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DNA-guided DNA cleavage at moderate temperatures by *Clostridium butyricum* Argonaute

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ABSTRACT

Prokaryotic Argonaute proteins (pAgos) constitute a diverse group of endonucleases of which some mediate host defense by utilizing small interfering DNA guides (siDNA) to cleave complementary invading DNA. This activity can be repurposed for programmable DNA cleavage. However, currently characterized DNA-cleaving pAgos require elevated temperatures ($\geq 65^\circ\text{C}$) for their activity, making them less suitable for applications that require moderate temperatures, such as genome editing. Here, we report the functional and structural characterization of the siDNA-guided DNA-targeting pAgo from the mesophilic bacterium *Clostridium butyricum* (CbAgo). CbAgo displays a preference for siDNAs that have a deoxyadenosine at the 5'-end and thymidines at nucleotides 2–4. Furthermore, CbAgo mediates DNA-guided DNA cleavage of AT-rich double stranded DNA at moderate temperatures (37°C). This study demonstrates that certain pAgos are capable of programmable DNA cleavage at moderate temperatures and thereby expands the scope of the potential pAgo-based applications.

INTRODUCTION

Eukaryotic Argonaute proteins (eAgos) play a key role in RNA interference (RNAi) processes (1–3). As the core of the multiprotein RNA-induced silencing complex (RISC), eAgos bind small non-coding RNA molecules as guides to direct the RISC complex towards complementary RNA targets (3–5). Reflecting their physiological function, variation

among eAgos is observed with respect to the presence or absence of a catalytic site, and to their potential to interact with other proteins (6). Depending on the eAgo and on the sequence complementarity between guide and target RNA, eAgo-guide complexes either catalyze endonucleolytic cleavage of the target RNA (7) or indirectly silence the target RNA by repressing its translation and promoting its degradation through recruitment of additional silencing factors (8). Independent of the mechanism, eAgo-mediated RNA binding generally results in sequence-specific silencing of gene expression. As such, eAgos can coordinate various cellular processes by regulating intracellular RNA levels.

Prokaryotes also encode Argonaute proteins (pAgos) (9,10). Various pAgos share a high degree of structural homology with eAgos as both pAgos and eAgos adopt the same four domain (N-PAZ-MID-PIWI) architecture (9–12). Despite their structural homology, several recently characterized pAgos have distinct functional roles and different guide and/or target preferences compared to eAgos. For example, several pAgos have been implicated in host defense by directly targeting DNA instead of RNA (13–16). One of the best characterized mechanisms that pAgos utilize is DNA-guided DNA interference, which is demonstrated for pAgos from *Thermus thermophilus* (TtAgo), *Pyrococcus furiosus* (PfAgo) and *Methanocaldococcus jannaschii* (MjAgo) (13–15,17–20). These pAgos use 5'-end phosphorylated small interfering DNAs (siDNAs) for recognition and successive cleavage of complementary DNA targets. This mechanism enables both TtAgo and PfAgo to mediate host defense against invading nucleic acids. Prokaryotes lack homologs of eukaryotic enzymes that are involved in guide biogenesis (21). Instead, both TtAgo and MjAgo—besides the canonical siDNA-

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dependent target cleavage termed ‘slicing’—exhibit an alternative nuclease activity termed ‘chopping’ (14,17). Chopping facilitates autonomous generation of small DNA fragments from dsDNA substrates. Subsequently, these DNA fragments generated during chopping can serve as siDNAs for canonical slicing (14,17).

TtAgo and *PfAgo* can be programmed with short synthetic siDNA which allows them to target and cleave dsDNA sequences of choice *in vitro* (13,15). This activity has enabled the repurposing of *PfAgo* as an universal restriction endonuclease for *in vitro* molecular cloning (22). In addition, a diagnostic *TtAgo*-based application termed NAVIGATER (Nucleic Acid enrichment Via DNA Guided Argonaute from *T. thermophilus*) was developed which enables enhanced detection of rare nucleic acids with single nucleotide precision (23). In analogy with the now commonly used CRISPR-Cas9 and CRISPR-Cas12a enzymes (24–26), it has also been suggested that pAgos could be repurposed as next-generation genome editing tools (27). However, due to the thermophilic nature (optimum activity temperature $\geq 65^\circ\text{C}$) and low levels of endonuclease activity at the relevant temperatures (20–37°C), it is unlikely that the well-studied *TtAgo*, *PfAgo* and *MjAgo* are suitable for genome editing. The quest for a pAgo that can cleave dsDNA at moderate temperatures has resulted in the characterization of the Argonaute protein from *Natronobacterium gregory* (*NgAgo*), which was claimed to be the first pAgo suitable for genome editing purposes (28). However, the study reporting this application has been retracted after a series of reproducibility issues (28–30). Instead, it has been suggested that *NgAgo* targets RNA rather than DNA (31).

Although considerable efforts have been made to elucidate the mechanisms and biological roles of pAgos, efforts have mainly focused on pAgo variants from (hyper)thermophiles. This has left a large group of mesophilic pAgos unexplored. We here report the characterization of the Argonaute protein from the mesophilic bacterium *Clostridium butyricum* (*CbAgo*). We demonstrate that *CbAgo* can utilize siDNA guides to cleave both ssDNA and dsDNA targets at moderate temperatures (37°C). In addition, we have elucidated the macromolecular structure of *CbAgo* in complex with a siDNA guide and complementary ssDNA target in a catalytically competent state. *CbAgo* displays an unusual preference for siDNAs with a deoxyadenosine at the 5'-end and thymidines at nucleotides 2–4. The programmable DNA endonuclease activity of *CbAgo* provides a foundation for the development of pAgo-based applications at moderate temperatures.

MATERIALS AND METHODS

Plasmid construction

The *CbAgo* gene was codon harmonized for *Escherichia coli* BL21 (DE3) and inserted into a pET-His6 MBP TEV cloning vector (obtained from the UC Berkeley MacroLab, Addgene #29656) using ligation independent cloning (LIC) using oligonucleotides oDS067 and oDS068 (Supplementary Table S1) to generate a protein expression construct that encodes the *CbAgo* polypeptide sequence fused to an N-terminal tag comprising a hexahistidine sequence,

a maltose binding protein (MBP) and a Tobacco Etch Virus (TEV) protease cleavage site.

Generation of the double mutant

CbAgo double mutant (D541A, D611A) was generated using an adapted Quick Directed Mutagenesis Kit instruction manual (Stratagene). The primers were designed using the web-based program primerX (<http://bioinformatics.org/primerx>).

CbAgo expression and purification

The *CbAgo* WT and DM proteins were expressed in *E. coli* BL21(DE3) Rosetta™ 2 (Novagen). Cultures were grown at 37°C in LB medium containing 50 $\mu\text{g ml}^{-1}$ kanamycin and 34 $\mu\text{g ml}^{-1}$ chloramphenicol until an OD_{600 nm} of 0.7 was reached. *CbAgo* expression was induced by addition of isopropyl β -D-1-thiogalactopyranoside (IPTG) to a final concentration of 0.1 mM. During the expression cells were incubated at 18°C for 16 h with continuous shaking. Cells were harvested by centrifugation and lysed by sonication (Bandelin, Sonopuls, 30% power, 1 s on/2 s off for 5 min) in lysis buffer containing 20 mM Tris–HCl pH 7.5, 250 mM NaCl, 5 mM imidazole, supplemented with a EDTA free protease inhibitor cocktail tablet (Roche). The soluble fraction of the lysate was loaded on a nickel column (HisTrap Hp, GE Healthcare). The column was extensively washed with wash buffer containing 20 mM Tris–HCl pH 7.5, 250 mM NaCl and 30 mM imidazole. Bound protein was eluted by increasing the concentration of imidazole in the wash buffer to 250 mM. The eluted protein was dialysed at 4°C overnight against 20 mM HEPES pH 7.5, 250 mM KCl and 1 mM dithiothreitol (DTT) in the presence of 1 mg TEV protease (32) to cleave of the His6-MBP tag. Next, the cleaved protein was diluted in 20 mM HEPES pH 7.5 to lower the final salt concentration to 125 mM KCl. The diluted protein was applied to a heparin column (HiTrap Heparin HP, GE Healthcare), washed with 20 mM HEPES pH 7.5, 125 mM KCl and eluted with a linear gradient of 0.125–2 M KCl. Next, the eluted protein was loaded onto a size exclusion column (Superdex 200 16/600 column, GE Healthcare) and eluted with 20 mM HEPES pH 7.5, 500 mM KCl and 1 mM DTT. Purified *CbAgo* protein was diluted in size exclusion buffer to a final concentration of 5 μM . Aliquots were flash frozen in liquid nitrogen and stored at -80°C .

Co-purification nucleic acids

To 500 pmol of purified *CbAgo* in SEC buffer CaCl₂ and proteinase K (Ambion) were added to final concentrations of 5 mM CaCl₂ and 250 $\mu\text{g/ml}$ proteinase K. The sample was incubated for 4 h at 65°C. The nucleic acids were separated from the organic fraction by adding Roti phenol/chloroform/isoamyl alcohol pH 7.5–8.0 in a 1:1 ratio. The top layer was isolated and nucleic acids were precipitated using ethanol precipitation by adding 99% ethanol in a 1:2 ratio supplied with 0.5% linear polymerized acrylamide as a carrier. This mixture was incubated overnight at -20°C and centrifuged in a table centrifuge at 16 000 g for 30 min. Next, the nucleic acids pellet was washed with

70% ethanol and solved in 50 μ l MilliQ water. The purified nucleic acids were treated with either 100 μ g/ml RNase A (Thermo), 2 units DNase I (NEB) or both for 1 h at 37°C and resolved on a denaturing urea polyacrylamide gel (15%) and stained with SYBR gold.

Single stranded activity assays

Unless stated otherwise 5 pmol of each *CbAgo*, siDNA and target were mixed in a ratio of 1:1:1, in 2 \times reaction buffer containing 20 mM Tris-HCl pH 7.5 supplemented with 500 μ M MnCl₂. The target was added after the *CbAgo* and siDNA had been incubated for 15 min at 37°C. Then the complete reaction mixture was incubated for 1 h at 37°C. The reaction was terminated by adding 2 \times RNA loading dye (95% formamide, 0.025% bromophenol blue, 5 mM EDTA) and heating it for 5 min at 95°C. After this, the samples were resolved on a 20% denaturing (7 M urea) polyacrylamide gel. The gel was stained with SYBR gold nucleic acid stain (Invitrogen) and imaged using a G:BOX Chemi imager (Syngene).

Double stranded activity assay

In two half reactions 12.5 pmol of *CbAgo* was loaded with either 12.5 pmol of forward or reverse siDNA in reaction buffer containing 10 mM Tris-HCl pH 7.5, 10 μ g/ml BSA and 250 μ M MnCl₂. The half reactions were incubated for 15 min at 37°C. Next, both half reactions were mixed together and 120 ng target plasmid was added after which the mixture was incubated for 1 h of 37°C. After the incubation the target plasmid was purified from the mixture using a DNA clean and concentrate kit (DNA Clean & Concentrator™-5, Zymogen) via the supplied protocol. The purified plasmid was subsequently cut using either EcoRI-HF (NEB) or SapI-HF (NEB) in Cutsmart buffer (NEB) for 30 min at 37°C. A 6 \times DNA loading dye (NEB) was added to the plasmid sample prior to resolving it on a 0.7% agarose gel stained with SYBR gold (Invitrogen). In the double stranded assays in which linear plasmid was targeted, 300 ng pUC IDT, linearized by EcoRI, was added to both half reactions and incubated for 1 h of 37°C. The reaction products were resolved on a 0.7% agarose gel.

Crystallization

To reconstitute the *CbAgo* DM-siDNA-target DNA complex, siDNA and target DNA were pre-mixed at a 1:1 ratio, heated to 95°C, and slowly cooled to room temperature. The formed dsDNA duplex (0.5 M) was mixed with *CbAgo* DM in SEC buffer at a 1:1:4 ratio (*CbAgo* DM:duplex DNA), and MgCl₂ was added to a final concentration of 5 mM. The sample was incubated for 15 min at 20°C to allow complex formation. The complex was crystallized at 20°C using the hanging drop vapour diffusion method by mixing equal volumes of complex and reservoir solution. Initial crystals were obtained at a *CbAgo* DM concentration of 5 mg/ml with a reservoir solution consisting of 4 M sodium formate. Data was collected from crystals grown obtained using a complex concentration of 4.3 mg/ml and reservoir solution containing 3.8 M Sodium Formate and 5 mM NiCl₂ at 20°C. For

cryoprotection, crystals were transferred to a drop of reservoir solution and flash-cooled in liquid nitrogen.

X-ray diffraction data were measured at beamline X06DA (PXIII) of the Swiss Light Source (Paul Scherrer Institute, Villigen, Switzerland). Data were indexed, integrated and scaled using AutoPROC (33). Crystals of the *CbAgo*-siDNA-target DNA complex diffracted to a resolution of 3.55 Å and belonged to space group *P*6₃ 2 2, with one copy of the complex in the asymmetric unit. The structure was solved by molecular replacement using Phaser-MR (34). As search model, the structure of *TtAgo* in complex with guide and target DNA strands (PDB: 5GQ9) was used after removing loops and truncating amino acid side chains. Phases obtained using the initial molecular replacement solution were improved by density modification using phenix.resolve (35) and phenix.morph_model (36). The atomic model was built manually in Coot (37) and refined using phenix.refine (37). The final binary complex model contains *CbAgo* residues 1-463 and 466-748, guide DNA residues 1-16, and target DNA residues (-18)-(-1).

Structure analysis

Core root means square deviations (rmsd) of structure alignments were calculated using Coot SSM superpose (38). Hydrogen bonding interactions were analysed using PDBePISA (39), while other interactions with the Mg²⁺ ion were determined manually in Coot (37). Figures were generated using PyMOL (Schrödinger).

Single-molecule experimental set-up

Single-molecule fluorescence FRET measurements were performed with a prism-type total internal reflection fluorescence microscope. Cy3 and Cy5 molecules were excited with 532 and 637 nm wavelength, respectively. Resulting Cy3 and Cy5 fluorescence signal was collected through a 60 \times water immersion objective (UplanSApo, Olympus) with an inverted microscope (IX73, Olympus) and split by a dichroic mirror (635dcxr, Chroma). Scattered laser light was blocked out by a triple notch filter (NF01-488/532/635, Semrock). The Cy3 and Cy5 signals were recorded using a EM-CCD camera (iXon Ultra, DU-897U-CS0-#BV, Andor Technology) with exposure time 0.1 s. All single-molecule experiments were done at room temperature (22 \pm 2°C).

Fluorescent DNA and RNA preparation

The RNAs with amine-modification (amino-modifier C6-U phosphoramidite, 10-3039, Glen Research) were purchased from STPharm (South Korea) and DNAs with amine-modification (internal amino modifier iAmMC6T) Ella biotech (Germany). The guide and target strands were labeled with donor (Cy3) and acceptor (Cy5), respectively, using the NHS-ester form of Cy dyes (GE Healthcare). 1 μ l of 1 mM of DNA/RNA dissolved in MilliQ H₂O was added to 5 μ l labeling buffer of (freshly prepared) sodium tetraborate (380 mg/10ml, pH 8.5). 1 μ l of 20 mM dye (1 mg in 56 μ l DMSO) was added and incubated overnight at room temperature in the dark, followed by washing and ethanol precipitation. The labeling efficiency was ~100%.

Single-molecule sample preparation

A microfluidic chamber was incubated with 20 μl Streptavidin (0.1 mg/ml, Sigma) for 30 s. Unbound Streptavidin was washed with 100 μl of buffer T50 (10 mM Tris-HCl pH 8.0, 50 mM NaCl buffer). The 50 μl of 50 pM acceptor-labeled target construct were introduced into the chamber and incubated for 1 min. Unbound labeled constructs were washed with 100 μl of buffer T50. The *CbAgo* binary complex was formed by incubating 10 nM purified *CbAgo* with 1 nM of donor-labeled guide in a buffer containing 50 mM Tris-HCl pH 8.0 (Ambion), 1 mM MnCl_2 and 100 mM NaCl (Ambion) at 37°C for 20 min. For binding rate (k_{on}) measurements, the binary complex was introduced into the fluidics chamber using syringe during the measurement. The experiments were performed at the room temperature ($23 \pm 1^\circ\text{C}$).

For fluorescence Guide Loading Experiments before immobilizing *CbAgo* on the single-molecule surface, 1 μl of 5 μM His-tagged apo-*CbAgo* was incubated with 1 μl of 1 $\mu\text{g/ml}$ biotinylated anti-6x His antibody (Abcam) for 10 min. Afterward, the mixture was diluted 500 \times in T50 and 50 μl were loaded in the microfluidic channel for 30 s incubation, followed by washing with 100 μl of T50 buffer. Cy3-labeled ssDNA (0.1 nM) was applied to the microfluidic chamber in imaging buffer (50 mM Tris-HCl pH 8.0, 100 mM NaCl, 1 mM MnCl_2 , 1 mM Trolox ((\pm)-6-hydroxy-2,5,7,8-tetramethylchromane-2-carboxylic acid), supplemented with an oxygen-scavenging system (0.5 mg/ml glucose oxidase (Sigma), 85 mg/ml catalase (Merck) and 0.8% (v/v) glucose (Sigma)).

Single-molecule data acquisition and analysis

CCD images of time resolution 0.1 or 0.3 s were recorded, and time traces were extracted from the CCD image series using IDL (ITT Visual Information Solution). Colocalization between Cy3 and Cy5 signals was carried out with a custom-made mapping algorithm written in IDL. The extracted time traces were processed using Matlab (MathWorks) and Origin (Origin Lab).

The binding rate (k_{on}) was determined by first measuring the time between when *CbAgo* binary complex was introduced to a microfluidic chamber and when the first *CbAgo*-guide docked to a target; and then fitting the time distribution with a single-exponential growth curve, $A(1 - e^{-k_{\text{on}}t})$. The dissociation rate was estimated by measuring the dwell time of a binding event. A dwell time distribution was fitted by single-exponential decay curve ($Ae^{-t/\Delta\tau}$).

Fluorescence competition experiments

MBP-tagged *CbAgo* was immobilized on the quartz surface using an anti-MBP antibody. An equimolar mixture of let7 DNA guide (Cy3 labeled) and let7 RNA guide (Cy5 labeled) in imaging buffer was introduced to the microfluidic chamber. After 5 min, 10 snapshots of independent fields of view with simultaneous illumination were collected to estimate the amount of guide molecules bound to protein. Movies were taken for 200 s (2000 frames) at continuous illumination of Cy3 and Cy5 molecules to determine the dwell

times of the binding events. Dwell times were binned in a histogram and fitted with a single exponential decay curve.

FRET targeting experiments of ATTT and AAAA guide target combinations

100 pM of target construct annealed with biotin handle were flushed in the microfluidic chamber. After incubation of 1 min, the microfluidic chamber was rinsed with 100 μl T50 buffer. 10 nM of apo-*CbAgo* was loaded with 1 nM of ATTT seed DNA guide or with AAAA seed DNA guide at 37°C for 30 min in imaging buffer after which the mixture is introduced inside the microfluidic chamber. Movies of 200 s were taken at continuous illumination of the Cy3 signal. Site specific protein target interactions were identified as FRET signals and were further analyzed.

RESULTS

CbAgo mediates siDNA-guided ssDNA cleavage

CbAgo was successfully expressed in *E. coli* from a codon-optimized gene using a T7-based pET expression system and purified (Supplementary Figure S1A). To determine the guide and target binding characteristics of *CbAgo*, we performed single-molecule experiments using Förster resonance energy transfer (FRET). We immobilized either Cy5-labeled single stranded RNA or DNA targets (FRET acceptor) on a polymer-coated quartz surface (Figure 1A). Next, we introduced *CbAgo* in complex with either a Cy3-labeled siRNA or siDNA guide (FRET donor) and recorded the interactions. Strikingly, *CbAgo* could utilize both siRNAs and siDNAs to bind DNA or RNA targets (Figure 1B). To test which guide is preferentially bound by *CbAgo* we performed a competition assay in which *CbAgo* was immobilized into the microfluidic chamber, and an equimolar mixture of siDNA and siRNAs was introduced. While only short-lived interactions (average dwell time: 0.48 s) were observed for siRNA, siDNA was strongly bound (average dwell time: 44 s) by *CbAgo* (Figure 1C). This results suggests that *CbAgo* utilizes siDNA rather than siRNA as a guide.

CbAgo is phylogenetically closest related to the clade of halobacterial pAgos, among which also pAgo from *Natronobacterium gregoryi* (*NgAgo*) can be found (Figure 1D and Supplementary Figure S2). A multiple sequence alignment of *CbAgo* with other pAgos (Supplementary Figure S1B) suggests that *CbAgo* contains the conserved DEDX catalytic residues (where X can be a D, H or N) which are essential for nuclease activity in 'slicing' Agos (40). In the case of *CbAgo*, this concerns residues D541, E577, D611 and D727.

To confirm whether *CbAgo* indeed is an active nuclease, we performed *in vitro* activity assays in which *CbAgo* was loaded with either synthetic siDNAs or siRNAs (21 nucleotides in length). Next the complexes were incubated at 37°C with 45-nucleotide complementary single stranded RNA or DNA target oligonucleotides. While no activity was found in any of the combinations in which siRNAs or target RNAs were used, *CbAgo* was able to cleave target DNAs in a siDNA-dependent manner (Figure 1E). In

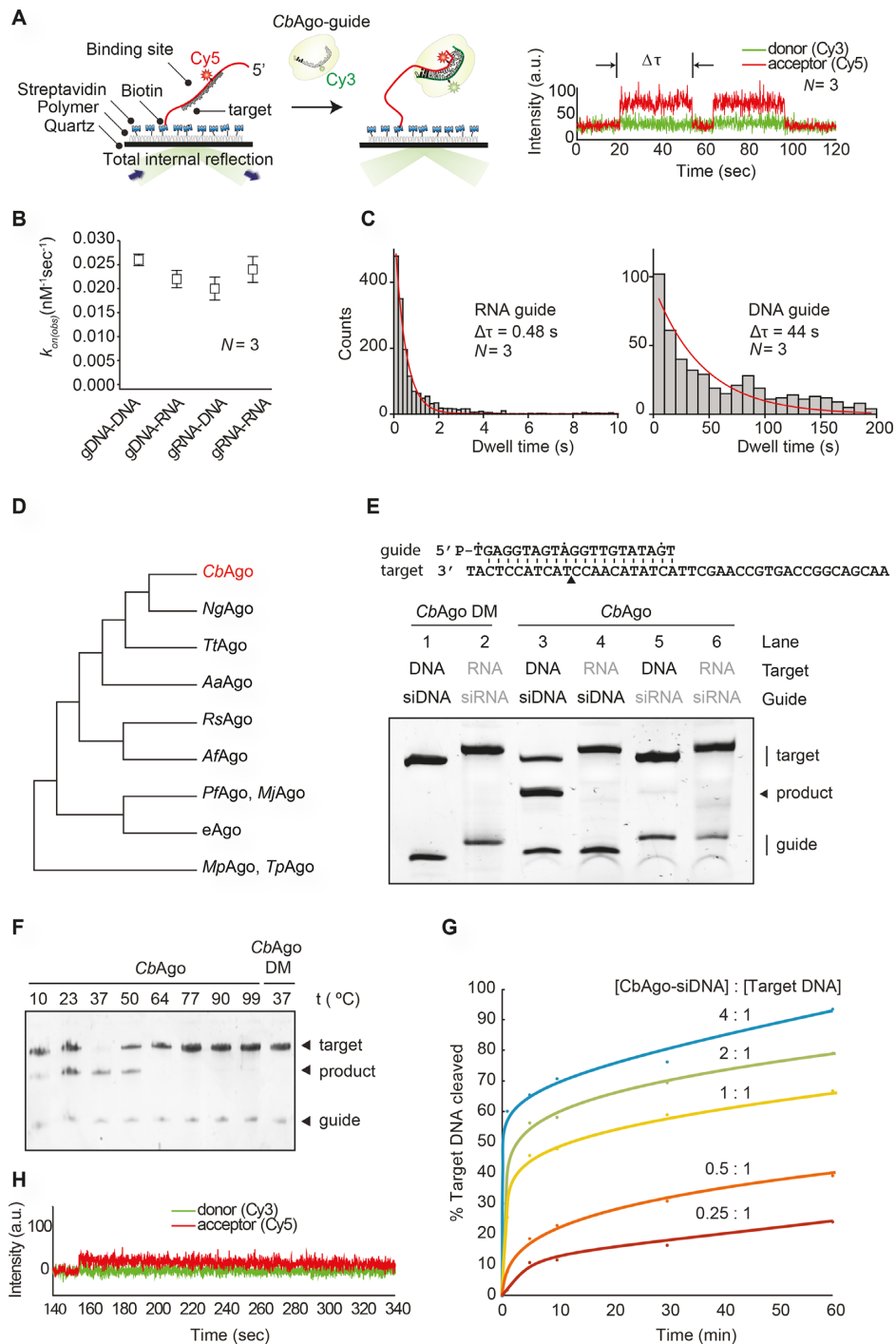


Figure 1. *CbAgo* exhibits DNA-guided DNA endonuclease activity at 37°C. (A) Left: Overview of the single molecule assay to determine the binding characteristics of *CbAgo*. Right: FRET diagram of a *CbAgo*-siDNA complex that has three complementary base pairs (2–4nt) to the DNA target. Indicated is the dwell time ($\Delta\tau$). (B) Comparison of the binding rates (k_{on}) of *CbAgo* in complex with siDNA or siRNA to bind DNA or RNA targets. The rates are similar for each nucleic acid type guide and target. N is the number of base paired nucleotides. (C) Dwell time histograms showing *CbAgo* preferentially binds siDNAs in siDNA-siRNA competition experiments. (D) Schematic phylogenetic tree of characterized pAgos. (E) *CbAgo* exhibits DNA-guided DNA endonuclease activity. Upper panel: Sequence of the synthetic let7 miRNA-based siDNA guide and target DNA sequences that were used for the *in vitro* cleavage assays. Lower panel: *CbAgo*, guides and targets were mixed in a 1:1:1 molar ratio and incubated for 1 h at 37°C. Catalytic mutant *CbAgoDM* was used as a control. Cleavage products were analysed by denaturing polyacrylamide electrophoresis. (F) *CbAgo* displays highest activity at 37°C. *CbAgo* and siDNA were mixed and pre-incubated at various temperatures for 10 min. Next, target DNA was added and the sample was incubated for 1 h at the same temperature. *CbAgoDM* was used as a control. Cleavage products were analysed by denaturing polyacrylamide electrophoresis. (G) Quantified data of a *CbAgo*-mediated siDNA-guided ssDNA cleavage turnover experiment using 5 pmol target DNA and increasing concentrations of *CbAgo*-siDNA (1.25–20 pmol). (H) FRET diagram showing that a cleavage compatible *CbAgo*-siDNA remains bound to a fully complementary target DNA ($N = 21$) during the entire the measurement (340 seconds).

agreement with the predicted DEDD catalytic site (Supplementary Figure S1B), alanine substitutions of two of aspartic acids (D541A, D611A) in the expected catalytic tetrad abolished the nuclease activity, demonstrating that the observed siDNA-guided ssDNA endonucleolytic activity was indeed catalyzed by the DEDD catalytic site. To further investigate the full temperature range at which *CbAgo* is active, we performed additional cleavage assays at temperatures ranging from 10 to 95°C. While *CbAgo* displayed the highest activity at its physiologically relevant temperature (37°C), *CbAgo* also catalyzed siDNA-guided target DNA cleavage at temperatures as low as 10°C and as high as 50°C (Figure 1F).

When *CbAgo*–siDNA complexes and target ssDNA substrates (45 nt) were mixed in equimolar amounts, cleavage of the target DNA was not complete after one hour incubation (Figure 1E). Therefore, we investigated the substrate turnover kinetics of *CbAgo* by monitoring the cleavage assays in a time course using variable *CbAgo*:siDNA:target DNA ratios (Figure 1G). A rapid burst of activity was observed during the first minute, likely indicating the first target binding and cleavage event. This stage was followed by a slow steady state, suggesting that under these conditions the *CbAgo*–siDNA complex slowly dissociates from the cleaved target DNA product before being able to bind and cleave a new target DNA strand. The cleavage kinetics were confirmed using single-molecule assays which demonstrated that the *CbAgo*–siDNA complex remains bound to the DNA target ($N = 21$) for several minutes (Figure 1H), which prevents *CbAgo*–siDNA complexes from binding and cleaving new DNA targets. Thus, while *CbAgo* functions as a multi-turnover nuclease enzyme, its steady-state rate is limited by product release.

Structure of *CbAgo* in the cleavage-competent conformation

To investigate the molecular architecture of *CbAgo* in light of its biochemical activity, we crystallized *CbAgo*DM in complex with both a 21 nt siDNA and a 19 nt DNA target, and solved the structure of the complex at 3.54 Å resolution (Figure 2 and Supplementary Table S1). Like other Agos, *CbAgo* adopts a bilobed conformation in which one of the lobes comprises the N-terminal, linker L1, and PAZ domains, which are linked by linker L2 to the other lobe comprising the MID and PIWI domains. Nucleotides 2–16 of the siDNA constitute a 15 base-pairs A-form-like duplex with the target DNA (Figure 2A). The 5'-terminal nucleotide of the siDNA is anchored in the MID domain pocket, where the 5'-phosphate group of the siDNA makes numerous interactions with MID domain residues and the C-terminal carboxyl group of *CbAgo* (Supplementary Figure S3), as observed for in structures of *TtAgo* and *MjAgo* bound to siDNAs (12,19,41). To test whether the interactions with the 5'-phosphate group of the siDNA are important for *CbAgo* activity, we performed target DNA cleavage assays in which we used siDNAs with a 5' phosphate or a 5' hydroxyl group (Supplementary Figure S4). As observed for other pAgos (42,43), *CbAgo* is able to utilize both siDNAs for target DNA cleavage, but it cleaves target DNA much more efficiently when the siDNA contains a 5'-phosphate

group. This is in agreement with the siDNA–protein interactions predicted from the crystal structure. Furthermore, the backbone phosphates of the siDNA seed segment appear to form hydrogen-bonding and ionic interactions with specific residues in the MID, PIWI and L1 domains (Supplementary Figure S3). At the distal end of the siDNA–target DNA duplex, the N-domain residue His35 caps the duplex by stacking onto the last base pair. After this point, the remaining 3'-terminal nucleotides of the siDNA are disordered, while the target DNA bends away from the duplex and enters the cleft between the N-terminal and PAZ domains. In agreement with other ternary pAgo complexes (18,44,45), the PAZ domain pocket, which normally binds the 3' end of the guide in a binary Ago–guide complex, is empty.

CbAgo is phylogenetically closely related to *TtAgo* (Figure 1D). However, *CbAgo* is 63 amino acids (9.2%) longer than *TtAgo* (748 amino acids versus 685 amino acids) and *CbAgo* and *TtAgo* share only 23% sequence identity. Superposition of the *CbAgo* complex structure with the structure of *TtAgo* bound to a siDNA and DNA target (PDB: 4NCB) (Figure 2C) reveals that the macromolecular architecture and conformation of these *TtAgo* and *CbAgo* structures are highly similar (Core root mean square deviation of 3.0 Å over 563 residues). In the *TtAgo* structure, which is thought to represent a catalytically competent state, a 'glutamate finger' side chain (Glu512^{*TtAgo*}) is inserted into the catalytic site completing the catalytic DDED tetrad (44). Similarly, the corresponding residue in *CbAgo* (Glu577) is located within a flexible loop and is positioned near the other catalytic residues (Figure 2D; Asp541, Asp611 and Asp727). All pAgos and eAgos characterized to date cleave the target strand in between nucleotide 10 and 11 of the target strand. In line with the consensus, the catalytic residues of *CbAgo* perfectly align with the scissile phosphate linking these nucleotides in our structure (Figure 2D). This observation implies that this structure represents the cleavage competent conformation of *CbAgo*.

Only 15 siDNA–target DNA base pairs are formed in the complex, which suggests that additional siDNA–target DNA binding is not essential for target DNA cleavage. To determine the minimum siDNA length that *CbAgo* requires for target binding, we performed single-molecule fluorescence assays. First, *CbAgo* was immobilized on a surface and next it was incubated with 5'-phosphorylated Cy3-labeled siDNAs (Figure 2E). These assays demonstrate that *CbAgo* can bind siDNAs with a minimal length of 12 nucleotides. Next, we determined the minimum siDNA length for *CbAgo*–siDNA mediated target DNA cleavage (Figure 2F). In line with the observation that the *CbAgo* adopts a cleavage-competent conformation when only 14 base pairs are formed, *CbAgo* can cleave target DNAs when programmed with siDNAs as short as 14 nt (forming 13 siDNA–target DNA base pairs) under the tested conditions. This resembles the activity of *PfAgo*, *MjAgo* and *MpAgo*, which require siDNAs with a minimal length of 15 nt to catalyze target DNA cleavage (14,15,43). Only *TtAgo* has been reported to mediate target DNA cleavage with siDNAs as short as 9 nt (12).

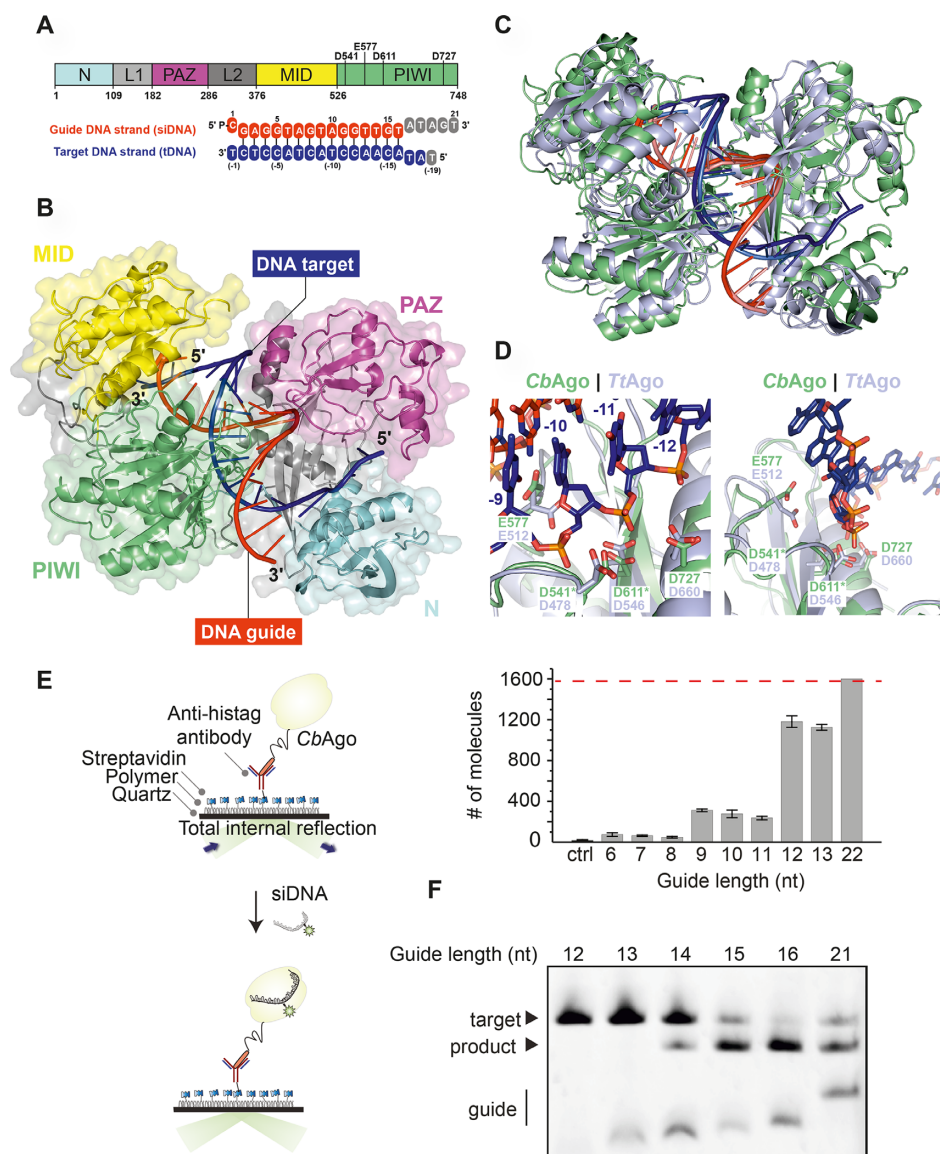


Figure 2. Structure of *CbAgo* in complex with a siDNA and a DNA target. (A) Upper panel: Schematic diagram of the domain organization of *CbAgo*. L1 and L2 are linker domains. Lower panel: Sequences of the siDNA (red) and target DNA (blue). Nucleotides that are unordered in the structure are coloured grey. See also Supplementary Table S1. (B) Overall structure of the *CbAgo*-siDNA-target DNA complex. Domains are coloured according to the colour scheme in panel A. (C) Structural alignment of *CbAgo* (green) and *TtAgo* (light purple; PDB: 4NCB). Core Root Mean Square Deviation of 3.0 Å over 563 residues. (D) Close-up views of the aligned DDED catalytic sites of *CbAgo* (green) and *TtAgo* (light purple; PDB: 4NCB). Modelled side chains of D541 and D611 in *CbAgo* are indicated with green asterisks. The glutamate finger of both pAgos (E512 in *TtAgo* or E577 in *CbAgo*) are inserted into the catalytic site. The scissile phosphate between nucleotide –10 and –11 of the target DNA strand (blue) is indicated with a black asterisk in the left panel. (E) Total internal reflection microscopy (TIRM) was used to determine the minimal length for siDNA to be bound by *CbAgo*. Left panel: Graphical overview of the TIRM method. Right panel: Histogram with TIRM results demonstrated that synthetic siDNAs of at least 12 nt in length are efficiently bound by *CbAgo*. The red line indicates the total number of countable molecules within the microscope image. The raw microscope images are given in Supplementary Figure S5. (F) *CbAgo* mediates target DNA cleavage with siDNAs as short as 14 nucleotides. *CbAgo* was incubated with siDNA and target DNA in a 1:1:1 ratio. Cleavage products were analyzed by denaturing polyacrylamide electrophoresis.

CbAgo associates with plasmid-derived siDNAs *in vivo*

It has previously been demonstrated that certain pAgos co-purify with their guides and/or targets during heterologous expression in *Escherichia coli* (13,16). To determine whether *CbAgo* also acquires siDNAs during expression, we isolated and analyzed the nucleic acid fraction that co-purified with *CbAgo*. Denaturing polyacrylamide gel electrophoresis revealed that *CbAgo* co-purified with small nucleotides with

a length of ~12–19 nucleotides (Figure 3A). These nucleic acids were susceptible to DNase I but not to RNase A treatment, indicating that *CbAgo* acquires 12–19 nucleotide long siDNAs *in vivo*, which fits with its observed binding and cleavage activities *in vitro* (Figures 1 and 2).

We cloned and sequenced the siDNAs that co-purified with *CbAgo* to determine their exact length and sequence. The majority of the siDNAs had a length of 16 nucleotides and are complementary to the plasmid used for expres-

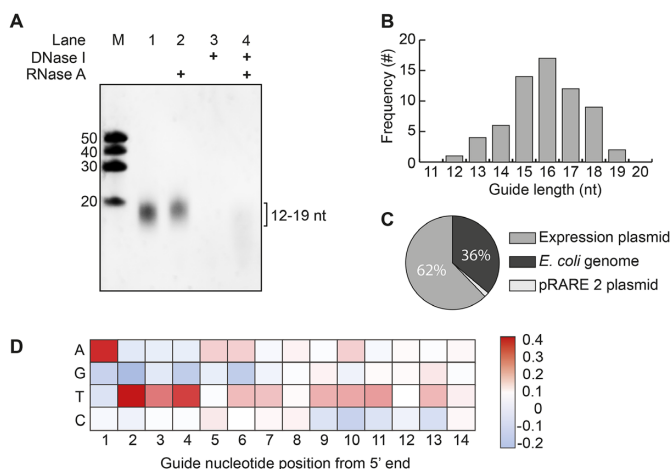


Figure 3. *CbAgo* associates with small plasmid derived siDNA *in vivo*. (A) Nucleic acids that co-purified with *CbAgo* were treated with either RNase A, DNase I or both, and were analyzed by denaturing polyacrylamide gel electrophoresis. (B) Histogram displaying the length of DNA co-purified with *CbAgo* as determined by sequencing. (C) Sequenced nucleic acids that co-purified with *CbAgo* are mostly complementary to the *CbAgo* expression plasmid. (D) Heat map showing the base preference of the co-purified nucleic acids at each position. The red squares indicate bases that were more often found compared to a random distribution (25%); blue squares indicate bases that were less frequently found.

sion of *CbAgo* (Figure 3B and C). Likewise the siRNAs and siDNAs that co-purify with respectively *Rhodobacter sphaeroides* (*RsAgo*) and *TtAgo* are also mostly complementary to their expression plasmids (13,16). As both *TtAgo* and *RsAgo* have been demonstrated to interfere with plasmid DNA, this suggests that also *CbAgo* might play a role in protecting its host against invading DNA. However, no significant reduction of plasmid content could be detected during or upon expression of *CbAgo* in *E. coli* (Supplementary Figure S6). We also investigated whether *CbAgo* co-purified with nucleic acids that were enriched for certain motifs. Sequence analysis revealed that most siDNAs co-purified with *CbAgo* contain a deoxyadenosine at their 5' ends (Figure 3D). In addition, we observed an enrichment of thymidine nucleotides in the three positions directly downstream of the siDNA 5' end (nt 2–4) (Figure 3D).

The sequence of the siDNA affects *CbAgo* activity

To investigate if the 5'-terminal nucleotide of the siDNA affects the activity of *CbAgo*, we performed cleavage assays. *CbAgo* was loaded with siDNA guides with varied nucleotides at position 1 (g1N) and incubated with complementary target DNAs (Figure 4A). Surprisingly, the highest cleavage rates were observed when *CbAgo* was loaded with siDNAs containing a 5'-T, followed by siDNAs containing 5'-A. *CbAgo* bound 5'-G or 5'-C siDNAs displayed slightly lower initial cleavage rates. Also for other pAgos the g1N preference observed *in vivo* is not reflected in the *in vitro* activities; *TtAgo* (which preferentially co-purifies with g1C siDNAs) as well as *PfAgo* and *MpAgo* (of which the *in vivo* g1N preferences are unknown) demonstrate no clear preference for a specific g1N during *in vitro* cleavage reactions

(13,17,43). Instead, the preference of *TtAgo* for 5'-C siDNAs is determined by specific recognition of a guanosine nucleotide in the corresponding position t(-1) in the target DNA (17). Indeed, *TtAgo* structures and models have revealed base-specific interactions with target strand guanine, while base-specific interactions with the 5'-terminal cytosine in the siDNA are less obvious (17). Similarly, we predict no obvious base-specific interactions with the 5'-terminal cytosine in the structure of the *CbAgo* complex (Supplementary Figure S7). When we investigated potential base-specific interactions with the base at the opposing target strand t(-1) position, we observed that the t(-1) thymine base is not placed in the t(-1) binding pocket as has been observed in *TtAgo*, *RsAgo* and hAGO2 (17,46,47). Instead, the thymine bases is flipped and stacks on Phe557 that also caps the siDNA-target DNA duplex (Supplementary Figure S7). At present, we are unable to rationalize the preferential co-purification of 5'-adenosine siDNAs with *CbAgo*.

In order to characterize the seed segment of *CbAgo*, and to test whether the seed length changes depending on the nature of the guide and the target (i.e. DNA versus RNA), we performed additional single-molecule binding assays. The length of seed was determined based on the minimal number of complementary nucleotide pairs between guide and target that were required to achieve a stable binding event. When only nt 2–4 of the siDNA are complementary to the DNA and RNA targets, *CbAgo*–siDNA complexes bound to the DNA target with an average dwell time 58-fold longer compared to RNA target-binding (Figure 4B). When nt 2–7 of the siDNA were complementary to the target, the *CbAgo*–siDNA complex stably bound to both to target DNA and RNA beyond our observation time of 300 s. This suggests *CbAgo* prefers DNA targets above RNA targets and that the seed segment of the siDNAs bound by *CbAgo* comprises nt 2–7.

Next, we set out to investigate whether *CbAgo* displays a preference for siDNAs with thymidines at nt 2–4 *in vitro*, similar to the observed sequence preference for siDNAs that co-purified with *CbAgo* *in vivo*. *CbAgo* was incubated with siDNAs in which nt 2–4 were varied and complementary target DNAs were added. In contrast to the 5'-base preference, the preference for thymidines in the nt 2–4 segment that we observed *in vivo* is also reflected *in vitro*: *CbAgo* displays the highest target cleavage rates when programmed with siDNAs containing thymidines at nt 2–4 (Figure 4C). To confirm these findings, we performed single-molecule assays in which we compared the target binding properties of *CbAgo*–siDNA complexes containing siDNAs with either thymidines or a deoxyadenosines at position 2–4. These assays demonstrate that the dwell time of *CbAgo* loaded with siDNA containing thymidines at nt 2–4 on a target was 18-fold longer compared to *CbAgo* loaded with siDNA containing deoxyadenosines at nt 2–4 (Figure 4D). Combined, these data indicate that *CbAgo* displays a preference for siDNAs containing thymidines at position 2–4.

A pair of *CbAgo*–siDNA complexes can cleave double stranded DNA

Thermophilic pAgos have successfully been used to generate double stranded DNA breaks in plasmid DNA (13,15).

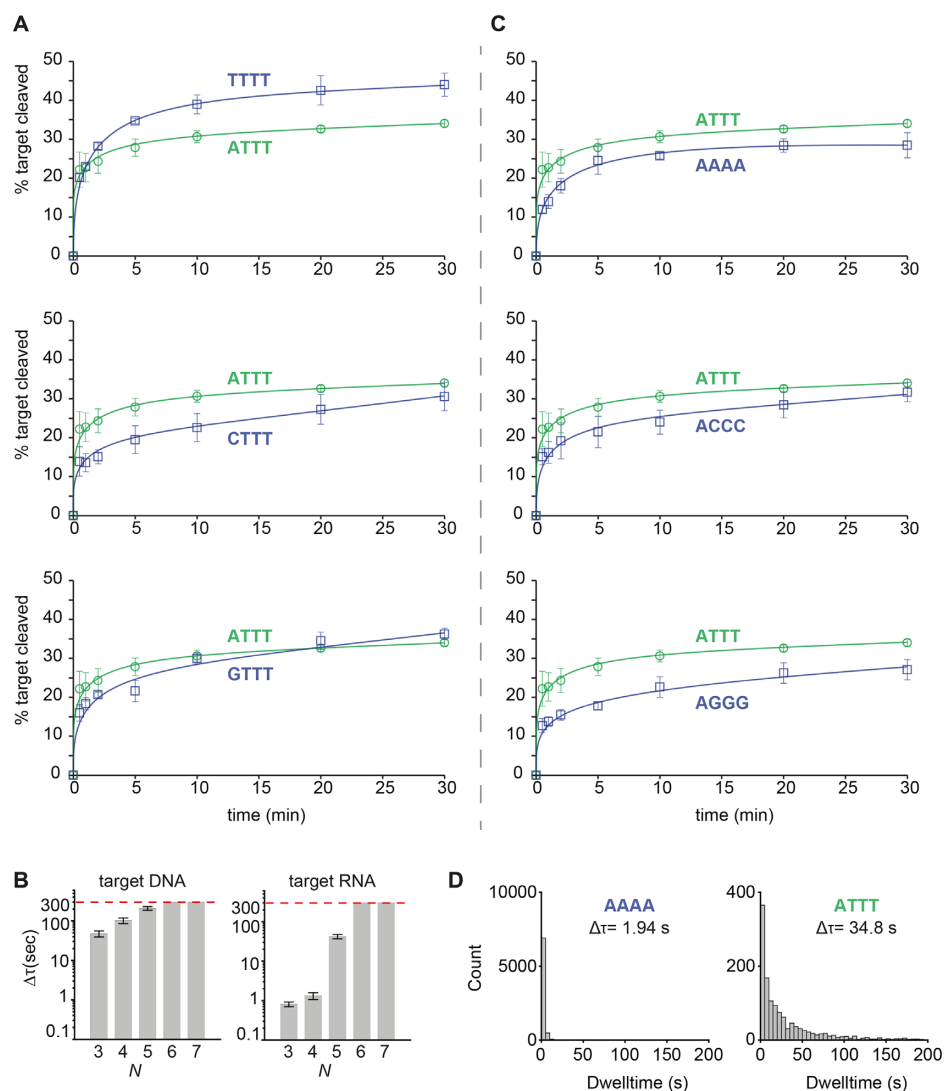


Figure 4. The siDNA sequence affects *CbAgo* activity. **(A)** *CbAgo* has no strong 5'-end nucleotide preference. *CbAgo* was incubated with siDNA with varied 5'-end and incubated with complementary DNA targets. Cleavage products were analysed by denaturing polyacrylamide electrophoresis and quantified. Graphs display the amount of target DNA cleaved. Error bars indicate the standard variation of three independent experiments. **(B)** Histograms displaying dwell time of *CbAgoDM*-siDNA complexes binding either DNA or RNA targets with a varied sequence complementarity (N = number of complementary nucleotides between the siDNA and the target, starting at nt 2. Thus $N3 = nt\ 2-4$). The photobleaching limit is reached where the signal is deactivated (300s). **(C)** *CbAgo* preferentially utilizes siDNAs with thymidines at position 2–4. *CbAgo*-siDNA complexes with siDNA in which nt 2–4 varied were incubated with complementary DNA targets. Cleavage products were analysed by denaturing polyacrylamide electrophoresis and quantified. Graphs display the amount of target DNA cleaved. Error bars indicate the standard variation of three independent experiments. **(D)** Histograms displaying dwell time of *CbAgoDM* in complex with a 5'-ATTT siDNA or 5'-AAAA siDNA binding to a target DNA. Interactions of *CbAgo* are on average ~18-fold longer with the siDNA containing a 5'-ATTT motif compared to interactions with siDNAs containing a 5'-AAAA motif.

As each pAgo–siDNA complex targets and cleaves a single strand of DNA only, two individual pAgo–siDNA complexes are required for dsDNA cleavage, each targeting another strand of the target dsDNA. Although all pAgos characterized so far appear to lack the ability to actively unwind or displace a dsDNA duplex substrate, it has been proposed that, at least *in vitro*, thermophilic pAgos rely on elevated temperatures ($>65^{\circ}\text{C}$) to facilitate local melting of the dsDNA targets to target each strand of the DNA individually. However, *CbAgo* is derived from a mesophilic organism and we therefore hypothesize that it is able to mediate protection against invading DNA at moderate temperatures (37°C). To test if *CbAgo* can indeed cleave ds-

DNA targets at 37°C , we incubated apo-*CbAgo* and pre-assembled *CbAgo*-siDNA complexes with a target plasmid. Previous studies showed that the ‘chopping’ activity of siDNA-free apo-*TtAgo* and apo-*MjAgo* can result in plasmid linearization or degradation, respectively (14,17). We observed that apo-*CbAgo* converted the plasmid substrate from a supercoiled to open-circular state, possibly by nicking one of the strands, but did not observe significant linearization or degradation of the plasmid DNA (Figure 5A). When the plasmid was targeted by *CbAgo* loaded with a single siDNA, we also observed loss of supercoiling (Figure 5A). As this activity was not observed with nuclease-deficient *CbAgoDM*, we conclude that apo-*CbAgo* and

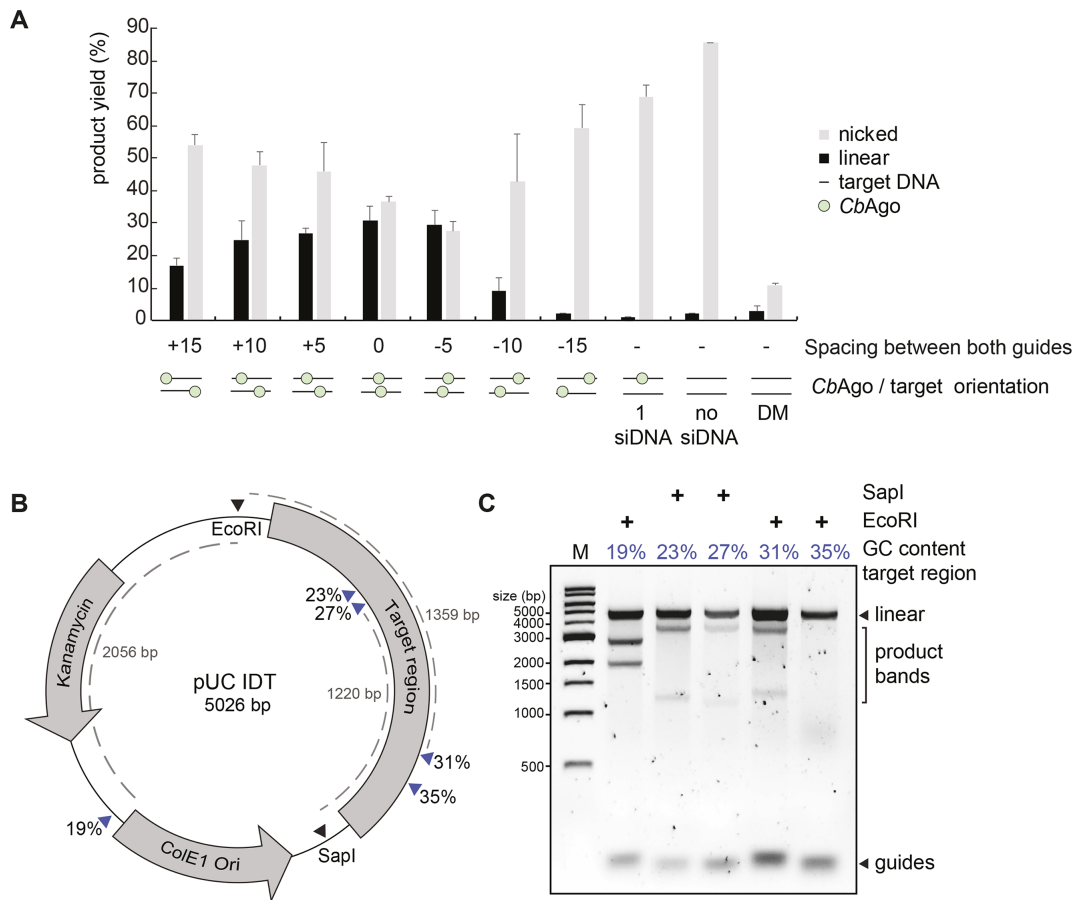


Figure 5. Double stranded plasmid DNA cleavage by *CbAgo*. (A) Two *CbAgo*–siDNA complexes can generate double stranded DNA breaks in plasmid DNA. *CbAgo*–siDNA complexes were pre-assembled and incubated with target plasmid DNA. Cleavage products were analysed by agarose gel electrophoresis (Supplementary Figure S8B) and quantified. The spacing between both *CbAgo*–siDNA target sites affects the linearization efficiency (nucleotide spacing between the predicted cleavage sites: +15 nt, +10 nt, +5 nt, 0 nt, –5 nt, –10 nt, –15 nt, a single siDNA, no siDNA). With 0 nt spacing, both *CbAgo*–siDNA complexes are exactly on top of each other. (B) Schematic overview of the pUC IDT target plasmid. Blue arrows indicate target sites while percentages indicate the GC-content of the 100 bp segments in which these target sites are located. (C) Pre-assembled *CbAgo*–siDNA complexes targeting various pUC IDT segments were incubated with pUC IDT. Cleavage products were incubated with EcoRI or SapI and were further analysed by agarose gel electrophoresis. The GC-content of the segments in which the target sites were located are indicated by the percentage (in blue).

CbAgo–siDNA complexes are generating nicks in dsDNA plasmid targets with their DEDD catalytic site. When using two *CbAgo*–siDNA complexes, each targeting one strand of the plasmid, we observed that a fraction of the target plasmid DNA becomes linearized (Figure 5A). This implies that *CbAgo*–siDNA complex-mediated nicking of each of the target plasmid DNA strands resulted in the generation of a double stranded DNA break. Next, we investigated if the spacing between the two siDNAs affects the ability of *CbAgo* to cleave the plasmid. The most efficient plasmid linearization was achieved when the siDNAs were orientated exactly or almost opposite to each other (Figure 5A).

Finally, we investigated whether the GC-content of the target DNA plays a role during DNA targeting by *CbAgo*. For *TtAgo*, it has been observed that AT-rich DNA is cleaved more efficiently than GC-rich DNA (17), possibly because AT-rich DNA is more prone to unwinding. To test if such preference also exists for *CbAgo*, we designed a target plasmid containing 16 gene fragments of 100 bp complementary to sequences from the human genome, with an increasing GC content (Figure 5B). The target plasmid

was digested with a restriction enzyme (SapI or EcoRI) either before or after the incubation with two *CbAgo*–siDNA complexes (Figure 5B and Supplementary Figure S8A). Interestingly, *CbAgo* could cleave the negatively supercoiled plasmid, but not the linearized plasmid, suggesting that, like *TtAgo*, *CbAgo* relies on the negative supercoiled state of the target plasmid to facilitate local DNA melting before targeting and cleavage can take place (13). Furthermore, *CbAgo*–siDNA complexes were only able to generate dsDNA breaks in gene fragments with a GC-content of 31% or lower (Figure 5C). We hypothesize that, at moderate temperatures, the supercoiled state of the target plasmid allows for local DNA unwinding especially in AT-rich DNA regions. Taken together, this suggests that, like thermophilic pAgos, *CbAgo* relies on unwinding of the dsDNA before targeting and cleavage can take place.

DISCUSSION

Several prokaryotic Argonaute proteins have been demonstrated to protect their host against invading nucleic acids,

such as plasmid DNA (13,15,16). Similar to *TtAgo* and *RsAgo*, *CbAgo* co-purifies with guides which are preferentially acquired from the plasmid used for its heterologous expression in *E. coli*. In addition, *CbAgo* mediates programmable DNA-guided DNA cleavage *in vitro*. This suggests that, similar to the phylogenetically related *TtAgo*, also *CbAgo* can interfere with plasmid DNA via DNA-guided DNA interference.

Sequencing of the nucleic acids that co-purified with *CbAgo* revealed that *CbAgo* preferentially associates with siDNAs with a 5'-ATTT-3' sequence at their 5' end. It was previously shown that RNA guides utilized by eAgos can be divided into functional segments. These segments are (from 5' to 3') the anchor nucleotide (nt 1), the seed (nt 2–8) and sub-seed segments (nt 2–4/2–5), the central (nt 9–12), the 3' supplementary (nt 13–16) and the tail (nt 17–21) segments (48,49). Extending this knowledge to the siDNAs that co-purified with *CbAgo*, *CbAgo* preferentially associates with siDNAs that have a 5'-terminal adenosine anchor (nt 1) and a thymidine-rich sub-seed. In RNAi pathways, the preference for a specific 5'-terminal nucleotide is important for guide RNA loading into a subset of eAgos (50–52). Similarly, several pAgos including *RsAgo*, *TtAgo*, and now *CbAgo* also preferentially associate with specific 5'-terminal nucleotides *in vivo* (13,16). However, for both *CbAgo* and *TtAgo*, there is no clear preference for siDNAs with that specific 5'-base during cleavage assays *in vitro*. Rather than having a functional importance, the preference of pAgos for a specific nucleotide at the siDNA 5' end might be a consequence of siDNA generation and/or loading, as has been demonstrated for *TtAgo* (17). Several studies on human Ago2 have described the importance of the sub-seed segment (nt 2–4/2–5) in its RNA guides (48,53,54). For hAgo2, a complete match between the guide RNA sub-seed segment and the target RNA triggers a conformational change that first exposes the remainder of the seed (nt 5–8/6–8), and eventually the rest of the guide. This facilitates progressive base pairing between the guide RNA and the target (55). However, a specific nucleotide preference in the sub-seed segment, as we have observed for *CbAgo*, has not been described for any other Argonaute protein. The preference for the T-rich sub-seed is not only observed in the *in vivo* acquired siDNAs, but also plays a clear role during target binding and cleavage assays *in vitro*. This may reflect a structural preference for these thymidines in the cleft of the PIWI domain. We have not been able to obtain diffracting crystals of *CbAgo* in complex with siDNAs that have a 5'-ATTT-3' sequence at the 5'-end. Future research will thus be necessary to determine the structural basis the apparent preference for these nucleotides at these positions. We hypothesize that this bias might reflect the mesophilic nature of *CbAgo*, which might have better access to AT-rich dsDNA fragments, both for siDNA acquisition and for target cleavage.

Several DNA-targeting pAgos have been repurposed for a range of molecular applications among which a cloning, recombineering and nucleic acid-detection method (22,23,56,57). Additionally, the potential repurposing of pAgos for genome editing applications has previously been discussed (27). However, all characterized DNA-cleaving pAgos to date originate from thermophilic prokaryotes

and are solely active at elevated temperatures, which limits the potential repurposing of pAgos for applications that require moderate temperatures, such as genome editing. The biochemical characterization of *CbAgo* reported herein is the first example of a pAgo that catalyzes siDNA-guided dsDNA cleavage at 37°C, indicating that the pool of mesophilic pAgos contains candidates that—in theory—can be utilized for potential applications that require moderate temperatures, such as genome editing. If *CbAgo* or other mesophilic pAgos could be harnessed for genome editing, they will have certain advantages over the currently well-established genome editing tools CRISPR-Cas9 and CRISPR-Cas12a; While CRISPR-based genome editing tools can be programmed with a guide RNA to target DNA sequences of choice, target DNA cleavage additionally requires the presence of a protospacer adjacent motif (PAM) next to the targeted sequence (5'-NGG-3' for Cas9 and 5'-TTTV-3' for Cas12a) (58). This limits the possible target sites of Cas9 and Cas12a. In contrast, pAgos do not require a PAM for DNA targeting, which would make them much more versatile tools compared to CRISPR-associated nucleases. However, PAM binding by Cas9 and Cas12a also promotes unwinding of dsDNA targets (59–61) which subsequently facilitates strand displacement by the RNA guide, and eventually R-Loop formation. The absence of such mechanism in pAgos might explain their limited nuclease activity on dsDNA targets.

Here, we have demonstrated that *CbAgo* does not strictly rely on other proteins when targeting AT-rich dsDNA sequences *in vitro*. As such, this study provides a foundation for future efforts to improve double stranded DNA target accessibility of pAgos and to facilitate the further development of pAgo-based applications at moderate temperatures.

DATA AVAILABILITY

Atomic coordinates and structure factors reported in this paper have been deposited in the Protein Data Bank (PDB) with the accession number PDB: 6QZK.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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Authors contributions: J.W.H. and J.v.d.O. conceived the project and designed the biochemical experiments, which were performed by J.H. and J.K. Single-molecule experiments were designed by S.C., T.J.C. and C.J. and performed by S.C. and T.J.C. X-ray crystallographic analysis was designed and performed by D.C.S. under the supervision of M.J., J.W.H., D.C.S., C.H., M.J., C.J. and J.v.d.O. wrote the manuscript. All authors read and approved the manuscript.

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