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$^{13}$C-Methyl isocyanide as an NMR probe for cytochrome P450 active sites

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Abstract:

The cytochromes P450 (CYPs) play a central role in many biologically important oxidation reactions, including the metabolism of drugs and other xenobiotic compounds. Because they are often assayed as both drug targets and anti-targets, any tools that provide: (a) confirmation of active site binding and (b) structural data, would be of great utility, especially if data could be obtained in reasonably high throughput. To this end, we have developed an analog of the promiscuous heme ligand, cyanide, with a $^{13}$CH$_3$-reporter attached. This $^{13}$C-methyl isocyanide ligand binds to bacterial (P450cam) and membrane-bound mammalian (CYP2B4) CYPs. It can be used in a rapid 1D experiment to identify binders, and provides a qualitative measure of structural changes in the active site.

Keywords: NMR, Heme, P450cam, CYP2B4, Screening.

Introduction

The cytochromes P450 (CYPs) are a family of heme-containing monooxygenase enzymes that use molecular oxygen to hydroxylate specific substrates, often with high stereoselectivity (Poulos and Johnson 2005). While CYPs are present in both eukaryotes and prokaryotes, the prokaryotic enzymes are soluble and therefore easier to study. A prototypical and well-studied bacterial CYP is P450cam (CYP101), from *Pseudomonas putida* (Katagiri et al. 1968; Poulos et al. 1986). P450cam is a 46 kDa soluble enzyme that catalyzes the 5-exo-hydroxylation of camphor (Gelb et al. 1982). A prototypical and well-studied mammalian CYP is CYP2B4. CYP2B4 is a 55 kDa membrane-bound protein from rabbit liver that either hydroxylates or demethylates a variety of foreign compounds, such as the drug benzphetamine (Kanaeva et al. 1992). The mammalian CYPs, being the enzymes predominantly responsible for drug and xenobiotic metabolism in humans, are of particular interest in the pharmaceutical industry. They are relatively large, membrane-bound proteins with a typical molecular weight of ~65 kDa (Furge and Guengerich 2006). Consequently, it is difficult to obtain structural information on the mammalian CYPs, either via X-ray or current multidimensional NMR techniques. Although, there have been some recent successes in crystallizing mammalian CYPs that lack the membrane-binding N-terminal region, and have other solubilizing mutations (Williams et al. 2000; Williams et al. 2003; Williams et al. 2004; Scott et al. 2003).
But, the mammalian CYPs are too large to structurally characterize with NMR, and membrane-bound CYPs have yet to be crystallized. Hence, there is a need to develop new techniques that rapidly probe at least part of the active site structure, to better define binding interactions with substrates or inhibitors.

With the above goal in mind, we have decided to capitalize on the fact that cyanide is a promiscuous ligand for heme iron (binding through its carbon), and that alkyl isocyanides (alkyl-NC) are known to be ligands for a number of CYPs (Tomita et al. 2001; Lee et al. 2001). Their absorption spectra are typical of Type II binding spectra, where the Soret band is shifted upon binding of the alkyl isocyanide, with absorption maxima occurring at either or both 430 and 455 nm, depending upon the extent of back-bonding from iron into the isonitrile triple bond. Indeed, resonance Raman experiments (Tomita et al. 2001; Lee et al. 2001) have established that the heme-bound isocyanide binds in two distinct orientations, linear or bent depending on C–Fe bond order; these forms are in slow exchange on the resonance Raman timescale, but would normally be in fast exchange on the NMR timescale. These two orientations are illustrated in Fig. 1, for the methyl isocyanide ligand that is the topic of this paper. The linear and bent forms are related to the two resonance forms of methyl isocyanide (Scheme 1), although the imine form is highly disfavored unless there is electron donation to stabilize it via formation of a dative bond (as in Fig. 1; in this case, electron donation is via backbonding from the iron).
Fig. 1 Methyl isocyanide bound to the heme iron. Schematic representation of methyl isocyanide bound to heme with (a) an iron-carbon bond order of two ("bent") and (b) an iron-carbon bond order of one ("linear"). The heme pyrrole rings, along with the CN triple (or double) bond, produce a highly anisotropic environment that can produce large chemical shift perturbations for the labeled methyl group.

Scheme 1 Resonance structures for methyl isocyanide

Alkyl isocyanides bind particularly strongly when the iron of the heme is in its reduced form ($K_d \sim 10^{-5}-10^{-6}$ M), although it can be shown that with increasing alkyl chain length, even the ferric form can
exhibit dissociation constants in the micromolar range (Simonneaux and Bondon 2000).

Given the established value of $^{13}$CH$_3$-methyl probes for NMR structural and dynamical studies of large proteins (Pellecchia et al. 2002; Showalter et al. 2007; Tugarinov et al. 2006), our interest in alkyl isocyanides is limited to methyl isocyanide ($^{13}$CH$_3$-NC). As the smallest member of the alkyl isocyanides, it is the one least likely to interfere with substrate or inhibitor binding (being only slightly larger than the native ligand, dioxygen). Also, the lack of adjacent methylene protons avoids dipolar relaxation effects, which would lead to line broadening. The $^{13}$C-label allows for selective observation of the attached methyl protons, which provides a chemical shift perturbation probe of structural (and potentially dynamical) changes in the binding pocket, due to binding of substrate or inhibitor. Moreover, because of the short internal correlation time of the rapidly rotating methyl group, even when bound to the heme, the proton linewidth should remain narrow—like those of a small molecule. Taken together, these features potentially allow structural and dynamic information to be extracted from NMR experiments such as: 2D $^1$H-$^{13}$C HSQC, $^{13}$C-filtered $^1$H 1D, $^{13}$C-edited (or filtered) NOESY, methyl dynamics (ex. using $^{13}$CHD$_2$, $^{13}$CH$_2$D (Showalter et al. 2007; Cavanagh et al. 2007), and various other experiments (Cavanagh et al. 2007). Given the large size of CYP enzymes, especially when complexed to micelles, most of these NMR experiments would not normally be feasible. Herein is presented the synthesis, initial characterization and application of the $^{13}$C-methyl isocyanide probe, to enable such studies. This probe will permit the first such targeted NMR studies of CYP binding sites, and provide evidence for active site changes near the heme, due to the presence or absence of substrates, inhibitors or other cellular components, especially lipids or electron donating protein partners.

**Materials and methods**

**CYP2B4, myoglobin and biochemical reagents**

Myoglobin (horse heart, >90% pure), benzphetamine, DEAE resin, and other reagents were from Sigma-Aldrich, unless specified otherwise. CYP2B4 was expressed in C41 E. coli cells using the pLW01-P450 2B4 mem plasmid, and purified as described previously (Miroux
and Walker 1996; Bridges et al. 1998). \(^{13}\)C-methyl iodide and \(d_8\)-glycerol were from Isotec, and \(d_{38}\)-DPC (dodecylphosphocholine-\(d_{38}\)) was from Cambridge Isotope Laboratories.

**Expression and purification of P450cam**

P450cam was expressed in *E. coli* (BL21DE3*) using a pCWori expression construct (a kind gift from Dr. Paul Ortiz de Montellano) and purified largely as described previously (Sibbesen et al. 1996; Yao et al. 2007), with some changes to the protocol. Briefly, 6 L of cell culture were grown at 37°C in 1 L batches to an OD\(_{600}\) of 0.7–0.9, before adding 1 mM IPTG (Isopropyl-\(\beta\)-D-Thiogalactopyranoside) for induction and 30 \(\mu\)M \(\delta\)-ALA (\(\delta\)-aminolevulinic acid) as a heme precursor, and allowing protein expression to continue for 20–22 h. After the cells were harvested they were frozen at –80°C for several hours, before resuspending in lysis buffer (50 mM Tris*HCl, pH 7.5 (room temp.), 25 mM KCl, 1–2 mM camphor, 1 mM EDTA, 1 mM PMSF, 1 IM leupeptin, 1 \(\mu\)M pepstatin, 32 U/ml DNase, 3 U/ml RNase, 1 mg/ml lysozyme) and sonicating. The cell debris was spun down at 50,000g and the supernatant was collected to be loaded onto a 50-ml bed volume DE-52 anion exchange column that had been pre-equilibrated with running buffer (50 mM Tris*HCl, pH 7.5 (4°C), 25 mM KCl, 2 mM camphor). After washing with two column volumes of running buffer, protein was eluted using a linear gradient from 0 to 100% of elution buffer (50 mM Tris*HCl, pH 7.5 (4°C), 300 mM KCl, 2 mM camphor). All fractions with an \(R_z\) ratio (\(A_{392}/A_{280}\)) > 0.5 were pooled and concentrated.

Ammonium sulfate was added to the pooled fractions to 25% saturation (~143 mg/ml), and they were then loaded onto a phenyl sepharose column (50 ml bed volume), after it had been pre-equilibrated with running buffer (50 mM Tris*Cl, pH 7.5, 50 mM KCl, 25% NH\(_4\)SO\(_4\), 100 \(\mu\)M camphor). After washing, elution was achieved with a linear gradient of 0 to 100% of elution buffer (50 mM Tris*HCl (pH 7.5), 50 mM KCl, 1 mM camphor). All fractions having an \(R_z\) ratio ≥ 1.4 were pooled, and purity was confirmed with an SDS gel (Supplementary Fig. S1). Protein (500 \(\mu\)M) was dialyzed into 30 mM phosphate buffer (pH 7.4), 100 mM KCl, 2 mM camphor, and flash frozen in 12 1-ml aliquots and stored at –80°C. Typical yield was ~ 100 mg/6 L cell culture.
Synthesis of $^{13}$C-methyl isocyanide ($^{13}$CH$_3$NC)

$^{13}$CH$_3$NC was synthesized with modifications to a previously reported procedure (Freedman and Nixon 1972). 1 mmol of $^{13}$C-methyl iodide and 2 mmol of silver cyanide were placed in a thick walled glass tube and sealed. The mixture was heated for 1 h at 95°C in an oil bath and the tube was then brought to room temperature slowly. The sealed tube was opened and the brown solid was quenched for 12 h in 5 ml of an 8 M KCN solution. The resultant mixture was distilled under vacuum to obtain crude $^{13}$C-methyl isocyanide, which was purified by bulb-to-bulb distillation. Yield was 21%, and the $^1$H NMR spectrum (Fig. 2) confirmed the structure. The pair of 1:1:1 triplets resulting from $^{13}$C-$^1$H scalar coupling ($^1$J coupling, producing the large doublet) and $^{14}$N-$^1$H scalar coupling ($^2$J coupling, producing the triplet) was only apparent in d$_4$-methanol and D$_2$O; in other solvents that were used, including d$_3$-acetonitrile, d$_1$-chloroform and d$_6$-benzene, the spectrum only showed the large doublet splitting from the $^{13}$C-$^1$H coupling. The proton chemical shift also varied depending upon the solvent used, and generally shifted upfield with increasing nonpolarity of the solvent (for example, the methyl protons had a chemical shift of 1.1 ppm in d$_6$-benzene).

![Fig. 2](image-url) NMR spectrum of $^{13}$C-methyl isocyanide in d$_4$-methanol. The $^{13}$C-coupled $^1$H resonances from $^{13}$CH$_3$NC flank the solvent resonance. The inset shows the further splitting (1:1:1 triplet) of the two $^{12}$C-split proton resonances. This triplet splitting is due to $^2$J coupling between
the quadrupolar $^{14}$N nucleus ($I = 1$) and the methyl protons. Spectrum acquired at 300 MHz

**NMR spectral analysis of $^{13}$CH$_3$NC bound to heme proteins**

All NMR spectra were collected at 25°C on a 600 MHz Varian spectrometer equipped with a triple-resonance cryoprobe. $^1$H-$^{13}$C HSQC spectra were collected with a spectral width of 12,000 and 28,000 Hz and digitized with 1920 and 256 complex points for the $^1$H and $^{13}$C dimensions, respectively. Spectra were referenced directly to H$_2$O ($\delta = 4.76$ ppm at 25°C) in the $^1$H dimension and indirectly in the $^{13}$C dimension. For the samples with P450cam as the heme protein, the protein concentration was 150 μM, as confirmed spectrophotometrically, using an extinction coefficient of 105 μM$^{-1}$ cm$^{-1}$ for the Fe$^{3+}$-substrate free protein with $\lambda_{max} = 418$ nm (O'Keeffe et al. 1978). Buffer conditions were 30 mM potassium phosphate (pH 7.4), 100 mM KCl, with saturating camphor (for samples that contained substrate) at 2 mM. When CYP2B4 was used as the heme protein, the concentration was 75 μM, determined using the extinction coefficient for the CYP2B4-CO complex of $\Delta\varepsilon_{450-490} = 91$ mM$^{-1}$ cm$^{-1}$ (Omura and Sato 1964). Buffer was 100 mM potassium phosphate (pH 7.4) with either 20% (v/v) d$_8$-glycerol or 50 mM d$_{38}$-DPC, with maximum benzphetamine concentrations (for samples that contained substrate) of 1.6 mM, close to its solubility limit at pH 7.4. All samples, after having argon gas bubbled through the solution for 15 min to purge any oxygen, were reduced with dithionite, followed by addition of an equimolar amount of $^{13}$CH$_3$NC to form the ferrous heme protein-$^{13}$CH$_3$NC complex.

**Results and discussion**

**Synthesis and characterization of $^{13}$CH$_3$NC: coupling to $^{14}$N and effects of solvent polarity**

The $^{13}$CH$_3$NC probe was prepared from $^{13}$C-methyl iodide and silver cyanide. Surprisingly, the proton NMR spectrum reveals that in some solvents (esp. polar solvents), a 2-bond coupling to the cyanide $^{14}$N atom is present. Such coupling to nitrogen is not typically observed
due to quadrupolar relaxation from $^{14}$N (which is $I = 1$). But the axial symmetry of the molecule, along with a very short correlation time, appears to be decreasing the extent of quadrupolar relaxation. This is because the quadrupolar relaxation rate is proportional to the correlation time, and also has a squared dependence on the asymmetry parameter $\eta = (E_{XX} - E_{YY})/E_{ZZ}$ (Nelson 2003), which should be zero ($E_{XX} - E_{YY} = 0$) for an axially symmetric molecule like CH$_3$NC. Perhaps in nonpolar solvents, where this coupling is no longer observed, there is some sort of intermolecular interaction between molecules that breaks this symmetry ($E_{XX} - E_{YY} \neq 0$), thereby restoring the quadrupolar relaxation effects that usually decouple $^{14}$N coupling. In this regard, it is interesting that gas phase (argon matrix) infrared spectroscopy studies (Freedman and Nixon 1972) have indicated that methyl isocyanide forms a dimer, whereby the large CN dipoles (see the isonitrile resonance structure in Scheme 1) align in an antiparallel orientation. Such pairing to neutralize dipole moments might also be favored in a nonpolar solvent, and could produce an upfield shift for the methyl protons, as shown in Scheme 2.

Scheme 2 Proposed methyl isocyanide dimer in nonpolar solvent

However, the situation could certainly be different in solution relative to the gas phase. One must also consider the possibility of
higher order structures, at least in solution, and even cyclic trimers that might form due to stabilizing dative bonds between molecules in the imine resonance form, shown in Scheme 1. Whatever the source of this interesting phenomenon, it is not relevant for protein studies, since the $^{14}\text{N}$ coupling is no longer observed when $^{13}\text{CH}_3\text{NC}$ is bound to heme iron.

**Application of the $^{13}\text{CH}_3\text{NC}$ probe to studies of a bacterial CYP: P450cam**

Initial studies were done with cytochrome P450cam (P450cam) which, being a soluble and well-studied P450 protein, serves as an excellent model system. In Fig. 3a is shown the $^1\text{H}-^{13}\text{C}$ HSQC spectrum of $^{13}\text{C}$-labeled methyl isocyanide ($^{13}\text{CH}_3\text{NC}$) in buffer, without protein present. The appearance of the spectrum is the same in the presence or absence of camphor, indicating that no significant interactions occur between substrate and probe in solution, in the absence of P450cam. The $^1\text{H}-^{13}\text{C}$ HSQC spectrum of $^{13}\text{CH}_3\text{NC}$ in the presence of an equimolar concentration of reduced (ferrous) P450cam (Fig. 3b) shows a large upfield shift due to heme binding. The signal of free $^{13}\text{CH}_3\text{NC}$ has disappeared and a new signal with a $^1\text{H}\delta$ of 1.2 ppm and $^{13}\text{C}\delta$ of 21.4 ppm has appeared. The upfield shift in both dimensions presumably reflects the fact that the methyl group of the probe, when bound to the heme-iron, now sits directly above the aromatic porphyrin ring, resulting in increased shielding of the relevant resonances due to ring current effects (Fig. 1). Figure 3c is a spectrum of the same sample shown in Fig. 3b, but now with excess $^{13}\text{CH}_3\text{NC}$, clearly showing that the free and bound forms are in slow exchange on the chemical shift timescale. Finally, the $^1\text{H}-^{13}\text{C}$ HSQC spectrum of the $^{13}\text{CH}_3\text{NC}/\text{Fe}^{2+}$-P450cam/camphor ternary complex (Fig. 3d) reveals two new major crosspeaks (and a third less intense resonance), distinct from both free and bound (-camphor). It appears therefore that camphor is either binding in two orientations, or that the probe itself occupies two major orientations (possibly linear and bent, but now in slow exchange on the NMR timescale, due to steric clash with camphor?). In any case, this experiment clearly identified camphor as binding in the active site, thereby validating the use of $^{13}\text{CH}_3\text{NC}$ for identifying such molecules in NMR-based screening.
**Fig. 3** P450cam $^1$H-$^{13}$C HSQC spectra showing (a) unbound $^{13}$CH$_3$NC, (b) stoichiometric $^{13}$CH$_3$NC bound to Fe$^{2+}$-P450cam, (c) $^{13}$CH$_3$NC bound to Fe$^{2+}$-P450cam in the presence of excess $^{13}$CH$_3$NC, and (d) $^{13}$CH$_3$NC bound to Fe$^{2+}$-P450cam in the presence of a saturating amount of camphor. The additional peaks in panels B, C, and D at 60–75 ppm are due to natural abundance glycerol $^1$H-$^{13}$C signal.

**Application of the $^{13}$CH$_3$ NC probe to studies of a mammalian CYP: CYP2B4**

The same binding studies were done with native, full-length CYP2B4 in 100 mM potassium phosphate buffer (pH 7.4) and 20% (v/v) d$_8$-glycerol (see Fig. S2). This time, upon addition of an equimolar amount of CYP2B4, the bound form of the $^{13}$CH$_3$NC probe had a $^1$H$\delta$ of 3.13 ppm and a $^{13}$C$\delta$ of 76.8 ppm. That is, there is now a dramatic downfield shift for carbon (but not hydrogen). When substrate (benzphetamine) was then added, the methyl crosspeak shifted to have a $^1$H$\delta$ of 3.13 ppm and a $^{13}$C$\delta$ of 49.5 ppm. The chemical shifts of the unbound $^{13}$CH$_3$NC probe (not shown) are the same as those reported above, in the absence of d$_8$-glycerol (Fig. 3a).
These data indicate that $^{13}$CH$_3$NC is an extremely sensitive chemical shift probe that can be used to identify occupancy of active sites for solubilized mammalian CYPs like CYP2B4.

While the above studies are of great value for identifying active site binders, they still suffer from the same limitations that most structural studies of mammalian CYPs do—they are of the enzyme in a relatively artificial state, unbound to lipid. Because CYP2B4 is normally bound to a phospholipid bilayer in vivo, we repeated the above experiments, but now in the presence of d$_{38}$-DPC micelles. In the control spectrum taken in the absence of both protein and substrate, there now appeared a new peak for the methyl of $^{13}$CH$_3$NC, in addition to the one previously observed for free $^{13}$CH$_3$NC (Fig. 3a). This new resonance is shifted upfield in the proton dimension with a $^1$H$\delta$ of 1.17 ppm, but is largely unchanged in the carbon dimension ($^{13}$C$\delta$ of 30.3 ppm). This new peak is likely due to $^{13}$CH$_3$NC that has been taken up into the nonpolar interior of the micelle, and is similar to the chemical shift change we saw for probe in nonpolar solvents like d$_6$-benzene (vide supra). When CYP2B4 protein is then added, one new peak appears, presumably from $^{13}$CH$_3$NC that binds the heme iron of micelle-bound CYP2B4. This peak is upfield shifted (Table 1). Finally, when benzphetamine was added, the intensity of this crosspeak decreased and three new peaks appeared with $^1$H$\delta$s of 2.64, 2.73 and 2.80 ppm and $^{13}$C$\delta$s of 37.5, 27.15 and 38.5 ppm, respectively (Table 1). These data suggest that $^{13}$CH$_3$NC-bound CYP2B4 can bind benzphetamine, and that multiple conformations exist for this CYP2B4:$^{13}$CH$_3$NC:benzphetamine ternary complex. Most importantly, these chemical shift changes are very different than what was observed for the free CYP2B4 (with glycerol present), indicating at least qualitatively that the heme is in a very different electronic state when CYP2B4 is bound to micelles versus when it is solubilized with glycerol. Currently, there are no structural methods to probe active site changes in mammalian CYPs, due to binding to lipid. Given that all current crystal structures are of free CYPs, it is therefore vitally important to determine if the active site structure changes in the presence of lipid.
Table 1 1H-13C HSQC crosspeak chemical shifts for various 13CH3NC complexes
*Buffer A: 100 mM KCl, 30 mM potassium phosphate, pH 7.4; Buffer B: 20% (v/v) glycerol, 100 mM potassium phosphate, pH 7.4; Buffer C: 50 mM DPC, 100 mM potassium phosphate, pH 7.4

Application of 13CH3NC probe using 1D spectra—towards higher throughput

The above studies establish the utility of the 13CH3NC probe for screening to identify molecules that bind in CYP active sites. But, 2D 1H-13C HSQC experiments are too time-consuming (1–4 h) for higher throughput screening efforts, especially if lower protein concentrations are used (≤150 μM). Fortunately, since only one crosspeak is being monitored, 2D experiments are not necessary. Instead, a much shorter 13C-filtered 1H 1D spectrum (Fig. 4) could be acquired, or a 13C
1D using a cryoprobe with cryocooled $^{13}$C preamp (Kovacs et al. 2005), depending upon whether the change of interest is occurring in the $^1$H or the $^{13}$C dimension. This, coupled with the fact that very little chemical probe is actually needed (~5 μg for a 0.5 ml sample of 250 μM protein), makes its use in a high-throughput assay even more attractive, as it should be able to provide reliable, fast and inexpensive information about a compound’s ability to bind to a given heme protein. Furthermore, advances in microflow probe technology (Olson et al. 2004) would permit assays to be done on volumes as low as 10 μl, making such screening even more feasible in terms of protein consumption (requiring only ~100 μg protein/assay, so ~10 mg to screen a 96-well plate of compounds), at least as a secondary assay. Of course, since protein concentration would be ~150 μM, one would need to have at least 50 nM of the compound to be screened present to see a significant effect on the methyl chemical shift. Finally, subsequent use of the $^{13}$CH$_3$NC probe to obtain structural information from NOE data will be limited to molecules that bind in fast exchange (i.e. using transferred NOE experiments), because it is usually not practical to obtain concentrated solutions of mammalian CYP enzymes, given their hydrophobic nature.

Fig. 4 $^{13}$C-filtered $^1$H 1D spectra of (a) unbound $^{13}$CH$_3$NC and (b) $^{13}$CH$_3$NC bound to Fe$^{2+}$-P450cam

**Interpretation of chemical shift changes**

Besides the qualitative information on a compound’s ability to bind to CYP active sites, the magnitude of the chemical shift perturbations could possibly be used to calculate a binding orientation for the probe relative to the central heme iron, and the largely planar
and aromatic porphyrin ring system (Fig. 1). Furthermore, anisotropy effects from a bound CYP substrate or inhibitor (which are often aromatic) could help to position it relative to probe. Indeed, chemical shift values are increasingly being used to refine protein structures (Xu and Case 2001). Unfortunately, these algorithms are not adequately parameterized at present to deal properly with the highly anisotropic effects from the CH$_3$NC triple bond, the heme pyrrole rings and the diverse array of aromatic ring systems likely to be present in CYP substrates or inhibitors.

At one level, since it is reasonable to assume the dominant resonance structure of $^{13}$CH$_3$NC free in solution is the isonitrile (Scheme 1), then any deviation of the $^1$H$\delta$ from that of its unbound solution form must be due to effects from the heme anisotropy and/or from increasing the population of the bent orientation. In the case of CYP2B4 in 20% glycerol, the $^1$H$\delta$ remained the same as that of the unbound probe, both in the absence and presence of saturating amounts of substrate, while the $^{13}$C$\delta$ was first shifted dramatically downfield ~45 ppm and then back upfield ~25 ppm, still distant from its original position. One possible interpretation of these unusual shifts is that, due to the structural/steric constraints of the binding pocket (or heme electronic changes leading to more backbonding), the probe is forced to adopt a more bent orientation relative to the porphyrin ring. It is surprising, though, that the methyl protons wouldn't shift downfield like the methyl carbon does, if the change is simply due to a more bent orientation. The reasons for these large chemical shift changes are not entirely clear, but will be a function of: (a) the high level of anisotropy coming from the heme pyrrole rings and the cyanide triple bond (Fig. 1), (b) the extent of backbonding from iron (a function of heme electronic state), which would alter the relative population of the linear and bent forms of the probe, (c) the physical perturbation/steric clash between a substrate or inhibitor and the $^{13}$CH$_3$NC probe, which could also alter the relative population of linear and bent forms (and its position in this highly anisotropic environment), and (d) any anisotropic effects from the substrate/inhibitor itself, if it has aromatic rings as benzphetamine does. The combination of these effects would make any structure-based calculation of chemical shift changes difficult at present, but clearly the probe's unique position in a highly anisotropic environment makes it exquisitely sensitive to even modest structural or electronic
perturbations, and thereby makes it a very sensitive chemical shift probe. As a final point on structural inferences that could be made, it is tempting to speculate that when the probe gives two new crosspeaks for a complex with substrate or inhibitor, this might indicate a steric clash is occurring, which slows the exchange between the linear and bent structures (Scheme 1 and Fig. 1). The data with camphor bound to P450cam and benzphetamine bound to micelle-bound CYP2B4 are consistent with this interpretation, but more experiments are needed to prove that this is in fact the source of the multiple crosspeaks.

**Application of the$^{13}$CH$_3$ NC probe to other heme proteins: myoglobin**

Regardless of the actual underlying structural changes occurring within these CYP complexes, and the source of the chemical shift changes, the NMR data do indicate that the $^{13}$CH$_3$NC probe will be useful as a screening tool for monitoring active site changes in CYPs. But, might the probe also be useful for exploring other heme proteins, beyond the CYPs? As a first step towards answering this question, spectra of $^{13}$CH$_3$NC bound to 500 μM myoglobin (a non-P450 heme protein) have been obtained (Fig. S3), and they show a similar upfield chemical shift in both $^1$H and $^{13}$C dimensions. This suggests the probe might be useful for studying other heme protein active sites as well.

In summary, we have developed $^{13}$CH$_3$NC as the first NMR-based probe of cytochrome P450 active sites. This probe binds to both bacterial and mammalian P450s, even membrane-bound P450s, and may have broader utility with other heme proteins. It can be used to detect structural and electronic changes in active sites. In particular, it can be used to screen for P450 substrates or inhibitors using a reasonably high throughput assay, by acquiring a simple $^{13}$C-filtered 1D spectrum and monitoring chemical shift changes for the methyl protons. Since this could be done with microflow NMR (ex. Protasis CapNMR) at volumes as low as ~10 μl, the assay would consume only ~100 μg protein, making it reasonably sensitive for an NMR assay.
Supplementary Material

Supplementary Data

Figure S1. SDS-PAGE gel electrophoretic analysis of P450cam expressed in E. coli. Lane 1 contains crude cell lysate; lanes 2 & 4 contain the final purified protein (Rz = 1.4) with protein concentration in lane 4 two-fold higher than that in lane 2; lane 3 contains pooled protein fractions eluted after the first purification step (DE52 column); and lane 5 contains standards with molecular weights of 15, 25, 35, 50, 75, 100, and 150 kDa, respectively (bottom to top).

Figure S2. CYP2B4 ¹H-¹³C HSQC spectra showing (A) ¹³CH₃NC bound to Fe²⁺-CYP2B4, or (B) ¹³CH₃NC bound to Fe²⁺-CYP2B4 in the presence of 1.6 mM benzphetamine. The additional peaks are due to natural abundance glycerol signal.
Figure S3. (A) $^1$H-$^{13}$C HSQC and (B) $^{13}$C-filtered $^1$H 1D spectrum of excess $^{13}$CH$_3$NC bound to Fe$^{2+}$-myoglobin.

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Abbreviations

δ-ALA δ-Aminolevulinic acid  
cam Camphor  
CYP Cytochrome P450  
DEAE Diethylaminoethyl  
DPC Dodecylphosphocholine  
EDTA Ethylenediaminetetraacetic acid  
HSQC Heteronuclear single quantum coherence  
IPTG Isopropyl-β-D-thiogalactopyranoside
PMSF  Phenylmethanesulfonyl fluoride  
Tris  tris(hydroxymethyl)aminomethane  

Footnotes

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