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The pH-Dependent Conformational States of Kyotorphin: A Constant-pH Molecular Dynamics Study

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ABSTRACT An extensive conformational study of the analgesic dipeptide kyotorphin (L-Tyr-L-Arg) at different pH values was performed using a constant-pH molecular dynamics method. This dipeptide showed a remarkable pH-dependent conformational variety. The protonation of the N-terminal amine was identified as a key element in the transition between the more extended and the more packed conformational states, as monitored by the dihedral angle defined by the atoms $1C\beta$ - $1C\alpha$ - $2C\alpha$ - $2C\beta$. The principal-component analysis of kyotorphin identified two major conformational populations (the extended *trans* and the packed *cis*) together with conformations that occur exclusively at extreme pH values. Other, less stable conformations were also identified, which help us to understand the transitions between the predominant populations. The fitting of kyotorphin's conformational space to the structure of morphine resulted in a set of conformers that were able to fulfill most of the constraints for the μ -receptor. These results suggest that there may be strong similarities between the kyotorphin receptor and the structural family of opioid receptors.

INTRODUCTION

The structure, function, and dynamics of most biomolecules in solution are very dependent on pH (1). pH effects occur via electrostatic interactions, one of the strongest forces at the molecular level, and can have a direct influence on molecular structure. However, a small peptide does not exhibit a preferred structure in aqueous solution, but rather an ensemble of interconverting conformations (2). In this case, pH has a more subtle effect by changing the distribution of the conformer populations (3).

Until recently, computational studies dealt with pH using mostly two simplified approaches. The first consists of setting the solute (e.g., protein) protonable groups to the states they would have in a solution at the intended pH and then performing molecular mechanics/dynamics (MM/MD) simulations. This means that any desolvation and site-site interaction effects caused by the solute environment upon protonation are entirely neglected. The second way consists of using a rigid solute structure and simplified electrostatic-oriented models, such as Poisson-Boltzmann (PB) (4–12), generalized Born (13–15), or protein dipoles and Langevin dipoles (16,17). (Several other approaches are discussed in the literature as well (18–24).) The protonation free energies thus obtained can then be used for sampling protonation states by Monte Carlo (MC) (7,25–27) or other approximate methods (7,16,28–31), making possible the calculation of pK_a values. This approach explicitly considers the effect of desolvation and site-site interactions, but the use of a single solute conformation makes it impossible to account for

structural reorganization and protonation-conformation coupling in general (32–35). These problems can be attenuated by the use of linear response approximations that combine MD with simplified approaches (16,17,32,36). Another, more general route is to use recently developed methods for constant-pH MD (37–50), some of which explore the complementarity between standard MD and simplified methods (see Machuqueiro and Baptista (50) for a more detailed comparison). One of these constant-pH MD methods, called stochastic titration (39), assumes that the dynamics of a solute with a fixed protonation state can be obtained from MM/MD, as usual, and that its electrostatics at fixed conformation can be captured by a PB model (followed by MC sampling of protonation states). Stochastic titration could successfully predict pK_a s for succinic acid (a ΔpK_a of 1.75 for an experimental value of 1.45) (39) and decalysine (pK_a of 10.2 for both simulation and experiment) (50), showing that the method can properly model the coupling between protonation and conformation in small and relatively large molecules.

Kyotorphin (KTP) was first isolated from bovine brain by Takagi and co-workers (51,52). The name was assigned to reflect an endorphin-like substance and the fact that it was discovered in Kyoto, Japan. This endogenous dipeptide (L-Tyr-L-Arg) belongs to the neuropeptide family (53–58) due to its opiate-like activity. KTP is distributed unevenly in regions of rat brain (52) and may be formed both by processing precursor proteins and by the biosynthesis from Tyr and Arg (57). The Tyr-Arg motif exists widely throughout the brain, not only as KTP, but also as the N-terminal part of several endogenous analgesic peptides (59,60). Also, this peptide is very rapidly degraded by aminopeptidases (61). Many of these properties are typical of neurotransmitters and it is not surprising that KTP has also nonopioid

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actions independent of enkephalin release (62). There is evidence suggesting that KTP does not bind the opiate receptors (μ , δ , κ) but that instead it exerts a Met-enkephalin-releasing force (53). These results led to the suggestion that the dipeptide binds to a specific receptor (KTP receptor, KTPr) (56), triggering a cascade of events that leads to strong analgesia in the brain (57,63). Two mechanisms of action have been proposed: 1), direct activation of the KTPr, inducing Met-enkephalin release (which can activate the δ -receptor), followed by $G\alpha_i$ and phospholipase C activations (64,65); 2), a fast degradation of KTP, resulting in L-Arg, which is a potent substrate for nitric oxide synthase, with the NO thus formed inducing the release of Met-enkephalin (66). Despite the fact that several studies (45,53,54) confirm the existence of a KTPr, it has not yet been identified. There is still the question of whether the KTPr is specific (45) or the result of a mixed oligomerization of μ - and δ -opioid receptors (67). Recent results suggest that KTP can also release opioid peptides from rat cardiac muscle and have an indirect regulatory role in β -adrenergic action through cross-talk with opioid receptors (68).

The molecular interaction between KTP and its receptor remains unknown. In fact, not much is known besides the fact that the receptor seems to be functionally coupled to $G\alpha_i$ and that it is strongly antagonized by the dipeptide L-Leu-L-Arg (64). There are obvious similarities between KTP and most of the endogenous opioid peptides and even with morphine. The L-Tyr residue at the first position of the peptide is present in most of the opioid peptides and is believed to be crucial for receptor recognition (69,70) due to both π -stacking (71) and hydrogen-bonding with the alcohol of the phenol group (72). A protonable N-terminal group is also present in KTP and, when protonated, can form a salt bridge with an anionic group in the receptor, which is typical for the morphine/ μ -receptor interaction (72). Also, when KTP's N-terminal group is protonated, the molecule becomes positively charged, which is an important characteristic present in both dynorphin A and nociceptin/orphanine FQ. These cationic peptides bind to two receptors that exhibit a highly acidic second exofacial loop (κ -receptor for dynorphin A and ORL1 receptor for nociceptin/orphanine FQ) (69,70). On the other hand, there is an important difference between KTP and the opioid peptides. The introduction of two methyl groups (*ortho* position) in the phenol ring of the Tyr residue in KTP (H-Dmt-Arg-OH) is detrimental to the interaction with the KTPr (62), unlike in the case of opioid peptides, where replacement of Tyr with Dmt produces a very significant potency enhancement (73). Thus, the binding site of KTPr interacting with the Tyr residue of KTP has structural requirements that differ from those of the binding sites of opioid receptors interacting with the Tyr residue of opioid peptides (62).

Recently, the structural constraints needed for KTP-receptor interactions were addressed with a combination of experimental and theoretical approaches (74). The study

showed that pH has a strong influence on the conformational behavior of KTP and an acylated KTP derivative, both in water and in model systems of membranes. In fact, the conformational space of KTP is expected to depend strongly on pH, due to its four titrable sites. Around physiological pH, the N-terminus protonation state should have a strong effect on both the conformation and total charge of the molecule.

The aim of this work is threefold. First, this work intends to study KTP using the stochastic titration method for constant-pH MD (39,50), which is theoretically more rigorous than the linear response approximation (LRA) method used previously (74); analogously to stochastic titration, that LRA method combines PB/MC and MM/MD, but it introduces further assumptions (32). The second aim of this work is to carry out a more extensive conformational analysis of the system than the one presented by Lopes and co-workers (74). Third, this work also intends to further test the performance of the stochastic titration method, which has so far only been applied to succinic acid (39) and decalysine (50). Overall, our general objective is to elucidate KTP's preferred conformations at different pH values and how they correlate with known structural constraints needed for opioid ligand-receptor interactions.

COMPUTATIONAL DETAILS AND METHODS

Constant-pH MD method

The constant-pH MD method implemented is the stochastic titration method previously proposed (39). The current implementation is described in a recent publication (50), which should be consulted for further details. The method relies on different sequential blocks (Fig. 1). The first block is a PB/MC calculation. The protonation states resulting from the last MC step are assigned to the protein. The second block is the solvent-relaxation dynamics, a short MM/MD simulation of the system with frozen protein, which allows the solvent to adapt to the new protonation states (the duration of this block is hereafter designated as τ_{relax}). The last block is a full MM/MD simulation of the unconstrained system (the duration of this block is hereafter designated as τ_{prot}). A variant of the reduced titration method is used (31,50), in which we run a full PB/MC calculation on the system every 10th step to assign a fixed state to all the sites whose mean occupancies fall outside a predefined threshold (0.001). This approach is very useful, because, when simulating pH values too far away from certain site pK_a s, these become temporarily excluded from the calculations, hence strongly decreasing the computational cost.

For the PB and MC calculations we used the programs MEAD (75) and MCRP (27), respectively. The atomic charges and radii used in the PB calculations were derived from the GROMOS 43A1 force field (76,77). All PB calculations consisted of finite-difference linear Poisson-Boltzmann calculations performed with the program MEAD (version 2.2.0) (75) at a temperature of 300 K, a molecular surface defined with a solvent probe radius of 1.4 Å, and a Stern (ion-exclusion) layer of 2.0 Å. The dielectric constants were 2 for KTP and 80 for the solvent. A two-step focusing procedure (78) was used, with consecutive grid spacings of 1.0 and 0.25 Å. The MC runs were performed using 10^5 MC cycles, one cycle consisting of sequential state changes over all individual sites and also all pairs of sites with at least one interaction term above 2.0 pK_a units (27). In all cases, the computed average protonations had errors lower than 0.0001; for each site, this error is computed as the correlation-corrected standard deviation of the average, taking as correlation time the value at which the autocorrelation of the protonations goes below 0.1.

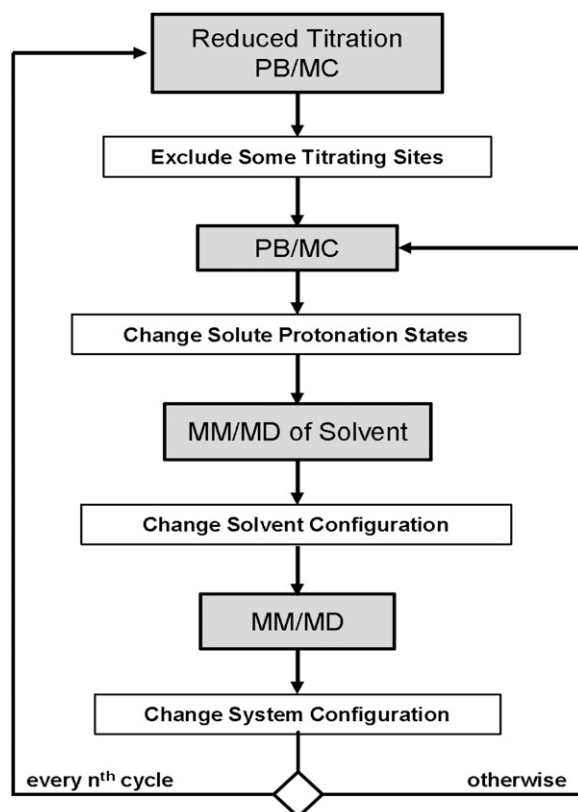


FIGURE 1 Schematic representation of the algorithm used.

MM/MD settings

All simulations were performed with the GROMOS 43a1 force field (76,77) for the GROMACS 3.2.1 distribution (79,80) (with some modifications (50)) and with the SPC water model (81). Periodic boundary conditions were used with a rhombic dodecahedral unit cell. A twin-range cutoff was used, with short- and long-range cutoffs of 8 Å and 14 Å, respectively, and with neighbor lists updated every five steps. Long-range electrostatics interactions were treated using the generalized-reaction-field method (82). The ionic strength in the simulations was set as an external parameter using a modified version of the GROMACS 3.2.1 distribution (50). All bond lengths were constrained using the LINCS algorithm, and a time step of 2 fs was used. Two Berendsen's temperature couplings to baths at 300 K, and with a relaxation time of 0.1 ps (83), were used for both the solute and solvent. A Berendsen's pressure coupling (83) was used with a relaxation time of 0.5 ps and a compressibility of $4.5 \times 10^{-5} \text{ bar}^{-1}$.

The dipeptide was built in an extended conformation, placed in the center of the box and filled with 1199 water molecules. The system was minimized first with ~ 40 steps of steepest descent followed by ~ 700 steps using the low-memory Broyden-Fletcher-Goldfarb-Shanno algorithm. The initiation was achieved by harmonically restraining all atoms in a 50-ps MD simulation, followed by another 50 ps simulation with only the CA atoms restrained. Note that this initiation was done for two different startup systems, one with the N-terminus deprotonated and the other with the N-terminus protonated (Tyr and Arg side chains were protonated and the C-terminus was deprotonated). The charged N-terminus conformation was used as the starting structure for the simulations at pH 2–7, whereas simulations at pH 8–12 started with the neutral N-terminus conformation.

Runs of 100 ns were performed and the first 5 ns were discarded to get the system equilibrated. The relaxation of the solvent (τ_{rlx}) and the full system dynamics segment (τ_{prt}) were done for 1.0 ps each. If τ_{prt} is too short, there will

be an increase of computational cost, and if it is too long, we will have a slower conformational sampling. In this work, the τ_{rlx} and τ_{prt} values were optimized to get high conformational sampling and good system stability at the expense of a high computational cost (we “waste” 50% of our MM/MD time).

The main simulations were performed at ionic strength of 0.1 M. Two extra sets of simulations of 30 ns were performed at ionic strengths 0.0 and 1.0 M and at pH values 3, 5, 6, 7, 8, 9, and 11.

General analysis

All structural analyses were done using GROMACS (79,80) or in-house tools. Errors were computed using standard methods (84).

Principal-component analysis

Principal-component analysis (PCA) is a method that can be used to simplify a dataset (85). It is a linear transformation that changes a set of (possibly) correlated variables into a set of uncorrelated variables called principal components (PCs). The first PC (i.e., the PC with the largest eigenvalue) accounts for as much of the variability in the data as possible, and each succeeding PC (in order of decreasing eigenvalues) accounts for as much of the remaining variability as possible, while being uncorrelated with the previous ones. The strongest correlations in the dataset tend to be captured by the first PCs. PCA can be used for reducing dimensionality by retaining only a few of the first PCs, hopefully without much loss of information. PCA is algebraically equivalent to the diagonalization of the covariance matrix: each PC is the variable associated with an eigenvector, and its variance is given by the corresponding eigenvalue.

In our analysis, we fitted separately all our trajectories at different pH values to a fixed KTP structure; changing this structure had no major effect on the results (data not shown). The atoms subsequently used to calculate the covariance matrix were CA, CB, CG, and C from Tyr, and N, CA, CB, CG, and NE from Arg. The trajectories were projected on the first three principal components, which account for $\sim 75\%$ of the information available. The set of all points (conformations) thus projected was used to obtain a kernel estimate of the corresponding probability density (86), which was computed on a grid of $(0.1 \text{ Å})^3$ bins, using a Gaussian kernel and a bandwidth of 0.2 Å. This probability density is the projection of the original isothermal-isobaric distribution onto the subspace spanned by the first three PCs, and is expected to retain the major features of the original distribution. The most populated peaks of the density (usually the higher ones) were then selected using an energy cutoff of $2RT$, where the energy was computed according to

$$E(\mathbf{x}) = -RT \ln \frac{P(\mathbf{x})}{P_{\text{max}}},$$

where \mathbf{x} is the coordinate in the 3D space of the first PCs and $P(\mathbf{x})$ is the probability density at point \mathbf{x} . A representative conformation was chosen for each peak, namely the one closest to the peak top.

RESULTS AND DISCUSSION

Titration with constant-pH MD

The titration of KTP (Figs. 2 and 3) was computed by averaging at each pH value the occupancy states of all four titrable sites over 95 ns for the simulations at ionic strength 0.1 M, and over 25 ns for the simulations at ionic strengths 0.0 and 1.0 M.

From the titration curves, we observe that the ionic-strength effect is very weak (Fig. 2). These results are not surprising because ionic strength works by shielding the

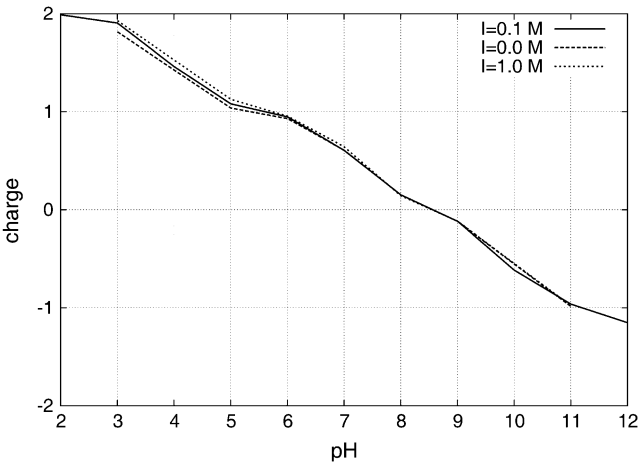


FIGURE 2 Titration curves of KTP at different values of ionic strength.

interactions between close charged groups, and KTP is a small molecule without a unique conformation in water. According to these results, KTP is only mildly positively charged at physiological pH, reaching the isoelectric point around pH 8.5. When looking at each site specifically (Fig. 3), we obtain pK_a values for the three relevant titrable sites (Arg guanidinium is titrating very far away from physiologic conditions) (Table 1).

It is interesting to compare the N-terminus titration observed here with the theoretical and experimental ones in Lopes et al. (74). The pK_a of 7.14 computed here with constant-pH MD is higher than the 6.2 value previously computed with the LRA method (74). Given that both methods rely on the same PB/MC and MM/MD approaches (but combined differently), this discrepancy may be due to sampling problems in either method or to failure of the linear response assumption. In particular, we show below (see “Detailed conformational analysis with PCA”) that some conformational classes show sudden changes upon N-termi-

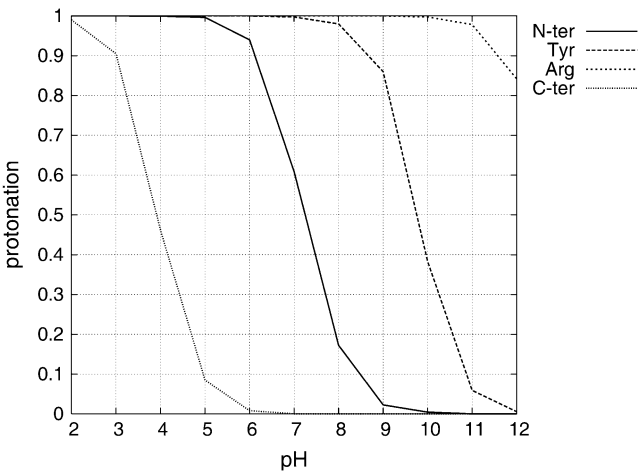


FIGURE 3 Titration curves of KTP titrable sites at 0.1 M.

TABLE 1 pK_a values obtained from the titration curves at $I = 0.1$ M

Titration site	pK_a values
N-ter	7.14 ± 0.14
Tyr	9.94 ± 0.26
Arg	>12
C-ter	4.04 ± 0.15

pK_a values were computed using a third-order polynomial interpolation.

nus titration, which could result in poor distribution overlap and consequent breakdown of the linear response assumption (see Eberini et al. (32) for details). On the other hand, 6.2 is also the pK_a value estimated from the ion exchange experiment (74) (we note that Fig. 6 a in Lopes et al. (74) shows the release of positively-charged KTP from an anionic resin, thus corresponding only to the N-terminus group. This value seems too low for such an exposed group and actually makes problematic the interaction with the negatively charged membrane (74). In any case, this issue could probably be clarified by further experimental studies.

Conformational mapping with constant-pH MD

As anticipated, our simulations clearly show that KTP does not have a unique structure in solution. In fact, this dipeptide showed a remarkable conformational freedom, especially at low pH values. We started by measuring the molecule packing at different pH values. This was done by computing the radius of gyration and the perimeter of the triangle formed by the following atoms: N from the N-terminus; O from Tyr hydroxyl; and C from Arg guanidinium. Both the radius of gyration and the perimeter are expected to be higher when the molecule is in an extended conformation and lower when the molecule is more packed. As seen in Fig. 4, there

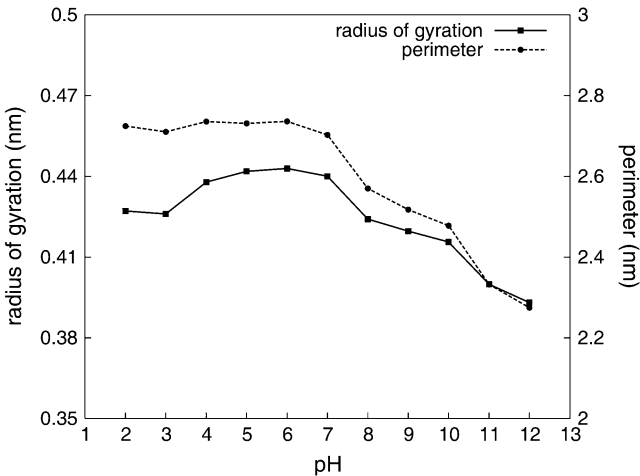


FIGURE 4 Average radius of gyration and average perimeter (triangle formed between N-terminus, OH (Tyr), and CZ (Arg)) over pH, at 0.1 M ionic strength.

is a clear gain of packing with pH increase. Also, the radius of gyration indicates a slight structural packing at lower pH values, suggesting that some structure other than the extended one is being preferred.

From these results, an obvious question arises, namely what kind of structures are being formed at lower and higher pH values. A possibility would be that conformational changes involve an interaction between the two side chains. For this interaction to take place, the correct alignment between the C_α and C_β atoms of both residues would be of major importance. With this in mind, we monitored the dihedral angle formed between the atoms $C\beta_{Tyr}-C\alpha_{Tyr}-C\alpha_{Arg}-C\beta_{Arg}$. When KTP is in an extended conformation (Fig. 5 *a*), this dihedral angle takes values around 180° (*trans*), and when the side chains interact with each other (Fig. 5 *b*), the dihedral angle takes values around 0° (*cis*).

Our simulations indicate that KTP has a clear preference for the *cis* conformation upon pH increase (Fig. 6). Interestingly, at relatively low pH both the N- and C-terminus are charged and KTP is preferably in the *trans* conformation, keeping both termini far apart. Upon pH increase, the N-terminus loses the charge and KTP prefers the *cis* conformations. These results might seem contradictory at first glance because opposite charges should attract each other. However, these groups are very exposed to water, and thus solvation can easily overcome this attraction (see Suppl. Mat.). In fact, salt bridges between solvent-exposed groups

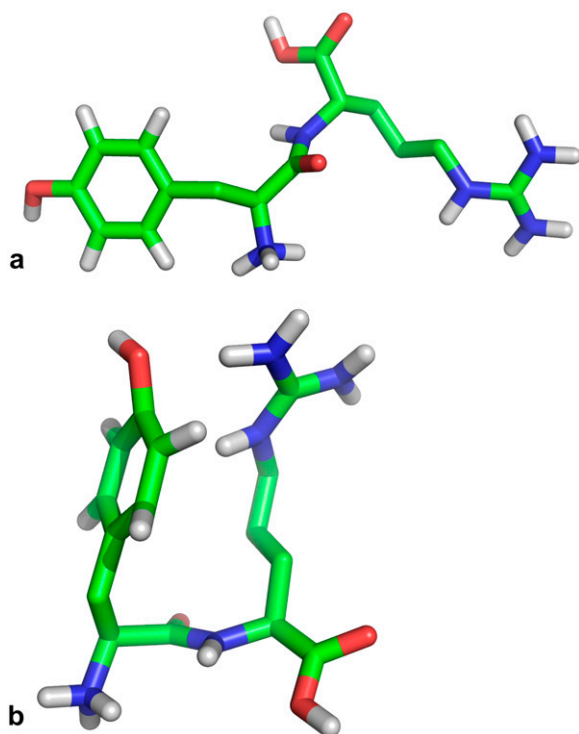


FIGURE 5 Two typical conformations of KTP in water. (*a*) *trans* extended conformation; (*b*) *cis* packed conformation. The fully protonated form is represented.

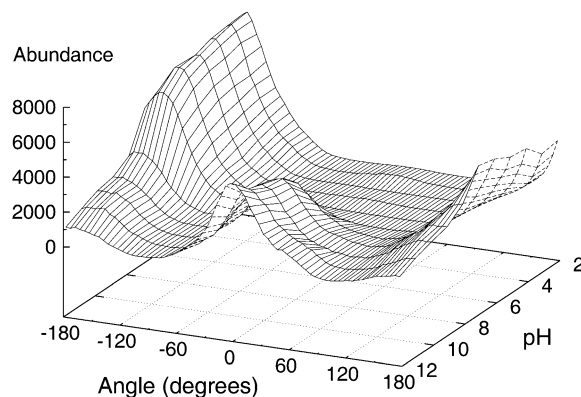


FIGURE 6 Histogram surface of the dihedral angle CB-CA-CA-CB for KTP at ionic concentration of 0.1 M at different pH values. Ninety-five thousand conformations are used for each pH value. Similar data is observed for 0.0 and 1.0 M ionic strength (data not shown).

are usually very weak (87,88), often breaking during simulations (89,90).

When looking at the histogram surface (Fig. 6), two distinct populations can be separated from each other. We considered a conformation to be *cis* if the dihedral angle is between -70° and 100° , and *trans* otherwise. In this way, we can quantify the amount of *cis* KTP at different pH values (Fig. 7, *solid squares*). From these results, it is clear that the *trans/cis* transition is induced not only by the deprotonation of the N-terminus (pH 7–8) but also by the deprotonation of the Tyr-OH (pH 10–11).

The interaction of the side chains was monitored by following the distance between Tyr-OH and the closest Arg-NH atoms. From the minima in the histograms (data not shown), we chose a cutoff of 6 Å to define an interaction, and used this criterion to determine the number of *cis* conformations with packed side chains (Fig. 7, *dashed circles*).

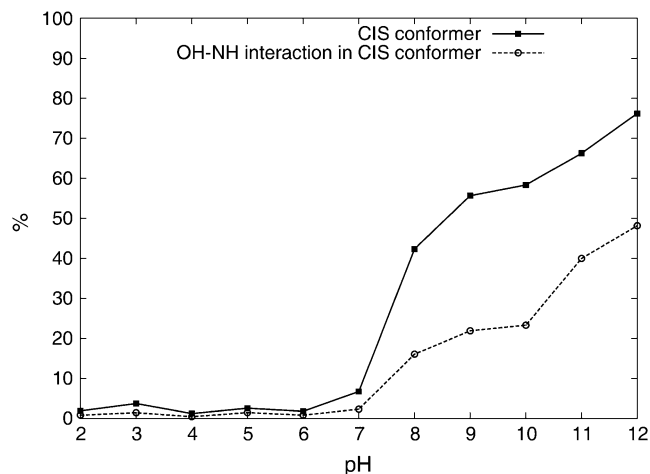


FIGURE 7 Percentage of *cis* conformations over pH (*solid squares*) and the percentage of *cis* conformations with Tyr-Arg interactions (cutoff = 6 Å) (*dashed circles*).

These results show that there is a very relevant number of *cis* conformations in which one or both of the side chains prefer to be solvated rather than interact with each other.

Detailed conformational analysis with PCA

From analysis of the dihedral angle CB-CA-CA-CB we observed that there is a clear transition from *trans* to *cis* upon pH increase, but the question remains whether all *trans* and *cis* conformers are in extended or packed conformations. In Fig. 7, we showed that there are *cis* conformers with noninteracting side chains. To better characterize the occurring conformations, we did a PCA analysis at different pH values, whose results are shown in Fig. 8 and Table 2. We observe that at very low pH values, the C-terminus is protonated, leading to the preference for a *trans* conformation with Tyr-OH interacting with the C-terminus (T_{t-c}). The T_{swch} conformation seems to be a particular case of the

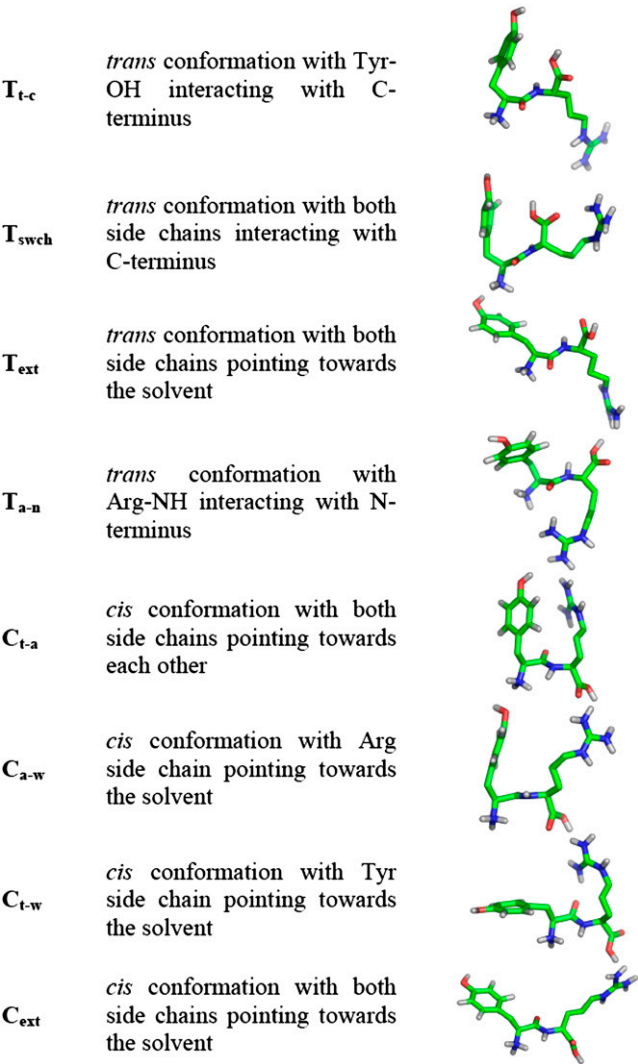


FIGURE 8 Representative conformations from most populated peaks. The fully protonated forms are represented.

TABLE 2 Population percentage of each conformational peak at different pH values

	pH											
Structure	2	3	4	5	6	7	8	9	10	11	12	
T _{l-c}	80.7	71.1	51.4	29.3	25.5	12.8	2.4	—	—	—	—	
T _{swch}	—	6.0	—	—	—	—	—	—	—	—	—	
T _{ext}	17.1	18.2	47.7	68.5	72.3	84.4	56.6	43.6	42.9	33.1	20.8	
T _{a-n}	—	—	—	—	—	—	—	—	—	2.4	4.5	
C _{t-a}	1.6	4.2	0.8	1.0	0.6	1.9	34.0	42.2	44.7	54.1	66.8	
C _{a-w}	—	—	—	—	—	0.5	3.9	2.7	8.0	10.1	4.7	
C _{t-w}	—	—	—	—	—	—	1.2	2.2	2.3	—	2.1	
C _{ext}	—	—	—	—	—	—	1.6	8.8	1.7	—	1.0	

T_{t-c} peak where the Arg's side chain is also interacting with the C-terminus. With the increase of pH, the C-terminus deprotonates and prefers to become solvated, favoring the *trans* conformation with both side chains pointing toward the solvent (T_{ext}). Above pH 8.0, the N-terminus is mainly deprotonated, which favors the formation of *cis* conformers. PCA was able to identify at least four distinct *cis* peaks, with the differences lying in the positions of the side chains. By far the largest peak is the one where both side chains are pointing toward each other (C_{t-a}). The conformations where one or both the side chains are pointing toward the solvent (C_{t-w} , C_{a-w} , and C_{ext}) can be seen as transient. At very high pH values, the Arg side chain deprotonates and allows an interaction between Arg-NH and the N-terminus in a *trans* conformation (T_{a-n}).

These PCA results are in good agreement with the cluster-analysis study of Lopes et al. (74), which reports that the main conformation clusters obtained for the charged and neutral N-terminus states were, respectively, the T_{ext} and the C_{t-a} conformers (using the nomenclature discussed above).

Conformational fit to morphine

Morphine is a pentacyclic compound where almost all atoms are constrained with little freedom. Part of the structure of KTP has a strong resemblance to morphine. In fact, the Tyr residue at the N-terminus of KTP can be mapped on morphine, fully matching its aromatic ring, as well as part of its cyclohexane and piperidine rings. For the conformational fit, we computed the non-mass-weighted root mean-square deviation (RMSD) between all the conformations at all pH values (1.1×10^6), and the morphine structure taken from its complex with an antimorphine antibody (PDB code 1Q0Y) (91). In this calculation, the atoms N, CA, CB, CG, CD1, CD2, CE1, CE2, CZ, and OH were considered. Due to the possible rotation of the phenol ring, two RMSD calculations were done for each conformation (CD1-CE1 exchanged with CD2-CE2), and the lowest value was chosen. Since most of the superimposed atoms are geometrically constrained by the phenol ring, the RMSD value essentially measures the different orientation of the N and CA atoms in both molecules. In this way, we obtained an RMSD value

of 1.083 Å for the worse fit and a value of 0.142 Å for the best fit. When a cutoff of 0.2 Å was applied, we obtained 156 structures (0.014%) which showed an almost perfect overlap with morphine (Fig. 9). This shows that KTP can fulfill the structural constraints present in morphine. In this set of conformers, there are conformations coming from all pH values, with no clear preference for any particular one (data not shown). There is also no preference for any *cis* or *trans* conformation (data not shown). Fig. 9 also shows that a very good fit to morphine turns out to select conformers in which the Arg side chain never approaches the phenol ring, nor even the morphine template.

CONCLUSIONS

We have reported a full conformational study of KTP at different pH values, using a constant-pH MD simulation method. We showed that KTP does not have a unique structure in aqueous solution and that the pK_a values are not significantly shifted from those of the individual amino acids. Furthermore, ionic strength does not have a relevant effect on the titration curves.

The protonation of the N-terminus amine is a preponderant factor in the definition of both the charge and the conformation of KTP for receptor interaction. Even though we predict a pK_a of 7.14 for the amine, it is generally accepted that the pK_a can be slightly higher close to the negatively charged membrane, and that the local pH should be lower than that of the cytoplasmatic (92). There-

fore, it is likely that, when interacting with its receptor, KTP is mainly protonated at the N-terminus.

We observed that pH has a strong influence on KTP's conformational preferences. From the packing and the side-chain position studies, we were able to identify one of the most important conformational transitions in KTP: the preference for the *cis* in detriment of the *trans* conformations upon pH increase. The key points for this transition are the deprotonation of both the N-terminus and the Tyr-OH sites. With PCA, we were able to characterize the different populations within the *cis* and *trans* groups. The two main populations were the *trans* extended conformation (T_{ext}) and the *cis* packed conformation with both side chains interacting with each other (C_{t-a}). Besides the low-populated peaks with a transient character (T_{swch} ; C_{t-w} ; C_{a-w} and C_{ext}), the method was able to identify two populations that are only present at extreme pH values (T_{t-c} and T_{a-n}).

The structural information about the KTP_r is scarce. We know that its binding pocket has to be different from the opioid receptors (53,62,73). Nevertheless, there are certain features in KTP that are common to the opioid analgesics, and which may direct the interaction with its receptor: 1), the phenolic ring can interact with the receptor both by π -stacking and hydrogen bonding; and 2), the N-terminus amine group, when protonated, can make an ionic bond with an acidic residue in the receptor. Another important aspect of the KTP-receptor interaction is orientation. The question of whether KTP can adopt the correct orientation for receptor binding is difficult to answer due to the absence of structural information on the receptor. By fitting the KTP conformations to the structure of morphine, we tried to exclude the hypothesis that the KTP_r pocket is similar to that of the μ -receptor. Interestingly, we observed exactly the opposite. The best-fitted KTP conformations revealed several nonobvious aspects: 1), there was no preference for any N-terminus charge or *cis/trans* conformation in the set; and 2), the predominance of the extended conformations (both *cis* and *trans*) results in the Arg side chain not invading the vicinities of the phenol group of the Tyr, allowing this to be presented to the receptor without any hindering effects. These results suggest that, despite the fact that KTP does not bind to the opioid receptors, there is no indication for the KTP_r pocket to be very different from those of the opioid receptors.

SUPPLEMENTARY MATERIAL

An online supplement to this article can be found by visiting BJ Online at <http://www.biophysj.org>.

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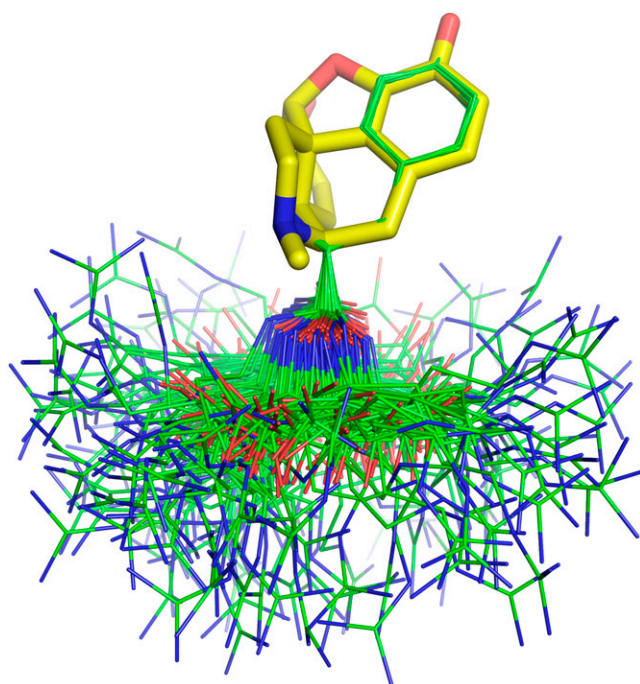


FIGURE 9 Fit of the 156 best conformations of KTP (lines, carbons in green) to the structure of morphine (sticks, carbons in yellow). All protons are omitted for clarity.

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