Study of Chemical Exchange by 2D NMR

Product used : Nuclear Magnetic Resonance (NMR)

The NMR signal of a spin reflects its local magnetic environment. If a spin due to chemical exchange samples two magnetically different states then its NMR signal would reflect both states. Its appearance on a NMR spectrum would be determined by the dynamics of the exchange event. In the case of chemical exchange that is slow on the NMR time scale, it is possible to observe two distinct signals for the same spin, one signal for each state under exchange. Presence of chemical exchange is often demonstrated with the exchange experiment (2D NOESY experiment). A exchange peak (cross peak) of the same sign can be observed between the two autopeaks (diagonal peaks) that represent the two states under exchange. Because the same results can be interpreted in a different way based on NOE, further evidence is desirable. ZZ exchange experiment is a 2D NMR experiment for the quantitative study of slow chemical exchange in proteins. It was originally reported as a $^1$H$^{15}$N HSQC that uses as NMR probe the relatively isolated $^1$H$^{15}$N spin system in the peptide bond of uniformly $^{15}$N labeled proteins. A $^1$H$^{13}$C HSQC version of the above experiment was prepared and tested on a small compound.

ZZ exchange experiment

Because of the partial double bond character of the amide C-N bond in N,N dimethylformamide (DMF), free rotation around C-N is hindered and a slow exchange can be established between the two distinct methyl proton signals (states A and B). Cross peaks (exchange peaks) with the same phase with the diagonal peaks can be observed in 2D NOESY and 2D ROESY experiments (Fig.1).

Fig 1: 2D NOESY (left) and 2D ROESY (right) of DMF in DMSO-d$_6$ at 70°C.

This slow exchange system was further examined with the newly prepared ZZ exchange experiment. Briefly, in the $^1$H$^{13}$C ZZ exchange experiment, following $^{13}$C frequency-labeling during $t_1$ (indirect dimension), magnetization is changed into in phase $^{13}$C Z magnetization and is allowed to exchange. Then magnetization is transferred to proton and labeled with its proton frequency during FID (direct dimension) (Fig.2).

Fig 2: $^1$H$^{13}$C ZZ exchange experiment.
The results with DMF are shown in Fig. 3 for different values of mixing time. Exchange peaks were observed between the two $^1$H-$^{13}$C cross peaks (auto peaks) representing the two exchanging states. Auto peaks represent magnetization that didn’t exchange during mixing time. Exchange peak denoted as B->A in Fig. 3 represents B magnetization at the start of ZZ exchange (labeled with the B carbon frequency during $t_1$) that was exchanged during mixing time and therefore it was transferred to A state (labeled with A proton frequency during $t_2$). Similarly, the A->B exchange peak represents A state carbon magnetization that was transferred to B state and labeled with the B proton frequency.

Reference: J Biomol NMR, 4, 727-734, 1994

It is necessary in the above experiment that the exchange states A and B show different chemical shifts for both proton and carbon. If this is not the case then the exchange peaks will overlap with the autopeaks. In the special case where exchange states show same proton but different carbon chemical shifts, exchange might be demonstrated with a $^{13}$C-$^{13}$C NOESY type of experiment (Fig. 4).

Phase cycling combined with proton decoupling and in phase carbon magnetization during mixing time ensure that there is no NOE contribution to exchange peaks. Therefore, the above experiment can confirm the presence of slow exchange or even provide under favorable conditions, quantitative description of the exchange event.

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