Introduction

Duplex formation between complementary oligomers is one of the most important processes in biology, and has found a wide range of applications in nanotechnology. The nucleic acid duplex provides a robust architecture for encoding molecular information and for replication, transcription and translation of this information through template-directed synthesis.\(^1\)\(^2\) These properties are currently unique to nucleic acids, so it is not surprising that the first approaches to synthetic sequence-controlled macromolecules use biopolymers. A range of nucleic acid analogues that form duplexes have been prepared modifying the bases,\(^3\) the sugar,\(^4\) and the backbone.\(^5\) Recently, DNAtemplated polymerization and in vitro selection was used to evolve oligomers with chemically diverse side chains.\(^6\)

Synthetic oligomers that bear no relation to nucleic acids have also been shown to form duplexes via non-covalent interactions such as hydrogen bonding,\(^7\) aromatic interactions,\(^8\) salt bridges,\(^9\) and metal–ligand coordination.\(^10\) Sequence-selective duplex formation has been demonstrated for short sequences, showing that it is possible to read and write sequence information encoded into synthetic oligomers.\(^11\) We have been exploring the blueprint shown in Fig. 1 as a template for the design of duplex forming molecules. The recognition units that form the base-pairs in the duplex are displayed as side chains on the oligomers, so that the three key elements of the blueprint (the synthesis, backbone and recognition modules) can independently be optimised. A two-letter alphabet is used to encode information in an oligomer as a sequence of H-bond donor and acceptor sites. Provided the backbone does not contain any polar functional groups that could compete with H-bond interactions between the recognition units, reliable duplex assembly can be achieved in non-polar conditions.

![Fig. 1](image1.png)
solvents like toluene. We recently showed that the trifluoromethylphenol–phosphine oxide base-pair illustrated in Fig. 1 is sufficiently stable to allow duplex formation in more polar solvents like chloroform.

The properties of the backbone are also important, because there are different self-assembly channels that compete with duplex formation (Fig. 2). The parameters that determine the outcome of the self-assembly of recognition-encoded oligomers are the effective molarities for intramolecular folding (EM$_f$), the effective molarities for duplex formation (EM$_d$), the strength of the H-bond interaction ($K$) and the operating concentration ($c$). In the absence of geometric constraints, the values of EM are usually in the 10–100 mM range, so operating at millimolar concentrations avoids the intermolecular networks channel. If the backbone is too flexible, the folding pathway dominates due to intramolecular H-bonding interactions between donors and acceptors that are adjacent in sequence. The use of the long-short phenol–phosphine oxide base-pair illustrated in Fig. 1 reduces the probability of these 1,2-folding interactions and promotes duplex formation. It is also possible to decrease EM$_f$ by using a rigid backbone, but if the backbone is too rigid, duplex assembly fails. Duplex initiation, formation of the second base-pair, is governed by EM$_1$ (see Fig. 2) and is favourable in all of the systems that we have studied to date. However, duplex propagation, formation of subsequent base-pairs (EM$_2$ ... EM$_n$), is highly dependent on the conformational properties of the backbone. Thus, a semi-flexible backbone is required to ensure that the molecule can adjust to a conformation compatible with duplex formation.

Here we describe a new family of oligomers that form stable duplexes in chloroform solution. The long-short trifluorophenol–phosphine oxide base-pair illustrated in Fig. 1 is used for the recognition module, because we have previously shown that this system reduces folding and increases duplex stability. The backbone is assembled from two different components, a dialdehyde that carries the recognition units and a diamine linker (Fig. 1). Imine formation or reductive amination can be used for oligomer synthesis, and the two-component approach provides the opportunity to introduce variation in the properties of the backbone by changing the diamine without the need to synthesise new monomer building blocks. The reversibility of imine bond formation also opens up possibilities in dynamic covalent chemistry and template synthesis. Reduction of the imines will result in a backbone that contains secondary amines, which are both H-bond donors and acceptors. However, the use of aniline nitrogens ensures that there will be no competition with the base-pairing interactions between the recognition units: the H-bond acceptor parameter (β) for aniline is 5 compared with 10 for phosphine oxide, and the H-bond donor parameter (α) for aniline is 2 compared with 4 for trifluorophenol.

Results and discussion

Synthesis

The synthetic route to the mono-aldehyde and di-aldehyde phenol building blocks is shown in Scheme 1. 5-Bromosalicylaldehyde was alkylated with racemic 2-ethylhexyl bromide to give 1. 4-Bromo-2-(trifluoromethyl)phenol was converted to the boronic ester 2 under microwave irradiation with palladium catalysis. Suzuki–Miyaura coupling of 1 with 2 under microwave irradiation using palladium catalysis gave the mono-aldehyde.

Fig. 2 Self-assembly channels for recognition-encoded oligomers. The first base-pairing interaction can take place in an intramolecular fashion leading to the folding channel or in an intermolecular fashion. The second base-pairing interaction can take place in an intramolecular fashion to initiate duplex formation or in an intermolecular fashion leading to the networks channel. The outcome depends on the concentration, $c$, the association constant for the intermolecular base-pairing interaction, $K$, and the effective molarities for folding, EM$_f$, duplex initiation, EM$_1$ and duplex propagation, EM$_n$.

Scheme 1 Synthesis of phenol building blocks D' and D (R = 2-ethylhexyl).
phenol $D'$. 5-Bromo-2-hydroxyisophthalaldehyde $3$ was synthesised via a Duff reaction, by reacting $p$-bromophenol with an excess of hexamethylenetetramine. Subsequent alkylation with racemic 2-ethylhexyl bromide gave $4$, which was coupled with 2 under Suzuki–Miyaura conditions to give the di-aldehyde phenol $D$.

The synthetic route to the mono-aldehyde and di-aldehyde phosphine oxide building blocks is shown in Scheme 2. Treatment of diethyl phosphite with butylmagnesium chloride gave $5$. The mono-aldehyde phosphine oxide $A'$ was synthesised from 5-iodosalicylaldehyde by alkylation with racemic 2-ethylhexyl bromide and then coupling with 5 using palladium and XantPhos under microwave irradiation. Similarly, the di-aldehyde phosphine oxide $A$ was synthesised from 5-bromo-isophthalaldehyde by coupling with 5 using palladium and XantPhos under microwave irradiation.

Two different 1,3-phenylenediamines were used as linkers for the synthesis of oligomers: 5-(trifluoromethyl)-1,3-phenylenediamine, which is commercially available; and compound $7$, which was prepared by treatment of 1-(chloromethyl)-3,5-dinitrobenzene with racemic 2-ethylhexyl thiol to give $6$, and subsequent reduction with tin(ii) chloride to give $7$ (Scheme 3).

Scheme 4 shows the synthesis of 2-mers DD and AA. Reductive amination of $D'$ or $A'$ with 5-(trifluoromethyl)-1,3-phenylenediamine and NaBH(OAc)$_3$ gave DD and AA respectively in good yield.

Schemes 5 and 6 show the synthesis of 3-mers DDD and AAA. The phosphine oxide building blocks carry additional solubilising groups on the phosphorus, so 5-(trifluoromethyl)-1,3-phenylenediamine was used as the linker for synthesis of the acceptor 3-mer. In all cases, racemic 2-ethylhexyl solubilising groups were used, because the diastereoisomeric mixture improves solubility. However, the donor 3-mer required additional solubilising groups, so $7$, the diamine equipped with a solubilising group, was used as the linker for this oligomer. This group is too remote from the recognition groups to affect duplex formation. Treatment of the di-aldehyde building block ($D$ or $A$) with nine equivalents of the corresponding 1,3-phenylenediamine gave the di-imine, which was reduced with NaBH$_4$ to give $8$ (Scheme 5) or $9$ (Scheme 6). Treatment of $8$ with mono-aldehyde $D$ followed by reduction with NaBH$_4$ gave 3-mer DDD, and treatment of $9$ with mono-aldehyde $A'$ followed by reduction with NaBH$_4$ gave 3-mer AAA.

**NMR binding studies**

Complexation of length-complementary oligomers was studied using $^1$H and $^{19}$F NMR titration experiments. The association constant $K$ for formation of the $A-D$ complex, which makes a single intermolecular hydrogen bond, was measured by titrating $A$ into $D$ in toluene. Addition of $A$ caused an upfield change in the $^{19}$F NMR chemical shift of the signal due to the $D$ CF$_3$ group and a downfield change in the $^1$H-NMR chemical shift of the signal due to the $D$ NO$_2$ group.
shift of the signal due to the \( \text{DOH} \) group. The titration data fit well to a 1:1 binding isotherm, giving an association constant of \( 3 \times 10^3 \text{ M}^{-1} \) (Table 1). Although the aldehyde substituents on \( \text{A} \) and \( \text{D} \) differ from the reductive amination products, these groups have no effect on the H-bonding properties of the phosphine oxide and phenol recognition groups, and the association constant measured in toluene is identical to the value measured previously for the corresponding monomers with dialkylamino substituents.

Addition of \( \text{AA} \) into \( \text{DD} \) caused an upfield shift of the \( ^{19}\text{F} \) NMR signal due to the \( \text{CF}_3 \) group on the phenol recognition unit of \( \text{DD} \) and a small downfield shift of the signal due to the \( \text{CF}_3 \) group on the diamin linker unit. The data fit well to a 1:1 binding isotherm, giving an association constant of \( 5 \times 10^5 \text{ M}^{-1} \). The association constant measured for \( \text{AA} \cdot \text{DD} \) is two orders of magnitude higher than the value measured for \( \text{A} \cdot \text{D} \), which indicates that \( \text{AA} \cdot \text{DD} \) forms a fully-assembled duplex with cooperative formation of two H-bonds. The association constants were used to determine the effective molarity (EM) for formation of the second intramolecular H-bond in the \( \text{AA} \cdot \text{DD} \) duplex (eqn (1)).

\[
K_N = 2K_1^N EM^{N-1}
\]

where \( K_N \) is the association constant for duplex formation between two oligomers with \( N \) interaction sites, \( K_1 \) is the association constant for formation a single intermolecular H-bond in \( \text{A} \cdot \text{D} \), and two is the statistical factor 2 that takes into account the parallel–antiparallel degeneracy of the duplex.

For the \( \text{AA} \cdot \text{DD} \) duplex, EM is 31 mM, which is consistent with the values of EM that we have measured for duplex forming oligomers with different backbones and base-pairing systems (10–100 mM).\(^{12b,18}\) The chelate cooperativity associated with duplex formation is expressed as the product \( K_1 \) EM, which is equal to 80, implying that the doubly H-bonded closed duplex is almost exclusively populated (98%).\(^{12b} \) The complex formed by the two 3-mers \( \text{AAA} \) and \( \text{DDD} \) was too stable in toluene for measurement of the association constant by NMR titration, so all of the binding studies were repeated in chloroform.

Fig. 3 shows the NMR spectra for titration of \( \text{AA} \) to \( \text{DD} \) in chloroform. Addition of \( \text{AA} \) caused a large upfield change in the \( ^{19}\text{F} \) NMR chemical shift of the signal due to the \( \text{DD} \) \( \text{CF}_3 \) group and a large downfield change in the \( ^{1}H\)-NMR chemical shift of the signal due to the \( \text{DD} \) \( \text{OH} \) group. The titration data fit well to a 1:1 binding isotherm for all three length complementary complexes, and the results are collected in Table 1.

The association constants are significantly lower than the values measured in toluene, but the association constant for \( \text{AA} \cdot \text{DD} \) is still an order of magnitude higher than the value measured for \( \text{A} \cdot \text{D} \), indicating cooperative assembly of the \( \text{AA} \cdot \text{DD} \) duplex. Similarly, the association constant for \( \text{AAA} \cdot \text{DDD} \)

### Table 1: Association Constants (\( K \)), Effective Molarities (EM), Limiting NMR Chemical Shifts (\( \delta_{\text{free}} \) and \( \delta_{\text{bound}} \)), and Complexation-Induced Changes in Chemical Shift (\( \Delta \delta \)) for the Formation of Duplexes at 298 K\(^{16}\)

| Solvent | Complex | \( \log K \) (M\(^{-1}\)) | EM (mM) | \( \delta_{\text{free}} \) | \( \delta_{\text{bound}} \) | \( \Delta \delta \) | \( \delta_{\text{free}} \) | \( \delta_{\text{bound}} \) | \( \Delta \delta \) |
|---------|---------|------------------|--------|----------------|----------------|----------|----------------|----------------|----------|
| CDCl\(_3\) | \( \text{D} \cdot \text{A} \) | 2.3 ± 0.1 | — | −61.1 | −62.8 | −1.7 | 5.7 | 11.3 | 5.6 |
| | \( \text{DD} \cdot \text{AA} \) | 3.5 ± 0.1 | 35 | −61.0 | −62.3 | −1.2 | 5.5 | 10.3 | 4.8 |
| | \( \text{DDD} \cdot \text{AAA} \) | 4.4 ± 0.1 | 35 | −61.0 | −62.2 | −1.2 | 5.6 | 10.3 | 4.7 |
| Toluene | \( \text{D} \cdot \text{A} \) | 3.5 ± 0.1 | — | −61.8 | −62.3 | −0.5 | 4.9 | 11.6 | 6.6 |
| | \( \text{DD} \cdot \text{AA} \) | 5.7 ± 0.1 | 31 | −61.5 | −61.8 | −0.3 | 4.9 | 11.6 | 6.7 |

\( ^{19}\text{F} \) NMR titrations, with errors at the 95% confidence limit.\(^{16}\) Data for signals due to the \( \text{OH} \) and \( \text{CF}_3 \) groups on the phenol recognition units.
is an order of magnitude higher than the value measured for AA-DD, indicating that all three H-bonds are formed in the AAA-DD duplex. For all three complexes, the values of Δδ for the OH and CF₃ groups on the trifluorophenol recognition units are similar, which confirms that the donor recognition units are fully H-bonded in all of the complexes (Table 1). Using eqn (1) to determine EM gives a value of 35 mM for both duplexes in chloroform, which is similar to the value in toluene. However, the reduction in the association constant for H-bond formation in chloroform means that the chelate cooperativity associated with duplex formation ($K_{1}$ EM) is reduced by an order of magnitude to 7.

The fact that the EM for formation of the AAA-DD duplex is the same as the value for formation of the AA-DD duplex is an important result, which suggests that it should be possible to assemble stable duplexes using longer oligomers. As we have shown previously, duplex initiation is relatively insensitive to changes in the chemical shift at different temperatures between 253 and 363 K. The changes in the chemical shifts of the ¹⁹F NMR signals due to the phenol CF₃ groups are indicative of an increase in the population of H-bonded complexes at lower temperatures and disruption of the duplex at higher temperatures (Fig. 5). These melting data were fit to a two-state model, assuming that only duplex and denatured single strands are present, that the enthalpy and entropy changes for the formation of the N-mer duplex ($ΔH_m$ and $ΔS_m$) are temperature independent, and that the change in heat capacity between free and bound states is zero (eqn (2)).

$$\delta = \delta_i + (\delta_h - \delta_i) \times \frac{1}{1 + 4e^{-\frac{\Delta H_m}{R} \left(\frac{1}{T} - \frac{1}{T_{m,N}}\right)}} \frac{1}{1 + 8e^{-\frac{\Delta H_m}{R} \left(\frac{1}{T} - \frac{1}{T_{m,N}}\right)}}$$

(2)

Fig. 5 Experimental ¹⁹F NMR chemical shift plotted as a function of temperature for 1 : 1 mixtures (2 mM) of A-D (black), AA-DD (blue), and AAA-DD (red) in 1,1,2,2-tetrachloroethane. The lines are the best fit to eqn (2) (total rmsd < 0.01 ppm), and the transition melting temperatures are indicated with a bar. The optimised values of $T_{m,N}$ and $ΔH_m$ are 298 K and -37 kJ mol⁻¹ for A-D, 318 K and -51 kJ mol⁻¹ for AA-DD, and 331 K and -55 kJ mol⁻¹ for AAA-DD.

Thermal denaturation experiments

Thermal denaturation experiments were carried out to extract thermodynamic parameters for duplex assembly. ¹⁹F NMR spectra of 1 : 1 solutions of length-complementary oligomers at 2 mM concentrations in 1,1,2,2-tetrachloroethane were recorded at different temperatures between 253 and 363 K. The changes in the chemical shifts of the ¹⁹F NMR signals due to the

![Fig. 4 Structure of the AAA-DD duplex.](image)
series of homo-oligomers were synthesised, and duplex formation was characterised by NMR titration experiments. When length complementary oligo-trifluorophenols and oligo-phosphine oxides were combined, an order of magnitude increase in stability was observed for every base-pair added to the duplex. The effective molarity for the intramolecular H-bonds responsible for zipping up the duplex is about 30 mM in toluene and in chloroform. The uniform increase in duplex stability with oligomer length suggests that the backbone structure and geometry is likely to be compatible with the formation of extended duplexes in longer oligomers. This two-component backbone is more versatile than previous designs, because it provides an opportunity for varying the dopamine component without the need to resynthesise complex monomer building blocks. The properties of mixed sequence oligomers and template-directed synthesis using dynamic imine chemistry are currently under investigation.

Conflicts of interest

There are no conflicts to declare.

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