared with the corresponding non-modified sequence (DNA7). CD spectra were measured at 5 °C.

~285 nm to ~275 nm together with increasing its intensity. Ethyl-linked phenyl modifications only cause increase in negative spectral band at ~250 nm compared to non-modified DNA7, however, overall change in the spectra are minimal indicating structural likeness with corresponding non-modified duplex.

Dynamic light scattering measurements

To qualitatively assess the aggregation state of the presented hyper-modified ONs, dynamic light scattering (DLS) measurements were performed, and the derived count rate (DCR) was compared. The DCR normalizes the measured scattered light intensity by accounting for instrument attenuation, allowing for

Table 2 Comparison of melting temperatures (T_m [°C]) of non-modified (DNA) and modified DNA duplexes (DNA^{A/E*}, E stands for ethynyl linker, A stands for alkyl/ethyl linker) determined by UV-vis spectroscopy at 260 nm, their sequences (X stands for A/E), the absolute difference of T_m between modified DNAs and non-modified DNAs (DNA^{A/E*} ΔT_m [°C]), and the difference of T_m between modified DNAs and non-modified DNAs per modification (DNA^{A/E*} ΔT_m /modification, DNA^{A/E*} ΔT_m /modification [°C])

Code	Sequence	DNA ^{A*} ΔT_m	DNA ^{E*} ΔT_m ^b	DNA ^{A*} ΔT_m / mod	DNA ^{E*} ΔT_m / mod ^b
DNA1 ^{A/EIn}	5'-ATCTCAGA ^{XIn} GAAGCTGC-3' 3'-TAGAGTC T CTTGACG-5'	-5.5	-2.2	-5.5	-2.2
DNA2 ^{A/EiPr}	5'-ATCTCAGG ^{XiPr} AAGCTGC-3' 3'-TAGAGTC C TTCGACG-5'	-4.4	-1.2	-4.4	-1.2
DNA3 ^{A/EiPr}	5'-ATGTCAG ^{XiPr} G ^{XiPr} G ^{XiPr} AGCTGC-3' 3'-TACAGT C C C TCGACG-5'	-8.1	-1.9	-2.7	-0.6
DNA4 ^{A/EPr}	5'-GCTCCGTC ^{XPr} GATTGAA-3' 3'-CGAGGCA G CTAACCT-5'	-5.3	+0.2	-5.3	+0.2
DNA5 ^{A/EPr}	5'-GCTCCC ^{XPr} C ^{XPr} C ^{XPr} GATTGAA-3' 3'-CGAGG G G G CTAACCT-5'	-4.6	+7.2	-1.5	+2.4
DNA6 ^{A/EPh}	5'-GCTCCGTU ^{XPh} GATTGAA-3' 3'-CGAGGCA A CTAACCT-5'	-7.5	-1.1	-7.5	-1.1
DNA7 ^{A/EPh}	5'-GCTCCGU ^{XPh} U ^{XPh} U ^{XPh} ATTGAA-3' 3'-CGAGG C A A A TAACCT-5'	-11.2	-1.0	-3.7	-0.3
DNA8 ^{A/E*}	5'-GCTCCG ^{XPr} U ^{XPh} C ^{XPr} U ^{XPh} ATTGAA-3' 3'-CGAGC G A G A TAACCT-5'	-10.8	+6.3	-2.7	+1.6
DNA9 ^{A/E*}	5'-G ^{XiPr} U ^{XPh} A ^{XIn} G ^{XiPr} A ^{XIn} G ^{XiPr} U ^{XPh} G ^{XiPr} C ^{XPr} A ^{XIn} C ^{XPr} U ^{XPh} C ^{XPr} G ^{XiPr} U ^{XPh} C ^{XPr} -3' 3'-CATCTACGTGAGCAG-5'	ND ^a	+7.1	ND ^a	+0.5
DNA10 ^{A/E*}	5'-G ^{XiPr} A ^{XIn} C ^{XPr} G ^{XiPr} A ^{XIn} G ^{XiPr} U ^{XPh} G ^{XiPr} C ^{XPr} A ^{XIn} U ^{XPh} C ^{XPr} U ^{XPh} A ^{XIn} C ^{XPr} -3' 3'-CTGCTCACGTAGATG-5'	ND ^a	+7.4	ND ^a	+0.5
DNA11 ^{A/E*}	5'-G ^{XiPr} U ^{XPh} A ^{XIn} G ^{XiPr} A ^{XIn} G ^{XiPr} U ^{XPh} G ^{XiPr} C ^{XPr} A ^{XIn} C ^{XPr} U ^{XPh} C ^{XPr} G ^{XiPr} U ^{XPh} C ^{XPr} -3' 3'-C ^{XPr} A ^{XIn} U ^{XPh} C ^{XPr} U ^{XPh} A ^{XIn} C ^{XPr} G ^{XiPr} U ^{XPh} G ^{XiPr} A ^{XIn} G ^{XiPr} C ^{XPr} A ^{XIn} G ^{XiPr} -5'	ND ^a	+6.6	ND ^a	+0.2

^a ND: not determined. ^b Taken from ref. 30.



a direct comparison of the overall scattering signal between samples.

Hyper-modified **ON9^{A*}** and **ON10^{A*}** exhibited a higher DCR (2555 ± 107.1 kcps and 6139 ± 278.7 kcps, respectively) compared to the non-modified **ON9** (297 ± 40.2 kcps) and **ON10** (568 ± 51.8 kcps). Given that the ONs were measured under the same conditions (0.5 mg per ml, in H₂O, 25 °C), the increase in light scattering intensity is characteristic for the presence of larger scattering species.

While the poor quality of the measured autocorrelation functions precluded the derivation of reliable quantitative size metrics (e.g., Z-average or Polydispersity Index, Table S7 in SI), the clear difference in normalized scattering intensity supports the hypothesis that the hydrophobic modification in **ON9^{A*}** and **ON10^{A*}** drives the formation of large, unstable aggregates in solution.

Duplex stability

Melting (T_m , Table 2) and annealing (T_a , see Table S4 in SI) temperatures of all modified and non-modified DNA duplexes were measured by UV-vis spectroscopy at 260 nm, and also by CD spectroscopy (see Table S5 in SI).

The T_m data from UV-vis spectroscopy show that the stability of the modified DNA duplexes are significantly lower compared to their unmodified variants. The most prominent decrease of stability is visible for **DNA6^{A^{Ph}}** bearing single ethylphenyl modification (-7.5 °C), followed by singly modified **DNA1^{Aⁱⁿ}** (-5.5 °C), **DNA4^{A^{Pr}}** (-5.3 °C) and **DNA2^{A^{Pr}}** (-4.4 °C). These results suggest one modification has higher destabilization potency than more incorporated modifications. In the case of hypermodified constructs **DNA9^{A*}**, **DNA10^{A*}** and **DNA11^{A*}** the melting temperatures could not be reliably measured as they did not form duplexes.

In comparison with previously published series with ethynyl linker (**DNA^{E*}**)³⁶ we can see that DNA duplexes formed from ONs containing modifications linked by ethynyl linker have generally higher T_m than in the ethyl-linked series (Table 2). The biggest difference is visible with singly phenyl-substituted U-modified **DNA6^{A/E^{Ph}}** where with ethynyl linker the T_m is lower by -1.1 °C in comparison with corresponding non-modified **DNA6** whereas by -7.5 °C when ethyl linker is present. In some cases, the T_m of ethynyl-linked modified DNAs is higher compared to the corresponding non-modified DNAs. Namely, those bearing modified C or U (**DNA5^{E^{Pr}}**, **DNA6^{E^{Pr}}**, **DNA8^{E*}**) and hyper-modified DNAs with either one or both strands fully modified (**DNA9^{E*}**, **DNA10^{E*}**, **DNA11^{E*}**) are showing higher stability than the non-modified version. We suggested that the stabilizing effect of multiple ethynyl-linked modifications might be caused by stabilizing π - π interactions of the ethynyl-linked modifications next to each other, whereas the flexible ethyl linkers cannot provide such stabilization and rather destabilize the duplex due to adding more energy to the system.

The previous study of nuclease resistance of hydrophobically modified ONs with the same modifications differing in ethynyl linkers³² showed no significant increase or decrease in enzymatic stability in comparison with non-modified ONs. Another study with 2'-O-alkylcarbamoyl ethyl modifications revealed

similar findings,⁴⁷ that length of the alkyl chain (C1–C5) did not significantly alter the nuclease resistance until longer chain was introduced (C8). We assume that the enzymatic stability of our ONs with ethyl-linked hydrophobic modifications would be showing similar trend.

Conclusions

We have synthesized a set of alkyl and aryethyl-linked pyrimidine and 7-deazapurine 2'-deoxyribonucleoside phosphoramidites and used them in solid-supported synthesis of modified and hypermodified oligonucleotides bearing flexible hydrophobic groups. The yields of the ONs bearing the flexible alkyl groups were mostly better than the previously reported ethynyl-linked ONs. The annealing of the partially modified ONs with non-modified complementary strands successfully formed DNA duplexes, whereas the hypermodified strands did not form double-stranded B-DNA. CD spectra for all partially modified DNA duplexes showed B-form like pattern. Apparently, the hypermodified ONs bearing multiple hydrophobic modifications in a row form aggregates in aqueous solution, as supported by DLS, that prevent hybridization and forming of dsDNA. However, partially modified ONs did not suffer from this problem, hence the potential usage of such hydrophobic modifications for partial modification of ONs and in selections remains to be feasible. The melting temperatures studies showed, that the overall stability of the modified duplexes with ethyl linker is lower than the stability of non-modified corresponding DNAs and of the ethynyl-linked modified DNA duplexes. The reported phosphoramidites will be useful reagents in solid-phase automated synthesis of partially modified ONs in aptamer engineering⁶ and other functional nucleic acids, as well as potential oligonucleotide therapeutics. For potential applications as therapeutics further studies on cellular uptake, nuclease resistance, and immunogenicity would need to be performed.

Conflicts of interest

The authors declare no competing financial interest.

Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information: synthetic protocols and characterization of new compounds, additional tables, chromatograms, melting curves, additional CD spectra and copies of NMR and MS spectra. See DOI: <https://doi.org/10.1039/d5ra06147d>.

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