

Neuroglobin dynamics observed with ultrafast 2D-IR vibrational echo spectroscopy

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Neuroglobin (Ngb), a protein in the globin family, is found in vertebrate brains. It binds oxygen reversibly. Compared with myoglobin (Mb), the amino acid sequence has limited similarity, but key residues around the heme and the classical globin fold are conserved in Ngb. The CO adduct of Ngb displays two CO absorption bands in the IR spectrum, referred to as N₃ (distal histidine in the pocket) and N₀ (distal histidine swung out of the pocket), which have absorption spectra that are almost identical with the Mb mutants L29F and H64V, respectively. The Mb mutants mimic the heme pocket structures of the corresponding Ngb conformers. The equilibrium protein dynamics for the CO adduct of Ngb are investigated by using ultrafast 2D-IR vibrational echo spectroscopy by observing the CO vibration's spectral diffusion (2D-IR spectra time dependence) and comparing the results with those for the Mb mutants. Although the heme pocket structure and the CO FTIR peak positions of Ngb are similar to those of the mutant Mb proteins, the 2D-IR results demonstrate that the fast structural fluctuations of Ngb are significantly slower than those of the mutant Mbs. The results may also provide some insights into the nature of the energy landscape in the vicinity of the folded protein free energy minimum.

myoglobin mutants | protein dynamics | energy landscape

Neuroglobin (Ngb) is a recently discovered family of vertebrate globin proteins. Ngb is expressed predominantly in nerve tissue. It has been hypothesized that Ngb facilitates O₂ diffusion to protect neuronal cells from hypoxia and ischemia (1). Ngb concentration in the brain is fairly low, too low to play the role that myoglobin (Mb) plays in the red muscles. However, because the expression level of Ngb is enhanced under hypoxic conditions *in vitro* as well as ischemia *in vivo* (2, 3), Ngb may be involved in neuronal responses to ischemia. A much higher concentration of Ngb is found in the retina. The concentration is high enough to deliver O₂ to mitochondria (4) and may facilitate O₂ diffusion in a manner similar to Mb. Another possibility is that Ngb is involved in redox-coupled sensor regulation for signal transduction in the brain. Ngb binds to the GDP-bound form of the α -subunit of heterotrimeric G protein under oxidative stress (5).

The 3D structures of Ngb from human and mouse, both having 151 amino acids, have been published (6, 7). Comparisons of Ngb with vertebrate Mb and Hb sequences show only minor similarities at the amino acid level (1), but Ngb features the conserved classical globin fold and contains heme (6). The heme iron in both the ferrous and ferric forms of Ngb is hexacoordinated (8), in contrast to mammalian Mb and Hb, which contain pentacoordinated heme iron. Although hexacoordinated heme has been reported in plant, bacteria, and invertebrate globins, its physiological significance is not yet understood (9). In Ngb, an external gaseous ligand must compete with the sixth ligand, the distal histidine (E7 in helix notation), for binding. Several key residues in the distal pocket for Mb, such as His-64 (E7), Val-68 (E11), and Phe-42 (CD1), are conserved in Ngb, although Leu-29 (B10) is replaced by Phe in Ngb (see Fig. 1). Phenylal-

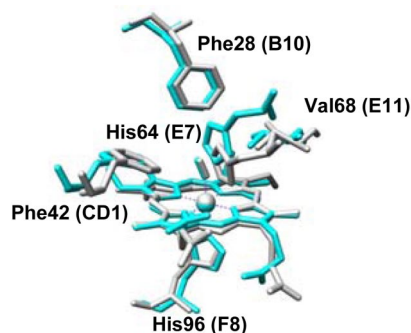


Fig. 1. Crystal structure of the active site of CO-bound ferrous mouse Ngb (gray) and L29F Mb (blue) mutant protein taken from the Protein Data Bank. The heme and some selected amino acid residues are shown.

anine is found at position B10 in many nonsymbiotic plant Hbs that have hexacoordinated binding to the heme (9). Previous studies revealed that Phe at position B10 influences the distal histidine and facilitates stable oxygen binding (10). Another feature of Ngb is that the structural analysis has shown the presence in human Ngb of an intramolecular disulfide bond, which affects its oxygen affinity (11).

Here we report measurements of the fast equilibrium structural dynamics of Ngb and comparisons of the Ngb dynamics with that of Mb mutant proteins. The Mb mutants mimic the structure of the active site of Ngb. The measurements are made by using 2D-IR vibrational echo spectroscopy, which is an ultrafast IR analog of 2D-NMR but operates on time scales many orders of magnitude shorter than NMR. 2D-IR vibrational echo spectroscopy can directly characterize protein motions on time scales from subpicosecond to 100 ps or longer (12–22).

A folded protein at equilibrium resides in a state that is in a free energy local minimum. A protein can have several such local minima, substates, separated by relatively high barriers. The substates are distinguished by differences in some well defined aspects of the protein's structure. Ngb and Mb are examples of proteins with several substates. As discussed below, the substates differ by the configuration of the distal histidine (19, 23–25). One way to view the local minimum of a substate is that it is relatively

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Abbreviations: CLS, center line slope; FFCF, frequency–frequency correlation function; Mb, myoglobin; Ngb, neuroglobin.

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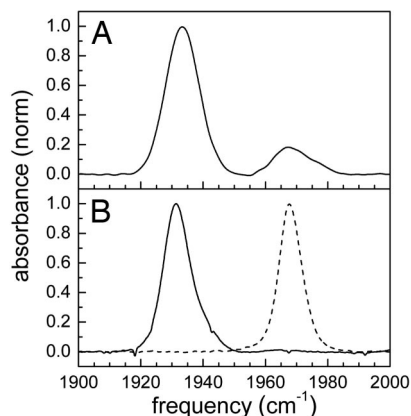


Fig. 2. Normalized FTIR spectra of the CO stretching mode bound to Ngb (A) and Mb mutant proteins L29F (solid curve) and H64V (dashed curve) (B).

broad with a rough energy landscape (26). This landscape has numerous minima separated by relatively low barriers. Changes in protein structure are caused by transitions from one landscape minimum to another. A substantial fraction or all of these transitions occur on subpicosecond to 100-ps time scales, depending on the protein. These structural fluctuations can be observed by using 2D-IR vibrational echo spectroscopy.

For heme proteins, 2D-IR vibrational echo experiments use the heme-ligated CO vibration as a direct sensor of protein dynamics (15, 16). The IR linear absorption spectrum of the CO stretching mode of heme proteins generally displays several bands. The different bands reflect structural differences, i.e., distinct structural substates, and the width of the bands reflects the range of CO vibrational energies that are associated with the distribution of protein structural configurations of each substate (15, 23, 24). The linear IR absorption spectrum cannot provide information on a protein's structural dynamics. The dynamical information is obtained from the time evolution of the 2D-IR line shapes that are acquired with the vibrational echo experiments (16).

The peak positions of the stretching bands of CO bound to Ngb in the absorption spectrum are almost identical with those of particular Mb mutant proteins. However, it will be shown below that, despite the great similarity in the absorption spectra, heme pocket structures, and backbone folds, the structural dynamics of Ngb and Mb are distinct. Ligand rebinding studies of Ngb show that CO rebinding kinetics after optical excitation-induced dissociation of the CO from the heme is different from that of Mb (27, 28). The results presented below demonstrate that the equilibrium structural fluctuations of Ngb are considerably slower than those of Mb. The results also provide insights into the nature of the energy landscape near the folded protein's free energy minimum.

Results and Discussion

Linear FTIR Spectroscopy. The background-subtracted linear FTIR spectrum of CO bound to human Ngb is shown in Fig. 2A. There are two CO absorption bands at $1,932\text{ cm}^{-1}$ and $1,968\text{ cm}^{-1}$. These bands have been called the N_3 (lower frequency) and N_0 (higher frequency) conformers, respectively, in analogy to the A_3 and A_0 bands of Mb (27, 29). The significant width of the CO stretching bands of Ngb and their Gaussian shapes imply structural heterogeneity associated with each protein substate that is sensed by the CO bound to the active site. Although the range of frequencies that make up a substate's absorption band is sensitive to the details of the heme pocket, it is also determined by the global structural variations of the protein (19, 20, 30, 31). The frequency of N_3 corresponds closely to the MbCO A_3

conformation, which has the distal histidine localized in the heme pocket (19), whereas the frequency of N_0 is almost identical with the MbCO A_0 conformer, which has the distal histidine rotated out of the pocket (25). Substitution in Ngb of the distal histidine by the nonpolar amino acid residues, valine or alanine, causes a substantial increase in the population of N_0 conformer (27).

The A_1 substate of wild-type MbCO is predominately populated and dominates the absorption spectrum. It does not correspond to either the N_3 or N_0 states of Ngb in vibrational frequency. The A_1 and A_3 bands of MbCO arise from different conformations of the distal histidine in the heme pocket (19, 20). The A_3 conformer places the hydrogen on the protonated ϵ -nitrogen of the distal histidine, much closer to the CO than the A_1 conformer (19). Because of the similarities of the N_3 and A_3 spectra and the N_0 and A_0 spectra, it is desirable to compare their dynamics by using 2D-IR vibrational echo spectroscopy. However, both the A_3 and A_0 bands are too small in wild-type MbCO to be useful for the 2D-IR experiments.

To compare the dynamics of Ngb and Mb, the mutant Mb proteins L29F and H64V were used. The dynamics of MbCO and MbCO mutants have been measured by using vibrational echo spectroscopy (see refs. 16, 18–21, 32–34). The background-subtracted normalized FTIR spectra of CO bound to the mutants are shown in Fig. 2B. The L29F mutant has one major band at $1,932\text{ cm}^{-1}$ corresponding closely to the N_3 spectrum of Ngb. Leu-29 (B10) in Mb is substituted by Phe in Ngb, although other key residues in the distal pocket of Mb are preserved in Ngb. Therefore, the L29F mutant mimics the heme pocket structure of Ngb (Fig. 1). Experiments, theory, and molecular dynamics simulations have shown that both the local and global coupling between protein structure and the CO vibrational transition frequency is mainly through the electric field along the CO bond (19, 20, 24, 29, 30, 35–37). The protein residues range from charged, to polar, to nonpolar. All of these, as well as, to some extent, the solvent, contribute to the electric field. The vibrational states couple to the electric field via the Stark effect (37). Differences in structure produce changes in the electric field at the CO and, therefore, changes in frequency. The range of frequencies associated with the inhomogeneous line widths of the CO bands is determined by the range of structures that produce the electric field at the CO. 2D-IR vibrational echoes monitor the interconversion of the structures via the time evolution of the CO frequency.

The spectrum of the Mb A_0 band has a frequency almost the same as the Mb mutant, H64V, which has the distal histidine replaced by a valine. H64V mimics the situation in which the distal histidine is rotated out of the heme pocket. Both H64V and the N_0 conformation for Ngb absorb at $1,968\text{ cm}^{-1}$. The spectral positions and line widths of the Ngb bands and the mutant bands are given in Table 1.

2D-IR Spectroscopy. Ultrafast 2D-IR vibrational echo spectroscopy can measure fast protein dynamics under thermal equilibrium conditions through the changes in the 2D-IR line shapes with time (13, 15–17, 19). In the vibrational echo experiments, three IR excitation pulses, ≈ 110 fs in duration, tuned to the vibrational absorption frequency are used. The times between pulses 1 and 2 and pulses 2 and 3 are called τ and T_w , respectively. The vibrational echo pulse emerges from the sample in a unique direction at a time $\leq \tau$ after the third pulse. A 2D spectrum is recorded by scanning τ at fixed T_w . The vibrational echo pulse is heterodyne detected through a monochromator. Heterodyne detection provides phase information in addition to signal amplitudes. Taking the spectrum of the heterodyned echo pulse performs one of the two Fourier transforms and gives the ω_m -axis (m for monochromator) in the 2D frequency domain spectrum. When τ is scanned, time interferograms are generated

Table 1. Parameters for CO bound to Ngb and Mb and mutants

Protein	Line center, cm^{-1}	FWHM, cm^{-1}	T_1 , ps
Ngb N ₃	1,932.7	12.1	19.3
Ngb N ₀	1,968.1	9.8	18.4
Mb L29F	1,931.8	9.6	15.8
Mb H64V	1,968.0	8.8	24.1

at each ω_m . The interferograms are numerically Fourier transformed to give the ω_τ -axis of the 2D spectrum. Two-dimensional vibrational echo spectra are recorded as a function of T_w .

As T_w increases, the shape of the 2D spectral bands change. These changes are directly related to the structural evolution of the protein through the influence of the structural changes on the frequency of the CO vibrational mode. The experiment can be viewed qualitatively as follows. The first and second pulses act to label the initial frequencies of the molecules. Between the second and third pulses, structural evolution of the proteins causes the initially labeled frequencies to change (spectral diffusion). The third pulse ends the evolution period, and the vibrational echo pulse reads out the final frequencies. As T_w increases, there is more time for structural evolution and, therefore, larger frequency changes. The frequency changes are reflected in the change in shapes of the 2D-IR vibrational echo spectra. Full descriptions of the method have been presented (16, 38–40).

Fig. 3*A* shows 2D-IR spectra of CO bound to Ngb at several values of T_w . The red bands are positive going and correspond to the 0–1 vibrational transition. The blue bands are negative going. They arise from vibrational echo emission at the frequency of the 1–2 vibration transition (41) and are shifted along the ω_m -axis by the CO stretching mode's anharmonicity. As T_w increases, all of the peaks decay at a rate determined by the

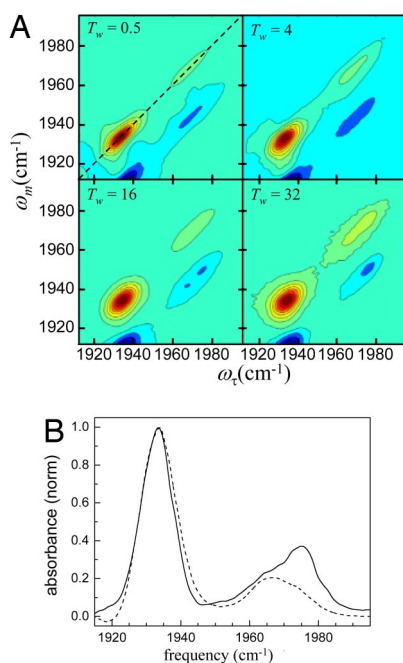


Fig. 3. 2D-IR vibrational echo spectra of NgbCO. (A) 2D-IR spectra at various times, T_w . Each contour corresponds to a 10% signal change. The red bands (positive going) correspond to the 0–1 vibrational transition. The blue bands (negative going) arise from vibrational echo emission at the 1–2 transition frequency. The dashed line in *Upper Left* is the diagonal. (B) Diagonal slice spectra of CO-bound Ngb at $T_w = 0.25$ ps (dashed curve) and 50 ps (solid curve).

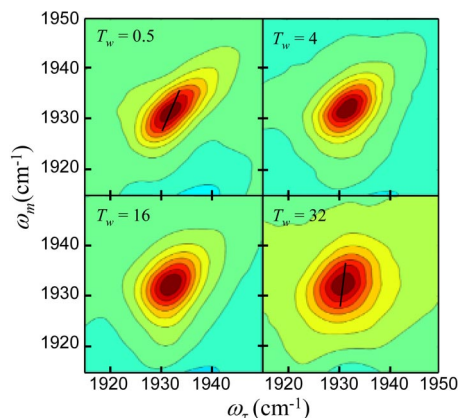


Fig. 4. 2D-IR spectra of CO bound to L29F Mb mutant protein. Each contour corresponds to a 10% signal change. The dashed lines are the center lines.

vibrational lifetime T_1 . In Fig. 3*A*, the peaks are normalized to the largest peak in each panel. Therefore, the decay of the vibrationally excited population manifests itself as a relative reduction in the amplitude compared with the largest peak. The dashed line from upper right to lower left in Fig. 3*A Upper Left* is the diagonal. The two bands on the diagonal correspond to the 0–1 transitions of the two peaks in the absorption spectrum shown in Fig. 2*A*. These two bands are located at $(\omega_\tau, \omega_m) = (1,932 \text{ cm}^{-1}, 1,932 \text{ cm}^{-1})$ and $(\omega_\tau, \omega_m) = (1,968 \text{ cm}^{-1}, 1,968 \text{ cm}^{-1})$ for the N₃ and N₀ conformers, respectively.

Fig. 3*A* illustrates two important features. As T_w increases, the bands go from highly elongated along the diagonal to less elongated and increasingly broad along the ω_τ -axis. In the long time limit, they would become round. The change in shape is a manifestation of spectral diffusion caused by protein structural evolution. Detailed analysis presented below of the time dependence of the shapes quantifies the time evolution of the protein structures.

As T_w increases, the shape of the N₃ changes but the peak position remains fixed. However, not only does the N₀ band change shape, but the center frequency also changes, as can be seen in Fig. 3*B*, which displays two diagonal cuts through the 2D spectrum. The dotted-dashed curve is the cut at $T_w = 0.25$ ps, and the solid curve is the cut at $T_w = 50$ ps. The spectra have been normalized to the N₃ band maximum. The N₀ line clearly shifts. The shift occurs because the N₀ line is produced by two distinct protein conformers with overlapping spectra and different vibrational lifetimes. The lower-frequency conformation in the N₀ band has a shorter CO lifetime than the higher-frequency band, so at long time, the high-frequency band is revealed. In Fig. 2*A*, the N₀ absorption is somewhat elongated to the high-frequency side, suggesting that there may be two peaks. In Fig. 3*B* it is clear that there are two peaks.

To compare the dynamics of Ngb and Mb, the 2D-IR spectra of the Mb mutants, L29F and H64V, were also measured. Fig. 4 shows 2D-IR spectra of the CO stretch for the L29F mutant at several T_w points. Only the 0–1 transition region is shown. The band is located on the diagonal at $(\omega_\tau, \omega_m) = (1,933 \text{ cm}^{-1}, 1,933 \text{ cm}^{-1})$ for L29F and at $(\omega_\tau, \omega_m) = (1,968 \text{ cm}^{-1}, 1,968 \text{ cm}^{-1})$ for H64V (not shown). The peak positions of the Mb mutants are time-independent. The structural evolution of L29F is manifested by the spectral-diffusion-induced change in shape of the 2D-IR band from elongated to close to round as T_w increases.

The changes in the shapes of the 2D bands with increasing T_w can be used to determine the time scales and amplitudes of various contributions to the fast structural dynamics of the proteins by using methods based on diagrammatic perturbation theory (42). The frequency–frequency correlation function

(FFCF) connects the experimental observables to the underlying dynamics. The FFCF is the probability that a vibrational oscillator with a given initial frequency still has the same frequency at time t later, averaged over all starting frequencies. Once the FFCF is known, all linear and nonlinear optical experimental observables can be calculated by using the time-dependent diagrammatic perturbation theory (42). Conversely, the FFCF can be extracted from 2D vibrational echo spectra with additional input from the linear absorption spectrum. In general, to determine the FFCF from 2D and linear spectra, full calculations of linear and nonlinear response functions are performed iteratively until the calculated results converge to the experimental results (40, 43, 44).

Here the center line slope (CLS) method, a new approach for extracting the FFCF from the T_w dependence of the 2D spectra, is used (39, 45). It is accurate and much simpler numerically to implement than the full diagrammatic perturbation theory. Furthermore, it provides a more useful quantity to plot for visualizing differences in the dynamics of the various proteins than a series of 2D line shapes (45). A slice through the 2D spectrum at a particular ω_m is a spectrum along the ω_r -axis. The peak of this spectrum is at a particular ω_r -value. So the peak is a point with ω_m , ω_r -coordinates. Slices are taken over a range of ω_m values, and the resulting set of points forms the center line. Two such center lines are shown in Fig. 4 for $T_w = 0.5$ ps and 32 ps. In the absence of a homogeneous contribution to the spectrum (see below), at $T_w = 0$ the 2D line shape would be a line along the diagonal at 45°. Then, the slope of the center line would be 1. At very long time, the 2D line shape becomes symmetrical, and the center line is vertical (the slope is infinite). It has been shown theoretically that the T_w -dependent part of the FFCF equals the T_w dependence of the inverse of the slope of the center line (45), which is referred to as the CLS. Therefore, the CLS will vary from a maximum of 1 at $T_w = 0$ to a minimum of 0 at sufficiently long time. In Fig. 4, it can be seen that for $T_w = 32$ ps the slope of the center line is steeper than at $T_w = 0.5$ ps. It has also been shown theoretically that combining the analysis of the CLS with the linear absorption spectrum, the full FFCF can be obtained, including the T_w -independent homogeneous component (45).

A homogeneous contribution to the 2D line shape and the linear absorption line shape can arise from three sources: very fast structural fluctuations that produce a motionally narrowed contribution (16, 43, 44, 46); the vibration lifetime, T_1 ; and orientational relaxation. Because the proteins are so large, the orientational relaxation contribution is negligible. The T_1 contribution was measured independently by using IR pump-probe experiments. The T_1 contribution to the homogeneous line width ($1/\pi 2T_1$) is small. It varies for the different bands but is ≈ 0.3 cm^{-1} . The T_1 values are listed in Table 1. The homogeneous contribution mainly comes from the motionally narrowed component. Motional narrowing occurs when some portion of the structural fluctuations are extremely fast, such that $\Delta\tau < 1$, where τ is the time scale of the fluctuations and Δ is the range (amplitude) of the frequency fluctuations. Motionally narrowed fluctuations produce a Lorentzian contribution to both the linear absorption spectrum and the 2D-IR spectrum. The T_w -independent homogeneous contribution manifests itself by broadening the 2D spectrum along the ω_r -axis even at $T_w = 0$. This broadening reduces the initial value of the CLS to a number < 1 , which permits the determination of the homogeneous component (45).

A multiexponential form of the FFCF, $C(t)$, was used to model the multi-time scale dynamics of the protein structural fluctuations. This form has been widely used, and in particular, it has been found applicable in studies of the structural dynamics of heme-CO proteins (see refs. 15, 16, 18–20, 31–33, 47). The FFCF has the form

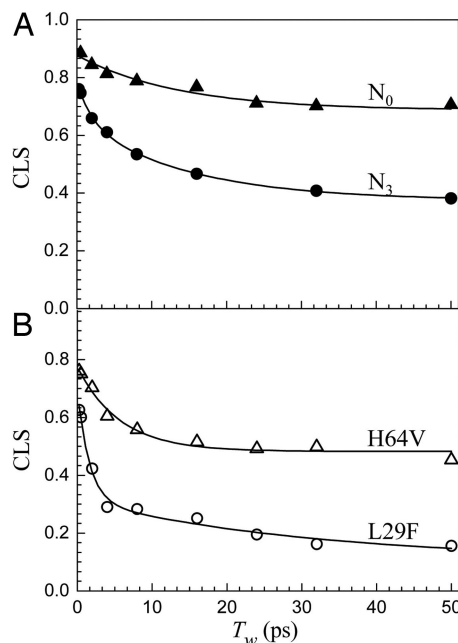


Fig. 5. T_w -dependent CLS for N_0 conformer of Ngb (A, filled triangles) and N_3 conformer of Ngb (A, filled circles) and for Mb mutant proteins H64V (B, open triangles) and L29F (B, open circles).

$$C(t) = \sum_{i=1}^n \Delta_i^2 e^{-t/\tau_i} + \Delta_s^2 \quad [1]$$

The Δ_i and τ_i terms are the amplitudes and correlation times, respectively, of the frequency fluctuations induced by protein structural dynamics. τ_i reflects the time scale of a set of structural fluctuations, and Δ_i is the range of CO frequencies sampled because of the structural fluctuations. The experimental time window is $\approx 5T_1$, because lifetime decay reduces the signal to zero. The 2D-IR vibrational echo experiment is sensitive to fluctuations a few times longer than this window (48), i.e., several hundred picoseconds, because some portion of slower fluctuations will occur in the experimental window if their time scale is not too slow. Protein structural dynamics that are sufficiently slow will appear as static inhomogeneous broadening, which is reflected in $C(t)$ by Δ_s , a static term. In obtaining the FFCF from the data, the Δ_i and the τ_i are determined. However, for a motionally narrowed term ($\Delta\tau < 1$), only the product, $\Delta^2\tau = 1/T_2^*$, can be obtained. T_2^* is called the pure dephasing time. The pure dephasing time is determined by using $1/T_2 = 1/T_2^* + 1/T_2$, where T_2 is determined from the CLS with use of the linear absorption spectrum (45) and T_1 is obtained from IR pump-probe experiments (see Table 1).

The T_w -dependent CLS for the N_3 , N_0 , L29F, and H64V bands are presented in Fig. 5. The N_0 band arises from two highly overlapped signals as is evident from Fig. 3B. Given the small amplitude of the N_0 signal, within the signal-to-noise ratio we can see no difference in the dynamics for the two underlying substates of the N_0 band. Therefore, we will take the dynamics of the two components of the N_0 line to be the same and present the data for N_0 as if they are a single entity.

Fig. 5A shows the CLS data for the Ngb N_3 and N_0 bands. It is clear from inspection of the data that the dynamics of the two Ngb substates are very different. The FFCF parameters are given in Table 2. The deviation from the maximum possible CLS value at $T_w = 0$ is indicative of a motionally narrowed term. T_2^* is more than a factor of 2 faster for N_3 than N_0 . The N_0 band has an

Table 2. FFCF parameters obtained from 2D-IR and absorption spectra

Protein	T_2^* , ps	Δ_1 , cm^{-1}	τ_1 , ps	Δ_2 , cm^{-1}	τ_2 , ps	Δ_s , cm^{-1}
Ngb N_3	5.0	1.9	2.0	2.7	14	3.1
Ngb N_0	11.9	1.8	11.5			3.5
Mb L29F	4.8	2.8	1.7	2.2	66	
Mb H64V	7.7	2.1	5.2			2.7

intermediate decay, 11.5 ps, followed by a static component. As discussed above, the static component shows that there are protein structural fluctuations occurring on time scales longer than several hundred picoseconds, which are outside of the experimental window. In contrast, the N_3 line has a fast (2 ps) decay followed by another, intermediate decay of 14 ps, and then a static term. Although the intermediate decay times are similar for the two Ngb substates, N_3 has a fast-decay component, reflecting fast structural fluctuations, which is not present for the N_0 substate.

The CLS data for the Mb mutants L29F and H64V are displayed in Fig. 5B. The difference between the CLS decays of L29F and H64V is qualitatively similar to the difference between the N_3 and N_0 decays shown in Fig. 5A. The decay of L29F is significantly faster than that of H64V in the same manner that the decay of N_3 is faster than the decay of N_0 . The fundamental difference between L29F and H64V is the distal histidine. L29F has the distal histidine in the heme pocket. Its spectrum is almost identical with the spectrum of the A_3 band of wild-type Mb. By combining vibrational echo experiments and molecular dynamics simulations, it has been shown that the A_3 configuration of Mb has the protonated ϵ -nitrogen of the imidazole side group of the distal histidine directed toward the CO (19). The conformation of the distal histidine results in a strong interaction between it and the CO bound at the active site. In contrast, H64V has the distal histidine replaced with a nonpolar valine, almost eliminating the interaction between residue 64 and the active site. Vibrational echoes and molecular dynamics simulations have shown that the reduction of CO vibrational spectral diffusion in H64V compared with the A_3 (and A_1) band of Mb is caused mainly by the elimination of the distal histidine (32). In N_3 the distal histidine is in the heme pocket but in N_0 it is swung out of the pocket. The IR CO vibrational absorption spectra of the N_0 and H64V bands, as well as the A_0 band of Mb, are almost identical. Elimination of the distal histidine (H64V) has the same effect on the absorption spectrum as its being swung out of the pocket. Therefore, it is reasonable to ascribe the difference between the N_3 and N_0 spectral diffusion to the difference in the configuration of the distal histidine. As with Mb, the fast component of the dynamics of N_3 can be substantially ascribed to motions of the distal histidine strongly interacting with the active site, which are absent for N_0 .

Although the nature of the relationship of the dynamics of N_3/N_0 to L29F/H64V is the same, the dynamics of Ngb is fundamentally different from the dynamics of Mb. The dynamics of N_3 is substantially slower than that of L29F, and the dynamics of N_0 is significantly slower than that of H64V. The fastest dynamics values of N_3 and L29F are similar. They have essentially identical T_2^* values. L29F has a somewhat faster τ_1 with somewhat greater amplitude, Δ_1 , but overall the fast dynamics values are similar. However, there is a major difference on the slower time scales. N_3 has an intermediate time scale component and a static component of significant amplitude. L29F does not have a static component. Instead, it has a single 66-ps decay. Thus, the equilibrium structural fluctuations of L29F have a slowest component that is relatively fast, only 66 ps. In L29F, all possible structures that give rise to the inhomogeneous broad-

ened CO absorption line are sampled in a few times at 66 ps. Although the heme pockets of N_3 and L29F are similar and they are both globin proteins, their amino acid sequences are substantially different. The 2D-IR vibrational echo experiment is sensitive to both local and global structural fluctuation. The similarities in the heme pockets indicate that the differences in dynamics arise from the different global structural fluctuations of the two proteins.

The dynamics of the N_0 substate of Ngb is also slower than the dynamics of H64V. Both have static components. The major difference between the two is in the moderately fast fluctuations. The τ_1 values for N_0 and H64V are 11.5 ps and 5.2 ps, respectively. Neither of these species has a distal histidine in the heme pocket, supporting the proposition that the differences in dynamics between Ngb and Mb involve differences in global structural fluctuations rather than differences in very local interactions at the active site.

Concluding Remarks

Ultrafast 2D-IR vibrational echo experiments have determined the time scales and relative magnitudes of the fast equilibrium structural fluctuations of the two substates, N_3 and N_0 , of the protein neuroglobin by observing the vibrational dephasing dynamics of the stretching mode of CO bound at the heme active site. It is found that the structural fluctuations of the N_0 substate are much slower than those of the N_3 substate. The heme pocket structures of Ngb and Mb are similar despite the lack of similarity in their amino acid sequences. The linear IR spectra of CO bound at the active sites of the N_3 and N_0 conformers of Ngb are almost identical with the linear IR spectra of L29F and H64V Mb mutant proteins, respectively. N_3 and L29F have a distal histidine in the active site pocket, whereas N_0 and H64V do not. H64V displays substantially slower vibrational spectral diffusion than L29F. Therefore, the reduction in structural fluctuations detected by CO bound at the active site of Ngb in the N_0 substate relative to the N_3 substate is substantially due to the distal histidine being swung out of the heme pocket.

The vibrational echo experiments demonstrate that the equilibrium structural fluctuations of Ngb are significantly different from those of Mb despite the globin structure of the proteins, the near identity of their heme pocket structures, and the almost identical CO FTIR spectra. However, away from the heme pocket, the primary structures of Ngb and Mb are quite different. Ngb has a disulfide bond not present in Mb. Even in the absence of an exogenous bound ligand, Ngb is hexacoordinated with the distal histidine in the sixth ligand, whereas Mb is pentacoordinated. As revealed by the 2D-IR vibrational echo experiments, the net result of all structural differences is that the Ngb structure is substantially more constrained on fast time scales than Mb.

As shown in Fig. 3B, the N_0 band is actually composed of two substates, N_0' and N_0'' , each of which has an absorption line width that is narrower than the total line width reported in Table 1. Thus, the range of structural variations of the N_0' and N_0'' substates is small, and as can be seen in Fig. 5A, there is not a great deal of structural dynamics out to ≈ 100 ps. The reduced structural sampling on fast time scales in Ngb, particularly the N_0' and N_0'' substates, may slow ligand migration through the protein to the active site compared with Mb.

A folded protein in a particular substate occupies a minimum on the free energy landscape (26). However, the minimum is broad and rough with many local minima separated by low barriers. Transitions among these minima give rise to the structural fluctuations that cause the spectral diffusion measured by the 2D-IR vibrational echo experiment. In L29F, all of these local minima are sampled rapidly. Spectral diffusion is complete, and therefore all structures within the broad free energy minimum are sampled, in a few hundred picoseconds. In

contrast, N₃ has a static component in the FFCF that indicates that some of the dynamics occur on time scales that are long compared with several hundreds of picoseconds. The implication is that the rough landscape of N₃ around the folding minimum has higher barriers than L29F.

The FFCF obtained for L29F has three terms (see Table 2). The motionally narrowed term does not permit the determination of the time constant, τ . However, simulations of Mb show that the motionally narrowed τ is tens of femtoseconds (32). With this value, L29F has structural fluctuations on three time scales: tens of femtoseconds, 1.7 ps, and 66 ps. The tens-of-femtosecond dynamics values are too fast for true structural changes. They reflect thermally excited oscillatory motions of small groups that are local and highly damped. The slower time constants are the time scales for true structural changes, such as changes in conformations of side chains. Although other explanations may exist, the fact that L29F has two well separated time constants suggests a two-tier energy landscape near the free energy minimum of the folded state. There is a set of low barriers that are crossed relatively rapidly and a set of higher barriers that are crossed more slowly. For the Ngb substates and H64V, the static terms in the FFCF imply that there is an even higher tier or tiers of barriers that push the time scale for part of the

structural evolution out of the ≈ 100 -ps time window of the experiments.

Materials and Methods

Expression and purification of His₆-tagged human Ngb was performed as described in ref. 5. Purity of protein and contamination of disulfide-dependent formation of dimers were checked by SDS/PAGE under reduced and nonreduced conditions. The mutant sperm whale Mb proteins L29F and H64V were expressed and purified as described in ref. 49.

The CO forms of Ngb and mutant Mb proteins were prepared according to protocols published in refs. 19, 20, and 31. For both the linear FTIR and vibrational echo measurements, $\approx 20 \mu\text{l}$ of the sample solution was placed in a sample cell with CaF₂ windows and a 50- μm Teflon spacer.

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