The nuclease domain of the SPP1 packaging motor coordinates DNA cleavage and encapsidation

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ABSTRACT

The large terminase subunit is a central component of the genome packaging motor from tailed bacteriophages and herpes viruses. This two-domain enzyme has an N-terminal ATPase activity that fuels DNA translocation during packaging and a C-terminal nuclease activity required for initiation and termination of the packaging cycle. Here, we report that bacteriophage SPP1 large terminase (gp2) is a metal-dependent nuclease whose stability and activity are strongly and preferentially enhanced by Mn²⁺ ions. Mutation of conserved residues that coordinate Mn²⁺ ions in the nuclease catalytic site affect the metal-induced gp2 stabilization and impair both gp2-specific cleavage at the packaging initiation site pac and unspecific nuclease activity. Several of these mutations block also DNA encapsidation without affecting ATP hydrolysis or gp2 C-terminus binding to the procapsid portal vertex. The data are consistent with a mechanism in which the nuclease domain bound to the portal switches between nuclease activity and a coordinated action with the ATPase domain for DNA translocation. This switch of activities of the nuclease domain is critical to achieve the viral chromosome packaging cycle.

INTRODUCTION

Nucleases hydrolyze the phosphodiester linkages present in nucleic acids. These enzymes are implicated in numerous DNA transactions such as DNA replication, repair and recombination. Many nucleases possess other associated catalytic properties such as DNA polymerase, ligase, helicase or kinase activities that combine together to achieve specific biological functions. One of the diverse biological processes where nucleases play a fundamental role is viral maturation. In most tailed double-stranded DNA (dsDNA) bacteriophages and herpes viruses, DNA is replicated as concatemers that are the substrate for encapsidation into preformed procapsids (1–3). Specific recognition and cleavage of the viral concatemeric DNA during the packaging process is mediated by the terminase, a hetero-oligomeric complex composed in most phages of small and large subunits. Besides cleavage of the concatemeric DNA substrate, the terminase assembles at the procapsid portal vertex and possesses an ATPase activity necessary to fuel DNA translocation into the procapsid during the packaging process (4). The reaction is normally terminated by endonucleolytic cleavage of the concatemer and initiation of a subsequent packaging cycle at another procapsid. The process yields up to 12 or more encapsidation cycles per concatemer (5,6) revealing the high processivity of the viral packaging motor. The terminase large subunit is a multifunctional enzyme whose nuclease activity is required for an initial, most frequently sequence-specific DNA cleavage, followed by a switch to ATPase activity that powers dsDNA translocation. A switch back to nuclease activity in response to the level of DNA filling of the capsid achieves the dsDNA termination cut ending the packaging cycle. This cleavage is either sequence-specific or sequence-independent (headful packaging mechanism) depending on the viral system (7).

Structural information is available for the complete large terminase from phage T4 (8) and for nuclease domains from phages RB49 (8), SPP1 (9), P22 (10) and also nuclease domain from the human cytomegalovirus (HCMV-1) (11). The N-terminal ATPase domain is related to monomeric helicases, while the carboxyl
terminal nuclease domain exhibits an RNase H fold. Structural conservation suggests a common, yet unknown mechanism, for coordination of the critical switches between nuclease and ATPase activities during the viral genome packaging cycle of tailed phages and herpes viruses.

The Bacillus subtilis bacteriophage SPP1 is a well-established model system for viruses that package their dsDNA by a headful packaging mechanism. The SPP1 terminase small subunit gp1 provides specificity for binding to the pac sequence in the DNA concatemer (12,13) whereas the large subunit, gp2, displays nuclease and ATPase activities (14–16). Gp2, which is a 422-amino acid long polypeptide, is a monomer in solution consisting of an N-terminal ATPase domain and a C-terminal nuclease domain (14). Both ATPase and nuclease activities are modulated by gp1 and the portal protein gp6 (15,17). The gp6 portal, at the procapsid, provides the interface for assembly of the packaging motor through a direct interaction with gp2 (18). Motor pumping of dsDNA to the procapsid interior through the gp6 central channel (19) relies on a complex cross-talk between gp1, gp2 and gp6 and conformational changes in the portal region surrounding the central DNA conduit channel (17,20). Genetic evidence shows that gp6 is also central for the sensor mechanism that measures the level of DNA capsid filling to trigger the switch of gp2 activity to exert the sequence-independent mechanism, for coordination of the critical switches between nuclease and ATPase activities during the viral genome packaging cycle (21,22).

We report that gp2 nuclease activity and stability are modulated by divalent cations being preferentially enhanced by Mn<sup>2+</sup> ions. We show that amino acid substitutions in the nuclease catalytic site impair both gp2-specific cleavage at pac and its unspecific nuclease activity but neither affect binding to the capsid portal vertex nor its ATPase activity. Significantly, several of these mutations block linear DNA packaging in vitro revealing that a correctly structured gp2 nuclease center is critical for the mechanism of DNA translocation into viral capsids.

**MATERIALS AND METHODS**

**Enzymes and reagents**

All chemicals used in this study were of p.a. grade. Sybr Gold was from Molecular Probes. Protein molecular weight markers and DNA restriction enzymes were from Biolabs. Proteinase K was from Roche. Lysozyme, DNase and Dextran were from Sigma. Adenosine-5'-triphosphate was purchased from Roche. [γ-<sup>32</sup>P]ATP was from Perkin Elmer.

**Bacterial strains, bacteriophages and plasmids**

SPP1 suppressor-sensitive mutants (sus70, sus19 and sus70sus115) were described previously (12,21,22). Bacillus subtilis HA101B (sup-3) and YB886 (sup') were the permissive and non-permissive strains used for SPP1 amplification (12). The Escherichia coli strains used for production of the terminase subunits gp2 and gp1 were BL21(DE3) and BL21(DE3) (pLysS), respectively (23). Plasmids pCB191 (24), pBT115 (12,16) and pBT163 (12) have been described. Expression was controlled by the LacI repressor coded by plasmid pREP4 (QIAGEN) in E. coli. The E. coli XL-1 Blue strain (Stratagene) was used for gp2 site-directed mutagenesis. The gene 2 alleles coding for gp2 single and double mutants were engineered by site-directed mutagenesis using the QuickChange Site-Directed Mutagenesis Kit (Stratagene). The templates for mutagenesis were plasmid pCB191 for expression in E. coli and pBT163 for expression in B. subtilis. Residues 1–198 of gp2 were cloned into pET28a using the NdeI/ HindIII restriction sites [N-terminal domain (NTD) with a C-terminal His tag]. The C-terminal gp2 construct (residues 232–422) was described previously (9). Complete sequencing of gene 2 alleles was carried out (GATC, Konstanz). Phage production with the different gp2 mutants was analyzed by in vivo complementation assays in the B. subtilis strain YB886 carrying the pBT163 constructs with different gene 2 alleles.

**DNA manipulations**

Plasmid DNA was purified with a plasmid purification kit from Qiagen. Mature SPP1 and T5 DNAs were extracted from CsCl-purified phage particles by phenol–chloroform extraction followed by precipitation with isopropanol. DNA concentration was determined using molar extinction coefficient of 6500 M/cm at 260 nm.

**Protein purification**

Gp1 and gp2 were purified as described previously (16). Protein concentration was determined by the Bradford method (25), using bovine serum albumin as a standard. Gp1 and gp2 concentrations were expressed as moles of protein decamers (13) and monomers (14), respectively.

**Production and purification of SPP1 procapsids**

Wild-type (wt) SPP1 procapsids or procapsids lacking the portal protein, gp6, were produced by infection of the B. subtilis YB886 strain with SPP1 sus70 and sus70sus115 suppressor-sensitive phages, respectively, and purified as described (17). The purity of the procapsid preparations was analyzed by SDS–PAGE and the presence or absence of gp6 was checked by western blotting using anti-gp6 antibody. Blots were developed using the ECL detection system (Amersham Pharmacia Biotech). The quality of the procapsid preparations was also checked by electron microscopy of negatively stained samples.

**Fluorescence-based thermal shift assay**

Thermal stability assays were performed with 1 μM gp2 in 20 mM Tris–HCl pH 7.5, 300 mM NaCl and the SYPRO Orange dye as a fluorescent probe (diluted ~800-fold from a 5000-fold stock solution, Invitrogen). Fluorescence measures the binding of the SYPRO orange probe to exposed protein hydrophobic regions. This dye becomes highly fluorescent when bound to protein hydrophobic
sites, which become accessible upon thermal unfolding (26). Reaction mixtures were made in duplicate in a 96-well fast PCR plate at a final volume of 10 μl. The temperature gradient was carried out in the range of 15°–95°C at 1.5°C/min with a StepOnePlus real-time PCR system (Applied Biosystems). Fluorescence was recorded as a function of temperature in real time (excitation with a blue LED source and emission filtered through a 5-carboxy-X-rhodamine emission filter). The melting temperature \( T_m \) was calculated with StepOne software v2.2 as the maximum of the derivative of the resulting SYPRO Orange fluorescence curves. Data from duplicates were very reproducible (error in the \( T_m \)s ranging from 0.0°C to 0.02°C). Then the mean from the duplicates was considered as the value obtained for each condition in each experiment.

**In vitro nuclease assay**

The *in vitro* nuclease activity assay was performed as described previously (16). Unless indicated elsewhere DNA was incubated with gp2 proteins (1 μM) in 50 mM Tris–HCl pH 7.5, 50 mM NaCl with or without metal ions at 37°C for 1 h. The reactions were stopped by addition of EDTA and the samples deproteinized by incubation with proteinase K and SDS for 45 min at 37°C. Analytical gel electrophoresis was carried out in 0.8% (w/v) agarose/Tris–borate EDTA horizontal slab gels.

**In vivo pac cleavage assays**

The assay was adapted from Chai *et al.* (12). Exponentially growing \( \text{OD}_{600} = 0.8 \) *B. subtilis* YB886 strains carrying plasmid pBT163 (co-expressing gene 1 and different gene 2 alleles) were centrifuged and the pellet was resuspended in 1/10 of the initial culture volume in lysis solution (100 mM Tris–HCl pH 7.5, 10 mM EDTA, 400 μg/ml RNase, 1 mg/ml lysozyme). After 1-h incubation at room temperature, SDS was added to a final concentration of 0.5% and cell lysates incubated 10 min, followed by addition of pronase and further incubation for 30 min at 37°C. Total DNA was extracted with phenol/chloroform and dialyzed extensively against 10 mM Tris–HCl pH 8.0, 10 mM EDTA. DNA was then EcoRI-cleaved and the digests were electrophoresed in 1% agarose and deproteinized. DNA was resolved by gel electrophoresis in 0.8% agarose gels and stained with Sybr Gold according to the manufacturer’s instructions.

**DNA packaging reaction in vitro**

DNA packaging reactions were performed using the SPPl *in vitro* DNA packaging system as described (16). Briefly, unless stated otherwise, a standard 20 μl reaction mixture contained 1 mM ATP, 1 mM linear plasmid DNA (pBT115), 10 nM procapsids and terminase proteins (1 μM of gp2 and 1 μM of gp1) in 50 mM Tris–HCl, pH 7.8, 50 mM NaCl, 10 mM MgCl₂, 10% dextran. Reactions were incubated for 1 h at 30°C. DNase was then added at 20 μg/ml and incubation continued for 10 min at 30°C. Reactions were stopped by addition of 50 mM EDTA and deproteinized. DNA was resolved by gel electrophoresis in 0.8% agarose gels and stained with Sybr Gold according to the manufacturer’s instructions.

**RESULTS**

**Metal ion and substrate specificity of gp2 nuclease**

Gp2 has sequence-independent endonucleolytic activity in the presence of the divalent cation Mg²⁺, converting supercoiled circular plasmid DNA to nicked open circular DNA and to linear DNA, but is not active in the presence of Ca²⁺, Zn²⁺ or Ni²⁺ (Figure 1A, lanes 3, 4, 7 and 8) (14–16). The gp2 nuclease activity is strongly increased by Mn²⁺ or Co²⁺ relatively to the effect observed with Mg²⁺, leading to complete DNA degradation in similar experimental conditions (Figure 1A, compare lanes 3, 5 and 6). Gp2 also displays a strong nuclease activity on linear dsDNA in the presence of Mn²⁺ and Co²⁺, degrading DNA to a smear of DNA fragments (Figure 1B, lanes 5 and 6). None of the other metal ions tested, including Mg²⁺, led to detectable cleavage of linear DNA (Figure 1B). Experiments performed with a circular DNA substrate and diverse Mn²⁺ concentrations ranging from 100 μM to 10 mM showed that as the Mn²⁺ dose increases, gp2 converts supercoiled circular plasmid DNA to nicked open circular DNA, subsequently to linear DNA and finally to completely degraded DNA (Figure 1C). In the presence of Mn²⁺, gp2 also digests long linear dsDNA substrates such as plage SPPl (45 kb) and T5 (125 kb) DNAs (Figure 1D) in a dose-dependent manner.
The gp2-isolated carboxyl-domain has nuclease activity in the presence of Mn\(^{2+}\)

The gp2 nuclease center resides in the protein C-terminal domain (CTD) (9). To check whether the gp2 CTD alone [residues 232–422, as defined in Smits et al. (9); Figure 2A and Supplementary Figure S1] could cleave DNA, its nuclease activity was compared with that of the full-length (FL) protein. The isolated gp2 CTD showed no detectable nucleolytic activity in the presence of Mg\(^{2+}\) as metal ion (Figure 2B compare lanes 2 and 3). Interestingly, in the presence of Mn\(^{2+}\), the isolated gp2 CTD gained the capacity to cleave and degrade different DNA substrates (Figure 2C, circular DNA, left panel and linear DNA, right panel), although less efficiently and at a higher Mn\(^{2+}\) concentration, than found for the FL protein. The results confirm that the gp2 CTD harbors the enzyme nuclease activity and strengthens the gp2 Mn\(^{2+}\) preference to support nucleolytic cleavage of circular and linear dsDNA. They also reveal that optimal gp2 nuclease action requires the FL gp2 protein, as reported for other phage terminases (10,28).

Binding of the gp2 nuclease domain to portal-containing procapsids

Besides its nuclease and ATPase activities, gp2 also plays a crucial role during assembly of the DNA packaging motor by directly docking at the procapsid portal vertex (18). Previous biochemical and genetics experiments in phages T3, T4 and λ suggested that residues of the CTD of the large terminase subunits are involved in the interaction

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**Figure 1.** Metal ion specificity of gp2 nuclease. (A) Effect of divalent cations on gp2 circular dsDNA nuclease activity. Supercoiled pBT115 DNA was incubated with purified gp2 alone or with gp2 in the presence of 10 mM Mg\(^{2+}\), Ca\(^{2+}\), Mn\(^{2+}\), Co\(^{2+}\), Zn\(^{2+}\) or Ni\(^{2+}\). After 1 h at 37°C, the reactions were stopped and DNA analyzed as described under ‘Materials and Methods’ section. R, relaxed circular; L, linear; CCC, covalently closed circular DNA. (B) Effect of divalent cations on gp2 linear dsDNA nuclease activity. Linear plasmid pBT115 (PstI-cleaved) was incubated with gp2 with or without different divalent cations and analyzed as above. (C) Dose-dependent effect of Mn\(^{2+}\) on gp2 nuclease activity. Supercoiled pBT115 DNA was incubated with gp2 in the presence of increasing concentrations of Mn\(^{2+}\). (D) Nuclease activity of gp2 with long linear DNA substrates in the presence of different Mn\(^{2+}\) concentrations. SPP1 (45 kb) or T5 (125 kb) phage DNAs were incubated with gp2 in the presence of increasing concentrations of Mn\(^{2+}\).
with the portal (29–32). To address the question of whether the gp2 CTD mediates portal binding during motor assembly, we carried out affinity pull-down assays with the isolated gp2 CTD and NTD and purified SPP1 procapsids. The gp2 CTD is able to pull-down procapsids, although less efficiently than the FL protein (Figure 3, lanes 2 and 3). The specific interaction between procapsids and the gp2 CTD is mediated by the portal protein, as shown by parallel experiments with procapsids lacking the portal protein (Figure 3, compare lanes 3 and 7). A very faint signal corresponding to procapsid pull-down with the gp2 NTD was also detected. However, this binding to procapsids is unspecific as no difference was found between procapsids with or without the portal protein (Figure 3, lanes 4 and 8). The CTD of gp2 thus mediates anchoring of the packaging motor at the portal vertex in addition to its nuclease function.

**Divalent metal ions stabilize the nuclease domain and the gp2 overall structure**

To investigate whether the effect of metal ions on gp2 nuclease activity results of an exclusive requirement for catalysis or if their binding correlates also with a stabilization effect on the gp2 structure we monitored the protein thermal stability by a fluorescence-based thermal shift assay (26). This method measures fluorescence emission upon binding of a probe to hydrophobic
regions that become exposed during protein thermal denaturation. Gp2 melting curves are shown in Figure 4A. An increase in thermal stability was found upon addition of 1 mM Mn$^{2+}$ leading to a $T_m$ (apparent melting temperature) rise of $\sim$8°C, followed by Co$^{2+}$, Ni$^{2+}$ and, to a less extent, Ca$^{2+}$ and Mg$^{2+}$. Dose-dependent experiments with metal concentrations ranging from 320 nM to 5 mM confirmed these results (Supplementary Figure S2). We then compared the stability of the isolated gp2 NTD and CTD, in the absence or in the presence of Mg$^{2+}$ or Mn$^{2+}$, relatively to FL gp2 in the same experimental conditions. No significant difference was detected between the $T_m$ of the metal-free NTD and FL gp2 (Supplementary Figure S3A). The metal-free gp2 CTD shows reproducibly a more complex denaturation behavior (black curve in Figure 4B, right panel) leading to double minima for the derivative of the fluorescence signal curve, which hampers an accurate determination of the $T_m$ (Supplementary Figure S3B). Nevertheless, its denaturation occurs at $\sim$39°C, as the NTD and FL gp2. In contrast to the FL gp2, no increase in $T_m$ of the NTD and CTD was detected following Mg$^{2+}$ or Mn$^{2+}$ addition (Figure 4 and Supplementary Figure S3). However, addition of Mn$^{2+}$ at 200 μM led reproducibly to a unimodal denaturation curve of the gp2 CTD in contrast to the behavior found in absence of metal ion or at the same Mg$^{2+}$ concentration (Figure 4B and Supplementary Figure S3).

Mutations in the nuclease active site have distinct effects on metal-induced gp2 stabilization

Soaking gp2 nuclease domain crystals with Mn$^{2+}$ revealed two peaks in the anomalous difference maps, corresponding to bound metal ions (9). One ion is coordinated by D266, H400 and D403 [metal ion A (33), while the second ion is coordinated by D266 and D321 [metal ion B]] (9) (Figure 5A and Supplementary Figure S4). These residues are highly conserved in gp2-related proteins and a similar organization for coordination of metal ions is found in all four terminase nuclease domains whose structure is available (Figure 5A). They create an electronegative environment for two metal ion-dependent catalysis of endonucleolytic cleavage in a strand of dsDNA that is accommodated in the active site cleft (9). The finding that two Mn$^{2+}$ are coordinated at the gp2 catalytic center led us to investigate if these two bound ions play a structural role responsible for the gp2 metal-induced enhanced stability. The thermal stability of gp2 carrying single and double mutations in residues D266, D321, H400 and D403 was analyzed with or without metal ions. Gp2 carrying a mutation to alanine in the conserved E134 of the Walker B motif in the N-terminal ATPase domain was characterized as a control. None of the mutations affected gp2 stable production (data not shown). Likewise, none of the mutations reduced protein thermal stability in the absence of divalent metal ions (Figure 5B, black bars). Two of the gp2 double mutants, D266N/D403N and D321N/D403N, were actually more stable than the wt protein in these experimental conditions ($\sim$4°C increase in $T_m$), and a slight increase of $T_m$ was observed for the H400A and D403N mutants. Addition of Mg$^{2+}$ (light to dark blue bars) or Mn$^{2+}$ (orange to red bars) caused distinct effects on the thermal stability of the different gp2 proteins. Wild-type gp2 and the N-terminal E134A mutant exhibited a similar behavior, i.e. a modest increase in $T_m$ caused by Mg$^{2+}$ and a marked Mn$^{2+}$-induced $T_m$ raise for both proteins. Addition of Mn$^{2+}$ also increased the thermal stability of the D321N, H400A and D403N gp2 mutant proteins, although the effect was less pronounced than in wt and in the E134A gp2. In contrast, neither Mg$^{2+}$ nor Mn$^{2+}$ stabilized the D266N and D266N/D321N mutants. Instead, Mn$^{2+}$, at the higher concentration used in these experiments (5 mM, dark orange bar) seems to render the two proteins slightly less stable. This small decrease in stability caused by high Mn$^{2+}$ associated with D266N mutation was abolished when combined with D403N substitution. The latter mutation, isolated or combined with other mutations of aspartic acids in the active site, leads to a higher stability of gp2 in absence of metal ions but renders also gp2 stability poorly responsive to this type of ions (Figure 5B).

Gp2 nuclease mutations impair non-specific DNA cleavage

Gp2 proteins carrying amino acid substitutions D266, D321, H400 and D403 in the nuclease active site lost the capacity to digest linear dsDNA, contrarily to wt gp2, demonstrating that these residues are critical for gp2 non-specific nuclease activity (Figure 5C, left panel, lanes 4–7). The same holds true for the three double mutants D266N/D321N, D266N/D403N and D321N/D403N (Figure 5C, right panel, lanes 3–5). The fact that all the substitutions affected the gp2 nuclease shows in addition that the cleavage seen in the in vitro assays with the wt nuclease is inherently associated with gp2 and not with other eventual nuclease contaminants. In contrast, the N-terminal mutation E134A has no effect on nuclease activity, as gp2 E134A mutant protein could...
Gp2 nuclease mutations drastically reduce in vivo pac cleavage

Initiation of SPP1 DNA packaging in vivo occurs by specific recognition and cleavage of the pac site at the DNA concatemer. Cleavage at pac requires the presence of the small and large terminase subunits, gp1 and gp2 (12,34). To evaluate the role of the gp2-mutated residues in protein-specific nuclease activity, we set up a pac cleavage assay in vivo, using a plasmid system [pBT163, ~7.4 kb; (12)] co-expressing genes 1 and 2. As schematized in Figure 6A, the SPP1 pac cleavage site is located within gene 1. Cleavage at pac divides a 958-bp EcoRI DNA fragment of pBT163 into the 242- and 716-bp DNA fragments. Total DNA from exponentially grown B. subtilis strains carrying the pBT163 constructs co-encoding gp1 and the wt or mutant gp2 proteins was isolated, digested with EcoRI and separated in agarose gels. Southern blots were then hybridized with the probe indicated in Figure 6A. As shown in Figure 6B, a fragment of ~700 bp is detected in the presence of wt gp2, in agreement with the predicted fragment size following pac cleaved and EcoRI digestion of the pBT163 plasmid (Figure 6A). As a negative control, the ~700-bp fragment was not present with a truncated gp2 protein (A232stop) lacking the gp2 CTD. As reported previously for pac cleavage during SPP1 infections (12), we
Figure 5. Thermal stability assays and in vitro nuclease activity of gp2 mutant proteins. (A) Conservation of the terminase nuclease catalytic site of phage SPP1 (gp2; PDB accession code 2WC9), of HCMV (cytomegalovirus) (UL89; PDB 3N4Q) and of phage T4 (gp17; PDB 3CPE). The side chains of residues involved in metal coordination/catalysis are shown as sticks. The Mn$^{2+}$ ions (magenta spheres) in SPP1 gp2 and in HCMV UL89 structures are 4 and 3.6 Å far apart, respectively. The structures were aligned with DALI and the figure prepared with PyMol. (B) Thermal stability of wild-type (wt) and gp2 mutant proteins analyzed as in Figure 4 in the absence of divalent metal ion (black bars) or in the presence of different concentrations (200 μM, 1 or 5 mM) of Mg$^{2+}$ (light to dark blue bars) or Mn$^{2+}$ (orange to red bars). The $T_m$ values (°C) shown are the mean ± SD of results from at least three independent experiments. All proteins showed a single-transition denaturation curve. (C) In vitro nuclease activity of gp2 mutants. The nuclease activity of the gp2 single (left panel) and double (right panel) mutants was checked in an in vitro nuclease assay, as described under 'Materials and Methods' section, using a linear plasmid DNA substrate in the presence of 2.5 mM Mn$^{2+}$.
failed to detect the small 242-bp fragment that could be expected following cleavage at pac (Figure 6A). This asymmetry may reflect the formation after or during pac cleavage of a complex between phage DNA and the terminase proteins that protected packageable DNA against nucleolytic attack from cellular exonucleases (12). The nuclease(s) that could be responsible for the observed degradation of the 242-bp fragment remain(s) unknown. The 958-bp fragment detected in the Southern blot corresponds to the pBT163 EcoRI DNA fragment with uncut pac whereas upper bands arise from uncut and partially digested plasmid DNA. The presence of the 958-bp fragment shows that the plasmid assay system reproduces faithfully also the control mechanism of pac cleavage observed during SPP1 infection that is not a frequent event, occurring only once 'per' SPP1 packaging series in a phage DNA concatemer (6,34).

No detectable cleavage at pac occurred with the single mutant protein D266N and the three double mutants D266N/D321N, D266N/D403N and D321N/D403N. In case of mutations D321N, H400A and D403N we observed a strong reduction in the generation of the ~700-kb fragment when compared to gp2 wt, revealing a residual nuclease activity. None of the mutations

Figure 6. Gp2 nuclease mutations inhibit pac cleavage in vivo. (A) Schematic representation of the pBT163 plasmid (7.4 kb) (12) which contains the 2.6-kb Banl-HindIII fragment from SPP1 DNA, comprising genes 1 and 2. The vertical arrow denotes the SPP1 pac cleavage site, situated within gene 1. EcoRI sites as well as position of gp2 mutations inserted by site-directed mutagenesis in pBT163 are indicated. The hybridization region of the Southern-blot probe used in the in vivo pac cleavage assay was from pBT115 carrying gene 1 with the pac cleavage site and part of gene 2) is shown. SPP1 DNA is depicted in light grey. (B) Southern blot of EcoRI-digested DNA isolated from B. subtilis strain YB886 carrying pBT163 coding gene 1 and different gene 2 alleles. The blot was hybridized with the DNA fragment probe schematized in panel (A). The band with an electrophoretic mobility close to the 5-kb marker is attributed to intact 7.4-kb supercoiled plasmid that was observed to exhibit this migration behavior. (C) Effect of gp2 mutations on phage yield. The effect of gp2 mutations on phage production was estimated in complementation assays by titration of SPP1sus19 phages [the position of the sus19 mutation in gene 2 is shown in panel (A)] in strains bearing plasmids coding for gene 2 mutant alleles and estimated as percentage of the titer (pfu, plaque-forming units/ml) obtained with the strain coding for wt gp2. The phage yield for the wt gp2 (considered as 100%) was $2.0 \times 10^{11}$ pfu/ml. Values are the average of three independent experiments.
affected gp2 production in the *B. subtilis* cells (data not shown). The deficiency of *in vivo* pac cleavage of gp2 nuclease mutants in the pBT163 system (Figure 6B) correlated with the inability of the plasmid-borne gp2 mutants to complement SPP1 sus19, a mutant defective for gp2 production (Figure 6C). The N-terminal gp2 E134A mutation, which affects neither the unspecific gp2 nuclease (Figure 5C) nor specific cleavage at pac (Figure 6B), strongly impairs phage production (Figure 6C). This can be explained by the lack of ATPase activity of this mutant gp2 (see below).

**Nuclease mutants retain gp1-stimulated gp2 ATPase activity and gp2 portal binding**

To control that mutations in the gp2 nuclease active center do not have a general impact on gp2 structure we assayed their ATPase activity, a requirement for gp2 function in DNA packaging. All nuclease mutants exhibited a low-basal ATPase activity similar to wt gp2 (Figure 7, light bars) and retained the gp1-stimulated ATPase activity (dark bars). No significant differences were observed when Mn$^{2+}$ is used in place of Mg$^{2+}$ as the metal ion (data not shown). On the contrary, the E134A mutation completely abrogated ATPase activity, as expected from its localization in the gp2 ATPase center. All mutant gp2 proteins exhibited normal binding to procapsids in a portal-mediated manner (Supplementary Figure S5, top and bottom panels).

**Gp2 nuclease mutants show distinct DNA packaging phenotypes**

The gp2 nuclease activity is required *in vivo* for packaging initiation at the pac sequence and to terminate the packaging cycle by a non-specific endonucleolytic cleavage. After both cleavage events, this activity shall be switched off to allow for a DNA translocation step into the viral procapsid driven by the ATPase activity of the gp2 NTD. We hypothesized that mutations in the nuclease center could impact on the DNA translocation mechanism because the substrate DNA might remain in the vicinity of the gp2 nuclease center cleft during its pumping to the capsid interior. To address this question, we took advantage of the *in vitro* defined SPP1 DNA packaging system that, contrarily to the *in vivo* situation, neither requires cleavage at pac nor headful cut if a linear DNA molecule shorter than the capsid packaging capacity is used as substrate (16). The system only requirement for packaging is a free end on a linear DNA molecule. Here DNA encapsidation was monitored by a DNase protection assay (16). Purified gp2 NTD or CTD, either alone or in combination, exhibited no DNA packaging activity, showing that SPP1 DNA packaging requires a FL gp2 (Figure 8A, lanes 3–5 versus lane 2), as also found for the large terminase of bacteriophage T4 (28). As could be expected from the ATPase assays (Figure 7), DNA packaging was fully abrogated when packaging reactions were carried out with the gp2 ATPase mutant E134A (Figure 8B, left panels, lane 7). Interestingly, diverse packaging phenotypes were registered with gp2 proteins carrying the nuclease mutations in packaging reactions carried out in the presence of Mg$^{2+}$ or Mn$^{2+}$. D321N and D266N/D321N substitutions do not significantly affect DNA packaging (Figure 8B). In contrast, mutations D266N, H400A (Figure 8B, left panels) and D266N/D403N (right panels) block DNA packaging in the presence of Mg$^{2+}$ (top panels) or of Mn$^{2+}$ (bottom panels). Interestingly, mutation D321N suppresses D266N yielding a double mutant competent for DNA packaging. Finally, mutant proteins D403N

![Figure 7](image-url)  
**Figure 7.** Effect of gp2 nuclease mutations on the terminase ATPase activity. The ATPase activity of wt and gp2 mutant proteins (1 µM) was determined in the absence (light bars) and in the presence (dark bars) of gp1 (1 µM) as described in ‘Materials and Methods’ section. The graphic shows the mean and SD of the results obtained for three independent experiments.
and D321N/D403N cannot package DNA in the presence of Mg\(^{2+}\) (left and right top panels in Figure 8B) but both mutants gain the capacity to package DNA in the presence of Mn\(^{2+}\) (lower left and right panels). Structural organization of the terminase nuclease active center and metal coordination thus plays an essential role in viral DNA encapsidation.

**DISCUSSION**

The two-domain terminase large subunit of tailed phages and herpesviruses is a dual activity enzyme of the viral DNA packaging motor. Its ability to switch between endonuclease and ATP-driven translocase activities is essential to achieve viral DNA packaging that occurs processively along a dsDNA concatemer substrate. The present study addressed the role of the nuclease catalytic center of SPP1 gp2 throughout the complete encapsidation cycle. Mutagenesis and functional analyses show that metal coordination at the nuclease center impacts on gp2 stability (Figures 4 and 5B) and, in addition to the expected function in DNA cleavage, it also plays a role in DNA encapsidation (Figure 8). The gp2 nuclease domain is thus the coordination center of the viral DNA packaging motor.
packaging motor, a function that fits well to the finding of its direct binding to the capsid portal system through which DNA is pumped (Figure 3).

The gp2 nuclease activity shows a strong preference for Mn$^{2+}$ and Co$^{2+}$ over Mg$^{2+}$ while the enzyme is inactive in the presence of Ca$^{2+}$, Zn$^{2+}$ or Ni$^{2+}$ (Figure 1). Mn$^{2+}$ has also a substantial and preferential stabilization effect on FL gp2 structure (Figure 4A). Mutagenesis of aspartate residues that coordinate metal ions for catalysis in the nuclease active center impaired or reduced the Mn$^{2+}$-enhancement of gp2 stability (Figure 5). Binding of the metal at the catalytic center thus plays a structural role. This role depends on the inter-domain cross-talk with the ATPase domain of gp2 since strong stabilization by Mn$^{2+}$ is observed only in the context of FL gp2 (Figure 4 and Supplementary Figure S3A). A similar dependence is found for the gp2 nuclease activity. The isolated gp2 CTD cleaves DNA in the presence of Mn$^{2+}$ but not of Mg$^{2+}$ (Figure 2). Mn$^{2+}$-dependent DNA cleavage with the purified CTD is, however, much less efficient than with the FL protein, in agreement with previous data (9). The reduced Mn$^{2+}$-enhancement of gp2 CTD stability relatively to FL gp2 may have an effect on the nuclease activity of the CTD domain which is weaker than in FL protein. However, high Mn$^{2+}$ concentrations enhance the CTD nuclease activity without a major effect in the protein thermal stability. Altogether the results with the gp2 CTD suggest that although the ATPase NTD has no detectable nuclease activity, its presence is required for optimal gp2 conformation and/or catalysis, as proposed for other phage terminases (10,28). A strong preference for Mn$^{2+}$ as co-factor in highly efficient nucleic acid cleavage appears to be a widespread feature of viral enzymes displaying DNA or RNA nuclelease activities such as the large terminase of the herpes virus HCMV UL-89 (11), the influenza virus RNA-dependent polymerase (35,36) or the influenza-like endonuclease domains of Bunyaviridae and Arenaviridae RNA polymerases (37,38). A divalent metal cation requirement (Mg$^{2+}$ or Mn$^{2+}$) was also reported for the nuclease activity of phages λ (39), T5 (40) and T4 terminases (28).

Despite poor sequence conservation, the core of the gp2 nuclease domain has a fold similar to the nuclease domains of RNase H family proteins and binds two Mn$^{2+}$ ions in its active site (9), suggesting a common two metal-ion-dependent catalysis mechanism to hydrolyze the phosphate ester bond of the nucleic acid chain (33). Conserved acidic residues are found in the catalytic centers of these nucleases (40). They coordinate a pair of divalent metal ions, often Mg$^{2+}$, spaced approximately by 4 Å that are proposed to activate the attacking nucleophile and stabilize the transition state during the nucleic acid cleavage reaction (33,41–43). Three of these acidic residues are positioned at equivalent positions in previously determined structures of RNase H family proteins and correspond to gp2 D266, D321 and D403 (9). In addition, the active site histidine H400 is also conserved in SPP1 closely related terminases (9) or is replaced by aspartate in the structures of the T4, HCMV and P22 terminases (8,10,11). H400 could be important in conferring the Mn$^{2+}$ preference found in our cleavage and protein stability assays because Mn$^{2+}$ ions can be favorably coordinated by both acidic residues and histidines, whereas ligation by Mg$^{2+}$ by histidines is less common (44,45). This is due to the higher affinity of Mn$^{2+}$ to nitrogen relatively to Mg$^{2+}$, which prefers oxygen (44). Mutation of the gp2 conserved residues (D266N, D321N, D403N and H400A) impaired the enzyme nuclease activity (Figures 5C and 6). All mutant gp2 proteins retained ATPase activity, gp1-stimulated ATPase and binding to portal-containing procapsids, showing that the mutations do not affect the gp2 overall folding. The most drastic effect was observed for the D266N substitution that completely abolishes gp2 cleavage at pac (Figure 6). This observation is consistent with the critical topology of D266 in the gp2 catalytic site (Figure 5A and Supplementary Figure S4). It is possible that defective metal-binding due to the other point mutations is poorly compensated by the presence of the remaining acidic residues in the active site, explaining some residual pac cleavage activity for the D321N, H400A and D403N gp2 proteins. In conformity with this possibility, double mutations of the active site aspartic acids totally abolished nuclease activity (Figure 6).

The precise location of Mn$^{2+}$ ions found in the active site of the gp2 nuclease domain (Figure 5A) (9) might undergo some change during catalysis because full metal occupation in the correct geometry at the nuclease catalytic site is only expected upon substrate binding (33). However, structural conservation and our functional analyses provide a plausible organization for the gp2 nuclease catalytic site (Figure 5A). Its architecture is compatible with a two-metal ion catalytic pathway (33) in which metal B is coordinated by D266 and D321 and metal A by D266, H400 and D403 (Supplementary Figure S4). D266 is the aspartate residue strictly conserved in nucleases of the RNase H family that coordinates the two metal ions required for the phosphoryl transfer reaction leading to cleavage of the scissile phosphate of a nucleic acid or nucleotide (33). Its substitution in gp2 fully inactivates nuclease activity and Mn$^{2+}$-enhancement of gp2 stability (Figure 5). This effect, which likely results of the inability of gp2 D266N to bind metal ions, is also found when combined with D321N. In contrast, mutation D403N and its combination with substitution of aspartates D266N and D321N lead to a high thermal stability independent of metal ions that is attributed to relief of local destabilization of the structure due to repulsive forces between D403 and neighbor carboxylate residues (D266 and D396). This destabilization effect might be one of the roles of D403 to support metal coordination and catalysis. Coordination of metal ions to the gp2 catalytic center thus has the expected role in catalysis but plays also a specific structural role to stabilize the gp2 structure. The concentration of Mg$^{2+}$ in the cytoplasm of exponentially growing B. subtilis was reported to be ~50mM, although a large fraction is complexed with cellular components primarily rRNA, while Mn$^{2+}$ is found in micromolar amounts (46). A similar situation was described for eukaryotic cells (47). The precise role of both ions on terminase activity in the cell is thus difficult to assess. It is important to note, however, that Mn$^{2+}$
increases gp2 stability at concentrations as low as 8 μM (Supplementary Figure S3) and that the endonuclease is active at 100 μM Mn²⁺ in our in vitro plasmid nicking/linearization assays (Figure 1C). The millimolar Mn²⁺ concentrations that induce a strong gp2 nuclease activity leading to complete degradation of linear DNA are not found in the cell. On the contrary, the terminase has an endonuclease activity that is well regulated and not frequent event during the encapsidation cycle which is compatible with the gp2 properties found at physiological, micromolar, concentrations of Mn²⁺. Interestingly, the affinity of the endonuclease domain of the influenza virus polymerase PA subunit for Mn²⁺ ions (Kds for the two sites of 0.3 and 6.5 μM) was found to be 500- to 600-fold higher than for Mg²⁺ ions (35). A role for both ions in viral nucleases activity strict regulation and structure stabilization is thus plausible.

Our study showed that the gp2 nuclease domain plays essential roles at different steps of viral DNA encapsidation. The novel plasmid assay developed to measure in vivo the initiation event of SPP1 DNA packaging showed that the pac binding protein gp1, the terminase small subunit (13), and gp2 are necessary and sufficient to reproduce faithfully sequence-specific cleavage at pac (12) in absence of other phage products (Figure 6). Gp1 creates the structural context for gp2 cleavage at pac (13,48) and prevents non-specific nuclease attack to one of the DNA ends, the substrate for packaging. The SPP1 DNA end destined for packaging is protected from cellular exonucleases most likely by the gp1–gp2 complex where the nuclease activity is strictly shut off (15). The low frequency of pac cleavage observed (Figure 6) is consistent with the requirement that the sequence pac is used for cleavage only once ‘per’ packaging series along a substrate concatamer while the other pac sites remain intact (6,12,49). The gp1–gp2–pac nucleoprotein complex thus fully ensures the auto-regulated level of pac cleavage and directionality of the subsequent encapsidation event by selectively protecting one DNA end.

The gp2 nuclease CTD is then involved in specific docking to the procapsid portal vertex (Figure 3) as found in previous genetic and biochemical studies in phages T3, T4 and λ (29–32). As observed for the nuclease activity, binding of gp2 CTD to the portal is nevertheless less efficient relatively to FL gp2. It is possible that other gp2 residues, probably comprising the linker region between the NTD and CTD (8,50), are also involved in direct portal binding or are required to keep gp2 in an optimal conformation for the interaction with the portal. Docking of gp2 at the procapsid portal vertex leads to significant stimulation of the gp1:gp2 ATPase activity (17) consistent with switching on the gp2 translocase activity and suppression of any nuclease activity while DNA moves through the packaging motor. Pumping of the viral concatemeric DNA to the procapsid interior requires the hydrolysis of ATP that is catalyzed by the gp2 NTD, as observed for other viral terminases (28,51). Remarkably, some mutations in the conservative gp2 nuclease catalytic site residues (D266N, H400A and D266N/D403N) block DNA encapsidation. This was demonstrated in DNA packaging assays in vitro with a linear substrate, which does not involve a DNA cut [(16); Figure 8]. All the nuclease mutations have regular ATPase activity and binding to the procapsid portal vertex (Figure 7 and Supplementary Figure S5). In addition, the nature of the metal ion present affects the DNA packaging capacity of two gp2 nuclease mutant proteins D403N and D321N/D403N that are active in presence of Mn²⁺ but not of Mg²⁺. The role of the nuclease center in DNA encapsidation is not strictly dependent on binding of metal ions to the nuclease active site as shown by the observation that gp2 D266N/D321N supports DNA encapsidation. It rather results from the structural organization of the nuclease active site and probably of its harboring cleft to support DNA translocation that likely occurs in their vicinity.

After filling of the viral capsid with DNA, gp2 integrates signaling from the portal (21,52) to stop translocation and switching to endonucleolytic activity that cleaves the DNA substrate (the unspecific headful cut) to end the genome packaging cycle. A mutant in the SPP1 portal protein (gp6T319A) that packages small size DNA (22) leads to less efficient ATP hydrolysis by gp2 and slow packaging kinetics (17). Gp6 can thus render the gp2 DNA translocation machine less effective and cause a premature switch from gp2 translocase to headful nuclease that yields packaged viral chromosomes shorter than wt mature length. It is likely that this intimate cross-talk is mediated by interaction of the gp2 CTD with gp6. Our study together with available data show that the nuclease domain is the coordination center of the DNA packaging motor that integrates signals from the phage pac recognition terminase subunit, the ATpase domain, and the portal protein to switch between nuclease and translocase activities that are essential to achieve the viral genome encapsidation cycle in tailed phages and herpesviruses.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online: Supplementary Figures 1–5.

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