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# ENZYMATIC FORMATION OF AN ARTIFICIAL BASE PAIR USING A MODIFIED PURINE NUCLEOSIDE TRIPHOSPHATE

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## ABSTRACT

The expansion of the genetic alphabet with additional, unnatural base pairs (UBPs) is an important and long standing goal in synthetic biology. Nucleotides acting as ligands for the coordination of metal cations have advanced as promising candidates for such an expansion of the genetic alphabet. However, the inclusion of artificial metal base pairs in nucleic acids mainly relies on solid-phase synthesis approaches and very little is known on polymerase-mediated synthesis. Herein, we report the selective and high yielding enzymatic construction of a silver-mediated base pair (**dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>**) as well as a two-step protocol for the synthesis of DNA duplexes containing such an artificial metal base pair. Guided by DFT calculations, we also shed light into the mechanism of formation of this artificial base pair as well as into the structural and energetic preferences. The enzymatic synthesis of the **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** artificial metal base pair provides valuable insights for the design of future, more potent systems aiming at expanding the genetic alphabet.

## INTRODUCTION

The expansion of the genetic alphabet with additional, artificial base pairs is a long standing goal in synthetic biology.<sup>1-2</sup> The design of these artificial base pairs mainly relies on complementary hydrogen bonding,<sup>3-4</sup> hydrophobic interactions<sup>5</sup> or geometric complementarity.<sup>6-7</sup> Expanded genetic systems have served as foundation of semi synthetic organisms able to store and retrieve increased information in their DNA which in turn could potentially permit the creation of proteins with new properties and structures.<sup>8-10</sup> This increased chemical diversity has also been used to develop modified aptamers with improved affinity and selectivity.<sup>11-15</sup> Metal ion coordination has lately emerged as an alternative strategy for the creation of UBPs.<sup>16-28</sup> Such artificial metal base pairs are indeed orthogonal to the natural Watson-Crick base pairs, are water soluble, possess a high thermal stability and do not cause any perturbation to the structure of canonical duplexes.<sup>29-30</sup> Moreover, unlike some of the cyclic  $\pi$ -conjugated analogues which can be photoactivated by near-visible light, metal base pairs do not necessarily absorb ultraviolet light and thus do not produce any reactive oxygen species which are highly toxic for DNA and cells.<sup>31-32</sup> The presence of a metal also offers the possibility of constructing novel devices such as metal nanowires or DNA-based logic gates.<sup>33-35</sup> Yet, most metal base pairs are incorporated into synthetic oligonucleotides and surprisingly, very little is known about their enzymatic formation.<sup>18-19, 36-38</sup>

In this context, we have recently investigated the enzymatic construction of two imidazole containing nucleotides (**1** and **2** in Fig. 1).

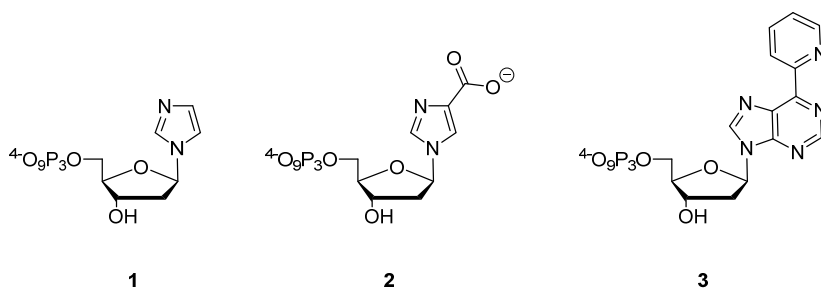


Figure 1. Chemical structures of the modified **dImTP 1**, **dIm<sup>C</sup>TP 2** and **dPur<sup>P</sup>TP 3** nucleotides.

The biochemical analysis carried out with nucleotide **dImTP 1** revealed that, in addition to the requirements of a strong thermal stability of the metal UBP and a low structural perturbation, multiple parameters affect the enzymatic construction of artificial metal base pairs. Additional effects that need to be taken into consideration for the design of novel candidates include size-complementarity,  $\pi$ -stacking, hydrophobic and minor groove interactions.<sup>39</sup> In view of these findings, we designed nucleotide **dIm<sup>C</sup>TP 2**, which has an

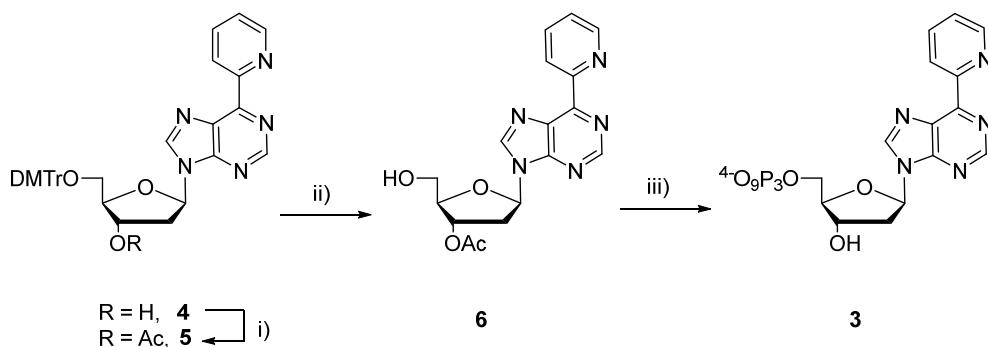
additional carboxy group on position 4 of the imidazole moiety. This was meant to increase the coordination environment and to reduce hydrogen-bonding to the canonical nucleotides.<sup>26</sup> This second imidazole generation nucleotide was shown to be able to form an artificial metal base pair with Ag<sup>I</sup> in a [2+1] coordination environment under enzymatic synthesis conditions. However, no multiple incorporations could be observed and the efficiency of the formation of **2**-M<sup>n+</sup>-**2** base pairs remained modest, which prevents the use of **2** for the construction of libraries to be used in selection experiments.<sup>40</sup> Here, we evaluate the effect on the efficiency of the enzymatic construction of artificial metal base pairs of increased stacking and minor groove interactions. To do so, we focused on the 6-pyridylpurine nucleoside **dPur<sup>P</sup>** (**3** in Fig. 1), which has been shown to form extremely stable metal base pairs with Ni<sup>II</sup> cations<sup>41</sup> and which presents a larger aromatic ring system and sustains interactions with polymerases through the minor groove acceptor nitrogen at N3.<sup>42</sup>

Herein, we demonstrate the efficient enzymatic synthesis of a **dIm<sup>C</sup>**-Ag<sup>I</sup>-**dPur<sup>P</sup>** artificial metal base pair. We also have developed a two-enzyme strategy that permits bypassing of the artificial metal base pair and allows enzymatic synthesis of DNA containing this metal UBP. Surprisingly, the expected formation of a **dPur<sup>P</sup>**-Ni<sup>II</sup>-**dPur<sup>P</sup>** base pair was not observed and only a **dPur<sup>P</sup>**-Ag<sup>I</sup>-**dPur<sup>P</sup>** base pair could be constructed albeit in moderate yields. Quantum mechanics calculations showed that such metal base pairs could form and gave an insight into their mechanism of formation in the context of enzymatic synthesis.

## RESULTS AND DISCUSSION

### Synthesis of the modified triphosphate and modified oligonucleotides

In order to evaluate the possibility of using the 6-pyridylpurine nucleoside **dPur<sup>P</sup>** to form artificial metal base pairs using polymerases we needed to synthesize the corresponding triphosphate and phosphoramidite building blocks. To do so, we synthesized the modified **dPur<sup>P</sup>TP** triphosphate by application of the Ludwig-Eckstein triphosphorylation conditions<sup>43-44</sup> on the suitably 3'-*O*-Ac protected nucleoside **6** which was obtained by a sequential acetylation-debenzylation protocol starting with the known nucleoside **4** (**Scheme 1**).<sup>41</sup>



Scheme 1. Synthesis of **dPur<sup>P</sup>TP 3**. Reagents and conditions: (i) Ac<sub>2</sub>O, DMAP, NEt<sub>3</sub>, pyridine, 0°C, 1 h; (ii) TCA 3% in CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, rt, 30 min, 70%; (iii) 1. 2-chloro-1,3,2-benzodioxaphosphorin-4-one, pyridine, dioxane, rt, 45 min; 2. (nBu<sub>3</sub>NH)<sub>2</sub>H<sub>2</sub>P<sub>2</sub>O<sub>7</sub>, DMF, nBu<sub>3</sub>N, rt, 45 min; 3. I<sub>2</sub>, pyridine, H<sub>2</sub>O, rt, 30 min; 4. NH<sub>3</sub> (aq.), rt, 2 h, 8% (4 steps).

The corresponding phosphoramidite building block for solid-phase synthesis of **dPur<sup>P</sup>**-containing DNA oligonucleotides was obtained by application of known literature procedures.<sup>45</sup> All the templates containing modified nucleotides were obtained using automated solid-phase synthesis and are summarized in Table 1. Template **T1**, containing a single **dPur<sup>P</sup>** nucleotide located immediately after the 3'-end of the FAM-labelled primer **P1** was synthesized to evaluate the possibility of forming the artificial metal base pairs **dPur<sup>P</sup>-M<sup>n+</sup>-dPur<sup>P</sup>** or **dIm<sup>C</sup>-M<sup>n+</sup>-dPur<sup>P</sup>** by enzymatic synthesis. In addition, the template **T2** containing a **dIm<sup>C</sup>** nucleotide previously synthesized in our laboratory<sup>40</sup> was used to study the formation of a **dPur<sup>P</sup>-M<sup>n+</sup>-dIm<sup>C</sup>** base pair.

Table 1. Sequences of the DNA primer and templates used for the primer extension reactions.

<b>P1</b>	5'-FAM-TAC GAC TCA CTA TAG CCT C
<b>T1</b>	5'-GGA GPur <sup>P</sup> G AGG CTA TAG TGA GTC GTA
<b>T2</b>	5'-GGA GIm <sup>C</sup> G AGG CTA TAG TGA GTC GTA
<b>T3-T6<sup>a</sup></b>	5'-GGA GNG AGG CTA TAG TGA GTC GTA

<sup>a</sup>N = C (**T3**); N = G (**T4**); N = A (**T5**); N = T (**T6**).

#### Initial polymerase screen for the construction of **dPur<sup>P</sup>-M<sup>n+</sup>-dPur<sup>P</sup>** and **dIm<sup>C</sup>-M<sup>n+</sup>-dPur<sup>P</sup>** base pairs

The enzymatic construction of a **dPur<sup>P</sup>-M<sup>n+</sup>-dPur<sup>P</sup>** base pair, which involves the incorporation of **dPur<sup>P</sup>TP** opposite a templating **dPur<sup>P</sup>** in the strict presence of a metal cation, was first evaluated with 5 different polymerases known to accept modified nucleotides<sup>18-19</sup> (i.e. *Bst*, Deep Vent, Taq, Vent (*exo*<sup>-</sup>) and the Klenow

fragment of DNA polymerase I *exo*- (Kf (*exo*<sup>-</sup>)) in the presence of 12 different metal cations (see the Supporting Information). First analysis of the primer extension (PEX) reactions using primer **P1** and template **T1** revealed the complete inefficacy of Deep Vent at recognizing the nucleotide as a substrate since only exonucleolytic degradation of the primer could be observed (data not shown). Similarly, Kf (*exo*<sup>-</sup>) did not accept **dPur<sup>P</sup>TP** as a substrate since no extension of the primer could be observed (Fig. S1). On the other hand, *Bst* and Vent (*exo*<sup>-</sup>) polymerases permitted a full incorporation of the modified nucleotide but without any significant metal dependence (Fig. S1). The reactions catalysed by Taq led to the partial incorporation of the modified nucleotide in the presence of Ag<sup>I</sup>, Cu<sup>II</sup> and Mn<sup>II</sup> but also to the incorporation of **dPur<sup>P</sup>MP** units in the absence of metal cations (Fig. S1), while all other PEX reactions conducted in presence of other metal cations only led to low yields of the n+1 product (Table S1). When we lowered the triphosphate concentration from 200 μM down to 20 μM in order to reduce misincorporation events in the absence of metal cofactors, no n+1 product formation could be detected with Vent (*exo*<sup>-</sup>). However, the reactions performed with the Taq polymerase showed complete formation (91% yield) of the expected n+1 product when Mn<sup>II</sup> was added to the reaction mixture with clear cation dependence since very little incorporation could be observed with Cu<sup>II</sup> (yield of n+1 formation of 5 % at 30 μM, 6% at 60 μM, 5% at 100 μM) Ni<sup>II</sup> or in the control reaction (yield of 6%) (Fig. 2). While a number of metal ions such as Cd<sup>II</sup>, Ni<sup>II</sup>, Cu<sup>II</sup> or Ga<sup>III</sup> support polymerase activity,<sup>46-47</sup> the addition of Mn<sup>II</sup> is known to relax the substrate specificity and the general polymerase fidelity and hence this result cannot be interpreted as the formation of a true **dPur<sup>P</sup>-Mn<sup>II</sup>-dPur<sup>P</sup>** base pair.<sup>48-49</sup>

Reactions carried out with the Kf (*exo*<sup>-</sup>) polymerase and 20 μM of **dPur<sup>P</sup>TP** led to an efficient incorporation of the modified nucleotide when the reaction mixtures were supplied with Ag<sup>I</sup> (71% yield of n+1 product). The presence of silver cations seems to be required for the incorporation of **dPur<sup>P</sup>MP** units since only a small fraction of n+1 products can be detected in the control reaction carried out in the absence of metal cations (conversion yield of 13%). Moreover, formation of the n+1 product appears to be selective with respect to the nature of the metal cation used since lower incorporation yields occurred with Ni<sup>II</sup>, Cd<sup>II</sup> or Fe<sup>II</sup> especially at higher triphosphate concentrations (Fig. 2 and Fig. S2). Surprisingly, we did not observe the formation of n+1 product under any of the experimental conditions that we tested when Ni<sup>II</sup> was included in the reaction mixtures (Fig. S2), even though this metal cation was shown to form very stable metal base pairs with short synthetic oligonucleotides.<sup>41</sup>

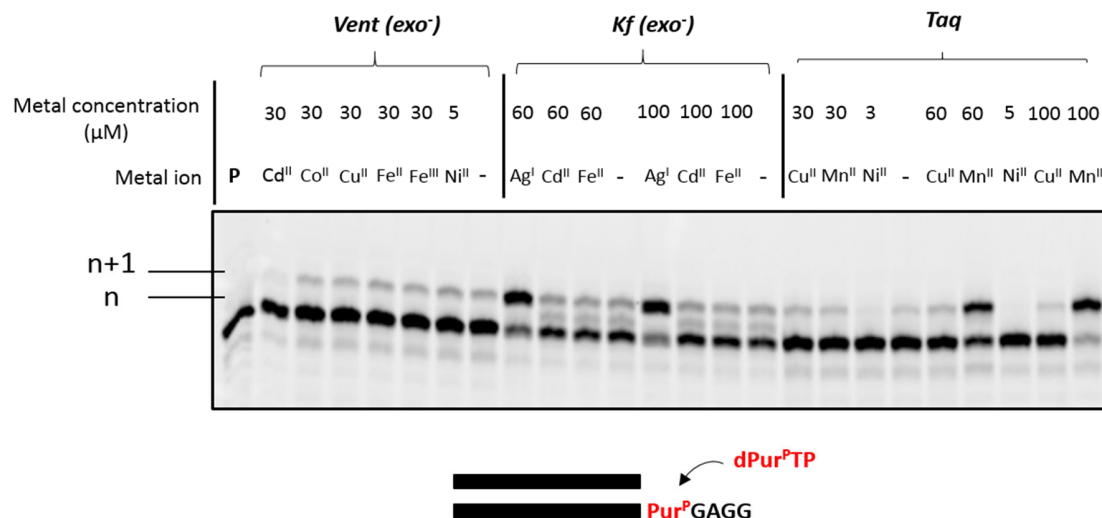


Figure 2. Gel analysis of the incorporation of **dPur<sup>P</sup>TP** (20 μM final concentration) opposite a single templating **dPur<sup>P</sup>** nucleotide in the presence of various metal cations using primer **P1** and template **T1** with *Vent (exo<sup>-</sup>)* (2 U), *Kf (exo<sup>-</sup>)* (5 U) and *Taq* polymerases (5 U). Reaction mixtures were incubated at 60 °C or 37 °C for 1 hour. **P** indicates the unreacted primer. Ag<sup>I</sup> reaction conditions were performed with AgNO<sub>3</sub> and 1X reaction buffer without any Cl<sup>-</sup> ions.

This initial screen revealed the capacity of *Kf (exo<sup>-</sup>)* at incorporating a modified nucleotide opposite a templating **dPur<sup>P</sup>** unit with ~ 80% efficiency in the strict presence of Ag<sup>I</sup>. We have previously shown<sup>40</sup> that **dIm<sup>C</sup>** sustained the formation of artificial Ag<sup>I</sup>-mediated base pairs under primer extension reaction (PEX) conditions and we therefore rationalized that **dIm<sup>C</sup>TP** could be used instead of **dPur<sup>P</sup>TP** to further improve the specificity and yield of metal base pair formation. Consequently, we used the *Vent (exo<sup>-</sup>)* and *Kf (exo<sup>-</sup>)* polymerases in conjunction with 12 different metal cations to evaluate the possibility of forming **dIm<sup>C</sup>-M<sup>n+</sup>-dPur<sup>P</sup>** base pairs. PEX reactions were carried out with primer **P1** and template **T1** and revealed the formation of the expected **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair for both polymerases and no incorporation of the modified nucleotide (i.e. yields < 10%; Table S2) in the presence of any other metal cation or in the control reactions (Fig. S3). However, despite these promising results, the yield of formation of the extended product remained modest (~50% conversion of the primer). We thus tried to optimize the reaction conditions in order to improve the yield of n+1 product formation (Fig. 3). In the case of *Vent (exo<sup>-</sup>)*, the incorporation of **dIm<sup>C</sup>MP** opposite a templating **dPur<sup>P</sup>** nucleotide does not seem to be dependent on the silver concentration as similar incorporation yields were obtained when the reactions were supplied with 60 and 30 μM Ag<sup>I</sup>. A



slight improvement was observed when the reaction time was increased to 4 hours. For the Kf (*exo*<sup>-</sup>) polymerase, the yield of incorporation of the modified nucleotide was increased to 71% after 2 hours of incubation but the formation of n+1 products could also be detected in the control reaction albeit to a lesser extent (25% conversion of the primer).

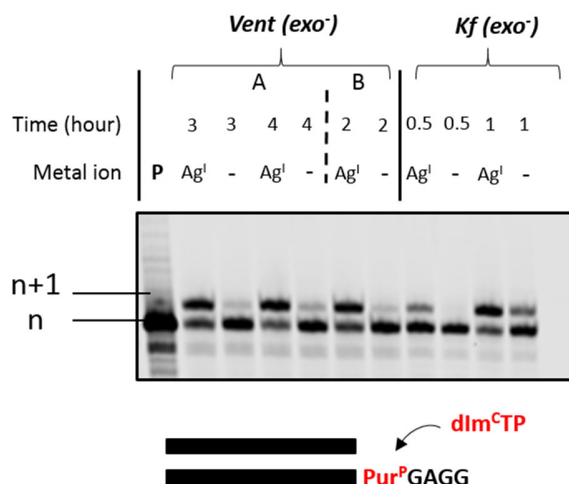


Figure 3. Gel analysis of the time dependence on the incorporation of **dIm<sup>C</sup>TP** (20 μM final concentration) opposite a single templating **dPur<sup>P</sup>** in the presence of Ag<sup>I</sup> using primer **P1** and template **T1** with *Vent (exo<sup>-</sup>)* (2 U) and *Kf (exo<sup>-</sup>)* (5 U) polymerases. Mixtures were incubated at 60 °C or 37 °C for different times. For *Vent (exo<sup>-</sup>)*: A) 60μM of silver, B) 30μM of silver. For *Kf (exo<sup>-</sup>)*: 30 μM of silver. **P** indicates the unreacted primer. Ag<sup>I</sup> reaction conditions were performed with AgNO<sub>3</sub> and 1X reaction buffer without any Cl<sup>-</sup> ions.

Taken together, this analysis revealed the higher propensity of the *Kf (exo<sup>-</sup>)* polymerase at sustaining the enzymatic formation of the **dPur<sup>P</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair and that of *Vent (exo<sup>-</sup>)* for the **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair. Having identified conditions and polymerases that permit the formation of these artificial metal base pairs, we next sought to further optimize the reaction conditions to improve the yields of formation. Full conversion of the primer to the n+1 product (86% yield) could be achieved when **dIm<sup>C</sup>TP** was used in combination with *Vent (exo<sup>-</sup>)*, long reaction times (up to 6h), and 60 μM Ag<sup>I</sup> (Fig. 4). For **dPur<sup>P</sup>TP**, up to 63% yields of n+1 product formation could be observed and the incorporation of the modified nucleotide was minimal in the absence of metal cations. Surprisingly, when modified triphosphate and modified template were inverted (i.e. using **dPur<sup>P</sup>TP** as the incoming triphosphate and **dIm<sup>C</sup>** as the templating nucleotide) under the same experimental conditions, a **dIm<sup>C</sup>-dPur<sup>P</sup>** mispair was formed irrespective of the presence and the nature of the metal cation (Fig. S4).

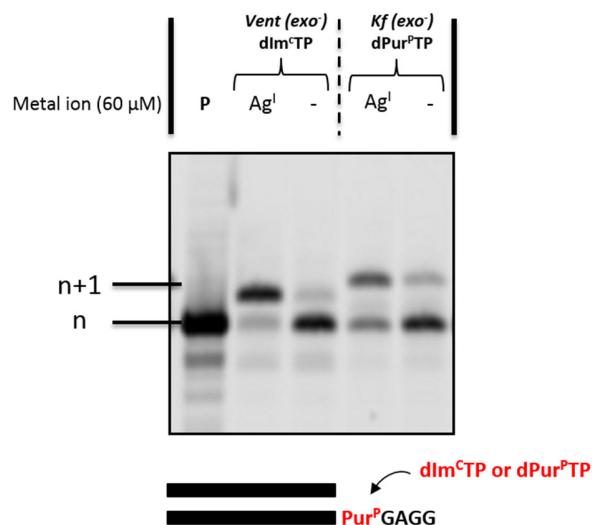


Figure 4. PAGE (20%) analysis of the optimal conditions for the formation of a **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair or **dPur<sup>P</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** in buffer 1 or buffer 2, using the primer **P1**/template **T1** system. For **dIm<sup>C</sup>TP**, the mixtures contained 2 U of Vent (*exo*<sup>-</sup>) polymerase, 20  $\mu$ M final **dIm<sup>C</sup>TP** and were incubated at 60  $^{\circ}$ C for 6 hours. For **dPur<sup>P</sup>TP**, the mixtures contained 5 U of Kf (*exo*<sup>-</sup>) polymerase, 40  $\mu$ M final **dPur<sup>P</sup>TP** and were incubated at 37  $^{\circ}$ C for 30 min. **P** indicates unreacted primer.

#### DFT calculations for modelling possible **dPur<sup>P</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** and **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pairs

The above experimental findings clearly highlighted the formation of non-natural metal base pairs by homopairing between two **dPur<sup>P</sup>** ligands and heteropairing between a **dPur<sup>P</sup>** and a **dIm<sup>C</sup>** nucleotide in the presence of Ag<sup>I</sup> ions. However, we could not observe any formation of the **dPur<sup>P</sup>-Ni<sup>II</sup>-dPur<sup>P</sup>** base pair, although it had been previously reported with synthetic oligonucleotides in UV melting experiments.<sup>41</sup> A similar observation was made previously with **dIm<sup>C</sup>** which forms stable **dIm<sup>C</sup>-Cu<sup>II</sup>-dIm<sup>C</sup>** base pairs with synthetic oligonucleotides in UV melting experiments<sup>26</sup> and a **dIm<sup>C</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** UBP under enzymatic conditions.<sup>40</sup> Also, we were surprised by the fact that a simple inversion of the roles of **dPur<sup>P</sup>** and **dIm<sup>C</sup>** nucleotides led to a completely different outcome during PEX reactions. This prompted us to carry out a DFT analysis in an attempt at rationalizing these findings. Indeed, quantum mechanics calculations are performed on systems involving short synthetic oligonucleotides<sup>50</sup> to confirm the formation of metal base pairs as well as on canonical nucleic acids to further refine our understanding of base pair formation.<sup>51</sup> Therefore, such an analysis was deemed to shed some light into these experimental findings. Thus, we first computationally modelled the possible base pairing geometries involving the homopairing of **dPur<sup>P</sup>** bases mediated by Ag<sup>I</sup>. Three possible base pair models were generated and optimized by DFT calculations, with the 'cis' relative orientation of their glycosidic bonds of the interacting nucleobases relative to the hydrogen

bonds<sup>52</sup>: a canonical Watson-Crick like geometry (cWW-like), a Watson Crick-Hoogsteen like geometry (cWH-like, where a **dPur<sup>P</sup>** uses its Watson Crick edge, while the other one uses its Hoogsteen edge in a cis orientation of the glycosidic bond), and a Hoogsteen-Hoogsteen like geometry (cHH-like, where both the **dPur<sup>P</sup>** bases use their Hoogsteen edges, see Fig. S6a). However, the default glycosidic bonds of the two nucleobases was assumed to be in an anti configuration. We carried out calculations in the presence of one or two coordinating Ag<sup>I</sup> cations, where the two metal cations are believed to be preferred, due to the coordination environment created by both modified nucleotides and to literature precedents.<sup>26, 29, 35, 53</sup>

In case of a single Ag<sup>I</sup> cation, in the optimized cWW-like geometry of the **dPur<sup>P</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** pair, the distance between the C1' atoms of the nucleotides, which gives an indication of their isostericity (i.e. the geometrical equivalence) with natural base pairs<sup>54</sup> is of 11.48 Å, that is only 0.88 Å longer than the average value of dA:dT and dG:dC pairs (10.48 and 10.69 Å, respectively, see Fig. S5). For the cWH- and cHH-like geometries, it is instead of 9.02 Å and 6.68 Å, i.e. shorter by 1.58 Å and 3.92 Å, respectively, as compared to natural A:T/G:C pairs (see Fig. S6, b,c). Further, energy calculations showed that the cWW-like geometry is more stable by 6.9 (5.0) kcal/mol and by 15.3 (10.6) kcal/mol in gas phase (water) as compared to the cWH- and cHH-like geometries, respectively. It is to be remarked that the above geometries resulted from imposing a Cs symmetry during the geometry optimizations, in order to maintain the planarity of the pair. Using no symmetry constraints (a C1 symmetry) resulted indeed in non-planar geometries, possibly due to the electronic repulsion between C2-H on the purine moiety and the C-H group of the pyridine moiety of the **dPur<sup>P</sup>** bases facing each other (see Fig. S7). However, the optimized unconstrained **dPur<sup>P</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** cWW base pair results to be only 4.91 kcal/mol less stable as compared to the constrained planar geometry. This small energy difference is expected to be easily compensated by the stacking interactions that the planar base pair can establish with adjacent bases in the DNA duplex and also by the energy gain due to absence of perturbation of adjacent base pair geometries.<sup>55</sup> We also tried to constrain the C1'-C1' distance of the cWW-like base pair to the ideal value of 10.6 Å (averaged over the two canonical dA:dT/dG:dC base pairs). This resulted in a geometry that is only 2.02 kcal/mol less stable than the Cs symmetry constrained optimized relaxed pair, in the gas phase. The above analyses support the experimental finding that **dPur<sup>P</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pairs can be readily formed in a geometry that perfectly fits a regular DNA duplex.

Then, we modelled the **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** base pair. The gas-phase optimized model structure is shown in Fig. S6d. The resulting C1'-C1' distance is 11.79 Å, that is ~1.2 Å longer than in canonical base pairs. Interestingly, also in the absence of symmetry constraints (C1 symmetry), no significant structural deformation is observed for the **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** geometry, which remains substantially planar. The energy difference between the Cs symmetry constrained and unconstrained base pair geometries is indeed negligible and amounts to 0.06 kcal/mol. Fixing the C1'-C1' distance to the ideal value of 10.60 Å results in

an energy loss of 3 kcal/mol. Both optimized geometries and energetics also support the successful formation of **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** base pairs with a duplex-compatible geometry.

In order to rationalize the difference observed when **dPur<sup>P</sup>** served as the incoming triphosphate from when it is integrated in the template, we considered two possible pathways for the formation of the **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** heterobase pair (see Fig. 5). In the former case, the base pairing energies were computed under the assumption that the bidentate **dPur<sup>P</sup>** ligand binds to the Ag<sup>I</sup> ion first, leading to the formation of **dPur<sup>P</sup>-Ag<sup>I</sup>**, with the subsequent addition of the **dIm<sup>C</sup>** ligand, resulting in the **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** complex (Scheme 1 in Fig. 5). In the latter modelled pathway, a bidentate **dIm<sup>C</sup>** binds first to Ag<sup>I</sup>, leading to the formation of **dIm<sup>C</sup>-Ag<sup>I</sup>**, with the subsequent addition of the **dPur<sup>P</sup>** ligand, resulting in the same **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** complex (Scheme 2 in Fig. 5). Our energy calculations indicate that silver-mediated base pair formation seems to be favourable for both modelled reactions (see Table S3). However, the former pathway is clearly favoured over the latter one with an energy difference of 90.5 kcal/mol and 3.6 kcal/mol in the gas phase and in water, respectively. Hence, metal coordination to **dPur<sup>P</sup>** appears to be the first step required for metal base pair formation which occurs preferentially when this analogue serves as a templating nucleotide. While this analysis does not fully explain the preference that was observed, it clearly demonstrates that metal coordination to the templating nucleotide is an important prerequisite for and an important first step in the formation of artificial metal base pairs under polymerase-mediated synthesis conditions.

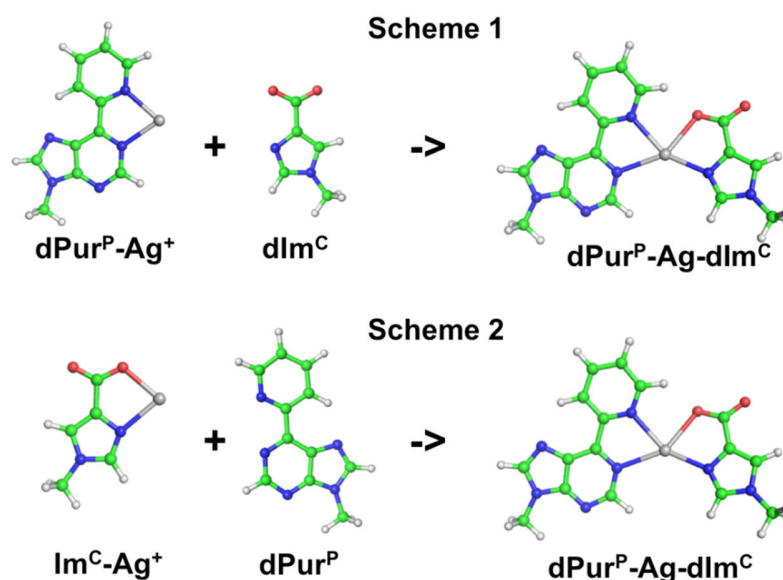


Figure 5: Formation of **dPur<sup>P</sup>-Ag<sup>I</sup>-dIm<sup>C</sup>** complex followed by two possible pathways depicted in Scheme 1 and Scheme 2.

As mentioned above, in addition to the mononuclear  $\text{Ag}^{\text{I}}$  mediated base pairs, we have also investigated the above base pair geometries mediated by two coordinating  $\text{Ag}^{\text{I}}$  ions (Fig. 6 a-d).

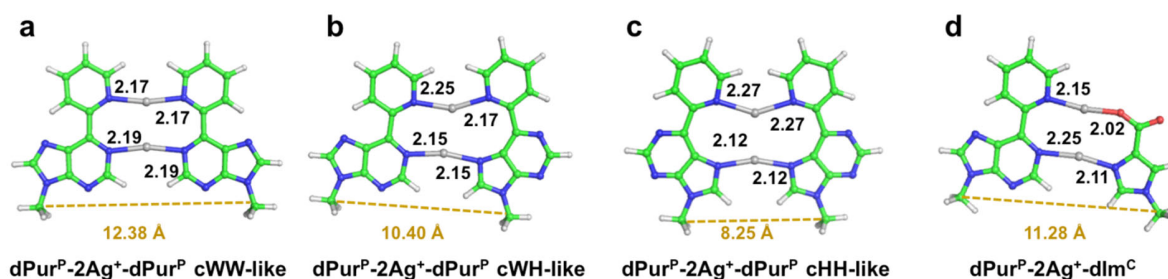


Figure 6. a-d) Optimized  $\text{dPur}^{\text{P}}\text{-2Ag}^{\text{I}}\text{-dPur}^{\text{P}}$  base pairs in different geometries: cWW like (a), cWH like (b) and cHH like (c). d) Optimized structure of the  $\text{dPur}^{\text{P}}\text{-2Ag}^{\text{I}}\text{-dIm}^{\text{C}}$  base pair. Optimal distances in gas phase of the key atoms from the coordinated  $\text{Ag}^{\text{I}}$  ion are reported. C1'-C1' distances are also shown for all the base pairs. A  $\text{C}_s$  symmetry constraint was imposed during the geometry optimizations to retain planarity.

For the  $\text{dPur}^{\text{P}}$  homopairing, our modeling studies show that the insertion of an additional  $\text{Ag}^{\text{I}}$  ion increases the C1'-C1' distances, as compared to the corresponding mono  $\text{Ag}^{\text{I}}$  mediated base pairs, for all the geometries (by 0.90 Å, 1.38 Å and 1.57 Å for cWW-, cWH- and cHH-like geometries, respectively, see Fig. 6a-c). As a consequence, the  $\text{dPur}^{\text{P}}\text{-2Ag}^{\text{I}}\text{-dPur}^{\text{P}}$  cWH-like geometry, with a C1'-C1' distance of 10.40 Å, results to be highly isosteric to the canonical A:T/G:C base pairs (10.48/10.69 Å). As for the energies, the dinuclear  $\text{Ag}^{\text{I}}$  mediated base pairs are less stable than the mononuclear  $\text{Ag}^{\text{I}}$  mediated ones by 1.38, 2.41 and 6.83 kcal/mol for the cWW-, cWH- and cHH-like geometries in the gas phase. When the pairing of  $\text{dPur}^{\text{P}}$  with  $\text{dIm}^{\text{C}}$  is considered; its C1'-C1' distance gets shortened by 0.51 Å as a consequence of the coordination of the second  $\text{Ag}^{\text{I}}$  cation, thus getting closer to the canonical 10.6 Å distance. Furthermore, the pairing appears to be more stable than in the case of the mono  $\text{Ag}^{\text{I}}$  coordinated base pair by 77.35 kcal/mol and 10.48 kcal/mol in the gas phase and in water, respectively. Lastly, calculations carried out in the absence of  $\text{Ag}^{\text{I}}$  suggest the possible formation of weak H-bonding pairing between  $\text{dPur}^{\text{P}}$  with  $\text{dIm}^{\text{C}}$  (Fig. S9) which might account for some results observed during our initial polymerase screen where complete formation of the n+1 product was observed even in the absence of metal cations with the Taq polymerase (Fig. S1).

Taken together, our analyses of geometries and energies suggest that both the single and the dinuclear  $\text{Ag}^{\text{I}}$  coordination modes are in principle possible for the homopairing between  $\text{dPur}^{\text{P}}$  bases. However, both for

energetic and geometrical reasons the cWW- and cWH-like geometries seem to be favored over the cHH-like ones. As for the heteropairing between **Pur<sup>P</sup>** and **Im<sup>C</sup>**, it seems to be favored both energetically and geometrically in the presence of a second coordinating Ag<sup>I</sup> ion.

## By-pass experiments

Having established conditions that permit the enzymatic construction of a **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair we next assessed whether enzymatic DNA synthesis could be resumed with canonical nucleotides after the construction of the metal base pair. By-passing a UBP is crucial required for the construction of modified libraries with view of their use in selection experiments but also for the general construction of modified oligonucleotides.<sup>56-58</sup> To do so, we first installed a **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair by PEX reaction using the **P1/T1** system and the experimental conditions described in Fig. 4. After addition of the four canonical dNTPs, only one additional dC nucleotide was incorporated immediately after the artificial metal base pair and DNA synthesis halted at this stage (Fig. S8a). We first thought that the inability of the polymerase to further extend the primer might be ascribed to the presence of remaining Ag<sup>I</sup> in the reaction mixture, which could disturb the formation of natural base pairs. Therefore, we added 1.2 equivalents of NaCl or EDTA after the formation of the **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair and prior to the addition of natural dNTPs but this had no positive impact on the bypass efficiency (Fig. S8b). We next conducted bypass experiments using the more permissive polymerase *Therminator*<sup>59-60, 61</sup> which is known to recognize UBPs.<sup>62</sup> Under these conditions, the formation of the n+6 product was observed by gel electrophoresis (Fig. 7) which corresponds to the expected full length sequence with an additional nucleotide stemming from a non-templated addition, showing that the **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair can efficiently be by-passed by the *Therminator* polymerase (68% formation of the n+6 product). To exclude the formation of any mispairs (especially dG:dT pairs) by the error prone *Therminator* polymerase and to confirm the identity of the formed product, an LC-MS analysis was performed. To do so, we first conducted by-pass experiments with a 5'-phosphorylated template **T1** (obtained by PNK treatment of **T1**), subjected the resulting dsDNA product to a  $\lambda$ -exonuclease-mediated degradation, and analyzed the ssDNA product by LC-MS (Fig. S19 and Fig. S20). This analysis univocally proved the formation of the expected full length product and further confirmed the untemplated addition of an additional dA nucleotide, which has previously been observed with *Therminator* (Fig. S20).<sup>61, 63</sup> Thus, this represents a convenient method for the synthesis of DNA duplexes containing a **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** UBP.

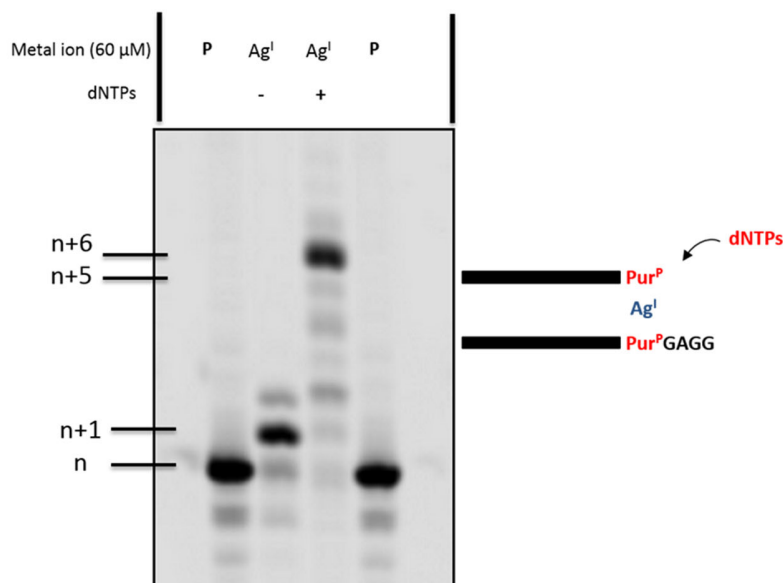


Figure 7. Gel (PAGE 20%) image analysis of the modified base pair by-pass experiments. All the reactions were carried out with primer **P1** and phosphorylated template **T1** and first incubated in the presence of **dIm<sup>C</sup>TP** (20 μM) and 2 U of Vent (*exo*<sup>-</sup>) polymerase at 60°C for 6h. dNTPs (20 μM) were added with 10 U of Therminator polymerase and incubated at 60°C for 30 min. **P** indicates unreacted primer.

### Specificity of the incorporation

Next, we sought to determine the specificity of the incorporation of canonical dNTPs opposite a templating **dPur<sup>P</sup>** nucleotide and conversely the specificity of the incorporation of the modified triphosphate **dPur<sup>P</sup>TP** opposite wild-type nucleotides. These experiments are important to evaluate the possibility of using the **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair system in SELEX experiments, specifically in the PCR amplification and sequencing steps. For both Vent (*exo*<sup>-</sup>) and Kf (*exo*<sup>-</sup>) polymerases, we carried out primer extension reactions with template **T1** and each individual dNTP, in the presence and absence of silver ions (Fig. 8). Even though synthetic duplexes were shown to be strongly destabilized in the presence of **dPur<sup>P</sup>** mismatches with all the natural bases,<sup>41</sup> the efficient incorporation of a dT nucleotide opposite **dPur<sup>P</sup>** was observed with both polymerases. Yet, more surprisingly, Vent (*exo*<sup>-</sup>) misincorporated dA opposite the modification with around 35% yield of the n+1 product regardless on the presence or absence of Ag<sup>I</sup> (38% yield without addition of silver, 32% yield in the presence of silver). In addition, n+1 and n+2 products can also be observed with dCTP corresponding to the formation of dC-**dPur<sup>P</sup>** and dC-Ag<sup>I</sup>-**dPur<sup>P</sup>** mispairs. The latter was not unexpected since silver is known to mediate the formation of dC-Ag<sup>I</sup>-dA base pairs.<sup>64</sup> When Vent (*exo*<sup>-</sup>) was used in PEX reactions, misincorporation of dG opposite the modification occurred at much lower levels

than with dC, dA, and dT regardless on the presence or absence of  $\text{Ag}^{\text{I}}$ . In the case of the Kf (*exo*<sup>-</sup>) polymerase, while dC (6% incorporation with or without silver) or dG (6% incorporation with or without silver) nucleotides are not misincorporated opposite **dPur<sup>P</sup>**, dAMP units are efficiently incorporated with or without  $\text{Ag}^{\text{I}}$  (80% n+1 product formation without silver and 75% with silver).

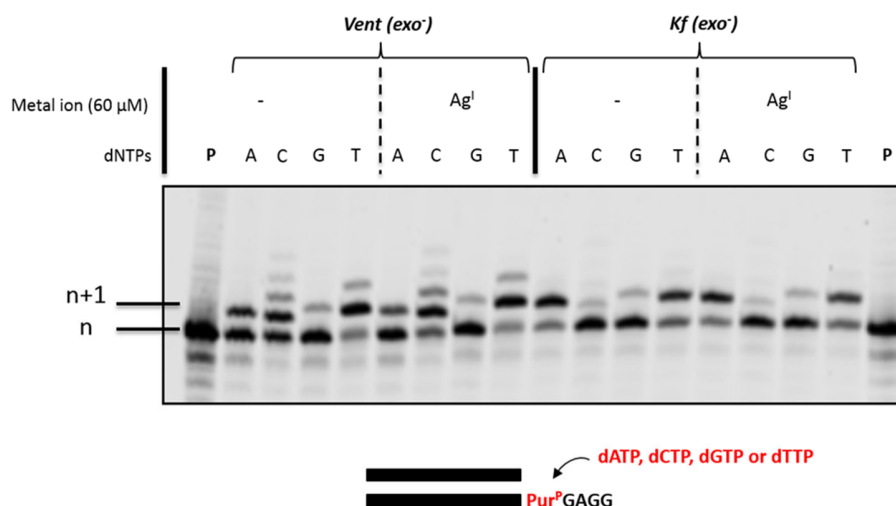


Figure 8. Extension reactions of the **dPur<sup>P</sup>** containing template **T1** with each individual dNTP. All reactions were conducted using the primer **P1** and Vent (*exo*<sup>-</sup>) (2 U) or Kf (*exo*<sup>-</sup>) (5 U) polymerase. The reaction mixtures contained natural dNTPs (200  $\mu\text{M}$  final concentration) and were incubated for 30min at 60°C or 37 °C. **P** indicates unreacted primer.

In order to investigate the specificity of the incorporation of **dPur<sup>P</sup>TP**, we carried out PEX reactions using templates **T3-T6** (Table 1) that contain each one of the canonical bases at the site of the modification in **T1**. This analysis clearly revealed that **dPur<sup>P</sup>MP** was readily incorporated opposite templating dC and dT nucleotides by the Vent (*exo*<sup>-</sup>) polymerase, even in the absence of silver cations. A partial incorporation of the modified nucleotide opposite dG was also observed (Fig. 9). The presence of silver seemed to inhibit the incorporation of **dPur<sup>P</sup>TP** opposite a dA nucleotide as 81% incorporation occurred in the absence of  $\text{Ag}^{\text{I}}$ , compared to 63% incorporation in its presence. For the Kf (*exo*<sup>-</sup>) polymerase, partial incorporation of the modified nucleotide was detected opposite dA and dG in the absence of  $\text{Ag}^{\text{I}}$  but not opposite the other two canonical nucleotides. Addition of  $\text{Ag}^{\text{I}}$  seems to negatively affect the incorporation opposite a dA nucleotide but to favor the formation of base pairs with dC, dG and dT. Therefore, the incorporation of **dPur<sup>P</sup>MPs** does not exclusively occur when the templating nucleotide is **dPur<sup>P</sup>** since misincorporation events are observed.



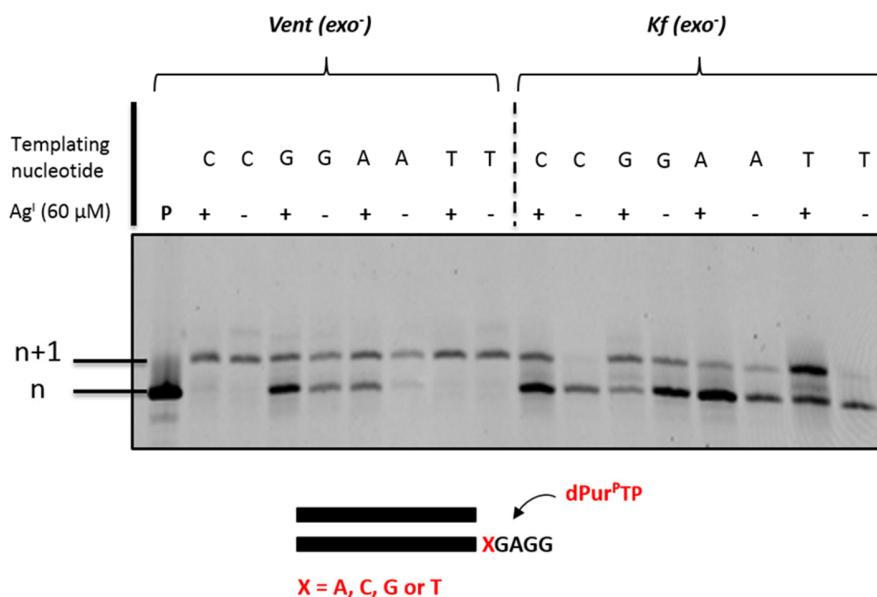


Figure 9. PEX reactions with **dPur<sup>P</sup>TP** modified nucleotide using the templates **T3-T6** and Vent (*exo*-) (2 U) or Kf (*exo*-) (5 U) as polymerases. The reaction mixtures contained **dPur<sup>P</sup>TP** at 20 μM final concentration and were incubated for 30min at 60°C or 37 °C. **P** indicates unreacted primer.

### DFT calculations for modelling the mispairs of dPur<sup>P</sup> and canonical nucleobases

UV melting experiments performed on dodecamers containing a single **dPur<sup>P</sup>** modification have shown that mismatches between **dPur<sup>P</sup>** and canonical nucleotides led to strong destabilization of the resulting duplexes. However, PEX reactions indicated that **dPur<sup>P</sup>** nucleotides (both as incoming triphosphate or templating nucleotide) misincorporated particularly well opposite canonical dT. In order to try to rationalize these findings we modelled pairing of **dPur<sup>P</sup>** with the four natural nucleobases, with and without the Ag<sup>I</sup> ion coordination. As the Ag<sup>I</sup> coordination is well known to occur with the nitrogen atoms of the heterocycles for both the natural and unnatural bases, in our modelling study we assumed the coordination to occur at the Watson-Crick edge of all the bases and, in addition, at the Hoogsteen edge of A and G (to N7) (cWH-like geometries, Fig. 10).

The C1'-C1' distances of the possible Ag<sup>I</sup> mediated base pairs are larger than the average C1'-C1' distance of canonical dA:dT/dG:dC base pairs for the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dA and **dPur<sup>P</sup>**-Ag<sup>I</sup>-dT base pairs (by 1.8 Å and 1.5 Å, respectively) and shorter by 0.9 Å for the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dC base pair. For the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dG base pair, an elongation of the C1'-C1' distance by 4.35 Å is observed, which may be considered incompatible with a regular DNA duplex. However, considering the Ag<sup>I</sup> coordination to the N7 of the A and G bases, in their

cWH-like geometries, the C1'-C1' distance of the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dG pair gets elongated by 1.1 Å, while it is extremely similar (10.74 Å) to the canonical A:T/G:C pairs for **dPur<sup>P</sup>**-Ag<sup>I</sup>-dA. Regarding energies, except for the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dC cWW-like pair, which is more stable than the corresponding **dPur<sup>P</sup>**-Ag<sup>I</sup>-**dPur<sup>P</sup>** pair (by 0.64 and 2.97 kcal/mol in the gas phase and in water, respectively), all the other Ag<sup>I</sup> mediated pairs between **dPur<sup>P</sup>** and the Watson-Crick edge of the natural nucleobases are less stable, with energy differences within 16.27 kcal/mol and 9.28 kcal/mol in the gas phase and in water, respectively. However, looking at the cWH geometries, the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dA pair is further destabilized as compared to the cWW-like geometry (by 3.55/1.56 kcal/mol in gas phase/water). On the contrary, the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dG cWH-like pair results more stable than the corresponding cWW-like geometry (by 16.2/11.0 kcal/mol in gas phase/water), also due to the formation of an additional weak H-bonding interaction (see Fig. 10a).

The high stability (and isostericity) of the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dC cWW-like pair observed in this DFT analysis is reflected in PEX reactions conducted with the Kf (*exo*<sup>-</sup>) polymerase since, when **dPur<sup>P</sup>** acts as an incoming triphosphate, formation of the mispair is only observed in the presence of Ag<sup>I</sup> (Fig. 9). However, the formation of the **dPur<sup>P</sup>**-Ag<sup>I</sup>-dC pair could not be observed (<5% product formation) when the modified nucleotide was incorporated in the template (Fig. 8). On the other hand, the incorporation events mediated by the Vent (*exo*<sup>-</sup>) polymerase are more difficult to rationalize based on this DFT analysis.

We have also modelled the possible H-bonded base pairs that are formed in the absence of Ag<sup>I</sup> (relative geometries are shown in Fig. 10b). All the base pairs have at least one strong N-H...N H-bond, along with one or more weak C-H...O/N H-bonds. Compared to classical dA:dT/dG:dC cWW geometries, the C1'-C1' distances between the H-bonded mispairs is larger for the **dPur<sup>P</sup>**:dA and **dPur<sup>P</sup>**:dG cWW base pairs, by 1.87 Å and 1.95 Å respectively, and shorter by 0.93 Å for the **dPur<sup>P</sup>**:dT cWW base pair. As for the **dPur<sup>P</sup>**:dC pair, its C1'-C1' distance is quite close (10.22 Å) to the canonical one. Therefore, especially the **dPur<sup>P</sup>**:dA and **dPur<sup>P</sup>**:dG pairs seem geometrically unfit to be part of a regular DNA duplex in their cWW geometries. That notwithstanding, the **dPur<sup>P</sup>**:dG cWW pair results to be in principle slightly more stable than the dA:dT pair, both in the gas phase and in water (by -2.61 and 1.42 kcal/mol respectively). On the contrary, the **dPur<sup>P</sup>**-dA/dT/dC base pairs are less stable than the corresponding dA:dT base pair with energy differences within 5.57 and 4.50 kcal/mol, in the gas phase and in water, respectively (see Table S3).

As they appear unfit to match a regular duplex in their cWW geometries, we also modelled the **dPur<sup>P</sup>**:dA and **dPur<sup>P</sup>**:dG base pairs in their cWH geometries. Interestingly, in the cWH geometry, the C1'-C1' distance of the **dPur<sup>P</sup>**:dA pair is extremely close (10.33 Å) to the “canonical” value, while its energy remains similar to that of the cWW geometry (with energy differences within 0.14 kcal/mol). These findings clearly indicate that the preferred geometry for the **dPur<sup>P</sup>**:dA pair, to be incorporated in a DNA duplex, would be the cWH one. In contrast, for **dPur<sup>P</sup>**:dG a cWH geometry could not be modelled, because of the unavailability of

complementary H-bond donor/acceptor atoms on its facing edges (see Fig. 10). This analysis is in agreement with the experimental data showing that the **dPur<sup>P</sup>**:dG pairing is not observed under enzymatic DNA synthesis.

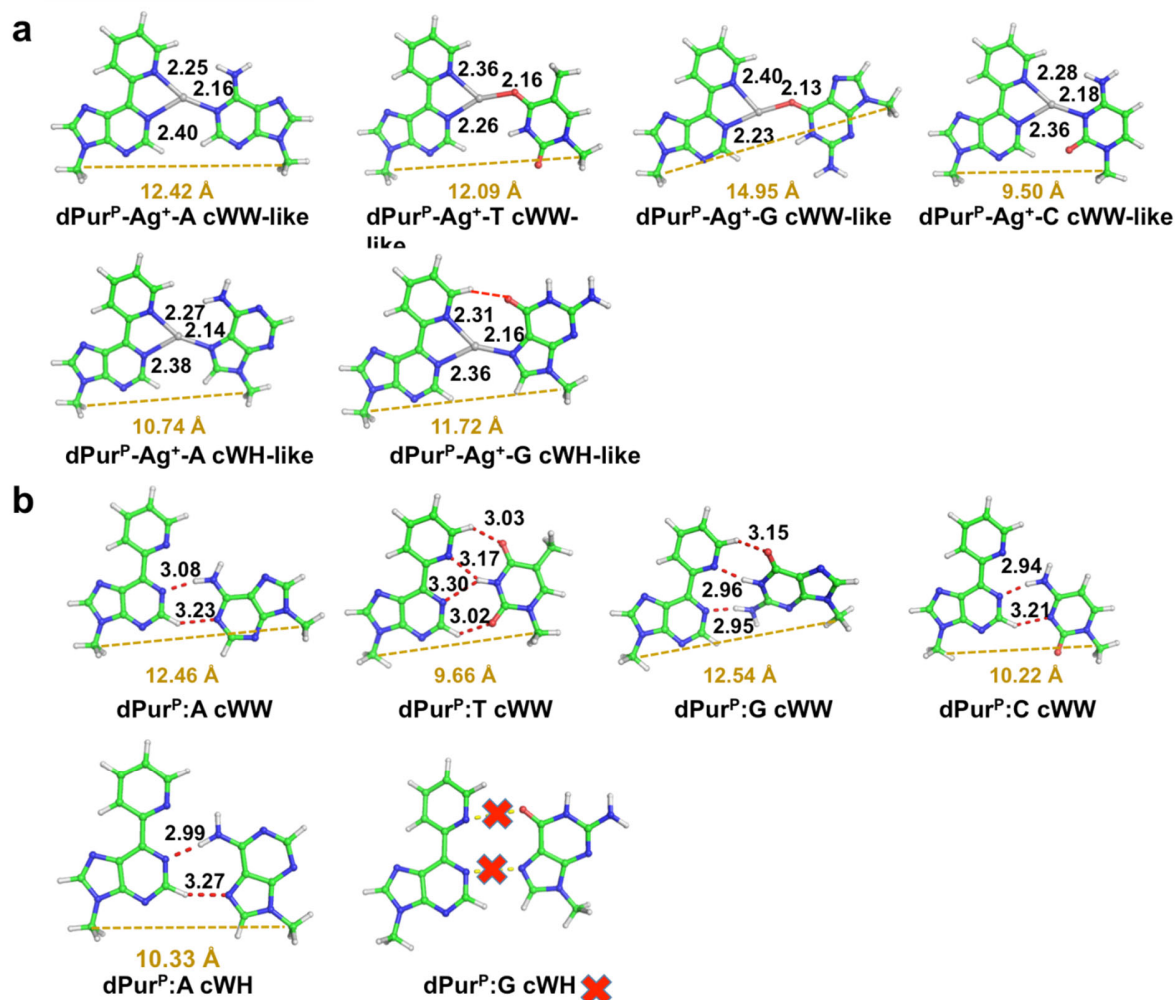


Figure 10. **a**) Geometry optimized structures of possible Ag<sup>I</sup> mediated base pairs between the non-natural **dPur<sup>P</sup>** and natural nucleobases using Watson-Crick edge (W) and Hoogsteen edge (H); **b**) Geometry optimized structures of mismatched H-bond mediated base pairs between **dPur<sup>P</sup>** and natural nucleobases. Modelling of pairing of **dPur<sup>P</sup>** and dG in a cWH geometry is not feasible due to the proximity of acceptor atoms opposite to each other, marked by red cross signs in figure. The natural nucleobases have been truncated at the C1' atom.

## CONCLUSIONS

The decoration of nucleic acids with metal ions through the formation of artificial metal base pairs is an alluring method for improving the chemical diversity and conferring additional properties to DNA and RNA. Nucleic acids containing metal base pairs have found applications in numerous fields including the construction of logic gate devices and the development of novel nanomaterials.<sup>65-68</sup> The enzymatic construction of artificial metal base pairs represents an alluring alternative method that alleviates the synthetic burden and size limitation impaired to traditional approaches involving solid-phase synthesis. Here, we demonstrate that the two synthetic analogues **dIm<sup>C</sup>** and **dPur<sup>P</sup>** can be used for the enzymatic synthesis of Ag<sup>I</sup>-mediated UBPs. In addition, we have developed a two-step enzymatic method that permits the installation of a **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair followed by natural DNA synthesis and could be extended to any type of oligonucleotide. Interestingly, the metal preference under enzymatic DNA synthesis conditions appears to be the opposite of that unraveled by UV melting experiments on short synthetic oligonucleotides since Ni<sup>II</sup> strongly stabilized a **dPur<sup>P</sup>-dPur<sup>P</sup>** mismatch while coordination to Ag<sup>I</sup> appeared to be possible but only mildly stabilizing.<sup>41</sup> The DFT calculations support formation of Ag<sup>I</sup>-mediated base pairs and shed some light into the mechanism of formation. Particularly, metal coordination of the templating nucleotide appears to be the first step in the formation of metal base pairs under enzymatic synthesis conditions. While PEX reactions performed with natural nucleotides reveal that the **dIm<sup>C</sup>** and **dPur<sup>P</sup>** system is not fully compatible with the construction of libraries for SELEX experiments due to misincorporation events, this study clearly demonstrates the possibility of synthesizing DNA duplexes containing artificial metal base pairs using PEX reactions. Collectively, the results demonstrate that the combination of a [2+2] metal coordination environment and increased stacking interactions offered by the **dIm<sup>C</sup>-Ag<sup>I</sup>-dPur<sup>P</sup>** base pair appear to be key elements for the design of nucleotide ligands for the enzymatic construction of metal UBPs.

## EXPERIMENTAL SECTION

### Materials and Methods

All reactions were performed under argon in flame-dried glassware. Anhydrous solvents for reactions were obtained from Sigma Aldrich. Flash chromatography was performed using silica gel (230–400 mesh) from Sigma Aldrich. Thin layer chromatography was carried out on pre-coated glass-backed plates of silica gel (0.25 mm, UV254) from Macherey-Nagel. All chemicals and solvents used were purchased from Sigma-Aldrich and Alfa Aesar. NMR spectra were recorded on a Bruker Avance 400 spectrometer (400.13 MHz for <sup>1</sup>H, 100.62 MHz for <sup>13</sup>C, and 161.62 MHz for <sup>31</sup>P) and all spectra were referenced to the signals of the corresponding solvent. Chemical shifts are given in ppm ( $\delta$  scale) and coupling constants ( $J$ ) in Hz.

Assignment of the NMR signals was performed by using a combination of  $^1\text{H}/^1\text{H}$ -COSY,  $^{13}\text{C}$ -DEPT-135, and  $^{13}\text{C}/^1\text{H}$ -HMBC experiments. Crude **dPur<sup>P</sup>TP** was purified by HPLC using an Äkta<sup>TM</sup> pure system (GE Healthcare) equipped with Thermo Scientific<sup>TM</sup> DNAPac<sup>TM</sup> PA100 semi preparative ion exchange column (13 $\mu\text{m}$ , 250 x 9.0 mm) with Buffer A) TEAB 10mM pH=8; Buffer B) TEAB 1M pH=8). High resolution electrospray ionization (ESI) mass spectra (MS,  $m/z$ ) were recorded on a Waters Q-Tof Micro MS in the positive-ion electrospray ionization (ESI<sup>+</sup>) mode. Solutions were prepared using 1:1 MeCN/H<sub>2</sub>O containing 0.1% formic acid or MeOH/water containing 10 mM ammonium acetate in the case of sensitive compounds. HPLC purification was performed using an Äkta<sup>TM</sup> pure system (GE Healthcare) equipped with Thermo Scientific<sup>TM</sup> DNAPac<sup>TM</sup> PA100 semi preparative ion exchange column (13 $\mu\text{m}$ , 250 x 9.0 mm). Unmodified DNA oligonucleotides were purchased from Microsynth. DNA oligonucleotides containing modified nucleotides were synthesized on an H-8 DNA synthesizer from K&A on a 1  $\mu\text{mol}$  scale. Natural DNA phosphoramidites (dT, dC4bz, dG2DMF, dA6Bz) and solid support (dA6Bz-lcaa-CPG 500Å) were all purchased from ChemGenes. The phosphoramidite was prepared by application of literature protocols.<sup>45</sup> Natural DNA phosphoramidites as well as the modified phosphoramidite were prepared as 0.07 M solutions in MeCN and were coupled using 50 sec and 490 sec steps, respectively. 5-(ethylthio)-1*H*-tetrazole (0.25 M in MeCN) was used as coupling agent. Capping, oxidation, and detritylation were performed using standard conditions. Cleavage from the solid support and deprotection of oligonucleotides was achieved by treatment with concentrated ammonia at 55°C for 16 h. After centrifugation, the supernatants were collected and the resulting solutions were evaporated to dryness on a speed-vac. Crude oligonucleotides were purified by anion exchange HPLC (Dionex - DNAPac PA200). Buffer solutions of 25 mM Tris-HCl in H<sub>2</sub>O, pH 8.0 (buffer A) and 25 mM Tris-HCl, 1.25 M NaCl in H<sub>2</sub>O, pH 8.0 (buffer B) were used. The purified oligonucleotides were then desalted with SepPack C-18 cartridges. Oligonucleotide concentrations were quantitated by UV spectroscopy using a UV5Nano spectrophotometer (Mettler Toledo). The chemical integrity of oligonucleotides was assessed by UPLC-MS analysis: UPLC was performed on a BEH C18 column (130 Å, 1.7  $\mu\text{m}$ , 2.1 mm x 50 mm) from Waters, installed on an ACQUITY UPLC H-Class System (SQ Detector 2). A Buffer containing 20 mM TEA and 400 mM HFIP in H<sub>2</sub>O was used with a linear gradient from 18 to 31% Methanol within 5 minutes and a flow rate of 0.3 mL/min. All the DNA polymerases (Therminator, Vent (*exo*<sup>-</sup>), Deep Vent, Bst, Taq, and Klenow fragment of DNA polymerase I *exo*<sup>-</sup> (Kf (*exo*<sup>-</sup>))) were purchased from New England Biolabs as well as the natural dNTPs. Acrylamide/bisacrylamide (29:1, 40%) was obtained from Fisher Scientific. Visualization of PAGE gels was performed by fluorescence imaging using a Storm 860 or a Typhoon Trio phosphorimager with the ImageQuant software (both from GE Healthcare). The following salts were used: AgNO<sub>3</sub>, CdCl<sub>2</sub>, CuSO<sub>4</sub>, CoCl<sub>2</sub>, EuCl<sub>3</sub>, FeSO<sub>4</sub>, FeCl<sub>3</sub>, HgCl<sub>2</sub>, MnCl<sub>2</sub>, NiSO<sub>4</sub>, Pb(OAc)<sub>2</sub>, ZnSO<sub>4</sub>.

## Reaction buffers

The following 10x buffers without any Cl<sup>-</sup> source were prepared for PEX reactions in presence of Ag<sup>I</sup>. Buffer 1 is used for Vent (*exo*<sup>-</sup>), buffer 2 for Kf (*exo*<sup>-</sup>), buffer 3 for Taq and buffer 4 for Bst.

Buffer 1: 200 mM Tris acetate, 100 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 100 mM KClO<sub>4</sub>, 20 mM MgSO<sub>4</sub> 0.1% Triton® X-100, pH 8.8 at 25°C; buffer 2: 500 mM NaClO<sub>4</sub>, 100 mM Tris acetate, 100 mM MgSO<sub>4</sub>, 100 μM DTT, pH 7.9; buffer 3: 100 mM Tris acetate, 500 mM KClO<sub>4</sub>, 15 mM MgSO<sub>4</sub>, pH 8.3 at 25°C; buffer 4: 200 mM Tris acetate, 100 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 500 mM KClO<sub>4</sub>, 20 mM MgSO<sub>4</sub>, 0.1% Tween® 20, pH 8.8 at 25°C.

### Synthesis of dPur<sup>P</sup>TP

**Synthesis of 3'-O-acetyl-1-(2-deoxy-β-D-ribofuranosyl)-adenine-6-pyridin-2-yl (6).** Nucleoside analogue **4** [35] (200 mg, 0.367 mmol, 1 eq) was dissolved in dry pyridine (2 mL) at RT under Ar. To this solution, 2-(dimethylamino)pyridine (9 mg, 0.073 mmol, 0.2 eq) was added followed by trimethylamine (128 μL, 0.919 mmol, 2.5 eq) and subsequently acetic anhydride (0.1 mL, 1.103 mmol, 3 eq). After 1h of stirring, the reaction mixture was quenched with NaHCO<sub>3</sub> sat. (20 mL) and extracted with DCM (3x 30 mL). The organic layers were combined, dried over MgSO<sub>4</sub>, and concentrated. The crude product was used in the second step without further purification. The starting material **5** (120 mg, 0.182 mmol, 1 eq) was dissolved in anhydrous DCM (5 mL) under Ar. To this solution, 5 mL of TCA 3% in DCM (5 mL) was added. The reaction mixture was stirred for 30 min at room temperature. The solvent was removed *in vacuo* and the residue was purified by flash chromatography (DCM/MeOH 96:4) to give 45 mg of a white solid (70%). HR-MS for C<sub>17</sub>H<sub>18</sub>N<sub>5</sub>O<sub>4</sub><sup>+</sup> m/z calcd: 356.1359; found: 356.1357. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ = 2.16 (s, 3H), 2.51-3.31 (m, 2H), 3.95-4.05 (m, 2H), 4.33 (s, 1H), 5.61-5.63 (m, 1H), 6.44-6.47 (m, 1H), 7.47-7.50 (m, 1H), 7.94-7.98 (m, 1H), 8.35 (s, 1H), 8.83 (d, *J* = 8.0 Hz, 1H), 8.99 (d, *J* = 4.0 Hz, 1H), 9.14 (s, 1H). <sup>13</sup>C NMR (101 MHz, CDCl<sub>3</sub>) δ = 21.0, 37.7, 63.2, 76.2, 87.4, 87.6, 125.2, 125.8, 133.1, 137.0, 145.0, 150.4, 151.9, 153.3, 154.5, 170.3.

**Synthesis of dPur<sup>P</sup>TP (3).** Nucleoside **6** (45 mg, 0.127 mmol, 1 eq) was dissolved in dry pyridine (0.2 mL) and dry dioxane (0.4 mL) at RT under Ar. To this clear solution, 2-chloro-1,3,2-benzodioxaphosphorin-4-one (38 mg, 0.185, 1.4 eq.) was added and the reaction mixture was stirred for 45 min. A solution of tributylammonium pyrophosphate (120 mg, 0.219 mmol, 1.7 eq) in dry DMF (0.1 mL) and tributylamine (60 μL) was added dropwise and the reaction mixture stirred for another 45 min. The reaction mixture was then oxidized by the addition of iodine (64 mg, 0.203 mmol, 1.6 eq) in pyridine (1 mL) and H<sub>2</sub>O (0.2 mL). After 30 min of stirring, the excess of iodine was quenched with a sodium thiosulfate solution (10% w/v in water) and the resulting clear solution was concentrated under reduced pressure at 30°C. The concentrated mixture was treated with ammonium hydroxide 30% (10 mL) for 2h. The yellow suspension was again concentrated under reduced pressure at 30°C. The yellow residue was dissolved in H<sub>2</sub>O (2 mL) and

precipitated by the addition of NaClO<sub>4</sub> 2% in acetone (12 mL). The crude product was purified by ion exchange HPLC (50% B in 20 min; Buffer A) TEAB 10mM pH=8; Buffer B) TEAB 1M pH=8) to yield 5.6 mg (8 %) of pure triphosphate **3**. HR-MS C<sub>15</sub>H<sub>17</sub>P<sub>3</sub>N<sub>5</sub>O<sub>12</sub><sup>-</sup> m/z calcd: 552.0087; found: 552.0083. <sup>1</sup>H NMR (400 MHz, D<sub>2</sub>O) δ = 2.55-2.86 (m, 2H), 2.94-3.26 (m, 2H), 4.11-4.14 (m, 2H), 4.24 (s, 1H), 6.62 (t, 1H), 7.57 (s, 1H), 8.04 (s, 1H), 8.30 (s, 1H), 8.69 (s, 1H), 8.81 (s, 1H), 8.96 (s, 1H). <sup>31</sup>P NMR (202 MHz, D<sub>2</sub>O) δ = -26.18 (t, *J* = 22.2 Hz 1P), -14.40 (d, *J* = 20.2 Hz 1P), -13.74 (d, *J* = 18.2 Hz 1P).

### Primer extension experiments

The 5'-FAM-labelled primer **P1** (10 pmol) was annealed to the appropriate template (15 pmol) in water by heating to 95 °C and then gradually cooling down to room temperature (over 30 min). The appropriate DNA polymerase and metal cation solutions were then added to the annealed oligonucleotides mixture on ice. Finally, natural dNTPs and/or modified triphosphate were added for a total reaction volume of 10 µL. Following incubation at the optimal temperature for the enzyme, the reactions were quenched by adding stop solution (10 µL; formamide (70%), ethylenediaminetetraacetic acid (EDTA; 50 mM), bromophenol (0.1%), xylene cyanol (0.1%)). The reaction mixtures were subjected to gel electrophoresis in denaturing polyacrylamide gel (20%) containing trisborate-EDTA (TBE) 1× buffer (pH 8) and urea (7 M). Visualization was performed by fluorescence imaging using a Storm 860 or a Typhoon Trio phosphorimager. Polygons were drawn around the bands corresponding to the unmodified primer and the reaction products and volumes were quantified using the ImageQuant software (GE Healthcare). Yields of product formation were then determined by application of the following equation:

$$yield = \frac{v_{n+1}}{(v_n + v_{n+1})} \times 100 \quad Eq. (1)$$

Where *v<sub>n</sub>* and *v<sub>n+1</sub>* correspond to the volumes of unmodified primer and of the *n*+1 product, respectively.

### Computational details

Geometry optimizations were performed within a DFT approach, based on the hybrid B3LYP functional<sup>69-70</sup> as implemented in the Gaussian09 package. The correlation-consistent polarized valence double-z-cc-pVDZ<sup>71</sup> basis set was used for all the geometry optimizations in gas phase. Since dispersion interactions might contribute differently to the stability of the base pairs under study, we also added an empirical dispersion term, with the Becke-Johnson damping scheme, to the electronic energy using empirical dispersion=GD3BJ keyword.<sup>72-73</sup> Interaction energies were calculated on the B3LYP-D3BJ/cc-pVDZ optimized geometries at the M06 level<sup>74</sup> of theory using the cc-pVTZ basis set. For the anionic oxygen atom as in Im<sup>C</sup>, an augmented aug-cc-pVTZ basis set is used. The energies have been calculated both in gas phase

and in water, modeled with the C-PCM continuum solvation model.<sup>75</sup> For calculation of the geometries and energies for Ag, we used the small-core, quasi-relativistic Stuttgart/Dresden effective core potential with an associated valence contracted basis set (standard SDD keywords in Gaussian 09).<sup>76-78</sup> Besides the optimizations with no symmetry constraints ( $C_1$  symmetry), we have also imposed  $C_s$  symmetry on the simulated non-natural base pairs in order to mimic the planarity of base pairs under study, situation in the DNA double helical structure.

Thus, the binding energy  $E_{\text{Bind}}$  is calculated as in Eq. 2:

$$E_{\text{Bind}} = E_{\text{Complex}} - (E_{\text{M1}} + E_{\text{M2}} + E_{\text{Metal}}) \quad (2)$$

where  $E_{\text{Complex}}$  is the electronic energies of the optimized complex (that is, M1-M2-metal base pair), and  $E_{\text{M1}}$  and  $E_{\text{M2}}$  and  $E_{\text{Metal}}$  are the electronic energy of the isolated optimized M1 and M2 bases, and respective complexed metal ion ( $\text{Ag}^+$ ). In addition, binding energies of H-bonded base pairs are also calculated. This is a rather standard approach used in this kind of calculations.<sup>79-84</sup>

## SUPPORTING INFORMATION

The supplementary information contains the full characterization data of all compounds and modified oligonucleotides, additional gel images of primer extension reactions, additional geometry optimizations, and the XYZ coordinates for metal base pairs.

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## CONFLICTS OF INTEREST

The authors declare no competing interests.





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