

# DNA-Organized Light-Harvesting Antennae: Energy Transfer in Polyaromatic Stacks Proceeds through Interposed Nucleobase Pairs

Caroline D. Bösch<sup>a</sup>, Elif Abay<sup>a</sup>, Simon M. Langenegger<sup>a</sup>, Maryam Nazari<sup>b</sup>, Andrea Cannizzo<sup>b</sup>, Thomas Feuerer<sup>b</sup>  
and Robert Häner<sup>\*,a</sup>

<sup>a</sup> Department of Chemistry and Biochemistry, University of Bern, Freiestrasse 3, CH-3012 Bern, Switzerland, E-mail: robert.haener@dcb.unibe.ch

<sup>b</sup> Institute for Applied Physics, University of Bern, Sidlerstrasse 5, 3012 Bern

Dedicated to Professor Philippe Renaud on the occasion of his 60<sup>th</sup> birthday

DNA-based light-harvesting antennae with varying arrangements of light-absorbing phenanthrene donor units and a pyrene acceptor dye were synthesized and tested for their light-harvesting properties. Excitation of phenanthrene is followed by rapid transfer of the excitation energy to the pyrene chromophore. A block of six light-absorbing phenanthrenes was separated from the site of the acceptor in a stepwise manner by an increasing number of intervening AT base pairs. Energy transfer occurs via interposed AT base pairs and is still detected when the phenanthrene antenna is separated by 5 AT base pairs.

**Keywords:** DNA • light-harvesting antenna • phenanthrene • pyrene • energy transfer

## Introduction

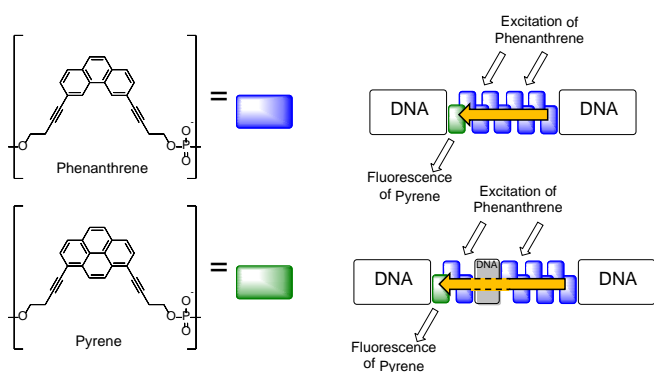
DNA is a useful framework for the precise arrangement of chromophores and can, thus, serve in the development of novel types of nanomaterials.<sup>[1-9]</sup>

Ordered arrays of chromophores are an essential requirement for the construction of artificial light-harvesting systems. In such systems, energy is absorbed by chromophores and subsequently transferred to an acceptor. The design and construction of efficient artificial systems is a continuing field of research.<sup>[10-23]</sup> In previous work, we described various light-harvesting systems based on DNA-organized chromophores.<sup>[21,24,25]</sup> In one approach, 3,6-dicarboxamide phenanthrene and 1,8-dicarboxamide pyrene were incorporated into DNA duplexes to build up light-harvesting antennae.<sup>[21]</sup> Light is absorbed by a stack of phenanthrenes and subsequently transferred to a phenanthrene-pyrene exciplex. The fluorescence intensity showed to be proportional to the number of light-absorbing phenanthrenes (up to eight phenanthrenes were used in a duplex). Another system describes the use of 1,8-dicarboxamide pyrene units and a cyanine dye (Cy5) in a DNA-assembled light-harvesting complexes (LHC).<sup>[24]</sup> Duplexes with different numbers of pyrene and a 5'-terminally attached Cy5 unit at one end were constructed. After excitation of pyrene, the excitation energy is transferred by FRET from pyrene excimers to the Cy5 acceptor. Furthermore, a DNA three-way junction was used for the arrangement of chromophores for the construction of light-harvesting antennae.<sup>[25]</sup> A 3,6-dicarboxamide phenanthrene antenna is located in one of the three stems and an exchangeable acceptor is placed in proximity through the annealing of the third strand. Here, we present a DNA based light-harvesting antenna consisting of a pyrene acceptor and a stack of light-harvesting phenanthrene units, which is interrupted by a varying number of DNA base pairs (see Figure 1 for illustration). In contrast to previous projects 3,6-dialkynyl phenanthrene is used as the light-

absorber as it has a higher absorption coefficient than the carboxamide derivative and was shown to stack more efficiently.<sup>[26]</sup> A 1,8-dialkynyl pyrene was used as the acceptor dye. A series of duplexes was prepared in which the phenanthrene antenna is split in two parts by 1-5 A-T base pairs. The study allows addressing the question if excitation energy transfer along an array of phenanthrenes also takes place over a certain number of intervening base pairs.

## Results and Discussion

DNA strands were synthesized by phosphoramidite chemistry using phenanthrene, pyrene and abasic site building blocks. Complementary DNA single strands were hybridized to yield duplexes with eight light-collecting phenanthrene units and one pyrene acceptor (Table 1, duplexes **0-5**). An abasic site analogue (**ϕ**) was introduced opposite the pyrene building block to fix its positioning at the end of the chromophore stack. Only A-T base pairs were used in the vicinity of the chromophores, as the quenching effect of G is well known.<sup>[27,28]</sup> However, G-C base pairs were used at the ends to increase the stability of the duplexes. In duplex **0**, the phenanthrenes form an uninterrupted stack, whereas in duplexes **1-5** the phenanthrenes are separated by one to five A-T base pairs. Additionally, two control duplexes were prepared, duplex **Ref1** containing only the acceptor part (two phenanthrenes and one pyrene) and duplex **Ref2** containing only a block of six continuous phenanthrenes.



**Figure 1.** A) Structures of DNA-incorporated phenanthrene and pyrene building blocks; B) Schematic illustration of DNA based light-harvesting antennae without (top) and with (below) separating base pairs between phenanthrene units. Arrows indicate energy transfer.

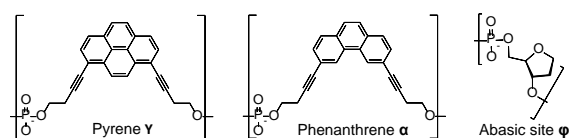
Fluorescence spectra of the duplexes were recorded by exciting phenanthrene at 316 nm (Figure 2). All duplexes exhibit mainly pyrene emission with maxima at 403 and 425 nm, indicating efficient energy transfer from phenanthrene to pyrene. The ratio of intensities of the two peaks varies within the series of duplexes. Duplexes **o-2** show higher intensity at 425 nm compared to 403 nm, duplex **3** shows roughly the same intensities, whereas for duplexes **4** and **5** the emission at 403 nm is highest. This observation can be explained by different environments (next neighbors) for pyrene which can lead to different vibronic band intensities. A similar effect was observed for pyrene fluorescence in different solvents.<sup>[29]</sup>

In general, the duplexes with zero, one and two separating base pairs have about the same fluorescence intensities. If the stack of phenanthrene is separated by three base pairs the intensity decreases. Four or five base pairs lead to a further decrease of pyrene emission, whereas there is no significant difference between these two duplexes. At the same time, phenanthrene monomer emission around 380 nm increases if the phenanthrenes are separated by more than three base pairs (see Figure 2, inset). Therefore, energy transfer is not much affected by interruption of the phenanthrene stack by up to two base pairs. Further elongation of the gap, however, is followed by a decrease of energy transfer.

**Table 1.** Sequences of phenanthrene and pyrene modified DNA strands and structures of unnatural building blocks.

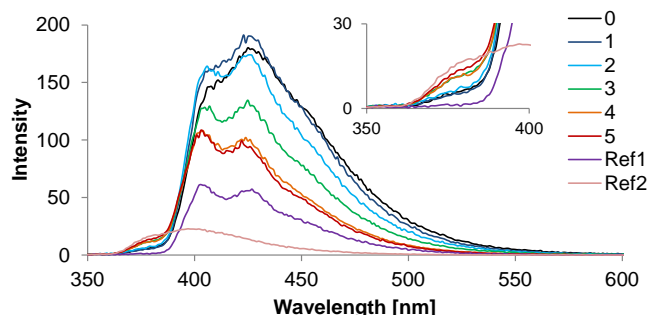
Duplex	Oligo	Sequences
<b>o</b>	<b>o_a</b>	5' GGC TAA $\phi\alpha\alpha$ $\alpha\alpha$ A TTA AAT CGC 3'
	<b>o_b</b>	3' CCG ATT $Y\alpha\alpha$ $\alpha\alpha$ T AAT TTA GCG 5'
<b>1</b>	<b>1_a</b>	5' GGC TAA $\phi\alpha$ T $\alpha\alpha\alpha$ TTA AAT CGC 3'
	<b>1_b</b>	3' CCG ATT $Y\alpha\alpha$ $\alpha\alpha\alpha$ AAT TTA GCG 5'
<b>2</b>	<b>2_a</b>	5' GGC TAA $\phi\alpha$ T $A\alpha\alpha$ $\alpha$ TA AAT CGC 3'
	<b>2_b</b>	3' CCG ATT $Y\alpha\alpha$ $T\alpha\alpha$ $\alpha$ AT TTA GCG 5'
<b>3</b>	<b>3_a</b>	5' GGC TAA $\phi\alpha$ T AT $\alpha$ $\alpha\alpha\alpha$ AAT CGC 3'
	<b>3_b</b>	3' CCG ATT $Y\alpha\alpha$ T $\alpha\alpha$ $\alpha\alpha$ T TTA GCG 5'

<b>4</b>	<b>4_a</b>	5' GGC TAA $\phi\alpha$ T ATA $\alpha\alpha\alpha$ AAT CGC 3'
	<b>4_b</b>	3' CCG ATT $Y\alpha\alpha$ TAT $\alpha\alpha\alpha$ TTA GCG 5'
<b>5</b>	<b>5_a</b>	5' GGC TAA $\phi\alpha$ T ATA T $\alpha\alpha$ $\alpha$ AT CGC 3'
	<b>5_b</b>	3' CCG ATT $Y\alpha\alpha$ TAT A $\alpha\alpha$ $\alpha$ TA GCG 5'
<b>Ref1</b>	<b>Ref1_a</b>	5' GGC TAA $\phi\alpha$ T ATA TTA AAT CGC 3'
	<b>Ref1_b</b>	3' CCG ATT $Y\alpha\alpha$ TAT AAT TTA GCG 5'
<b>Ref2</b>	<b>Ref2_a</b>	5' GGC TAA AT $\alpha$ $\alpha\alpha\alpha$ TTA AAT CGC 3'
	<b>Ref2_b</b>	3' CCG ATT T $\alpha\alpha$ $\alpha\alpha$ T AAT TTA GCG 5'

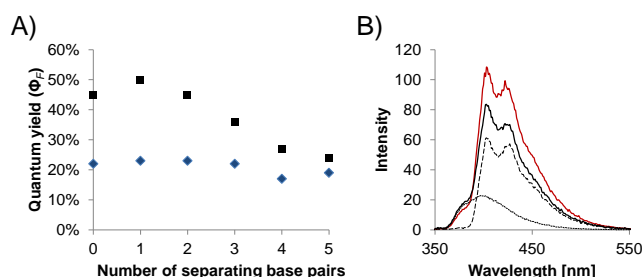


Quantum yields ( $\Phi_F$ , Table 2) were measured for all single strands and duplexes by phenanthrene excitation at 316 nm. Single strands containing only phenanthrene have values around 5%, whereas single strands with pyrene have significant higher quantum yields (29-46%), which again provides evidence for efficient excitation energy transfer from phenanthrene to pyrene. This increase in quantum yield cannot be explained by FRET theory.<sup>[30,31]</sup> The observed efficiency rather suggests that excitation energy transfer occurs by quantum coherence mechanism. It is feasible that electronic coupling between the stacked phenanthrenes in the framework of the DNA duplex is strong enough to enable coherent energy transfer.<sup>[32-37]</sup> This would also explain why the quantum yield of duplex **o** is lower than that of duplex **1**. In duplex **o** the phenanthrene units are not separated by base pairs, therefore, electronic coupling should be strongest in this duplex. Very strong electronic coupling was shown to favor relaxation rather than energy transfer.<sup>[38]</sup> Hence, energy transfer is optimal in duplex **1** and separation by additional base pairs results in reduced efficiency. Figure 3 A illustrates the measured (black) and calculated (blue) quantum yields of the duplexes. The deviation between the two values decreases with increasing number of separating base pairs. Using the quantum yields of duplexes **Ref1** and **Ref2** as a model for "infinite" separation of donor and acceptor, the theoretical quantum yield for no energy transfer was calculated to be 20% (see the *Experimental Section*). This is only slightly lower than the quantum yield of duplex **5** ( $\Phi_F = 24\%$ ) and a value around 20% seems to be approached in the series with a growing number of intervening base pairs. Nevertheless, the measured quantum yield of duplex **5** is higher than the average of both single strands. Additionally, as can be seen in Figure 3 B, duplex **5** (red) shows a higher pyrene and lower phenanthrene fluorescence than the arithmetic sum of duplexes **Ref1** and **Ref2** (black, solid). This indicates that, although the phenanthrene units are separated by five base pairs, excitation energy transfer still takes place, albeit at a reduced level. Table 2 also shows the melting temperatures ( $T_m$ ) of all duplexes. Generally, the hybrids in which A-T base pairs separate the phenanthrene units (hybrids 1-5) are less stable compared to the hybrid in which phenanthrenes and pyrene form a continuous stack (hybrid **o**). The lowest

$T_m$  values were determined for duplexes 2 and 3, in which the phenanthrenes are separated by two and three base pairs, respectively. All melting curves show no or only a negligible degree of hysteresis (see the Supplementary Material) indicating that duplex formation and disassembly are at thermodynamic equilibrium under experimental conditions.<sup>[39]</sup>



**Figure 2.** Fluorescence spectra of DNA-assembled light harvesting antennae. Conditions: 0.25  $\mu$ M each strand, 10 mM sodium phosphate buffer pH 7.0, 100 mM NaCl,  $\lambda_{exc}$  316 nm, 20°C. Excitation slit: 2.5 nm, emission slit: 5 nm.



**Figure 3.** A) Quantum yields of duplexes 0 – 5 (black) and the average quantum yield of the respective single strands 0\_b – 5\_b (blue, see Table 2). B) Comparison of the measured fluorescence spectrum of duplex 5 (red) with the fluorescence spectra of Ref1 (dashed) and Ref2 (dotted) and the arithmetic sum of Ref1 and Ref2 (black). Conditions: see Figure 2.

**Table 2.** Quantum yields of modified single strands and duplexes (average value of single strands and measured) and melting temperatures ( $T_m$ ) of duplexes.

Duplex	Oligo	$\Phi_F$ (single strands)	$\Phi_F$ (avg. of single strands) <sup>a</sup>	$\Phi_F$ (duplexes)	$T_m$
0	0_a	6%	22%	45%	51°C
	0_b	37%			
1	1_a	4%	23%	50%	49°C
	1_b	42%			
2	2_a	5%	23%	45%	45°C
	2_b	41%			
3	3_a	6%	22%	36%	45°C
	3_b	38%			

4	4_a	5%	17%	27%	46°C
	4_b	29%			
5	5_a	4%	19%	24%	48°C
	5_b	33%			
Ref1	Ref1_a	6%	26%	56%	51°C
	Ref1_b	46%			
Ref2	Ref2_a	5%	5%	8%	54°C
	Ref2_b	5%			

<sup>a</sup> Calculated from the quantum yields of individual complementary strands.

## Conclusions

In conclusion, a series of DNA-based light-harvesting antennae composed of light-collecting phenanthrenes and pyrene acceptors were synthesized and investigated. Excitation of phenanthrene leads to energy transfer to pyrene which then results in pyrene emission. Energy transfer is observed if the stack of phenanthrene units is interrupted by up to five DNA base pairs. The separation by one and two base pairs shows no decrease in energy transfer efficiency compared to a continuous phenanthrene stack. The light-harvesting antenna interrupted by one base pair has the highest efficiency of all tested duplexes. A separation by 3 to 5 base pairs leads to a gradual decrease in efficiency but energy transfer is still detected with 5 intervening base pairs. The data demonstrate that it is possible to assemble phenanthrene-based light-harvesting antennae which are mixed with natural DNA base pairs. This extends the range of potential designs and applications of DNA-based artificial light-harvesting complexes.

## Experimental Section

### General Methods

All reagents and solvents were purchased from commercial suppliers and used without further purification. Mass-spectrometric data were obtained on Thermo Fisher LTQ Orbitrap XL using Nano Electrospray Ionization (NSI). UV-vis spectra were measured on a Cary 100 Bio spectrophotometer. Fluorescence and excitation spectra were measured on a Cary Eclipse spectrofluorimeter, excitation and emission slit widths were set to 5 nm if not mentioned otherwise.

### DNA Synthesis and Purification

Syntheses of phenanthrene and pyrene phosphoramidites were described previously.<sup>[40,41]</sup> The phosphoramidite derivative of building block  $\phi$ , representing a stable analog of an abasic site, is commercially available (Glen Research). Modified DNA strands were assembled on an Applied Biosystems 394 DNA/RNA synthesizer. A standard cyanoethyl phosphoramidite coupling protocol was applied using a 3'-attached, nucleoside-loaded controlled pore glass (CPG) support. After synthesis, the CPG-bound strands were cleaved and deprotected by treatment with 28-30% NH<sub>4</sub>OH (aq) at 55°C overnight. The supernatants were collected. The solid supports were additionally washed three times with 1 ml EtOH/H<sub>2</sub>O 1:1. After combination and lyophilization of the supernatants, the crude oligomers were purified by reversed phase HPLC (Merck LiChroCART 250-4; LiChrospher 100, RP-18, 5  $\mu$ m); solvent A: 0.1 M aqueous ammonium acetate; solvent B: CH<sub>3</sub>CN; 1 ml/min; T = 40°C; B[%] ( $t_R$  [min]) = 0 (0); 5 (2); 50 (22). Identities of the oligomers were confirmed by ESI mass spectrometry (see the *Supplementary Material*). The samples were measured in negative ion mode in mixtures of water/acetonitrile/triethylamine. The purified oligomers were dissolved in 1 ml Milli-Q H<sub>2</sub>O. Samples of the stock solutions were diluted and the absorbance at 326 nm was measured to determine the concentrations. The molar absorption coefficients of the oligomers were calculated using the  $\epsilon_{326}$  value of 35'400 for phenanthrene.

### Quantum Yields

The quantum yields (Table 2) were determined using quinine sulphate in 0.5 M H<sub>2</sub>SO<sub>4</sub> as a standard according to the procedure described in the literature.<sup>[42]</sup> The theoretical quantum yield for 'no energy transfer' was calculated assuming that the duplexes **Ref1** and **Ref2** do not interfere with each other using the following equations:

$$(eq. 1) \Phi_{Ref1} = I_{Ref1}/A_{Ref1}$$

$$(eq. 2) \Phi_{Ref2} = I_{Ref2}/A_{Ref2}$$

$$(eq. 3) \Phi_{noET} = (I_{Ref1} + I_{Ref2})/(A_{Ref1} + A_{Ref2})$$

$$(eq. 4) I_{Ref1} = A_{Ref1} * \Phi_{Ref1} \quad I_{Ref2} = A_{Ref2} * \Phi_{Ref2}$$

$$(eq. 5) \Phi_{noET} = (A_{Ref1} * \Phi_{Ref1})/(A_{Ref1} + A_{Ref2}) + (A_{Ref2} * \Phi_{Ref2})/(A_{Ref1} + A_{Ref2})$$

$A_{Ref1} = 2$  and  $A_{Ref2} = 6$  (respective number of phenanthrenes)

$$(eq. 6) \Phi_{noET} = \Phi_{Ref1}/4 + 3 * \Phi_{Ref2}/4 = 20\%$$

### Supplementary Material

Supporting information for this article is available on the WWW under <http://dx.doi.org/10.1002/MS-number>.

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### Author Contribution Statement

C.D.B, E.A. and M.N. performed the experiments and analyzed the data.  
C.D.B, S.M.L., A.C., T.F. and R.H. conceived and designed the experiments, and wrote the paper.

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