

Expanded Genetic Alphabet

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The Expanded Genetic Alphabet

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All biological information, since the last common ancestor of all life on Earth, has been encoded by a genetic alphabet consisting of only four nucleotides that form two base pairs. Long-standing efforts to develop two synthetic nucleotides that form a third, unnatural base pair (UBP) have recently yielded three promising candidates, one based on alternative hydrogen bonding, and two based on hydrophobic and packing forces. All three of these UBPs are replicated and transcribed with remarkable efficiency and fidelity, and the latter two thus demonstrate that hydrogen bonding is not unique in its ability to underlie the storage and retrieval of genetic information. This Review highlights these recent developments as well as the applications enabled by the UBPs, including the expansion of the evolution process to include new functionality and the creation of semi-synthetic life that stores increased information.

1. Introduction and Overview

DNA encodes life, and no other molecule has captured the attention and imagination of scientists and the general public more. However, as a polymer, DNA is chemically and physically rather unremarkable, consisting of nothing more than a linear sequence of the deoxyribonucleotides dG, dC, dA, and dT. What makes DNA remarkable, and absolutely unique, is its sequence-specific formation of a duplex and its sequence-specific replication, as well as its sequence-specific transcription into RNA, which consists of the ribonucleotides G, C, A, and U. This sequence specificity is made possible by the selective pairing of the four letters of the genetic alphabet to form two base pairs; (d)G pairs with (d)C and (d)A pairs with dT or U. The same base pairing between mRNA and tRNA enables decoding of the nucleotide sequences into protein sequences. It is the selective formation of these two base pairs that allows this otherwise unremarkable molecule to encode the diversity of life.

The development of synthetic nucleotides that form a third, unnatural base pair (UBP), and thus expand the genetic alphabet, has been a goal of chemists since the idea was suggested over half a century ago.^[1] A UBP would have wide-ranging in vitro applications, especially if one or both of the constituent nucleotides could be derivatized with linkers for the site-specific attachment of groups with interesting physicochemical properties. Moreover, the development of a UBP that functions within a cell has emerged as a central goal of the developing field of synthetic biology. Such a UBP would allow for the creation of semi-synthetic organisms that store increased information in their DNA, and thus form the foundation of semi-synthetic life with the potential to possess new and useful attributes or functions.

While nucleotides with modified sugars^[2–5] or phosphates^[6,7] have been explored extensively and have important applications, the generation of a UBP requires nucleotides with modified nucleobases, as it is the nucleobases that mediate base pair formation. A wide variety of synthetic nucleotides bearing nucleobase analogues have been reported,^[8–10] and several have been identified that stably

pair with one another within an otherwise natural duplex.^[11–19] However, efficient and high-fidelity replication of DNA containing a UBP has proven much more challenging, and early studies by the Benner group yielded what were for over a decade the only promising candidates.^[20,21] These early UBPs were designed to interact through complementary hydrogen-bonding patterns not employed by the natural nucleotides, but there is no reason to assume a priori that hydrogen bonding is the only force sufficient to underlie the storage and retrieval of genetic information. For example, we^[22] and others^[12,23–26] have explored metal-dependent pairing. However, hydrophobic and packing forces (including ring-stacking) have been explored most intensively, and along with optimized versions of the original alternative hydrogen-bonding UBPs, several of these predominantly hydrophobic UBPs now form a small group of candidates that have been developed to a high level of proof-of-concept (Figure 1). Progress in the development of these UBPs through the middle of the last decade has already been reviewed.^[27–30] Here, we focus on more recent developments, with earlier studies included only for context. Also for context, we include a discussion of the forces that likely underlie the replication of DNA containing predominantly hydrophobic UBPs. Lastly, we present a discussion of currently emerging applications, culminating in the recent creation of the first semi-synthetic organism that stably harbors a UBP in its DNA.

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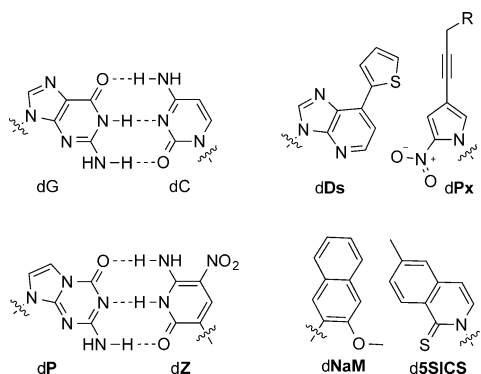


Figure 1. Natural dG–dC base pair and dZ–dP, dDs–dPx (R = H or –CH(OH)–CH₂OH), and dNaM–d5SICS UBPs. Sugar and phosphate groups are omitted for clarity.

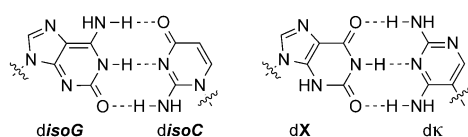


Figure 2. UBPs disoC–disoG and dX–dK. Sugar and phosphate groups are omitted for clarity.

2. Alternative Hydrogen-Bonding UBPs

In 1962, Alexander Rich proposed that *disoC* and *disoG*, constitutional isomers of dC and dG (Figure 2), should be capable of selective pairing through complementary hydrogen-bonding patterns similar to, but distinct from, those employed by the natural base pairs.^[1] However, it was not until three decades later that the Rappaport^[31,32] and Benner^[20,21] research groups independently explored the development of such UBPs. In particular, the early progress by Benner and co-workers, initially with *disoC*–*disoG*, and then with dK–dX, remained state-of-the-art for over a decade (Figure 2). Highlights of this seminal work include the demonstration that DNA containing *disoC*–*disoG* was at least reasonably well replicated by the Klenow fragment of *Escherichia coli* DNA polymerase I (Kf)^[20] and the later demonstration that *isoC* could be decoded at the ribosome with cell extracts and synthetic tRNA.^[33]

The major limitations of these early UBP candidates were the deamination and epimerization of *disoC* and dK; the

mispairing of *disoG* and dT/U, as a result of the tautomerization of *disoG*; and the poor transcription of DNA containing *disoG* into RNA containing *isoC*.^[34,35] The mispairing problem was later partially circumvented by the use of 2-thiothymidine deoxyribonucleoside triphosphate instead of natural dTTP, which made possible the first polymerase chain reaction (PCR) mediated amplification of DNA containing a UBP, a landmark in the development of UBPs.^[36] Nonetheless, the fidelity of UBP replication was only 98 %^[36] (throughout, fidelity refers to the retention of the UBP per doubling of the DNA), which translates into a third of the UBPs being lost after 20 PCR doublings ($0.98^{20} = 0.668$ retention) and is too low for most applications. More recently, Benner and co-workers reported the dZ and dP nucleotides (Figure 1),^[37,38] which are more stable, do not epimerize, and form a UBP that, when incorporated into DNA, is PCR amplified with a fidelity of 99.8 %.^[39] Very recently, the transcription and reverse transcription of dZ–dP was reported.^[40]

In 2000, the Hirao group reported the development of an orthogonal hydrogen-bonding UBP formed between the purine analogue dX and the pyridone analogue dY (Figure 3).^[41,42] To decrease the mispairing of dX with natural

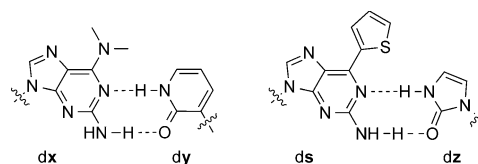
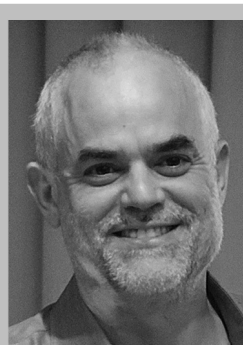


Figure 3. dX–dY and dS–dZ UBPs. Sugar and phosphate groups are omitted for clarity.

nucleotides, they developed the dS–dY pair the following year by replacing the dimethylamino group of dX with the more bulky and aromatic thiophenyl moiety of dS (Figure 3).^[43] Although mispairing of dY, mostly with dA, precluded PCR amplification, DNA containing dS was successfully transcribed into mRNA containing y, which was then translated with the aid of a synthetic tRNA containing s into a protein with the unnatural amino acid 3-chlorotyrosine.^[44] Conversely, DNA containing dY was transcribed into RNA containing s with low fidelity. To reduce the mispairing of dY with dA, dZ was developed to pair with dS by replacing the pyridone scaffold with a smaller cyclic dehydroimidazolone



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scaffold (Figure 3); however, both **dzTP** and **zTP** were recognized inefficiently by DNA and RNA polymerases, respectively.^[45]

3. Conceptual Interlude

The forces underlying the replication of natural DNA clearly include the complementary hydrogen bonds that link the pairing natural nucleobases. However, more than 15 years ago the Kool group reported the remarkable observation that Kf polymerase selectively pairs dA with dF, whose difluorotoluene nucleobase is a shape mimic of thymine, in which the hydrogen donor group N–H has been replaced with a C–H group and the hydrogen-accepting carbonyl groups have been replaced with C–F bonds.^[46,47] Although the ability of the fluorine atom to participate in hydrogen bonding was, and still is, actively debated,^[48,49] it is clear that the F...H hydrogen bond is much weaker than the bonds it replaces (3.8 kcal mol^{−1} less stable according to recent estimates).^[49] Although DNA primers terminating with the dF–dA pair are not efficiently extended, the selective pairing of dF with dA was interpreted as evidence of a “geometrical selection” mechanism of DNA replication (which was also independently proposed by Goodman)^[50,51] and it clearly demonstrates that forces other than hydrogen bonding also contribute to DNA synthesis. Such forces might be harnessed to control the replication of a UBP.

It has long been known that hydrogen bonding is not the sole force contributing to duplex stability.^[52–56] However, caution must be taken when considering how these forces might contribute to replication, because the polymerase active site provides a unique (and incompletely understood) environment. For example, it was noted decades ago that the destabilization associated with forming a mispair in duplex DNA could not account for the fidelity of DNA replication.^[57,58] Nonetheless, given that DNA synthesis involves triphosphates leaving their aqueous environment and developing at least some of the same intrabase interactions that stabilize a DNA duplex, it seems likely that some of the same forces may contribute to DNA synthesis.^[59]

One prominent force that favors duplex formation is that associated with the hydrophobic effect.^[54,60,61] Qualitatively, the hydrophobic effect may be understood as an interaction that causes hydrophobic moieties to aggregate or cluster in water, and it plays a central role in micelle formation, as well as in many biological processes such as membrane formation, protein folding, and molecular recognition. The “water ordering” or “iceberg” model, first proposed by Frank and Evans in 1945,^[62] still constitutes an important part of our understanding of the hydrophobic effect. This model posits that a nonpolar molecule has a tendency to induce the ordering of nearby water molecules into a crystalline- or clathrate-like cage structure such that a maximal number of hydrogen bonds may be retained, despite the production of the requisite cavity. This mechanism nicely explained the initially surprising result that the hydrophobic effect could be driven by entropy. However, with further studies it became clear that increasing the temperature or the size of the

hydrophobic solute could cause the hydrophobic effect to be driven predominantly by enthalpy changes. Thus, the original iceberg model was extended to allow for variable clathrate stability.^[63–66] Specifically, it has been postulated that the hydrogen bonds of the clathrate structure are disrupted before those of bulk water as the temperature is increased, and hydrogen-bonding networks cannot be maintained around large, hydrophobic molecules, especially those with low surface curvature. Thus, at high temperatures or with sufficiently large nonpolar molecules, less order is induced and the positive entropy of mixing or the negative enthalpy associated with hydrogen-bond rupture are more likely to dominate the associated change in free energy. Although a recent study has suggested that the binding of hydrophobic molecules to ordered environments such as proteins or DNA might also be favored by the displacement of high-energy water molecules,^[67] the revised iceberg model is currently the only widely accepted model of the hydrophobic effect.

A variety of studies have demonstrated that the hydrophobic effect may be harnessed for the stable pairing of nucleotides bearing predominantly hydrophobic nucleobases. The Kool group used a “dangling-end” assay, wherein the nucleotide of interest is attached to the end of one strand of a duplex where it can interact with its flanking nucleobase but has no hydrogen-bonding partner, to demonstrate that duplex stabilization was uniquely correlated with nucleobase surface area, which led them to conclude that it was mediated predominantly by the hydrophobic effect.^[68] In another study by Leumann and co-workers, an entropic origin for the stability of DNA containing pairs formed between nucleotides bearing fluorinated biphenyl rings was also interpreted as evidence of the hydrophobic effect.^[69] The importance of the hydrophobic effect in the pairing of nucleotides with hydrophobic nucleobases has been further supported by computational studies of Hobza and co-workers.^[70,71]

Although there is a consensus that the hydrophobic effect favors the sequestration of the aromatic faces of the nucleobases within the hydrophobic core of the duplex, the exact nature and role of the non-hydrogen-bonding forces that arise between the nucleobases have been more controversial. Early views centered around two opposing perspectives. The first emphasized nucleobase overlap and dispersive forces (i.e. electron correlation),^[72] while the second focused on interactions between the permanent dipoles associated with the exocyclic groups of one nucleobase and the delocalized electron density of the other (i.e. induction).^[73] The latter appeared consistent with structural data (which showed that polar exocyclic groups tend to be positioned above the rings of neighboring nucleobases),^[73,74] but increasingly sophisticated theoretical approaches have consistently identified dispersive interactions as being the most important.^[75–77] For example, computational studies from Hobza and co-workers suggest that while the hydrophobic effect is the dominant force driving nucleobase association, dispersion interactions determine the specific structure adopted.^[78] In something of a compromise between the original perspectives, recent theoretical studies based on the topological analyses of calculated electron density have suggested that the bonding interactions between flanking nucleobases are

manifest between specific atoms, especially the ring nitrogen atoms and the polar amino and carbonyl substituents.^[79–81] These results refocus attention on local interactions between specific ring atoms and substituents of the nucleobases.

The overlap of nucleobases within a DNA duplex, which are at least partially developed during replication, are an example of a broader class of molecular interactions observed between aromatic rings. These interactions have been studied intensively for decades, and are typically referred to as π - π stacking, although recent studies have suggested that this term is overused, and even potentially misused.^[82,83] The conventional model of Hunter and Sanders emphasized the repulsion between π clouds and focused on electrostatic quadrupole–quadrupole interactions that result in favorable interactions between the negatively charged π electrons of one ring and the positively charged σ framework of the other.^[84] These studies neatly explained the otherwise perplexing edge-to-face geometry observed with benzene, and the offset, or parallel-displaced, geometry observed with many other aromatic compounds. However, more recent theoretical studies have again emphasized the importance of the dispersion interactions that result from the close contact of the interacting rings, which were difficult to capture in early studies due to the high levels of theory required.^[85–88]

The effects of derivatizing the stacking aromatic rings with different substituents have also been studied intensively with small molecules. The traditional view, exemplified by the Hunter–Sanders model^[84,89–92] and the polar/ π model of Cozzi and Siegel^[93–97] describes the effects of substituents in terms of their nonlocal effect on the repulsive interactions between aromatic electrons through polarization and resonance effects within the ring to which they are attached. Thus, electron-withdrawing substituents are predicted to stabilize the interaction between two rings, and electron-donating substituents are predicted to destabilize it. However, high-level calculations have now decisively demonstrated that both electron-withdrawing and -donating substituents can stabilize the interaction of two aromatic rings.^[98–101] Calculations from the Wheeler and Houk groups^[102–105] explained these results by suggesting that substituent effects are not manifested through the aromatic rings and associated resonance effects, but instead arise from direct interactions of the local dipole associated with the substituent and the electric field of the other ring, as well as additional local dispersive interactions mediated by the polarized bond. The Wheeler–Houk model was recently supported and refined by high-level calculations from Parrish and Sherrill.^[106] Most recently, experimental studies have suggested that substituent effects are mostly additive, which seems inconsistent with their being mediated by resonance effects, and strongly supports the Wheeler–Houk model.^[107]

The interactions of heterocyclic aromatic compounds have also been studied. Calculations have suggested that the addition of nitrogen atoms favors electrostatic interactions, but that this stabilization is offset by increases in exchange repulsion, which along with only small changes in induction terms, make dispersion interactions dominant.^[108] In contrast, Sherrill reported calculations that suggest that aza substitution weakens both dispersion and exchange-repulsion forces,

and that electrostatic effects of aza substitution are governed by direct interactions between the localized electron density of the aza substituent and an electron-deficient hydrogen atom of the interacting ring.^[109] Liedl and co-workers recently reported the effects of heteroatom substitution on the stability of aromatic rings interacting through a parallel-displaced geometry.^[110] Generally, their results predict that the position of substitution has a large effect on the stability of the complex, as placing an electronegative atom, (especially a nitrogen atom) more proximal to the center of high electron density of the interacting ring was destabilizing. The stabilities and the specific geometries of the favored structures were interpreted in terms of nondirectional dispersive interactions and more directional electrostatics, and the differences in stabilities and structures were strongly correlated with the magnitude of the dipole moment associated with the heteroatom.

Finally, experimental studies with model systems have also been used to study the interactions between aromatic rings.^[111–115] Generally, these studies are consistent with the current perspective provided by theory: a dominant role for the hydrophobic effect, which is fine-tuned by dispersive and local electrostatic interactions.

Collectively, these studies suggest that in addition to complementary hydrogen bonding, the hydrophobic effect, desolvation, dispersive, and electrostatic interactions between nucleobases can contribute to the stability of a DNA duplex. What remained unclear was whether these forces can mediate efficient replication. Moreover, because the contribution of these forces to duplex stability is at least partially independent of sequence, it was even less clear if these forces could contribute to sequence-specific replication. For example, while Kool's results with dA with dF illustrate that, at least in some cases, these forces are sufficient to mediate triphosphate insertion, they also suggest that they may not be sufficient to prevent mispairing with a natural nucleotide (from the perspective of developing a UBP, the dF–dA pair is a mispair). Thus, it was unclear whether hydrophobic and packing forces would be sufficiently strong and specific to underlie the development of a functional UBP.

4. Predominantly Hydrophobic UBPs

Since 1999, our group has pursued the idea of using hydrophobic and packing forces to create a UBP. Our development strategy has been inspired by medicinal chemistry and involves the synthesis and evaluation of a wide variety of analogues as well as the elucidation of structure–activity relationship (SAR) data to guide optimization. During early development efforts, the most useful SAR data was generated from steady-state kinetics, which was used to determine the efficiency (i.e. second-order rate constant) and fidelity (i.e. the ratio of second-order rate constants for correct and incorrect pairings) of UBP synthesis (i.e. incorporation of the unnatural triphosphate opposite its cognate unnatural nucleotide in a template), as well as the subsequent extension step (i.e. incorporation of the next correct nucleotide). However, as the UBPs were optimized, their replication became too

efficient to be characterized by steady-state methods (because of rate-limiting product dissociation, a step that is not relevant to processive synthesis), and thus we turned to pre-steady-state kinetics and the efficiency and fidelity with which DNA containing the UBP is PCR amplified. During later stages of development, these studies employed multiple polymerases and sequence contexts to ensure the generality of the SAR data.

Our first generation analogues, ultimately including 37 different synthetic nucleotides, were designed to explore the ability of the hydrophobic effect and packing interactions to replace interstrand hydrogen bonding. The nucleobase analogues were based largely on the relatively large isocarboxtyrilyl, naphthyl, and indole scaffolds (we note that in many cases, the “nucleobases” are not actually basic, but the designation is still used for simplicity). The details of these early efforts have already been reviewed,^[28,29] but, in general, a variety of pairs were identified that were capable of relatively stable and selective pairing in duplex DNA.^[116] More importantly, several were identified that were synthesized with rates approaching those of a natural pair,^[117] or with at least modest selectivity, thereby suggesting that hydrophobic and packing forces could be harnessed for this step of replication.^[117,118] However, once synthesized, all of the first-generation UBPs were extended with very low efficiency. In fact, this pattern of large nucleobases and efficient synthesis but poor extension emerged as the most pronounced SAR from the first generation analogues.

Structural studies of duplex DNA suggested that these first generation UBPs do not pair in an edge-to-edge manner, but instead pair through cross-strand intercalation of the large aromatic nucleobases (see Section 5). We speculated that this mode of pairing might favor UBP synthesis, as a result of hydrophobic and dispersion forces, but result in a structure refractory to continued primer extension, because of the associated mispositioning of the primer terminus. Thus, our second generation analogues, ultimately consisting of 55 synthetic nucleotides, were mostly based on benzene,^[119–121] pyridine,^[122,123] or pyridone^[124] scaffolds to test whether smaller nucleobases that should be less prone to intercalate could be optimized for replication. Perhaps surprisingly, several of these second generation UBPs formed stably and selectively in duplex DNA, despite the absence of hydrogen bonding and extended aromatic surface area.^[121]

Among these second generation UBPs, several were identified that are synthesized and/or extended with at least reasonable efficiency.^[119,125–127] However, optimization efforts yielded only flat SAR and, problematically, the SAR identified an apparent conflict among the required physicochemical properties. The conflict revolved around the nature of the substituent positioned *ortho* to the glycosidic linkage, which is expected to be disposed into the developing minor groove during replication. Efficient UBP synthesis required this substituent to be hydrophobic in both the incoming and the templating nucleobase. However, efficient extension appeared to have contradictory requirements, as it required the substituent in the template to be hydrophobic and the substituent at the primer terminus to be able to accept a hydrogen bond.^[122,124,125] Indeed, hydrogen-bond acceptors

are positioned at the analogous positions of each natural nucleobase, and at the primer terminus they are known to accept hydrogen bonds from the polymerase.^[128] In addition, biochemical studies have shown that these interactions are critical for primer elongation.^[34,128–130]

The different physicochemical properties required for UBP synthesis and extension challenged the notion that a solution could be designed rationally. Thus, we designed and implemented two different screens to evaluate all possible pairs formed between 60 diverse nucleotides culled from our first and second generation efforts (resulting in ca. 1800 possible UBPs).^[131] Remarkably, the top hit identified by both screens was the UBP formed between **dMMO2** and **dSICS** (Figure 4). A particularly interesting aspect of the **dMMO2**–

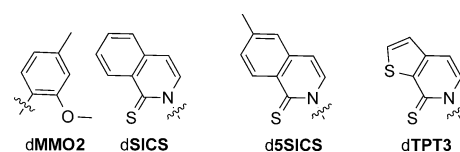


Figure 4. **dMMO2**–**dSICS** UBP and unnatural nucleotides **d5SICS** and **dTPT3**. Sugar and phosphate groups are omitted for clarity.

dSICS pair is the nature of the substituents at the position *ortho* to the glycosidic linkage, which appears to resolve the above-mentioned conflict. The sulfur atom of **dSICS** is softer and more polarizable than an oxygen atom, thus making it more hydrophobic, but it is still capable of accepting a hydrogen bond. A simple rotation of the *O*-methyl group of **dMMO2** allows this *ortho* substituent to present either a hydrophobic methyl group or an oxygen hydrogen-bond acceptor.

Thus, the **dMMO2**–**dSICS** UBP became the founding member of a third generation of candidate UBPs, reinvigorating development efforts and rapidly resulting in the identification of **dMMO2**–**d5SICS** (Figure 4). With **dMMO2**–**d5SICS**, the problematic self-pairing of **dSICS**–**dSICS** was eliminated,^[131] and in contrast to our first and second generation UBPs, this UBP is recognized with high fidelity by a variety of A, B, and X family DNA polymerases.^[132] The least efficient step in the replication of DNA containing **dMMO2**–**d5SICS** is the incorporation of **dMMO2TP** opposite **d5SICS**, and optimization of this step eventually yielded **dNaM**–**d5SICS** (Figure 1),^[133,134] which is synthesized an order of magnitude more efficiently.

Both **dMMO2**–**d5SICS** and **dNaM**–**d5SICS** represented landmarks in our development program, as they were the first of our UBPs that when incorporated into DNA are efficiently amplified by PCR.^[135] Indeed, massive amplification of DNA containing **dNaM**–**d5SICS** is possible with no significant loss of the UBP (i.e. 10^{24} -fold amplification with > 99.9% fidelity), and importantly, with no significant sequence bias.^[136] This efficient, high fidelity, and sequence-independent replication makes **dNaM**–**d5SICS** especially attractive for applications that require massive amplification of a randomized template, such as SELEX (see Section 6). Importantly, when using T7 RNA polymerase, both **dMMO2**–**d5SICS** and **dNaM**–**d5SICS** are efficiently transcribed into RNA (i.e. **dMMO2** or **dNaM**

directs the incorporation of **5SICS** into RNA, and **d5SICS** directs the incorporation of either **MMO2** or especially **NaM** into RNA) with efficiencies only about 30-fold reduced relative to natural nucleotides and with a fidelity of 93–99 %.^[137]

Despite the high efficiency and high-fidelity replication and transcription of DNA containing **dNaM-d5SICS**, the overall rate of its replication is still slower than that of natural base pairs. After efforts to further optimize **dNaM** resulted in only flat SAR,^[134,138–140] our efforts turned to the optimization of **d5SICS**, since continued primer extension after the incorporation of this unnatural nucleotide limits replication. These efforts eventually yielded **dTPT3** (Figure 4).^[141] We then supplemented this optimization effort with another screen of 111 unnatural nucleotides (many of them newly synthesized since our original screening efforts) that formed about 6000 candidate UBPs.^[142] This screen yielded an entire family of well-replicated UBPs (Figure 5). Although **dNaM-**

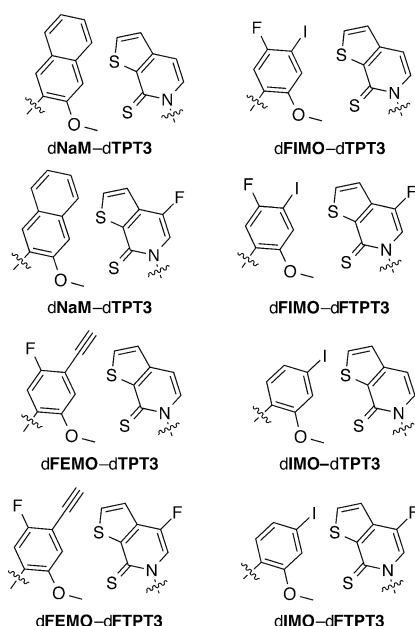


Figure 5. **dNaM-dTPT3**-like family of UBPs.

dTPT3 remains the most efficiently replicated UBP identified to date, each member of this family of UBPs is replicated more efficiently than **dNaM-d5SICS**, which, as described in Section 7, is sufficiently well-replicated for in vivo use. This suggests that all of these UBPs should be suitable for practical applications.

Since 2003, the Hirao group has also focused on the development of predominantly hydrophobic UBPs. Their approach was inspired by Kool's work with **dQ** and **dF**, which are hydrophobic isosteres of **dA** and **dT** (Figure 6).^[143,144] To reduce mispairing with **dA**, the Hirao group first developed **dPa** as a partner for **dQ**,^[145] and then further decorated **dQ** with a bulky thienyl moiety, thereby resulting in **dDs** (Figure 6). The resulting **dDs-dPa** UBP was the first that could be PCR amplified with an error rate below 1% (corresponding to >99% fidelity),^[146] although this required

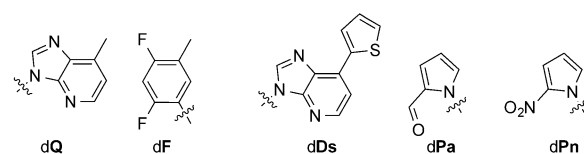


Figure 6. UBP **dQ-dF** and unnatural nucleotides **dDs**, **dPa**, and **dPn**. Sugar and phosphate groups are omitted for clarity.

the use of the γ -amidotriphosphates of **dDs** and **dA** (**dDsTP_N** and **dATP_N**, respectively) to reduce formation of the **dDs-dDs** and **dA-dPa** mispairs. A high fidelity of transcription (>95%) of the biotinylated analogue of **Pa** was also demonstrated.

Further development of the **dDs-dPa** pair resulted from a substitution of the aldehyde group of **dPa** with a nitro group, which yielded the **dPn** nucleoside (Figure 6).^[147] The **dDs-dPn** UBP exhibited improved fidelity during PCR and bypassed the need to use **dATP_N**. Finally, the addition of a propynyl moiety to **dPn** resulted in the **dDs-dPx** UBP (Figure 1, with **R**=H), which further improved the fidelity (>99.9%) and eliminated the need to use either **dDsTP_N** or **dATP_N**.^[148]

The efficient and selective replication of **d5SICS-dNaM**, its related analogues, and **dDs-dPx** clearly indicate that forces other than hydrogen bonding can also be harnessed to mediate stability and high-fidelity replication through the optimization of predominantly hydrophobic nucleobases. Although these studies largely proceeded empirically, several general trends are apparent. Most generally, the data are consistent with the primacy of the hydrophobic effect and local electrostatic and dispersive interactions between the nucleobases (natural and unnatural). These effects are subject to the steric constraints imposed by the polymerase active site. The specific positions of heteroatoms within the nucleobase scaffolds can have substantial positive or negative effects, largely as a result of electrostatic interactions and/or desolvation. The effects of modifications were typically greater with the triphosphates than with the templates. As already discussed, efficient UBP synthesis required the substituent *ortho* to the glycosidic linkage to mediate hydrophobic packing interactions, while UBP extension required it to be capable of hydrogen bonding, and efficient replication required substituents capable of both. At the *meta* and *para* positions, oriented toward the developing major groove, increased aromatic surface area and other substituents that appear to mediate favorable local electrostatic and dispersive interactions were found to optimize UBP synthesis but to disfavor extension, and efficient replication required an intermediate level of these forces. Indeed, many modifications were found to have opposite effects on UBP synthesis and extension, and optimization required compromises.

5. Structural Studies

The structures of duplex DNA containing nucleotides with synthetic nucleobases have provided invaluable information regarding the forces that stabilize UBP formation and which might contribute to their replication. The first structure of a predominantly hydrophobic UBP in duplex DNA was

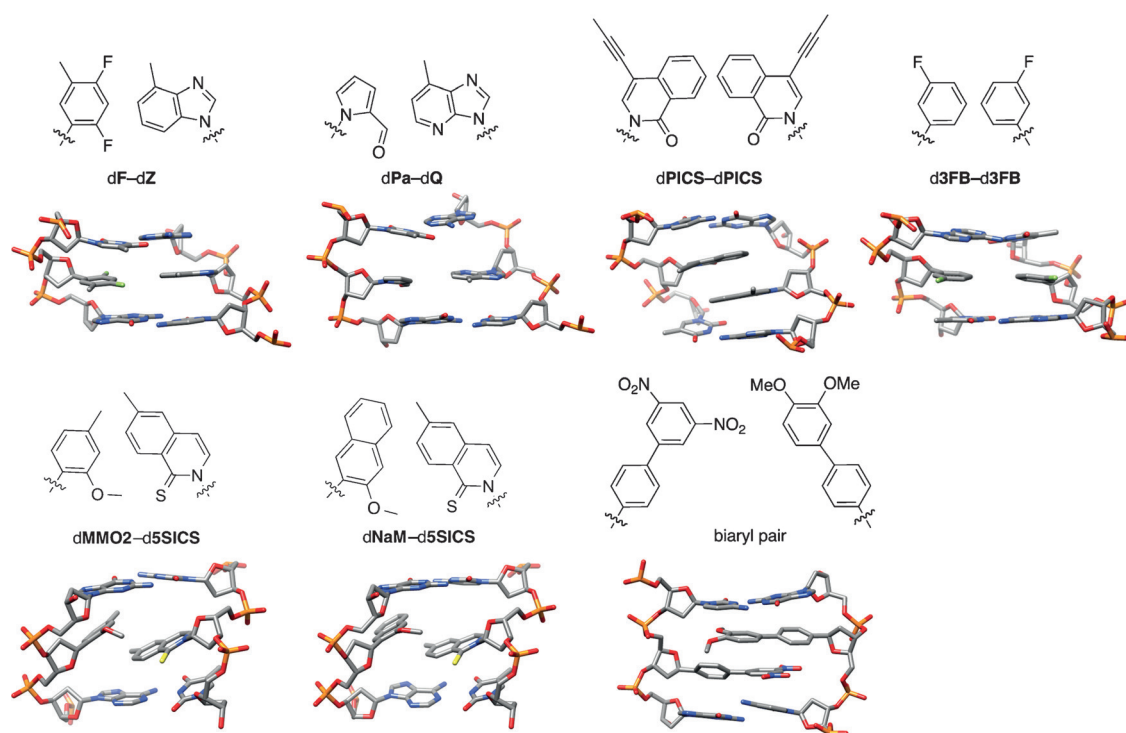


Figure 7. Structures of UBPs. Sugar and phosphate groups are omitted for clarity.

that of **dF** paired with a hydrophobic isostere of **dA** called **dZ** (not to be confused with Benner's analogue of the same name; Figure 7).^[149] The duplex adopted a regular B-form structure with the nucleobases packed within the duplex interior and paired in an edge-to-edge, planar manner. The C1'-C1' distance was found to be slightly increased (by 0.8 Å) compared to a natural base pair, presumably to accommodate the additional hydrogen atom at the interface between the nucleobase analogues. The structure of the **dQ-dPa** pair, which was also designed to structurally mimic a natural purine-pyrimidine pair, is also well-accommodated within a B-form duplex with a structure similar to that of a natural base pair (Figure 7).^[145] The structure of DNA containing the pyrene nucleobase paired against the abasic site revealed that even this large nucleobase analogue did not result in major structural perturbations of the duplex.^[150] Several other structures of DNA^[25,151–154] and RNA^[155] containing at least partially hydrophobic nucleobase analogues have been reported with similar results, thus demonstrating that the DNA duplex is able to accommodate these nucleobases, with more or less natural-like geometries.

The structures of duplexes containing **dPICS-dPICS** (one of our first generation UBPs, specifically a “self-pair” formed between two identical **dPICS** analogues, which was efficiently synthesized, but not extended; see Section 4),^[154] **d3FB-d3FB** (a second generation UBP),^[119] **dMMO2-d5SICS**,^[138] and **dNaM-d5SICS**^[156] have all been reported (Figure 7). Unlike previously reported structures, the NMR structure of DNA containing the **dPICS** self-pair showed that the nucleobases did not pair in an edge-to-edge, planar manner, but instead paired through cross-strand intercalation. Although the X-ray crystal structure of the **d3FB-d3FB** second generation UBP

revealed a planar, edge-to-edge mode of pairing, the NMR structures of **dMMO2-d5SICS** and **dNaM-d5SICS** again revealed intercalative modes of pairing. The *ortho* sulfur atoms and methoxy groups are oriented into the minor groove of the duplex, with the methoxy groups rotated out of planarity with respect to the aromatic ring, thereby allowing for the establishment of van der Waals contacts with the sulfur atom of **d5SICS**. The stacking interface between the nucleobases is formed by the methyl group and proximal portion of the aromatic ring of **d5SICS** and the *para* substituent of **dMMO2** or **dNaM**. However, relative to the **dPICS** self-pair, there is somewhat less overlap between the nucleobases, due to their geometry and/or reduced size. Based on molecular modeling studies, ¹H NMR analysis, and circular dichroism studies, Leumann and co-workers had previously proposed such an intercalative mode of pairing between biaryl-derivatized nucleotides,^[13,14,157] which was later confirmed by NMR structural studies (Figure 7).^[158] Finally, duplexes containing nucleotides bearing multiple aromatic chromophores instead of nucleobases associate in a zipper-like motif, wherein each individual chromophore intercalates into the complementary strand in a similar fashion (for reviews, see Refs. [159,160]). Thus, this intercalative mode of pairing appears to be general with large aromatic nucleobase analogues, and is consistent with an important role of hydrophobicity and dispersion forces.

Although the intercalative mode of pairing in duplex DNA clearly contributes to UBP stability, it results in a structure that is more similar to a mismatch than a correct natural pair. It was thus difficult to reconcile the efficient and high-fidelity replication of these UBPs with the accepted mechanism of polymerase recognition, in which binding of the

correct (natural) triphosphate induces a large conformational change involving closure of the fingers domain over the palm and thumb domains to produce a closed and rigid complex that geometrically selects for the conserved Watson–Crick structure of a natural base pair.^[50,51,161–164] To address this apparent contradiction and elucidate the mechanism of UBP synthesis, in collaboration with the Marx group, we used X-ray crystallography to characterize the structure of several KlenTaq complexes (KlenTaq is the large fragment of *Taq* DNA polymerase I; Figure 8).^[156] Comparison of the binary and ternary complexes revealed that with dNaM in the templating position, the addition of d5SICSTP induces the same large conformational changes in the polymerase as

induced by the addition of a cognate, natural triphosphate^[163,165,166] (Figure 8). In fact, comparison of the ternary complexes with natural and unnatural substrates reveals remarkable similarities, including the interactions between the triphosphate and polymerase, the orientations of the active-site side chains, and even the bound metal ions. Most remarkably, unlike the intercalated structure formed in a free duplex, the nucleobases of the developing UBP in the polymerase active site of the closed complex adopt a coplanar structure with nearly optimal edge-to-edge packing and a C1'–C1' internucleotide distance that is roughly the same as that of a natural base pair (11.0 versus 10.6 Å, respectively, compared to 9.1 Å for dNaM–d5SICS in a free duplex). In contrast, the addition of dNaMTP to a complex of KlenTaq bound to template containing d5SICS does not fully induce the formation of the catalytically competent closed state.^[167] However, it does induce the formation of a partially closed state, in which d5SICS moves toward the insertion site and dNaMTP is bound to the O-helix through its triphosphate moiety. This state is similar to the ajar state, which is thought to allow the polymerase to test for correct base pairing during the synthesis of natural DNA.^[164,168,169]

The structures of several post-incorporation complexes were also characterized, wherein the UBP strand context and flanking sequences were varied.^[167] Although structural differences were observed, in each case the UBP was positioned in the correct post-insertion site, where it again assumed an intercalated structure, as it does in the free duplex.

These structural studies, along with the analysis of UBP replication in different sequence contexts,^[133] allowed us to propose a mechanism for UBP replication (Figure 9). Upon binding of an unnatural triphosphate to the O-helix, the polymerase samples different conformations, and with sufficient UBP stability, populates the catalytically active closed conformation. With d5SICSTP incorporation, the intermediate states are populated only transiently, but with dNaMTP incorporation, the series of conformational changes are halted at the ajar-like state, either as a consequence of the

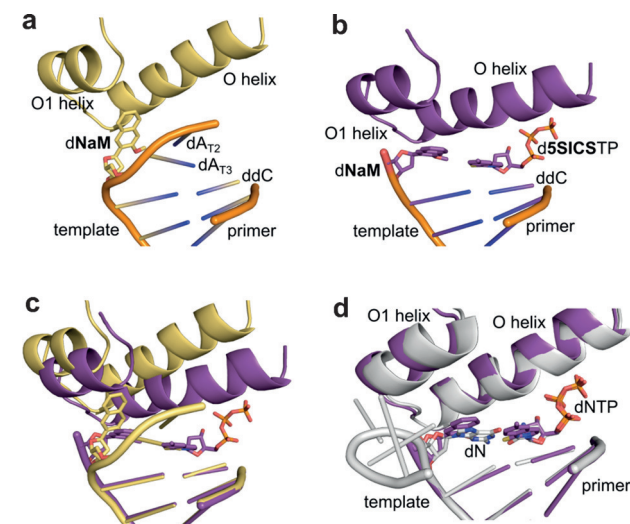


Figure 8. KlenTaq polymerase induces the UBP dNaM–d5SICS to adopt a natural, Watson–Crick-like structure. a) Structure of the polymerase–template binary complex; b) structure of the polymerase–template–d5SICSTP ternary complex; c) a superposition of binary and ternary complexes; and d) a superposition of the ternary complex where d5SICSTP is paired with dNaM and the ternary complex, where dGTP is paired with dC. Reproduced from Ref. [156].

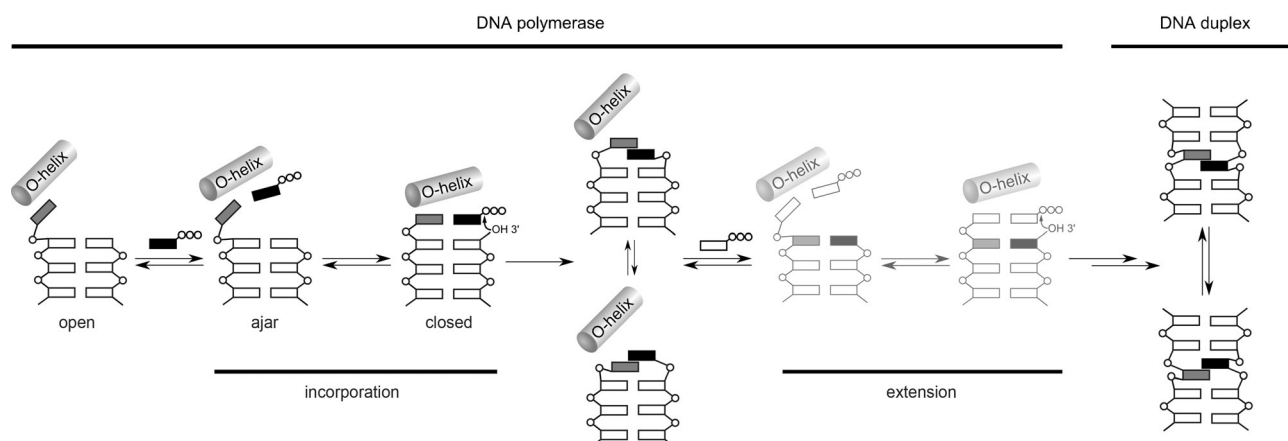


Figure 9. Proposed mechanism of replication of UBP d5SICS–dNaM. Intermediates based on solved structures are shown in a darker color and those not yet validated by structural studies are shown in a lighter color. The steps corresponding to incorporation of the unnatural triphosphate and subsequent extension of the nascent UBP are indicated. Phosphates are indicated with open circles, natural nucleotides are indicated with open rectangles, and the unnatural nucleotides are indicated with gray and black boxes. Reproduced from Ref. [167].

stability of this complex or the instability of the corresponding closed complex, and further progress towards the incorporation of **dNaMTP** requires thermal fluctuations to populate the closed state. In either case, with the population of the closed complex, the developing UBP adopts a Watson–Crick-like structure that facilitates covalent attachment of the incoming unnatural nucleotide. After incorporation, the polymerase returns to the open conformation, a molecule of pyrophosphate is released, and the UBP translocates to the post-insertion site, where it intercalates in the complementary strand. UBP extension then requires thermal fluctuations to both de-intercalate the UBP and reorganize the polymerase active site. This intercalation/de-intercalation mechanism explains the balance of electrostatic and dispersive interactions that was required for the simultaneous optimization of UBP synthesis and extension. In fact, more than just being consistent with their mediating replication, this mechanism suggests that hydrophobic and packing forces may be ideal due to their relatively strong but nondirectional, and thus plastic, nature.

6. In Vitro Applications

With the development of replicable UBPs, interesting applications soon followed. Many early applications took advantage of the inherent increase in hybridization specificity associated with three base pairs, relative to two (reviewed by Kimoto et al., 2011^[170]). A more recent and potentially revolutionary application is the use of UBPs in SELEX for the evolution of aptamers or nucleic acid catalysts with unnatural functionality. This important milestone was first reached by the Hirao group, who used a derivative of the **dDs-dPx** UBP (Figure 1, with $R = \text{CH(OH)CH}_2\text{OH}$) to evolve aptamers that use the unique functionality of **dDs** to recognize VEGF-165 or IFN- γ .^[171] Since no straightforward sequencing method exists for determining the position of multiple UBPs, one or multiple **dDs** analogues were incorporated into defined positions of a 43-nt randomized region, with their positions identified by a two- or three-nucleotide barcode. After multiple rounds of selection and PCR amplification, all unnatural nucleotides were converted into natural nucleotides by PCR, and the resulting population was analyzed by Ion Torrent deep sequencing. The positions of the **dDs** nucleotides were then reconstituted from the identity of the barcodes. Both selections with the UBP yielded aptamers that bound their target with approximately 100-fold greater affinity than aptamers evolved from selections with only natural nucleotides. The highest affinity unnatural aptamer for VEGF-165 had two **dDs** nucleotides, neither of which could be replaced with dA without a decrease in affinity, and the highest affinity aptamer for IFN- γ had three, two of which could not be replaced with dA.

Soon after the study by Hirao and co-workers, the Benner group evolved aptamers containing the unnatural nucleotides **dP** and **dZ** that recognize breast cancer cells with a dissociation constant of 30 nM.^[172] To identify specific binders after 12 rounds of in vitro selection, the authors sequenced the resulting unnatural library by controllable mutation of **dZ** to

dC and dT (developed according to Ref. [39]), followed by standard deep sequencing with an Ion Torrent instrument. Importantly, as in the previous example, aptamer affinity for the target cells was significantly reduced when unnatural nucleotides were mutated to a natural counterpart, thus highlighting the importance of the unnatural functionality.

Although the functionality made available by the inclusion of the **dDs-dPx** or **dP-dZ** UBPs was clearly utilized in the evolved aptamers, a much greater range of physicochemical properties would be accessible by using the UBPs to site-specifically attach other functionality to DNA or RNA. The attachment of such cargo would allow for an almost limitless range of functionality to be subjected to the evolution process. For this to be accomplished, the unnatural nucleotides must be modified with a linker that does not interfere with replication or transcription and that can be used to attach the cargo of interest either to the triphosphate or to the unnatural nucleotide after its incorporation into DNA or RNA.

The first example of such a site-specific labeling of an oligonucleotide was reported by Tor and Dervan, who used a template containing **disoC** to direct the incorporation of 6-aminohexyl**isoG** into RNA, which was followed by post-transcription labeling with *N*-hydroxysuccinimidobiotin (NHS-biotin) or EDTA dianhydride.^[173] More recently, the **dDs-dPa/dPn/dPx** family of UBPs has been used by Hirao and co-workers to site-specifically label DNA^[146,148] and RNA.^[146,174,175] Several of the RNA labeling experiments relied on direct incorporation during transcription of the unnatural ribonucleoside triphosphate covalently attached to the functional group of interest. However, the attachment of larger moieties, such as fluorophores, sometimes interfered with transcription. To overcome this limitation, **dDs** was used to template the transcription of the alkyne-modified ribonucleoside triphosphate Eth-C4-**PaTP**, with the resulting RNA then modified with a biotin tag or a fluorophore through Click chemistry.^[176] The Hirao group has also reported a series of linker-modified **dPx** analogues that are well-replicated,^[177] one of which was used in the above-mentioned selection, and the use of other linkers or linkers with interesting functionality attached will no doubt increase the range of activities that can be sampled.

We synthesized and evaluated UBP replication and transcription with a variety of (d)**5SICSTP**, (d)**MMO2TP**, and (d)**NaMTP** derivatives modified with linkers bearing free or protected amines for coupling to cargo through NHS esters,^[178] or alkynyl groups for coupling to cargo through Click chemistry.^[179] With the **d5SICS** scaffold, the addition of a propynyl ether linker at the free *meta* position is accommodated better than its aliphatic analogue, but the protected propargylamine linker is best tolerated. Interestingly, the **dTPT3** scaffold is more tolerant of linker derivatization than the **d5SICS** scaffold.^[141] Although we have not yet identified linker modifications that are well-tolerated by the **dNaM** scaffold, the *para* and *meta* positions of **dMMO2** are tolerant of linker attachment, and while aliphatic and ether-based linkers are accommodated, the direct attachment of an ethynyl group to the nucleobase core is best tolerated. In general, these results may be rationalized on the basis of the

ternary structure of KlenTaq polymerase bound to 5'-propargylamido dUTP.^[180] The structure revealed a hydrogen bond between Arg660 and the nitrogen atom of the propargylamido linker in the developing major groove. In the absence of the linker, Arg660 interacts with the phosphate backbone of the primer terminus,^[181,182] and thus the linkers must be either small enough not to perturb this interaction, or capable of replacing it with a stabilizing interaction.

To demonstrate the application of the linker-modified UBPs, DNA containing dNaM or d5SICS was used to direct the transcription of *Methanococcus jannaschii* tRNA^{Tyr} with the amino-linker-modified variants of 5SICS or MMO2, respectively, at the third position of the anticodon, which in the latter case was used to site-specifically biotinylate the tRNA.^[178] Towards material-like applications, DNA containing the UBP formed between dNaM and propynyl ether derivatized d5SICS was amplified and used to site-specifically array two nSH3 domains (Src homology 3 domain from the human CrkII adaptor protein), with the expected topology confirmed by atomic force microscopy.^[179] We also synthesized the α -phosphorothioate variant of d5SICSTP and demonstrated its use in thiolation of the DNA backbone and postamplification labeling.^[178] Together, these modified nucleotides allow for the site-specific modification of nucleic acids with up to three different functional groups with unprecedented spatial control.

7. UBPs and Semi-synthetic Organisms

Perhaps the ultimate goal of developing UBPs is for their use in vivo as the foundation of semi-synthetic organisms that store and retrieve increased genetic information (Figure 10). With the demonstration that dNaM-d5SICS is replicated with high efficiency and fidelity in any sequence context, and that it is also efficiently transcribed into RNA, we initiated efforts towards this goal. With the wealth of knowledge and tools available, *E. coli* was the obvious candidate for the creation of

the first semi-synthetic organism. However, the two membranes possessed by this organism complicated the first challenge of how to make the requisite unnatural triphosphates available within the cell. We first explored a strategy employed by many nucleoside drugs, namely, the passive diffusion of the free nucleoside into the cell followed by its triphosphorylation through the sequential action of the kinases of the nucleoside salvage pathway.^[183] Although we found that a variety of nucleosides with predominantly hydrophobic nucleobases were monophosphorylated by the nucleoside kinase from *D. melanogaster*, we were unable to identify monophosphate kinases that catalyzed conversion into the diphosphates, which are more specific for their natural substrates.^[184] In addition, overexpression of kinases, in an attempt to compensate for their low activity, resulted in significant levels of toxicity, presumably due to misregulation of the natural triphosphate pool.

Challenged with having to identify a novel route to make the unnatural triphosphates available intracellularly, we noted that the genomes of a variety of intracellular bacteria and algal plastids do not encode the enzymes required for nucleoside triphosphate synthesis. Instead, they encode dedicated nucleoside triphosphate transporters (NTTs) and scavenge the requisite triphosphates from their cellular milieu.^[185–192] We examined the plasmid-based expression of eight different NTTs in *E. coli*, and found that one from *Phaeodactylum tricornutum* (PtNTT2)^[191] was active and able to import both d5SICSTP and dNaMTP (Figure 10). The addition of 250 μ M d5SICSTP and dNaMTP to the media resulted in steady-state intracellular levels of approximately 90 μ M and 30 μ M, respectively. These concentrations are significantly above the sub-micromolar K_M values of the unnatural triphosphates for DNA polymerases,^[134] thus setting the stage for replication of the UBP in a living organism.

To determine whether *E. coli* equipped with PtNTT2 could use the imported unnatural triphosphates to replicate DNA containing the UBP, we used solid-phase synthesis and circular extension PCR to replace a single dA-dT in the pUC19 plasmid with the dNaM-dTPT3 UBP, thereby resulting in pINF (information plasmid). *E. coli* cells expressing a plasmid-borne PtNTT2 were grown in media supplemented with dNaMTP and d5SICSTP and transformed with pINF (Figure 10). As controls, cells were also transformed with the parent pUC19 plasmid, or grown either without induction of the transporter or without the unnatural triphosphates. After overnight growth, pINF, which had been amplified approximately 10^7 -fold, was recovered, digested, and dephosphorylated to free nucleosides, and analysis by LC-MS/MS demonstrated the presence of approximately one d5SICS per plasmid. In contrast, no d5SICS was detected when cells were transformed with the fully natural pUC19 plasmid, when they were transformed with pINF but the transporter was not induced, or when the unnatural triphosphates were not provided. Importantly, since the synthetic pINF contained dNaM-dTPT3, and d5SICS was only provided as triphosphate to the media, the detection of d5SICS in the recovered plasmid unambiguously demonstrated in vivo replication. This conclusion was confirmed by sequencing and by using a biotin shift assay after amplification of the recovered pINF

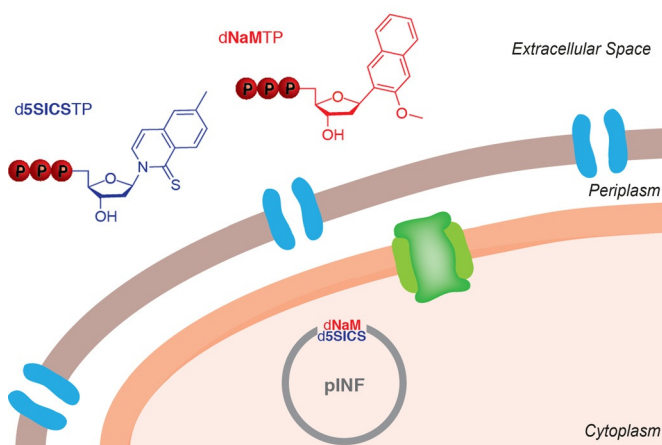


Figure 10. Recently developed semi-synthetic organism. Porins in the outer membrane and PtNTT2 in the inner membrane are illustrated in blue and green, respectively.

with **d5SICS** and a biotinylated analogue of **dNaM**. Based on the lower limit of detection provided by the analysis of UBP retention and the amplification level, the fidelity of replication is at least 99.4%. Thus, this modified *E. coli* represents the first semi-synthetic organism that propagates a UBP in its DNA.

8. Implications and Future Directions

It has been more than 50 years since Alex Rich first proposed the development of UBPs, and 25 years since Steve Benner's group produced the first viable candidates. Today, three classes of UBPs have been developed and validated to a high level of proof-of-concept: one based on alternative hydrogen bonding, exemplified by **dZ-dP**, and two based on hydrophobic and packing forces, exemplified by **dNaM-d5SICS** and **dDs-dPx**. The data demonstrate that at least for a single UBP embedded within an otherwise natural environment, hydrogen bonds are not unique in their ability to mediate stable and selective pairing, replication, and transcription into RNA, and that they may be replaced by the hydrophobic effect and dispersive interactions mediated by nucleobase scaffolds with complementary structures, combined with desolvation and electrostatic interactions mediated by judiciously positioned heteroatoms. Continued optimization of all three classes is expected to yield UBPs that are replicated with rates and fidelities that are indistinguishable from those of a natural base pair.

Future efforts should also focus on developing methods to sequence DNA containing the UBPs, especially DNA containing multiple UBPs. This could include modification of the nucleobase with fluorophores for Sanger sequencing, modification of the nucleobase and sugar with fluorophores and reversible terminators, respectively, for sequencing with the Illumina platform,^[193] or modification of the triphosphate moiety with a fluorophore for sequencing by the Pacific Biosciences method.^[194] Sequencing with the Ion Torrent^[195] or Oxford Nanopore^[196] platforms would also be attractive, as they would not require any nucleotide modification. In any case, modifications to the hardware and software will be required. Efforts should also include an increased focus on the systematic exploration and optimization of linkers for the site-specific attachment of different cargo to DNA. Such efforts will continue to facilitate the application of the UBPs in different technologies that draw upon the potential physicochemical properties of the cargo, and importantly, in an evolvable context. The use of UBPs for the evolution of novel devices or materials represents perhaps their most significant in vitro application.

It is our opinion that the most exciting application of UBPs is their use for the creation of semi-synthetic organisms that store and retrieve increased information. Again drawing upon well-established tenets of medicinal chemistry, the availability of the **dNaM-dTPT3** family of UBPs is especially important, as the different nucleotides are expected to possess a range of different pharmacokinetic-like properties, such as triphosphate uptake, stability, and off-target activity. Correspondingly, SAR studies should move toward including these

parameters, in addition to in vitro and in vivo replication efficiency and fidelity.

The next step towards the creation of semi-synthetic organisms capable of retrieving increased information is the in vivo transcription of DNA containing the UBP into RNA. In conjunction with the orthogonal tRNA/aminoacyl synthetase pairs developed by Schultz and co-workers,^[197–199] this will set the stage to explore the retrieval of the information encoded by the UBP in the form of novel proteins. There is no reason such efforts should be limited to bacterial cells, and the engineering of eukaryotic cells to store and retrieve increased information brings with it its own potentially transformative applications. Such semi-synthetic bacterial and eukaryotic organisms will form the foundation of semi-synthetic life, with the potential to possess and evolve new and useful attributes or functions. DNA underlies all that life is, has been, and might evolve to be, and it would appear that UBPs now promise to dramatically increase the potential of this already remarkable molecule.

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