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Temporary electrostatic impairing of DNA recognition. Light-driven DNA-binding of peptide dimers derived from a bZIP transcription factor

Adrián Jiménez-Balsa, Elena Pazos, Borja Martínez-Albaronedo, José L. Mascareñas* and M. Eugenio Vázquez*

Gene expression relies on a myriad of carefully orchestrated interactions between specialized proteins called transcription factors (TFs) and regulatory DNA sequences.^[1] In general, such interactions are subtly regulated in time and space, so that many TFs remain inactive until receiving an appropriate activation signal.^[2] It is well established that the DNA readout by TFs largely relies on interactions between amino acid side chains and the DNA bases and phosphates.^[3] Among these contacts, those involving positively charged basic amino acids are critical for the thermodynamic stability of their DNA complexes.^[4] We reasoned that the temporary electrostatic deactivation of such contacts might provide for the development of TF-based systems whose DNA binding activity could be externally controllable, for instance by light. These systems could be useful tools for transcriptional control,^[5] or for probing spatiotemporal patterns of gene expression in living organisms.^[6] It is curious that despite the well established use of light-activated compounds in chemical biology,^[7,8] examples of photocontrolled DNA binding are certainly scarce. These include reversible switches based on the photostationary equilibrium of azo-modified DNA binders,^[5,9] special chromophores with poor DNA binding affinity and/or specificity,^[10] and single-use caging strategies for triggering minor groove binding or intercalation.^[11] Thus, inspired by the use of negatively-charged elements to modulate cell internalization,^[12] we sought to develop a general strategy for photocontrolling the sequence-specific DNA binding of TF peptide mimics

Herein we demonstrate that tethering polyanionic tails to basic DNA-binding bZIP peptides through a light-sensitive linker suppresses the DNA interaction. Upon irradiation, the negatively charged appendages are released, and the DNA-binding activity is thus restored. As reference system for implementing the strategy we

chose the GCN4 transcription factor, an archetypical bZIP TF that specifically binds to ATF/CREB (5'-ATGA(c/g)TCAT-3') or AP1 (5'-ATGA(c)TCAT-3') sites as a leucine zipper-mediated dimer of uninterrupted α helices. The N-terminal basic regions feature many positively charged amino acids that are key for the DNA recognition (10 Lys or Arg out of 31 residues in the basic region. Figure 1, GCN4br).^[13] It has been shown that the leucine zipper itself can be substituted by a number of dimerizing units without significant loss in the DNA binding properties.^[14] Therefore, in our first iteration for the design of the electrostatically impaired DNA-binding peptides, we selected the minimum sequence of the bZIP basic region (**br**) that it is known to retain the DNA binding ability when engineered as a disulfide dimer.^[15] This minimal peptide was extended at the N-terminus by adding acidic extensions with four or eight Glu residues, linked to the core **br** sequence through a photolabile 3-amino-3-(2-nitrophenyl)-propionic acid (ANP) group.^[16] The disulfide control peptide **br**₂ was constructed by dimerization of the monomers containing a C-terminal cysteine, using Ellman's reagent in phosphate buffer.^[17] Peptide dimers $\phi(\mathbf{br})_2$, $\phi(\mathbf{E}_4\mathbf{br})_2$ and $\phi(\mathbf{E}_8\mathbf{br})_2$ were obtained in good yield by direct alkylation of each of the cystein monomers with 1,4-bis(bromomethyl)benzene.

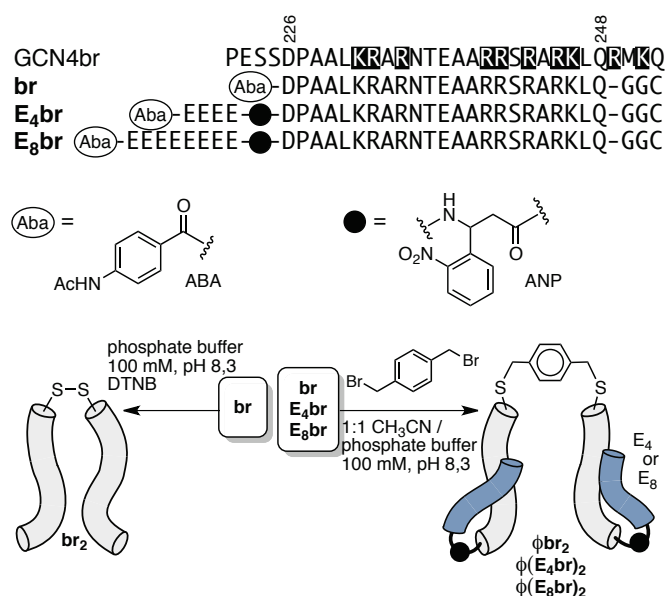


Figure 1. Top: Sequence of the GCN4 basic region highlighting the positively charged residues. Sequences of the acidic tail peptides, **E₄br** and **E₈br**. Structure of the Aba chromophore (used as internal standard), and the ANP photocleavable linker (shown as ●). Bottom: Dimerization reactions of **br**, **E₄br** and **E₈br** to give the corresponding control disulfide **br**₂, or benzylic dimers, $\phi(\mathbf{br})_2$, $\phi(\mathbf{E}_4\mathbf{br})_2$, and $\phi(\mathbf{E}_8\mathbf{br})_2$. The acidic tails (Glu₄ or Glu₈) are represented in the cartoons with darker grey, and the photocleavable element as a black circle.

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The DNA binding of ϕbr_2 , $\phi(\text{E}_4\text{br})_2$ and $\phi(\text{E}_8\text{br})_2$ was studied by electrophoretic mobility shift assays (EMSA) under non-denaturing conditions,^[18] and using SYBR gold for DNA staining. Thus, a short ds-oligonucleotide containing the ATF/CREB binding site was incubated with each of the three peptides at 4 °C. As a positive binding control we used the basic region disulfide dimer br_2 , which in the presence of the ATF/CREB oligo gave the expected slower migrating band (Figure 2, lanes 2-4, band b). The ϕbr_2 dimer displayed qualitatively similar binding properties as the positive control, albeit exhibiting a slightly reduced affinity (figure 2, lanes 5 and 6). Curiously, the dimer $\phi(\text{E}_4\text{br})_2$ also displayed measurable affinity for the target oligonucleotide, despite the presence of a significant number of negatives charges (two Glu₄ tails), as evidenced by the appearance of a retarded band in the gel similar that observed with the controls (figure 2, lanes 7 and 8). In contrast, the $\phi(\text{E}_8\text{br})_2$ peptide, featuring the longer Glu₈ acidic tails, was incapable of forming stable complexes in the electrophoretic gel, and only at high peptide concentrations it was possible to observe a faint, slower migrating band (lanes 9 and 10).

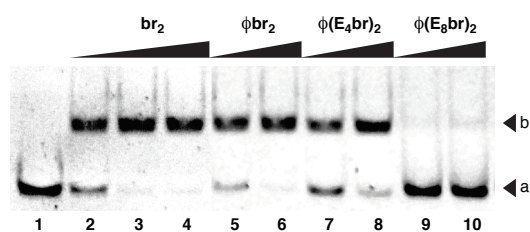
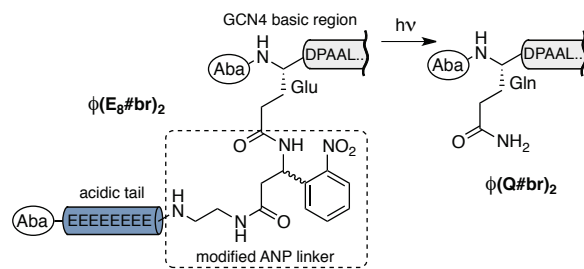


Figure 2. EMSA assays of DNA recognition. Lanes 1-10: target ATF/CREB dsDNA (50 nM). Lanes 2-4: 75, 150, 300 nM br_2 ; lanes 5, 6: 150, 300 nM ϕbr_2 ; lanes 7, 8: 150, 300 nM $\phi(\text{E}_4\text{br})_2$; lanes 9, 10: 150, 300 nM $\phi(\text{E}_8\text{br})_2$. Band **a** corresponds to the free ds-oligo; the slow-migrating band **b** corresponds to the DNA/peptide complexes.^[19] ATF/CREB (one strand shown, binding site in italics): 5'- TGGAG ATGA cg TCAT CTCGT -3'. Peptide and dsDNA in 5 mM Tris-HCl, pH 6.8, 50 mM NaCl, (4°C, 10 min) were added to 18 mM Tris-HCl pH 7.5, 90 mM KCl, 1.8 mM MgCl₂, 1.8 mM EDTA, 9% glycerol, 0.11 mg/mL BSA, 2.25% NP-40 (4°C, 10 min) and loaded into the gel.

Once we confirmed that the Glu₈ acidic appendages significantly impaired the DNA binding, we investigated the photocleavage of the ANP linker. Unfortunately, HPLC analysis of the irradiated solution of $\phi(\text{E}_8\text{br})_2$ (30 sec. $\lambda = 300\text{-}375$ nm) showed a complex mixture of products that could not be characterized (figure 3, left), and from which it was not possible to isolate the expected photocleaved dimer. Addition of commonly used reagents to capture reactive photolysis byproducts, such as DTT or hydrazine,^[20] did not result in any significant improvement. Considering that the complex product mixture could arise from the degradation of highly reactive phenylnitroso ketone groups photoreleased at the N-terminus of the active peptide fragments, we modified the design, reversing the orientation of the ANP linker.^[21] The new $\phi(\text{E}_8\#\text{br})_2$ peptide contained a reconfigured ANP linker attached to the side chain of a Glu residue at the N-terminus of the basic region; cleavage of this inverted ANP would release the intact native peptide leaving a natural Gln residue at the N-terminus ($\phi(\text{Q}\#\text{br})_2$, Scheme 1).

The revamped $\phi(\text{E}_8\#\text{br})_2$ peptide was synthesized following a similar procedure to that described previously for the synthesis of $\phi(\text{E}_8\text{br})_2$. Gratifyingly, we found that photolysis of $\phi(\text{E}_8\#\text{br})_2$ was much cleaner than that of $\phi(\text{E}_8\text{br})_2$, as shown by the HPLC analysis of the reaction mixture (figure 3, and supporting information).



Scheme 1. Structure of the modified photolabile basic regions with the reversed ANP linker connected through a Glu side chain ($\phi(\text{E}_8\#\text{br})_2$), and the expected photodissociation product, $\phi(\text{Q}\#\text{br})_2$.

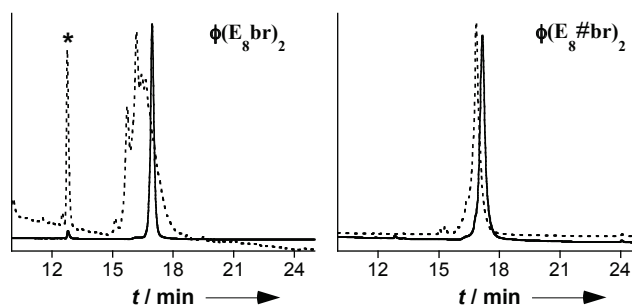


Figure 3. HPLC traces of the irradiated buffered solutions of $\phi(\text{E}_8\text{br})_2$ (left) and $\phi(\text{E}_8\#\text{br})_2$ (right), showing the improved photolysis in the reversed design (5 to 75%, 0.1% TFA CH₃CN/H₂O). Solid lines represent the starting dimers, dashed lines are the chromatograms after 30 sec irradiation. The acidic tail released upon irradiation of $\phi(\text{E}_8\text{br})_2$ is marked with an asterisk. The major peak in the trace at the right corresponds to the expected uncaged dimer $\phi(\text{Q}\#\text{br})_2$. The photoreleased acidic tail containing the reactive nitrosoketone group is not observed in this case, probably because it decomposes and the degraded products are eluted with the injection peak; traces of the dimer with one acidic tail were also observed in the MALDI spectra (supporting information). Y-axis scale in arbitrary absorbance units.

As expected, the inverted peptide $\phi(\text{E}_8\#\text{br})_2$ qualitatively reproduces the DNA-binding behavior observed with the original $\phi(\text{E}_8\text{br})_2$ peptide. Thus incubation of $\phi(\text{E}_8\#\text{br})_2$ with a double stranded oligonucleotide containing the ATF/CREB target sequence did not show significant retarded bands in the PAGE experiments (figure 4, top, lanes 2-5). Irradiation of a 50 μM solution of $\phi(\text{E}_8\#\text{br})_2$ in 10 mM Tris-HCl pH 6.8 buffer, for 30 sec with a standard gel transilluminator lamp, and subsequent incubation with the same oligonucleotide, resulted in the appearance of retarded gel bands, consistent with a specific peptide-DNA complex (figure 4, top, lanes 6-9).^[22] Moreover, these bands were similar to those observed using the $\phi(\text{Q}\#\text{br})_2$ peptide dimer, which was purposely synthesized *de novo* as a true uncaged control. As expected, incubation of the irradiated sample of $\phi(\text{E}_8\#\text{br})_2$ with a random DNA did not result in any new band (figure 4, top, lane 10). In addition to the experiments with the symmetrically caged $\phi(\text{E}_8\#\text{br})_2$, we synthesized a single-caged $\phi(\text{Q}\#\text{br})(\text{E}_8\#\text{br})$ peptide, in which only one of the basic regions was modified with the acidic tail. Remarkably, this peptide retained a significant DNA binding affinity, showing in the EMSA gels as a slightly slower migrating band than that of the complex with the uncaged control $\phi(\text{Q}\#\text{br})_2$ (figure S17, supporting information).

Importantly, the photochemical activation can be carried out in the presence of the DNA. Therefore, irradiating a mixture of

$\phi(\text{E}_8\#\text{br})_2$ and the target oligonucleotide allowed a substantial recovery of the DNA binding, as shown by EMSA (figure 4, bottom, lanes 3-5). An irradiation control experiment with a random oligonucleotide confirmed the specificity of the interaction, as in this case we did not observe the formation of new retarded bands in the gel (figure 4, bottom, lanes 8-10).

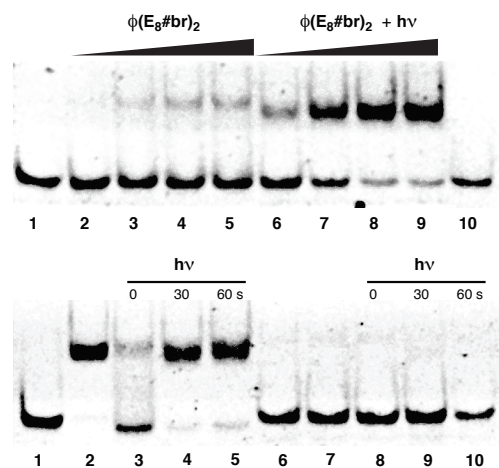


Figure 4. EMSA analysis of DNA binding by $\phi(\text{E}_8\#\text{br})_2$ at 4 °C. **Top:** Lanes 1-9: target ATF/CREB oligo (50 nM). Lanes 2-5: 125, 250, 400, 600 nM $\phi(\text{E}_8\#\text{br})_2$, no irradiation; lanes 6-9: 125, 250, 400, 600 nM $\phi(\text{E}_8\#\text{br})_2$, after 30 sec. irradiation with UV light (10 mM Tris-HCl, pH 6.8); lane 10: 600 nM $\phi(\text{E}_8\#\text{br})_2$ after 30 sec. irradiation and addition of a random oligo (*rndDNA*, 50 nM, see supporting information for full sequence). **Bottom:** DNA binding of $\phi(\text{E}_8\#\text{br})_2$; irradiation in the presence of the oligonucleotide (10 mM Tris-HCl, pH 6.8). Lanes 1-5: target ATF/CREB oligo (50 nM). Lane 2: control peptide ϕbr_2 (300 nM). Lanes 3-5: $\phi(\text{E}_8\#\text{br})_2$ (300 nM) at increased irradiation times; lanes 6-10: *rndDNA* (50 nM); lane 7: control, ϕbr_2 (300 nM); lanes 8-10: $\phi(\text{E}_8\#\text{br})_2$ (300 nM) at increasing irradiation times.

It is well known that the DNA binding of bZIP dimers is coupled to their folding into α -helical conformations.^[23] Therefore we made complementary circular dichroism studies to gain some structural insight into the DNA binding process. These experiments showed that the caged dimer $\phi(\text{E}_8\#\text{br})_2$ is essentially unstructured in the absence of DNA (figure 5, left, dashed line). Addition of the target ATF/CREB oligonucleotide induced a significant increase in the negative CD signal at 222 nm, consistent with an increase in the α -helical content of the peptide that could result from the high concentrations required to run the CD experiments (5 μM of peptide and DNA), or from non-specific interactions with the dsDNAs.^[24] More importantly, UV irradiation of this $\phi(\text{E}_8\#\text{br})_2$ /oligonucleotide mixture led to a large increase in the negative ellipticity, in the range of what has been observed when this type of dimers fold upon interacting with their target DNA sites (figure 5, left).^[23]

In order to further confirm these results and rule out any distortions associated with the gel shift experiments, we performed fluorescence anisotropy titrations that provided a more quantitative characterization of the DNA recognition process.^[25] Incubation of a fluorescein-labeled ATF/CREB oligonucleotide with increasing amounts of $\phi(\text{Q}\#\text{br})_2$ resulted in a progressive increase of the anisotropy, which was consistent with the formation of a DNA complex with higher molecular weight and reduced mobility.^[26] Fitting of the resulting isotherm to a 1:1 binding model,^[27] allowed to calculate its dissociation constant ($K_D \approx 38$ nM). As expected, the caged derivative $\phi(\text{E}_8\#\text{br})_2$ showed drastically weaker binding than

the control peptide under the same conditions ($K_D \approx 1$ μM , figure 5, right). Moreover, the uncaging process could be monitored by fluorescence anisotropy. Thus, irradiation of the caged $\phi(\text{E}_8\#\text{br})_2$ peptide in the presence of the fluorescently-labeled ATF/CREB oligonucleotide resulted in an anisotropy value similar to that obtained when the target oligonucleotide is incubated with the $\phi(\text{Q}\#\text{br})_2$ control peptide (bar graph, figure 5, right).

In conclusion, we have demonstrated that the attachment of a negatively charged tether to an Arg/Lys-rich bZIP based peptide can effectively hamper its DNA recognition ability. Moreover, connecting this appendage through a photocleavable ANP linker allows to restoring the DNA binding upon simple UV irradiation. The negatively charged patch might disturb the binding both, by interfering with electrostatic pairing, or by generating repulsive contacts with the DNA. Typical photocaging or switching strategies are based on the modification of specific key residues, and usually require linear synthetic processes as well as detailed structural knowledge of the interaction. Our “*electrostatic turn-off*” strategy, by relying on simple tethering of highly charged appendages to the natural recognition elements, should provide a facile, versatile and general route to temporarily control specific biological interactions involving highly charged partners such as nucleic acids. Furthermore, this method seems particularly appropriate for modulating interactions involving multivalent contacts that might tolerate a single caging group.

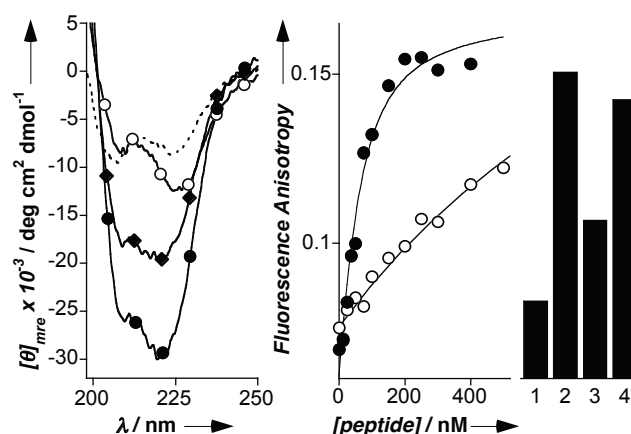


Figure 5. **Left:** CD spectra of 5 μM $\phi(\text{E}_8\#\text{br})_2$ in 10 mM Tris-HCl, pH 6.8, at 4 °C (dashed line). Solid lines: $\phi(\text{E}_8\#\text{br})_2$ after 30 sec irradiation (○); $\phi(\text{E}_8\#\text{br})_2$ in the presence of 5 μM of the target oligo ATF/CREB before (◊), and after 30 sec. irradiation (●). **Center:** Fluorescence anisotropy titrations at 518 nm of a fluorescein-labeled ATF/CREB (50 nM) with $\phi(\text{E}_8\#\text{br})_2$ (○), with the $\phi(\text{Q}\#\text{br})_2$ control (●), and best fit to a simple 1:1 binding model (10 mM Tris-HCl buffer, 10 mM NaCl, pH 6.8 at 10 °C). **Right:** fluorescence anisotropy values: oligo ATF/CREB (50 nM) (bar 1); oligo with 300 nM $\phi(\text{Q}\#\text{br})_2$ (bar 2); same oligo with 300 nM $\phi(\text{E}_8\#\text{br})_2$ (bar 3), and with 300 nM $\phi(\text{E}_8\#\text{br})_2$ after 60 sec irradiation (bar 4). Same scale as the titrations.

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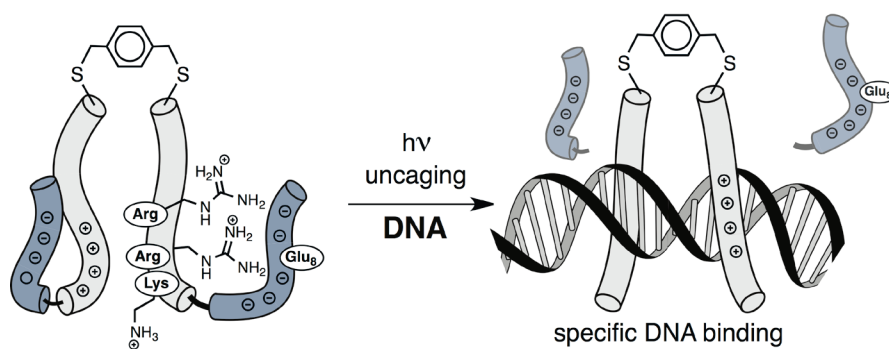
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(Electrostatic caging)

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Temporary electrostatic impairing of
DNA recognition. Light driven DNA-
binding of peptide dimers derived from a
bZIP transcription factor



Appending negatively charged Glu₈ tails to a peptide dimer derived from the GCN4 transcription factor leads to an effective suppression of its DNA binding. The specific DNA recognition can be restored by irradiation with UV light, owing to the use of a photolabile linker between the acidic tail and the DNA binding peptide.