Towards Quantitative Classification of Folded Proteins in Terms of Elementary Functions
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A comparative classification scheme provides a good basis for several approaches to understand proteins, including prediction of relations between their structure and biological function. But it remains a challenge to combine a classification scheme that describes a protein starting from its well organized secondary structures and often involves direct human involvement, with an atomary level Physics based approach where a protein is fundamentally nothing more than an ensemble of mutually interacting carbon, hydrogen, oxygen and nitrogen atoms. In order to bridge these two complementary approaches to proteins, conceptually novel tools need to be introduced. Here we explain how the geometrical shape of entire folded proteins can be described analytically in terms of a single explicit elementary function that is familiar from nonlinear physical systems where it is known as the kink-soliton. Our approach enables the conversion of hierarchical structural information into a quantitative form that allows for a folded protein to be characterized in terms of a small number of global parameters that are in principle computable from atomary level considerations. As an example we describe in detail how the native fold of the myoglobin 1M6C emerges from a combination of kink-solitons with a very high atomary level accuracy. We also verify that our approach describes longer loops and loops connecting α-helices with β-strands, with same overall accuracy.

I. INTRODUCTION

Comparative protein classification schemes such as CATH [1] and SCOP [2] are among the most valuable and widely employed tools in bioinformatics based approaches to protein structure. These schemes classify folded proteins in terms of their geometric shape, starting from prevalent secondary structures such as α-helices and β-strands. But at the moment the final stages of the classification usually involve manual curation, and consequently these schemes are best suited for qualitative analysis of folded proteins.

The goal of the present article is to develop novel tools that we propose can eventually provide a firm quantitative basis for the existing protein classification schemes. Ultimately we hope to close gaps between bioinformatics based protein structure classification and physics based atomary level approaches to protein folding, to comprehensively address wide range of issues such as protein structure prediction and relations between shape, function and dynamics. In this way we hope to open doors to new ways to perform evolutionary, energetic and modelling studies.

Our approach is based on the recent observation [3], [4] that the geometric shape of helix-loop-helix motifs can be captured by a single elementary function that is familiar from the physics of nonlinear systems where it describes the kink-soliton. This function involves only a relatively small set of global parameters but still characterizes an entire super-secondary structure involving two (α-)helices and/or (β-)strands in addition of the loop that connects them. In [3] only individual supersecondary structures in relatively simple proteins and with quite short loops were considered. The approach proposed there did not work very well for entire protein chains, involving several helices and loops, it was essentially limited to a relatively short single loop with adjoining helices. The purpose of the present article is to show that the method can be developed to describe an entire protein and not just its helix-loop-helix segments. The protein can also be quite complex, it can involve several loops, both short and long and including those that connect α-helices with β-strands. Furthermore, the original Ansatz can be even simplified without affecting its accuracy. Remarkably we observe no loss of accuracy even when the length and complexity of the protein chain increases. Indeed, there does not appear to be any limitations whatsoever that have to be imposed on the complexity of the protein, for our approach to remain practical.

Our motivation derives from an investigation of nonlinearities that are generic in the force fields employed in...
classical molecular dynamics, a technique that is widely used in various theoretical studies of the structure, dynamics and thermodynamical properties of proteins, and in determining their folding patterns in x-ray crystallography and NMR experiments [5]. A classical molecular dynamics approach like AMBER [6] and GROMACS [7] describes the evolution of a folding protein in terms of Newton’s law that determines the time dependence of the atomary spatial coordinates $\mathbf{X}(t) = \{\mathbf{x}_i(t)\}$

$$m_i \ddot{x}_i(t) = - \nabla_i U(\mathbf{X})$$  \hfill (1)

Here $i = 1, \ldots, N$ catalogue the individual atoms both in the protein molecule and its environment, and $U(\mathbf{X})$ is an empirically constructed potential energy that governs the relevant mutual interactions between all atoms involved.

Generically the potential energy is written as the sum of two terms [6]

$$U(\mathbf{X}) = \sum_i U_{\text{covalent}}(\mathbf{X}) + \sum_i U_{\text{rest}}(\mathbf{X})$$ \hfill (2)

The first term describes the covalent two-, three-, and four-body interactions between all covalently bonded atoms. The second term describes the non-covalent interactions between all atoms. For example, in the widely used harmonic approximation the two-body contribution to potential energy that describes the vibrational motion of all pairs of covalently bonded atoms acquires the familiar form

$$U^{(2)}_{\text{bond}} = \sum_{\text{bonds}} k_{ij} (|x_i - x_j| - r_{ij})^2 \hfill (3)$$

where $r_{ij}$ are the equilibrium distances between the pairs of covalently bonded atoms $i$ and $j$, and $k_{ij}$ are the ensuing spring constants.

But there are also nonlinear corrections to the potential energy such as [8], albeit in practice they may be difficult to account for in a systematic manner. The study of these nonlinearities forms a basis of the present work.

We start with a Gedanken experiment where we scrutinize a highly simplified version of an improvement to the harmonic approximation [4], with only a single (relative) coordinate on a line $x$ so that Newton's equation is mere

$$m \ddot{x} = - \frac{dV}{dx}$$

where the potential has the form

$$V(x) = \frac{1}{2} k(x - a)^2 \approx \frac{1}{4} \kappa (x + b)^2 \cdot (x - a)^2$$

That is we account for nonlinear deviations from the harmonic approximation by promoting the spring constant to a $x$-dependent quantity. The equilibrium position $x = a$ of the harmonic approximation is recovered when $|x| \approx |a| << |b|$, but here we retain the full potential. We introduce

$$c = - \frac{1}{2} (b + a)$$

and define

$$y = x - \frac{1}{2} (a - b)$$

to arrive at the familiar "$\lambda \phi^4$" (kink) equation of motion

$$\ddot{y} = - \frac{\kappa}{m} y(y^2 - c^2)$$

with the explicit dark soliton solution

$$y(t) = c \cdot \tanh\left(c \sqrt{\frac{\kappa}{2m}} (t - t_0)\right)$$
\[ x(t) = y(t) + \frac{1}{2}(a - b) \]

\[ = -\frac{b \cdot e^{\varepsilon \sqrt{\frac{c t}{2m}}(t-t_0)} - a \cdot e^{-\varepsilon \sqrt{\frac{c t}{2m}}(t-t_0)}}{\cosh[c \sqrt{\frac{c t}{2m}}(t-t_0)]} \]  (4)

This is the hallmark dark soliton (kink) configuration that interpolates between the two uniform ground states at \( x = a \) and \( x = -b \) when \( t \to \pm \infty \). The parameters \( a, b, t_0 \) and the combination \( c \sqrt{\frac{c t}{2m}} \) are the canonical ones that characterize the asymptotic values of \( x(t) \) i.e. minima of the potential, and the size and location of the soliton. It is also noteworthy that for finite \( t \) the soliton \([4]\) describes a configuration with an energy above the uniform ground state \( x = a \) (or \( x = b \)) but that nevertheless cannot decay into \( x = a \) (or \( x = b \)) through any kind of continuous finite energy transformation: A soliton configuration such as \([4]\) cannot be obtained from any approach that only accounts for perturbations that describe small localized fluctuations around the uniform background ground state.

We argue that our example is not just an academic exercise but can be developed into a systematic tool to quantitatively characterize the geometrical shape of super-secondary structures such as helix-loop-helix motifs. In fact, we propose that the very same function \([4]\) with \( t \) a length parameter that measures distance along a static protein backbone, together with its asymmetric generalization of the form

\[ \hat{x}(t) = \frac{b \cdot e^{c_1(t-t_0)} - a \cdot e^{-c_2(t-t_0)}}{e^{c_1(t-t_0)} + e^{-c_2(t-t_0)}} \]  (5)

which becomes handy e.g. when we consider loops connecting an \( \alpha \) helix with a \( \beta \) strand, can describe the geometry of native folds of proteins in Protein Data Bank (PDB) \([8]\). Besides the four canonical soliton parameters that we have specified, we need to introduce only two additional independent global parameters to characterize a given super-secondary structure such as a helix-loop-helix motif and even an entire folded protein, with an atomary level accuracy that matches the resolution in experimental data.

As an explicit example we have chosen myoglobin, a widely studied oxygen-binding protein of both historical and biological interest that has been discussed extensively in most biochemistry texts. Specifically, we have selected the 153 amino acid myoglobin with Protein Data Bank code 1M6C whose all-atom structure is known to an all-atom resolution of 1.90 Å in root-mean-square distance (RMSD) from x-ray diffraction measurements \([8]\). We analyze it in detail, to show that its entire fold can be encoded into the global parameters of the elementary function \([4]\), \([5]\) with a RMSD accuracy of 1.27 Å for the central \( C_{\alpha} \) carbons. Moreover, as the myoglobin only involves super-secondary structures with \( \alpha \) and 3/10 helices that are connected by relatively short loops, we also verify that our approach can be extended to longer loops, and loops that connect \( \alpha \) helices with \( \beta \) strands. For this we analyze an \( \alpha \) helix - loop - \( \beta \) strand segment in the HIV-1 reverse transcriptase protein with PDB code 3DLK. The loop is now clearly longer than those in myoglobin, nevertheless we find that it can be described with comparable RMSD accuracy by the profile \([5]\).

**II. MYOGLOBIN AS MULTISOLITON**

In order to describe the PDB fold of a relatively complex protein such as the 153 amino acid 1M6C in terms of the single elementary function \([4]\), we start by computing the values of its discrete Frenet curvature \( \kappa_i \) and Frenet torsion \( \tau_i \) from the PDB data. The relevant equations are as follows (for detailed derivation, see \([9]\)): From PDB we get the three dimensional coordinates \( r_i \) of the central \( \alpha \)-carbons \((i = 1, ..., N)\). With these we compute the tangent vector \( t_i \) and the binormal vector \( b_i \) using

\[ t_i = \frac{r_{i+1} - r_i}{|r_{i+1} - r_i|} \quad \& \quad b_i = \frac{r_{i-1} \times r_i}{|r_{i-1} \times r_i|} \]  (6)

and the normal vector is given as

\[ n_i = b_i \times t_i \]

These three vectors are subject to the discrete Frenet equation

\[ \begin{pmatrix} n \\ b \\ t \end{pmatrix}_{i+1} = \exp\{-\kappa_i \cdot T^2\} \cdot \exp\{-\tau_i \cdot T^3\} \begin{pmatrix} n \\ b \\ t \end{pmatrix}_i \]  (7)
Here $T^2$ and $T^3$ are two of the standard adjoint generators of three dimensional rotations, explicitly in terms of the permutation tensor we have

$$(T^i)_{jk} = \epsilon^{ijk}.$$ 

From (6), (7) we can compute $\kappa_i$ and $\tau_i$ as the bond angles and the torsion angles in terms of the PDB data for $r_i$. Alternatively, if we know these angles we can compute the coordinates $r_i$ up to global rotations and translations. The common convention is to select the range of these angles so that $\kappa_i$ is non-negative. In the continuum limit where (7) becomes the standard Frenet equation for a continuous curve, $\kappa_i \to \kappa(x)$ then corresponds to local curvature which is by convention defined to be non-negative.

For 1M6C we take $i$ to take values $i = 3, \ldots, 149 = N$; We leave out three (four) sites at both end as we need three sites to initiate the computation of the $\kappa_i$ and $\tau_i$ along the polygon, and the end points are anyway presumed to be subject to relatively large conformational fluctuations. In Figure 1 (top) we display the $\kappa_i$ and $\tau_i$ along the myoglobin backbone, using the standard differential geometric convention that $\kappa_i$ is non-negative. This Figure displays the geometric structure of the 1M6C backbone fold: At the location of the $\alpha$ and 3/10 helices both $\kappa_i$ and $\tau_i$ have pretty constant values, as expected for helical geometry. The difference between these two types of helices is visible in the Figure, in (slight) difference in the corresponding constant values of $\kappa_i$ and $\tau_i$. At the location of loops, we note small variations in $\kappa_i$ while the values of $\tau_i$ are fluctuating quite wildly. In order to identify the locations of the inflection points that determine the center of the loops i.e. solitons, we follow [3] and subject the data in Figure 1 (top) to local $\mathbb{Z}_2$ gauge transformations in the loop regions; these transformations leave the solution of (7) intact and thus have no effect on the geometry of the space polygon. The result is shown in Figure 1 (bottom): the two data point sets in the top and bottom of Figure 1 describe the same space polygon. But from the bottom Figure 1 we conclude that in terms of $\kappa_i$ we may interpret the backbone as a space polygon with eleven helices that are separated by ten inflection points (soliton centers), these are the points where $\kappa_i$ changes its sign. Consequently we divide the backbone into ten super-secondary structures, each consisting of a helix-loop-helix soliton motif. These motifs are identified in Table I.

We note that PDB lists 1M6C as an eight-helix protein. But Figure 1 reveals that there is an advantage to interpret it in terms of a curve with ten inflection points, so that for a match with the functional form (4) we need to introduce ten overlapping segments. Furthermore, an examination of the PDB data reveals that there are four different types

![Figure 1](image_url)

**FIG. 1:** The values of $\kappa_i$ and $\tau_i$ for 1M6C, obtained from PDB. In the top picture we present these values using the standard convention that $\kappa_i$ is non-negative. In the bottom picture we have resolved the soliton structure using $\mathbb{Z}_2$ gauge structure of the Frenet equation, by allowing $\kappa_i$ to change sign whenever there is an inflection point. This identifies the soliton structures (loops) along the backbone. The indexing refers to the position of amino acids along the backbone, counting from the N-terminus.
of loops \textit{i.e.} solitons: Those that connect two \( \alpha \) helices, those that connect an \( \alpha \)-helix with a 3/10-helix or \textit{vice versa}, and finally those that connect two 3/10-helices.

In order to describe a motif consisting of a loop together with the two similar types of helices that it connects, we use the Ansatz (4) with the symmetric \((a = b)\) relation for the two parameters in (4). But for motifs where a loop connects different types of helices \((\alpha \text{ with } 3/10)\) we allow these parameters to be independent, reflecting the difference in the helices. Thus our Ansatz for the entire backbone is the modification (5) of the Ansatz introduced in [3]. For the bond angles we introduce the dark soliton profile

\[
\kappa_i = (-1)^{r+1} \frac{m_{r1} \cdot e^{c_r(i-s_r)} - m_{r2} \cdot e^{-c_r(i-s_r)}}{2 \cosh[c_r(i-s_r)]}
\]

and we obtain the torsion angles from this soliton profile using the relation

\[
\tau_i = -\frac{1}{2} \frac{b_r}{1 + d_r \kappa_i^2}
\]

Here \( r = 1, ..., 10 \) labels the ten helix-loop-helix motifs of 1M6C and \((c_r, m_{r1}, m_{r2}, s_r)\) are the canonical parameters for a kink-soliton, and \((b_r, d_r)\) are additional parameters needed to express \( \tau_i \) in terms of \( \kappa_i \).

Note that in [3] we have simplified the Ansatz of [3] for torsion angles. Now there is no contribution from \( \kappa_i \) in the numerator, thus there is one less parameter. The reason for this simplification is, that (8), (9) is not an \textit{ad hoc} Ansatz but can be firmly justified in terms of the equations of motion in an underlying Hamiltonian model which is based on the Abelian Higgs Model [10]. The additional term used in [3] version of (9) does not have any natural interpretation in terms of the Abelian Higgs Model and thus there is no geometric reason for including it. Here we confirm that it can be safely removed, with no adverse effect in accuracy. In fact, despite the additional increased complexity in protein structure that we consider, the accuracy reported here is even better than that in [3].

We also emphasize that the parameters are all \textit{global} parameters that are specific to a given helix-loop-helix motif and as such have no direct reference to the amino acids even though they should eventually become computable from an atomic level set-up. At the level of the Abelian Higgs Model [10] each of the parameters has a well established interpretation in terms of charge, mass, self-coupling \textit{etc}. Here, these parameters characterize the global attributes such as the location and the size of the soliton-loop in terms of \( \kappa_i \) and \( \tau_i \), the nature of the adjacent helices, and the chirality of the protein. Moreover, since all the solitons except 2 and 5 connect similar helices, whenever \( r \neq 2, 5 \) we can set \( m_{r1} = m_{r2} \) while for solitons number 2 and 5 that connect two different kind of helices we choose \( m_{r1} \neq m_{r2} \). We also emphasize that the Ansatz involves only the \textit{single} function (4), in its discrete form. This means that for each helix-loop-helix superstructure we only need to determine the five (or six in case the helices are different) global parameters. In our computations we determined these parameters using a standard Metropolis algorithm in combination with simulated annealing, to minimize the RMSD between the polygon described by our Ansatz and the \( C_\alpha \) backbone of the 1M6C protein in PDB. The actual algorithm is a very simple and straightforward application of standard Monte Carlo minimization that runs with PC.

In Table I we display the parameters that yield the smallest RMSD value \((\text{RMSD} = 1.27 \text{ Å})\) that we have obtained when we have subjected the entire 1M6C backbone to a RMSD minimization. We also give the lowest RMSD values that we have obtained when we have separately optimized the parameters for each of the individual soliton. For the solitons 1, 2, 3, 4, 5, 9 and 10 we find very low RMSD values, clearly smaller than the radius \((\sim 0.7 \text{ Å})\) of an individual carbon atom. However, the number of sites that appear in the solitons 3, 4, 5 are also quite small. This is due to the proximity of the ensuing solitons along the backbone. For solitons number 6, 7, 8 the RMSD values are somewhat larger, but the solitons are also longer. However, even in these cases our RMSD values are clearly below the overall 1.90 Å resolution in the underlying PDB data. In Figure 2 we display the \( C_\alpha \) backbone of 1M6C, together with its reconstruction in terms of the Ansatz [5, 6].

We remind that even though our description involves five (six) free parameters for each helix-loop-helix motif, there is only one \textit{single} function, the kink-soliton (5). These parameters can in principle be determined from a first principle atomary level approach to protein folding, even though in practice this is not yet possible. For this we recall [3] that as such, (8), (9) is an approximate solution to a definite discrete nonlinear equation of motion in a Hamiltonian system that provides an effective description of a more fundamental atomary level model.

### III. LONG LOOPS

The previous interpretation and construction of the myoglobin 1M6C backbone clearly demonstrates that the method proposed in [3] can be extended from helix-loop-helix super-secondary structures to entire proteins, even for relatively long proteins and with several helix-loop-helix combinations and both \( \alpha \) and 3/10 helices. However,
The solitons have some overlap with their nearest neighbors, to enable us to combine them into a single multi-soliton profile. The type identifies whether the soliton consists of a loop that connects $\alpha$-helices and (or) $3/10$-helices.

The question remains whether the quality of the method becomes adversely affected if the loop length increases, and whether the method also describes loops that connect an $\alpha$-helix with a $\beta$-strand. We address these issues by considering a protein loop with 12 $C_\alpha$-carbons connecting an $\alpha$-helix with a $\beta$-strand. More specifically, we consider the sites 398-416 in the HIV-1 reverse transcriptase protein with PDB code 3DLK. In line with the construction of the solitons in the case of myoglobin, we describe the super-secondary structure with the following variant \(^{(5)}\) of the Ansatz \(^{(8)}, (9),\)

$$\kappa_i = \frac{m_1 \cdot e^{c_1(i-s_0)} - m_2 \cdot e^{-c_2(i-s_0)}}{e^{c_1(i-s_0)} + e^{-c_2(i-s_0)}}$$  \hspace{1cm} (10)

and we again obtain the torsion angles from this soliton profile using the relation

$$\tau_i = -\frac{1}{2} \frac{b}{1 + d\kappa_i^2}$$  \hspace{1cm} (11)
The asymmetric choice \((m_1, c_1) \text{ vs. } (m_2, c_2)\) reflects the difference between the \(\alpha\)-helix and \(\beta\)-strand, and we now start the indexing by choosing \(i = 1\) for the site 398. With the choice of parameters in Table II we find that the Ansatz describes the 3DLK segment with a RMSD accuracy of 1.13 Å; Notice that due to the presence of exponentials, for high accuracy it is imperative to include sufficiently many decimal points in the parameters. In Figure 3 we display the original 3DLK segment, together with its soliton approximation.

| \(m_1\) | \(c_1\) | \(m_2\) | \(c_2\) | \(s_0\) | \(b\) | \(d\) |
|---------|---------|---------|---------|--------|------|-----|
| 57.626008 | 1.836469 | 58.05348 | 1.8462217 | 10.43150 | 0601165.9 | -0.000101 |

We conclude, that the present approach is suitable not only for long protein chains such as myoglobin, but it also describes long loops and loops that connect very different kind of helices such as \(\alpha\) helices, \(3/10\) helices and \(\beta\) strands. However, if the loop length increases substantially, we propose that a more accurate prescription is obtained by describing these loops as bound states of several short loops, each with the profile \([10], [11]\). This is consistent with the well known fact that short supersecondary structures are known to recur many times in PDB proteins. A detailed analysis of long loops as bound states of short loops (multi-soliton states) will be presented elsewhere.

**IV. CONCLUSION**

Using the myoglobin 1M6C as an example, we have demonstrated that the entire native fold of a long protein can be described with high accuracy as a combination of kink-solitons, in a manner that involves only one single elementary function. In this picture, each of the solitons describe a loop configuration that interpolates between two different helices. By inspecting a longer loop that connects an \(\alpha\)-helix with a \(\beta\)-strand we have verified, that the approach remains valid with no loss of accuracy as the loop size increases. However, for substantially longer loops, we expect that an interpretation in terms of a multi-soliton configuration becomes more accurate both mathematically and phenomenologically. The parameters that characterize a particular protein fold are all global, and specific to its supersecondary helix-loop-helix motifs. Consequently the determination of these parameters becomes synonymous to a quantitative classification of proteins. The presence of an underlying Hamiltonian interpretation at the level...
of motifs also strongly suggests that our approach could eventually provide a bridge between comparative protein classification schemes such as CATH and SCOP, and the atomary level physics based approaches to protein folding and structure prediction, including folding pathways and various other dynamical issues that presently can not be easily addressed in qualitative protein classification schemes. This should open doors to new ways of performing evolutionary, energetic and modelling studies.

Acknowledgement

We thank D. van der Spoel and R. Lavery for discussions and comments. This research has been supported by the Vetenskapsrådet grant number 2009-4099.

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