Chemical-Shift Perturbations Reflect Bile Acid Binding to Norovirus Coat Protein: Recognition Comes in Different Flavors

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Dedicated to Prof. Jesús Jiménez-Barbero on the occasion of his 60th birthday.

Bile acids have been reported as important cofactors promoting human and murine norovirus (NoV) infections in cell culture. The underlying mechanisms are not resolved. Through the use of chemical shift perturbation (CSP) NMR experiments, we identified a low-affinity bile acid binding site of a human GII.4 NoV strain. Long-timescale MD simulations reveal the formation of a ligand-accessible binding pocket of flexible shape, allowing the formation of stable viral coat protein–bile acid complexes in agreement with experimental CSP data. CSP NMR experiments also show that this mode of bile acid binding has a minor influence on the binding of histo-blood group antigens and vice versa. STD NMR experiments probing the binding of bile acids to virus-like particles of seven different strains suggest that low-affinity bile acid binding is a common feature of human NoV and should therefore be important for understanding the role of bile acids as cofactors in NoV infection.

Introduction

Norovirus infections are the leading cause of viral gastroenteritis infections worldwide.[1] There is compelling evidence that Norovirus infections are the leading cause of viral gastroenteritis infections in cell culture. The underlying mechanisms are not resolved. Through the use of chemical shift perturbation (CSP) NMR experiments, we identified a low-affinity bile acid binding site of a human GII.4 NoV strain. Long-timescale MD simulations reveal the formation of a ligand-accessible binding pocket of flexible shape, allowing the formation of stable viral coat protein–bile acid complexes in agreement with experimental CSP data. CSP NMR experiments also show that this mode of bile acid binding has a minor influence on the binding of histo-blood group antigens and vice versa. STD NMR experiments probing the binding of bile acids to virus-like particles of seven different strains suggest that low-affinity bile acid binding is a common feature of human NoV and should therefore be important for understanding the role of bile acids as cofactors in NoV infection.

Besides HBGA binding sites, bile acids are another important cofactor for the promotion of norovirus infection.[4] Two crystallographic studies addressed the binding of bile acids to human and to murine norovirus capsid protein.[4a,c] For particular strains of human NoVs two symmetrical bile acid binding pockets with affinities in the low-micromolar range have been identified adjacent to the HBGA binding sites.[4c] In the case of a rare GII.1 strain, binding of bile acids promoted attachment to HBGA. Surprisingly, for the dominant epidemic causing GII.4 and GII.17 strains no ligand binding to these pockets was observed, leaving open the question of why bile acids are essential for replication of GII.17 viruses and significantly promote replication of GII.4 viruses in human intestinal enteroids.[4f] Another study based on X-ray crystallography and cryo-electron microscopy has identified two symmetrical bile acid binding pockets at the P-domain dimer interface of a CW3 mouse NoV.[4d] Interestingly, these binding pockets are highly specific for glycochenodeoxycholic acid (GCDCA) with low-micromolar affinity. For human NoVs inspection of available crystal structures of P-domains shows that this binding pocket is inaccessible to bile acid molecules.

Long-timescale molecular dynamics (MD) simulations can reveal transient ligand binding pockets tailored for small molecule binding.[5] A ligand may selectively bind to one or to an ensemble of such pre-existing conformations.[6] The sites of binding are referred to as sub-pockets, adjacent pockets, breathing motion, channel/tunnel, or allosteric pockets.[7] MD simulations sample conformational space and provide snapshots of relevant protein conformations for docking, improving the accuracy of virtual screening over rigid protein docking.[8] The method also facilitates discovery of allosteric sites.[9]
Recently, we obtained a nearly complete backbone assignment of the P-domain of a human epidemic genotype norovirus, GII.4 Saga.\cite{10} We also obtained a complete methyl group assignment of a MILV-la labe l sample of GII.4 Saga P-domain (unpublished data). Based on this work, it is now possible to investigate ligand binding using protein NMR experiments. It is well established that chemical shift perturbation (CSP) NMR experiments either based on $^1$H,$^15$N TROSY HSQC or on methyl TROSY experiments provide exhaustive information on ligand binding sites under near-physiological conditions.\cite{11} Therefore, we used CSP NMR experiments to study bile acid binding to human GII.4 Saga P-domains uncovering a low-affinity binding site for bile acids (cf. Scheme 1). This site is not present in published crystal structures but becomes ligand-accessible during long-timescale MD trajectories. Ensemble-based docking of various bile acid molecules to a large number of conformations plus additional MD refinement of high-ranked poses reveals the plasticity of this site, yielding binding poses in very good agreement with CSP data. Finally, STD NMR experiments using VLPs of seven different human NoV strains suggest that low-affinity bile acid binding is a common feature of human NoVs.

**Results**

**CSP NMR experiments identify a bile acid binding pocket**

Binding of bile acids to the P-domain of VP1 of GII.4 Saga NoV was studied by using $^1$H,$^15$N TROSY HSQC spectra as well as methyl TROSY spectra, identifying perturbations of backbone NH signals and of side chain methyl groups, respectively. Samples were uniformly $^2$H,$^15$N-labeled ([$U-^2$H,$^15$N]) or specifically $^{13}$C-methyl (MIL ProSVProSA)-labeled. Binding of four bile acids, CA, DCA, GCDCA, and CDCA (Scheme 1) was tested with backbone $^1$H,$^15$N HSQC TROSY experiments (Figure 1 and Figure S1 in the Supporting Information). For comparison, CA-induced CSPs were also derived from a methyl TROSY experiment (see below). Both types of CSP experiments yielded two symmetrical binding sites close to the C termini of the dimeric P-domain (P dimer) and at large distance to the HBGA binding pockets as this is shown in Figure 1 C for CA as representative example.

Binding of DCA, GCDCA, and CDCA yields very similar CSPs reflecting binding to the same site (cf. Figure S1). NMR signals of amino acids in the HBGA binding pocket or at the sites matching the high-affinity bile acid binding pockets reported for rare genotypes of human NoV\cite{4a} or murine NoV\cite{4c} remain unaffected.

**STD NMR experiments demonstrate bile acid binding to P dimers and to VLPs from different norovirus strains**

Using STD NMR experiments, we tested binding of CA, GCA, GCDCA, TCA, and TCDCA to viral capsids. For these experiments we used VLP samples from different laboratories and from different NoV strains (GI.1 Norwalk, GII.4 Saga, GII.4 Ast6139, GII.7 RKI, GII.10 Vietnam026, GII.17 Kawasaki308, and GII.17 Saitama/T87). In each case STD NMR spectra indicate binding of bile acids to VLPs (Figure S2). The STD NMR spectra did not indicate any significant differences across the strains suggesting similar binding modes. Therefore, we chose CA as a representative bile acid for further STD NMR experiments.

For a sample of GII.4 Saga VLPs we recorded STD buildup curves and determined a binding epitope (Figure 2) from initial STD growth rates, complementing the topological information about the binding site from CSP NMR (Figure 1). Almost all CA protons receive saturation from the protein but due to signal overlap STD amplification factors were obtained only for a subset of protons. The corresponding binding epitope suggests that one side of the steroid skeleton makes closer contact with protons located in the binding pocket.

**NMR titration experiments provide relative affinities for CA binding**

We performed CSP and STD NMR titration experiments to study the binding of CA and GCDCA to NoV P dimers and VLPs. CA was chosen, as its solubility in water is higher than other bile acids. GCDCA is less water soluble, but was included because GCDCA has been the focus of previous investigations.\cite{4a,4c}

**Scheme 1.** Chemical structures and abbreviations of ligands used for NMR experiments.
It is well established that bile acids form micelles when dissolved in water above a critical concentration. These micelles differ from conventional micelles in that the micellar assembly consists of only few molecules (<10–20). Formation of such aggregates has been reported to begin at higher concentrations for CA than for DCA or TCA, as this is reflected by critical micelle concentrations of about 5 mM obtained from diffusion ordered NMR experiments. From chemical shift changes observed in simple 1H NMR spectra of CA and GCDCA at increasing concentrations we conclude that the formation of aggregates begins at concentrations above about 4 mM for CA, and above about 1 mM for GCDCA (cf. Figure S4). Therefore, interpretation of binding isotherms (Figure 3) requires caution. Our data may well reflect binding of free CA or GCDCA molecules and of aggregates at the same time. On the other hand, exchange between free and micelle-associated bile acid molecules should be rapid, not limiting the amount of free bile acid molecules binding to P dimers. Nevertheless, analysis of binding isotherms cannot provide true dissociation constants as saturation is only reached at ligand concentrations well above estimated critical micelle concentrations, making an assessment of the contribution of aggregates impossible. Consequently, we only report apparent dissociation constants and translate these values into relative affinities for meaningful comparisons (Table 1). Of note, all apparent dissociation constants were of the same order of magnitude in the low-milli-molar range.

Figure 1. A) Regions of a 1H,15N TROSY HSQC spectrum showing backbone NH signals of a [U-2H,15N] labeled sample of P dimers of GII.4 Saga norovirus (100 μm) being disturbed by the presence of 8 mM cholic acid (CA). The spectrum was recorded at 500 MHz and 298 K. B) Chemical shift perturbations (CSPs, calculated as Euclidean distances) of backbone NH signals as a function of amino acid position. CSPs larger than mean ± σ are shown in orange, and values larger than mean ± 2σ in red. C) Mapping of CSPs onto the crystal structure of P dimers (PDB ID: 4X06) using the color coding in panel (B). The remote HBGA binding site is highlighted with a blue ball (position of C6 of the fucose moiety of B-trisaccharide).

Figure 2. Binding epitope of CA bound to GII.4 VLPs from STD NMR buildup curves (cf. Figure S3). Almost all protons receive saturation, but due to overlap STD amplification factors (AFs) could only be determined for a subset of protons. Where STD amplification factors could be obtained respective protons are color coded. Experiments were performed at 600 MHz, with the temperature set at 277 K.
molar range. Assuming the contribution of aggregation is constant for all titrations with CA, a comparison of relative binding affinities for different strains is possible, with lower relative values indicating increased affinities. The $^{1}$H,$^{15}$N TROSY HSQC spectra of GII.4 Saga P dimers.

Table 1. Relative affinities for binding of CA to NoV P-domains and VLPs.[a]

| No. | Ligand | NoV strain | Protein | Isotopic labeling | NMR experiment | T [K] | $K_{app}$ [mM] | $R^{2}$ | Relative affinity[b] |
|-----|--------|------------|---------|-------------------|----------------|-------|----------------|-------|---------------------|
| 1   | CA     | GI.1 Norwalk | VLPs | – | STD | 277 | 12.2 ± 1.8 | 0.9933 | 1.2 |
| 2   | CA     | GII.4 Saga | P dimers | U-2H,15N | HSQC TROSY | 298 | 10.0 ± 0.6 | 0.9924 | 1.0 |
| 3   | CA     | GII.4 Saga | P dimers | MIL-ProSVProSA | methyl TROSY | 298 | 9.5 ± 1.1 | 0.9981 | 1.0 |
| 4   | CA     | GII.4 Saga N373D | P dimers | – | STD | 277 | 6.2 ± 1.1 | 0.9929 | 0.6 |
| 5   | CA     | GII.4 Saga | VLPs | – | STD | 277 | 4.7 ± 0.4 | 0.9912 | 0.5 |
| 6   | CA     | GII.4 Saga | VLPs | – | STD | 277 | 3.6 ± 0.7 | 0.9907 | 0.4 |
| 7   | CA     | GII.4 MI001 | P dimers | U-2H,15N | HSQC TROSY | 298 | 11.1 ± 0.9 | 0.9882 | 1.1 |
| 8   | CA     | GII.4 Ast6139 | VLPs | – | STD | 277 | 26.5 ± 8.0 | 0.9830 | 2.7 |
| 9   | CA     | GII.7 RKI | VLPs | – | STD | 277 | 10.9 ± 4.3 | 0.9775 | 1.1 |
| 10  | CA     | GII.10 Vietnam026 | VLPs | – | STD | 277 | 12.9 ± 2.9 | 0.9881 | 1.3 |
| 11  | CA     | GII.17 Kawasaki308 | P dimers | MIL-ProSVProSA | Methyl TROSY | 298 | 20.1 ± 2.9 | 0.9895 | 2.0 |
| 12  | CA     | GII.17 Kawasaki308 | VLPs | – | STD | 277 | 31.5 ± 2.8 | 0.9990 | 3.1 |
| 13  | CA     | GII.17 Saltama/T87 | VLPs | – | STD | 277 | 6.4 ± 0.5 | 0.9888 | 0.6 |
| 14  | GCDCA | GII.4 Saga | P dimers | U-2H,15N | HSQC TROSY | 298 | 1.5 ± 0.3 | – | – |
| 15  | GR     | GII.4 Saga | P dimers | U-2H,15N | HSQC TROSY | 298 | 13.8 ± 0.4 | 0.9977 | – |

[a] Relative affinities for GCDCA and GR cannot be compared because the contribution of ligand aggregates to binding is different (cf. main text). Samples used for STD experiments and for $^{1}$H,$^{15}$N TROSY HSQC experiments contained 10% D$_{2}$O and the solution was adjusted to pH 7.3. Samples for methyl TROSY experiments contained > 99% D$_{2}$O and the solution was adjusted to pH 7.4 to create comparable conditions. [b] Apparent dissociation constants were calculated from STD initial growth rates (STD-AF$_{0}$). [c] Arbitrary units.
based titration of GII.4 Saga P dimers with CA was used as a reference with the relative affinity set to 1.0 (Table 1, No. 2).

For binding of CA to GII.4 Saga P dimers we performed three different and independent types of NMR titration experiments. One set of titrations used CSPs from $^{1}H_{3}^{15}N$ TROSY HSQC spectra, a second set of titrations was based on methyl TROSY spectra, and a third data set was based on STD NMR spectra. In all three cases we obtained similar relative affinities (Table 1, No. 2–4; Figure 3). For the STD NMR titration we used the N373D mutant, which cannot undergo deamidation. The experiment was performed at a lower temperature (277 K) than the CSP titrations (298 K), which may account for the slightly lower relative affinity (Table 1, No. 4).

To compare P dimers to viral capsids, we also obtained STD NMR titration curves for CA binding to GII.4 Saga VLPs (Figure 3D; Table 1, No. 5 and 6). In one set of experiments we performed titrations at single saturation times of 2 s$^{15}$ (cf. Figure S5 and Table S1). Another titration experiment made use of initial STD growth rates$^{16}$ determined from STD build up curves (Figure S6 and Table S2). The initial growth rates were also used to derive a binding epitope for CA (Figures 2 and S3). The apparent $K_{D}$ values obtained are similar, with the value resulting from initial growth rates being slightly smaller, as expected.

We also studied binding of CA to $[U-^{2}H,^{15}N]$-labeled P dimers of a related GII.4 strain (MI001) that infects humans as well as mice.$^{17}$ As we have no backbone assignment for this strain yet, we compared backbone chemical shifts of MI001 P dimers to Saga P dimers, yielding some tentative assignments (Figure S7). Based on these preliminary data, comparison of CSPs measured in $^{1}H_{3}^{15}N$ TROSY HSQC spectra suggests that the binding pockets of MI001 and Saga are rather similar, and titration curves (Figure 3) yield a very similar relative affinity compared to Saga (Table 1, No. 2 and 7).

A sample of MI1 ProSVProSA labeled P dimers of a GII.17 Kawasaki308 strain was also subjected to a methyl TROSY based CSP titration showing specific effects of binding. In this case, assignments are not available yet. Nonetheless, a relative affinity can be obtained from determining CSPs of unassigned cross-peaks (Table 1, No. 11, Figure S8).

STD NMR titrations were performed for VLPs of the NoV strains GII.4 Ast6139, GII.7 RKI, and GII.10 Vietnam026, GII.17 Kawasaki308 and GII.17 Saitama/T87 as well as for VLPs of the GI.1 Norwalk virus, using a single saturation time of 2 s (Table 1, No. 1, No. 8–10, and No. 12–13). Relative affinities for GI.1 Norwalk, GII.10 Vietnam026, GII.7 RKI, GII.17 Saitama/T87 and GII.4 Saga VLPs are very similar. For GII.4 Ast6139 and for GII.17 Kawasaki308 VLPs as well as for Kawasaki308 P dimers slightly larger relative affinities are found.

To study binding of GCDCA to GII.4 Saga P dimers we finally determined an apparent dissociation constant from a CSP titration based on $^{1}H_{3}^{15}N$ TROSY HSQC spectra (Table 1, No. 14). However, the apparent dissociation constant derived has to be treated with great caution, as only the Val508 NH signal could be used for calculating a dissociation constant (Figure S9). All other CSPs were too low to justify global fitting. Moreover, as discussed above, this value cannot be directly compared with the CA titrations, and, therefore, no relative affinity is reported in Table 1.

Deamidation of Asn373 does not affect binding of CA

For GII.4 Saga P dimers it has been shown that spontaneous deamidation of Asn373 and subsequent formation of an iso-aspartate residue at this position abrogates HBGA binding.$^{10}$ This process is likely to be relevant for about 66% of all GII.4 strains. Therefore, we tested the influence of deamidation on CA binding to GII.4 Saga P dimers. Native Saga P dimers (NN P dimers) and completely deamidated Saga P dimers (dID P dimers) were purified using an ion-exchange chromatography protocol and immediately subjected to CSP NMR experiments, keeping conversion of NN P dimers at a minimum. Comparison of CSPs in corresponding methyl TROSY spectra upon titration with CA demonstrates that deamidation has practically no influence on binding. Almost identical apparent $K_{D}$ values were obtained (Figure 4).

We also applied STD NMR titrations to compare binding of CA to the N373D mutant of GII.4 Saga P dimers. This mutant does not undergo deamidation and does not convert into the iso-aspartate form. Within experimental error, the apparent $K_{D}$ values are identical to those obtained for wild-type P dimers (Table 1, No. 4 and Figure S10).

HBGA binding revisited

Using CSP titrations we have shown that the affinity of L-fucose, which constitutes the minimal recognition element of HBGAs, has been significantly overestimated in preceding studies. We also showed that the same holds true for blood group B-trisaccharide (B-Tri) by reevaluating data from a competitive STD NMR experiment.$^{19}$ Therefore, we determined the dissociation constant $K_{D}$ for B-Tri binding to Saga P dimers using a CSP titration based on methyl TROSY experiments before addressing the question whether there is mutual cross talk between HBGA-binding and bile acid binding. Using a freshly purified sample of NN P dimers we obtained a $K_{D}$ value of 5.6 mM (cf. Figure S5), in excellent agreement with the value of 5.5 mM from a competition STD NMR experiment.$^{10,19}$

Bile acid binding does not affect HBGA binding and vice versa

To answer the question whether bile acid binding and HBGA binding have a mutual impact, we performed methyl TROSY-based CSP NMR titrations with CA in the presence of saturating amounts of B-Tri and with B-Tri in the presence of near-saturating amounts of CA. The results are summarized in Figure 5 and show that binding affinities of CA and B-Tri are unaffected by the presence of B-Tri or CA, respectively.

Glycyrrhizic acid binds to an adjacent site

To shine some more light on the specificity of the bile acid P-domain interaction we chose glycyrrhizic acid (GR) as a test
compound. GR belongs to the class of saponins, which are amphipathic glycosides, containing a triterpene ring system with structural similarities to the steroid backbone of bile acids. From $^1$H,$^1$N TROSY HSQC based CSPs it is clear that GR binds to GII.4 Saga P dimers at a site adjacent to the bile acid site, as this can be seen from Figure S11. Although a comparison with CA binding is impossible, it can be stated that the affinity of GR is of the same order of magnitude as found for CA (Table 1, No. 15).

Microsecond MD combined with docking reveals a dynamic cavity in the binding region

Crystal structures of GII.4 Saga P dimers (PDB IDs: 4X06 and 4OOX) exhibit no accessible binding pocket of sufficient volume to accommodate bile acid molecules. Therefore, the P dimer was subjected to a long all-atom MD simulation (1 µs), revealing significant dynamics of the backbone as reflected by the root-mean-square deviation (RMSD) of backbone atoms in Figure 6A and the different protein conformations shown in Figure S12. The RMSD relative to the crystal structure fluctuates around 0.10 nm in the first 200 ns simulation time, then increases to 0.20 nm during the next 600 ns, and finally converges to 0.25 nm during the last third of the simulation.

The root-mean-square fluctuations (RMSF) of the backbone atoms are used to locate regions of high flexibility. As shown in Figure 6B the RMSF is especially large at both termini (0.6 nm) and within flexible loop regions (0.3 nm), which can be expected. However, in the crystal structure the C terminus occupies the experimentally identified binding site, preventing bile acid binding. During the MD simulation the binding site volume increases significantly relative to the crystal structure (Figures 6A and S13). Volume fluctuations between 20 and

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**Figure 4.** A) Regions of methyl TROSY spectra of a $^{13}$C-methyl (MIL ProSVProSA)-labeled sample of GII.4 Saga NN and iDiD P dimers. Two representative cross-peaks demonstrate that perturbations upon addition of CA are unaffected by deamidation of Asn373. B) CSPs as a function of amino acid position. CSPs at 8 mM CA larger than mean $\pm$ s (red: NN P dimers; blue: iDiD P dimers). C) $^{13}$C methyl CSPs mapped onto the surface of GII.4 Saga NN P dimers (PDB ID: 4X06). The deamidation site Asn373 is highlighted. D) Binding isotherms from chemical shift titrations of NN P dimers with CA. E) Binding isotherms from chemical shift titrations of iDiD P dimers with CA.
130 nm³ during the first 400 ns of the simulation can be assigned to small conformational changes of mainly amino acid side chains. After 400 ns, fluctuations increase in amplitude and several distinct changes can be detected due to larger structural, that is, backbone rearrangements, especially concerning the C-terminal residues 527–530.

Ensemble docking identifies initial binding modes

The high conformational flexibility of the binding site and the associated fluctuations of the pocket volume do not allow unambiguous identification of accurate bile acid binding poses. Therefore, a large ensemble of \( N = 2000 \) protein conformations was generated by systematically extracting protein conformations at intervals of 0.5 ns. These 2000 receptor conformations were subsequently used for docking of CA, DCA, CDCA and GCDCA. The docking scores (arithmetic mean of the docking scores of CA, DCA, CDCA, and GCDCA to GII.4) for the 2000 docked conformations are shown in Figure 6A (cf. lowest panel) together with corresponding RMSD values and pocket volume differences \( \Delta V \). The five lowest scores (highest affinities) are found only after long simulation times of 747.0, 464.5, 817.5, 460.0, and 718.5 ns, with average docking scores of \( /C_0 \), \( /C_0 \), \( /C_0 \), \( /C_0 \), and \( /C_0 \) kcal mol\(^{-1}\), respectively. For these five protein conformations, all of the four bile acid species exhibit lowest docking scores, with GCDCA systematically showing the lowest ones (Figure S14 and Table S3).

It has to be emphasized again, that the best scores were achieved only after 400 ns of MD simulation, correlating with an increase of backbone RMSD values from 0.1 to 0.2 nm. However, the best scoring protein-ligand complexes neither exhibit a distinctly high protein RMSD nor a particularly large pocket volume. Therefore, the use of such a large ensemble of protein conformations is superior to the selection of a small set of conformations via RMSD values or pocket-shape based clustering as suggested in earlier studies.^[20] Otherwise, impor-
tant protein conformations critical for ligand binding may remain unresolved.

In the five best-scoring protein–ligand poses the C-terminal residues are more solvent exposed as compared with the crystal structure, forming a novel cavity of sufficient size and allowing interactions of the C terminus with the carboxylate groups of the bile acids, as exemplarily shown for CA in Figure 7 (for the other bile acids, see Figure S15). The bile acid orientation is quite similar for all poses with the carboxylate being close to Leu527. In pose 1, the hydroxy groups are facing His505, Leu506 and Val508. In Pose 2, the hydrophilic site is oriented away from the protein, pointing to the solvent. Poses 3–5 are almost identical and feature interactions of the hydroxy groups with protein residues Leu486 and Phe487.

Equilibrium MD of protein–ligand complexes reflects weak binding with multiple binding modes

As the poses from docking only approximate true binding modes, we used them as initial configurations for subsequent MD refinement simulations. For each of the five top-ranked poses of each bile acid molecule, ten independent MD runs of 20 ns each with different initial velocity distributions were performed. The average ligand RMSDs of the 50 MD runs were used as a criterion for the receptor–ligand complex stability (for the full ligand RMSDs, see Figure S16). Here, RMSD > 1.0 nm indicates dissociation of the complex, a value between 0.3 and 1.0 nm corresponds to a stable complex with a ligand orientation rearrangement relative to the docked structure, and a RMSD < 0.3 nm belongs to a stable binding pose with minor refinement during the MD simulation. Depending on the initial coordinates and velocity distributions, the ligand RMSDs range from 0.2 to 7.1 nm as shown in Figure 8A. Taking into account the low affinity of CA and the nonideal, approxi-
ing site, leading to a dynamic binding equilibrium with multiple, transient binding modes. This is experimentally reflected by the broad spatial distribution of residues with significant CSP and the low affinity in the millimolar range of CA binding.

Discussion

Protein-based CSP NMR experiments combined with ligand-based NMR experiments and long-timescale all-atom MD simulations have provided a detailed picture of low-affinity binding.
of bile acids to NoV capsid protein. For the translation of experimental CSPs into docking models\textsuperscript{21} it was crucial to engage long MD simulations because the low-affinity bile acid binding site cannot be represented by a single conformation. For instance, CA–protein contacts for the red filled circle trajectories are shown in (B) and (D) in detail. B) Contact occupancies of CA with backbone nitrogen atoms during the last 10 ns of trajectories Pose 3 Rep 5 (B) and Pose 4 Rep 3 (D). Contact criterion is a distance $<0.6$ nm between the backbone N and at least one heavy atom of CA. Contact amino acids that exhibit significant CSPs are highlighted in red (backbone HSQC) and gold (methyl TROSY), respectively. Proline residues are not considered as they show no NMR signals. Only amino acids with an occupancy $>0.02$ are shown. C,E) Representative snapshots of stable protein–CA complexes for Pose 3 Rep 5 (C) and Pose 4 Rep 3 (E).
the true binding mode does not necessarily correlate with pocket volume and shape or clustered conformations. Rather, the conformational ensemble should be sufficiently large and not limited to few cluster representatives, and the docking algorithm should take into account side chain flexibility, as this is nicely supported by our results.

Our results highlight that good docking scores do not necessarily lead to stable protein–ligand complexes in MD trajectories during further refinement as this has been discussed by others recently. Depending on ligand affinity, many replicates may be necessary to obtain stable complexes with structural integrity and convergence, even after very careful equilibration. Finally, a converged complex with a low RMSD does not necessarily represent an accurate binding mode but can also appear due to improbable or overly strong interactions in the initial configuration. Our study shows once more that careful inspection and rationalization of the computational predictions by comparison with experimental restraints is vital for generating a realistic binding model. Following these arguments, our approach revealed a binding pocket of sufficient volume to accommodate different bile acids, thus suggesting dynamic binding with multiple transient modes that all share backbone contacts with Leu486, Phe487, Leu507, Val508 and Ile509, as well as ligand orientations with the bile acid methyl groups being buried.

Our studies suggest that low-affinity binding of bile acids is a common feature of NoV capsid proteins independent of genogroup or genotype (Table 1). Dissociation constants are in the low millimolar range and are thus similar to affinities for HBGAs. For GII.4 Saga, an almost complete assignment of backbone NH signals of the P-domain of the VP1 capsid protein is available, allowing location of the bile acid binding site from specific CSPs. We hypothesize that this binding site is also present in the other NoV strains where we have observed low-affinity bile acid binding. This hypothesis is supported by the fact that for GII.4 MI001, where we have some tentative backbone NH assignments available, the same region seems to be affected. We are currently working on the backbone and 13C-methyl assignments of other NoV strains to further substantiate this hypothesis.

Importantly, for GII.4 Saga P dimers no CSPs have been observed at sites corresponding to two other bile acid binding sites described before. As replication of GII.4 NoVs in human intestinal enteroids has been reported to be significantly enhanced by the presence of bile acids, and because we can exclude binding to one of the putative high-affinity sites, we suggest that the low-affinity interaction with viral capsids provides the molecular basis for understanding the role of bile acids in promoting infection.

To test the possibility that there is mutual cross-talk between HBGA and bile acid binding we performed NMR binding experiments in the absence and presence of the respective other ligand (Figure 5). Our experiments show that binding of bile acids to the low-affinity site and binding of HBGAs to the HBGA site are independent events. Likewise, we have shown that the influence of spontaneous deamidation in the HBGA binding site on the binding affinity for bile acids is negligible. These findings suggest that there is no or very minor cross-talk between these binding events.

For GII.10 Vietnam026 we expected affinities for bile acids in the micromolar range due to binding to a high-affinity bile acid binding site adjacent to the HBGA site as described recently. In this case we have no isotope-labeled P dimers available yet, but we have performed STD NMR experiments with GII.10 Vietnam026 VLPs. Unexpectedly, binding isotherms from STD NMR titrations only reflect low-affinity binding, at least showing that the low-affinity binding site is present. The failure of observing STD effects resulting from binding to the high-affinity site is likely due to low rates of dissociation of bile acids from that site, leading to weak or no saturation transfer. Therefore, we conclude that this high-affinity site is invisible to STD NMR experiments. We are currently working on labeling GII.10 Vietnam026 P-domains with stable isotopes to further explore this binding site with CSP NMR experiments.

In this respect it is of note that the low-affinity bile acid site is located close to the C-terminus of NoV P-domains. In our studies we have used C-terminally truncated P-domains of the VP1 capsid protein because the presence of the highly conserved so-called arginine tail leads to aggregation and significantly impedes NMR experiments. On the other hand, all VLP preparations contain the arginine tail. Therefore, if the arginine tail played a role in bile acid binding this should be reflected by different affinities of bile acids for P dimers versus VLPs. From CSP NMR experiments the relative binding affinities of CA binding to Saga P dimers or to Saga VLPs are only about a factor of two apart (Table 1). Such small differences might well be due to slightly different experimental setups and likely do not indicate involvement of the arginine tail in binding to bile acids. In fact, the temperatures for the two types of experiments, CSP versus STD NMR, were different and may explain the observed small alterations. On the other hand, as discussed above for the high-affinity site present in GII.10 Vietnam026 P-domains one can speculate that the arginine tail induces conformations that bind to bile acids with higher affinity. This would be invisible to STD NMR and would require CSP NMR experiments using either VLPs or P-domains including the arginine tail. Neither stable isotope labeling of VLPs nor preparing non-aggregating samples of P-domains including the arginine tail is a trivial task. We are currently working on these problems in our laboratory.

At present, we have no complete backbone NH assignments for P dimers of NoV strains other than GII.4 Saga. Therefore, we cannot yet directly identify bile acid binding pockets of these strains from CSP NMR experiments. However, the observation of similar binding affinities of various NoV strains (Table 1) indirectly suggests similar binding modes and sites in all cases. A structure based sequence alignment of NoV VP1 domains available from our previous study shows that within a given genotype most amino acid positions are highly conserved. Therefore, it is not surprising that some of the amino acids identified to be affected by bile acid binding are also highly conserved. For instance, the stretch of amino acids from R484 to K490 is almost identical for all GII.4 sequences (> 98% sequence identity). In general, the complete C terminus is highly
conserved with an average sequence identity of >95%. Therefore, a comparison among different genogroups and genotypes is more informative. Evaluation of the VP1 C-terminal sequences of the NoV strains studied in this work demonstrates that mutations of amino acids affected by bile acid binding are possible without impeding binding (cf. Figure S18). This observation matches the results from MD simulations, showing considerable plasticity of the low-affinity bile acid binding site and thus allowing different poses of bile acids (Figure 8C, E). It appears that this low-affinity binding site is rather promiscuous, able to accommodate various bile acids with similar affinities. It will be interesting to further scrutinize the consequences of our findings in cell culture systems and animal models.

Conclusion

This study highlights the complex role of bile acids in norovirus infection from a structural point of view and underlines the potential of protein- and ligand-based NMR binding experiments in combination with long-timescale MD simulations to portray low-affinity binding events. A low-affinity bile acid binding site appears to be a common feature of a variety of human NoV strains. As affinities in the millimolar range match portrait low-affinity binding events. A low-affinity bile acid binding should be saturated when the virus is residing in the intestine. It would be very interesting to find NoV strains lacking the ability of binding bile acids at the C terminus and to study the consequences for infection. We speculate that “soft recognition” of bile acids affects the stability of viral capsids and in turn modulates infection. In the light of the development of novel and better accessible NoV cell culture systems, this hypothesis seems to be verifiable in the near future.

Experimental Section

Protein biosynthesis and purification: Non-deamidated [U-2H,15N] Saga 2006 (GenBank accession number BAG70518.1) and GII.4 Saga 2006 (GenBank accession number AGQ57036.1) were gifts from Prof. Stefan Taube (University of Lübeck, Germany). GII.4 Ast6139 VLPs (GenBank accession number CAE47529.1) were donated by Prof. Francisco Parra (University of Oviedo, Spain).

Bile acids and other ligands: Cholic acid (CA), glycochenodeoxycholic acid (GCDCA), sodium taurochenodeoxycholate (TDCDA), deoxycholic acid (DCA), chenodeoxycholic acid (CDCA), taurocholic acid (TCA), sodium glycodeoxycholate hydrate (GCA) and glycuronic acid (GR) were purchased from Sigma–Aldrich. Blood group B-trisaccharide (α-L-Fuc-(1,2)-[β-D-Gal-(1,3)-]β-D-Gal-(1,4)-N3) was a gift from Dr. Hanne Peters in our laboratory and had been obtained via enzymatic synthesis from (α-L-Fuc-(1,2)-[β-D-Gal-(1,4)-N3], which was a gift from Prof. Javier Pérez-Castells (CEU San Pablo, Madrid, Spain).

NMR spectroscopy: STD NMR experiments were acquired at 277 K on a Bruker Avance III HD 600 MHz NMR spectrometer equipped with a TXI room temperature probe. All other NMR experiments were recorded on a Bruker AV III 500 MHz NMR spectrometer equipped with a TCI cryogenic probe at 298 K if not specified otherwise. Data were processed with TopSpin 3.6, and peak positions were extracted using CCP NMR 2.4.2.21 A backbone assignment is available for Saga GII.4 P dimers and is deposited with the BioMagResBank with the accession code 27445.

Backbone and side-chain chemical shift perturbation experiments:22 H,15N TROSY HSQC spectra were acquired with 8–24 scans with 2048 data points in t2 and 430 increments in the indirect dimension t1. The acquisition time was 128 ms in t2 and 121 ms in t1. The relaxation delay was set at 1.5 s. The NMR samples contained 100–120 μM [U-2H,15N] Saga 2006 or M001 P dimers, 200 mM D2O, 300 μM imidazole, 100 mM NaCl, 75 mM sodium phosphate buffer (pH 7.3), 0.02% NaN3, and 10% D2O. Ca has been titrated up to final concentrations of 15 mM (Saga 2006) and 18 mM (M001). GR has been titrated up to 25 mM, GCDCA up to 15 mM. Other bile acids have only been added with a single concentration of 2 mM. Bile acid titration stocks were prepared with final concentrations up to 300 mM in 75 mM sodium phosphate buffer, 100 mM NaCl (pH 7.3). The pH of the titration stocks was increased by titration of NaOH until all solid dissolved and subsequently re-adjusted to pH 7.3. Imidazole signals were used as internal standard to monitor the pH during titrations as described previously (see supplementary data section 2.3 of ref. [10]).

Methyl TROSY spectra22,23 were acquired with 4–48 scans with 1024 data points in t2 and 512 increments in the indirect dimension t1. The acquisition time was 137 ms in t2 and 120 ms in t1. The relaxation delay was set at 1.5 s. The NMR samples contained 16.5–31 μM of MIL ProSVProSA-labeled GII.4 Saga or GII.17 Kawasakii308 P dimers in 75 mM sodium phosphate buffer (pH 7.4) in D2O (>99%). The deamidation status of individual P dimer samples depended on the storage time and is described in the main text. Samples contained 100 mM NaCl, 100 μM D2O, 100 μM imidazole and 0.02% NaN3. Samples were titrated to a final concentration of CA of 14 or 20 mM. Bile acid titrations were performed as described above, with the only difference that NaOD in D2O was used to prepare bile acid titration stocks with a final pH 7.4. Saturation transfer difference (STD) NMR: A train of 50 ms Gaussian-shaped radio frequency pulses separated by 1 ms for a total duration of 2 s were used for protein irradiation.21 For extracting Kd values from STD initial growth rates, 0.25, 0.5, 0.75, 1, and 2 s saturation times were used. In all cases, an attenuation of 40 dB...
was chosen resulting in a 680.5° flip angle. The water signal was suppressed using excitation sculpting,[32] and for protein signals of P dimers were attenuated applying a 20 ms spinlock filter before acquisition. The acquisition time was set at 1.96 s with an additional relaxation delay of 5–20 s. On and off resonances were set at −4 ppm and 200 ppm, respectively.[33] The number of scans ranged from 200 to 2400.

Samples containing VLPs were prepared at 0.41 to 1 mg mL⁻¹ VP1 concentration (6.9 to 17.7 μm binding sites) in PBS pH 7.3, 100 μm [D]3DSS, 0.02% NaN₃ and 10% D₂O. Samples were titrated with cholonic acid to a final concentration of 15–20 μm, with GCA and TDCA (1 mM each) or with TCA and GCDCA (1 mM each). Bile acid stock solution were prepared as described above for backbone chemical shift perturbation experiments. For N373D Saga P dimers, one sample containing 45 μm P dimers, 100 μm [D]3DSS in 20 mM fully deuterated sodium phosphate buffer (pH 7.45) was prepared and titrated with CA up to a final concentration of 12 μm.

Assessing critical micelle concentrations: Critical micelle concentrations of CA and GCDCA have been estimated from chemical shift perturbations of the C19 methyl group in a series of 1D ¹H NMR spectra of the bile acids at different concentrations in 75 mM sodium phosphate buffer, 100 mM NaCl (pH 7.3) with 10% D₂O. Determination of (apparent) dissociation constants K₀: Dissociation constants K₀ were calculated either from CSPs, STD-AF or STD-AF₀ by using Equation (1):

\[
O = \frac{K₀ + |P| + |L|}{2|P||L|} \quad O_{\text{max}}
\]

in which O is the experimentally observed CSP, STD-AF, or STD-AF₀ at each ligand concentration, |P| and |L| are the total protein and ligand concentrations, respectively, and O_{\text{max}} is the observable at saturation with ligand.

For the experiments based on backbone and side-chain chemical shift perturbations, CSPs were calculated as Euclidean distances ΔP_{\text{sat}} in Hz according to Equation (2):[34]

\[
\Delta P_{\text{sat}}(Hz) = \sqrt{\Delta v^{2} + \Delta v_{D}^{2}}
\]

for which X is either ¹¹N or ¹³C, Δvₓ, and Δvₓ, the CSPs of ¹H and ¹³N or ¹⁵N resonances, respectively, at a given ligand concentration. Global nonlinear least-squares fitting of Eq. (1) to CSPs (mean ± 2σ for ¹H NMR TROSY HSQC and mean ± σ for methyl TROSY experiments) furnished dissociation constants K₀.

To derive K₀ values from STD NMR titrations, STD amplification factors (AF) were calculated by using Equation (3):[31]

\[
\text{STD-AF} = \frac{l_0 - l_{\text{sat}}}{l_0} \times \text{ligand excess}
\]

where l₀ and l_{sat} are the signal intensities in the off- and on-resonance spectra, respectively. Ligand excess refers to the ratio of ligand concentration over protein concentration. STD-AF values were plotted against cholic acid concentration. Fitting Equation (1) to the data delivered dissociation constants K₀.

Calculation of K₀ values from STD initial growth rates,[36] STD-AF values were measured as a function of the saturation time t_{sat} and fitted to Equation (4):

\[
\text{STD-AF}(t_{\text{sat}}) = \text{STD-AF}_{\text{max}}[1 - \exp(-k_{0}t_{\text{sat}})]
\]
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Conflict of Interest

The authors declare no conflict of interest.

Keywords: chemical shift perturbation · ensemble docking · long-timescale MD · molecular recognition · STD NMR spectroscopy

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Contact analysis was performed with MDTraj. [51] For each frame, for each amino acid, a contact was counted if at least one heavy atom of CA was in proximity of 0.6 nm or less to its backbone nitrogen. The contact occupancy of an amino acid is the number of counted contacts divided by the number of frames. Molecular images were rendered with VMD ver. 1.9.3. [54]
