19-Fluorine nuclear magnetic resonance chemical shift variability in trifluoroacetyl species

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Abstract: This review examines the variability of chemical shifts observed in 19-fluorine (19F) nuclear magnetic resonance spectra for the trifluoroacetyl (TFA) functional group. The range of 19F chemical shifts reported spectra for the TFA group varies generally from −85 to −67 ppm relative to CFCl3. The literature revealed several factors that impact chemical shifts of the TFA moiety—substrate topology, electronic environment, solvent polarity, and concentration effects. Often these effects conspire to cause deshielding of the TFA group by up to 15 ppm. These factors will be examined for a series of TFA-functionalized acyclic, cyclic, aromatic, and heterocyclic molecules.

Keywords: 19F NMR, chemical shift, shielding, trifluoroacetyl group, fluorinated diketones, triketones

Fluorinated molecules have found wide application in many areas, including the pharmaceutical and agrochemical industries and as precursors to surfactants, polymers, and flame retardants.1–3 The properties exhibited by molecules incorporating fluorine often include tuned lipophilicity or lipophobicity as well as enhanced bioactivity.4–7 As a biochemical marker and precursor to heterocyclic species, the trifluoroacetyl (TFA) group has gained exceptional versatility; its incorporation into molecular architecture is effected via its use as a synthon or through acetylation strategies.

19-Fluorine (19F) nuclear magnetic resonance (NMR) has become widely used as a means of verifying successful introductions of specific TFA moieties in organic molecules.8,9 In the pivotal work Guide to Fluorine NMR for Organic Chemists, Dolbier described the use of 19F NMR spectroscopy as a tool for structural elucidation and characterization, especially when combined with other spectroscopic methods.10 An examination of the scientific literature showed that while the variation demonstrated in the chemical shifts exhibited by the TFA group can be of tremendous assistance in elucidating the molecular topology of fluorinated materials, factors, such as electronic environment, solvent polarity, and concentration, often modulate 19F NMR resonance frequencies observed for the TFA group.11–13

This review examines 19F NMR chemical shift variability for nine classes of molecules that contain TFA groups: isolated α,α,α-trifluorocarbonyl species, α,β-unsaturated TFA species, acyclic trifluoromethyl-β-dicarboxyls, 2-trifluoroacetylcycloalkanones, fused-ring trifluoroacylated ketones, α,α,α-trifluoromethyl-1,3,5-triketones, cyclic trifluoroacylated triketones, trifluoroacylated aromatics, and trifluoroacylated heterocycles. The species chosen in this study are of broad importance in the field of organic synthesis as building blocks to highly functionalized, industrially,
and medicinally valuable molecules that incorporate the trifluoromethyl (CF₃) moiety. The effects of substrate topology, electronic environment, the influence of solvent—especially when related to modulation of tautomeric equilibria and concentration, will be discussed for each TFA class.

**Factors influencing ¹⁹F NMR chemical shifts of the TFA functional group**

**Shielding of the fluorine nucleus — general considerations**

The chemical shifts exhibited by CF₃ group-containing molecules in the ¹⁹F NMR range from −5.1 ppm (relative to CFCl₃) in the case of CF₃I to more than −84 ppm for hexafluoroacetone, (CF₃)₂C=O (Figure 1).

An important aspect of ¹⁹F NMR chemical shift interpretation for the −CF₃ group involves an understanding of the effect of shielding on the fluorine nucleus. As Figure 1 shows, as field strength increases, shielding of the CF₃ group increases. This is perhaps not intuitive since, for most persons who are familiar with proton NMR, the reverse is observed.

Of course, there is an explanation for this difference. For NMR active nuclei, the effective magnetic field experienced by the nucleus (B_eff) is given by Equation 1:

\[ B_{\text{eff}} = B_0 - \sigma B_0 \]

where B₀ is the applied magnetic field and σ is the shielding constant. The shielding constant arises from the contributions of the terms shown in Equation 2:

\[ \sigma = \sigma_{\text{dia}} + \sigma_{\text{para}} + \sigma_i, \]

where σ_dia is defined as the opposing field generated by the applied magnetic field’s effect on the atom’s electron cloud, σ_para is the contribution to shielding resulting from the applied magnetic field’s interaction with the atom’s π orbital electrons, and σ_i is the contribution arising from the influence that neighboring groups have on the atom of interest. Whereas for the hydrogen atom that contains a lone s orbital electron, the principal contribution to shielding is σ_dia; σ_para dominates in the case of fluorine due to the large number of p electrons present. The σ_para contribution is opposite in its shielding effect from that of σ_dia, hence the typical deshielding effect observed in proton spectra does not apply to ¹⁹F spectra. Presented below are factors that impact the chemical shifts observed specifically for the TFA group. These include substrate topology, electronic environment, solvent polarity and concentration effects.

**Substrate topology**

As Figure 1 shows, the carboxyl moiety of the TFA group shields −CF₃ fluorines to a greater extent than other functionalities. Certain TFA groups attached to molecular residues that extend conjugation or permit tautomerization can, however, be deshielded by as much as 15 ppm.

As the Figure 2 shows, α,α,α-trifluoro-β-diketone tautomerization from the diketo form (π_Csp2-Osp2) to a keto-enol form (π_Csp2-Csp2) results in a shift of the ¹⁹F NMR resonance for the −CF₃ group to a lower field as a result of diminished shielding by the C=C enolic topology. To illustrate this trend, the ¹⁹F NMR spectra of 1,3-ditrifluoroacetyl-2-indanone, 2-trifluoroacetyl-1-tetralone and 3-trifluoroacetyl-2-naphthol (Figure 3) show how keto-enolic and ring-size substrate topology influences chemical shift. As indicated, the smaller ring size and exocyclic keto-enol structure in 1,3-ditrifluoroacetyl-2-indanone, 2-trifluoroacetyl-1-tetralone and 3-trifluoroacetyl-2-naphthol (−68.5 ppm) than the endocyclic keto-enol TFA-substituted naphthol (−72.1 ppm) aromatic TFA-substituted naphthol (−74.3 ppm) examples. The aromatic ring “locks” the COCF₃ group into an endocyclic keto-enol configuration, shielding the fluorines further. A similar effect is also observed in conjugated CF₃ ketones, such as the trifluoroacetophenones, where chemical shifts can vary up to 10 ppm.

**Electronic environment**

Shielding diminution due to resonance effects likewise influences the TFA group. This phenomenon is exhibited by the carboxylic acid derivative family of compounds and can result in deshielding of the fluorine nucleus by −6−8 ppm relative to chemical shifts observed for aliphatic CF₃ ketones. In Figure 4, the contributors to the overall resonance hybrid highlight a redistribution of electron density that serves to
mitigate the shielding effect of the carbonyl on the fluorine atoms of the CF₃ group.

**Solvent polarity**

In general, solvent polarity impacts chemical shifts in the ¹⁹F NMR by no more than ~2 ppm.¹⁰ In most situations, shielding decreases as solvent polarity increases, with highly polar aprotic solvents, such as dimethyl sulfoxide (DMSO) and dimethyl formamide (DMF), having the largest deshielding effect.¹⁰ One may rationalize this phenomenon on the basis of increased solvent clustering around the highly electron-withdrawing −CF₃ group, ameliorating the shielding effect of neighboring functional groups.

An exception to the rather limited effect noted above occurs when a change in solvent polarity influences
Concentration effects
In some limited situations, concentration can alter tautomeric equilibria exhibited by the β-diketone classes of TFA-bearing molecules. As Reichardt and other researchers noted, highly concentrated solutions of β-diketones favor a tautomeric shift toward the diketo form (Figure 2). In a few cases where the subject compounds investigated are neat liquids, 19F NMR studies showed that small proportions of the diketo tautomer (usually <10%) are present in addition to the keto-enol form. This effect is linked to substrate topology and is observed more commonly in cyclic examples than in acyclic cases. For example, variable concentration studies of 4,4,4-trifluoro-1-phenyl-1,3-butanedione (0.0016 M–0.20 M, neat liquid) reveal no chemical shift variability in the 19F NMR spectra, while neat liquid 19F NMR studies of 2-trifluoroacetylcyclopentan-1-one, 2-trifluoroacetylcyclohexan-1-one, 2-trifluoroacetylcycloheptan-1-one, and 2-trifluoroacetylcyclooctan-1-one show both diketo and keto-enol forms present, with the diketo form being present in less than 10% proportion to the keto-enol form.

Temperature effects
While temperature changes are noted to impact chemical shifts of complex biomolecules by altering the three-dimensional nature of the molecules, simple TFA-bearing molecules of interest in this review do not show any chemical shift variability. Variable temperature studies performed on 1,1,1-trifluoro-2,4-pentanedione (253 K to 298 K) and 4,4,4-trifluoro-1-phenyl-1,3-butanedione (298 K to 348 K) show no chemical shift change in their 19F NMR spectra as a function of temperature.

Discussion and analysis of chemical shift variations for TFA-bearing molecules
We begin our review by considering the range of 19F chemical shift values associated with the simplest examples of the TFA functionality, those bearing TFA groups that are isolated from functional groups that exhibit conjugating effects. This will be followed by an examination of more complex trifluoroacetylated molecules. For direct comparative purposes, all fluorine NMR chemical shifts (δ) are reported in ppm relative to CFCl3.

Isolated α,α,α-trifluorocarbonyl species
Carbonyl species in this group and shown in Figure 6 include ketones, aldehydes, pyruvates, and carboxylic acid derivatives. As can be seen, the CF3 group in aldehydes and ketones is shielded relative to that of the pyruvates and carboxylic acid derivatives. In the case of the pyruvate family, the α-carbonyl...
residue can be considered a de facto conjugating group to the TFA functionality, having a net deshielding effect on the CF$_3$-19F chemical shift. For the carboxylic acid derivatives, the chemical shifts observed are largely a function of resonance deshielding. One notes that with the improved donor capacity of the nitrogen, the deshielding effect is enhanced, albeit to a small degree. Solvent and concentration effects, if any, would be small for these examples.

$\alpha,\beta$-Unsaturated trifluoromethyl ketones

The presence of an alkenyl unit in conjugation with the TFA group (compound series 10) deshields the $\text{CF}_3$ group relative to isolated CF$_3$ ketones (compound series 2) (Figure 7). This effect is smaller than the resonance deshielding observed for the trifluoroacetic acid derivatives (series 5–9), with chemical shifts for compound series 10 differing by only $\sim$1 ppm. Even when $Y = \text{Ph}$, the additional conjugation afforded by the benzene ring does not serve to deshield the $\text{CF}_3$ group to any greater extent ($\delta = -78.1$ ppm). Solvent and concentration effects on the $^{19}$F NMR chemical shifts would be minimal for this series of compounds.

Acyclic trifluoromethyl-$\beta$-dicarboxyls

This class of TFA-containing compounds has been thoroughly investigated. As shown in Figure 8, the $\beta$-diketones (compound series 11) show a marked preference for the keto-enol tautomeric forms a and b in nonpolar solvents. For cases where R = alkyl groups, the a= b dynamic equilibrium is near unity and only one $^{19}$F NMR resonance for the $\text{CF}_3$ group is observed, usually ranging from $-78$ to $-76$ ppm. In polar aprotic and polar, protic solvents the equilibrium can shift in certain cases toward the diketo form, but the keto-enol forms remain dominant. When $R = \text{aryl}$, recent evidence suggests that keto-enol form b predominates because of extended conjugation afforded by the aromatic ring, and a single $^{19}$F NMR resonance for the $\text{CF}_3$ group is observed, again ranging from $-78$ to $-76$ ppm. Solution studies of selected trifluoromethyl-$\beta$-diketones ($R = \text{aryl}$) have found little evidence of a discernible concentration effect over a wide range of concentrations.

Conversely, in the cases of compound series 12 and 13, the only form observed is the diketo form. For series 12, this preference is rationalized on the basis of Frontier Molecular Orbital theory. The presence of a single 2-fluoro substituent destabilizes the keto-enol form, leading to observations of the more highly shielded diketo tautomer, which has chemical shifts above $-80$ ppm. This preference holds true in all examined solvent systems, whether nonpolar, polar aprotic, and polar protic. It also appears to be independent of concentration. For compound series 13, only the diketo form is possible, yielding chemical shift values $> -80$ ppm as well.

The $\beta$-keto ester and $\beta$-ketoamide series 14 enolizes in the direction of the $\text{CF}_3$ group, producing an enol form reminiscent of the keto-enol form in series 11. As expected, this enolization deshields the $\text{CF}_3$ group; chemical shift values range from $-77$ to $-75$ ppm.
Trifluoroacetyl-cycloalkanones

In cases where the TFA group is attached to a ring system, such as those shown in Figure 9, substrate topological constraints serve to further deshield the $-\text{CF}_3$ group relative to the acyclic trifluoromethyl-$\beta$-dicarbonyls discussed earlier.

A neat liquid $^{19}$F NMR study of these compounds, conducted by Ebraheem, helped establish that ring size can modulate chemical shifts and that highly concentrated samples of these compounds shift the keto-enol $\rightleftharpoons$ diketo equilibrium to permit observation of the diketo form.

According to Ebraheem, the $^{19}$F NMR data for compound 15 is indicative of an exocyclic keto-enol form as the major tautomer present (98%). It is possible that the observed chemical shift value for 15 ($\delta = -74.95$ ppm) is partially attributed to the limited flexibility of the smaller cyclopentane ring system. The diketo form (2%) was also observed for 15 ($\delta = -80.96$ ppm). When the ring size is increased from 5 to 6–8 carbons (compounds 16–18), however, the dominant form is the endocyclic keto-enol (90.5%–93.9%) with a slightly larger proportion of the diketo form also found in the neat liquids (6.1%–9.5%). The chemical shifts recorded for compounds 16–18, (keto-enol: $\delta = -71.81$ to $-72.99$ ppm; diketo: $\delta = -79.90$ to $-80.50$ ppm) are indicative of how substrate topology in larger, more flexible ring systems can cause slight deshielding of the fluorine nucleus.

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure8}
\caption{Acyclic trifluoromethyl-$\beta$-dicarbonyls. Abbreviations: CF$_3$, trifluoromethyl; R, alkyl, aryl; OR, alkoxy; NH$_2$, NHR, NR$_2$, amino.}
\end{figure}

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure9}
\caption{Trifluoroacetyl-cycloalkanones.}
\end{figure}
Fused-ring trifluoroacetylated ketones
The molecules depicted in Figure 10 are trifluoroacetylated derivatives of indanone and tetralone, fused-ring systems incorporating benzene and five- or six-membered rings, respectively.\(^{13,14}\) As in the previous section, the variation of ring size and topology influences the shielding of \(^{19}\)F NMR resonances for these compounds.

In a dilute solution phase study, Salman found that 2-trifluoroacetyl-1-indanone (19) followed a similar trend observed for compound 15, namely that the exocyclic keto-enol form was dominant.\(^{13}\) Compound 20, 2-trifluoroacetyl-1-tetralone, was found to exist as a 1:1 ratio of endocyclic: exocyclic keto-enols. The chemical shifts for 19 and 20 in the highly polar aprotic solvent dimethylformamide (DMF) show greater deshielding of the CF\(_3\) group relative to CDCl\(_3\), in accordance with other solvent-effects studies mentioned earlier. It is interesting to note that the diketo form (~6%) was only identified for compound 20, indicating that the polarity of DMF is sufficient to shift the keto-enol=diketo equilibrium slightly for the six-membered ring system. The cyclopentane ring of the indanone (19) may lack the structural flexibility to overcome the keto-enol preference even in highly polar solvents.

In more recent work, compounds 21 and 22 were investigated and found to exist solely as the exocyclic keto-enol and the endocyclic keto-enol forms, respectively.\(^{14}\) The higher level of deshielding of the CF\(_3\) group in the case of these compounds relative to 19 and 20 is likely due to a combination of substrate topology and the extended conjugation of the keto-enol forms.

Compound 23 is an unusual case, the study of which is ongoing. Initial \(^{19}\)F NMR spectroscopic results indicate that the molecule exists as the triketone (\(\delta = -75.90\) ppm), while ab initio computations show the triketone tautomer to be lower in energy by approximately 1 kcal/mol than either the exocyclic or endocyclic keto-enol tautomers (Sloop et al, unpublished data, 2013).\(^{19}\)

Compound 24 was found to exist solely as the endocyclic keto-enol tautomer shown. The structure was verified by X-ray crystallography and the \(^{19}\)F NMR chemical shift observed is consistent with that of other keto-enol tautomeric forms.

In one of the first reported examples of an exocyclic dienol tautomer, compound 25 exhibits a higher degree of deshielding of the CF\(_3\) group (\(\delta = -68.50\) ppm), again a consequence of the substrate topology and the extended cross-conjugation noted in this structure. Like 24, the unique keto-enol structure has been corroborated by X-ray crystallography.

\(\alpha,\alpha,\alpha\)-trifluoromethyl-1,3,5-triketones
The trifluoromethyl-1,3,5-triketone class of compounds enjoys greater conjugation than \(\beta\)-diketones owing to the possibility of a variety of keto-enol tautomeric forms that
could potentially be present. Representative examples are shown in Figure 11.

Compound 26, 1,1,1,7,7,7-hexafluoro-2,4,6-heptanetrione, is a simple example of this class of triketone. The observed chemical shift at δ = −75.2 ppm as the sole 19F NMR resonance is fully consistent with the presence of the symmetric, cross-conjugated, dienolic tautomer depicted in Figure 11.20

As demonstrated in earlier examples, when an additional ring system is incorporated in the substrate topology, deshielding of the CF3 group increases. Recent studies of compound 27 show the extension of conjugation into the tetralone ring system causes the 19F NMR resonance to shift upfield slightly by ≈3 ppm. In this case, the observation of a resonance at δ = −72.04 ppm in the 19F NMR indicates a single, cross-conjugated, dienol tautomer.14

This deshielding trend is even more evident in compound 28, where we constrain the topology and electronic effects with the bicyclic, aromatic, naphthol system.14 In this instance, the hydroxyl group cannot participate in tautomerization to a ketonic form since that would require loss of aromaticity. The additional aromatic ring system in 28 further deshields the CF3 group, resulting in a shift upfield of 0.4 ppm relative to 27. Observation of a single 19F NMR resonance at δ = −71.60 ppm is indicative of the characteristic cross-conjugated dienolic tautomer depicted in Figure 11.

Cyclic trifluoroacetylated triketones

As our examination of cyclic trifluoromethylated β-diketones has shown, ring systems can modulate 19F chemical shifts of the CF3 group. For the cyclic trifluoromethylated triketones depicted in Figure 12, the 19F chemical shifts observed for the CF3 groups align with deshielding effects already discussed.

While compounds 29 and 30 are currently under investigation, preliminary NMR data collected on 29 suggest that the same type of substrate topology effects found in compounds 15, 19, and 21 is in operation – an exocyclic keto-enol form predominates in CDCl3 solution. For compound 30, the endocyclic keto-enol form is substantiated by the 19F NMR resonance at δ = −76.30 ppm, a fact consistent with data discussed for compounds 16, 20, and 22. Consistent with previously discussed solvent effects, NMR studies of compounds 29 and 30 in the highly polar DMSO further deshield the CF3 group by ≈2 ppm.19

Sevenard et al prepared and studied compound 31. Based on a combination of 1H, 13C, and 19F NMR data, they concluded that an 80:20 equilibrium exists between the exocyclic dienol and the endocyclic keto-enolic structures in the nonpolar solvent C6F6.21 A thorough review of the literature has revealed no other examples of a cyclohexanone-based bis-exocyclic enol tautomer. The most interesting finding in this study was that the exocyclic dienol was present in a larger proportion than the endocyclic tautomer, an observation that has not been supported previously in the literature for trifluoroacetylated cyclohexanones. Clearly, however, the observation of distinct 19F NMR resonances for each type of −CF3 group found in the tautomeric mixture as well as the chemical shift positions support the presence of the two separate keto-enol tautomers.

Trifluoroacetylated aromatics

As discussed earlier, extended conjugation tends to deshield the 19F NMR resonance of the TFA group. When a TFA group is attached directly to an aromatic ring system, the −CF3 19F NMR resonance is shifted upfield relative to nonconjugated, alkyl-substituted, trifluoroacetylated species, a consequence of deshielding. Examples of aromatics containing the TFA functionality are shown in Figure 13.

For compound 32, the topology of the aromatic ring system “locks” the TFA group into a keto-enol form that cannot tautomerize. The presence of the carbonyl adjacent to the −CF3 group and the ortho-substituted hydroxyl group that hydrogen bonds with the carbonyl combine effects for a 19F NMR resonance that is slightly more shielded than the other examples in our series.

Figure 11 α,α,α-Trifluoromethyl-1,3,5-triketones.
Abbreviation: CF3, trifluoromethyl.
Compound series 33, prepared and studied by Maekawa et al, demonstrate that the $^{19}$F NMR chemical shifts vary little for substituted (R = alkyl, aryl) trifluoroacetophenone derivatives. Therefore, the principal deshielding effect observed in this series is governed by the direct attachment of the benzene ring to the TFA group.

Compound 34 is an unusual example of a steric-structural topology deshielding effect on the $^{19}$F NMR $-\text{CF}_3$ chemical shift. The $-\text{CF}_3$ groups in the 2,2'-bis(trifluoroacetyl) biphenyl structure are evidently close enough in proximity to the biphenyl ring system to allow for greater deshielding of the fluorine nuclei than would normally occur.

Trifluoroacetylated heterocycles

Trifluoroacetylated heterocyclic compounds comprise a very important class of molecules, serving as synthons for natural products and numerous combinatorial processes important to the general chemical industry as well as the pharmaceutical and agrochemical industries. For the purposes of this review, we will limit our scope to heterocyclic molecules containing nitrogen and oxygen. Some representative examples are shown in Figure 14.

Fused bicyclic ring system examples include nitrogen-containing indole and benzotriazole derivatives 35–37. The nonaromatic heterobicyclic compound 35 has conjugation limited to the heterocyclic ring, resulting in a greater level of $-\text{CF}_3$ group shielding ($\delta = -74.3$ ppm) than that found for the heteroaromatic indole 36 ($\delta = -73.4$ ppm). The attachment of the TFA group on the heteroatom coupled with the aromaticity of the bicyclic ring system in compound 37 creates the largest deshielding of the $-\text{CF}_3$ group, causing an upfield shift of the $^{19}$F resonance of more than 3 ppm.

Single ring systems examined in this review include the pyridine, pyrrole, and pyrazole classes of heterocycles. The heteroaromatic nature of compound 38, 

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Figure 12 Cyclic trifluoroacetylated triketones. 
Abbreviations: $\text{CF}_3$, trifluoromethyl; DMSO, dimethyl sulfoxide; CDCl$_3$, deuteriochloroform; C$_6$F$_6$, hexafluorobenzene.

Figure 13 Trifluoroacetylated aromatics. 
Abbreviations: $\text{CF}_3$, trifluoromethyl; R, hydrogen, alkyl, acyl.
2-chloro-3-trifluoroacetylpyridine, deshields the $-\text{CF}_3$ group ($\delta = -73.4$ ppm) to a lesser extent than the TFA-substituted benzenes (compound series 33) discussed earlier. $^{19}F$ chemical shifts for the trifluoroacetylated pyrrole 39 and pyrazole 40 indicate a greater deshielding effect than found in 38, possibly a consequence of the reduction in ring size.

The oxygen-containing 2-trifluoroacetylfuran (41) $^{19}F$ chemical shift is similar to those of the pyrrole and indole compounds. The five-membered heteroaromatic ring systems evidently give rise to similar electronic effects on the $\text{CF}_3$ group. The upfield $^{19}F$ chemical shift (−3 ppm) of the nonaromatic dihydropyran derivative (42), which is more highly deshielded than the other heterocycles in this group, may be rationalized using resonance theory as shown in Figure 15. The extension of conjugation by delocalization of electron density in the resonance contributor may, in part, account for the observed upfield chemical shift observed in this case.

Conclusion

This review of nine major classes of TFA-containing molecules provides new comparisons of how $^{19}F$ NMR may be utilized in structural connectivity analyses of these fluorinated species. While the $^{19}F$ NMR chemical shifts for the $-\text{COCF}_3$ groups in this study range from approximately $-85$ ppm for the parent hexafluoroacetone to $-67$ ppm for certain TFA-bearing aromatics, the narrow band of variability can nonetheless yield valuable information to aid in understanding the structural topology, electronic environment, solvent, and concentration effects that are often examined by NMR spectroscopy.

The chemical shift variability observed for the $-\text{COCF}_3$ group results from the interplay of substrate structure, electronic environment, solvent polarity, and in a few important instances, concentration. Key structure-related influences include constraints imposed by the introduction of ring systems. Structural, electronic, and solvent effects can conspire to change tautomeric equilibria in TFA-substituted di- and triketones, producing enolic and keto forms for which the chemical shifts can vary from $-82$ ppm to $-68$ ppm ($\Delta\delta_F = -16$ ppm). Steric deshielding of fluorine, like that found in 2,2′-dinitrofluorodiphenyl, is a phenomenon that occurs when large groups in close proximity to one another disturb the p orbital electrons of the fluorine atom.

Distinct electronic effects arising from neighboring groups, conjugation, and resonance can also have substantial
deshielding effects on the $\text{–COCF}_3$ group. The presence of a multiple bond in conjunction with the TFA group produces a $\Delta \delta_f \approx -7$ ppm, while incorporation of an aromatic system conjugated with the TFA can extend the chemical shift range another 6 ppm. Heteroatoms present in carboxylic acids have a deshielding effect of similar magnitude brought about by resonance, often resulting in a $\Delta \delta_f \approx -7$ ppm.

Solvent effects on the $^{19}\text{F}$ chemical shift of the $\text{–COCF}_3$ group are normally $<2$ ppm, unless solvent polarity leads to a shift in tautomeric equilibria. Highly polar solvents cause an upfield shift of the $\text{–CO}^\text{F}$ $^{19}\text{F}$ NMR resonance. In cases where keto-enol=diketo or keto-enol=triketo equilibria are modulated by solvent polarity (as in the case of TFA-substituted $\beta$-diketones where the proportion of the diketo form increases with highly polar solvents), $\Delta \delta_f$ can range from $-5$–$9$ ppm.

The concentration of TFA-bearing $\beta$-diketones in solution can also influence the proportions of tautomers present. In dilute solutions the keto-enol tautomer is usually the only discernible species present, while the diketo form proportion can increase to $-10$% of the mixture composition in neat solutions of the $\beta$-diketone.

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### Disclosure

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