Biochemical and Biophysical Properties of a Putative Hub Protein Expressed by Vaccinia Virus

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Background: Vaccinia protein H5 is a multifunctional protein involved in several aspects of viral replication. H5 is a nucleic-acid binding, elongated rod-like tetramer with an intrinsically disordered N terminus and an α-helical C terminus. H5 functions as a hub protein in vaccinia virus replication linking otherwise unique viral processes. Understanding the mechanism of H5 function in vaccinia virus replication is crucial for studying vaccinia biology.

Results:

H5 is a constitutively expressed, phosphorylated vaccinia virus protein that has been implicated in viral DNA replication, post-replicative gene expression, and virus assembly. For the purpose of understanding the role of H5 in vaccinia biology, we have characterized its biochemical and biophysical properties. Previously, we have demonstrated that H5 is associated with an endoribonucleolytic activity. In this study, we have shown that this cleavage occurs in a 3’-OH end suitable for polyadenylation of the nascent transcript, corroborating a role for H5 in vaccinia transcription termination. Furthermore, we have shown that H5 is intrinsically disordered, with an elongated rod-shaped structure that preferentially binds double-stranded nucleic acids in a sequence nonspecific manner. The dynamic phosphorylation status of H5 influences this structure and has implications for the role of H5 in multiple processes during virus replication.

Conclusion: Vaccinia virus is the prototypical and best studied member of the poxvirus family. Members of this family are large (~200 kb) double-stranded DNA viruses that replicate exclusively in the host cell cytoplasm and therefore encode all the enzymes and factors required for DNA and RNA metabolism (1). Viral transcription is regulated as a temporal cascade with three different classes of transcripts (early, intermediate, and late) (1, 2). Transcription requires a multi-subunit RNA polymerase, several stage-specific transcription factors, and multi-subunit enzymes for cap formation and polyadenylation. Early transcription occurs before viral DNA replication and early gene products serve as factors for viral DNA replication and intermediate transcription. Intermediate and late transcription (collectively termed post-replicative) are coupled to viral DNA replication. Each stage of transcription employs different mechanisms of transcription termination. Early transcription termination requires a cis-acting sequence and several trans-acting factors yielding 5’ and 3’ co-terminal transcripts (3–9). Post-replicative transcription termination results in heterogeneous length transcripts, 1–4-kb-long, that are 5’ co-terminal (10). Although several factors have been shown to modulate post-replicative transcription termination, no cis-acting sequence similar to early transcription termination has been identified (11–13). However, a pyrimidine-rich sequence in the coding strand has been identified immediately preceding the 3’ ends of these post-replicative transcripts (14). Viral DNA replication requires a viral DNA polymerase (E9), a heterodimeric processivity factor (A20 and D4), and several other accessory proteins (I3, D5, H5, B1, J2, F4/I4, G5, A22, A50, I6, and A32) that support viral DNA replication either directly or indirectly (1).

Viral DNA replication, postreplicative transcription, and virion assembly take place in discrete cytoplasmic sites termed virus factories. Virion assembly and morphogenesis involves the formation of crescent shaped viral membrane structures within factories, followed by packaging of condensed viroplasm and DNA within the viral membranes to form spherical immature virions and immature virions with nucleoids. Proteolysis of major virion structural proteins and metamorphosis of the immature virions with nucleoids into mature virions is required before transport of the mature virions out of the factory. Several viral proteins are required for all of these events (15).

Ongoing investigations of poxvirus RNA metabolism, DNA replication, and virion morphogenesis have consistently highlighted a role for the vaccinia H5 protein in all of these processes. H5 is absent from the entomopoxviruses but present in all chordepoxviruses. H5 is a constitutively expressed, 22.3-kDa phosphoprotein that is localized in viral factories and packaged into virions (16–21). H5 is phosphorylated by both viral kinases (F10 and B1) and by cellular kinases; the pattern of its phosphorylation appears to be dynamic and temporally regulated (15, 18). Although the native molecular mass of the H5 polypeptide based on its amino acid sequence is only 22 kDa, this protein behaves aberrantly in several biochemical estimations of molecular weight, suggesting that it may exhibit transient polymorphic structures. H5 fractionates on SDS-PAGE at 35 kDa; this anomaly has been attributed to a rigid amphipathic...
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helix located in the C terminus of the protein and/or to a proline-rich domain in the N terminus capable of producing kinks in the protein structure (18, 19). Another unusual characteristic of the purified protein is that it elutes from gel filtration columns with an extremely large apparent molecular mass (>400 kDa) regardless of the source of purified protein (vaccinia-expressed or Escherichia coli-expressed), suggesting multimerization of the protein and possibly an unusual shape/structure (22).

There is substantial evidence to suggest that H5 is involved in viral DNA replication. H5 interacts with viral DNA replication enzymes A20, the viral DNA polymerase processivity factor and B1, the viral protein kinase that inhibits the host antiviral protein, BAF, during viral DNA replication (23, 24). A temperature-sensitive mutant (Dts57) that maps to the H5 gene has been characterized to have a block in viral DNA replication (DNA-negative) confirming a direct role for H5 in viral DNA replication (25).

In addition to playing a role in DNA replication, H5 has been shown to function in all aspects of post-replicative transcription: initiation, elongation, and termination. Kovacs and co-workers (20) have shown that H5 possesses a post-replicative stimulatory activity in vitro. In addition, several studies (23, 26, 27) have shown that this protein interacts with viral late transcription initiation (A2 and G8), elongation (G2), and termination (A18) factors. Importantly, an H5 mutant has been isolated that is resistant to isatin β-thiosemicarbazone, an antivopoxvirus drug known to affect vaccinia postreplicative gene elongation, further cementing a role for H5 in postreplicative gene transcription (28). As mentioned earlier, post-replicative transcription termination results in 3' heterogenous length transcripts. However, at least four late genes have been identified to yield transcripts that have homogenous 3' ends (29–33). For two of these exceptions (ATI and F17R), the uniform 3' ends are generated by post-transcriptional site-specific endoribonucleolytic cleavage. We have characterized the vaccinia H5 protein as being associated with this activity, suggesting a role for H5 in late transcription termination (22). This cleavage was characterized to be possibly RNA sequence or structure specific, and phosphorylation was determined to be essential for cleavage activity.

Consistent with an apparent role for H5 in viral DNA replication and post-replicative transcription, published data suggest that H5 is a nucleic acid binding protein with affinity for both DNA and RNA (18, 20, 34, 35). Vaccinia-expressed H5 was purified using either single-stranded or double-stranded DNA cellulose columns in these studies (18, 20, 34). However, an extensive study characterizing this binding has not been performed.

Finally, a role for H5 in virion morphogenesis is evident by phenotypic analysis of two temperature-sensitive mutants, tsH5–4 and Dts57 (25, 36). Data from these studies indicate that H5 is involved in multiple steps of viral morphogenesis, including membrane biogenesis, encapsidation of viroplasm to form immature virions, and maturation of immature virions into mature virions.

In the current study, we have characterized the biochemical and biophysical properties of H5 and propose that H5 functions as a hub/scaffold protein that links otherwise discrete vaccinia virus processes. To understand the role of H5 in post-replicative transcription termination, we have determined the nature of 3' ends during the endoribonucleolytic cleavage reaction and determined that H5-associated cleavage results in a 3'-OH end. We have further defined the nucleic acid binding properties of the H5 protein and shown that this protein can bind nucleic acids with high affinity in a sequence nonspecific manner requiring a minimum of 40 nt. Using structure modeling, circular dichroism, sedimentation equilibrium, and small-angle x-ray scattering (SAXS),2 we have demonstrated that the basic structure of unphosphorylated H5 protein is an elongated rod-like tetramer with the intrinsically disordered N terminus of each subunit contributing to its rod-like structure. Phosphorylation of the protein likely causes aggregation of H5 tetramers to form higher order complexes.

EXPERIMENTAL PROCEDURES

Eukaryotic Cells, Viruses, and Bacterial Hosts—BSC40 cells, wild type vaccinia strain Western reserve (WR), and the conditions for their growth and infection have been described previously (37–41). HeLa S3 cells were grown in suspension culture in Eagle’s minimal essential medium (Joklik’s modification containing 5% calf serum and 2% fetal calf serum. E. coli BL21(DE3) harboring the plasmid pET14b and containing the H5 ORF was a gift from Dr. Paula Traktman.

EhisH5 is a recombinant vaccinia virus that contains the 10x His and factor Xa sequences (MGHHHHHHHHHHHHSH-GHIEGRH) from plasmid pET16b at the N terminus of the endogenous H5R. For the construction of ehisH5, BSC40 cells were infected with Dts57-WR virus (Dts57-WR is a reconstituted temperature-sensitive virus in the WR background that contains the Dts57 lesion, G189R in the C terminus of the WR-H5 protein) and transfected with a PCR product that contained the H4R-H5R region of vaccinia WR with the pET16b polyhistidine and factor Xa sequence inserted at the N terminus of H5R. Infection transfection was carried out at 37 °C to select for recombinant WT virus. The harvested lysates were then plaque assayed. Several plaques were picked and analyzed by Western blotting using an anti-His monoclonal antibody (Novagen). Plaque lysates that tested positive by Western blotting were then propagated in cell culture and the sequence of His-tagged H5 was confirmed by Sanger sequencing.

Purification of His-H5 Protein—The purification schemes for vaccinia-expressed and E. coli-expressed His-tagged H5 proteins were as described previously (22). For further purification by gel filtration, the H5 protein eluted from the nickel affinity column (His-Trap, GE Healthcare) was pooled, concentrated using an Amicon concentrator (Ultra cel 5K; Millipore) to ~500 μl and applied to a Superdex 200 gel filtration column (GE Healthcare) equilibrated in 25 mM Tris-HCl, pH 7, containing 1 M NaCl and protease inhibitor mixture (1:100, Sigma). (This buffer also contained 5% glycerol and/or 2 mM DTT in situations where these reagents did not affect downstream events.) Molecular weight standards (high and low molecular

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2 The abbreviations used are: SAXS, small-angle x-ray scattering; TAP, tobacco acid pyrophosphatase; nt, nucleotide; ts, temperature-sensitive; pCP, cytidine-3',5'-bis-phosphate.
weight standards; Amersham Biosciences) were used to calibrate the gel filtration column.

**Determining the Nature of 3' Ends of H5 Cleaved RNA**—The cleavage substrate (430-nt ssRNA) was prepared by *in vitro* transcription and uniformly labeled with $^{32}$P α-CTP as described previously (33). The 430-nt RNA substrate was either directly used in the cleavage assay or 30 pmol of RNA were incubated at 30 °C for 10 min with 30 units of tobacco acid pyrophosphatase (TAP; Epicenter, 10 units/μl) in a total volume of 6.6 μl. Cleavage was carried out under standard cleavage conditions as described previously by incubating 10 fmol of RNA substrate with vaccinia-expressed nickel affinity-purified H5 protein in a reaction containing 25 mM Tris-HCl, pH 8, 7.5 mM MgCl$_2$, and 300 mM NaCl for 1 h at 30 °C (33). The reactions were then treated with either 1 unit of Terminator exonuclease (Epicenter) or buffer (as control) for 2 h at 37 °C. Following Terminator exonuclease treatment, 30 μl of formamide-loading buffer was added, and the samples were denatured at 90–95 °C for 5 min, flash cooled, centrifuged, and loaded on a 6% polyacrylamide gel containing 8 μm urea (Sequagel, National Diagnostics). The gels were then fixed, dried, and analyzed using the STORM PhosphorImager and ImageQuant software (GE Healthcare). For $[^{32}\text{P}]$pCp labeling, the cleavage substrate was prepared using the RNA megascript kit (Ambion) to yield large quantities of “cold” substrate. The substrate was then cleaved with vaccinia-expressed nickel affinity-purified H5 protein under the conditions mentioned above. The residual substrate and the cleaved products were then purified using phenol-chloroform extraction and ethanol purification and resuspended in nuclease-free water. This RNA (and RNA not subjected to the cleavage reaction) was then labeled in a reaction containing 150 μCi of $[^{32}\text{P}]$pCp (300 Ci/mm mole; PerkinElmer Life Sciences) and 10 units of RNA ligase (New England Biolabs) at 4 °C overnight. The labeled RNA was then purified using phenol-chloroform extraction and ethanol purification. The pellets were resuspended in 10 μl of formamide-loading dye and denatured at 90–95 °C for 5 min, flash cooled, centrifuged, and loaded on a 6% polyacrylamide gel containing 8 μm urea (Sequagel, National Diagnostics). The gels were then processed as described above.

**Filter Binding Assays**—Double sandwich filter binding assays were carried out following instructions of Lohman and Wong (42) with some modifications. One hundred microliter reactions were prepared in duplicate by equilibrating various concentrations of *E. coli*-expressed H5 (or vaccinia-expressed H5) with 10–100 pm nucleic acid (1 μl/reaction containing 1–10 fmol nucleic acid of different lengths and sequence as described under “Results”) on ice in rinse buffer (25 mM HEPES, pH 7.5, 50 mM NaCl, 1 mM EDTA, 5 mM DTT) containing 10 μg of BSA. After 30 min, 90 μl of the reaction mixture was vacuum filtered through a nitrocellulose (Bio-Rad) and nylon membrane (Hybond N+, GE Healthcare) sandwiched within a Schleicher and Schuell minifold. An equal volume of rinse buffer was then filtered through to rinse out the wells. The membranes were then separated, dried, and analyzed using a STORM PhosphorImager and ImageQuant software (GE Healthcare). The percentage of nucleic acid binding to the nitrocellulose membrane was computed from these results and plotted against increasing concentrations of H5 protein using GraphPad Prism software to estimate a dissociation constant ($K_D$) for each binding reaction. The $K_D$ for each experiment was determined using technical duplicates and the results shown are from at least two different experiments. All of the optimization data shown in Fig. 2 were obtained using 100 pm $^{32}$P-labeled ssRNA (430 nt; uniformly labeled) and vaccinia-expressed H5 protein (ehisH5). Experiments at pH 6 were done in MES buffer, those at pH 7–8 were done in HEPES, and those at pH 10 were done in 3-(cyclohexylamino)propanesulfonic acid buffer. The pH of all buffers was adjusted either with HCl or NaOH. 5'-Cy5-tagged nucleic acids (Sigma Genosys) of varying lengths were used in the binding experiments that determined length dependence of H5 binding. All $^{32}$P-labeled substrates were gel-purified; 5'-Cy5-tagged substrates were obtained PAGE-purified from the manufacturer. For the preparation of double-stranded 5'-Cy5-tagged nucleic acids, complementary sequences of equal lengths and concentration were mixed in 1× TE buffer containing 50 mM NaCl and annealed by heating to 100 °C for 10 min in a Thermocycler. The temperature was then decreased to 4 °C gradually using a gradient of 1 °C/min. The hybrids were then analyzed on non-denaturing polyacrylamide gels. Greater than 95% of the annealed oligonucleotides were observed as double-stranded molecules and, therefore, these hybrids were not gel-purified.

**Intrinsic Disorder Prediction and Homology Modeling**—The PONDR-Fit algorithm was used to calculate the intrinsic disorder disposition in vaccinia H5 protein (43). The PONDR-Fit intrinsic disorder disposition versus residue number plot was scaled by sequence alignment to the published sequence of H5 (vaccinia WR) to better compare the different regions on the H5 monomer. Homology models of H5 were also made based on the published H5 sequence (WR) using the ROBETTA server (44). These were combined and a composite H5 model made, H5model.

**CD**—The CD data were collected on an Aviv 414 CD spectrometer. The data were collected between 200 and 260 nm at 25 °C. Three hundred and fifty microliters of sample were used at 0.4 mg/ml of H5. The data, collected in triplicate, were averaged across the 50 scans and the triplicate experiments to improve the signal to noise ratio. The averaged data were then scaled to molar ellipticity values to account for concentration differences and to compare with other CD spectra. The collected data were processed using in-house algorithms to determine the secondary structural propensity of H5 (45).

**SAXS**—The SAXS data were collected at the Cornell High Energy Synchrotron Source MacCHESS G1 beamline. The data were collected at a wavelength of 1.444 Å (8.58 keV). Twenty-five second exposures were used at a detector distance of 800 mm to collect data between q values of 0.008 and 0.3 nm$^{-1}$. Two milligrams per milliliter sample concentrations were used with 30 μl of sample volume in plastic cuvettes. 10× and 20× attenuation levels were used to achieve optimum signal to noise ratio while at the same time having minimal radiation damage to the sample. The images were processed for intensity and q values using the DATASQUEEZE (University of Pennsylvania) and BioXTAS RAW programs (46).
The ATSAS suite of programs was used to further process the data after raw image processing (47). GNUMPLOT was used for data plotting and radius of gyration ($R_g$) calculations. Pairwise distribution functions and radius of gyration values were calculated using the GNOM program from the ATSAS suite. DAMMIN was used to generate the three-dimensional ab initio model. Ten DAMMIN simulations were averaged using the DAMAVER program to generate a final density profile. This was converted to a SITUS volume map for docking purposes and the H5model was docked manually into the map using the CHIMERA program (48). Theoretical SAXS curves from the H5model structure of the monomer and possible multimers were computed using the CRYSTOL. GNOM, DAMMIN, DAMAVER, and CRYSTOL are all located within the ATSAS suite. All visualizations were made using University of California, San Francisco Chimera.

Sedimentation Equilibrium Ultracentrifugation—Sedimentation equilibrium were performed using absorption optics in a Beckman XLA analytical ultracentrifuge at 4 °C in 25 mM Tris, 1 mM NaCl, and 2 mM DTT, pH 7, and analyzed as described previously (49, 50). Absorption was measured at 280 nm as a function of radius for samples of 100 and 150 μl. The gradient was invariant for 24 h after equilibrium was achieved at 6,200 rpm, and then additional equilibrium data were obtained at 13,200. Samples were examined after the run by SDS-PAGE and showed no evidence of degradation. The subunit partial specific volume was estimated from the amino acid content. The four sets of data (two samples of different volumes, each at two different angular velocities) were fit simultaneously with the Marquard-Levenberg method was employed for nonlinear, least-squares curve fitting.

RESULTS

The Nature of 3' Ends Generated by H5 Cleavage of RNA

We have shown that vaccinia virus protein, H5 is associated with endo-ribonucleolytic cleavage of at least two homogenous late transcripts (ATI and F17) in vitro (22). Furthermore, because H5-associated cleavage activity is postulated to function at the level of vaccinia post-replicative termination (22), we hypothesize that cleavage would result in a free 3'-hydroxyl end unlike most RNases, including RNase A and T1, which cleave RNA to yield a 3' phosphate. The substrate for H5-associated cleavage is a 430-nt uniformly labeled, in vitro-synthesized ssRNA with a 5'-triphosphate. Cleavage of this RNA yields two products: the upstream product is 190 nt in length, and the downstream product is 240 nt in length (Fig. 1A). The cleaved products and the residual substrate are routinely fractionated on a polyacrylamide gel, and the results are analyzed by phosphorimaging analyses as shown in Fig. 1B. To determine the nature of the 3' ends during the H5-associated cleavage reaction, we treated the H5 cleavage products with Terminator exonuclease, an RNase that degrades 5'-monophosphate. 5'-monophosphate 430-nt ssRNA. If H5-associated cleavage of the 430-nt ssRNA substrate results in a 5'-PO4 (scenario I) on the downstream cleavage product, treatment with Terminator exonuclease (-TAP) will cause degradation of the 240-nt downstream product of H5 cleavage but not of the 430-nt substrate and the 190-nt upstream product (outcome I). This result was observed in lane I (Fig. 1B), indicating that H5-associated cleavage results in a 5'-PO4. However, if the 430-nt ssRNA substrate is first treated with tobacco acid pyrophosphatase (+TAP) prior to the H5-associated cleavage, the 430-nt ssRNA substrate will also have a 5'-monophosphate. Under these conditions, if H5-associated cleavage of the 430-nt ssRNA substrate results in 5'-PO4 (scenario I), treatment with Terminator exonuclease will degrade both the substrate and the two cleavage products (outcome I). This result was observed in lane 4 (Fig. 1B), confirming that H5-associated cleavage of the ssRNA substrate results in a 5'-PO4. To confirm these data, we also

![FIGURE 1. Endoribonucleolytic cleavage with purified H5 protein results in a 3'-OH. A, schematic of potential cleavage products for the TAP-treated (+TAP) or untreated (−TAP) 430-nt ssRNA substrate. I and II, possible scenarios depending on the nature of 3' ends after cleavage.

Ends denoted in boldface type indicate possible results of cleavage. B, schematic of the potential and actual products after treatment of the cleavage reaction with Terminator exonuclease (Term). I and II, possible outcomes depending on the nature of 3' ends after cleavage. Gray line, RNA degraded by Terminator; black line, RNA not degraded by Terminator. C, schematic of the potential and actual products after [5'-32P]pCp treatment of unlabeled, purified RNA substrate and products from a cleavage reaction. Black line, RNA labeled with [5'-32P]pCp; gray line, RNA not labeled with [5'-32P]pCp. Lane 1, RNA substrate treated with H5 in cleavage assay, purified, and labeled with [5'-32P]pCp; lane 2, RNA substrate treated with buffer in cleavage assay, purified, and labeled with [5'-32P]pCp; lane 3, RNA substrate labeled with [5'-32P]pCp.](image-url)
analyzed the H5-cleaved products for the presence of a 3′-OH on the upstream H5 cleavage product (190 nt). To do so, we prepared unlabeled (cold) 430-nt ssRNA substrate in vitro and treated it with cleavage-competent H5 or enzyme dilution buffer under the normal conditions for H5-associated cleavage as mentioned previously. The residual substrate and the two cleaved products from H5 or control reactions were then purified and labeled with [5′-32P]pCp using RNA ligase (Fig. 1C, lanes 1 and 2). Cleavage substrate not subjected to the cleavage reaction was also labeled as described above (Fig. 1C, lane 3). Because H5-associated cleavage results in a 3′-OH as determined in Fig. 1B, we observed 32P-labeling of residual substrate and both cleaved products as shown in Fig. 1C, lane 1, confirming that H5-associated cleavage of the ssRNA substrate results in a 3′-OH.

Protein–Nucleic Acid Interaction

There is some evidence that H5 is a nucleic acid binding protein (18, 20, 34). Given the role of H5 in both DNA and RNA metabolism, we were interested to determine the parameters of H5-nucleic acid interaction. In addition, to better understand the function of its nucleic acid binding characteristics, we determined whether H5 has a preference for one type of nucleic acid over another. To determine the binding affinity of H5 for each nucleic acid, we employed the double-sandwich filter binding assay based on the method of Wong and Lohman (42) and as described under “Experimental Procedures.” Given that vaccinia-expressed H5 (but not E. coli-expressed H5) is associated with endoribonucleolytic cleavage, we first analyzed the binding of vaccinia-expressed H5 to the in vitro-generated 430-nt ssRNA cleavage substrate and to its reverse complement, which is not a substrate for cleavage (33). Both substrates bound vaccinia-expressed (cleavage competent) affinity-purified H5 protein comparably and with high affinity (K_D 4–45 nM), suggesting that binding of H5 to nucleic acid is nucleic acid sequence-independent (data not shown). Additionally, we showed that both vaccinia-expressed (cleavage competent) and E. coli-expressed (cleavage incompetent) affinity-purified H5 bound the 430-nt ssRNA cleavage substrate with comparable affinities (K_D 10–50 nM), suggesting that nucleic acid binding and cleavage activity are discrete events (data not shown).

Interaction Parameters

Effect of Temperature on Binding—To determine the optimal temperature for studying the nucleic acid binding properties of H5, we analyzed the binding of affinity-purified, vaccinia-expressed H5 to the 430-nt ssRNA cleavage substrate at three different temperatures (Fig. 2A). The K_D of binding at the three temperatures were as follows: 20.5 ± 6.0 nM (ice); 45.5 ± 11.1 nM (room temperature); and 50.0 ± 27.2 nM (30 °C). A plot of these dissociation constants as a function of temperature (Fig. 2B) suggests that temperature did not drastically affect the affinity of H5 for this RNA. Because the K_D of the reaction was lowest when the binding reaction was incubated on ice, we have used this temperature for the remainder of the filter binding assays.

Effect of Salt on Binding—Given that binding of H5 to the cleavage substrate was sequence-independent; we examined the role of electrostatic interactions in the binding of H5 to the 430-nt ssRNA substrate as a function of ionic strength using 50–500 mM NaCl. No binding was observed in 500 mM salt at the concentrations of H5 tested (data not shown). The binding of H5 to the RNA substrate at salt concentrations between 50–300 mM NaCl. The data plotted are from four replicates, and the mean and S.E. are plotted. B, temperature dependence of K_D. Dissociation constants for H5 binding were plotted as a function of the temperatures tested in the assay. C and D, effect of salt on nucleic acid binding. C, equilibrium binding curves for H5 binding to 430-nt ssRNA between 50–300 mM NaCl. The data plotted are from four replicates, and the mean and S.E. are plotted. D, salt dependence of K_D. Dissociation constants for H5 binding were plotted as a function of the salt concentration tested in the assay. E and F, effect of pH on nucleic acid binding. E, equilibrium binding curves for H5 binding to 430-nt ssRNA between pH 6–10. The data plotted are from four replicates, and the mean and S.E. for each point are plotted. F, pH dependence of K_D. Dissociation constants for H5 binding were plotted as a function of the pH tested in the assay.
nucleic acid. HEPES buffer, pH 7.5, was chosen as the buffer for characterizing the rest of the binding conditions.

**Substrate Specificity**—Next, we compared the affinity of H5 for single-stranded and double-stranded DNA and RNA. Fig. 3 shows the saturation curves of H5 binding to 20-, 40-, 68-, and 91-nt length oligomers of ssDNA, ssRNA, dsDNA, dsRNA, and DNA:RNA hybrids. Because the binding of cleavage competent vaccinia-expressed H5 and cleavage incompetent *E. coli*-expressed H5 to nucleic acid was comparable, we used *E. coli* expressed H5 for the remainder of the binding experiments. Additionally, the nucleic acid substrates for this set of experiments were synthesized with a Cy5 label on the 5′ end of the nucleic acid oligomer. For all of the nucleic acids tested, H5 was unable to bind with high affinity to any of the 20-nt nucleic acid oligomers. Additionally, the binding of H5 to the 20-mer of every set of nucleic acid oligomers tested was significantly different from the binding to the 40–91-mers within the same set (p ≤ 0.0001). The average dissociation constants (between 6 and 18 replicates were averaged for each data set) for H5 binding to each oligomer in all five sets of nucleic acids are shown in Table 1. Using the unpaired Student’s t test, we determined that with the exception of ssRNA, within a given set, the dissociation constants for H5 binding to lengths of 40–91-nt oligomers were not significantly different from each other based on 95% confidence (p ≥ 0.16). For ssRNA, there appeared to be a significant difference in dissociation constants (p = 0.01) between H5 binding 40-nt ssRNA versus 91-nt ssRNA only. Therefore, excluding the data obtained for the 20-mers, we calculated a mean $K_D$ for H5 binding to each of the nucleic acid polymers as shown in Table 1 and in supplemental Fig. 1 using row mean statistics. The binding affinity of H5 to the five different sets of nucleic acid polymers sorted themselves into three categories. The $K_D$ was the lowest for binding of H5 to dsDNA (13.9 ± 1.6 nM) and dsRNA (11.7 ± 2.1 nM), intermediate for DNA:RNA hybrids (21.4 ± 3.9 nM) and the highest for H5 binding to ssDNA (62.9 ± 8.8 nM) and ssRNA (44.2 ± 10.4 nM). The binding of H5 to double-stranded nucleic acids (DNA and RNA) was significantly different from binding to single-stranded nucleic acids and both these categories were significantly different from binding to DNA:RNA hybrids (supplemental Fig. 1). The data indicate that H5 preferentially binds homoduplexed nucleic acids best, followed by heteroduplexed nucleic acid. The binding to single-stranded nucleic acids in comparison was weakest.

**Biophysical Properties of H5**

**Structure and Modeling**—H5 is phosphorylated and consists of 203 amino acids. Potential sites of phosphorylation include 18 serines, 19 threonines, and 1 tyrosine interspersed through the length of the protein. A single cysteine is present at the C terminus of H5 (Fig. 4A). Within the poxvirus family, H5 is present in the chordopoxviruses but absent from the entomopoxviruses. Among the chordopoxviruses, different H5 orthologs within the orthopoxvirus genera show strong conservation (data not shown). However, only a modest conservation is observed for the H5 orthologs within different genera of the chordopoxviruses. The H5 polypeptide sequence for representatives from each chordopoxvirus genera were aligned using the multiple sequence alignment software, MUSCLE at the European Bioinformatics Institute website (supplemental Fig. 2). Significant conservation was observed in the most N-terminal region and in the C terminus of the protein, whereas the middle region of H5 (between residues 50–120) appeared non-conserved but high in charged residues. A tertiary structure of H5 was generated using the homology modeling program ROBETTA (Fig. 4, B–G). Five models were generated based on the sequence (Fig. 4, B–F). The models all showed an α-helical propensity at the C terminus and a mixed/helix motif loop throughout the remainder of the structure. Fig. 4G shows a structural superimposition of three best fit models (models depicted in Fig. 4, B–D) and highlights the conserved α-helices predicted.

**TABLE 1**

Dissociation constants for H5 binding to different lengths of nucleic acid

The 20-nt curves did not reach saturation at the concentrations of H5 tested (ND, not determined).

| Nucleic acid | Mean $K_D$ (nM) |
|-------------|----------------|
| ssRNA       | 94.8 ± 16.2     |
| ssDNA       | 69.5 ± 10.8     |
| dsRNA       | 16.4 ± 1.5      |
| dsDNA       | 13.7 ± 1.0      |
| DNA:RNA     | 35.5 ± 6.7      |

**FIGURE 3.** Equilibrium curves for H5 binding to different lengths of nucleic acid oligomers. The $K_D$ for each curve is computed with data obtained from at least three individual experiments with each experiment done in duplicate, and the mean and the S.E. for each data point is plotted. A, ssRNA; B, ssDNA; C, dsRNA; D, dsDNA; E, DNA:RNA hybrids.

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in the C terminus of H5 and ~ 50% of the N terminus as unstructured or disordered. Previously, we observed that purified H5 (both cleavage-competent and -incompetent) eluted from a gel filtration column with a significantly large apparent molecular mass (~460 kDa), suggesting either complex formation or a non-globular structure (22). To further analyze these findings, the protein sequence of H5 was examined using the PONDR-Fit algorithm that predicts the degree of disorder in the secondary structure of a protein (Fig. 4). The PONDR-fit algorithm predicted a high intrinsic disorder disposition in the N-terminal region of the H5 protein, shown as loops in the homology model (Fig. 4I). Experimentally, these data were confirmed using circular dichroism (Fig. 5). Secondary structural propensity as determined by the CD data processing indicated a 50% α-helix and a 50% random coil propensity, which are in agreement with the homology models. Interestingly, the CD data also showed a profile consistent with an intrinsically disordered or unstructured protein.
tendency to form higher order complexes as a result of phosphorylation.

Given the discrepancy between the apparent molecular mass of H5 on a gel filtration column and by sedimentation equilibrium ultracentrifugation, we were interested in analyzing the structure of H5 by yet another independent biophysical measurement, SAXS (Fig. 7). The SAXS data indicated that the protein showed a tendency to aggregate. The apparent intrinsic disorder of the protein might have a role in the aggregation process. A low resolution three-dimensional reconstruction based on SAXS data showed an elongated rod-like structure that fitted four of the homology model monomers. Consistent with the sedimentation equilibrium data, the SAXS data suggests that H5 exists as a tetramer in solution. The radius of gyration determined from the data is also in agreement with the dimensions of a tetramer.

**DISCUSSION**

In this study, we have characterized the biochemical and biophysical properties of the multi-functional vaccinia protein, H5. We have shown that H5 endoribonucleolytically cleaves the ssRNA cleavage substrate *in vitro* to yield a 3′-OH end. H5 binds both single-stranded and double-stranded nucleic acids, requiring at least 40 nt for efficient binding but prefers double-stranded nucleic acids over single-stranded as follows: dsDNA/dsRNA > DNA:RNA > ssDNA/ssRNA. The interaction of H5
with nucleic acid is sequence nonspecific and electrostatic: salt affects binding and pH has no effect on binding, suggesting binding to the phosphodiester backbone. The N terminus of H5 is intrinsically disordered but the C terminus has an α-helical structure. The purified E. coli-expressed H5 protein is a stable and homogenous tetramer (90 kDa) with an elongated rod-like structure. We suggest that the unusual rod-like shape of the tetrameric protein in conjunction with the intrinsic disorder of the N terminus of each monomer contributes to its large apparent molecular mass upon gel filtration. Gel filtration data consistent with apparent mass greater than actual mass are expected for intrinsically disordered proteins because of increased hydrodynamic volume relative to that of ordered proteins of the same mass (51). However, the molecular mass of purified vaccinia-expressed H5 appears to be more complex with a heterogeneous mix of protein structures ranging from tetramers to octamers and even higher order structures. We predict that the dynamic phosphorylation pattern of vaccinia-expressed H5 plays a key role in this difference.

Viral proteins, in general, are very abundant in intrinsically disordered regions (52–55). Intrinsically disordered proteins have been identified in both DNA and RNA viruses with a larger number present in RNA viruses (52–55). However, unlike the intrinsically disordered proteins found in eukaryotes, viral intrinsically disordered proteins are characterized to have short, disordered segments. Because viral genomes are unusually compact with overlapping reading frames, a single mutation can affect more than one protein. Additionally, the viral lifestyle requires multiple interactions with proteins, membranes, DNA, and RNA of the host in conjunction with the ability to adapt to new/alternative hosts. Thus, the ability to interact with different and multiple partners accompanied with the high rate of evolution can be linked to disorder and conformational flexibility.

The cleavage of ssRNA by H5 to yield a 3′-OH is consistent with its role in post-replicative transcription termination. Although most RNases such as RNase A and RNase T1 cleave RNA to form a 3′-PO₄, other specialized RNases, including β-lactamases (e.g. CPSF73, the cleavage factor of the eucaryotic cleavage and polyadenylation complex, and RNase Z, the tRNA maturation nuclease), RNase III (Dicer), and RNase H (cleaves DNA:RNA hybrids) produce a 3′-OH during cleavage (56–59). A 3′-OH in the cleavage reaction suggests that the nascent transcript can be a substrate for vaccinia poly(A) polymerase during vaccinia post-replicative transcription. Alternatively, the endoribonucleolytic function of H5 may be involved in viral DNA replication. H5 interacts with a vaccinia protein of unknown function, A49 (23). Blast analysis of the A49 protein sequence against the non-redundant database indicates the presence of a DnaI conserved motif in the C terminus of the protein. DnaI proteins are components of the Bacillus subtilis replication restart primosome (60). Another vaccinia protein required for viral DNA replication, D5, has been characterized as a primase, suggesting the formation of Okazaki fragments on the lagging strand during viral replication (61). Given the role of H5 as a viral endoribonuclease and the fact that H5 binds DNA:RNA hybrids with high affinity, it is possible that H5 has RNase H-like activity and cleaves the RNA primer in the lagging strand during DNA replication. However, preliminary experiments to characterize RNase H activity in H5 were negative (data not shown). Additionally, other nucleases such as the FEN-1-like endonuclease (vaccinia protein G5), the nicking and joining enzyme (K2), and the holiday junction resolvase (A22) are pres-
ent in vaccinia and may be involved in this role for vaccinia DNA replication (62–64).

Independent evidence suggests that H5 exists as a multimeric protein. Wright et al. (27) observed self-interaction of H5 both by yeast two-hybrid analyses and by in vitro pulldown assays. DeMasi et al. (36) have shown that a temperature-sensitive mutant of H5 (tsH5–4) has a dominant negative phenotype: when both a WT and mutant copy of H5 are expressed on the same viral genome, the function of H5 in morphogenesis is affected, suggesting that WT and mutant H5 protein interact with each other affecting function. Additionally, Boyle and Traktman3 have shown that unfrictionated extracts from another temperature-sensitive mutant of H5 (Dts57, affected in DNA replication) elutes with a smaller apparent molecular mass from gel filtration (150 versus 460 kDa for WT protein), suggesting the mutant protein has an altered ability to form multimers. Unlike tsH5–4, the phenotype for Dts57 is not dominant negative (the phenotype of this mutant was rescued using wild type H5). The mutation in tsH5–4 is in the N-terminal region of the protein (EEE to AAA at amino acid positions 73–75), and in Dts57, the C-terminal region of the protein is altered (G189R). Taken together, these observations suggest that the higher order structure of H5 is key to its function both in morphogenesis and DNA replication, and we suggest that this multimerization of H5 is possibly via its C-terminal domain.

Extrapolating from data in this study and previous studies, it is clear that H5 functions as a hub protein in vaccinia virus replication. Hub proteins are highly connected nodes that are capable of binding several partners and linking several independent biological functions (65). These partners can be both nucleic acid and protein. Several hub proteins, including the family of high mobility group proteins (HMGA), have been shown to bind bent or folded DNA in a sequence-nonspecific manner via the minor groove. Most hub proteins are known to have at least some degree of intrinsic disorder in their structure to achieve flexibility while interacting with several binding partners. Finally, post-translational modifications such as phosphorylation, acetylation, and methylation have been shown to add to this flexibility. H5 fits the criteria for all of the above characteristics of hub proteins: it is dynamically phosphorylated, has a highly disordered N-terminal region, can self-interact, and also bind several viral proteins (A20, B1, A2, G2, G8, A18, and A49) and nucleic acids (23, 26, 27, 66).

A recent literature search revealed another protein with surprisingly similar characteristics - coilin (67). Coilin is an intrinsically disordered, 576-amino acid, constitutively expressed protein that is a marker for Cajal bodies but is also present in high abundance in the nucleoplasm. Cajal bodies are sub-nuclear sites of RNA-related metabolic processes, including small nuclear ribonucleoprotein biogenesis and maturation, histone mRNA processing, and telomere maintenance (68). Cajal bodies are assembled during interphase and disassembled during mitosis. Coilin is thought to function as a scaffold to seed Cajal body formation. This protein interacts with itself and with other proteins important for small nuclear ribonucleoprotein biogenesis. Coilin is dynamically phosphorylated: it is hyper-phosphorylated during mitosis causing a decrease in self-interaction and association with its binding partners. This hyper-phosphorylation is thought to cause disassembly of the Cajal bodies. Broome et al. (67) showed that purified coilin binds non-specifically to dsDNA and cleaves U2 pre-snRNA at a C-urach site in vitro. Furthermore, depletion of coilin mRNA in vivo caused an increase in both U1 and U2 pre-snRNA, suggesting that coilin binds non-specifically to chromatin and is required for the maturation of U1 and U2 pre-snRNA in Cajal bodies. Similar to H5, coilin does not bear resemblance to any known ribonucleases. The disordered N terminus of coilin was found to be required for its nuclease activity. Superposing of the H5_model with a coilin_model did not reveal any common structural features.

We have constructed a model that posits H5 as a hub via which several viral functions can be linked (Fig. 8). This model makes several predictions. 1) The minimal structure of H5 is tetrameric and shaped like an elongated rod. 2) The intrinsically disordered N terminus is free and interacts with several protein partners, including A20, B1, G2, and A18 causing an induced-fit conformation change. 3) The α-helical C-terminal domain of each monomer is required for interaction of the protein with itself and for interaction with the sugar-phosphate backbone of nucleic acid. 4) The dynamic phosphorylation status of H5 changes the structure of H5 from tetrameric to higher order structures and also determines the affinity of H5 to its binding partners.

With these predictions in place, we propose that the tetrameric form of the protein requires at least 40 nt of nucleic acid to bind with high affinity. We also propose that H5 binds and coats double-stranded nucleic acids and recruits several viral replication partners, transcription factors, and viral proteins effecting morphogenesis, depending on its phosphorylation status. We propose that membrane formation and early stages of virion assembly, including crescent formation and virospore encapsidation, is facilitated by or occurs on an H5 matrix. The phosphorylation of H5 by F10 may play a critical role in this process. Finally, we believe the role of H5 in cleavage may be unusual. We propose that H5 recognizes structure rather than sequence of the late mRNA around the cleavage site and binds to the double-stranded region to cause cleavage mechanically by kinking the RNA or enzymatically via endoribonucleolytic cleavage. This would cause destabilization of the transcription ternary complex and allow for termination catalyzed by

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3 K. Boyle and P. Traktman, personal communication.
A18, the vaccinia termination factor. This hypothesis is predicated on the torpedo model of transcription termination in eukaryotes (69).

In conclusion, H5 is an intrinsically disordered, sequence-nonspecific double-stranded DNA (or RNA) binding, dynamically phosphorylated protein that has endoribonucleolytic activity in vitro and is capable of self-interaction as well as transient interaction with several other vaccinia viral proteins.

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