Solution Structure of LC4 Transmembrane Segment of CCR5

Kazuhide Miyamoto1*, Kayo Togiya2

1 Department of Pharmaceutical Health Care, Faculty of Pharmaceutical Sciences, Himeji Dokkyo University, Himeji, Hyogo, Japan, 2 15-203 Hirano-machi, Himeji, Hyogo, Japan

Abstract

CC-chemokine receptor 5 (CCR5) is a specific co-receptor allowing the entry of human immunodeficiency virus type 1 (HIV-1). The LC4 region in CCR5 is required for HIV-1 entry into the cells. In this study, the solution structure of LC4 in SDS micelles was elucidated by using standard 1H two-dimensional NMR spectroscopy, circular dichroism, and fluorescence quenching. The LC4 structure adopts two helical structures, whereas the C-terminal part remains unstructured. The positions in which LC4 binds to the HIV-1 inhibitory peptide LC5 were determined by docking calculations in addition to NMR data. The poses showed the importance of the hydrophobic interface of the assembled structures. The solution structure of LC4 elucidated in the present work provides a structural basis for further studies on the HIV-1 inhibitory function of the LC4 region.

Introduction

CC-chemokine receptor 5 (CCR5) is a member of the G-protein-coupled receptor superfamily and is comprised of seven transmembrane segments [1]. The N-terminus of the extracellular region is associated with binding of the chemokines, while the C-terminus of the intracellular region serves as a binding site for β-arrestin [2]. Furthermore, CCR5 and CXCR4 are specific co-receptors for human immunodeficiency virus type 1 (HIV-1) entry, which is the causative agent for AIDS [3,4]. To gain entry into cells, HIV-1 requires a CD4 receptor and co-receptors such as CCR5 and CXCR4 [5,6]. Blocking HIV-1 entry into the cell naturally leads to the inhibition of its infection and replication [7].

Recently, a novel synthetic LC5 peptide (157VFASLPGIIFTRSQKEGL174) corresponding to the LC4 region (15-203) that inhibits HIV-1 infection of MT-4 cells was reported [8]. It is suggested that the LC5 peptide interacts with the LC4 region (157VFASLPGIIFTRSQKEGL174) corresponding to the fourth transmembrane segment of CCR5 [5]. Gly163 in the LC4 region plays an important role in the formation of the complex between the gp120 envelope glycoprotein of HIV-1 and sCD4, and its mutation results in reduced susceptibility to HIV-1 infection. The three-dimensional structure of the LC5 peptide was determined by using NMR methods in our previous study [10]. LC4 is an attractive target for the inhibition of HIV-1 infection. The three-dimensional structure of the LC5 peptide is crucial for understanding the functional mechanism of the LC4 region.

The micellar environment of sodium dodecyl sulfate (SDS) micelles has been utilized to establish a reasonable model for the conformation of KcsA potassium channels in the membranes [11]. SDS micelles produce a membrane-mimetic environment allowing the structural study of the peptide LC5 [10] and Slc11a1 [12] in the membrane. Moreover, the membrane-mimetic environment of SDS micelles facilitates the characterization of the structural conformation of the transmembrane segments of membrane proteins [13].

Thus, in this study, we investigated the solution conformation of the LC4 region in the membrane-mimicking environment of SDS micelles using 1H-NMR spectroscopy, circular dichroism, and fluorescence quenching. In addition, the possible binding sites between the LC4 region and the LC5 peptide, which inhibits HIV-1 infection, were determined using docking calculations. This is the first conformational study of LC4 in the micellar environment.

Materials and Methods

Peptide synthesis

The LC4 peptide (157VFASLPGIIFTRSQKEGL174) corresponding to the LC4 region was synthesized with N-acetylated and C-amidated termini. Chemicals for peptide assembly, including amide resin, were obtained as SynProPep products from Shimadzu Corp. (Kyoto, Japan). After cleavage with trifluoroacetic acid, the peptide was purified on a reversed-phase HPLC using a Shim-pack C18 column (Shimadzu Corp.). The peptide purity was greater than 98%, and its molecular mass was assessed by MALDI-TOF MS on a Shimadzu AXIMA-TOF.

* E-mail: miyamoto@himeji-du.ac.jp
Circular dichroism (CD) spectroscopy

Far-UV CD spectra were recorded on a JASCO J-810 spectropolarimeter after calibration using d-camphor-10-sulfonate. The sample was dissolved in a buffer solution (80 mM phosphate buffer) or SDS solution (80 mM phosphate buffer, 10 mM SDS). A 0.1 mm path length quartz cuvette was used for 50 μM sample solution. Spectra at room temperature were acquired from 190 to 250 nm with a scanning speed of 30 nm/min, a response time of 4.0 s, and a bandwidth of 1 nm. Each CD spectrum was the average of four scans. After subtraction of the solvent spectrum, the CD data were obtained by converting from CD signal into mean residue molar ellipticities.

Fluorescence

The fluorescence emission spectra were examined using a RF5300PC spectrophotometer (Shimadzu Corp.). The sample was dissolved in a buffer solution or SDS solution, as described for the CD measurements. A 3×3 mm quartz cuvette was used for 50 μM sample solution. Fluorescence quenching of phenylalanine (Phe) was performed with 5–85 mM acrylamide of an extrinsic sample solution. Fluorescence quenching data were used to calculate the Stern–Volmer constants (Ksv) as follows [14,15]:

\[ \frac{F_0}{F} = 1 + K_{sv}Q \]

Where, \( F_0 \) and \( F \) are the fluorescence intensities in the absence and presence of the quencher, respectively, and \( [Q] \) is the quencher concentration.

NMR spectroscopy

For the NMR sample, the LC4 peptide (2 mM) was dissolved in \( ^{1}H_2O/^{2}H_2O \) (9:1) 80 mM phosphate buffer (pH 4.5) containing 200 mM sodium dodecyl-\( \beta \)-sulfate (d\( \beta \)-SDS) (Sigma-Aldrich). NMR measurements were made at 20°C on a Bruker AVANCE 500 MHz. Two-dimensional NOESY spectra were recorded with 80, 100, 120, 150, and 200 ms mixing times [16]. Different mixing times were used to evaluate the linearity of the NOE build-up, as described previously [10]. Phe was excited at 257 nm and its fluorescence emission at room temperature was acquired from 270 to 450 nm. After subtraction of the solvent spectrum, the fluorescence quenching data were used to calculate the Stern–Volmer constants (Ksv) as follows [14,15]:

\[ \frac{F_0}{F} = 1 + K_{sv}Q \]

Where, \( F_0 \) and \( F \) are the fluorescence intensities in the absence and presence of the quencher, respectively, and \( [Q] \) is the quencher concentration.

Structure calculation

A peak list for the NOESY spectrum with a mixing time of 150 ms was generated by the peak picking and integration functions of NMRView [22]. Automated NOE cross-peak assignments and structure calculations with torsion angle dynamics were performed using the software package CYANA 2.1 [23]. Structure calculations were started from 100 randomized conformers, and the standard CYANA-simulated annealing protocol was used with 15,000 torsion angle dynamics steps per conformer. The 20 conformers with the lowest final CYANA target function values were validated using the program PROCHECK-NMR [24]. The Connolly surfaces of the structures were simulated using the program Discovery Studio 2.1 (Accelrys Software Inc.). The program MOLMOL [25] was used to analyze the resulting 20 conformers.

The atomic coordinates (code 2RRS) have been deposited in the Protein Data Bank, Research Collaboratory for Structural Bioinformatics.

ZDOCK calculations

The docking calculation was performed using ZDOCK 3.0.1 [26]. ZDOCK is an initial-stage rigid-body molecular docking algorithm that uses a fast Fourier transform (FFT) algorithm to improve performance for searching in translational space. The precision of ZDOCK in predicting the docking structure has been proved in the Critical Assessment of Prediction of Interactions (CAPRI) Challenge [26]. All of the available structures from NMR were used to calculate the docking poses and the structures obtained were subjected to energy minimization using the Smart Minimizer algorithm (Max steps 200, RMS gradient 0.01) in the program Discovery Studio 2.1. The resulting top 5 conformers with the lowest energy were used as appropriate candidates.

Results

Secondary structure of LC4

To gain information on the secondary structure of LC4, CD spectra of LC4 were recorded at pH 4.5 and 7.4 in an aqueous environment and in SDS micelles. As shown in Figure 1, the CD spectra of LC4 were little affected by the variation in pH in both the aqueous environment and SDS micelles. The CD spectra in phosphate buffers showed a single broad negative ellipticity centered at approximately 200 nm, indicative of a random-coil conformation. In contrast, in SDS micelles, a helical conformation was induced with spectral minima occurring at approximately 225 nm (n-π* transitions) and 207 nm (π-π* transitions) along with a strong positive ellipticity at approximately 195 nm characteristic of a typical helical structure [27].
Insertion of LC4 into SDS micelles

To characterize the degree of insertion of LC4 into SDS micelles, its solvent accessibility was evaluated by the fluorescence quenching of Phe with acrylamide. Stern–Volmer plots (Figure 2A) provided information on its solvent accessibility. The $K_{sv}$ values in phosphate buffers at pH 4.5 and pH 7.4 are approximately $58.3 \text{ M}^{-1}$ and $67.1 \text{ M}^{-1}$, respectively, while in the presence of SDS micelles, the $K_{sv}$ values are approximately $14.5 \text{ M}^{-1}$ and $15.6 \text{ M}^{-1}$, respectively. The smaller $K_{sv}$ values in SDS micelles indicate that the side chain of Phe is buried in the SDS micelles (Figure 2B). The aromatic rings of the residual Phe of LC4 are embedded into SDS micelles, irrespective of the pH of the solution.

Resonance assignments

Two-dimensional $^1$H-NMR spectra of LC4 were obtained in SDS micelles at pH 4.5, because the exchange rates of amide protons at pH 4.5 are lower than those at pH 7.4 [28,29], and the conformation of LC4 is not affected by pH, as indicated by the CD spectra. Assignment of the proton spectra was accomplished by standard sequential assignment techniques utilizing the identification of spin systems and NOE connectivities (Figure 3). The

![Figure 2](https://www.plosone.org/figure/10.1371/journal.pone.0020452.g002)

**Figure 2. Insertion of LC4 into SDS micelles monitored by acrylamide quenching of LC4 fluorescence.** (A) Stern–Volmer plot. (○), in phosphate buffer at pH 4.5; (●), in phosphate buffer at pH 7.4; (□), in SDS micelles at pH 4.5; (○), in SDS micelles at pH 7.4. Peptide-coupled SDS micelles were preincubated for 15 min before measurements. (Q) is the quencher concentration (5–85 mM). The SDS micelles concentration was 10 mM, and the peptide concentration was 50 μM. (B) Stern–Volmer constants calculated from the acrylamide quenching experiments in (A). Each error bar shows the SEM from six measurements.

doi:10.1371/journal.pone.0020452.g002
backbone resonance assignments were complete, with the exception of the amide group of Leu174. The signal of the amide proton of Leu174 was not observed in clean-TOCSY and NOESY spectra. With regard to the signals of the side-chain of Leu174, although they were very weak due to line broadening, their assignments were made by interactive interpretation of DQF-COSY, clean-TOCSY, and NOESY spectra. The proline side-chain resonances were assigned from the clean-TOCSY cross-peaks between $H_d$ protons and their respective $H_c$, $H_b$, and $H_a$ protons.

**Estimation of the secondary structure from chemical shifts and NOE connectivities**

The differences in the $H^\alpha$ chemical shift ($\Delta\delta$ in ppm) between the observed and the random coil values are shown in Figure 4.

![Figure 4. Chemical shift differences of $H^\alpha$ between the experimental values and random coil values. The experimental values were observed as the $H^\alpha$ chemical shift for 2 mM LC4 in the presence of 200 mM $d_{25}$-SDS micelles. The random coil values were obtained from Chemical Shift Index [30]. The negative $\Delta\delta$ values showed the presence of the helical conformation.](http://doi.org/10.1371/journal.pone.0020452.g004)

Negative $\Delta\delta$ values were observed at two regions, from V157 to A159 and from F166 to K171. A series of upfield $H^\alpha$ chemical shifts associated with random coil values showed the presence of an $\alpha$-helical structure [30]. Thus, LC4 is considered to assume an $\alpha$-helical conformation in these regions. Analysis of the NOESY spectrum provided the sequential and the medium range NOEs ($|i-j|$ < 5) as summarized in Figure 5. The presence of $d_{\alpha}N_{(i,i+3)}$ and $d_{\alpha}N_{(i,i+4)}$ NOEs suggests an $\alpha$-helical conformation; if $d_{\alpha}N_{(i,i+2)}$ NOEs are also observed, its region assumes a $3_{10}$-helical conformation [28]. Figure 5 shows that LC4 assumes an $\alpha$-helical conformation from F166 to K171, as suggested by the $H^\alpha$ chemical shift. In addition, from F158 to S160, characteristic $d_{\alpha}N_{(i,i+2)}$ NOEs were also observed, and thus this region adopts a $3_{10}$-helical conformation.

**NMR structure**

The three-dimensional structure of LC4 was determined using the torsion angle dynamics of the program CYANA [23]. A best fit superposition of an ensemble of the 20 lowest energy structures of LC4 in SDS micelles at pH 4.5 is shown in Figure 6A. The statistics of the structure as well as the distance restraints used for the program CYANA are summarized in Table 1. The 20 lowest energy structures in the well-ordered region (Val157–Glu172) are superimposed over the backbone (N, $C^\alpha$, $C'$) atoms and the non-hydrogen atoms, with rms deviations of 0.52 Å and 1.17 Å, respectively. The C-terminal region of residues Gly173–Leu174 is not well ordered. The quality of the structures was checked using the program PROCHECK-NMR [24]. In a Ramachandran plot, 87.1% of the non-glycine residues were located within the most favored regions as shown in Table 1. The NMR results show that LC4 possesses two helical regions ($\alpha_1$ helix: Phe158–Ser160; $\alpha_2$ helix: Phe166–Lys171) in SDS micelles (Figure 6B). $\alpha_1$ and $\alpha_2$ adopt $3_{10}$-helical and $\alpha$-helical structures, respectively, as indicated by the analysis of NOEs connectivities. Pro162 exists around the end of $\alpha_1$. It adopts a trans conformation, which was confirmed by the NOEs between the $H_a$ and $H_d$ protons [28]. The residue Pro is referred to in a negative sense, it lacks an NH group, and often disrupts the formation of the secondary structure [31]. It appears that Pro162 links $\alpha_1$ and $\alpha_2$, and promotes the bend conformation between two helices. The Connolly surface of LC4 is shown in Figure 6B. The residues Val157, Phe158, Ala159, Leu161, Pro162, Ile164, Ile165, and Phe166 contribute to the formation of the hydrophobic cluster, which stabilizes the ordered structure of LC4 bound to micelles. Arg168, Lys171, and Glu172...
Figure 5. Patterns of sequential and medium range NOE cross-peaks of LC4 in SDS micelles. The NOEs for 2 mM LC4 were derived from the NOESY spectrum with a mixing time of 150 ms in the presence of 200 mM d_{25}-SDS micelles. The NOE patterns are used to characterize the region of the helical structure [28].

Figure 6. Overall structure of LC4 in SDS micelles at pH 4.5 by 	extsuperscript{1}H-NMR. (A) Stereoview illustrating a trace of the backbone atoms for the ensemble of the 20 lowest energy structures, showing the heavy atoms of the side chains (residues Val157-Leu174). The structures in the well-ordered region (residues Val157-Glu172) was superimposed over the backbone atoms. (B) Surface representation and ribbon diagram of LC4 showing the side chains (residues Val157-Glu172). The helical regions (α1 and α2) are shown in red.

doi:10.1371/journal.pone.0020452.g005

doi:10.1371/journal.pone.0020452.g006
gather at the molecular surface, and thus the structure of LC4 exhibits an amphipathic character in the micellar environment.

Possible docking poses
Docking of LC4 and LC5 was calculated using ZDOCK and Discovery Studio. The possible binding poses were obtained as 1,000 structures with the contacting residue Gly163, which is considered to be important in susceptibility to HIV-1. Figure 7A shows a best-fit superposition of the ensemble of the 5 lowest energy structures. The rms deviation is 1.15 Å for backbone (N, Cα, C') atoms in all residues of the ordered region of LC4 and LC5. The a1 helical region of LC4 is spatially located in the vicinity of the helical region of LC5. Figure 7B shows the possible binding interface. Gly163, located next to residue Pro162 in LC4, is capable of contacting the Ile237 residue of LC5. In addition, in this position, the residues (Leu236, Ile237, and Phe238) of LC5 could interact with the residues (Ala159, Pro162, and Ile165) of LC4 which are included in the hydrophobic cluster. Therefore, the hydrophobic interactions between these residues preferentially contribute to the formation of the complex of LC4 and LC5.

Discussion
The solution structure of LC4 in SDS micelles presented here provides an insight into its functional interaction with CCR5 at an atomic level. LC4 bound to micelles clearly shows two helical regions (a1 and a2) with a flexible C-terminal region of the peptide. The structure possesses a hydrophobic cluster and several charged residues preferentially gather on the molecular surface (Figure 6B).

The CD spectra showed that LC4 assumes a random-coil conformation in an aqueous environment, but a helical conformation in SDS micelles. Furthermore, acrylamide quenching fluorescence showed that LC4 is inserted into SDS micelles through Phe158 and/or Phe166. Both of these Phe residues contribute to the formation of a hydrophobic cluster that stabilizes the LC4 structure (Figure 6B), suggesting that LC4 is embedded into micelles through these two Phe residues. The hydrophobic cluster of the N-terminal region that includes these residues might be inserted into the micelles. In the favored model of the transmembrane region of CCR5, it is considered that residues Val157-Arg168 of LC4 are inserted into the membrane and residues Ser169-Leu174 exist in the extracellular region [8]. The hydrophobic cluster of LC4 is formed by the residues for the insertion into the membrane. The conformation of LC4 and the degree of insertion of Phe into the micelles were hardly affected by a variation in pH, as indicated by the CD spectra and acrylamide quenching fluorescence.

LC5 peptide inhibits HIV-1 infection of MT-4 cells, and its three-dimensional structure determined by NMR methods was reported in our previous study [10].

Table 1. Summary of Structure Statistics of LC4 in SDS micelles

| NOE upper distance restraints | Total | 196 |
|-------------------------------|-------|-----|
| Intra residual and sequential |       | 110 |
| Medium range and long rage (i - i + 2, i - i + 3, i - i + 4) | 86 |
| CYANA target function value | 0.09Å² |

Distance constraints violations
| Number > 0.1 Å | 0 |
| Maximum        | 0.04Å |
| PROCHECK Ramachandran plot analysis² |
| Residues in favored regions | 87.1% |
| Residues in additionally allowed regions | 12.9% |
| Residues in generously allowed regions | 0.0% |
| Residues in disallowed regions | 0.0% |
| RMS deviation to the average coordinates² |
| Backbone atoms | 0.52Å |
| Heavy atoms    | 1.17Å |

²Except for the number of constraints, average values given for the set of 20 conformers with the lowest CYANA target function value.
³The values were calculated for residues 157–172.
doi:10.1371/journal.pone.0020452.t001

Figure 7. Possible docking positions of LC4 and LC5 calculated using the program ZDOCK. The NMR structure of LC5 was determined in the previous study [10]. The docking positions were calculated using the program ZDOCK and then the energy-minimization calculations were performed using the program Discovery Studio. (A) Ribbon diagrams of the 5 lowest energy structures (residues Val157–Glu172 of LC4; residues Lys229–Thr239 of LC5) (B) close-up view of the binding interface of the lowest energy structure, showing the heavy atoms of the side chains.
doi:10.1371/journal.pone.0020452.g007
LC4 with LC5 were performed using the program ZDOCK. The possible poses were obtained and superimposed with the rms deviation of 1.15 Å for backbone (N, Cα, C') atoms over the ordered regions. The LC5 structure possesses a hydrophobic cluster in the C-terminal region. The interaction between the hydrophobic clusters of LC4 and LC5 leads to the formation of a hydrophobic interface on the molecular surface. This might be an important key to the inhibitory function of LC5, when LC4 and LC5 interact with each other. Further studies are required to shed light on the role of the LC4 region of CCR5 in its HIV-1 inhibitory function.

**Acknowledgments**

The authors would like to thank Dr. Yoshitsugu Shiro, RIKEN SPring-8 Center, for the NMR instrumentation.

**Author Contributions**

Conceived and designed the experiments: KM. Performed the experiments: KM KT. Analyzed the data: KM. Contributed reagents/materials/analysis tools: KM. Wrote the paper: KM.

---

**References**

1. Agrawal I, VanHorn-Ali Z, Berger EA, Alkhatib G (2004) Specific inhibition of HIV-1 coreceptor activity by synthetic peptides corresponding to the predicted extracellular loops of CCR5. Blood 103: 1211-1217.
2. Schlecht HP, Schelhorn S, Desaube BJ, Jacobson JM (2008) New approaches in the treatment of HIV/AIDS - focus on maraviroc and other CCR5 antagonists. Ther Clin Risk Manag 4: 473-485.
3. Berger EA, Murphy PM, Farber JM (1999) Chemokine receptors as HIV-1 coreceptors: roles in viral entry, tropism, and disease. Annu Rev Immunol 17: 637-700.
4. Moore JP, Tikiola A, Dragic T (1997) Co-receptors for HIV-1 entry. Curr Opin Immunol 9: 551-562.
5. Wyant K, Sodroski J (1996) The HIV-1 envelope glycoproteins: fusogens, antigens, and immunogens. Science 280: 1804-1808.
6. Eckert DM, Kim PS (2001) Mechanisms of viral membrane fusion and its inhibition. Annu Rev Biochem 70: 777-810.
7. Toshiro AM, Kurtzkes DR (2007) Chemokine antagonists as therapeutics: focus on HIV-1. Annu Rev Med 58: 445-459.
8. Imai M, Baranyi L, Okada N, Okada H (2007) Inhibition of HIV-1 infection by synthetic peptides derived CCR5 fragments. Biochem Biophys Res Commun 353: 851-856.
9. Chill JH, Louis JM, Delaglio F, Bi (1999) Local and global structure of the rat brain sodium channel in SDS micelles. Biophys Acta 1768: 3260-3270.
10. Miyamoto K, Togaya K, Katsura H, Akaoka K, Kuroda Y (2010) Solution structure of LC5, the CCR5-derived peptide for HIV-1 inhibition. J Pept Sci 16: 165–170.
11. Chill JH, Louis JM, Delaglio F, Bax A (2007) Local and global structure of the monomeric subunit of the potassium channel KcsA probed by NMR. Biochim Biophys Acta 1768: 3260-3270.
12. Xue R, Wang S, Qi H, Song Y, Xiao S, et al. (2009) Structure and topology of Slc11a1(164-191) with G169D mutation in membrane-mimetic environments. J Struct Biol 165: 27–36.
13. Tulumello DV, Deber CM (2009) SDS micelles as a membrane-mimetic environment for transmembrane segments. Biochemistry 48: 12096-12103.
14. Li M, Cornea RL, Antye JM, Jones LR, Thomas DD (1996) Phosphorylation-induced structural change in phospholamban and its mutants, detected by intrinsic fluorescence. Biochemistry 35: 7869-7877.
15. Laskowski RA, Rose JA, Swallow WN, Thornton JM (1996) AQUA and PROCHECK-NMR: programs for checking the quality of macromolecular structures. J Mol Biol 238: 403-438.
16. Bednash K, Muggel M, Awad A, Thornton JM (1996) Improved spectral resolution in cosy 1H NMR spectra of proteins via double quantum filtering. Biochem Biophys Res Commun 227: 417-423.
17. Griesinger C, Otting, G, Wuthrich, K, Ernst, RR (1988) Clean TOCSY for proton spin system identification in macromolecules. J Am Chem Soc 110: 7870-7872.
18. Miyamoto K, Nakagawa T, Kuroda Y (2001) Solution structure of the cytoplasmic linker between domain III-86 and domain IV-81 (III-IV linker) of the rat brain sodium channel in SDS micelles. Biopolymers 59: 389-393.
19. Rance M, Sorensen OW, Bednash K, Wagner G, Ernst RR, et al. (1983) Improved spectral resolution in cosy 1H NMR spectra of proteins via double quantum filtering. Biochem Biophys Res Commun 117: 479-485.
20. Hwang TL, Shaka AJ (1995) Water Supression That Works. Excitation Sculpting Using Arbitrary Wave-Forms and Pulsed-Field Gradients. J Magn Reson 112: 273-279.
21. Delaglio F, Grzesiek S, Vuister GW, Zhu G, Pfeifer J, et al. (1995) NMRPipe: a multidimensional spectral processing system based on UNIX pipes. J Biomol NMR 6: 277-293.
22. Johnson BA (2004) Using NMRView to visualize and analyze the NMR spectra of macromolecules. Methods Mol Biol 278: 313-352.
23. Guitteri P (2004) Automated NMR structure calculation with CUYANA. Methods Mol Biol 278: 355-378.
24. Laskowski RA, Rullmann JA, MacArthur MW, Kaptein R, Thornton JM (1996) AQUA and PROCHECK-NMR: programs for checking the quality of protein structures solved by NMR. J Biomol NMR 6: 477-486.
25. Koradi R, Billeter M, Wuthrich K (1996) MOLMOL: a program for display and analysis of macromolecular structures. J Mol Graph 14: 51-55, 29-32.
26. Wache K, Pierce B, Mintseris J, Tong WW, Anderson R, et al. (2003) ZDOCK and RDOCK performance in CAPRI rounds 3, 4, and 5. Proteins 60: 207-213.
27. Yang JT, Wu CS, Martinez HM (1986) Calculation of protein conformation from circular dichroism. Methods Enzymol 130: 208-269.
28. Wuthrich K (1986) NMR of Proteins Nuclear Acids. New York: John Wiley.
29. Bai Y, Milne JS, Mayne L, Englander SW (1993) Primary structure effects on proton spin system identification in macromolecules. J Mol Graph 14: 51-55, 29-32.
30. Togaya K, Togaya, N, Murakami, K, Kuroda, Y, & Miyamoto, K (2010) Solution structure of LC5, the CCR5-derived peptide for HIV-1 inhibition. J Pept Sci 16: 165–170.
31. Bhattacharyya R, Chakrabarti P (2003) Stereospecific interactions of proline residues in protein structures and complexes. J Mol Biol 331: 925-940.