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Interaction of a Model Peptide on Gram Negative and Gram Positive Bacterial Sliding Clamps.

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SUMMARY.

Bacterial sliding clamps control the access of DNA polymerases to the replication fork and are appealing molecular targets for antibacterial drugs development. To this end, it is critical to decipher the polymerase-clamp binding mode across various bacterial species. We showed previously that synthetic peptides targeting the clamp binding pocket of Gram- bacteria poorly interact with that of Gram+ homologous proteins. Here we analyzed the interaction of a reference peptide with several *E. coli* and *B. subtilis* clamp variants. For both Gram- and Gram+ pockets, the peptide binds through an induced-fit process but the complex stability

varies according to a pocket specific network of interactions. Thermodynamic and molecular dynamics analyses identify a strategic position in the pocket where a mobile residue is necessary for an efficient peptide interaction. A residue at another position modulates the folding dynamics of the pocket upon ligand binding in *E. coli*, while in *B. subtilis*, this residue is essential for polymerase activity and might thus be a Gram+ specific molecular marker.

INTRODUCTION.

Bacterial resistance to antibiotics is a major threat for human health. According to several official reports, a return to the pre-antibiotic era during the 21st century is a realistic possibility¹. In order to tackle this challenge, one strategy aims at identifying new bacterial molecular targets and at developing efficient molecules that will block their physiological functions. The bacterial processivity factor, also referred to as the sliding clamp (SC), has been previously identified as a potential drugable new target^{2 3 4 5 6}. The ultimate proof of concept was recently brought by the natural cyclic peptides griselimycins that bind to *Mycobacterium tuberculosis* sliding clamp (*Mt*SC) and display good *in vivo* anti-bacterial activity in animal models⁷. However, the molecular mechanisms that govern the SC/ligand interaction are not well understood⁴⁵⁸⁹.

In bacteria, SC is a homodimer that encircles and slides along double stranded DNA¹⁰. It binds the replicative DNA polymerase, thus conferring a high processivity to the resulting holoenzyme^{11 12}. It also serves as a molecular hub on which all the other DNA polymerases bind (polI, II, IV and V in *E. coli*)^{13 14} as well as other enzymes involved in DNA metabolism¹⁵ and this interaction is required for these proteins to fulfill their functions¹³. Remarkably, in all cases, SC-protein interaction is mediated by a short peptide segment which encompasses the consensus sequence (QL[S/D]LF)¹⁶. This peptide portion binds SC within a hydrophobic pocket located between domains II and III of SC¹⁷³ and formed by two sub-sites joined by a shallow groove: sub-site 1, also referred to as the leucine-rich pocket, interacts with the C-terminal part of the peptide, whereas sub-site 2 binds the highly conserved Q residue (**SI.1**).

We have previously observed that short synthetic peptides designed to bind to *Escherichia coli* SC (*Ec*SC) with an increased affinity also interact efficiently with SC from other Gram- bacteria such as *Pseudomonas aeruginosa*⁵. However, they show a lower interaction with *Mt*SC and fail to interact efficiently with SC from other Gram+ strains such as *Staphylococcus aureus* and *Bacillus subtilis* (*Bs*SC), suggesting that peptide binding on SC occurs through a different process in Gram+ and Gram- bacteria¹⁸.

A careful analysis of the peptide binding process on *Ec*SC revealed that a fully efficient binding pocket folds upon peptide binding^{3 5 19}. Notably, two residues, S₃₄₆ and M₃₆₂, seem to play a strategic role in the interaction process as, upon peptide interaction, their side chains

undergo a large concerted movement that allow the opening of a groove in which the linear peptide can lie ⁵ (**SI.1**). Interestingly enough, these two residues are conserved in Gram-strains but not in Gram+ bacteria, where they are essentially replaced by proline and leucine/threonine, respectively. Noteworthy, in ^{Mt}SC only the residue corresponding to ^{Ec}SC S₃₄₆ is changed into proline, whereas the methionine residue (^{Mt}SC M₃₉₆) is conserved. Thus, as far as these two residues are concerned, ^{Mt}SC (P₃₆₂, M₃₉₆) stands as an intermediate between ^{Ec}SC (S₃₄₆, M₃₆₂) and ^{Bs}SC (P₃₅₇, L₃₇₃) (**Table 1**). These observations, in correlation with peptide binding analyses ¹⁸, suggest that the mode of peptide interaction might be linked to the gating function of these two residues.

In this work, we evaluate the respective contribution of residues ^{Ec}SC S₃₄₆ and ^{Ec}SC M₃₆₂ and their corresponding residues in ^{Mt}SC and ^{Bs}SC, to peptide binding. For each SC, every residue was mutated (M→L, S→P, L→M and P→S), leading to a series of single and double mutants (**Table 1**). The interaction of peptide **P**₇ (AcQXD_LLF, X = cyclohexylalanyl, Cha) (**SI.2**)⁵ with each of these SCs was analyzed by ITC, X-Ray crystallography, molecular dynamics and *in vitro* replication assays. Our results reveal the prominent role of these residues for the peptide interaction with Gram- and Gram+ SC, and will have strong implications for the design of strain-specific anti-replicative molecules.

MATERIAL AND METHODS.

Construction of mutant SC, expression and purification of processivity factors.

Plasmids pET15b containing the *dnaN* genes from *E. coli*, *M. tuberculosis* or *B. subtilis* ¹⁸ were site specifically mutagenized using the Quikchange® procedure (Stratagene). Oligonucleotides were from IDT. Selected recombinant plasmids were sequenced (GATC, Kontanz, Ge) and transformed in BL21 (DE3) pLys *E. coli* strains. For expression of *dnaN* proteins, cells were grown in LB at 37°C to OD 0.5, then induced by IPTG (0.1mM) at 28°C overnight. *dnaN* proteins fractions were first enriched on a Ni-NTA column, eluted with an histidine step (300 mM) and further purified on a Source Q column in buffer containing 20 mM Tris HCl pH 7.5, 0.5 mM EDTA and 10% (v/v) glycerol, using a gradient from 0 to 0.5 M NaCl. After a final ultracentrifugation (45K, 1 h, 20°C), soluble proteins were concentrated on a Centricon 30K (Millipore) in the same buffer and stored at 4°C in 2 M ammonium sulfate. Buffer exchange was performed on Centrikon 10K at 4°C before use and protein quality was assessed by DLS analysis (**SI.3**).

Peptide synthesis.

Procedure for the synthesis of peptide **P₇** has been described elsewhere⁵. The chemical formula is shown in supporting information (**SI.2**)

Isothermal Titration Calorimetry.

ITC was performed using an iTC₂₀₀ (Microcal Malvern Panalytical) or a PEAQ-ITC instrument (Microcal Malvern Panalytical). Peptides (300 or 600 μM) were titrated at different temperatures (15, 25 and 34°C) by sequential injections (usually 2 μl each) into a SC solution (30 or 60 μM). Data were corrected for heat of injection by subtracting the signal of the titration of peptides into protein-free buffer solution (Hepes 10 mM pH 7.4, NaCl 0.15 M, EDTA 3 mM) (**SI.4**). Each titration was performed at least twice. Analyses of experimental data were performed following a classical treatment with the AFFINImeter software (<https://www.affinimeter.com>; S4S, Santiago de Compostela, Spain). Both types of analyses yield similar data (data not shown). Non classical methods were also used. First a global thermodynamic treatment (GTT, see below) was performed. In addition, kinetic information was obtained in some cases with *kinITC*^{20 32} as implemented in the software AFFINImeter. A comparison of GTT and classical treatment analyses is presented in **SI.5**. All thermodynamic data are provided in **SI.6**, representative titration curves are presented in **SI.7** and all thermodynamic profiles are presented in **SI.8**.

Global thermodynamic treatment (GTT) of ITC experiments.

This procedure has been described in²⁰. Briefly, the method consists in using a single reduced set of parameters to fit at once all experimental titration curves obtained at different temperature. The free parameters are K_{d0} and ΔH_0 , the values of K_d and ΔH at a reference temperature T_0 (the mean temperature of all experiments) and $\Delta C_p = \partial\Delta H/\partial T$ governing the evolution of ΔH with the temperature. These three parameters alone, have to replace $2 \times N_{\text{exp}}$ parameters (one K_d and one ΔH for each experiment). This GTT yields theoretical curves describing the evolution of all thermodynamic parameters with the temperature. These curves are shown with the points corresponding to the results from the usual individual treatments: it is important to stress that they do not result from a fit of these points (see **Figure 1**). A comparison of the data obtained by GTT and classical treatment is presented in **SI.5**. When the peptide interaction with SC was weak, the deviation between the points from the

individual treatments and the curve from GTT was large (see **Figure 1B** and **SI.5 B**). In these cases, the reported results are from the individual treatments. Of course, the discrepancy just mentioned is a mark of poor quality.

Crystallogensis, X-ray diffraction, data collection and processing.

Crystals of $^{Ecwt}SC/P_7$ and $^{EcMt}SC/P_7$ complexes were obtained by screening the crystallization reagent kit PEG/Ion HT from Hampton Research. 200 nL sitting drops were prepared on plates (Greiner XTL round) by mixing 50 or 100 nL of preformed SC/P_7 complexes (protein concentration from 6 to 12 mg/ml; protein/peptide ratio: 1.5) with 150 or 100 nL of crystallization solution, using a Mosquito pipetting station (TTP Labtech). Plates were incubated at 20°C (^{Ec}SC) or 4°C (^{Mt}SC). Crystals were obtained in several crystallization conditions and were frozen in liquid ethane. Crystals with the higher resolution were obtained in the following conditions: $^{Ecwt}SC/P_7$: MES 50 mM pH 6, CaCl₂ 50 mM PEG400 30% (m/v) + PEG/Ion HT kit condition E6 (0.2 M sodium malonate pH 6, PEG 3350 20% (m/v)). $^{EcM1}SC/P_7$: MES 50 mM pH 6, CaCl₂ 50mM, PEG400 28% (m/v) + PEG/Ion HT kit condition B3 (0.2 M lithium nitrate, PEG 3350 20% (m/v), pH 7.1). $^{Mtw}SC/P_7$: Acetate de Na 3.2 M pH 6.9, PEG 3350 18% (m/v) + PEG/Ion HT kit condition A2 (0.2 M potassium fluoride, PEG 3350 20% (m/v), pH 7.3). $^{MtM1}SC/P_7$: Acetate de Na 3.2 M pH 6.9, PEG 3350 18% (m/v) + PEG/Ion HT kit condition B8 (Magnesium formate dihydrate 0.2 M; PEG 3350 20% (m/v), pH 7).

The data were collected at the X06DA PXIII beamline at the Swiss Light Source (Villigen, Switzerland), equipped with a PILATUS 2M detector and the multiaxis PRIGo goniometer ²¹. The wavelength was 1.00 Å, with a flux at full transmission of 4·10¹¹ photons·s⁻¹. The data were collected with two rotations of 360° at $\chi=0^\circ$ and $\chi=30^\circ$ (0.2° and 0.1s per frame). The crystal-to-detector distance was 300 mm. The data were processed and merged using the software autoPROC version 1.0.5 ²² and STARANISO (Cambridge, United Kingdom: Global Phasing Ltd) ²³. The structures were solved by molecular replacement using the software MOLREP version 11.4.04 ²⁴ and search models 1OK7 for the *E. coli* cases and 4TR7 for the *M. tuberculosis* cases. Refinement was performed with BUSTER version 2.10.3 (Cambridge, United Kingdom: Global Phasing Ltd) and phenix.refine version 1.10.1-2155 ²⁵, with NCS constraints for all structures and TLS for both higher resolution *E. coli* structures. Secondary structure restraints were additionally imposed in phenix.refine for the *Mtw*-P7 structure. 5% of reflections were set aside for the R_{free} test set.

Data processing and refinement statistics are available in **SI.9** The 4 structures ^{Ecwt}SC/P₇, ^{EcM1}SC/P₇, ^{Mtwt}SC/P₇ and ^{MtM1}SC/P₇ are deposited in the Protein Data Bank (PDB) under the respective accession codes 6FVL, 6FVM, 6FVN and 6FVO. No complex structure could be solved for any ^{Bs}SC.

The solvent accessible surface area was calculated using AREAIMOL in CCP4²⁶. The difference in accessible surface area of a binding pocket with and without P₇ was obtained by considering the structures from this work and from Wolff et al.¹⁸ (PDB codes 1OK7 for ^{Ecwt}SC and 4TR7 for ^{Mtwt}SC).

Replication assays.

Escherichia coli Pol III* purification. *E. coli* strain BLR transformed with pHOC2.6.1 (kindly provided by C. McHenry), coding for all the 8 polIII subunits, were grown at 37°C in LB buffer supplemented with 100 µg/ml carbenicillin to reach an OD_{600nm} of about 0.65, transferred to 30°C and then expression of Pol III* subunits was started by adjunction of 1 mM IPTG and extended for 3 h. Cell pellets (16.8 g) were stored at -20°C overnight. Purification of PolIII* was conducted essentially as described previously²⁷. Briefly, pellets thawed at 4°C for 30 min were resuspended in 50 ml of 50 mM Hepes-KOH pH 7.5, 10% (m/v) sucrose, 10 mM DTT. 5 ml of lysis buffer (50 mM Hepes-KOH pH 7.5, 10% (m/v) sucrose, 2 M NaCl, 0.3 M spermidine) were added to the cell suspension and Tris base 2 M was used to adjust to pH 8. Lysis was performed on ice for 1 h after addition of lysosyme to 0.2 mg/ml. After centrifugation at 20 000 rpm for 1 hour at 4°C (Beckmann 70 Ti rotor), proteins were precipitated by slow addition of 0.226 g/ml ammonium sulfate and incubation 30 min at 4°C with agitation. After centrifugation (45 min, 16 000 rpm, 4°C), pellet was successively extracted with 3.275 ml of buffer A (25 mM Hepes-KOH pH 7.5, 5% (v/v) glycerol, 100 mM KCl, 1 mM EDTA, 5 mM DTT) supplemented with 0.2 g/ml ammonium sulfate, centrifuged, and further extracted with 1.2 ml of buffer A with 0.7 g/ml ammonium sulfate. After centrifugation (20 min, 30, 000 rpm, 4°C), the pellet was resuspended in buffer A without KCl and conductivity was adjusted to the one of buffer B (25 mM Hepes-KOH pH 7.5, 5% (v/v) glycerol, 20 mM KCl, 1 mM EDTA, 5 mM DTT), then loaded onto a SP sepharose column (1 ml, GE Healthcare) equilibrated with buffer B and developed with a linear gradient of 0 to 30 % buffer C (25 mM Hepes-KOH pH 7.5, 5% (v/v) glycerol, 1 M KCl, 1 mM EDTA, 5 mM DTT) on AKTA smart chromatography system (GE Healthcare).

Finally, pooled fractions were buffer exchanged for buffer D (25 mM Hepes-KOH pH 7.5, 5% (v/v) glycerol, 20 mM KCl, 5 mM DTT) on a PD10 column (GE Healthcare) and concentrated by ultra-filtration on Amicon Ultra centrifugal units (MWCO 10K), adjusted to 50% (v/v) glycerol and to 0.21 mg of protein per ml and stored at -80°C.

Bacillus subtilis minimal replicase (BsP) was reconstituted by assembly of purified proteins (kindly provided by Charles McHenry and Paul R. Dohrmann) at ratio of 1 PolC, 5 tau, 1.25 delta and 1.25 delta' to provide a 1.25 μ M enzyme complex in buffer HD (40 mM Tris acetate pH=7.8, 10 mM Mg acetate, 3 μ M zinc sulfate and 0.015 % (v/v) IGEPAL CA-630).

In vitro primer extension assays. Single-stranded pUC118 DNA was annealed to a 5' radiolabeled 40 mer oligonucleotide (5'GCTGGCGAAAGGGGGATGTGCTGCAAGGCGA TTAAGTTGG3') in 50 mM NaCl and 15 mM Tris acetate (pH=7.8). All DNA synthesis assays were pre-assembled on ice in RB buffer (HD buffer supplemented with 2 mM DTT, 150 mM K glutamate, 160 μ g/ml BSA, 0.5 % (m/v) PEG 8000, 250 μ M ATP and 100 μ M each dNTP). Unless specified, each reaction contains 40 fmoles of primed DNA, 100 fmoles of polymerase complex and 100 fmoles, as a dimer, of beta sliding clamp (SC). DNA synthesis assay were performed for 6 min. at 37°C before addition of 15 μ l of 95% formamide dye loading buffer and heating at 95°C for 5 minutes. Radioactive reaction products were analysed by electrophoresis on 6 to 8 % denaturing (8 M urea) polyacrylamide gels followed by imaging using a Typhoon FLA 9500 device (GE Healthcare).

P7 primer extension inhibition assays were performed essentially as above except that mixtures of SC (100 fmoles in HD buffer, 1 μ l) and P7 peptide (diluted in HD buffer, 1 μ l) were first incubated for 5 min. at RT (20-25 °C) before adding 8 μ l of primed DNA plus polymerase complex in RB buffer.

Molecular Dynamics.

Initial Structures:

Experimental X-Ray structures of ^{Ecwt}SC were used as a starting point for the molecular dynamics simulations of the ^{Ecwt}SC in complex with a de-acetylated form of P₇ (QXDLF, X = Cha). Starting structure for the ^{EcM2}SC (M₃₆₂L) was constructed from the ^{Ecwt}SC by side-chain modification using PyMol. *B. Subtilis* simulations were constructed from the ^{Bswt}SC structure (PDBID 4TR6, ¹⁸) by inserting the appropriate side-chain replacements with PyMoL for the

mutant ^{Bs}M2SC (L₃₇₃M). For both ^{wt}SC and mutant ^{M2}SC, the peptide was positioned by superimposition onto P₁ (PDB 1OK7) and P₁ was subsequently removed.

Free-energy perturbation calculation:

The relative affinities of the ^{wt}SC and ^{M2}SC for the peptide were estimated using free energy perturbation implemented in NAMD²⁸ in the following thermodynamic cycle^{29,30}:

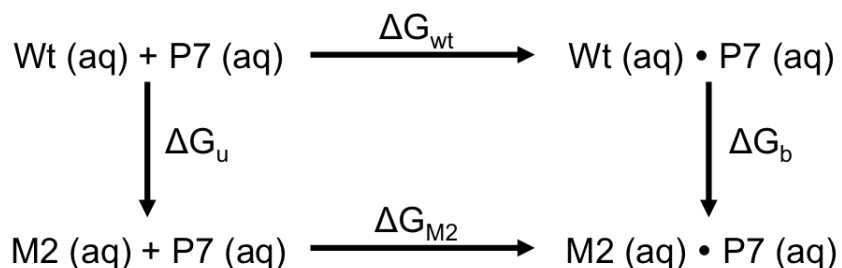


Figure 1: Thermodynamic cycle used in computing the free energy change in the virtual mutation of the ^{wt}SC into the mutant ^{M2}SC in the unbinding form (ΔG_{u}) and in the binding form (ΔG_{b}).

These relative affinities, $\Delta\Delta G$, can be calculated from the following equation:

$$\Delta\Delta G = \Delta G_{\text{M2}} - \Delta G_{\text{wt}} = \Delta G_{\text{b}} - \Delta G_{\text{u}} \quad (1)$$

ΔG_{u} and ΔG_{b} are calculated by transforming the Hamiltonian of the initial states (either the apo WT (ΔG_{u}) or the holo WT (ΔG_{b})) into the final states (either the apo mutant M2 (ΔG_{u}) or the holo mutant M2 (ΔG_{b})). The transformation is modeled by introducing a state variable, λ , which can vary from 0 (initial state) to 1 (final state). The free-energy calculation is split into small windows, each one involving a small variation of λ . A new residue is designed to simulate the transformation (for example of the residue M₃₆₂ into L₃₆₂). In that way, we have built a new residue, name M2L, which includes the atoms of the side chains of a methionine and of a leucine. The structure was then generated by using the force field (FF) charmm 27³¹ and was immersed in a tetragonal box filled with TIP3 water molecule and 24 Na⁺ ions to neutralize the total charge of the full system. The simulation was then heated at 300K and equilibrated before the free-energy perturbation calculation.

RESULTS AND DISCUSSION.

We have previously observed that peptides initially designed to bind Ec SC with high affinities poorly interact with Mt SC and Bs SC, although the binding pockets of all SC are structurally highly homologous¹⁸. In order to decipher the molecular basis of these different binding processes, we determined the thermodynamic parameters of the interaction between a reference ligand, namely peptide **P₇** (**SI.2 and SI.10**) and different variants of Ec SC, Mt SC and Bs SC (**Table 1**).

Peptide P₇ differentially interacts with natural SC from different bacteria.

We first determined the thermodynamic profiles at different temperatures of **P₇** binding on natural SC, namely Ecwt SC, Mrwt SC and Bswt SC, (**Figure 1**). For each SC/**P₇** interaction, global thermodynamic treatment (GTT)³² of all experiments at all temperatures yielded theoretical curves describing the evolution of each thermodynamic parameters with temperature (**Figure 1**, left column). Thermodynamic profiles are inferred from these curves (**Figure 1**, right column). The profiles observed with the three natural SC are very different (**Figure 1**, see also **SI.5** for a comparison between GTT and individual analysis of ITC raw data and **SI.6** for numerical data).

The Ecwt SC/**P₇** interaction is enthalpy driven, with a significant increase of the enthalpic contribution from -10 to -14 kcal/mole when temperature rises from 15 to 34°C. This enthalpic contribution is much weaker for the Mrwt SC (-4 to -8 kcal/mol), and for Bswt SC (-6 to -3 kcal/mol) over the same range of temperature (**Figure 1** right column, **SI.5**). Nevertheless, the binding process is still enthalpy-driven for Mrwt SC and Bswt SC. The ΔC_p ($\partial\Delta H/\partial T$) values are -210 cal/mol/deg for **P₇** binding onto Ecwt SC, -180 cal/mole/deg with Mrwt SC. The ΔC_p for Bswt SC was not reliably determined. **P₇** interaction with Ecwt SC shows an unfavorable entropic contribution (-T ΔS) which increases with temperature (**Figure 1A** and **SI.5**). For Bswt SC, this contribution is weaker and becomes favorable above 25°C (**Figure 1B**), while for Mrwt SC, it evolves in an opposite way (**Figure 1C**). The Gibb's free energy, ΔG , also varies significantly between the different complexes. For Ecwt SC, it reaches -9 kcal/mole and slightly increases with higher temperatures (**Figure 1** and **SI.5**). It reaches about -5 and -7 kcal/mole for Bswt SC and Mrwt SC, respectively, indicative of a weaker interaction between **P₇** and these two SC, and in both cases, it varies very slightly over the range of temperature. Interestingly, ΔG reaches a maximum at 18°C for Bswt SC/**P₇** and a minimum at 29°C for Mrwt SC/**P₇** interactions due to the null value of ΔS at these temperatures (**Figure 1** left

column), implying that around these temperatures, the ΔG variation is minimal. For $^{Ecwt}SC/P_7$, extrapolation of the curve $\Delta S = f(T)$ indicates that $\Delta S = 0$ at $T < 0^\circ C$.

	<i>E. coli</i>		<i>M. tuberculosis</i>		<i>B. subtilis</i>	
	Name	Amino acids	Name	Amino acids	Name	Amino acids
wt	^{Ecwt}SC	S ₃₄₆ ; M ₃₆₂	^{Mrwt}SC	P ₃₆₂ ; M ₃₉₆	^{Bswt}SC	P ₃₅₇ ; L ₃₇₃
M1	^{EcM1}SC	P ₃₄₆ ; M ₃₆₂	^{MtM1}SC	S ₃₆₂ ; M ₃₉₆	^{BsM1}SC	S ₃₅₇ ; L ₃₇₃
M2	^{EcM2}SC	S ₃₄₆ ; L ₃₆₂	^{MtM2}SC	P ₃₆₂ ; L ₃₉₆	^{BsM2}SC	P ₃₅₇ ; M ₃₇₃
M3	^{EcM3}SC	P ₃₄₆ ; L ₃₆₂	-	-	^{BsM3}SC	S ₃₅₇ ; M ₃₇₃
G1	^{EcG1}SC	G ₃₄₆ ; M ₃₆₂	-	-	^{BsG1}SC	G ₃₅₇ ; L ₃₇₃
G2	^{EcG2}SC	S ₃₄₆ ; G ₃₆₂	-	-	^{BsG2}SC	P ₃₅₇ ; G ₃₇₃

Table 1: Names and sequences characteristics of the different sliding clamps used in this study. *Ec*: *Escherichia coli*; *Mt*: *Mycobacterium tuberculosis*; *Bs*: *Bacillus subtilis*; wt: wild type sliding clamps; M1, M2, M3, G1 and G2: mutant sliding clamps.

The dissociation constant measured for $^{Ecwt}SC/P_7$ interaction increases from 100 to 450 nM between 15 and 34°C (**Table 2, Figure 1A and SI.6**). For the $^{Mtwt}SC/P_7$ interaction, the Kd is about 20 to 40 times lower (between 4 and 7 μM), while the lowest affinity is observed for ^{Bswt}SC , with Kd values ranging from 40 to 80 μM range.

All these data confirm our previous observations and show that **P₇** binding greatly vary with the SC bacterial origin¹⁸. To investigate the molecular basis of these differences, we constructed several mutants of each SC where the residues corresponding to ^{Ecwt}SC S₃₄₆ and ^{Ecwt}SC M₃₆₂ are singly (M1, M2, G1 and G2 respectively) or both (M3) mutated (**Table 1**). The interaction of **P₇** with these SC variants was analyzed using ITC, crystal structures, *in vitro* replication assays and molecular modeling analyses.

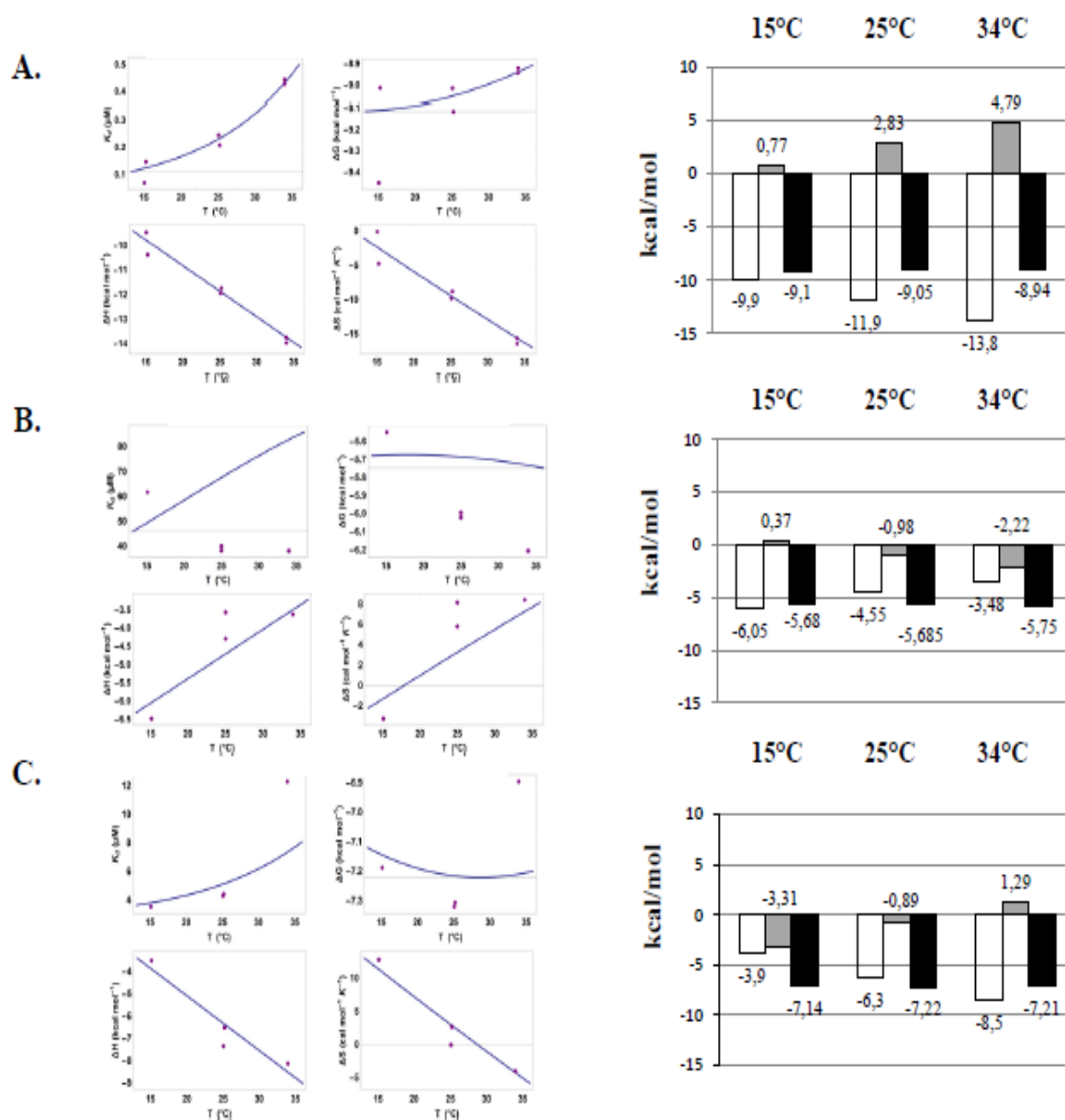


Figure 1: Thermodynamic analyses of P_7 interaction with $EcwtSC$ (A), $BswtSC$ (B) and $MlwtSC$ (C) at different temperatures. Global thermodynamic treatment (GTT) (left column): The curves represent the functions obtained by treating simultaneously all experimental ITC data at three temperatures. Dots represent the results of a classical treatment of each experimental ITC data processed individually. For weak interaction, such as $BswtSC/P_7$, the gap between individual treatment and the theoretical curve obtained by GTT is more important. Thermodynamics profiles (right column) displaying ΔH , $-T\Delta S$ and ΔG at three temperatures were obtained from the GTT derived curves. ΔH , $-T\Delta S$ and ΔG (all in kcal/mol) are white, grey and dark bars, respectively. All ITC data are presented in **SI.6** and typical ITC titration curves are shown in **SI.7**. ITC control experiments at three temperatures are presented in **SI.4**.

		15°C (288.15 K)	25°C (298.15 K)	34°C (307.15 K)
$E_{\text{cwt}}^{\text{SC}}$	Classical processing	0.09 (\pm 0.04)	0.16 (\pm 0.05)	0.42 (\pm 0.04)
	GTT processing	0.12	0.22	0.44
$B_{\text{swt}}^{\text{SC}}$	Classical processing	24 (\pm 18)	14 (\pm 0.6)	43(\pm 49)
	GTT processing	49	68	82
$M_{\text{rwt}}^{\text{SC}}$	Classical processing	3.5 (\pm 3)	5 (\pm 3.9)	7.23 (\pm 1)
	GTT processing	4	5	7

Table 2: Dissociation constants (K_D : μM) of the P_7 interaction with the three natural SC at different temperatures. The large standard deviations observed for $B_{\text{swt}}^{\text{SC}}$ and $M_{\text{rwt}}^{\text{SC}}$ are indicative of a poor interaction. Independent experiments were performed two to four times. All results were obtained through classical or GTT processing of experimental data, as indicated. Full thermodynamic data are in **SI.6** and examples of ITC titration curves are presented in **SI.7**.

Mutations $S_{346}P$ and $M_{362}L$ in $E_{\text{cwt}}^{\text{SC}}$ drastically modify the thermodynamics of SC/ P_7 interaction.

Complete thermodynamic profiles of P_7 interaction with the various SC mutants are presented in **SI.8** and **SI.11**. **Figure 2** presents a comparison of the thermodynamics profiles of the P_7 interaction with $E_{\text{cwt}}^{\text{SC}}$ and the double mutant $E_{\text{cM3}}^{\text{SC}}$. Details on the specific effects induced by single mutations are developed in **SI.11**. Introduction of the $S_{346}P$ and $M_{362}L$ mutations in the $E_{\text{cwt}}^{\text{SC}}$ binding pocket results in a large reduction of $|\Delta H|$ of about +6 kcal/mol at 15°C and +10 kcal/mol at 34°C. The resulting ΔC_p drops from -210 ($E_{\text{cwt}}^{\text{SC}}$) to -50 cal/mol/deg for $E_{\text{cM3}}^{\text{SC}}$, indicative of a change in the mode of interaction³³. As opposed to what is observed with $E_{\text{cwt}}^{\text{SC}}$, the entropic component ($-T\Delta S$) becomes favorable (\approx -2 to -3 kcal/mol) to the $E_{\text{cM3}}^{\text{SC}}/P_7$ interaction at all temperatures. The resulting free energy of interaction, ΔG , is stabilized around -6 to -7 kcal/mol, *i.e.* about 2-3 kcal/mol higher than those observed for $E_{\text{cwt}}^{\text{SC}}$, at all temperatures. Consequently, the mutations $S_{346}P$ and $M_{362}L$ in $E_{\text{cwt}}^{\text{SC}}$ trigger a drastic change in the P_7 mode of interaction, underlying the significant contribution of these residues to the efficient binding of P_7 on $E_{\text{cwt}}^{\text{SC}}$.

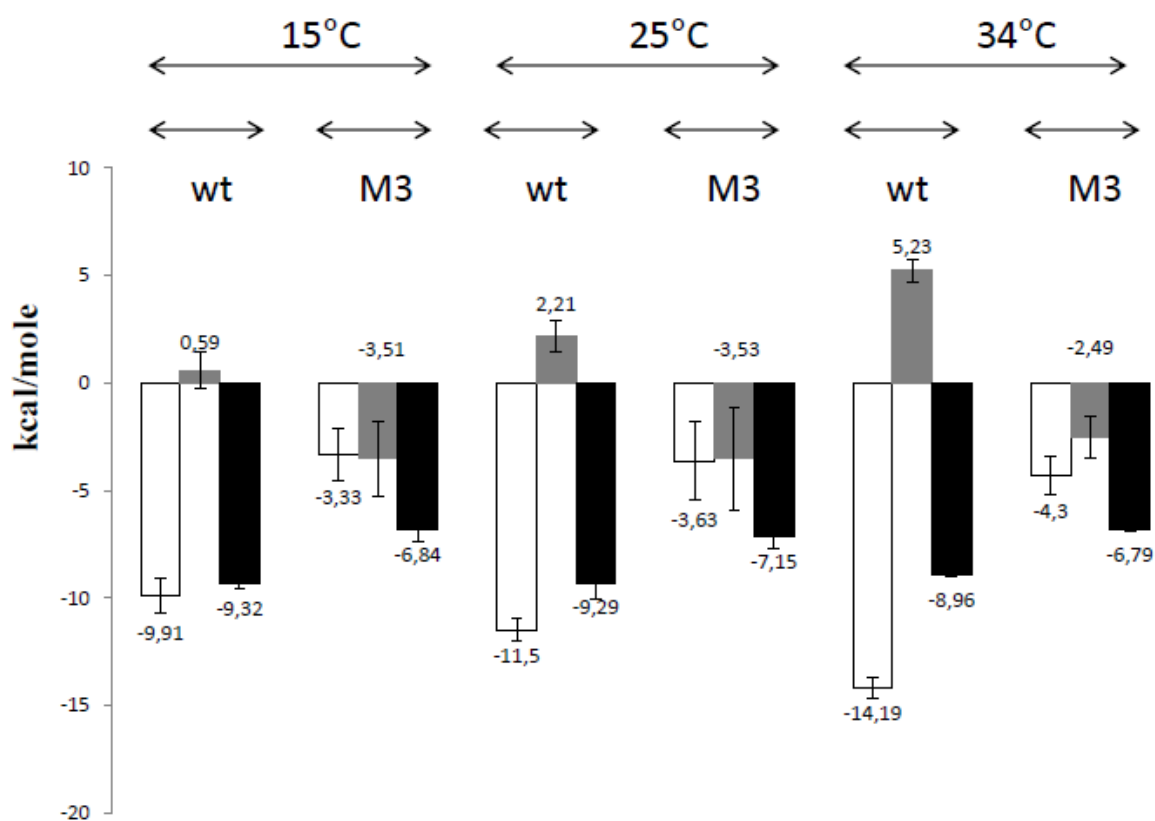


Figure 2: Thermodynamic profiles of P₇ interactions with ^{Ecwt}SC (wt) and ^{EcM3}SC (M3) at different temperatures. ΔH, -TΔS and ΔG (all in kcal/mol) are white, grey and dark bars, respectively. Data are means of at least two independent experiments. For the sake of comparison, all results are obtained through classical processing of experimental data (SI.6).

^{EcM3}SC and ^{Bswt}SC present similar thermodynamic profiles upon P₇ interaction.

Because the two mutations introduced in ^{EcM3}SC are equivalent to the natural residues found in ^{Bswt}SC (Table 1), we also compared the thermodynamic profiles obtained for P₇ interaction with ^{EcM3}SC and ^{Bswt}SC (Figure 3). These profiles are very similar at all temperatures, confirming that the introduction of S₃₄₆P and M₃₆₂L mutations in ^{Ecwt}SC is sufficient to convert the *E. coli* specific thermodynamic profile into that observed with ^{Bswt}SC. For both types of interactions, the enthalpic contribution is equivalent, ranging from -3 to -4 kcal/mole, and in both cases, the entropic contributions are favorable to the interaction, around -3 kcal/mol. The large variations of both parameters result from the poor efficiency of the peptide binding. The resulting free energy of interaction is slightly more favorable for ^{EcM3}SC/P₇ than for ^{Bswt}SC/P₇, suggesting that besides the contributions of S₃₄₆ and M₃₆₂

residues, the local environment of the Ec SC pocket stabilizes the peptide interaction more significantly than that of Bswt SC.

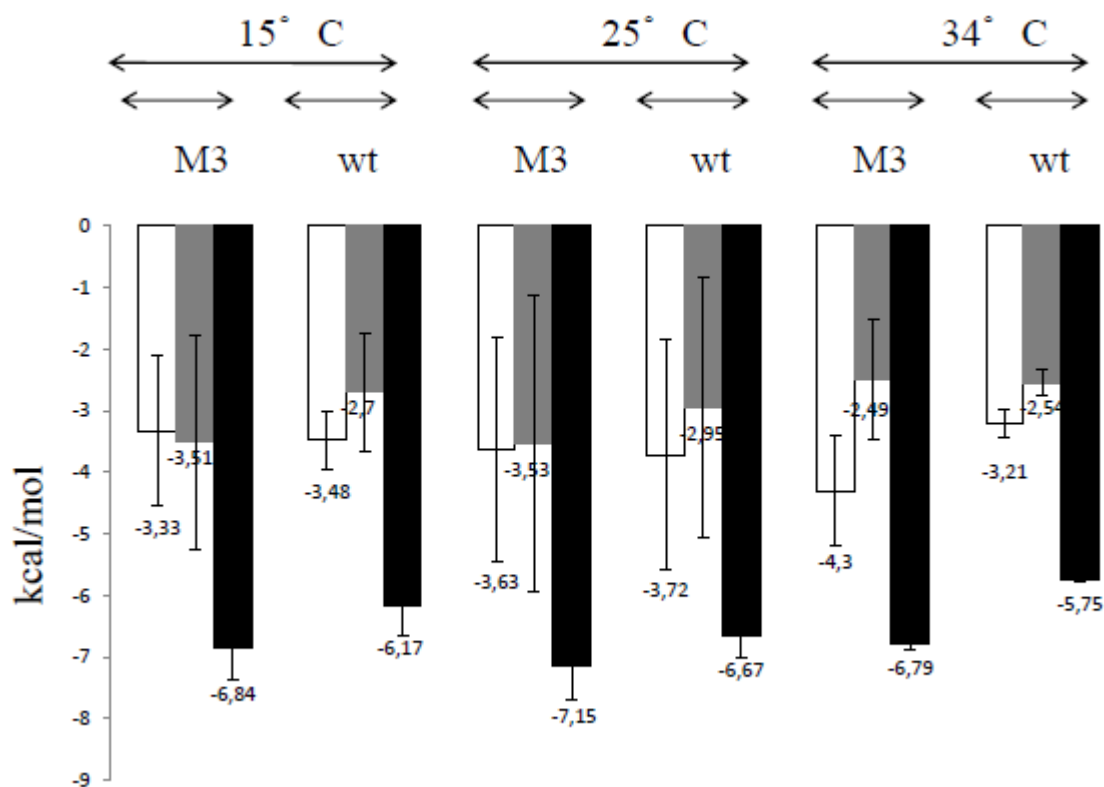


Figure 3: Thermodynamic profiles of P₇ interactions with EcM3 SC (M3) and Bswt SC (wt) at different temperatures. ΔH, -TΔS and ΔG (all in kcal/mol) are white, grey and black bars, respectively. Data are means of at least two independent experiments. Results are obtained through classical processing of experimental data (SI.6).

Mutations P₃₅₇S and L₃₇₃M residues in Bswt SC improve P₇ interaction.

We also analyzed the effects of the introduction of *Ec* natural residues (S₃₄₆ and M₃₆₂) in the *Bs* context (**Figure 4, SI.8 B and SI.11 B** for details on single mutations effects on P₇ interaction). As opposed to the major deleterious effect on complex formation triggered by mutations introduced in Ecwt SC (**Figure 2**), the mutations of the corresponding residues in Bs SC improve the efficiency of the interaction with P₇, as indicated by the systematic increase of ΔG values with BsM3 SC as compared to Bswt SC (**Figure 4**). However, the effect is limited as these ΔG values remain about 2 kcal/mole lower than those measured for Ecwt SC. This is in agreement with the above mentioned observation that the Bs SC pocket might be globally less favorable to P₇ binding. Consequently, as observed for the *Ec* context (SI.11 A) but with opposite effects, these mutations differentially trigger a change in the thermodynamic profiles

of the interaction (**SI.11 B**). At all temperatures, the P₃₅₇S mutation (mutant ^{BsM1}SC) triggers a reduction of the ΔH values, as compared to those measured in ^{Bswt}SC, and an increase of the entropic factor favorable contribution. Alternatively, the L₃₇₃M mutation (mutant ^{BsM2}SC) increases the ΔH value by -3 to -5 kcal/mol at each temperature and drastically reduces the entropic contribution, which becomes unfavorable at high temperature.

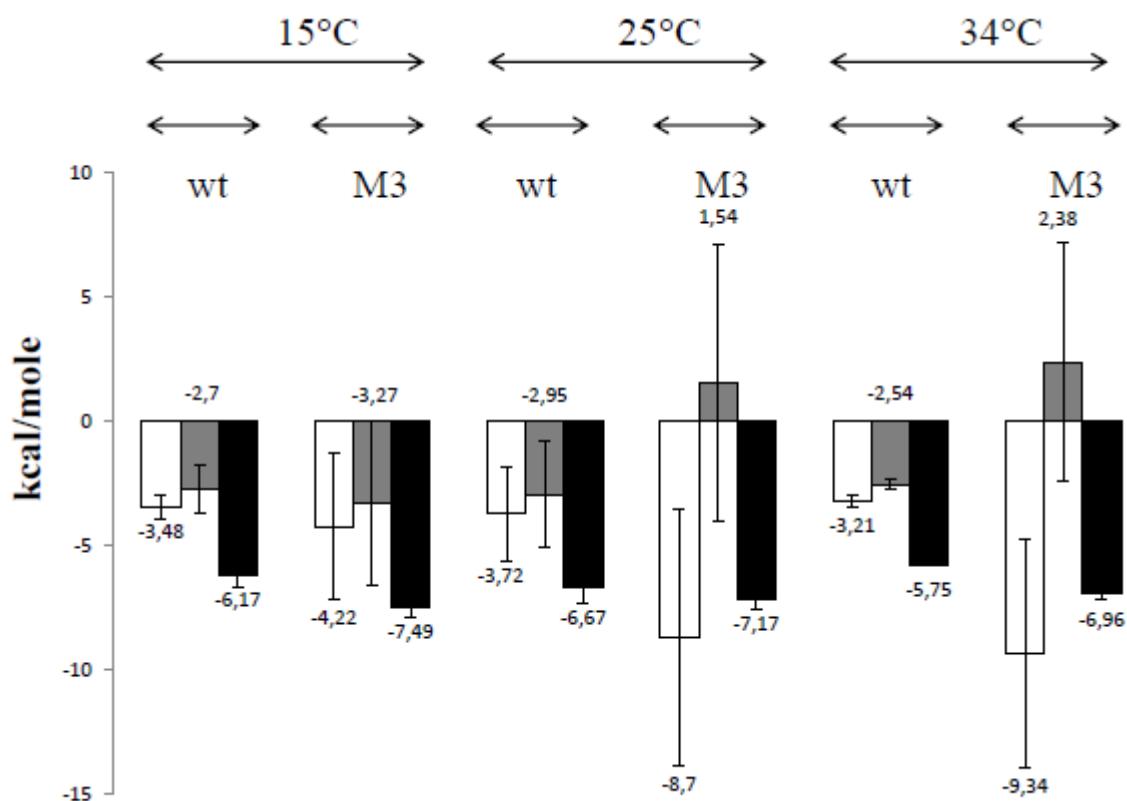


Figure 4 : Thermodynamic profiles of P₇ interactions with ^{Bswt}SC (wt) and ^{BsM3}SC (M3) at different temperatures. ΔH , $-T\Delta S$ and ΔG (all in kcal/mol) are white, grey and black bars, respectively. Data are means of at least two independent experiments. Results are obtained through classical processing of experimental data (**SI.6**).

The added contribution of both mutations, measured with mutant ^{BsM3}SC, combine the effects of each individual mutation. As a result, the ΔH evolution becomes strongly dependent on temperature and the entropic contribution turns to be unfavorable at 34°C for both M2 and M3 mutants (**SI.11 B**), but already at 25°C for the double mutant, which suggests a synergistic effect of both mutations (**Figure 4**). Indeed, the ΔC_p value determined for ^{BsM3}SC (-270 cal/mole/deg) is closed to that measured for ^{Ecwt}SC (-210 cal/mole/deg), whereas the values calculated for ^{BsM1}SC and ^{BsM2}SC are -10 and -40 cal/mol/deg, respectively. These

observations show that the two mutations, P₃₅₇S and L₃₇₃M, induce a complete change of the thermodynamic profile of the ^{Bs}SC/P₇ interaction. Moreover, in the two different pocket environments, *Ec* and *Bs*, the drastic changes in thermodynamic profiles are triggered by the M2 mutations (**Figure SI.11 A and B**) thus identifying a strategic position for what concerns the peptide interaction. The position of M1 mutations is not that crucial for the complex formation.

Specific contributions of the strategic residues to the P₇ interaction.

To further analyze the specific contributions of the S₃₄₆ (or P₃₄₆) and M₃₆₂ (or L₃₆₂) residues to the P₇ interaction, we compared the thermodynamic data obtained for ^{wt}SC (or ^{M1}SC or ^{M2}SC) with those obtained with ^{G1}SC and ^{G2}SC (**Table 1**), where the natural residues are replaced by glycine (**Figure 5 and SI.12**). Note that thermodynamic data obtained with the G mutants directly provide the whole pocket contribution to the interaction, excluding the S/P and M/L side chain effects. The specific contributions of the S/P and M/L residues to each thermodynamic parameter are obtained by the difference between the parameter measured for ^{wt}SC and that measured for the ^{Gn}SC (**SI.12**). The results are presented in **Figure 5**.

In the *Ec* context, the S₃₄₆ residue does not contribute significantly to the P₇ interaction, as indicated by the weak ΔG value, particularly at high temperatures (**Figure 5A**, left). The ΔH value reveals an endothermic contribution, while the favorable entropic contribution increases with temperature and might be linked, at least partly, to water release upon peptide interaction, as observed in crystal structures⁵. A very different profile is observed for the M₃₆₂ residue (**Figure 5A**, right), with essentially an exothermic specific contribution, no entropic effect and a constant Gibbs free energy (-1 kcal/mole) over the temperature range. Thus this residue contributes for about one tenth of the overall interaction of the peptide (see **Figure 1A**). Mutation S₃₄₆P (**Figure 5B**, left) presents a similar profile as that observed for the natural residue, with a very weak contribution to binding. In contrast, the profile observed for the M₃₆₂L mutation (**Figure 5B**, right) reveals how detrimental is the L residue at this position to the P₇ interaction in the *Ec* context. All together, these residue specific profiles reveal that position 362 in ^{Ec}SC significantly contributes to the binding, while position 346 does not. In addition, the nature of the residue, M vs L, at this very position appears to be crucial for the binding.

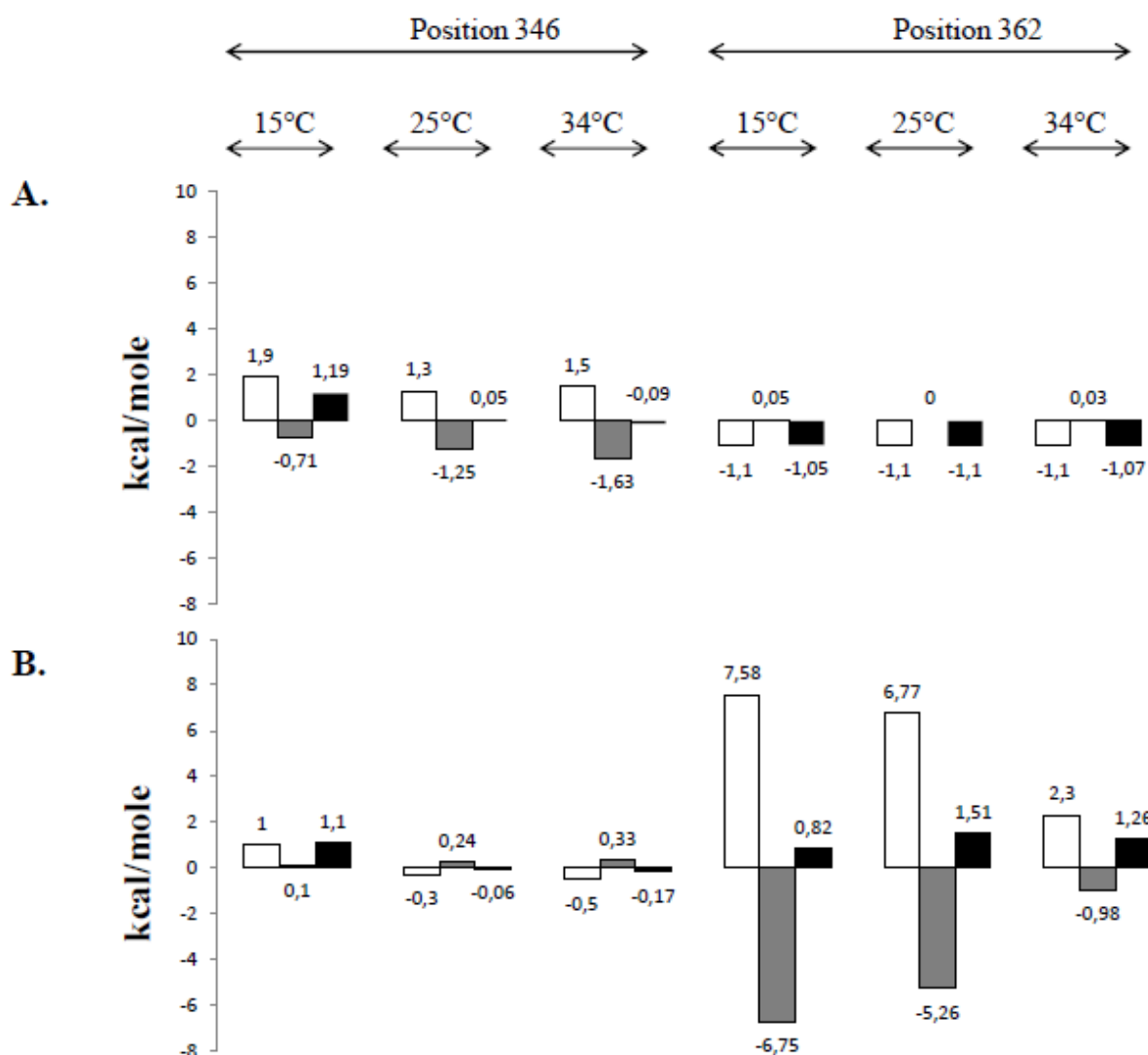


Figure 5: Specific contribution to the P₇ interaction of residues at positions 346 and 362 in ^{Ec}SC. (A): contribution of natural residues S₃₄₆ (left) and M₃₆₂ (right). The residue specific profiles result from the difference between thermodynamic values for ^{Ec}wt and ^{Ec}G1 (left) or ^{Ec}G2 (right). **(B):** contribution of non natural residues P₃₄₆ (left) and L₃₆₂ (right). The residue specific profiles result from the difference between thermodynamic values for ^{Ec}M1 and ^{Ec}G1 (left) or ^{Ec}M2 and ^{Ec}G2 (right) (SI.12). ΔH , $-T\Delta S$ and ΔG (all in kcal/mol) are white, grey and black bars, respectively. ΔH and $-T\Delta S$ numerical data are retrieved from GTT curves (SI.5), except for ^{Ec}M2 and ^{Ec}G2 (SI.6).

A similar approach aiming at defining the specific contributions of the same residues in the ^{Bs} context was hardly feasible as the large variations in thermodynamic data obtained with ^{Bs}SC (Figure 1 and 4, SI.6) preclude one to retrieve any reliable conclusion on the contribution of the P/S and L/M residues in this context. Nevertheless, the thermodynamic profiles obtained with ^{Bs}G1SC and ^{Bs}G2SC provide information on the contribution of the

exchanged residue (**Figure SI.12 B**). As for the S₃₄₆ residue in the *Ec* context, the P₃₅₇ residue does not seem to contribute to the P₇ interaction as indicated by the similarity of profiles observed with ^{B_swt}SC and ^{B_sG1}SC (**Figure SI.12 B**). On the contrary, the profiles obtained for the ^{B_sG2}SC/P₇ interaction reveal that the L residue, as observed for the M₃₆₂ residue in the *Ec* context, occupies a strategic position to impede the efficient interaction of P₇ with ^{B_swt}SC, as indicated by the fact that the L₃₇₃G mutation restores an interaction profile similar to that observed for ^{Ec}SC (**Figure SI.12B**). Indeed, the ΔC_p value calculated for the ^{B_sG2}SC/P₇ interaction is -226 cal/mol/deg, equivalent to that measured with ^{Ecwt}SC.

Specific contribution of the binding pocket to the P₇ interaction.

Beside these strategic positions, the whole pocket largely contributes to an efficient ligand binding, as revealed by the thermodynamic profiles obtained with G1 or G2 mutants (**SI.12**). In the *Ec* context, the pocket-specific thermodynamic profiles are only slightly modulated by the S₃₄₆G or M₃₆₂G mutation (**Figure SI.12 A**), while in the *Bs* context, the profiles obtained with G1 and G2 are very different (**Figure SI.12 B**). The G1 profile characterizes a weak interaction similar to that observed with ^{B_swt}SC. In contrast, the G2 profile, similar to those observed for the *Ec* context, reveals that the *Bs* pocket environment is also suitable for peptide binding, provided the L₃₇₃ side chain is removed. This observation confirms the strategic position of the *Bs*₃₇₃ (or *Ec*₃₆₂) residue for the control of peptide interaction and the deleterious effect of L for P₇ binding. However, as indicated by the K_D values (**SI.6**), P₇ interaction with ^{B_sG2}SC remains weaker than with ^{EcG2}SC (**SI.12**). Moreover, the introduction of a M₃₇₃ residue (^{B_sM2}SC) fails to restore an interaction as strong as that observed in the *Ec* context (**Figure 4** and **SI.11**), which may suggest that the side chain of this M residue is not adapted to the *Bs* pocket environment and that subtle pocket residue to residue interactions determine an optimal ligand interaction¹⁸.

In conclusion, thermodynamic analyses reveal that, for both *Ec* and *Bs* contexts, the same strategic position, corresponding to *Ec*M₃₆₂ or *Bs*L₃₇₃, controls the efficient interaction of the peptide with the SC binding pocket. Residues at position corresponding to *Ec*S₃₄₆ (or *Bs*P₃₅₇) are not actively contributing to the binding. Finally, both pockets contribute to the interaction but with different efficiency.

Mode of peptide interaction as deduced from the ΔC_p analysis.

ΔC_p analysis provides insights into the peptide mode of interaction with SC. It has been previously shown that a $\Delta C_p \neq 0$ is partly linked to a variation in the hydration pattern and to a variation in the solvent accessible surface area (ASA) of the protein³⁴. A ΔC_p^{calc} value can be retrieved by measuring the ASA variation from the crystal structure of the $^{E\text{cwt}}\text{SC}/\mathbf{P}_7$ complex (**SI.9**), as compared to the structure of the peptide free $^{E\text{cwt}}\text{SC}$ (PDB 1OK7 chain A). This approach characterizes the ΔC_p value resulting from the burying of amino acid residues upon peptide binding and is calculated using the empiric formula: $\Delta C_p^{\text{calc}} = 0.27 \Delta A_{\text{aromatic}} + 0.40 \Delta A_{\text{nonaromatic}}$ ³⁵. We also took into account the specific contribution of the free \mathbf{P}_7 by considering that the residues Q, L and F were fully buried and Cha, which stacks on the platform {Wolff, 2011 #60} was only 50% hidden after binding. As indicated by the crystal structures, the D residue, which points toward the solvent, does not contribute to the interaction. Accessible surfaces area values for each \mathbf{P}_7 residue were determined according to {Lins, 2003 #122}. A value of -276 cal/mole/deg is calculated for $^{E\text{cwt}}\text{SC}/\mathbf{P}_7$ which is close to the ΔC_p^{ITC} (-213 cal/mol/deg) considering the approximations on \mathbf{P}_7 . Similar close values are obtained for $^{E\text{cM1}}\text{SC}/\mathbf{P}_7$ ($\Delta C_p^{\text{ITC}} = -240$ cal/mole/deg and $\Delta C_p^{\text{calc}} = -281$ cal/mole/deg).

In absence of any $^{B\text{s}}\text{SC}/\mathbf{P}_7$ complex structure, such information is not available for the $B\text{s}$ context. However, the profiles obtained with $^{B\text{s}G2}\text{SC}$ (**SI.12 B**) yields a ΔC_p value of -226 cal/mol/deg, very close to that measured with $^{E\text{cwt}}\text{SC}$. It suggests that this mutated pocket binds \mathbf{P}_7 according to an induced-fit process and that the low ΔC_p values calculated for $^{B\text{swt}}\text{SC}$ and $^{B\text{sM1}}\text{SC}$ is more indicative of a weak interaction, due to the presence of the L residue, than of a different binding mode. Interestingly, the ΔC_p value measured for $^{B\text{sM2}}\text{SC}$ (-42 cal/mole/deg) remains weak, despite the L₃₇₃M mutation. Only for the double mutant $^{B\text{sM3}}\text{SC}$ is the ΔC_p value (-269 cal/mole/deg) in the range of that measured for $^{B\text{s}G2}\text{SC}$. This may reflect the fact that, in contrast to G₃₇₃, introducing a single M₃₇₃ residue in the $^{B\text{s}}\text{SC}$ pocket does not trigger a large structural change, presumably because of some steric hindrance between the M₃₇₃ side chain and the P₃₅₇ residue (or others). Only the double mutant $^{B\text{sM3}}\text{SC}$ allows this structural change, as indicated by the corresponding ΔC_p value, because the S₃₅₇ and M₃₇₃ side chains move along, as observed in $^{E\text{cwt}}\text{SC}$. This observation reveals that a subtle network of residue to residue interactions operate specifically in each pocket, as we noted before¹⁸, to ensure an optimal binding process.

Structure analysis of mutant SC/P₇ complexes.

We solved the structures of different complexes, namely ^{Ecwt}SC/P₇ (PDB ID: 6FVL), ^{EcM1}SC/P₇ (PDB ID: 6FVM), ^{Mtwt}SC/P₇ (PDB ID: 6FVN) and ^{MtM1}SC/P₇ (PDB ID: 6FVO) (**SI.9**). Until now, no structure of a SC/P₇ complex has been obtained using ^{EcM2}SC, ^{EcM3}SC or any ^{Bs}SC. Although crystals were obtained with these SC, no density corresponding to the peptide was observed in the electron density map. This observation reflects the weak peptide interaction, in agreement with thermodynamic and biochemical data, and suggests that P₇ is mobile in these specific complexes, or that the crystallization process has selected the peptide-free SC. The overlay of several complexes structures is presented in **SI.13**. ^{Ecwt}SC/P₇ and ^{EcM1}SC/P₇ complexes align nicely, yielding an rmsd = 0.37 Å over 327 C_α atoms (**Figure SI.13 A**). ^{Mtwt}SC/P₇ and ^{MtM1}SC/P₇ complexes also superimpose correctly, with an rmsd= 0.89 Å over 330 C_α atoms (**Figure SI.13 C**). Not surprisingly, ^{Ecwt}SC/P₇ and ^{Mtwt}SC/P₇ do not readily superimpose as indicated by an averaged rmsd = 1.9 Å over 280 C_α atoms (**Figure SI.13 B**). Nevertheless, the two P₇ peptides superimpose correctly. In all cases, we observed a full overlap of the ^{Ecwt}S₃₄₆ and ^{Ecwt}M₃₆₂ or their corresponding residues in ^{Mtwt}SC, with their cognate residues in the M1 mutant SC. This indicates that the conversion of this residue (from S to P, or the opposite) does not alter the pocket conformation, in agreement with the ITC and biochemical data for the *Ec* replicative system (**SI.11A** and **Figure 11** below).

Kinetics analysis of the P₇ binding onto SC.

The treatment of the ITC injection curves by the kinITC program^{20 32} provides the kinetic rate constants that govern the peptide interaction (**Figure SI.14 A and B**). **Figure 6** shows the Arrhenius plots for the association and dissociation constants obtained at different temperatures for the interaction of P₇ with ^{Ecwt}SC and ^{EcM1}SC. No kinetic information could be obtained, neither for ^{EcM2}SC/P₇ and ^{EcM3}SC/P₇ complexes formation (**SI.14 C and D**), nor for ^{Bswt}SC/P₇ and ^{Mtwt}SC/P₇ (data not shown), due to too weak signals.

The Arrhenius plots describing the ^{Ec}SC/P₇ interaction (**Figure 6**) reveal a significant difference between the two complexes: an unusual negative ΔH_{on}^{\ddagger} is observed for the ^{EcM1}SC/P₇ complex formation because the ON-rate decreases upon temperature increase. This can be interpreted in the frame of a two-step kinetic model, which was described in great

details in two seminal papers on DNA helix formation. A two steps process has also been proposed previously for the ligand-SC system. According to this model, an initial but labile binding of \mathbf{P}_7 by one of its two anchors, the N-ter Q residue or the C-ter LF region, is followed by a fast step, conditioned by the first one, and corresponding to the full binding of the peptide into the SC pocket. Therefore, the observed k_{on} is a global rate constant for the two successive steps, which explains that the global $\Delta H_{\text{on}}^\ddagger$ may be negative. A smaller effect of the $S_{346}P$ mutation is also observed for the $\Delta H_{\text{off}}^\ddagger$ which is reduced four times with ^{EcM1}SC as compared to ^{Ecwt}SC . All together, these effects suggest that the $^{Ecwt}SC/\mathbf{P}_7$ interaction proceeds through an induced-fit mechanism, in agreement with the ΔC_p analysis, and that the mutation affects the binding-induced dynamics of the pocket but not the overall efficiency of binding, as indicated by the similar thermodynamic data obtained with ^{Ecwt}SC and ^{EcM1}SC (SI.6).

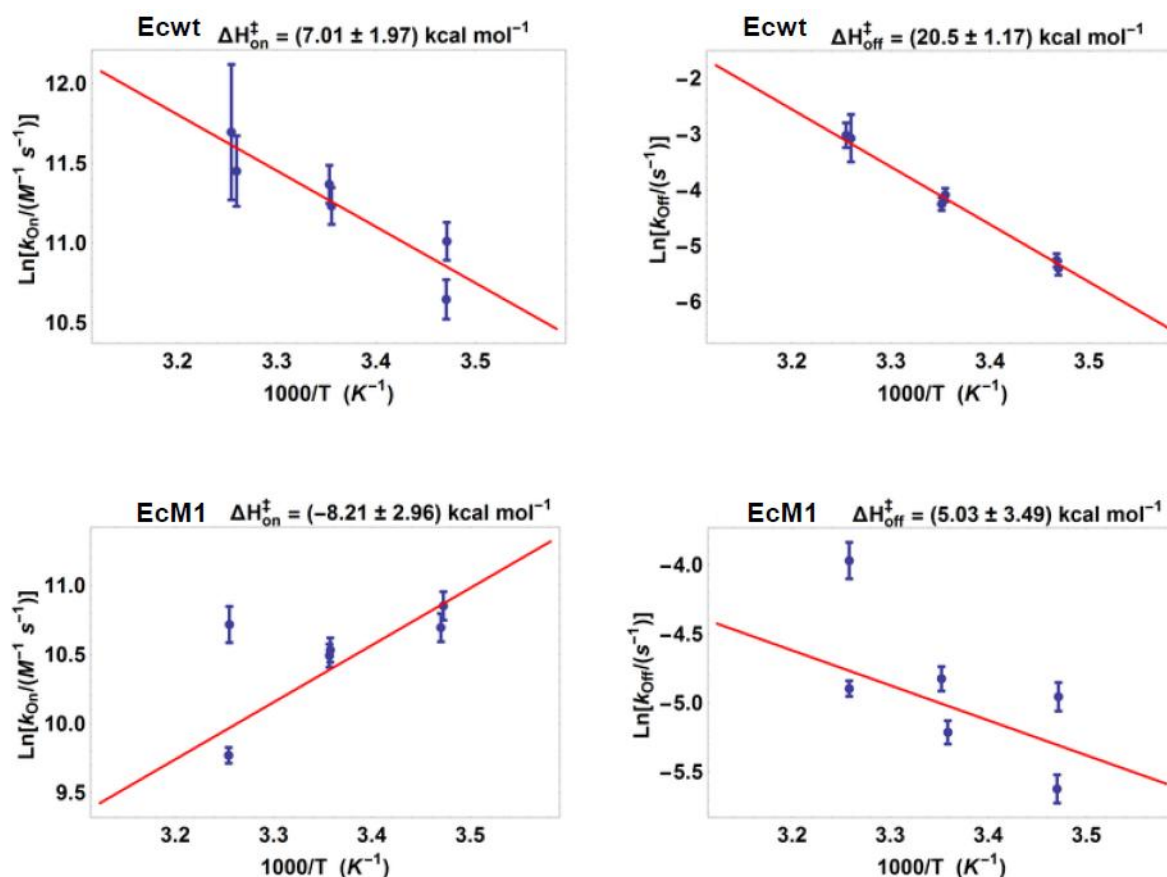


Figure 6: Arrhenius plots for the association and dissociation constants of $^{Ecwt}SC/P_7$ and $^{EcM1}SC/P_7$ complexes. k_{on} and k_{off} values measured for the $^{Ecwt}SC/P_7$ interaction are 3.4, 5.4 and 7.1 $10^4 M^{-1}.s^{-1}$ and 4.8, 15 and 47 $10^{-3} s^{-1}$ at 15, 25 and 34°C, respectively. Values measured for the $^{EcM1}SC/P_7$ interaction are 3.2, 2.4 and 2.1 $10^4 M^{-1}.s^{-1}$ and 5.3, 6.7 and 13 $10^{-3} s^{-1}$.

Molecular modeling studies.

Molecular dynamics simulations were performed to elucidate how the nature of residues $^{Ec}M/L_{362}$ and $^{Bs}L/M_{373}$ modulate the ligand interaction, as indicated by the ITC analyses. **Table 3** compares the results of ITC-derived experimental and calculated $\Delta\Delta G$ values. The simulation data are in good agreement with the experimental values and reveal that the $^{Ec}L_{362}$ residue is highly detrimental to the interaction while the $^{Bs}M_{373}$ residue slightly favors the complex formation, in full agreement with our ITC analyses. The simulated structures of the different complexes ($^{Bswt}SC/P_7$ and $^{EcM2}SC/P_7$ on one hand and $^{Ecwt}SC/P_7$ and $^{BsM2}SC/P_7$ on the other hand) were superimposed (**SI.15**). In both cases, the peptides are nicely aligned (rmsd (C_α) = 0.38 Å), while the two SC chains present a looser alignment (rmsd = 1.33 Å over 305 atoms). Nevertheless, the superimposition reveals that in ^{EcM2}SC and ^{Bswt}SC , the $L_{362/373}$ residue positions immediately below the peptide's third residue (**SI.15 A**). This position blocks the path where the peptide can lie and results in a poor interaction, as observed by ITC for ^{EcM2}SC and ^{Bswt}SC . In contrast, the M_{373} residue of ^{BsM2}SC adopts

Data for peptide binding	$\Delta\Delta G^{^{EcM2}SC \text{ vs } ^{Ecwt}SC}$	$\Delta\Delta G^{^{BsM2}SC \text{ vs } ^{Bswt}SC}$
Experimental (ITC)	$2,8 \pm 1.1$	$-0,3 \pm 0.5$
Calculated	$3,7 \pm 0,7$	$-0,4 \pm 0,04$

Table 3: Comparison of calculated and experimental $\Delta\Delta G$ values. $\Delta\Delta G$ (kcal/mole) results from the comparison between $\Delta G^{M2}SC$ and $\Delta G^{wt}SC$ (namely, $\Delta G^{M2}SC - \Delta G^{wt}SC$) at 25°C. When positive, binding to mutant is less favorable. In ^{Ec}SC , M2 corresponds to a $M_{362}L$ mutation (**Table 1**). In ^{Bs}SC , M2 corresponds to a $L_{373}M$ mutation (**Table 1**).

a position similar to the M_{362} position in ^{Ecwt}SC (**SI.15 B**), i.e a shift of its side chain that clear the way for the peptide to lie within the resulting groove, thus allowing an improved interaction.

Biochemical analysis of the various SC activities.

We used a primer elongation assay to analyze the biochemical properties of the various *Ec*SC and *Bs*SC to promote SC dependent polymerization by *in vitro* reconstituted cognate and non-cognate replicative DNA polymerase complexes. In this assay, the interaction between the replicase and the clamps is mediated by the polymerase specific peptide, namely QADMF for the *E. coli* polIII holoenzyme³⁶ and QLSLF for the *B. subtilis* PolC¹⁶. The ability of **P₇** to interact with these various SC in this biochemical tests is evaluated in competition assays.

As shown in **Figure 7**, both *Ec*^wSC and *Bs*^wSC associate with their cognate *in vitro* reconstituted *E. coli* (EcP) and *B. subtilis* (BsP) DNA polymerases to promote a SC dependent primer elongation. The reaction is concentration dependent and saturation is reached at equimolar concentrations of SC and polymerases (100 fmole/reaction). The elongation profiles show distinct bands of arrest specific for each type of polymerase, which may result from a different sensitivity of each complex for DNA sequences or secondary structures. Neither the *E. coli* nor the *B. subtilis* replicative complexes (comprised of at least the DNA polymerase catalytic subunit and the SC loading complex) are able to form productive complexes with the non cognate SC (**SI.16**), as previously observed by others when *E. coli* (Gram-) and *S. pyogenes* (Gram+) replicative machineries were analyzed³⁷. This underlines the specificity of interaction between the partners, mediated by the specific peptides but also others polymerase-SC contacts such as described for *Ec* PolIV¹⁷ {Beuning, 2006 #123}.

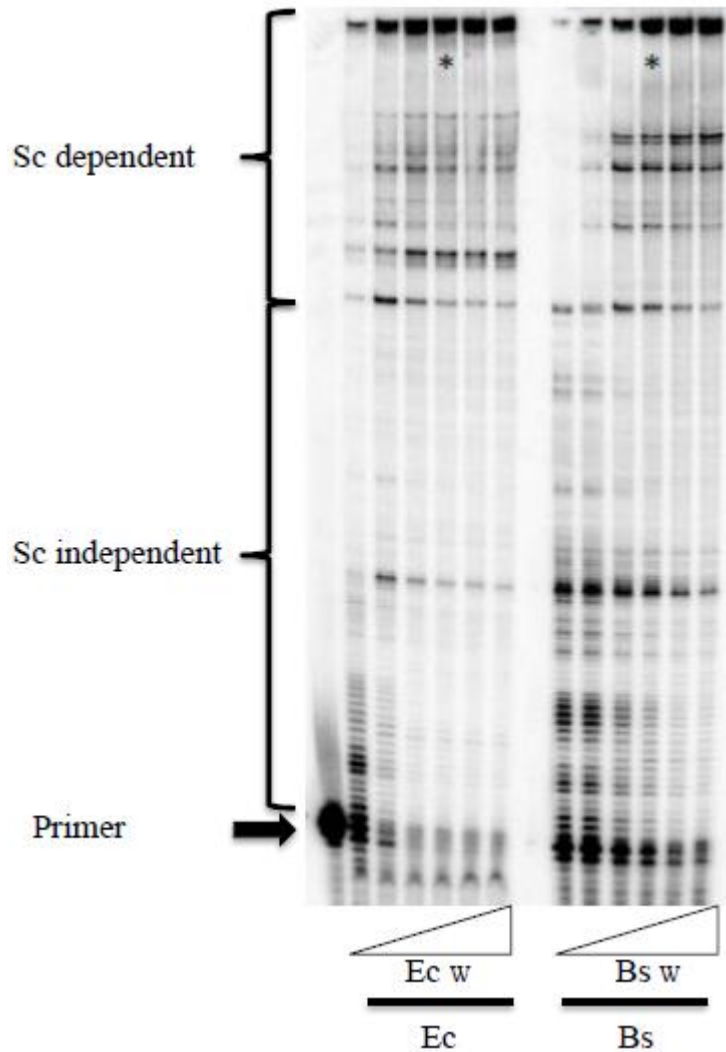


Figure 7 : Elongation profiles obtained with reconstituted *E. coli* and *B. subtilis* holoenzymes. The *E. coli* and *B. subtilis* polymerases (EcP or BsP, 100 fmoles) are supplemented with various amount of the cognate ^{wt}SC (from 0 to 400 fmoles). The labeled lanes (*) correspond to 100 fmoles ^{wt}SC. The primer position is indicated by the arrow.

Then we analyzed how EcP and BsP behave when associated with the different cognate mutant SC (**Figure 8**). Holoenzyme formed between EcP and ^{EcM1}SC (**Figure 8**, lane 4) displays the same primer elongation profile as that observed using ^{Ecwt}SC (**Figure 8**, lane 3). This indicates that the natural EcP peptide interaction is not altered by the S₃₄₆P mutation, nor in terms of elongation efficiency, as indicated by the equal bands intensities in lanes 3 and 4, nor in terms of specificity as indicated by their similar elongation profiles. This result is in agreement with the P₇ interaction data obtained in ITC experiments (**Figure SI. 8 A** and **S.I 11 A**). In contrast, holoenzymes formed by association of EcP with ^{EcM2}SC or ^{EcM3}SC (**Figure 8**, lanes 5 and 6) present a profile different from that observed for the natural holoenzyme,

which can be accounted for by a decrease in processivity and/or modifications of polymerase-SC interaction dynamics. This is in agreement with the ITC data and indicates that, regardless of the peptide, the M₃₆₂L mutation is deleterious to the interaction. The combination of both mutations (^{EcM3}SC, lane 6) shows a synergistic effect, as revealed by the decrease in full length band intensity in lane 6 as compared to lane 5. Surprisingly enough, these profiles, and particularly that obtained with ^{EcM3}SC, are very similar to that obtained with the natural BsP holoenzyme (**Figure 8**, lane 8).

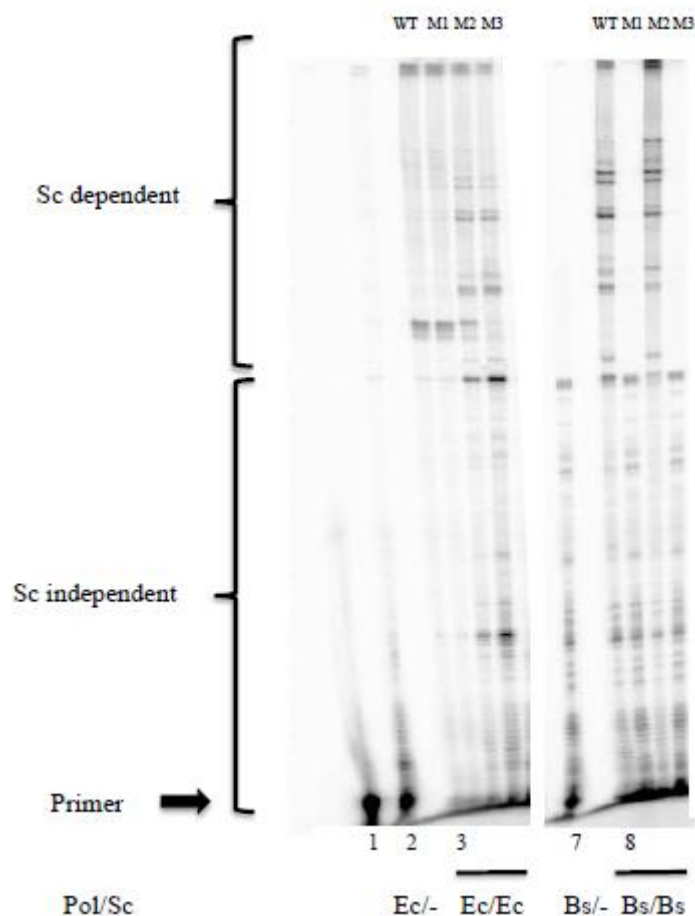


Figure 8: Elongation assays with the different ^{Ec}SC and ^{Bs}SC. EcP or BsP reconstituted polymerases (100 fmoles) were each tested with the various cognate SC (lanes 3 to 6 and lanes 8 to 11 for Ec and Bs polymerases, respectively) (100 fmoles) (**Table 1**). The primer position is indicated by the arrow (lane 1). Lanes 2 and 7 display the activity of Ec and Bs polymerases in absence of the cognate SC.

We observed that only ^{BsM2}SC associates productively with BsP and yields a similar pattern of elongation as the natural BsP holoenzyme (**Figure 8**, lane 10), while nor ^{BsM1}SC neither ^{BsM3}SC form a productive complex (**Figure 8**, lane 9 and 11). In this Gram+ context,

the P₃₅₇S mutation appears to be highly deleterious to the polymerase-SC interaction, as opposed to what is observed with the *Ec*SC, while the L₃₇₃M mutation is harmless and even favorable to the interaction, as indicated by the increased band intensities of the full length products (lane 10). The lack of elongation with *BsM3*SC is in contradiction with ITC data showing that the combination of the two mutations in *BsM3*SC improves the **P**₇ interaction as compared to *BsM1*SC/**P**₇ or *BsM2*SC/**P**₇ interactions (**Figure 4**). In the elongation assay, the effect of both mutations is not cumulative but rather reflects the effect of the P₃₅₇S mutation alone. These differences between the two approaches reveals that, in the *Bs* context, the S₃₅₇ residue blocks the interaction of the PolC polymerase natural peptide with *Bs*SC, but not that of **P**₇. Alternatively, the *Bs*P₃₅₇ position could have some unknown specific contribution to the whole polymerase function. This effect is not observed in the *Ec* context and could thus reveal this position as a specific marker for Gram+ SC/polymerase interaction. Finally, as observed for both natural SC, none of the mutants SC form a productive heterologous complex with their non cognate polymerase (data not shown).

To make sure that the effects observed with *BsP* reflect the polymerase activity and not the SC loading process, we performed the same assay using a linear template (**SI.17**). SC dependent elongation products are detected with the *Bswt*SC and *BsM2*SC, as observed when using circular templates, but none with *BsM1*SC, ruling out any defect in the SC loading process. Noteworthy, in this assay with linear templates, we observed *BsM3*SC dependent products, as opposed to what we obtained using circular DNA (**Figure 8**, lane 11). This suggests that the observed increased elongation activity brought by the L₃₇₃M mutation compensate the deleterious effect of the P₃₅₇S mutation for what concerns the polymerase interaction, but not the SC loading process. Elucidation of this point will require the study of the respective natural peptides with these SC variants.

Then, we challenged the different reconstituted holoenzymes for their primer elongation activity with increasing concentrations of **P**₇. The elongation profiles are displayed in **Figure 9**, and the data are analyzed in **Figure 10**. In accordance with our previous observations in **Figure 8** (lanes 3 and 4) showing that *Ecwt*SC and *EcM1*SC are similarly efficient in driving *EcP* SC-dependent elongation, equal concentrations of **P**₇, pre-incubated with *Ecwt*SC or *EcM1*SC, equally inhibit the SC dependent primer elongation activity of *EcP*, yielding a IC₅₀ of 1.5 μM (**Figure 10 A, D**). These values are close to our previous results yielding an IC₅₀ of 2.8 μM for the inhibition of *EcP* holoenzyme elongation by **P**₇⁵. These

experiments confirm that the S₃₄₆P mutation has no, or a limited influence on the peptides binding, either the natural peptide (QADMF) or P₇. We note a 4-fold difference between these biochemical results and those measured by ITC at 34°C (SI. 6), but both approaches describe the same trend despite their different sensitivity, as observed before⁵.

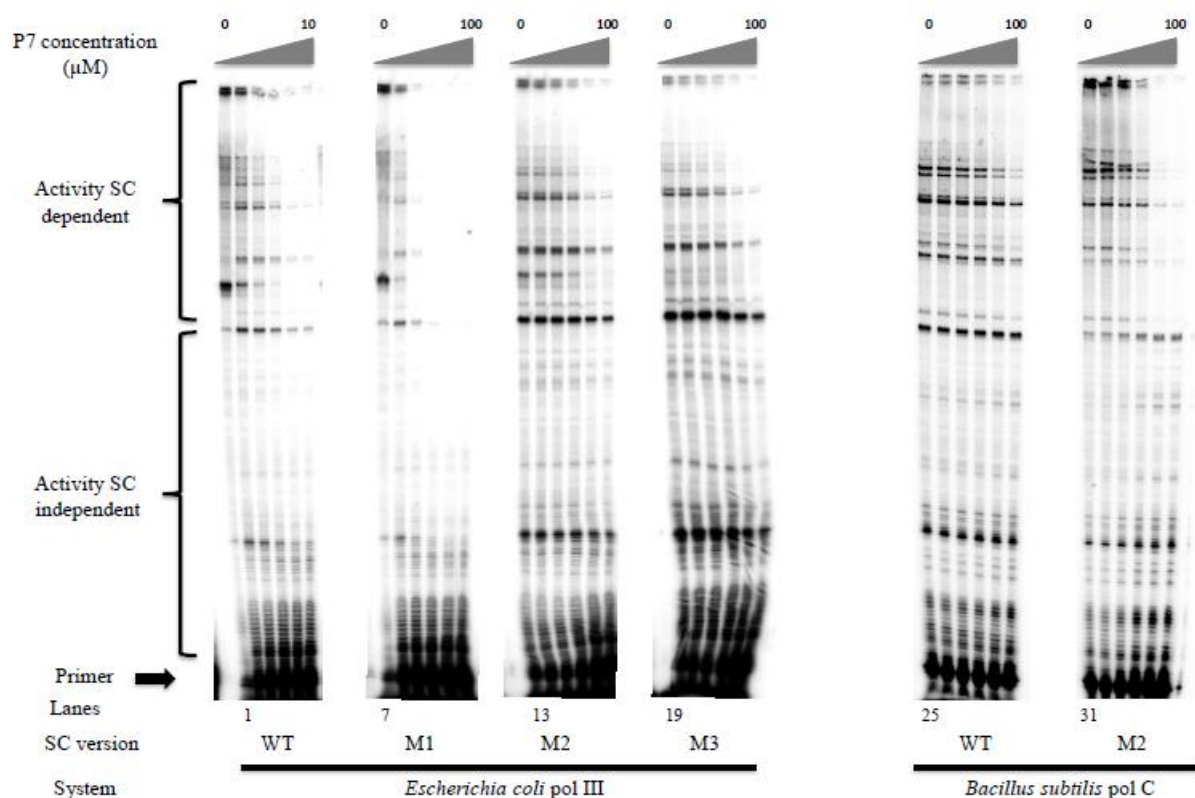


Figure 9 : Inhibition of primer elongation by P₇. EcP and BsP are complemented with their cognate wt or mutants SC pre-incubated with increasing concentrations of P₇ peptide (0, 1, 2, 4, 8 and 10 μM for ^{Ecwt}SC and 0, 2, 8, 25 and 75 μM for all the other SC). **Lanes 1 to 6:** EcP-^{Ecwt}SC. **Lanes 7 to 12:** EcP-^{EcM1}SC. **Lanes 13 to 18:** EcP-^{EcM2}SC. **Lanes 19 to 24:** EcP-^{EcM3}SC. **Lanes 25 to 30:** BsP-^{Bswt}SC. **Lanes 31 to 36:** BsP-^{BsM2}SC.

The same approach using ^{EcM2}SC or ^{EcM3}SC yield a much higher IC₅₀ value of 46 μM, indicating a much weaker interaction of these SC with P₇ (Figure 10 B, D), in agreement with our ITC data (SI.6). All the results from different analytical approaches indicate that for ^{Ec}SC, the introduction of a S₃₄₆P mutation is innocuous, whereas the M₃₆₂L mutation is strongly deleterious.

Finally, we challenged the natural BsP holoenzyme and its $BsM2$ SC supplemented version by **P₇** (Figure 10 C, D). While the interaction of **P₇** with Bs^{swt} SC is weak ($IC_{50} = 68 \mu M$), in agreement with our ITC data (SI.2), a 5 fold ($IC_{50} = 14 \mu M$) improvement is measured by introduction of the L₃₇₃M mutation in Bs SC, in good agreement with ITC and MD data.

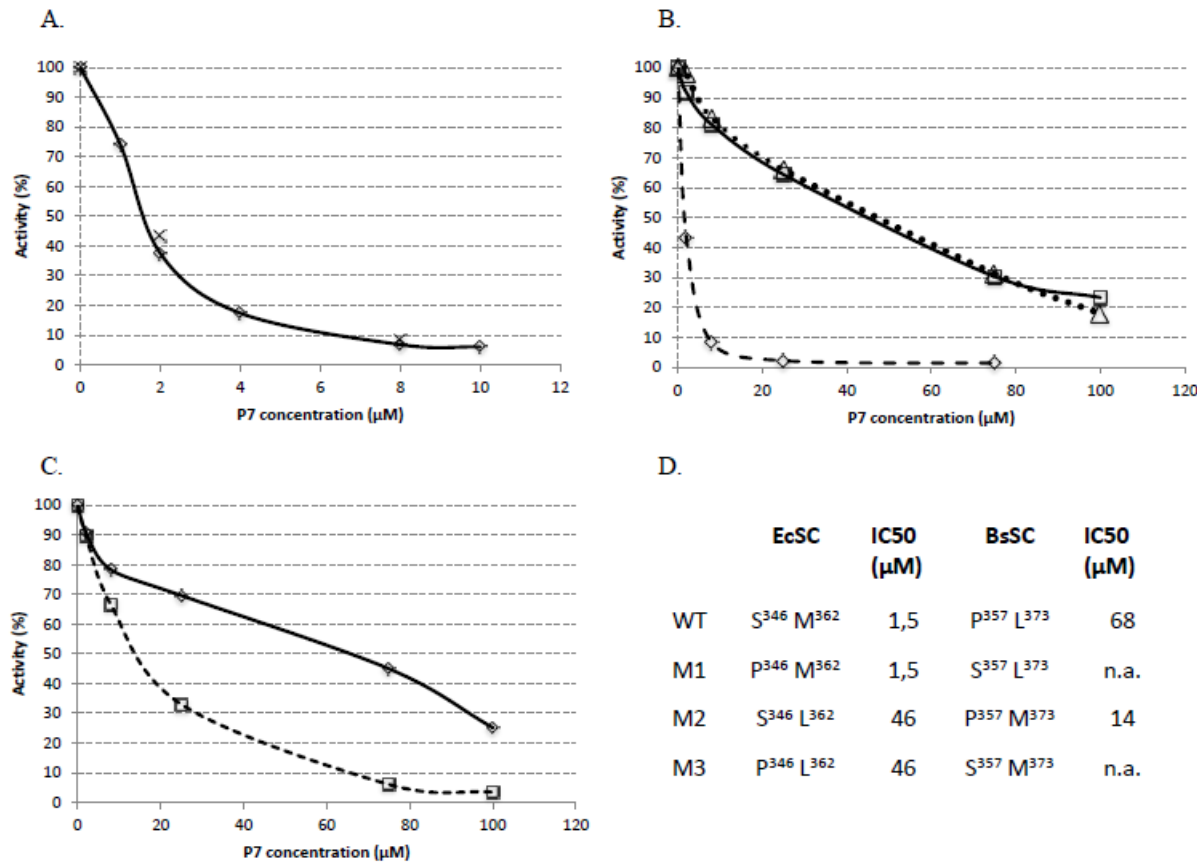


Figure 10: Quantification of EcP and BsP holoenzymes inhibition by P₇. A: Ec^{wt} SC (diamonds); $EcM1$ SC (crosses). B: $EcM1$ SC (diamonds, dotted line); $EcM2$ SC (squares, black line); $EcM3$ SC (triangles, dotted line). C: Bs^{swt} SC (diamonds); $BsM2$ SC (squares, dotted line). D: Determination of IC_{50} from the curves in A, B and C .

CONCLUSION.

The interaction of all DNA polymerases with the replicative processivity factor (or SC) is central for their activities and is mediated by a conserved peptide sequence which binds into a hydrophobic pocket located at the SC surface. In this piece of work, we analyzed the interaction of a reference peptide, **P₇**, with natural and mutant SC from *E. coli*, *B. subtilis* and *M. tuberculosis*. In particular, we focused on the contribution of two residues of the Ec SC binding pocket, namely S₃₄₆ and M₃₆₂, and their corresponding residues in Bs SC, P₃₅₇ and L₃₇₃.

ITC experiments confirm the differential interaction of **P₇** with the different natural SC. Single mutant analyses identify a strategic position in the *Ec*SC binding pocket: a single M₃₆₂L mutation is sufficient 1) to turn the thermodynamic profile of **P₇** interaction into that observed with *B^{swt}*SC, 2) to inhibit the SC dependant activity of the Pol III holoenzyme and 3) to reduce the competitive inhibitory effect of **P₇** on this enzyme activity. Conversely, the L₃₇₃M substitution in the *B^s*SC pocket strongly modifies the thermodynamic profile which becomes almost similar to that observed in the natural *Ec* context. All these results are in line with molecular dynamics data indicating that, contrary to M residues that, in both *Ec* and *B^s* contexts, shift upon peptide interaction, L residues remain in place, immediately below the peptide, providing a structural rationale for the peptide binding inhibition in *EcM2*SC and in *B^{swt}*SC. Δ Cp analyses reveal that all binding pockets endow a large structural change upon peptide binding, suggesting that **P₇** interacts through an induced fit process³³. In the case of *B^s*SC, this process is observed only with G2 and M3 mutants, because they lack the static L₃₇₃ residue which blocks peptide interaction. This indicates that the weak Δ Cp values calculated for *B^{swt}*SC, *EcM2*SC and others, reveal a poor peptide binding and not necessarily a change in binding mode.

Thermodynamic analyses suggest that the nature of the *Ec*₃₄₆ residue (S or P) is not as important for the binding as the M₃₆₂ residue, as deduced from the closely similar thermodynamic profiles obtained with *Ec^{wt}*SC, *EcG1*SC and *EcM1*SC. However, kinetics analyses reveal that the S₃₄₆P mutation strongly reduces the pocket dynamics. This suggests that the residue composition of each binding pocket is highly specific and has evolved to ensure an optimized binding process through a complex and specific network of interaction, as observed previously¹⁸. Another example is illustrated in the *B^s* context where biochemical analyses highlighted the P₃₅₇ residue as a *B^s* specific strategic position for productive elongation. These data underline the interest of a multidisciplinary approach in defining the binding characteristics of each binding pocket and will help in the future design of SC targeting compounds, with either large spectrum or strain specific activities.

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ABBREVIATIONS

ITC, Isothermal Titration Calorimetry; N-ter: N-terminal ; C-ter : C-terminal ; Ac: acetyl group; Cha: β -cyclohexyl-L-alanyl; (3,4-di-Cl)Phe : 3,4-dichloro-L-phenylalanyl.

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