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Measurement of Imino ^1H - ^1H Residual Dipolar Couplings in RNA

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Abstract

Imino ^1H - ^{15}N residual dipolar couplings (RDCs) provide additional structural information that complements standard ^1H - ^1H NOEs leading to improvements in both the local and global structure of RNAs. Here, we report measurement of imino ^1H - ^1H RDCs for the Iron Responsive Element (IRE) RNA and native *E. coli* tRNA^{Val} using a BEST-Jcomp-HMQC2 experiment. ^1H - ^1H RDCs are observed between the imino protons in G-U wobble base pairs and between imino protons on neighboring base pairs in both RNAs. These imino ^1H - ^1H RDCs complement standard ^1H - ^{15}N RDCs because the ^1H - ^1H vectors generally point along the helical axis, roughly perpendicular to ^1H - ^{15}N RDCs. The use of longitudinal relaxation enhancement increased the signal-to-noise of the spectra by ~3.5-fold over the standard experiment. The ability to measure imino ^1H - ^1H RDCs offers a new restraint, which can be used in NMR domain orientation and structural studies of RNAs.

Keywords

Residual Dipolar Couplings; RNA Structure; Partial Alignment; tRNA; Iron Responsive Element RNA

Solution NMR studies of nucleic acids are more challenging than those of similarly sized proteins due to the lower proton density and more extended structures generally found in nucleic acids. As a result, the local structure of nucleic acids can often be well-determined from NOE and J-coupling constant restraints, whereas the global orientations of helical secondary structural elements are usually much less precisely determined (Vermeulen, *et al.*, 2000). Residual dipolar coupling (RDC) data relate inter-nuclear vectors to a common molecular axis system (Bothner-By *et al.* 1981; Kung *et al.* 1995; Tolman *et al.* 1995; Bax *et al.* 2001), and provide long-range global information that complements short-range NOE and J-coupling constants. Thus, RDC constraints obtained under conditions of weak alignment (Tjandra and Bax, 1997) are extremely valuable for NMR solution structure determination of nucleic acids (Hansen *et al.* 2000; Mollova and Pardi 2000; Lukavsky *et al.* 2003; Davis *et al.* 2005).

The vast majority of RDCs routinely measured in biomolecules involve interactions between nuclei at a fixed known distance (Bax *et al.* 2001; Prestegard *et al.* 2004). The most commonly measured RDCs in nucleic acids are the ^1H - ^{15}N iminos of guanine and uracil, the $^1\text{H}_2$ - $^{13}\text{C}_2$ of adenine, the $^1\text{H}_8$ - $^{13}\text{C}_8$ of purines, the $^1\text{H}_6$ - $^{13}\text{C}_6$ and $^1\text{H}_5$ - $^{13}\text{C}_5$ of pyrimidines, and the $^1\text{H}_1$ '- $^{13}\text{C}_1$ ' of the ribose sugar. Experiments have also been developed to measure additional RDCs in nucleic acids, for example between other nuclei separated by one- and two-bonds in the nucleobase and ribose-phosphate backbone (Fiala *et al.* 2004; Latham *et al.*

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2005; Getz et al. 2007). Of all these nuclei, the imino protons generally offer the highest resolution, and thus represent important probes for collecting structural restraints.

Imino ^1H - ^{15}N RDCs have previously been used in domain orientation studies of helical regions in RNAs (Mollova et al. 2000; Al-Hashimi et al. 2002; Bondensgaard et al. 2002; Lukavsky et al. 2003; D'Souza et al. 2004; Vermeulen et al. 2005; Getz et al. 2007; Ying et al. 2007); however, there is generally only one imino ^1H - ^{15}N RDC for each base pair in helical regions leading to few restraints. Thus, ^1H - ^1H RDCs (D_{HH}) between the well-resolved imino protons represent attractive candidates for providing additional structural data. The high proton gyromagnetic ratio can yield large residual dipolar couplings (Hansen et al. 1998b; Lipsitz and Tjandra 2004), so various methods have been developed for measuring D_{HH} values to take advantage of these additional structural constraints (Hansen et al. 1998b; Tian et al. 2000; Wu and Bax 2002; Boisbouvier et al. 2003; Meier et al. 2003). However, a major obstacle in these measurements is the large network of simultaneously dipolar coupled ^1H probes in the biomolecule, making the measurement of D_{HH} values greater than 4 Å apart difficult (Hansen, *et al.*, 1998). Strategies for perdeuteration and/or homonuclear decoupling have therefore been developed to help overcome these problems (Scott et al. 2000; Tian et al. 2000; Wu and Bax 2002; Boisbouvier et al. 2003; Meier et al. 2003). For example, homonuclear decoupling methods have been used to measure D_{HH} values between the non-exchangeable H1' resonances to help improve the accuracy and precision of a DNA duplex structure (Boisbouvier, *et al.*, 2003). Bax and co-workers used a semi-selective pulse sequence to measure ^1H - ^1H RDCs between amide protons in proteins (Wu and Bax, 2002). This so-called SS-HMQC2 pulse sequence consists of two semi-selective heteronuclear multiple quantum coherence (HMQC) blocks surrounding a COSY-type mixing pulse, and the magnitudes of D_{HH} are determined from a quantitative J-correlation type approach (Wu and Bax, 2002). Recently, Schanda et al. (2007) employed modifications to this experiment, which resulted in 1.5 to 8 times higher signal-to-noise for measurement of amide ^1H - ^1H RDCs in proteins. This experiment is referred to as BEST-Jcomp-HMQC2, where BEST stands for: Band-selective-Excitation Short-Transient and Jcomp for: J-mismatch compensated (Schanda, *et al.*, 2007). The higher sensitivity is the result of two improvements: longitudinal relaxation enhancement (Pervushin et al. 2002; Schanda and Brutscher 2005; Schanda et al. 2005) and J-mismatch compensation (Schanda, *et al.*, 2007). Here we report the application of this BEST-Jcomp-HMQC2 technique for measuring D_{HH} interactions between the well-resolved imino protons in ^{15}N -labeled RNAs.

The original 2D ^{15}N - ^1H BEST-Jcomp-HMQC2 experiment (Schanda, *et al.*, 2007) was modified here so proton evolution occurs during the t_1 period allowing for measurement of D_{HH} couplings in a spectrum analogous to a 2D ^1H - ^1H NOESY data set (Supplementary Figure 1). In addition to increasing the sensitivity, the ^1H selective pulses employed by the BEST approach only excite the exchangeable imino protons; whereas, the other protons remain at equilibrium. This ensures that D_{HH} evolve between imino protons and not between imino and neighboring amino protons, which are close together in a base pair (~1.9–2.5 Å). The modified BEST-Jcomp-HMQC2 experiment was performed on a 1 mM uniformly ^{15}N -labeled IRE RNA (inset of Figure 1) sample containing ~17 mg/mL Pf1 bacteriophage (McCallum and Pardi, 2003). Figure 1 shows the imino proton region of the 2D ^1H - ^1H BEST-Jcomp-HMQC2 spectrum collected at 25 °C on a 600 MHz instrument equipped with a cryogenically cooled probe using HMQC constant time delays of $\Delta_1 = \Delta_2 = 35$ ms (Supplementary Figure 1), an inter-scan delay of 0.69 s and 400 scans per FID for a total experiment time of 45 hrs. As expected, the selective excitation results in only couplings between the imino groups with no other peaks observed in the spectrum (data not shown). The strongest cross peaks in the BEST-Jcomp-HMQC2 spectrum arise from imino protons in the G25-U5 wobble base pair, where the two imino protons are separate by ~3.1 Å (McCallum and Pardi, 2003). Cross peaks are also observed for the G22-U9 base pair (~2.2 Å imino proton distance), and the remaining cross peaks in Figure 1 involve imino protons on neighboring base pairs in the IRE. Some non-

linear phase error is observed in the indirect ^1H dimension, possibly due the selective pulses; however, it is still possible to obtain precise values for the imino ^1H - ^1H RDC (see below).

The absolute value of D_{HH} ($|D_{\text{HH}}|$) was calculated from the ratio of cross ($I_{\text{cross}}^{\text{AB}}$ and $I_{\text{cross}}^{\text{BA}}$) and diagonal peak ($I_{\text{diag}}^{\text{AA}}$ and $I_{\text{diag}}^{\text{BB}}$) intensities according to the following equation (Wu and Bax 2002; Schanda et al. 2007):

$$\frac{I_{\text{cross}}^{\text{AB}} I_{\text{cross}}^{\text{BA}}}{I_{\text{diag}}^{\text{AA}} I_{\text{diag}}^{\text{BB}}} = \frac{\lambda^2 \sin^2(\pi D_{\text{HH}} \Delta_1) \sin^2(\pi D_{\text{HH}} \Delta_2)}{\lambda^2 \cos^2(\pi D_{\text{HH}} \Delta_1) \cos^2(\pi D_{\text{HH}} \Delta_2) + 2\lambda(1-\lambda) \cos(\pi D_{\text{HH}} \Delta_1) \cos(\pi D_{\text{HH}} \Delta_2) + (1-\lambda)^2} \quad (1)$$

where Δ_1 and Δ_2 are the constant time delays for the first and second HMQC blocks ($\Delta_1 = \Delta_2 = 35$ ms), respectively, and λ is a correction for the percentage of protonation versus deuteration at each imino residue ($\lambda = 0.95$ for this IRE NMR sample) (Wu and Bax, 2002). The intensity ratios and experimental $|D_{\text{HH}}|$ values for the data set shown in Figure 1 are reported in Table 1. Error estimates in the intensity ratios and experimental $|D_{\text{HH}}|$ values were obtained from triplicate BEST-Jcomp-HMQC2 experiments. Two additional data sets were collected at 500 MHz at 25 °C with a room temperature probe using the same numbers of complex points in both dimensions as the 600 MHz data set, an inter-scan delay of 0.8 s and 256 scans per FID for a total experiment time of 20 hrs (data not shown). The average standard deviation of 0.1 Hz for the $|D_{\text{HH}}|$ values indicates a high level of precision.

The experimental $|D_{\text{HH}}|$ values were compared to $|D_{\text{HH}}|$ values predicted from the previously RDC-refined NMR solution structure for the IRE (Table 1) using the REDCAT program (Valafar and Prestegard, 2004). The alignment tensor used for these predictions was determined from the experimental ^1H - ^{15}N RDCs for the IRE (McCallum and Pardi, 2003). Figure 2 shows the plot of the predicted versus experimental (600 MHz and average values) imino ^1H - ^1H RDCs. The 600 MHz data have a Pearson's correlation coefficient (R_p) of 0.99 and fit well to the RDC-refined solution structure: average rmsd = 0.35 Hz and Q-factor = 0.11 (Cornilescu, *et al.*, 1998). The largest experimental $|D_{\text{HH}}|$ (5.4 Hz) was between the imino protons in the G25-U5 base pair and the next largest $|D_{\text{HH}}|$ (3.7 Hz) was for the inter-base pair G4-U5 interaction. The smallest magnitude imino-imino D_{HH} that was observed in the IRE was between the imino protons of G4-G25 (1.9 Hz). The smaller experimental than predicted D_{HH} value for the G22-U9 interaction results from overlap of the U9 diagonal peak with the imino resonance of G21 (see Eqn. 1 and Fig. 1). This overlap could be resolved by collecting the BEST-Jcomp-HMQC2 experiment as a 3D experiment with an additional ^{15}N dimension. With the exception of the G22-U9 wobble base pair, D_{HH} interactions are only observed between imino protons in the lower helical domain of the IRE RNA (Figure 1). The largest D_{HH} values predicted between imino protons on neighboring base pairs in the lower helix were for G1-G2 (3.1 Hz) and G2-U27 (1.6 Hz), but were not observed. The G1-G2 interaction is not observed because of rapid base pair opening and exchange of the imino proton with water in the terminal base pair. The G2-U27 $|D_{\text{HH}}|$ value of 1.6 Hz is less than the smallest resolved interaction observed (1.9 Hz for G4-G25), and thus is below the sensitivity of the current data set. The predicted $|D_{\text{HH}}|$ values for other inter base pair interactions in the upper stem, G21-U9, G21-G22 and G21-U20, were all too small to observe (< 1 Hz).

Additional structural information can be obtained in the IRE by comparing the BEST-Jcomp-HMQC2 spectrum (Figure 1) with the ^1H - ^1H NOESY spectrum (Figure 3). For example, the G4 imino proton shows 3 cross peaks to imino protons of U5, G25 and U27 in the BEST-Jcomp-HMQC2 spectrum (boxed in Figure 1). The G4-U5 interaction is the most intense of

these (Figure 1), has the largest ratio of intensities of cross peaks to diagonal peaks (Table 1) and therefore has a larger magnitude for D_{HH} (3.7 Hz) compared to what was observed for G4-U27 interaction (2.3 Hz). In contrast, the G4-U5 and G4-U27 crosspeaks are of comparable volumes (2.70×10^7 and 2.58×10^7 , respectively) in the NOESY spectrum (Figure 3). These data indicate that G4-U5 and G4-U27 have similar distances between imino protons but that these interactions have different orientations with respect to the alignment tensor of the IRE. This is confirmed in Figure 4, which highlights the 1H - 1H interactions involving G4 in the NMR structure of the IRE (McCallum and Pardi, 2003), where there are similar distances for the imino proton-imino proton interactions for G4-U5 and G4-U27 (Table 1), but where the angles of these interactions differ giving rise to different RDC values. Thus, the 1H - 1H RDCs for imino protons on neighboring base pairs complement the standard distance restraints obtained from NOESY data by providing information on the relative orientations of the observed interactions. Additionally, the imino 1H - 1H RDCs from neighboring base pairs also complement one-bond imino 1H - ^{15}N intra-base RDCs, because the imino D_{HH} interactions will generally point along the helical axis; whereas, the imino D_{NH} vectors are generally perpendicular to the helix axis. Thus, the measurement of imino D_{HH} values represent a valuable target for additional restraints to be used in high-resolution structure determination and domain orientation studies of RNAs.

In some cases, the absence of an observable RDC can still provide structural information, especially when combined with 1H - 1H NOESY data (Tjandra, *et al.*, 2000a, Tjandra, *et al.*, 2000b). For example, some short through-space 1H - 1H interactions that give rise to strong NOESY cross peaks showed no cross peaks in DQF-COSY or TOCSY spectra of a partially aligned DNA duplex because the 1H - 1H vectors are at angles that lead to very small RDCs (Hansen *et al.* 1998a; Hansen *et al.* 1998b). Thus, the 1H - 1H RDC data yield orientational restraints that are not available from NOESY spectra. For the IRE data presented above, through-space correlations between imino protons in the upper helix were observed in the 1H - 1H NOESY data, whereas the corresponding RDC correlations were all too small to detect in the BEST-Jcomp-HMQC2 spectra (< 1 Hz). However, we observed an additional issue when comparing the 2D BEST-Jcomp-HMQC2 and NOESY spectra on the IRE. This is illustrated in the 1D spectra for the first t_1 value (Supplemental Figure 2), where some imino resonances in the NOESY spectrum are not observed in the BEST-Jcomp-HMQC2 spectrum. These spectral differences cannot arise from solvent exchange because the NOESY experiment has a longer effective “mixing time” for solvent exchange than the BEST-Jcomp-HMQC2 experiment (~ 175 ms versus ~ 70 ms). Thus, the differences in intensities of the imino resonance likely arise from differential multiple-quantum relaxation properties of imino groups during the constant time periods required for build-up of the 1H - 1H RDCs in the BEST-Jcomp-HMQC2 experiment. This means it is not possible to directly compare sizes of cross peaks in the NOESY spectrum with those in the BEST-Jcomp-HMQC2 spectrum, and instead the magnitudes of the RDCs obtained from Equation 1 need to be compared with the NOESY data.

Tjandra *et al* (2000a) previously developed a protocol for incorporating 1H - 1H RDCs into simulated annealing structure calculations, using the protein ubiquitin as a model system. This protocol employs an alignment tensor derived from one-bond RDCs, such as D_{NH} values, and an energy minimization procedure to find values of 1H - 1H distance and orientation that satisfy the experimentally observed D_{HH} restraints. For ubiquitin, including the 1H - 1H RDCs increased the accuracy and precision of the structure when compared to previous X-ray and NMR solution structures (Tjandra, *et al.*, 2000a). A similar protocol was also used to incorporate 1H - 1H RDC restraints, measured between non-exchangeable protons, into the calculation of a high-resolution solution structure of the Dickerson dodecamer DNA (Tjandra, *et al.*, 2000b, Wu, *et al.*, 2003). These structure determinations utilized both the measured $|D_{HH}|$ and the absence of an observable 1H - 1H RDC as restraints in the refinement (Tjandra, *et al.*, 2000a, Tjandra, *et al.*, 2000b, Wu, *et al.*, 2003). Thus, the imino 1H - 1H RDC data obtained

here could be directly used in this type of high-resolution structure determination. Furthermore, the methods developed by Tjandra *et al* (2000a) for incorporating ^1H - ^1H RDC into structure refinements could also be used for domain orientation studies of RNAs.

Imino ^1H - ^1H RDCs were also measured at 25 °C on a 0.75 mM ^{15}N -labeled sample of native *E. coli* tRNA^{Val} containing ~12 mg/mL Pf1 bacteriophage. HMQC constant time delays of 35 ms were employed using the same experimental parameters as described above for the IRE RNA. Only two sets of imino D_{HH} cross peaks were observed: between the imino protons of the G50-U64 wobble base pair and the imino protons of U29 and G42 in neighboring base pairs (Supplemental Figure 3). A large imino D_{HH} was expected based on the short distance between the imino protons in a G-U wobble base pair (<2.4 Å). The cross and diagonal peak intensity ratio for the G50-U64 imino proton interaction is 3.31 which is >10-fold larger than the largest intensity observed in the IRE (G25 and U5, Table 1), corresponding to an experimental $|D_{\text{HH}}|$ value of ~49 Hz (Equation 1, $\lambda = 0.90$). This experimental $|D_{\text{HH}}|$ value agrees well with the calculated value of 46.6 Hz derived from the previously described native *E. coli* tRNA^{Val} model and alignment tensor ($D_{\text{a}}^{\text{NH}} = 20.4$ Hz, $R = 0.62$, $\alpha = 31.9^\circ$, $\beta = 3.9^\circ$, and $\gamma = -23.3^\circ$) (Mollova, *et al.*, 2000, Vermeulen, *et al.*, 2005). The intensity ratio of the inter base pair U29-G42 interaction is 7.5×10^{-3} corresponding to a $|D_{\text{HH}}|$ value of 2.7 Hz (Equation 1), which is much smaller than the predicted value of 8.4 Hz. However, both the U29 and G42 diagonal peaks are overlapped, leading to lower intensity ratios and therefore lower $|D_{\text{HH}}|$ values. In such cases of overlap, the constraints derived from these ^1H - ^1H imino RDCs could take into account that the measured $|D_{\text{HH}}|$ represents only a lower limit, due to overestimate of the intensity of the diagonal peaks in Equation 1. Additional ^1H - ^1H imino RDCs were initially expected, so the imino D_{HH} were predicted using the previously determined tRNA^{Val} model structure and Pf1 alignment tensor (Mollova, *et al.*, 2000, Vermeulen, *et al.*, 2005). However, many of the predicted imino D_{HH} interactions are to be too small observed in the BEST-Jcomp-HMQC2 experiment. In fact, after the U29-G42 interaction, the next largest $|D_{\text{HH}}|$ interaction occurs between the imino protons of G2-G3, G3-G4 and G52-G53 with predicted RDC values of ~3-4 Hz, and majority of the other predicted RDC values are < 1 Hz. The very small ^1H - ^1H RDCs arise because these imino ^1H - ^1H interactions generally point along the helical axis and the orientation of the two helical regions of this tRNA with respect to the alignment tensor is unfavorable for these RDCs (Mollova, *et al.*, 2000, Vermeulen, *et al.*, 2005). Since, both helices are oriented in a manner that only small imino ^1H - ^1H RDCs are observed, this information could still be used to help determine the global structure of this RNA. One issue that also needs to be considered when interpreting the absence of observable ^1H - ^1H RDC interactions is that the larger size of tRNA^{Val} (76 nucleotides) leads to faster multiple-quantum relaxation during the two HMQC periods, which reduces the sensitivity of the BEST-Jcomp-HMQC2 experiment.

One method for increasing the relative size of cross peaks in the BEST-Jcomp-HMQC2 experiment is to use a higher degree of molecular alignment. Thus the BEST-Jcomp-HMQC2 experiment was also performed on a ^{15}N -labeled sample *E. coli* tRNA^{Val} containing ~25 mg/mL Pf1 phage. However, this higher degree of alignment resulted in very broad lines in the direct ^1H dimension, which arise from the large imino proton to amino proton dipolar couplings. Therefore, higher alignment can actually reduce the signal-to-noise in the BEST-Jcomp-HMQC2 experiment. A second approach for increasing the relative size of the cross peaks in the BEST-Jcomp-HMQC2 experiment is to increase the two constant time delays in the two HMQC periods (Δ_1 and Δ_2 in Supplemental Figure 1), which allows more time for smaller couplings to evolve. Thus, a 2D ^1H - ^1H BEST-Jcomp-HMQC2 experiment was acquired on the IRE at 500 MHz varying the HMQC constant time delays from 30 to 60 ms. As expected from Equation 1, the ratio of the cross peaks and diagonal peaks increases with longer Δ delays; however, the signal-to-noise decreases because of increased multiple quantum relaxation. In fact, the weak cross peaks observed between imino resonances G4 and G25 in

Figure 1 were no longer observed in the 60 ms constant time spectrum (data not shown). Thus, no new D_{HH} interactions were observed with the 60 ms constant time periods.

Finally, to test the effect of the BEST methodology (Schanda, *et al.*, 2007) for measurement of imino 1H - 1H RDCs, the spectrum obtained with the BEST-Jcomp-HMQC2 pulse sequence (Supplementary Figure 1) was compared with a spectrum obtained using the original semi-selective HMQC2 pulse sequence (Wu and Bax, 2002). The SS-HMQC experiment was acquired at 500 MHz on the IRE with the same number of complex points in each dimension as the BEST experiment, but with a longer inter-scan delay (1.6 vs. 0.8 s) and fewer scans per FID (144 vs. 256) resulting in the same total experiment time (20 hrs). For the SS-HMQC experiment, D_{HH} interactions were observed for all of the imino proton-imino proton interactions except G22-U9. However, for the peaks observed in both the BEST and SS-HMQC2 experiments collected at 500 MHz, the average increase in signal intensity is ~3.5-fold when the BEST methodology is utilized. This ~3.5-fold signal increase is higher than the ~1.8-fold increase predicted from the difference in the number of scans collected in each experiment (256 vs. 144). For example, Figure 5 compares 1D traces through the G4 imino resonance for the BEST and SS-HMQC2 spectra, illustrating that the G25 – G4 cross peak is barely observable in the SS-HMQC2 experiment. Thus, the longitudinal relaxation enhancement afforded by the BEST methodology leads to an increase in signal-to-noise.

It is demonstrated here that accurate imino proton-imino proton $|D_{HH}|$ values can be obtained using the BEST-Jcomp-HMQC2 experiment collected on partially aligned RNA samples. Interactions were observed between the imino protons in G-U wobble base pairs as well as between the imino protons of neighboring base pairs in the 29 nucleotide IRE RNA and 76 nucleotide native *E. coli* tRNA^{Val}. These RDC data complement standard NOESY distance restraints due to their additional dependence on the relative orientation of the inter-nuclear interactions to the common alignment frame. The D_{HH} data also provide complementary orientation information to the imino 1H - ^{15}N RDCs, since the imino 1H - 1H RDC generally point along the helical axis whereas the imino 1H - ^{15}N RDCs generally point perpendicular to the helical axis. Therefore, these imino D_{HH} restraints contain valuable information for domain orientation studies of RNAs.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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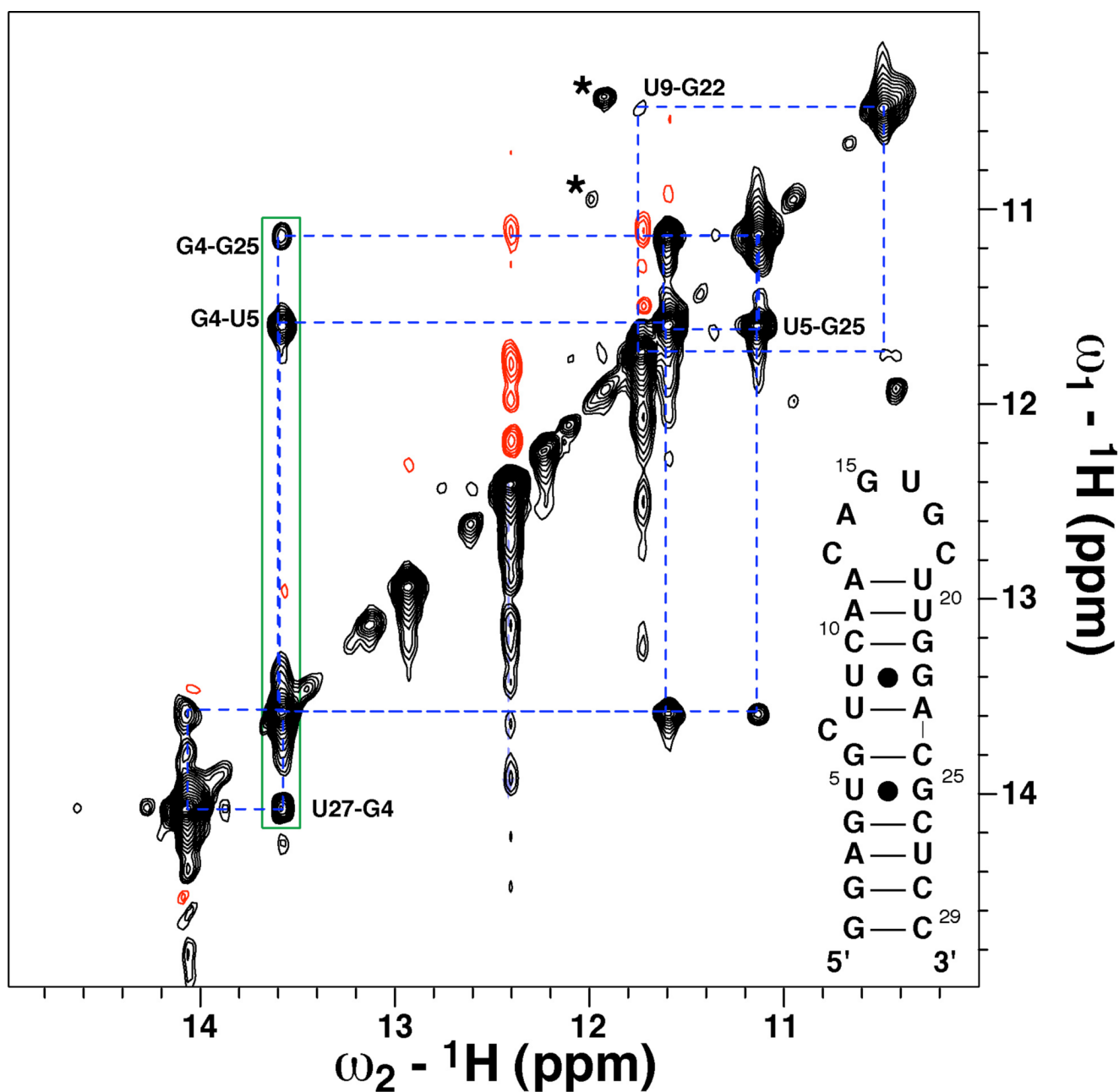


Figure 1.

Imino region of the 2D ^1H - ^1H BEST-Jcomp-HMQC2 experiment (Schanda, *et al.*, 2007) shown in Supplemental Figure 1 collected on a 1 mM ^{15}N -labeled IRE RNA containing ~17 mg/mL Pf1 bacteriophage. This spectrum was collected on a Varian INOVA 600 MHz spectrometer equipped with a cryogenically cooled triple resonance z-axis gradient probe operating at 25 °C. HMQC constant time delays were $\Delta_1 = \Delta_2 = 35$ ms. 140×2048 complex points were taken using sweep widths of 14006 Hz in the t_1 and t_2 dimensions. 400 scans per FID were collected with a 0.69 s inter-scan delay for a total experiment time of ~45 hrs. Black and red contours represent positive and negative intensity, respectively. The dashed blue lines illustrate the cross peaks which arise because of the transfer of magnetization due to imino ^1H - ^1H RDCs. The asterisks denote cross peaks for the two G-U wobble base pairs, which

are in an alternate conformation. The previously published imino assignments are indicated (Address, *et al.*, 1997), and the sequence and secondary structure of the IRE are shown in the lower right.

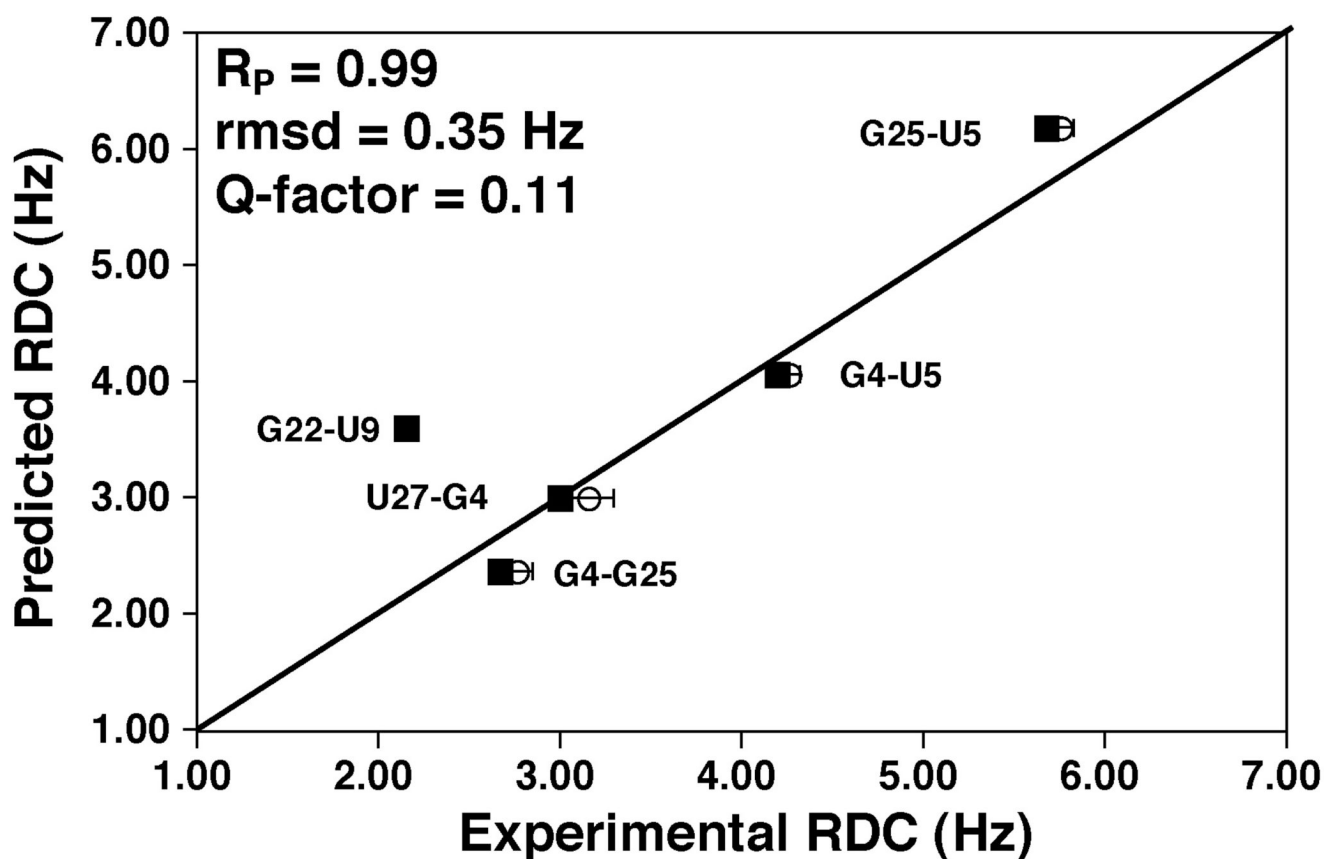


Figure 2.

Plot of the absolute value of the experimental and predicted imino ^1H - ^1H RDCs of the IRE. The predicted RDCs were obtained from the previously RDC-refined solution structure of the IRE. The experimental imino $|D_{\text{HH}}|$ values for data collected at 600 MHz (filled squares) are shown along with the average $|D_{\text{HH}}|$ values determined from triplicate measurements at 500 MHz (open circles). The predicted RDCs were obtained using an alignment tensor of $D_a^{\text{NH}} = -18.8 \text{ Hz}$, $R = 0.16$, $\alpha = 24.1^\circ$, $\beta = 108.4^\circ$, and $\gamma = 123.7^\circ$ and the refined solution structure (McCallum and Pardi, 2003) as input in REDCAT (Valafar and Prestegard, 2004). Assignments for the $|D_{\text{HH}}|$ interactions are given for each point, and error bars on the average $|D_{\text{HH}}|$ values represent the standard deviation calculated from triplicate experiments. The Pearson's correlation coefficient, R_p , average rmsd and Q-factor (Cornilescu, *et al.*, 1998) for the 600 MHz data set are listed in the upper left-hand corner. These parameters were calculated without the $|D_{\text{HH}}|$ value for G22-U9 because of overlap of U9 and G21 imino proton resonances (see text).

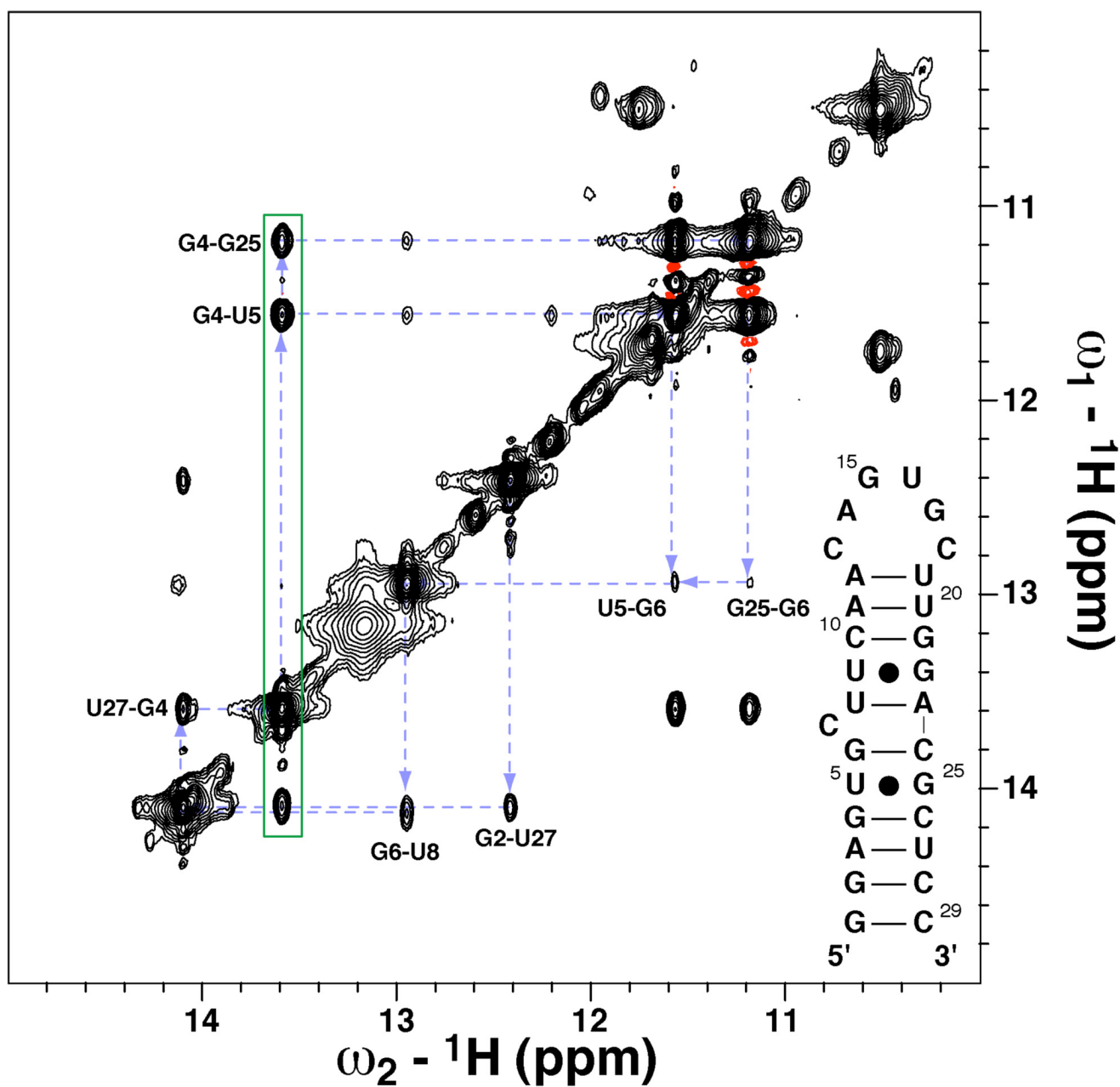


Figure 3.

Imino region of the 2D ^1H - ^1H NOESY experiment with 175 ms mixing time collected on a 1 mM ^{15}N -labeled IRE containing ~17 mg/mL Pf1 phage. This spectrum was collected at 5 °C on a Varian INOVA 600 MHz spectrometer equipped with a cryogenically cooled triple resonance z-axis probe. Water suppression was achieved with the gradient 11-echo scheme (Sklenár and Bax, 1987). Black and red contour lines represent positive and negative intensity, respectively. Dashed blue lines show the imino proton walk in the lower helix, and the green box highlights the three cross peaks to the imino proton of G4.

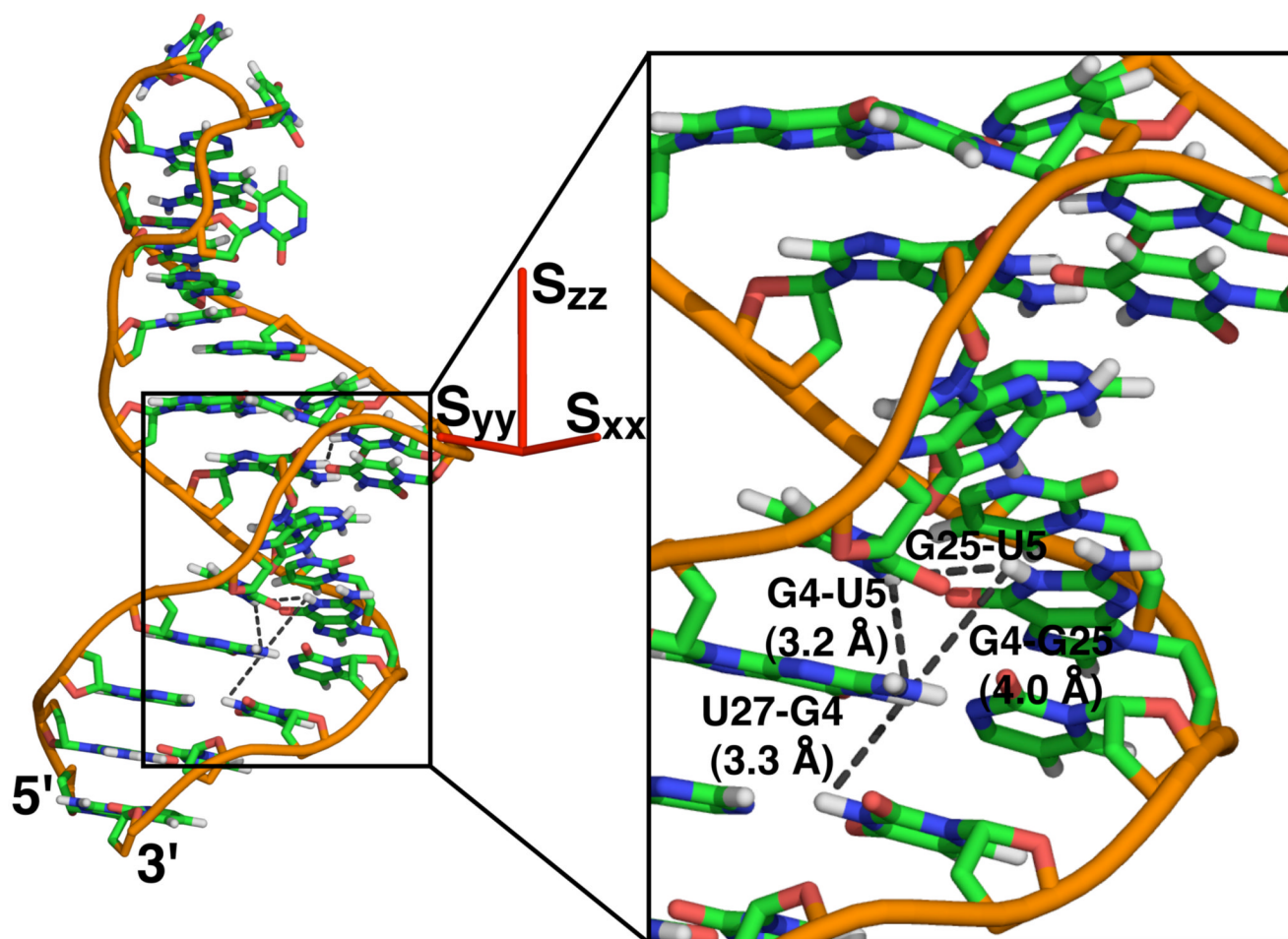


Figure 4. Experimentally observed imino D_{HH} interactions are highlighted on the RDC-refined solution structure of the IRE (pdb entry 1NBR) (McCallum and Pardi, 2003). The alignment tensor (red axis system) resulting from alignment in ~17 mg/mL Pf1 bacteriophage is shown next to the structure (McCallum and Pardi, 2003). The lower stem containing some of the D_{HH} interactions is expanded highlighting dipolar interactions for G4 labeled with their respective inter-nuclear distances.

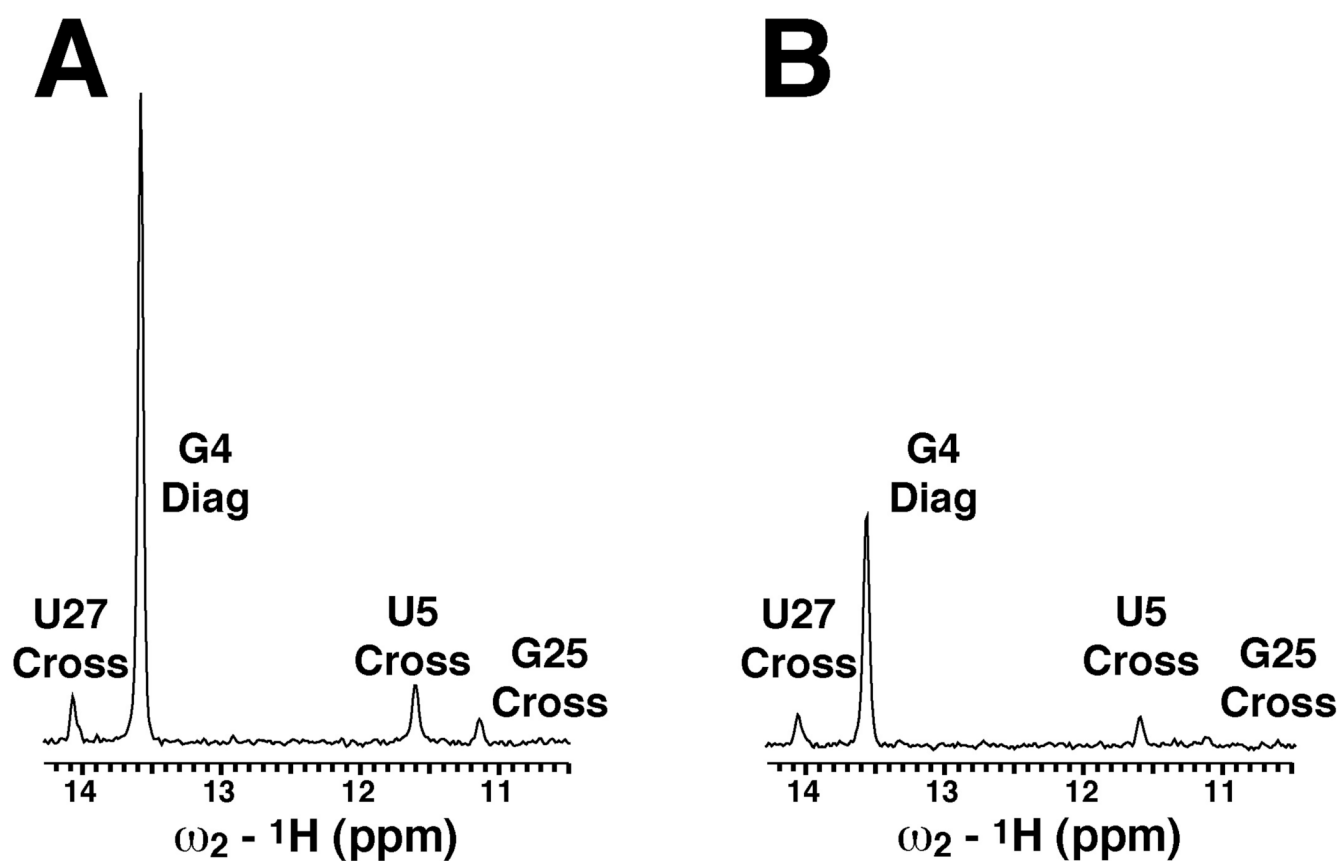


Figure 5.

1D traces for the G4 imino resonance taken from the 2D (A) BEST-Jcomp-HMQC2 and (B) SS-HMQC2 spectra on the IRE in Pf1 bacteriophage. Both spectra were collected at 500 MHz and 25 °C for ~20 hr, as described in the text. The assignments for G4 diagonal peak and U27, U5 and G25 cross peaks are given.

Table 1

Experimental and predicted imino ^1H - ^1H RDCs for ^{15}N -labeled IRE RNA containing ~17 mg/mL Pf1 phage.

Imino Assignment ^a	$\frac{I_{cross}^{AB} I_{cross}^{BA}}{I_{diag}^{AA} I_{diag}^{BB}} (\times 10^{-3})^b$	Experimental $ \text{D}_{\text{HH}} $ (Hz) ^c	Predicted $ \text{D}_{\text{HH}} $ (Hz) ^d	Internuclear Distance (Å) ^e
U27 – G4	3.6 ± 1.7	2.3 ± 0.2	2.3	3.3
G4 – U5	27 ± 4.8	3.7 ± 0.1	3.5	3.2
G4 – G25	1.7 ± 0.6	1.9 ± 0.1	1.5	4.0
G25 – U5	160 ± 30	5.4 ± 0.1	5.5	3.1
G22 – U9 ^f	0.37	1.3	3.9	2.2

^a Assignment of D_{HH} interactions observed in Figure 1 (Address, *et al.*, 1997).

^b Ratio of the intensities of the cross and diagonal peaks observed in the 2D ^1H - ^1H BEST-Jcomp-HMQC2 experiments. Uncertainties in the intensity ratios are the standard deviation calculated from triplicate experiments.

^c Experimental $|\text{D}_{\text{HH}}|$ values determined by fitting of the experimental intensity ratios using Equation 1. $\Delta_1 = \Delta_2 = 35$ ms and $\lambda = 0.95$ (i.e., 95% $\text{H}_2\text{O}/5\%$ D_2O). Errors in D_{HH} are the standard deviation calculated from triplicate experiments.

^d Predicted $|\text{D}_{\text{HH}}|$ values from the RDC-refined structure of the IRE RNA (McCallum and Pardi, 2003). The previously reported $^1\text{D}_{\text{HN}}$ values and solution structure (McCallum and Pardi, 2003) were used as input for the REDCAT program (Valafar and Prestegard, 2004) for SVD calculation of the alignment tensor. This tensor ($\text{D}_{\text{a}}^{\text{NH}} = -18.8$ Hz, $R = 0.16$, $\alpha = 24.1^\circ$, $\beta = 108.4^\circ$, and $\gamma = 123.7^\circ$) was used along with the structure to then predict the D_{HH} values.

^e Internuclear distances determined from the RDC-refined solution structure of the IRE RNA (pdb entry 1NBR)(McCallum and Pardi, 2003).

^f Cross peaks between the imino protons of G22 and U9 were not observed in the two data sets collected at 500 MHz.