

# Hydrogen Bonding in Human Manganese Superoxide Dismutase Containing 3-Fluorotyrosine

Idelisa Ayala,\* J. Jefferson P. Perry,<sup>†</sup> Jan Szczepanski,<sup>‡</sup> John A. Tainer,<sup>†</sup> Martin T. Vala,<sup>‡</sup> Harry S. Nick,<sup>§</sup> and David N. Silverman\*

\*Department of Pharmacology, University of Florida, Gainesville, Florida 32610; <sup>†</sup>Department of Molecular Biology, The Scripps Research Institute, La Jolla, California 92037; <sup>‡</sup>Department of Chemistry and Center for Chemical Physics, University of Florida, Gainesville, Florida 32611; and <sup>§</sup>Department of Neuroscience, University of Florida, Gainesville, Florida 32510

**ABSTRACT** Incorporation of 3-fluorotyrosine and site-specific mutagenesis has been utilized with Fourier transform infrared (FTIR) spectroscopy and x-ray crystallography to elucidate active-site structure and the role of an active-site residue Tyr<sup>34</sup> in human manganese superoxide dismutase (MnSOD). Calculated harmonic frequencies at the B3LYP/6-31G\*\* level of theory for L-tyrosine and its 3-fluorine substituted analog are compared to experimental frequencies for vibrational mode assignments. Each of the nine tyrosine residues in each of the four subunits of the homotetramer of human MnSOD was replaced with 3-fluorotyrosine. The crystal structures of the unfluorinated and fluorinated wild-type MnSOD are nearly superimposable with the root mean-square deviation for 198  $\alpha$ -carbon atoms at 0.3 Å. The FTIR data show distinct vibrational modes arising from 3-fluorotyrosine in MnSOD. Comparison of spectra for wild-type and Y34F MnSOD showed that the phenolic hydroxyl of Tyr<sup>34</sup> is hydrogen bonded, acting as a proton donor in the active site. Comparison with crystal structures demonstrates that the hydroxyl of Tyr<sup>34</sup> is a hydrogen bond donor to an adjacent water molecule; this confirms the participation of Tyr<sup>34</sup> in a network of residues and water molecules that extends from the active site to the adjacent subunit.

## INTRODUCTION

Superoxide dismutases catalyze the disproportionation of superoxide anion radicals to molecular oxygen and hydrogen peroxide and are classified according to the metal cofactor found in the active site (1,2). Human manganese superoxide dismutase (MnSOD) is a homotetramer of 22 kDa subunits (3) located in mitochondria, where it provides protection against oxidative damage. The crystal structure of human MnSOD shows an apparently hydrogen-bonded network of side chains and water that extends from the manganese-bound solvent molecule at the active site to solvent-exposed residues and to the interface between subunits (4). This network has been suggested to support proton transfer in catalysis (5). In the network, the side chain of Gln<sup>143</sup> appears to form hydrogen bonds with the manganese-bound solvent and with the side-chain hydroxyl of Tyr<sup>34</sup>. The crystal structure shows an ordered water molecule that also appears to form a hydrogen bond with the hydroxyl of Tyr<sup>34</sup> (4). The replacement of Tyr<sup>34</sup> with Phe (Y34F) has only minor effects on catalysis at low substrate concentrations (6,7); however, near maximal velocity the value of  $k_{\text{cat}}$  for Y34F is 10-fold lower than for wild-type (7). The ionization of the phenolic side chain of Tyr<sup>34</sup> has been identified as the origin of the pH dependence of several spectroscopic and kinetic properties of Mn(III)SOD (7,8).

In this report, Fourier transform infrared (FTIR) spectroscopy and x-ray crystallography have been applied to human

MnSOD to better understand the role of Tyr<sup>34</sup>. Each of the nine tyrosine residues in each subunit of wild-type MnSOD was replaced with 3-fluorotyrosine to help identify vibrational modes associated with tyrosine side chains. To identify vibrational modes that are specific to Tyr<sup>34</sup>, the FTIR spectrum of wild-type is compared with that of Y34F MnSOD. Our FTIR data show distinct vibrational modes arising from fluorinated tyrosine; based on the frequency of the C-O stretching vibrations in wild-type and Y34F MnSOD, the C-O stretching vibration of the phenolic group of Tyr<sup>34</sup> has been identified. The frequency of this absorption is consistent with a phenolic hydroxyl that is hydrogen bonded acting as a proton donor. These data complement the crystal structure and confirm the participation of Tyr<sup>34</sup> in the hydrogen-bonded network of residues and water molecules that extends from the active site to the adjacent subunit.

## MATERIALS AND METHODS

### Reagents

L-tyrosine and 3-fluorotyrosine were purchased from Sigma (St. Louis, MO) and Acros Organics (a subdivision of Fisher Scientific, Pittsburgh, PA).

### MnSOD expression, purification, and fluorotyrosine labeling

*Escherichia coli* cells that express wild-type and Y34F human MnSOD were grown in enriched and minimal media. For minimal media growth, the overnight culture was grown for 17 h at 37°C in 50 mL of minimal media until an OD<sub>595</sub> = 1 was reached. The minimal medium (M9) consisted of 0.04 M Na<sub>2</sub>HPO<sub>4</sub>, 0.02 M KH<sub>2</sub>PO<sub>4</sub>, 8.6 mM NaCl, and 0.02 M NH<sub>4</sub>Cl. The overnight culture was supplemented with 0.1 mM CaCl<sub>2</sub>, 1 mM MgSO<sub>4</sub>,

Submitted February 2, 2005, and accepted for publication August 11, 2005.

Address reprint requests to David Silverman, Box 100267, Health Sciences Center, University of Florida, Gainesville, FL 32610-0267. Tel.: 352-392-3556; Fax: 352-392-9696; E-mail: silvrnm@ufl.edu.

© 2005 by the Biophysical Society

0006-3495/05/12/4171/09 \$2.00

doi: 10.1529/biophysj.105.060616

0.2% glucose, 1  $\mu\text{g/mL}$  of thiamine, 0.2  $\text{mg/mL}$  of amino acids (except the aromatic amino acids), 1  $\text{mM}$  tryptophan, 1  $\text{mM}$  phenylalanine, and ampicillin. The glucose, thiamine, amino acids, tryptophan, and phenylalanine solutions were filter sterilized through a 0.2  $\mu\text{m}$  Corning filter system unit. The  $\text{CaCl}_2$  and  $\text{MgSO}_4$  were autoclaved. The overnight growth was then transferred to 11 L of minimal media and supplemented in the same manner as the overnight culture plus the addition of sterile 18  $\mu\text{M}$   $\text{MnSO}_4$ . The cells were allowed to grow for  $\sim 5$  h until an  $\text{OD}_{595}$  of 0.3–0.4 was reached. At this point, the cells were induced with 0.3  $\text{mM}$  isopropyl-beta-D-thiogalactopyranoside supplemented with 1  $\text{mM}$  3-fluorotyrosine (or unlabeled L-tyrosine as a control) and allowed to grow for an additional 4 h. Due to the low solubility of L-tyrosine and its fluorinated analog in water, these compounds were added as solids to the growing media. The cells were placed at  $4^\circ\text{C}$  overnight and harvested the next day by centrifugation. The resulting pellet was frozen at  $-70^\circ\text{C}$  until lysis was performed. The cells were lysed as previously described (9). The resulting protein solution was filtered, and fast protein liquid chromatography was performed to further separate MnSOD from some cellular proteins still present after the heating and dialysis steps. MnSOD was eluted at  $\sim 40$   $\text{mM}$  NaCl in the presence of 10  $\text{mM}$  Tris-HCl pH 8.2. The purified protein was concentrated and dialysed extensively against the final buffer 20  $\text{mM}$  potassium phosphate at pH 7.8. The content of Mn in MnSOD was determined using flame atomic absorption spectroscopy, and the concentration of MnSOD for kinetic study was taken as the manganese concentration.

Fluorotyrosine was incorporated into 67% of all tyrosine sites in human MnSOD, determined by amino acid analysis composition (Protein Chemistry Laboratory, Texas A&M University, College Station, TX), and corroborated by MALDI-TOF mass spectrometry.

## FTIR spectroscopy

A Midac (Costa Mesa, CA) FTIR spectrometer equipped with a deuterated triglycine sulfate detector was utilized to obtain infrared spectra of fluorinated and unlabeled wild-type and Y34F human MnSOD. FTIR data were acquired on samples prepared in a uniform manner; each contained 30  $\mu\text{L}$  of protein at 0.5  $\text{mM}$  in 20  $\text{mM}$  potassium phosphate at pH 7.8 that were dehydrated on a ZnSe window by applying a stream of dry nitrogen gas at room temperature. FTIR data were also collected on 1  $\text{mM}$  L-tyrosine and 1  $\text{mM}$  3-fluorotyrosine in 20  $\text{mM}$  potassium phosphate at pH 7.8. The samples were placed in the sample compartment, which was evacuated through a rotary pump to eliminate water vapor. The spectral parameters were as follows: apodization, Happ-Genzel; resolution, 4  $\text{cm}^{-1}$ ; number of scans, 215; levels of zero filling, one; and temperature, room temperature. The spectra acquired from protein samples were corrected for an amide I vibration of 0.35 to allow direct comparison. The data shown here represent an average of 860–1075 scans and 4–5 different samples.

## Crystallography

Orthorhombic crystals of fluorinated wild-type and Y34F human MnSOD were grown by vapor diffusion from protein solutions at 26  $\text{mg/mL}$ , buffered in 20  $\text{mM}$  potassium phosphate buffer at pH 7.8 and 22% poly(ethylene glycol) (PEG) 2000 monomethyl ether. X-ray diffraction data collection was conducted at 100 K on flash-frozen single crystals, supplemented with 20% ethylene glycol cryoprotectant in their mother liquor. Data sets were collected at beamline 8.3.1 of the Advanced Light Source, Lawrence Berkeley National Laboratory (Berkeley, CA) on a Q315 ADSC CCD detector. Data sets for the fluorinated wild-type and Y34F MnSOD were collected to 1.85  $\text{\AA}$  and 1.53  $\text{\AA}$ , respectively. Data sets were indexed and merged with Denzo and scaled with Scalepack (10). The fluorinated wild-type crystal belongs to space group  $P2_12_12$  with unit cell dimensions of  $a$ ) 74.0  $\text{\AA}$ ,  $b$ ) 75.4  $\text{\AA}$ , and  $c$ ) 67.8  $\text{\AA}$ . The data set was 97.8% complete (94.7% for the highest-resolution shell (1.85–1.97  $\text{\AA}$ ); 32,265 unique reflections measured) with an  $R_{\text{sym}}$  of 6.7% (30.8% for the highest-resolution shell).

The data have an overall  $I/\sigma I$  of 37.1 with an  $I/\sigma I$  in the highest resolution shell of 6.2. The fluorinated mutant Y34F crystal also belongs to space group  $P2_12_12$ , and has unit cell dimensions of  $a$ ) 73.7  $\text{\AA}$ ,  $b$ ) 75.3  $\text{\AA}$ , and  $c$ ) 68.0  $\text{\AA}$ . The data set was 99.9% complete (94.7% for the highest-resolution shell (1.58–1.53  $\text{\AA}$ ); 57,452 unique reflections measured) with an  $R_{\text{sym}}$  of 5% (28.4% for the highest-resolution shell). The data have an overall  $I/\sigma I$  of 57.9 with an  $I/\sigma I$  in the highest resolution shell of 5.3. Phases were obtained for both structures by molecular replacement against the wild-type human MnSOD structure using AMoRe (11). The two structures were refined with several cycles of rigid body and restrained refinement, using Crystallography and NMR Systems, version 1.1 (CNS) (12). Final refinement steps for the higher resolution Y34F-fluorinated MnSOD were completed using SHELX (13). Side-chain orientations were corrected where required, due to potential van der Waals clashes of their hydrogen atoms, using Reduce and Probe (14,15). The structures were fit against calculated  $2F_o - F_c$ ,  $F_o - F_c$  and composite omit electron density maps in the Xfit module of XtalView (16). The fluorinated wild-type MnSOD structure was refined to an  $R_{\text{work}}$  of 21.2% and an  $R_{\text{free}}$  (17) of 25.2% (with 5% of the reflections flagged for the test set) and the Y34F to a  $R_{\text{work}}$  of 15.3% and an  $R_{\text{free}}$  of 20.3%. Coordinates for the structures are deposited in the Protein Data Bank with accession numbers 1XDC for wild-type fluorinated MnSOD and 1XIL for the fluorinated Y34F mutant MnSOD.

## Density functional theory calculations

Geometry optimization and harmonic frequencies were calculated for L-tyrosine and 3-fluorotyrosine and their modified structures with density functional theory (DFT) applying the hybrid Becke3-Lee-Yang-Parr (B3LYP) exchange-correlation function and 6-31G\*\* basis set under the Gaussian 98 suite of programs (18). This level of theory has been demonstrated to be reliable in predicting vibrational frequencies and relative intensities of medium-sized polycyclic aromatic hydrocarbons (19). Zero point-corrected total energies were used to search for the global energy minima of the potential surfaces. A factor of 0.978 was used to scale the harmonic frequencies and zero point energies to account for anharmonicity effects and basis set deficiency. Such a factor was adopted earlier for neutral and cationic fluorene (a medium-sized polycyclic aromatic hydrocarbon) and found to give good agreement between calculated and observed mode frequencies (19).

Modeling of the atomic displacements in the various vibrational modes of the optimized molecular structures was viewed using AniMol software (Innovative Software, Gainesville, FL).

## RESULTS

### FTIR spectra of tyrosine and 3-fluorotyrosine

The FTIR absorption spectra of uniformly hydrated tyrosine and 3-fluorotyrosine solutions in phosphate buffer at pH 7.8 were measured at room temperature (Figs. 1 and 2). Only the 1200–1900  $\text{cm}^{-1}$  region is shown due to the broad OH band at 3000  $\text{cm}^{-1}$  and large phosphate contribution below 1200  $\text{cm}^{-1}$ . No clear band for the bending vibration of liquid water (at 1650  $\text{cm}^{-1}$ ) is observed in our experimental spectrum (Fig. 1 A). This indicates that there is no significant non-bonded (i.e., free) water left in our sample. The excellent agreement of the observed IR spectrum with that calculated for hydrogenated tyrosine, as displayed in Fig. 1 B, indicates our samples are nearly fully hydrated. (In our calculations we hydrogenated a carboxyl oxygen to simulate the presence of a water molecule hydrogen bonded to that group.) In

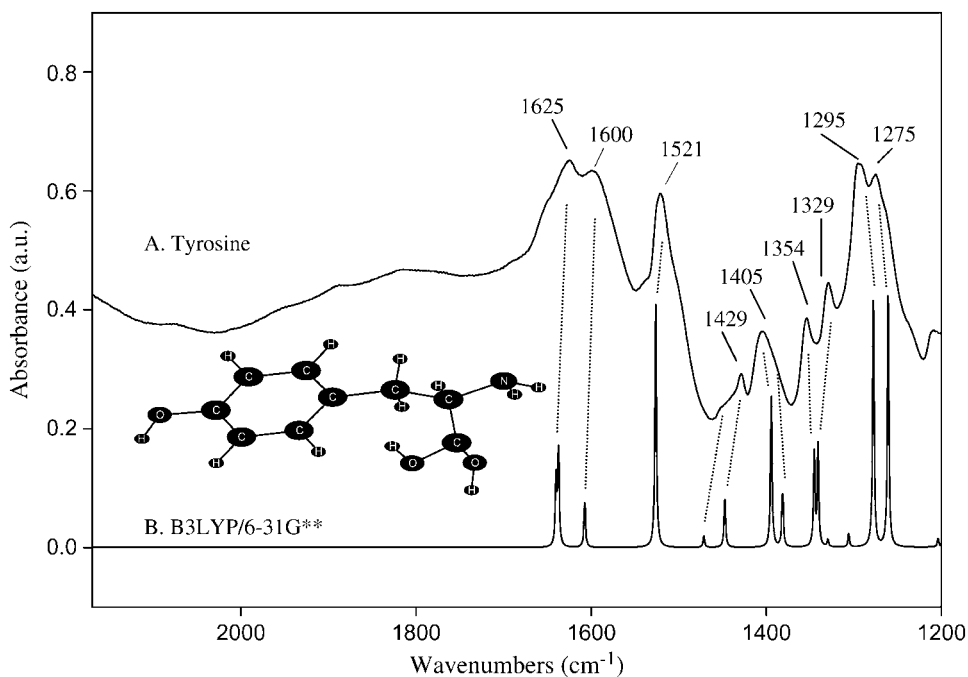


FIGURE 1 (A) Experimental room temperature infrared absorption spectrum of a partially hydrated sample of tyrosine in phosphate buffer (at pH 7.8) and room temperature, and (B) calculated (B3LYP/6-31G\*\*) infrared absorption spectrum of hydrogenated tyrosine at optimized (displayed) geometry. In the calculated spectrum, the harmonic band frequencies were scaled uniformly by a factor of 0.978. Bands of  $3\text{ cm}^{-1}$  bandwidth were constructed using 50% Gaussian and 50% Lorentzian functions. The vertical lines mark the proposed band assignments collected in Table 1.

addition, the  $3300\text{ cm}^{-1}$  peak, which contains contributions from the OH stretch of water (as well as the NH stretch from the protein) is 0.3 absorbance units in all the spectra. This suggests uniformity of hydration.

The lack of any strong bands in the  $1700\text{--}1800\text{ cm}^{-1}$  region indicates the absence of significant  $\text{C}=\text{O}$  stretching vibration from the carboxylic group of the tyrosines, consistent with ionization to carboxylate at the pH of 7.8 of the original solution. To mimic this in our calculations, hydrogen atoms were placed in proximity to the carboxyl

groups in the tyrosines, as displayed in Fig. 1. The figure also shows a comparison of the calculated and observed FTIR spectra for hydrogenated tyrosine. All bands in the experimental spectrum are now readily assignable, based on agreement of band frequencies and relative intensities. Table 1 gives a similar comparison for hydrogenated 3-fluorotyrosine.

A symmetric C-C ring-stretching vibration is assigned to the  $1625\text{ cm}^{-1}$  tyrosine band (and the  $1606\text{ cm}^{-1}$  band in 3-fluorotyrosine) (Figs. 1 and 2). DFT computations place this mode at  $1637\text{ cm}^{-1}$  in tyrosine (and  $1638\text{ cm}^{-1}$  in 3-fluorotyrosine). The asymmetric C-C ring-stretching mode is calculated at  $1607\text{ cm}^{-1}$  for tyrosine (and  $1615\text{ cm}^{-1}$  for fluorotyrosine) and observed at  $1600\text{ cm}^{-1}$  (tyrosine) and  $1587\text{ cm}^{-1}$  (3-fluorotyrosine; Table 1). Note that this region also has a contribution from the carboxylate anion near  $1600\text{ cm}^{-1}$ . In the case of 3-fluorotyrosine, a strong coupling of the C-F stretch oscillator to the C-C stretch ring oscillators was identified from viewing the atom motion during vibrations. Also, the electronegative F-atom causes a change in the atomic charge distribution in phenol, which likely affects the interaction of the phenol ring with the surrounding molecular bath (vide infra). These effects account for the change in the observed C-C stretch frequencies in 3-fluorotyrosine. However, in the B3LYP harmonic frequency prediction, the modes coupling and interaction with the surrounding bath are not incorporated into the calculation. This omission certainly affects the calculated band frequencies for the C-C stretch ring modes, causing higher predicted frequencies than observed (Table 1).

Both tyrosines are predicted to have a symmetric ring CC/CH stretching mode at  $1526\text{ cm}^{-1}$  (Table 1). A prominent band is observed at  $1521\text{ cm}^{-1}$  for tyrosine ( $1516\text{ cm}^{-1}$  for

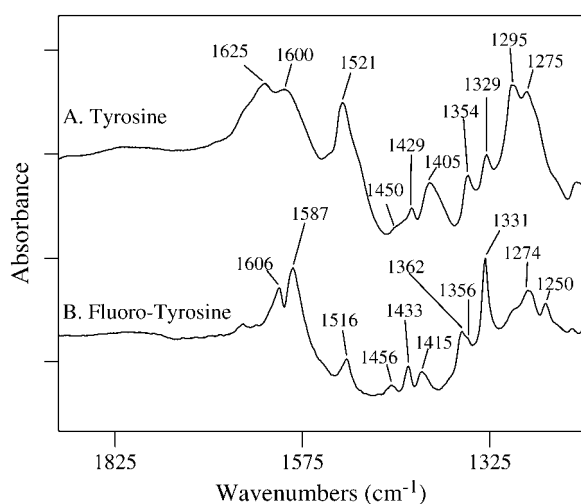


FIGURE 2 Infrared absorption spectra of uniformly hydrated samples of (A) tyrosine and (B) 3-fluorotyrosine. FTIR data were collected on 1 mM L-tyrosine and 1 mM 3-fluorotyrosine in 20 mM phosphate buffer at pH 7.8 at  $20^\circ\text{C}$ . The tick marks on the y axis represent 0.3 absorbance units.

**TABLE 1** Infrared absorption band frequencies (in  $\text{cm}^{-1}$ ) observed and calculated (B3LYP/6-31G\*\*) for tyrosine (Figs. 1 A and 2 A) and 3-fluorotyrosine (Fig. 2 B)

Mode description*	Tyrosine		3-Fluorotyrosine	
	Observed	Calculated <sup>†</sup>	Observed	Calculated <sup>†</sup>
HNH bend + CC symmetric ring stretch	1625	1640 + 1637 (0.71)	1606	1643 + 1638 (0.20)
CC asymmetric ring stretch	1600	1607 (0.18)	1587	1615 (0.12)
CC/CH symmetric ring stretch	1521	1526 (0.98)	1516	1526 (1.00)
HC <sub>α</sub> H bend	1450sh	1471 (0.05)	1456	1467 (0.03)
CC/CH asymmetric ring stretch	1429	1447 (0.20)	1433	1450 (0.14)
C <sub>β</sub> C <sub>γ</sub> stretch	1405	1394 (0.60)	1415	1393 (0.29)
HC <sub>β</sub> C <sub>γ</sub> bend	~1390sh	1381 (0.19)	1362	1378 (0.09)
CC asymmetric ring stretch	1354	1345 (0.38)	1356sh	1375 (0.12)
HC <sub>α</sub> C <sub>β</sub> bend	1329	1340 (0.39)	1331	1338 (0.19)
C <sub>1</sub> C <sub>α</sub> H <sub>α</sub> bend	–	1329 (0.02)	–	1320 (0.06)
C <sub>1</sub> C <sub>α</sub> H <sub>α</sub> bend	–	1305 (0.05)	–	1304 (0.24)
FC stretch	–	–	1293sh	1282 (0.20)
CO phenolic stretch	1275	1277 (0.99)	1250	1246 (0.65)
HC <sub>α</sub> H/HC <sub>β</sub> C <sub>γ</sub> /HOC <sub>γ</sub> bend	1295	1260 (1.00)	1274	1256 (0.52)

\*Major vibrations are noted only.

<sup>†</sup>Frequencies are scaled uniformly by a factor of 0.978. Relative integral intensities are given in parentheses.

fluorotyrosine) and is assigned to the symmetric ring CC/CH mode. Although higher in frequency than previously observed ( $1500 \text{ cm}^{-1}$ ) (20), this discrepancy could be due to different experimental conditions (see below). A low intensity HC<sub>α</sub>H bending mode is expected at  $\sim 1471 \text{ cm}^{-1}$  for tyrosine (and  $\sim 1467 \text{ cm}^{-1}$  for fluorotyrosine). A weak shoulder at  $1450 \text{ cm}^{-1}$  in tyrosine and more pronounced, though still weak, band in fluorotyrosine at  $1456 \text{ cm}^{-1}$  is assigned to this bending mode. These results are in agreement with the literature values for ring modes of the phenolic side chain of tyrosine (21) but are significantly different than ring modes in tyrosinate (20) and phenolate anionic species (22).

Several bands in the  $1430\text{--}1330 \text{ cm}^{-1}$  region arise from additional ring and carboxyl group vibrations as well as from modes containing the C<sub>α</sub>, C<sub>β</sub>, and C<sub>γ</sub> atoms. They are observed at 1429, 1405, 1390 (shoulder), 1354 and  $1329 \text{ cm}^{-1}$  for tyrosine (Figs. 1 A and 2 A), and 1433, 1415, 1362, 1356 (shoulder), and  $1331 \text{ cm}^{-1}$  for fluorotyrosine (Fig. 2 B).

The C-O stretching vibration of the phenolic group of tyrosine appears at  $1275 \text{ cm}^{-1}$  (Figs. 1 A and 2 A); this is characteristic of protonated phenols with the O-H group involved in hydrogen-bonding interactions and acting as a proton donor (21). In fluorotyrosine, this band is shifted significantly to lower frequencies (to  $1250 \text{ cm}^{-1}$ ). Substitution of an *o*-hydrogen atom with the more electronegative fluorine atom is expected to redshift the C-O stretch of the phenolic group, an expectation fully consistent with our DFT calculations, which predict a downshift of  $31 \text{ cm}^{-1}$  (observed shift:  $25 \text{ cm}^{-1}$ ). Modeling of the atomic displacements for this mode shows that the phenol C-O stretch requires a substantial ring deformation involving F atom movement as well. The large mass difference between fluorine and hydrogen is probably partially responsible for the lowering of the phenol C-O stretching vibrational frequency.

The spectrum of fluorotyrosine (Fig. 2 B) shows an additional band at  $\sim 1293 \text{ cm}^{-1}$  (shoulder), which is assigned to a ring mode that includes a large C-F displacement. As expected, this band is not observed in L-tyrosine.

### FTIR spectra of unlabeled and fluorinated wild-type human MnSOD

To identify the vibrational modes that arise from the tyrosine moieties, human MnSOD was prepared with each of the nine tyrosine residues in each subunit replaced with 3-fluorotyrosine. Fig. 3 A (*solid line*) shows the FTIR spectrum of unlabeled wild-type MnSOD in phosphate buffer at pH 7.8; the fluorinated version is shown superimposed as a dotted line. A difference spectrum, given in Fig. 3 B, shows only those vibrational modes that are sensitive to fluorination. Vibrations not altered in frequency or intensity upon fluorination will cancel and not be observed. Any differences in environment of the 3-fluorotyrosine residues will influence their normal modes, and thus affect the IR spectrum.

The difference spectrum shows the vibrational modes of unlabeled tyrosine as positive bands and those from fluorinated tyrosine as negative bands. Significant changes are observed in the aromatic ring-stretching regions around  $1600 \text{ cm}^{-1}$ . The amide I (C = O stretching of the backbone) absorbs at  $\sim 1650 \text{ cm}^{-1}$  and is expected to contribute absorption in this region if there are changes in the secondary structure of the protein upon fluorination. A positive line at  $1614 \text{ cm}^{-1}$  can be assigned to the symmetric CC stretching vibration of unlabeled tyrosine. This mode was observed at  $1625 \text{ cm}^{-1}$  for the model compound (Table 1). Upon fluorination of the tyrosines in MnSOD, this mode shifts to  $1605 \text{ cm}^{-1}$  and is shown as a negative band in Fig. 3 B. The asymmetric CC ring-stretching vibration for unlabeled tyrosine occurs most probably at  $1593 \text{ cm}^{-1}$ . The band

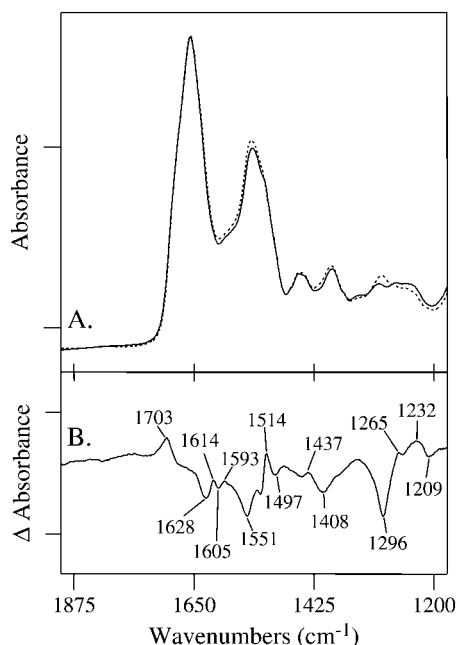


FIGURE 3 Infrared absorption spectra of (A, *solid line*) unlabeled human wild-type MnSOD, and (A, *dotted line*) wild-type MnSOD in which each tyrosine residue is labeled with fluorine at the 3-position of the phenol ring. The tick marks on the y axis represent 0.2 absorbance units. The difference spectrum (unlabeled-minus-fluorinated) is shown in B. The tick marks on the y axis represent 0.03 absorbance units. FTIR data were acquired at 20°C on 30  $\mu$ L of protein sample at 0.5 mM in 20 mM phosphate buffer at pH 7.8 and partially dehydrated onto a ZnSe window.

resulting from fluorination is not evident. The negative 1628 and 1551  $\text{cm}^{-1}$  bands are assigned to the amide I and amide II groups, respectively. These modes may indicate small changes in the secondary structure of MnSOD upon fluorination. The exact attribution of the 1703  $\text{cm}^{-1}$  band is uncertain, but it is characteristic of the carbonyl group of the carboxylic acid.

The positive 1514  $\text{cm}^{-1}$  band is assigned to the symmetric ring CC/CH stretching vibration of unlabeled tyrosine (Table 2). The red-shifted band is observed at 1497  $\text{cm}^{-1}$ . The  $\text{HC}_{\alpha}\text{H}$  bending and asymmetric counterparts of the ring CC/CH stretching mode are not evident but are predicted to be low intensity modes. Vibrational bands observed at 1437 and 1408  $\text{cm}^{-1}$  could represent vibrational changes involving the  $\text{C}_{\beta}$  and  $\text{C}_{\gamma}$  atoms. A negative band at 1296  $\text{cm}^{-1}$  is assigned to the C-F stretching modes of fluorotyrosine in wild-type MnSOD. It is observed at  $\sim 1293 \text{ cm}^{-1}$  in the model compound (Fig. 2 B). Two distinct bands are observed for unlabeled MnSOD in this region: at 1265 and 1232  $\text{cm}^{-1}$ . The mode at 1265  $\text{cm}^{-1}$  is consistent with the phenol group of tyrosine residues involved as proton donors in hydrogen-bonding interactions (21). Furthermore, the mode at 1232  $\text{cm}^{-1}$  can be assigned to tyrosine residues in which the phenol group is a proton acceptor in hydrogen-bonding interactions (21). These differences in environment give rise to different components of the C-O stretching vibration.

TABLE 2 Proposed assignment of infrared absorption band frequencies (in  $\text{cm}^{-1}$ ) observed for wild-type and Y34F human manganese superoxide dismutase

Mode description	Wild-type		Y34F	
	Unlabeled	Fluorinated	Unlabeled	Fluorinated
CC symmetric ring stretch	1614	1605	—	—
CC/CH asymmetric ring stretch	1593	—	—	—
CC/CH symmetric ring stretch	1514	1497	1514	1498
$\text{HC}_{\beta}\text{C}_{\gamma}$ bend	1437	1408	—	—
$\text{C}_{\beta}\text{C}_{\gamma}$ stretch	—	—	1406	—
$\text{HC}_{\alpha}\text{C}_{\beta}$ bend	—	—	1348	—
FC ring stretch	—	1296	—	1296
C-O phenolic stretch	1265	—	1257	—
C-O phenolic stretch	1232	1209	1228	1211

### FTIR spectra of unlabeled and fluorinated Y34F Human MnSOD

FTIR spectroscopy has been applied to the site-specific mutant of human MnSOD in which Tyr<sup>34</sup> was replaced with Phe (Y34F). Fig. 4 A shows the FTIR absorption spectra of unlabeled Y34F (*solid line*) and Y34F in which each of the eight remaining tyrosine residues is labeled with fluorine at the 3-position of the phenol ring (*dotted line*); the difference spectrum is shown in Fig. 4 B. Again, vibrational modes of unlabeled tyrosine are positive bands, and the modes of fluorinated tyrosine are negative bands. As was observed

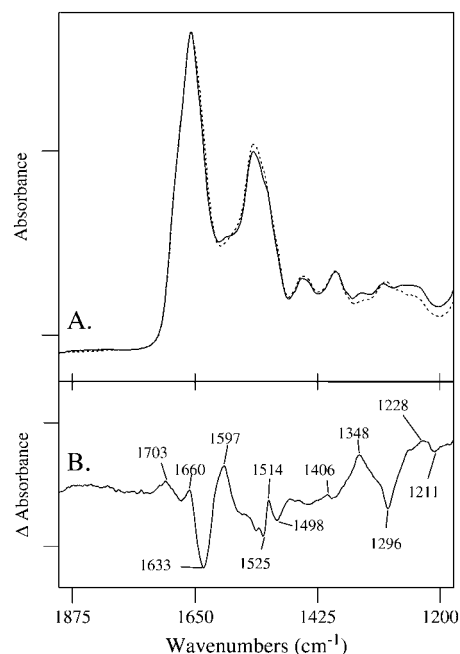


FIGURE 4 Infrared absorption spectra of (A, *solid line*) unlabeled Y34F and (A, *dotted line*) fluorinated Y34F MnSOD. The difference spectrum (unlabeled-minus-fluorinated) is shown in B. The tick marks on the y axis represent 0.03 AU. Spectral conditions were as described in Fig. 3.

for wild-type, fluorination of the tyrosine residues in Y34F causes changes in the vibrational modes of tyrosine. The  $1600\text{ cm}^{-1}$  region, which contains contributions from the tyrosine ring-stretching vibrations and from the amide I/II of the protein backbone, shows perturbations upon fluorination. Again, these changes are more likely due to ring vibrations than to changes in the backbone vibrations because of fluorine labeling on the phenol ring. Because of large and overlapping bands in the  $1600\text{ cm}^{-1}$  region, it is difficult to pinpoint particular ring-stretching vibrations that shift upon fluorination.

However, the CC/CH stretching vibrations observed at  $\sim 1500\text{ cm}^{-1}$  are quite apparent. A positive  $1514\text{ cm}^{-1}$  band is assigned to the symmetric ring CC/CH stretching vibration of unlabeled tyrosine (Table 2). The red-shifted band is observed at  $1498\text{ cm}^{-1}$ . The asymmetric counterpart of the ring CC/CH stretching mode is not seen, due to its expected low intensity. A negative mode at  $1296\text{ cm}^{-1}$  is attributed to the C-F stretching modes of 3-fluorotyrosine in Y34F MnSOD. Also, positive bands are observed at  $1406\text{ cm}^{-1}$  and  $1348\text{ cm}^{-1}$  and are tentatively assigned to vibrations involving the  $C_{\alpha}$ ,  $C_{\beta}$ , and  $C_{\gamma}$  atoms.

Note that the intensity of the  $1265\text{ cm}^{-1}$  band has decreased significantly and shifted to lower energy by  $8\text{ cm}^{-1}$  in Y34F compared with wild-type (Fig. 4 B). An expanded version of this part of the spectrum is shown in Fig. 5. The difference spectrum shown in Fig. 5 is the average of five spectra obtained on five different samples. Each one of the individual spectra shows the  $1265\text{ cm}^{-1}$  band. The spectra are corrected for protein concentrations by setting the amide

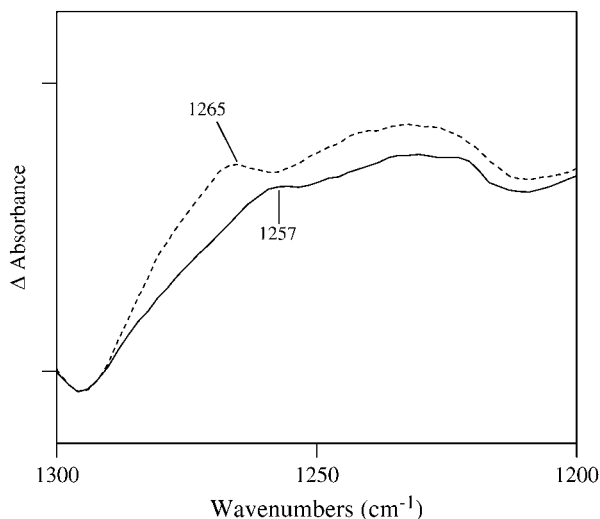


FIGURE 5 Comparison of the FTIR difference spectra of wild-type and Y34F MnSOD. The tick marks on the ordinate represent 0.02 absorbance units. Note that the intensity at  $1265\text{ cm}^{-1}$  has decreased significantly in Y34F compared with wild-type. These data show that the C-O stretching vibration of the phenolic group of the tyrosine side chain at residue 34 in wild-type makes an infrared contribution at  $1265\text{ cm}^{-1}$ . This frequency is characteristic of phenol groups that act as hydrogen bond donors (21).

I vibration of each absorbance spectrum to 0.35 absorbance units. This is an arbitrary number that was chosen to normalize the spectra that were used to obtain a difference spectrum. This method has been employed extensively for photosystem II (for an example, see Ayala and Barry (23)). By correcting for the amide I vibrational mode, small differences in protein concentration from sample to sample were taken into account. In addition to the correction for protein concentration, baseline correction of the absorbance spectra was performed using the software GRAMS (Galactic Industries, Salem, NH). The absorbance spectra shown in Figs. 3 A and 4 A were corrected using a two-point baseline correction. In this method two baseline points were selected at the ends of the spectral region. A multiple-point baseline correction was not practicable because our absorbance spectra show multiple overlapping bands and it is difficult to select points that belong to the baseline without biasing our data. Difference spectra were constructed using the baseline-corrected absorbance spectra; these difference spectra were indistinguishable from those obtained using no baseline correction.

One of the components of the C-O vibration at  $1228\text{--}1232\text{ cm}^{-1}$  is evident in both Figs. 3 B and 4 B; the other component at  $1265\text{ cm}^{-1}$  is very much smaller in Y34F. These data show that the C-O stretching vibration of the phenolic group of Y34 makes a contribution to the  $1265\text{ cm}^{-1}$  mode. This frequency is characteristic of phenol groups, which hydrogen bond and act as proton donors (21). This band has shifted to  $1257\text{ cm}^{-1}$  and has nearly disappeared in Y34F. A summary of the proposed vibrational mode assignments for wild-type and Y34F are shown in Table 2.

## Crystal structures

Crystal structures were determined for human wild-type MnSOD and Y34F MnSOD at  $1.85$  and  $1.53\text{ \AA}$ , respectively, in which the tyrosine residues were replaced with 3-fluorotyrosine. The electron density maps show the presence of 3-fluorotyrosine side chains in the two structures, with each of the nine 3-fluorotyrosine side chains in a single orientation. The fluorine groups were mostly refined at 70% occupancy. This is consistent with the percentage incorporation of fluorine tyrosine and resulted in reasonable  $B$  values for the fluorine atoms and the surrounding side chain atoms. We had expected to see a distribution of rotamers for 3-fluorotyrosine side chains, which would be clearly evident in these relatively high resolutions structures. However, it was immediately apparent from the initial electron density maps after molecular replacement that this was not the case. In the refined structures, each of the side chains of the nine 3-fluorotyrosine residues was observed clearly to adopt a preferred orientation, most likely reflecting local environments. The  $B$  values for these fluorine atoms were refined to reasonable  $B$  values, which were very similar to or the same as the surrounding atoms of the side chain.

Incorporation of 3-fluorotyrosine does not significantly alter the structures of the fluorinated wild-type and Y34F MnSOD, as they are structurally superimposable with their nonfluorinated counterparts; the average root mean-square deviation for all 198  $\alpha$ -carbon atoms between unfluorinated and fluorinated wild-type MnSOD is 0.3 Å. An apparent hydrogen-bonding scheme extending from the metal-bound solvent is maintained in the structure of the wild-type MnSOD containing 3-fluorotyrosines (Fig. 6). Analysis of potential van der Waals contacts of the hydrogen atoms (14,15) of Gln<sup>143</sup>, which is part of the hydrogen-bonding scheme, indicates that the amide group is oriented toward the hydroxyl of Tyr<sup>34</sup>; a 180° flip of the Gln<sup>143</sup> side chain (about  $\chi_3$ ) would cause steric clashes with neighboring atoms, including Asn<sup>73</sup>, Trp<sup>123</sup>, and the protein backbone. This suggests the directionality of the hydrogen bond chain emanating from the metal-bound solvent. The amide of the carboxamide side chain of Gln<sup>143</sup> acts as a hydrogen bond donor to the phenolic hydroxyl of fluorinated Tyr<sup>34</sup>, which in turn appears to be in a hydrogen bond with the water molecule between the side chains of Tyr<sup>34</sup> and His<sup>30</sup> (Fig. 6; Table 3). This is the same observation as made for the wild-type human MnSOD (4). However, this hydrogen-bonding network is altered in the Y34F mutant structure. The Y34F mutation results in another water molecule being present in the active site, in a position approximately equivalent to the hydroxyl group of the wild-type Tyr<sup>34</sup> (Fig. 7; Table 3). Thus, the His<sup>30</sup> ND1-water-water-Gln<sup>143</sup> NE<sup>2</sup> hydrogen bonds may partially restore the hydrogen-bonding relay in Y34F.

## DISCUSSION

Each of the nine tyrosine residues of wild-type human MnSOD was replaced with 3-fluorotyrosine. Interestingly, this had rather little effect on the structure of the enzyme; the

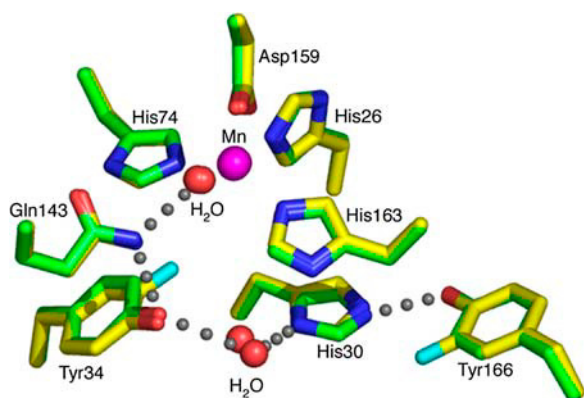


FIGURE 6 Crystal structure of the active site of human MnSOD, in which tyrosine residues have been replaced with 3-fluorotyrosine (yellow), superimposed upon wild-type human MnSOD (green). The apparent hydrogen-bond scheme between the manganese-bound solvent molecule, the side chains of Gln<sup>143</sup>, Tyr<sup>34</sup>, a water molecule, and the His<sup>30</sup> and Tyr<sup>166</sup> side chains is conserved in both structures and is depicted by gray spheres.

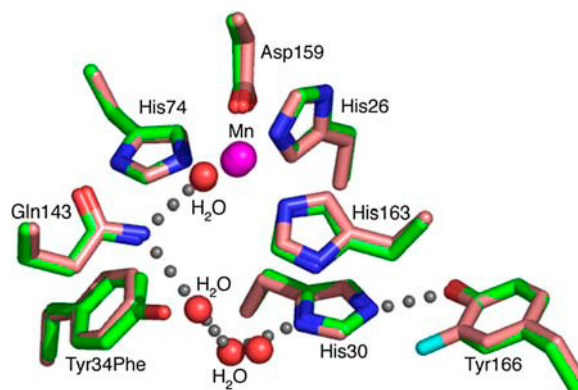


FIGURE 7 Crystal structure of the active site of Y34F mutant of human MnSOD that has tyrosines replaced by 3-fluorotyrosine (salmon), superimposed upon wild-type human MnSOD (green). The active-site hydrogen-bonding scheme, present in the wild-type enzyme, is altered in the Y34F mutation. Gray spheres depict the Y34F active-site hydrogen bonds, where an extra water molecule is observed in the active site, which is likely to partially restore the hydrogen bond relay.

fluorinated and unfluorinated structures are closely superimposable. Significantly, the apparent hydrogen-bonded network, extending from the manganese-bound solvent, appears unchanged in the fluorinated wild-type MnSOD (Fig. 6). The crystal structure of unfluorinated wild-type human Mn(III)-SOD with the highest resolution (Protein Data Bank accession code 1LUV (24)) shows that in subunit A of the asymmetric subunit dimer the hydroxyl oxygen of Tyr<sup>34</sup> is located 3.0 Å from the side-chain carboxamide (NE<sup>2</sup>) of Gln<sup>143</sup> and 2.8 Å from the oxygen of a water molecule, both suggesting hydrogen bonding. However, the corresponding

TABLE 3 Bond distances and angles of the active-site hydrogen bond network in fluorinated wild-type and fluorinated Y34F human MnSOD

Bond distances, Å	Fluorinated wild-type		Fluorinated Y34F	
	Subunit A	Subunit B	Subunit A	Subunit B
Mn-bound OH-Gln <sup>143</sup> NE <sup>2</sup>	2.8	2.8	2.7	2.7
Gln <sup>143</sup> NE <sup>2</sup> -Tyr <sup>34</sup> OH	2.9	3.0	—	—
Tyr <sup>34</sup> OH-H <sub>2</sub> O	2.9	2.8	—	—
Gln <sup>143</sup> NE <sup>2</sup> -H <sub>2</sub> O	—	—	3.1	3.0
H <sub>2</sub> O-H <sub>2</sub> O	—	—	2.7	2.7
H <sub>2</sub> O-His <sup>30</sup> ND1	2.7	2.5	2.8	2.8
His <sup>30</sup> NE <sup>2</sup> -Tyr <sup>166</sup> OH	2.8	2.7	2.8	2.7
Bond angles, degrees				
Mn-bound OH-Gln <sup>143</sup> NE <sup>2</sup> -Y <sup>34</sup> OH	116.0	112.2	—	—
Gln <sup>143</sup> -Y <sup>34</sup> OH-H <sub>2</sub> O	120.7	122.1	—	—
Ty <sup>34</sup> OH-H <sub>2</sub> O-His <sup>30</sup> ND1	147.8	144.7	—	—
Mn-bound OH-Gln <sup>143</sup> NE <sup>2</sup> -H <sub>2</sub> O	—	—	95.9	91.1
Gln <sup>143</sup> NE <sup>2</sup> -H <sub>2</sub> O-H <sub>2</sub> O	—	—	140.8	137.9
H <sub>2</sub> O-H <sub>2</sub> O-His <sup>30</sup> ND1	—	—	122.0	115.3
His <sup>30</sup> NE <sup>2</sup> -Tyr <sup>166</sup> OH-Ty <sup>166</sup> CZ	123.8	115.8	120.5	122.7



distances in subunit B are 2.9 Å for the carboxamide (Gln<sup>143</sup>) to hydroxyl (Tyr<sup>34</sup>) and 3.3 Å for the hydroxyl to water. The fluorinated wild-type human Mn(III)SOD structure shows that in subunit A the hydrogen bond distancing between hydroxyl oxygen of Tyr<sup>34</sup> and the side-chain carboxamide of Gln<sup>143</sup> is 2.9 Å and also 2.9 Å to the oxygen of the water molecule. Distances in subunit B are 3.0 Å for the Gln<sup>143</sup> carboxamide to Tyr<sup>34</sup> hydroxyl and 2.8 Å for Tyr<sup>34</sup> hydroxyl to water. Thus, these crystal structures suggest that the hydroxyl of Tyr<sup>34</sup> is a hydrogen bond acceptor from NE<sup>2</sup> of Gln<sup>143</sup>; however, the nature of the interaction of the hydroxyl of Tyr<sup>34</sup> with the adjacent water molecule is ambiguous.

In this work, FTIR spectroscopy is used to study MnSOD; thorough studies of MnSOD using resonance Raman spectroscopy are reported (8). To the best of our knowledge, this is the first FTIR study of fluorinated tyrosine in a protein with the express goal of assessing the structural characteristics of a particular tyrosine residue. The addition of the electronegative fluorine atom at the 3-position of the phenol ring of tyrosine caused the ring-stretching vibrations to be lower in frequency, indicating weaker bonds. Also, the phenolic C-O stretching vibration of 3-fluorotyrosine was found at lower frequency than in tyrosine, suggesting a weaker C-O bond in 3-fluorotyrosine than in tyrosine. This was confirmed in both tyrosine as a model compound and DFT calculations (Figs. 1 and 2; Table 1) and in the protein data (Figs. 3 and 4; Table 2).

The FTIR spectra revealed additional differences between fluorinated and unfluorinated MnSOD (Figs. 3 and 4). Although the crystal structures of the unfluorinated and fluorinated wild-type MnSOD were closely superimposable (Fig. 6), the FTIR data showed changes in the amide I and II vibrational bands, suggesting some changes in backbone conformation which are apparently not within the detection limits of the crystallography. Among other notable differences was the C-O stretch of the phenol group of tyrosine, the intensity and frequency of which is sensitive to the status of the hydroxyl as a proton donor or acceptor (21). The difference FTIR spectra of Figs. 3 and 4 for fluorinated and unfluorinated MnSOD identify this tyrosine C-O stretch, and the comparison of these spectra for wild-type and Y34F MnSOD identify this vibrational mode in the active-site residue Tyr<sup>34</sup>.

Although the phenolic side chain of Tyr<sup>34</sup> can participate as both a hydrogen bond acceptor and donor, the FTIR analysis of human Mn(III)SOD in this study shows a prominent absorption at 1265 cm<sup>-1</sup> (Fig. 5), indicating that the C-O stretch in the side chain of Tyr<sup>34</sup> is dominated by the OH group acting as a hydrogen bond donor. This then confirms the protonation state of Tyr<sup>34</sup> in Mn(III)SOD and the presence of a hydrogen bond between the hydroxyl of Tyr<sup>34</sup> and the adjacent water molecule. Moreover, the data complement the crystal structure, which suggests this hydrogen bond, although somewhat ambiguously. These hydrogen bond interactions of Tyr<sup>34</sup> have implications for the spectroscopic and catalytic properties of the enzyme. They

affect the visible absorption spectrum of MnSOD at 480 nm, with Tyr<sup>34</sup> being the source of the pK<sub>a</sub> near 9.5 (7,8,25). The ionization of Tyr<sup>34</sup> also explains the pH dependence of catalysis observed in wild-type and mutants in which  $k_{cat}/K_m$  shows a pK<sub>a</sub> near 9.5 (8). Tyrosine<sup>34</sup>, most likely through hydrogen-bonded interactions with Gln<sup>143</sup> and a water molecule, provides an active-site framework that supports proton transfer in catalysis, although Tyr<sup>34</sup> may not be the source of the protons that eventually form the H<sub>2</sub>O<sub>2</sub> product (5).

We thank Professor I. Fridovich for helpful suggestions on growing bacteria in minimal media and Max Jurcovich for excellent technical assistance.

This work was supported by grants from the National Institutes of Health: GM54903 to D.N.S. and AI054945 to J.A.T.

## REFERENCES

- Holm, R. H., P. Kennepohl, and E. I. Solomon. 1996. Structural and functional aspects of metal sites in biology. *Chem. Rev.* 96:2239–2314.
- Beyer, W., J. Imlay, and I. Fridovich. 1991. Superoxide dismutases. *Prog. Nucleic Acid Res. Mol. Biol.* 40:221–253.
- Fridovich, I. 1989. Superoxide dismutases: an adaptation to a paramagnetic gas. *J. Biol. Chem.* 264:7761–7764.
- Borgstahl, G. E. O., H. E. Parge, M. J. Hickey, W. F. Beyer, R. A. Hallewell, and J. A. Tainer. 1992. The structure of human mitochondrial manganese superoxide dismutase reveals a novel tetrameric interface of two 4-helix bundles. *Cell.* 71:107–118.
- Silverman, D. N., and H. S. Nick. 2002. The catalytic pathway of manganese superoxide dismutase by direct observation of superoxide. *Methods Enzymol.* 349:61–74.
- Whittaker, M. M., and J. W. Whittaker. 1997. Mutagenesis of a proton linkage pathway in *Escherichia coli* manganese superoxide dismutase. *Biochemistry.* 36:8923–8931.
- Guan, Y., M. J. Hickey, G. E. O. Borgstahl, R. A. Hallewell, J. A. Lepock, D. O'Connor, Y. Hsieh, H. S. Nick, D. N. Silverman, and J. A. Tainer. 1998. The crystal structure of Y34F mutant human mitochondrial manganese superoxide dismutase and the functional role of tyrosine 34. *Biochemistry.* 37:4722–4730.
- Maliekal, J., A. Karapetian, C. Vance, E. Yikilmaz, Q. Wu, T. Jackson, T. C. Brunold, T. G. Spiro, and A.-F. Miller. 2002. Comparison and contrasts between the active site pKs of MnSOD and those of FeSOD. *J. Am. Chem. Soc.* 124:15064–15075.
- Beck, Y., D. Bartfield, Z. Yavin, A. Levanon, M. Gorecki, and J. R. Hartman. 1988. Efficient production of active human manganese superoxide dismutase in *Escherichia coli*. *Biotechnology.* 6:930–935.
- Otwinowski, Z., and W. Minor. 1997. Processing of x-ray diffraction data collected in oscillation mode. In *Methods in Enzymology*, Vol. 276. C. W. Carter Jr. and R. M. Sweet, editors. 307–326. Academic Press, New York.
- Navaza, J. 2001. Implementation of molecular replacement in AMoRe. *Acta Crystallogr. D.* 57:1367–1372.
- Brunger, A. T., P. D. Adams, G. M. Clore, W. L. DeLano, P. Gros, R. W. Grosse-Kunstleve, J. S. Jiang, J. Kuszewski, M. Nilges, N. S. Pannu, R. J. Read, L. M. Rice, T. Simonson, and G. L. Warren. 1998. Crystallography & NMR system: a new software suite for macromolecular structure determination. *Acta Crystallogr.* D54:905–921.
- Sheldrick, G. M., and T. R. Schneider. 1997. SHELXL: high-resolution refinement. *Methods Enzymol.* 277:319–343.
- Word, J. M., S. C. Lovell, J. S. Richardson, and D. C. Richardson. 1999a. Asparagine and glutamine: using hydrogen atom contacts in the choice of side-chain amide orientation. *J. Mol. Biol.* 285:1735–1747.



15. Word, J. M., S. C. Lovell, T. H. LaBean, H. C. Taylor, M. E. Zalis, B. K. Presley, J. S. Richardson, and D. C. Richardson. 1999b. Visualizing and quantifying molecular goodness-of-fit: small-probe contact dots with explicit hydrogen atoms. *J. Mol. Biol.* 285:1711–1733.
16. McRee, D. E. 1999. XtalView/Xfit: a versatile program for manipulating atomic coordinates and electron density. *J. Struct. Biol.* 125: 156–165.
17. Adams, P. D., N. S. Pannu, R. J. Read, and A. T. Brunger. 1997. Cross-validated maximum likelihood enhances crystallographic simulated annealing refinement. *Proc. Natl. Acad. Sci. USA.* 94:5018–5023.
18. Frisch, M. J., G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, V. G. Zakrzewski, J. A. Montgomery Jr., R. E. Stratmann, J. C. Burant, S. Dapprich, J. M. Millam, A. D. Daniels, K. N. Kudin, M. C. Strain, O. Farkas, J. Tomasi, V. Barone, M. Cossi, R. Cammi, B. Mennucci, C. Pomelli, C. Adamo, S. Clifford, J. Ochterski, G. A. Petersson, P. Y. Ayala, Q. Cui, K. Morokuma, D. K. Malick, A. D. Rabuck, K. Raghavachari, J. B. Foresman, J. Cioslowski, J. V. Ortiz, B. B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I. Komaromi, R. Gomperts, R. L. Martin, D. J. Fox, T. Keith, M. A. Al-Laham, C. Y. Peng, A. Nanayakkara, C. Gonzalez, M. Challacombe, P. M. W. Gill, B. Johnson, W. Chen, M. W. Wong, J. L. Andres, C. Gonzalez, M. Head-Gordon, E. S. Replogle, and J. A. Pople. 1998. Gaussian 98, Revision A.3. Gaussian Inc., Pittsburgh, PA.
19. Szczepanski, J., J. Banisaukas, M. Vala, S. Hirata, R. J. Bartlett, and M. Head-Gordon. 2002. Vibrational and electronic spectroscopy of the fluorene cation. *J. Phys. Chem. A.* 106:63–73.
20. Ayala, I., K. Range, D. York, and B. A. Barry. 2002. Spectroscopic properties of tyrosyl radicals in dipeptides. *J. Am. Chem. Soc.* 124: 5496–5505.
21. Takeuchi, H., N. Watanabe, Y. Satoh, and I. Harada. 1989. Effects of hydrogen-bonding on the tyrosine Raman bands in the 1300–1150  $\text{cm}^{-1}$  region. *J. Raman Spectrosc.* 20:233–237.
22. Mukherjee, A., M. L. McGlashen, and T. G. Spiro. 1995. Ultraviolet resonance Raman spectroscopy and general valence force field analysis of phenolate and phenoxy radical. *J. Phys. Chem.* 99:4912–4917.
23. Ayala, I., and B. A. Barry. 2002. Histidine 190–D1 and glutamate 189–D1 provide structural stabilization in photosystem II. *Biochemistry.* 41:11456–11465.
24. Hearn, A. S., M. E. Stroupe, D. E. Cabelli, C. A. Ramilo, J. P. Luba, J. A. Tainer, H. S. Nick, and D. N. Silverman. 2003. Catalytic and structural effects of amino acid substitution at His 30 in human manganese superoxide dismutase. *Biochemistry.* 42:2781–2789.
25. Hsu, J.-L., Y. Hsieh, C. Tu, D. O'Connor, H. S. Nick, and D. N. Silverman. 1996. Catalytic properties of human manganese superoxide dismutase. *J. Biol. Chem.* 271:17687–17691.