

Figure 3. Drug binding to site 1 in HSA (defatted). The detailed binding conformations are shown for (a) CMPF, (b) oxyphenbutazone and (c) phenylbutazone. In each case the drug is shown in a stick representation with a semi-transparent van der Waals surface. Abbreviated names for drugs are defined in the footnotes to Table 1. Selected side-chains are shown as sticks colour-coded by atom type; yellow dashed lines indicate hydrogen bonds. Residue I264, which bifurcates the back-end of the pocket, is shown as grey spheres. Note that the tip of K195 is disordered in the HSA–CMPF complex so the side-chain amino group is not shown in (a). (d) Top view of the superposition of drugs bound to site 1 in defatted HSA. Drugs are shown in a stick representation with carbon atoms coloured orange, nitrogen atoms in blue and oxygen atoms (also shown as small spheres) in red. Oxygen atoms tend to cluster on either side of the binding pocket. (e) Side view of superposition of drugs shown in (d) along with a semi-transparent surface (orange) depicting the extent of the pocket as determined by combining the pseudo-atom output from PASS,⁵⁶ which maps potential pockets on the protein surface, with the superposed drugs bound to the pocket, to account for observed variation in pocket dimensions due to ligand binding.

presence of flexible side-chains provides significant room for manoeuvre.

The drugs occupy the apolar compartments of site 1 to different extents. All compounds access the right-hand sub-chamber to a greater (oxyphenbutazone, phenylbutazone, warfarin) or lesser (CMPF, thyroxine⁶) degree but only phenylbutazone and CMPF project hydrophobic moieties into the left-hand sub-chamber (Figure 3(a)–(c); Supplementary Figure 1). The front, lower sub-chamber is occupied by phenyl groups of oxyphenbutazone and warfarin, and one of the iodine atoms projecting from the outer phenyl ring of thyroxine.⁶

In addition to hydrophobic contacts the site 1 compounds make a number of specific interactions with residues belonging to the inner and outer polar clusters. All of the compounds are positioned to make a hydrogen bond interaction with the

hydroxyl group of Y150, as found previously for thyroxine,⁶ and this residue therefore assumes a central role in drug interactions. In total CMPF makes five hydrogen-bond or salt-bridge interactions with Y150, H242, K199 and R222 and appears particularly well adapted to the pocket (Figure 3(a)), an observation that probably explains the high-affinity binding of this compound ($K_d = 0.1 \mu\text{M}$),⁴¹ despite its relatively polar nature.¹² Clearly the drug binding defect observed *in vivo* as CMPF levels rise in kidney patients^{35,41} is due to specific steric blocking of drug site 1 by this compound. The *R*-(+) and *S*-(-) enantiomers of warfarin bind in essentially the same position as one another and appear capable of making a total of three hydrogen bonds with the residues that interact with CMPF (the acetyl oxygen atom being able to bond alternately to K199 or R222;

Supplementary Figure 1(e)); the similarity of the binding environments for the enantiomers helps to explain the poor stereoselectivity of HSA for this drug.⁴² Oxyphenbutazone and phenylbutazone both make just a single hydrogen bond interaction with Y150 in drug site 1 (Figure 3(b) and (c)). In each case an oxygen atom on the opposite side of the drug lies 4–6 Å from R222 and/or K199; it is possible that water molecules may bridge interactions with these residues but these are not evident at the present resolution of these structures (Table 1). Strikingly, although oxyphenbutazone is a derivative of phenylbutazone, possessing an additional hydroxyl group on one of the phenyl rings, it binds in a conformation that is rotated by about 180° with respect to phenylbutazone and places the hydroxyl group at the mouth of the pocket where it can interact with bulk solvent. This is a revealing example of the unpredictable effects that even minor structural modifications can have on drug binding.

The prevalence of basic residues and the absence of acidic ones define the specificity of the pocket. The observation that reagents that are specific for site 1 generally possess centrally located anionic or electronegative features^{2,9,28} is due to location of polar patches in the middle of the pocket flanked by apolar regions. In fact the structural data suggest a refinement of this view in that the pocket appears to

be specific for molecules with two anionic or electronegative features on opposite sides of the ligand that can simultaneously interact with the two polar patches (Figure 3(d)). The distance between these basic patches accounts for the finding that the presence of two electronegative groups separated by five to six bonds, as in CMPF, is particularly important for tight binding to site 1.^{12,27}

Superposition of the HSA–drug complexes for site 1 compounds reveals that there are only small side-chain movements associated with drug binding, in contrast to the displacements observed in the complex with thyroxine, a significantly larger molecule (M_r 777 Da)⁶ (Figure 3(d) and (e)). For CMPF and the drugs used in this study (M_r ~310 Da) the greatest movement is observed for Y150 and W214. Side-chain variability with this set of molecules seems rather modest given the scope provided by numerous aliphatic residues lining the pocket. It may be that the apparent adaptability of the pocket is more a product of its size, which does not place tight steric constraints on the binding of small drugs and allows co-binding of water molecules that can flexibly mediate interactions with the protein.

Drug site 1 in HSA-myristate

Upon binding of fatty acids, Y150 from

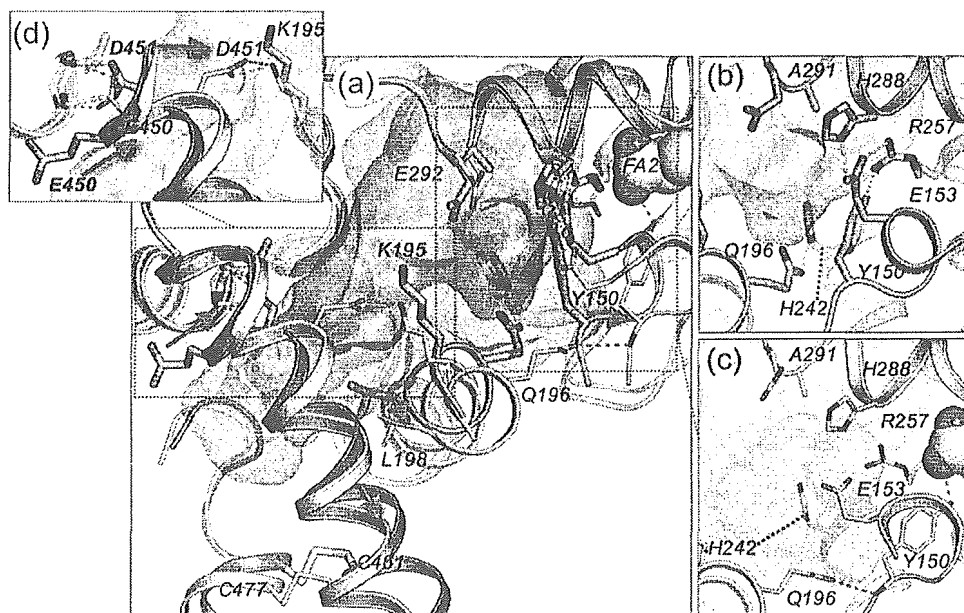


Figure 4. Conformational changes in drug site 1 as a result of fatty acid binding. (a) Superposition of HSA (secondary structure colour-coded by subdomain; selected side-chains coloured by atom type) and HSA-myristate (light-grey secondary structure with light grey carbon atoms in side-chains). Drug site 1 in HSA is depicted by a light-brown semi-transparent surface; binding of fatty acid results in expansion of drug site 1 (blue semi-transparent surface) as a result of concerted movement of several structural components. Red arrows indicate the direction of structural changes associated with fatty acid binding. (b) Close-up views of the region around Tyr150 in HSA and (c) HSA-myristate, colour-coded as in (a). (d) Rotated view of the vicinity of Glu450 and Asp451, which both rotate to new positions on fatty acid binding; Glu450 supplants Asp451 in interacting with the main-chain amides of residues 343–344 while Asp451 itself moves to form a salt-bridge with Lys195. Initial residue positions are labelled in boldface.

subdomain IB moves to interact with the carboxylate moiety of the lipid bound to the site that straddles domains I and II (fatty acid site FA2³) (Figure 4(a)–(c)). This helps to drive the relative rotation of domains I and II and has a large impact on one side of drug site 1 (Figure 2(a) and (c)). There is an extensive rearrangement of the H-bond network involving Y150, E153, Q196, H242, R257 and H288, which opens a solvent channel (between Y150 and Q196), thus increasing the volume of the pocket and altering its polarity distribution: the inner polar cluster is disrupted and partially neutralised by fatty acid binding; only H242 is relatively unaffected (Figure 4(b) and (c)). The helix containing L198 is also displaced outwards. This appears to impact an adjacent helix from subdomain IIIA (residues 442–466) and its disulphide-bonded neighbour. This latter helix is also twisted around its axis, since binding of myristate to site FA3 in IIIA replaces E450 in a salt-bridge interaction

with R348. As a result E450 rotates to replace D451 in interacting with the amide groups of residues 343–344 (Figure 4(d)). In turn, D451 relocates to a position that allows it to form a salt-bridge with K195. This cascade of interactions indicates one possible link between the two drug sites, at least in the presence of fatty acid.

To assess the impact of fatty acid-induced conformational changes on drug binding, we investigated the structure of HSA-myristate complexed with a range of site 1 drugs. The results are applicable to more physiologically relevant fatty acids, such as palmitate or oleate, since these exert the same conformational effects on the protein.^{4,5} Although drug site 1 is co-incident with a fatty acid binding site (FA7),⁴ this is likely to be a low-affinity site and in each case the drug was observed to displace the lipid.

In spite of the structural changes wrought by fatty acid binding, many of the features that emerged

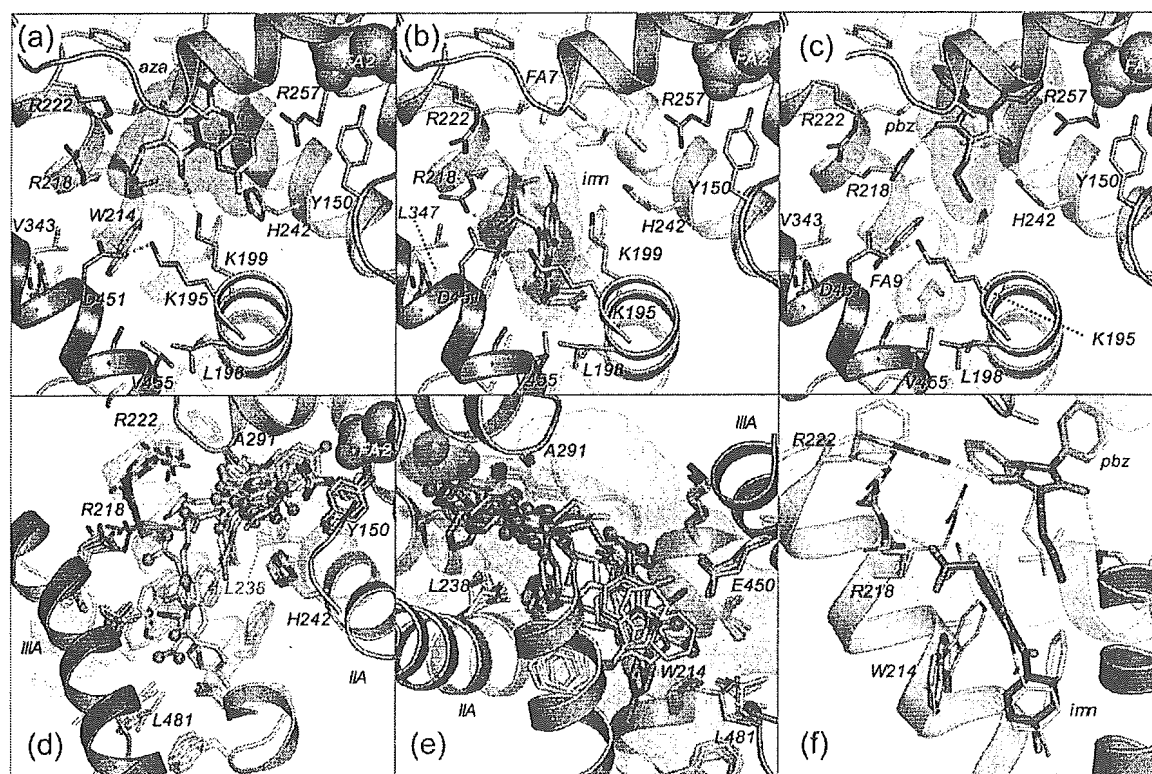


Figure 5. Drug binding to site 1 in HSA-myristate. The detailed binding conformations are shown for (a) azapropazone, (b) indomethacin and (c) phenylbutazone. In each case the drug is shown in a stick representation with a semi-transparent van der Waals surface (magenta). Bound fatty acids are depicted with a yellow semi-transparent van der Waals surface; otherwise colour-coding is as in Figure 3. The methylene tail of a molecule of myristate was observed co-bound with phenylbutazone in site FA9; this corresponds to a weak fatty acid site observed previously for medium-chain fatty acids.⁴ (d) Top view of the superposition of drugs bound to site 1 in HSA-myristate (coloured as in Figure 3). In this case the clustering of oxygen atoms is less pronounced. (e) Side view of superposition of drugs shown in (d) along with a semi-transparent surface (blue) depicting the extent of the pocket, determined as described in the legend to Figure 3. (f) Superposition of the structure of HSA-myristate–indomethacin–phenylbutazone (secondary structure coloured by domain with drugs and selected side-chains shown as sticks with grey carbon atoms) with HSA-myristate–indomethacin (drug and side-chains shown as thin sticks with cream carbon atoms) and HSA-myristate–phenylbutazone (blue carbon atoms in side-chains).

from the comparison of complexes of site 1 drugs with defatted HSA were also evident in the presence of myristate (Figure 5). For example, with the notable exception of indomethacin (Figure 5(b)), all the compounds studied were again found to bind in the central portion of the binding cavity, pinned between L238 and A291 (Figure 5(d) and (e)). As before there was considerable variability in the lateral positioning of drugs in the plane defined by this grouping, with different drugs occupying the pocket sub-chambers to different extents.

Nevertheless, some remarkable differences were also observed. Since Y150 is removed from the pocket to interact with fatty acid, it is no longer available to make the central contribution to drug binding that is observed in complexes with defatted HSA. Rather, different drugs make use of the various basic and polar ligands on both sides of the binding pocket. Most interactions are made with the side-chains of K199 and R222 on one side of the pocket and H242 on the other, though R218 and R257 both interact specifically with some compounds (e.g. indomethacin, phenylbutazone; Figure 5(b) and (c)).

For oxyphenbutazone, phenylbutazone and warfarin we have solved the structures of their complexes with HSA in the absence and presence of fatty acid. Comparison of the structures reveals only minor adjustments of the binding conformations of phenylbutazone (Figures 3(c) and 5(c)) and warfarin (Supplementary Figures 1(e) and 2(h)), a surprising result given that both drugs lose a specific interaction with Y150 on fatty acid binding, although interactions with H242 are retained. Interactions with solvent, as observed for warfarin,¹⁵ may also help to compensate for the loss of Y150. In the presence of fatty acid, phenylbutazone rotates to insert one of its two phenyl groups about 1.5 Å further into the hydrophobic sub-chamber at the back end of the pocket and position a carbonyl group within 3.4 Å of the guanidinium group of R218 (compare Figures 3(c) and 5(c)). Warfarin slides forward by 1 Å to accommodate the new position of R257. Addition of fatty acids to HSA reportedly increases the affinity of site 1 for warfarin^{28,31,32} but it is difficult to extract a precise molecular explanation for this effect from the structural data alone. One interesting difference is that the electron density for the coumarin ring of the drug at the back end of the pocket is significantly stronger in the HSA-myristate complex, suggesting that this moiety is more stably associated with the pocket in the presence of fatty acid. A similar observation was made for the phenyl rings of phenylbutazone, which also bind deep in the pocket, and we therefore suggest that fatty acid binding should also enhance the affinity of phenylbutazone.

In contrast to phenylbutazone and warfarin, oxyphenbutazone undergoes a re-orientation of about 180° due to fatty acid binding so that in the HSA-myristate complex this drug binds in a

conformation that corresponds closely to that found for phenylbutazone (compare Supplementary Figure 1(c) and (d) with Supplementary Figure 2(f) and (g)). Thus in the presence of fatty acids the addition of a hydroxyl group to a phenyl ring in phenylbutazone has a minimal effect on the binding orientation.

In their complexes with HSA-myristate, oxyphenbutazone and phenylbutazone occupy both the left-hand and right-hand sub-chambers with phenyl or phenolic moieties (the phenolic hydroxyl of oxyphenbutazone makes a hydrogen bond to the main chain carbonyl oxygen of R257). In the case of di-iodosalicylic acid (DIS), tri-iodobenzoic acid (TIB)³ and iodipamide, two of the iodine atoms on each ring overlie the positions of the aromatic rings in phenylbutazone and oxyphenbutazone (Supplementary Figure 2). This consistent positioning of iodine-substituted rings appears to be dictated by the shape of the binding site, in particular by the location of I264, which bifurcates the back end of the pocket into the two apolar sub-chambers. In addition, like oxyphenbutazone and phenylbutazone, the carboxylate groups of DIS and TIB both make hydrogen bonds with the side-chain of H242, which appears to assume a more prominent role in drug interactions than in defatted HSA.

Indomethacin is unusual in that it binds exclusively to the front, lower sub-compartment of site 1 and does not displace the fatty acid that is weakly bound to the centre of subdomain IIA (Figure 5(b)). In fact this drug can only be accommodated by inducing rotation of the side-chain of W214 through ~160°. This provides access to an additional cavity within IIA at the very base of the interdomain cleft that is largely delineated by L198, F206, A210, F211, W214 from IIA and L481 from subdomain IIIA; W214 also contacts residues V343 and L347 from IIB, so that the integrity of this cavity depends on contributions from three subdomains (Figure 5(b)). The chlorobenzoyl moiety of indomethacin binds at the bottom of this cavity while the indole ring is stacked between the flipped tryptophan and the apolar stem of K199. The indomethacin carboxylate group appears to make a bidentate salt-bridge to Arg218 (~2.8 Å) but there is only weak density for this moiety and an alternative conformation in which the carboxylate group flips over to interact with K199 may also be possible.

This expanded lower sub-chamber is also accessed by the contrast agent iodipamide (Supplementary Figure 2(e)), which is long enough to span the distance to the central portion of the site 1 pocket. Notably, iodipamide gains access to the lower chamber by inducing a much more modest ~20° χ_1 rotation of the W214 side-chain in the opposite direction to that induced by indomethacin, thereby placing the indole ring in a plane anti-parallel conformation (Supplementary Figure 2(d) and (e)).

Superposition of the HSA-myristate-indomethacin structure with those for other HSA-myristate-drug

complexes suggested that indomethacin would co-bind with some other site 1 compounds such as azapropazone, oxyphenbutazone, phenylbutazone, DIS and TIB. We tested this idea by performing azapropazone–phenylbutazone and azapropazone–indomethacin double-drug soaks with HSA-myristate crystals. In both cases, the resulting difference electron density maps indicated that indomethacin was binding in contact with the second drug. The occupancies refined to >80% for the two drugs in each complex, indicating that they were binding simultaneously to the pocket on HSA. This interpretation is supported by the finding that the two drugs are slightly shifted in the double drug soaks by comparison to their positions in the corresponding single drug soaks, presumably as a result of drug–drug contacts (Figure 5(f); Supplementary Figure 3). The most striking effect of co-binding of these two drugs is the concerted rearrangement of R218 and R222, the principal effect of which is to substitute R222 instead of R218 as a binding partner for the carbonyl group of phenylbutazone (Figure 5(f)).

The simultaneous accommodation of indomethacin and either azapropazone or phenylbutazone in drug site 1 of the crystal structure is supported by binding data, which show that these drugs do not displace one another from HSA (A. Annis, personal communication).¹⁸ These results were obtained using defatted HSA, indicating that co-binding also happens in the absence of fatty acid as expected from modelling experiments (data not shown). In contrast, comparison of the crystal structures suggests that indomethacin will not co-bind with every other site 1 drug. For example, we would predict steric clashes between indomethacin and iodipamide or warfarin (Supplementary Figure 2); this is consistent with binding data showing that indomethacin competes with warfarin.¹⁸

Superposition of all the drugs that have been analysed in complex with HSA-myristate reveals that site 1 extends significantly beyond the core of subdomain IIA (Figure 5(d) and (e)). The larger dataset of structures also reveals additional side-chain alterations associated with ligand binding. The greatest side-chain movements are again seen for residues at the mouth of the pocket, especially W214, the gatekeeper to the expanded lower sub-chamber, and the basic residues that make specific interaction with many of the bound drugs (K199, R218 and R222). This flexibility clearly contributes to the adaptability of the binding pocket. Within the pocket, packing constraints seem to restrict the side-chain variability in the HSA-myristate complex. Interestingly, although fatty acid binding displaces Y150 and Q196, thus opening up a new solvent channel with access to the protein exterior, none of the drugs studied here appears to take advantage of this new feature. The structural change nevertheless suggests ways in which compounds might be designed to specifically recognise the fatty acid-bound form of HSA.

For several drugs, secondary binding sites out-

side subdomain IIA were observed in the HSA-myristate complex. Azapropazone, indomethacin and warfarin all bind in subdomain IB. With the exception of azapropazone, which displaced the fatty acid from subdomain IB, these compounds bound, apparently co-operatively with the lipid, in contact with its methylene tail. Conceivably these interactions will be altered by the presence of fatty acids longer than myristate which are more prevalent *in vivo*.⁴³ Interestingly, evidence for weak secondary binding in subdomain IB by azapropazone and warfarin was also observed in the absence of fatty acid (data not shown). An alternative mode of co-operativity was found for oxyphenbutazone, which makes a hydrogen bond *via* its hydroxyl group to the carboxylate group of the fatty acid bound to subdomain IIIB (FA5). A secondary iodipamide site that accommodates only one half of the molecule was found within the interdomain cleft, in precisely the same locus as the thyroxine site identified in the HSA-myristate complex.⁶

Drug site 2

Drug site 2 is composed of all six helices of subdomain IIIA and is therefore topologically similar to site 1 (subdomain IIA). Although, like site 1, it also comprises a largely pre-formed hydrophobic cavity with distinct polar features, there are significant differences between the two drug pockets. Drug site 2 is smaller than site 1; the principal binding region corresponds to the central portion of the site 1 pocket and appears to possess just one sub-compartment, the rear right-hand hydrophobic sub-chamber, though in this case the sub-chamber is only accessed following ligand-induced side-chain movements (see below). To a large extent the left-hand sub-chamber is eliminated by the presence of Y411, which occurs in subdomain IIIA at the position corresponding to L219 in IIA (Figure 6(a)). A further difference arises because, although the two drug sites are in structurally similar subdomains, these are packed in different contexts with respect to the remainder of the protein. The entrance to drug site 1 is enclosed by subdomains IIB and IIIA; residues from these subdomains contribute to the formation of the front sub-chamber which binds indomethacin and accommodates portions of iodipamide, phenylbutazone and warfarin. However, the entrance to site 2 is not encumbered in this way: although IIIA is followed by IIIB, this subdomain is rotated further away from the drug site entrance (in comparison to drug site 1, domain II) and leaves the pocket entrance more exposed to solvent (Figure 2(a) and (c)).

In contrast to site 1, drug site 2 has a single main polar patch, located close to one side of the entrance of the binding pocket and centred on Tyr411 but also including R410, K414 and S489 (Figure 6(a)–(c)). Of these residues only R410 and K414 occur in equivalent positions to polar-patch

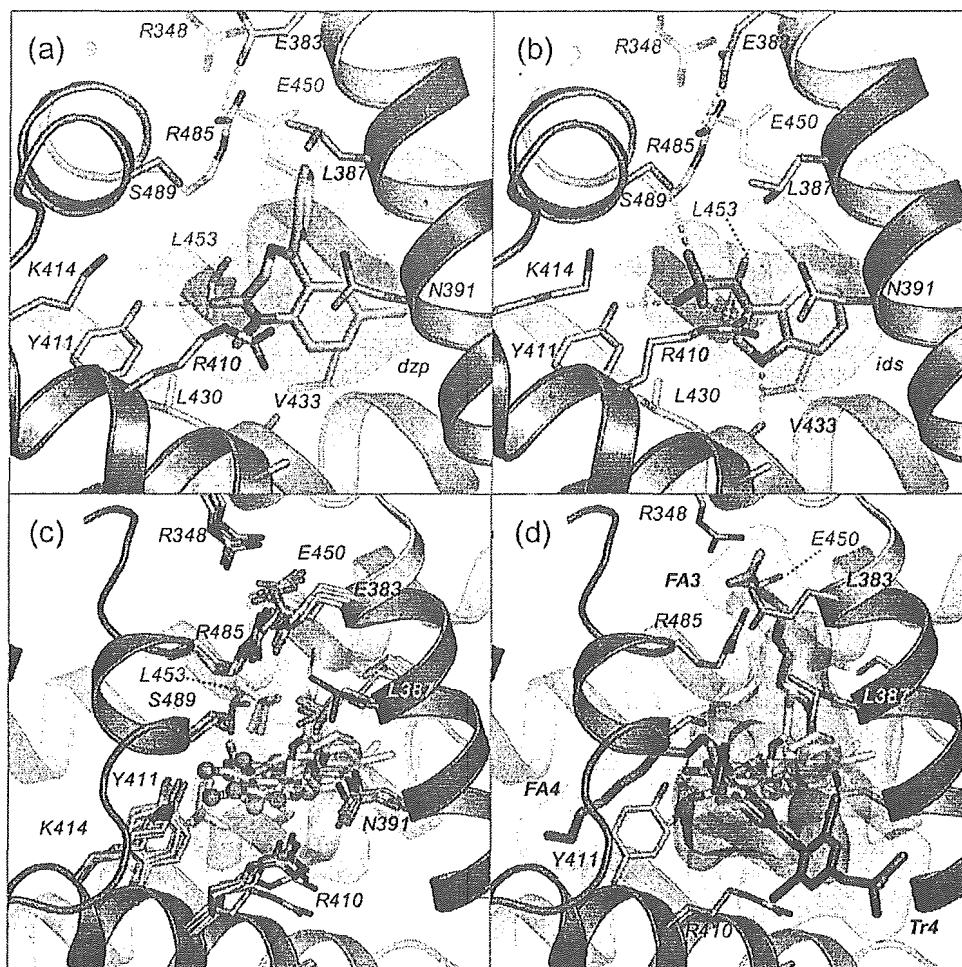


Figure 6. Drug binding to site 2 in HSA. The detailed binding conformations are shown for (a) diazepam and (b) indoxyl sulphate. In each case the drug is shown in a stick representation with a semi-transparent van der Waals surface (magenta). Colour-coding is as in Figure 3. (c) Top view of the superposition of drugs bound to site 2 in HSA along with a semi-transparent surface (orange) depicting the extent of the pocket. (d) Binding of endogenous ligands indicates possible expansion of drug site 2. Fatty acids (FA3 and FA4)^{3,4} and thyroxine⁶ which also bind to subdomain IIIA are added to the drug superposition shown in (c); the van der Waals surface defined for these endogenous ligands is coloured blue.

residues in drug site 1 (R218 and R222, respectively). Thus in terms of shape, size and polarity, drug sites 1 and 2 are clearly distinguishable and this helps to account for the different binding specificities of the two pockets.

Diflunisal, diazepam, ibuprofen and indoxyl sulphate all cluster in the centre of the binding pocket of subdomain IIIA, oriented with at least one oxygen atom in the vicinity of the polar patch (Figure 6(a)–(c); Supplementary Figure 4). In every case, there is an interaction with the hydroxyl group of Y410, whereas none of the drugs were found to interact with K414. R410 and S489 also contribute salt-bridge and hydrogen-bond interactions to drug binding, though not in the case of diazepam. Thus the observation that site 2 is generally selective for drugs with a peripherally located electronegative group² can be ascribed to the presence of a basic

polar patch located at one end of a generally apolar pocket in subdomain IIIA.

However, the uniform binding orientation of diflunisal, diazepam, ibuprofen and indoxyl sulphate contrasts with that of di-isopropyl phenol (propofol), a general anaesthetic drug. Due to steric effects of the isopropyl groups, the single polar hydroxyl group in the centre of the propofol molecule cannot interact with the main polar patch in drug site 2 and instead, propofol adopts a conformation that allows formation of a hydrogen bond to the carbonyl oxygen of Leu430.¹⁶ Interestingly this carbonyl group also interacts with the indole amide of indoxyl sulphate and the bromine atom of halothane¹⁶ and appears to constitute a secondary polar feature in the pocket (Supplementary Figure 4(d) and (g)).

There is comparatively little side-chain

movement associated with ligand binding if one considers just the smallest drugs (diflunisal, ibuprofen, halothane, indoxyl sulphate, and propofol; M_r 197–250 Da); V433 and R410 are the most susceptible to ligand-induced alterations (Figure 6(c)). However, binding of diazepam, which has a larger, branched structure (M_r 284.7 Da) is accompanied by large rotations of the side-chains of L387 and L453 that increases their separation from 5.4 Å to 7.7 Å and allows the phenyl ring of the drug to access the rear right-hand sub-chamber of the pocket. This pocket is closed off by R348-E450 and R485-E383 salt-bridges (Figure 6(c)). As in site 1, variations in the water structure, which was generally not visible at the resolutions of our structure determinations, may help to make the pocket more adaptable.

Further evidence of the adaptability of drug site 2 in subdomain IIIA derives from the fact that although it appears to be relatively small, it can bind two molecules of long-chain fatty acid (in fatty acid sites FA3 and FA4)⁴ or one of thyroxine.⁶ Comparison of drug and fatty acid binding reveals the very different ways in which these classes of ligand bind to a common locus on the protein (Figure 6(d)). In fact, drug site 2 is composed of the apolar region that is occupied by the methylene tails of fatty acids bound to FA3 and the polar patch that interacts with the carboxylate moiety of fatty acids bound to FA4. None of the drugs examined to date

is observed to access the long, narrow hydrophobic tunnel of FA4 that accommodates the methylene tails of lipids bound to this site. Moreover, fatty acids bound to FA3 do not interact with the polar patch centred on Y411. Instead, binding of the fatty acid opens access to a different polar patch by inducing the same rotations of L387 and L453 that are observed upon diazepam binding and the lipid carboxylate group supplants E450 in a salt-bridge interaction with R348 in subdomain IIB (Figure 6(d)). These observations suggest possible ways in which drugs might be modified in order to take advantage of this flexible binding facility in the pocket.

Fatty acid binding is also known to be associated with a large conformational change in HSA, involving rotations of domains I and III relative to domain II, which suggests a possible molecular mechanism for allosteric interactions between fatty acid binding sites.^{3,4,44} In contrast, the conformational changes observed for drug binding at sites 1 and 2 are more local; there is no evidence for the global conformational changes on the scale observed with fatty acid binding. The observed instances of allosteric interactions between drug sites 1 and 2^{28–30} may possibly be due to more subtle structural effects or to the presence of additional binding sites.

Several of the site 2 compounds analysed also bind to additional sites outside subdomain IIIA.

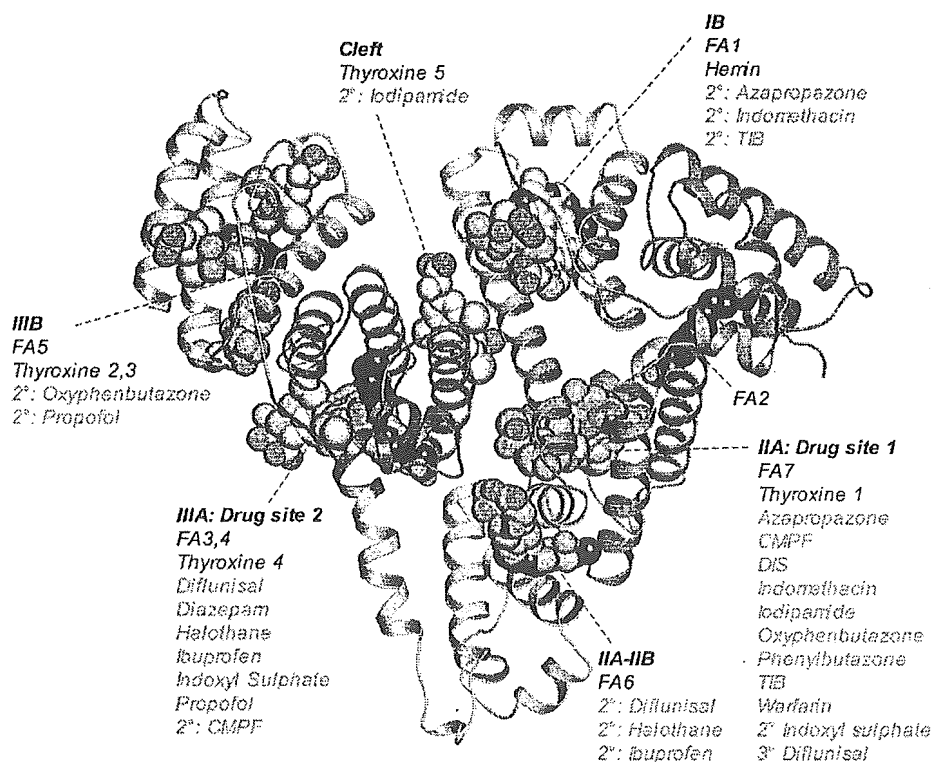


Figure 7. Summary of the ligand binding capacity of HSA as defined by crystallographic studies to date. Ligands are depicted in space-filling representation; oxygen atoms are coloured red; all other atoms in fatty acids (myristic acid), other endogenous ligands (hemin, thyroxin) and drugs are coloured dark-grey, light grey and orange, respectively.

A secondary binding site is observed for indoxyl sulphate in drug site 1 where two molecules of the compound appear to bind in overlapping and mutually exclusive conformations, one with the sulphate group positioned to interact with the inner polar patch and one in which the sulphate is salt-bridged to the outer patch at the pocket entrance. There is evidence to suggest that diflunisal and ibuprofen may also bind within site 1, though the density in the case of ibuprofen is rather weak and this drug was therefore not incorporated at this site in the refined model. In contrast the electron density maps clearly indicate that diflunisal and ibuprofen both occupy a previously undetected secondary site at the interface between subdomains IIA and IIB in a binding cleft that overlaps the fatty acid site FA6^{4,5} (Figure 7). The carboxylate groups of the drugs interact with the side-chains of K351 and S480 (of subdomain IIIA) and the amide groups of L481 and V482. In this locus both drugs pack against the helix (residues 209–223) which forms part of the entrance to drug site 1; conceivably binding of diflunisal or ibuprofen to this secondary site may therefore impact the binding of site 1 drugs^{28,29}. We did not observe a secondary site for diazepam.

Our structural data show that the two primary drug sites on HSA are highly adaptable binding cavities containing distinct sub-compartments, some of which are only accessed by local drug-induced conformational changes, and reveal a range of secondary binding sites distributed widely across the protein. In each case, the drug sites overlap with endogenous ligand-binding sites (Figure 7). The binding specificities of the pockets are determined by their shapes and the particular distributions of basic and polar residues on the largely hydrophobic interior walls that are involved in charge neutralization and hydrogen bonding interactions with acidic or electronegative small molecule ligands. The combination of shape-adaptability with specific polar interactions exhibited by HSA in these sites is reminiscent of the promiscuous binding site identified in QacR, a multi-drug binding protein from *Staphylococcus aureus*,⁴⁵ although, in contrast to HSA, QacR has a preference for cationic lipophilic drugs and its hydrophobic cavity is therefore studded with acidic glutamate side-chains. The detailed insights into HSA–drug interactions reported here provide an invaluable structural framework for the interpretation of drug binding data and will facilitate efforts to modify new therapeutic compounds to control their interaction with HSA and therefore optimise their distribution within the human body.

Materials and Methods

Protein purification, complex formation and crystallisation

Samples of purified recombinant HSA were kindly provided by Delta Biotechnology Ltd. (Nottingham, UK)

and Professor Eishun Tsuchida (Waseda University, Japan). Prior to crystallisation in the absence of fatty acid, the protein was defatted⁴⁶ and subjected to gel-filtration to ensure a purely monomeric preparation.^{3,16} Drugs were purchased as the highest purity preparations available from Sigma or Fluka. Azapropazone was kindly provided by Professor Ulrich Kragh-Hansen and CMPF was synthesised as described.³⁶ Crystals of defatted HSA generally do not tolerate soaking in ligand solutions so HSA–drug complexes were prepared before crystallisation by incubating the protein with a fivefold molar excess of drug at room temperature for 1–16 h. For example, 400 μ l of HSA at 100 mg/ml (1.5 mM) was mixed with 600 μ l of drug at 5 mM in 50 mM sodium phosphate buffer (pH 7). Where drug stock solutions were prepared in methanol or dimethyl sulphoxide, the maximum concentration of organic solvent at this stage was 7% (v/v). The free drug concentration was then fixed by repeated cycles of concentration and dilution in buffer containing 0.1 mM drug using a 10 kDa ultracentrifugation device (Millipore) and the protein concentration restored to 100 mg/ml; during this process any organic solvent present was reduced in concentration to less than 0.1% (v/v).

All crystals were grown by sitting-drop vapour diffusion using protein concentrations of around 100 mg/ml in 50 mM potassium phosphate (pH 7)^{4,16,47}. The HSA–drug complexes were crystallised typically by mixing 2.5 μ l of protein with 2.5 μ l of a reservoir solution containing 24–30% (w/v) polyethylene glycol 3350 (Sigma-Aldrich), 50 mM potassium phosphate (pH 7.0).

HSA–myristate complexes were prepared (without prior defatting of the protein) and crystallised as described.^{3,4} In all cases the largest crystals were obtained by streak- or micro-seeding into drops that had been allowed to equilibrate for 5–7 days.⁴⁸ Crystals were harvested into solutions containing slightly higher PEG 3350 concentrations than were used for crystallisation.⁴ In a deviation from our previous practice, myristate was omitted from the harvest buffer in order to favour drug displacement of the fatty acid. Ternary HSA–myristate–drug complexes were prepared by soaking crystals of HSA–myristate in a series of harvest buffer solutions containing increasing concentrations of the requisite drug; typically the starting concentration was 0.1 mM and this was doubled every few minutes or hours up to the maximum tolerable concentration (\sim 5 mM) (as judged by the fragmentation of crystals). Total soak times ranged from 2–48 h.

Data collection and structure determination

X-ray diffraction data were collected at room temperature using synchrotron radiation on station 9.6 at Daresbury SRS (UK) and stations BW7A, X11 and X13 at EMBL/DESY Hamburg (Germany) (Table 1). The data were indexed and measured with MOSFLM.⁴⁹ In all cases the HSA–drug complexes crystallised isomorphously with the P1 crystals of defatted HSA obtained previously in this laboratory.¹⁶ The protein model for this structure (PDB ID, 1e78) was used as a starting model for phasing of the X-ray data. The model, split into its six subdomains, was first refined as a rigid body using CNS (version 1.1)⁵⁰ and then subjected to cycles of positional and B-factor refinement interleaved with manual model corrections in O.⁵¹ Datasets for HSA–myristate–drug complexes were phased and refined in the same way using the original HSA–myristate structure (PDB ID 1e7g),^{3,4} stripped of all its ligands, as the starting model.

Following initial refinement of the protein structure, difference electron density maps showed clear density for bound drug molecules and in each case defined the orientation and conformation of the bound ligand. Where possible, models for the drug molecules or their constituent fragments were obtained from the Cambridge Structural Database via the Chemical Database Service⁵² and used to generate refinement dictionaries with XPLO2D.⁵³ In the case of CMPF, structure was generated using the Dundee PRODRG2 server.⁵⁴

The HSA–drug complexes (with or without myristate) were refined to resolutions of 2.25–3.2 Å; the models have R_{free} values in the range 24.3–29.2% and good stereochemistry (Table 1). Average B -factors for the different models are relatively high, at around 55 Å² for HSA–myristate models (C2 space group) and 74 Å² for HSA without fatty acid (P1 space group). For both crystal forms, subdomain IIIB consistently exhibits higher than average B -factors, indicative of greater mobility of this region.

Protein Data Bank atomic coordinates

Atomic co-ordinates have been deposited with the RCSB Protein Data Bank (ID codes are given in Table 1).

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Supplementary Data

Supplementary data associated with this article can be found at 10.1016/j.jmb.2005.07.075

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Effect of Nitric Oxide in Amyloid Fibril Formation on Transthyretin-Related Amyloidosis[†]

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ABSTRACT: Although oxidative stress is said to play an important role in the amyloid formation mechanism in several types of amyloidosis, few details about this role have been described. Amyloid is commonly deposited around the vessels that are the primary site of action of nitric oxide generated from endothelial cells and smooth muscle cells, so nitric oxide may be also implicated in amyloid formation. For this study, we examined the *in vitro* effect of S-nitrosylation on amyloid formation induced by wild-type transthyretin, a precursor protein of senile systemic amyloidosis, and amyloidogenic transthyretin V30M, a precursor protein of amyloid deposition in familial amyloidotic polyneuropathy. S-Nitrosylation of amyloidogenic transthyretin V30M via the cysteine at position 10 was 2 times more extensive than that of wild-type transthyretin in a nitric oxide-generating solution. Both wild-type transthyretin and amyloidogenic transthyretin V30M formed amyloid fibrils under acidic conditions, and S-nitrosylated transthyretins exhibited higher amyloidogenicity than did unmodified transthyretins. Moreover, S-nitrosylated amyloidogenic transthyretin V30M formed more fibrils than did S-nitrosylated wild-type transthyretin. Structural studies revealed that S-nitrosylation of amyloidogenic transthyretin V30M induced a change in its conformation, as well as instability of the tetramer conformation. These results suggest that the nitric oxide-mediated modification of transthyretin, especially variant transthyretin, may play an important role in amyloid formation in senile systemic amyloidosis and familial amyloidotic polyneuropathy.

Progress in molecular genetics and biochemical methodologies has led to the identification of various types of amyloidosis and related amyloidogenic precursor proteins (1, 2). However, the precise mechanism of amyloid formation remains to be elucidated. Familial amyloidotic polyneuropathy (FAP),¹ which is induced by amyloidogenic transthyretin (ATTR), is characterized by systemic accumulation of amyloid fibrils in the peripheral nerves and other organs (3, 4). Of the different types of ATTR-related amyloidosis, FAP

ATTR V30M is most common (3–5). Because most of the transthyretin (TTR) is produced by the liver, progression of the disease can be halted by liver transplantation (6–9). However, no other known therapy can prevent amyloid fibril formation *in vivo*, mainly because the mechanism of amyloid fibril formation in FAP is not well understood. In addition, amyloid deposits from elderly patients with systemic senile amyloidosis (SSA) have been proven to be derived from the wild-type (WT) TTR molecule. Details of the amyloid formation mechanism in SSA have also not been clarified (10).

Injury caused by free radicals has been implicated in the amyloid formation process in several types of systemic amyloidosis and localized amyloidosis, such as Alzheimer's disease (11, 12), β_2 -microglobulin amyloidosis (13), and FAP (14), and administration of scavengers of these free radicals has been suggested as a method of preventing amyloid fibril formation (15). Of the reactive oxygen species, reactive nitrogen species such as peroxynitrite and nitrogen dioxide have been proposed to be involved in the etiology of diverse pathophysiological conditions, including inflammation, neu-

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¹ Abbreviations: FAP, familial amyloidotic polyneuropathy; ATTR, amyloidogenic transthyretin; TTR, transthyretin; SSA, systemic senile amyloidosis; WT, wild-type; IAN, isoamyl nitrate; DTT, 1,4-dithiothreitol; DTNB, 5,5'-dithiobis(2-nitrobenzoic acid); cTTRs, TTRs that underwent the same procedure as the S-nitrosylated TTRs but without IAN; NOC18, 1-hydroxy-2-oxo-3,3'-bis(2-aminoethyl)-1-triazene; GSNO, S-nitrosoglutathione.

rodegenerative diseases, cardiovascular disorders, cancer, and amyloidosis (16, 17). S-Nitrosothiols are adducts of nitric oxide (NO[•]) and thiol-containing compounds that are found in many mammalian tissues and may affect various physiological functions, including signal transduction and immune responses (18). The bioactivities of low-molecular-weight S-nitrosothiols may be mediated either by their decomposition to NO[•] or by direct exchange of their nitrosonium cation (NO⁺) function with thiol-containing proteins (transnitrosylation) (19).

TTR forms tetramers with thyroxine and retinol-binding protein. Tetramers of amyloidogenic variants of TTR have been suggested to be more unstable than those of WT TTR (20), and partial acidic denaturation of the tetramers leads to formation of monomeric amyloidogenic intermediates (21). In addition, post-translational modification of TTR is also known to lead to amyloid fibril formation more easily than in native TTRs (22–24). TTR has one cysteine (Cys) at position 10, which is prone to undergoing several post-translational modifications such as cysteinylolation, sulfonation, glutathionation, and nitrosylation. In addition, structural analysis has revealed that the Cys residue of ATTR V30M is located more to the outside of the molecule, which suggests that ATTR V30M may undergo modifications more frequently than the WT TTR via the Cys moiety (25). The variant TTR is a more highly amyloidogenic protein than WT TTR; the difference in the degree of post-translational modification between WT TTR and variant TTR also explained the amyloid fibril formation in FAP and SAA (25).

We recently reported the existence of oxidative stress in deposits of amyloid in tissues of patients with FAP as well as in those with other amyloidoses (14). Nitrotyrosine, a product of nitration, has also been reported to be present around amyloid deposited in transgenic mice with ATTR V30M (26). These results suggest that oxidative stress, especially that induced by NO, may play an important role in amyloid formation in FAP.

The aim of this study was to elucidate the effect of NO on amyloid fibril formation in FAP and SSA. We used biochemical methods to compare, in the presence or absence of S-nitrosylation, the degree of amyloid formation induced by both types of TTRs after changes in TTR structure, as well as the stability of the tetrameric forms of both TTRs.

EXPERIMENTAL PROCEDURES

Materials. Both WT TTR and ATTR V30M were purified from serum samples obtained from healthy volunteers and homozygotic FAP ATTR V30M patients as described previously (27). A brief description of this method follows.

Cold Ethanol Fractionation. Pooled serum samples from normal volunteers and homozygotic FAP ATTR V30M patients (200 mL) were mixed for 1 h at –5 °C with 8% ethanol at pH 7.5 and were then centrifuged at 10000g for 30 min. The supernatant was mixed for 1 h at –5 °C with 21% ethanol at pH 6.8 and was then centrifuged at 10000g for 30 min. This supernatant was dialyzed against 50 mM Tris-HCl (pH 7.6) and was then concentrated to 2 mL by ultrafiltration by using a Minisette Model ultrafiltration apparatus (Pall Technical Center, Port Washington, NY) and an OMEGA 10 kDa cutoff membrane (Filtron, Carbondale, PA). The resulting crude TTR preparation was used as the

starting material in further purification steps. Affinity chromatography with 3 mg of human retinol-binding protein (Cortex Biochem, San Leandro, CA) was linked to 1.0 mL of HiTrap NHS-activated Sepharose 4FF (Amersham Pharmacia Biotech, Tokyo, Japan) according to Amersham Pharmacia Biotech's standard protocol. The column was equilibrated with 50 mM Tris-HCl (pH 7.6). A sample of 2 mL of starting material was applied to the column, and flow-through fractions with absorbance at 280 nm of more than 0.1 were collected as TTR-enriched fractions.

Anion-Exchange Column Chromatography. A 50 mL sample of the TTR-enriched fractions was passed through a 1.0 mL Resource Q column (Amersham Pharmacia Biotech) equilibrated with 50 mM Tris-HCl (pH 7.6). Unbound material was washed out with 50 mM Tris-HCl (pH 7.6) containing 70 mM NaCl. The bound fraction (crude TTR) was eluted with 50 mM Tris-HCl (pH 7.6) containing 200 mM NaCl, and was then concentrated to 5 mL by ultrafiltration with the Minisette Model ultrafiltration apparatus.

Removal of Albumin by Affinity Chromatography with Anti-Human Albumin Antibody. Anti-human albumin antibody (ICN, Tokyo, Japan) (25 mg) was linked to 5.0 mL of HiTrap NHS-activated Sepharose 4FF (Amersham Pharmacia Biotech) according to Amersham Pharmacia Biotech's standard protocol. The column was equilibrated with 50 mM Tris-HCl (pH 7.6) containing 0.3 M NaCl. An aliquot (5 mL) of the crude TTR-enriched fraction was passed through this column, and TTR devoid of albumin in the flow-through fractions was collected.

Gel Filtration Chromatography. The partially purified TTR from the anti-human albumin affinity chromatography column was concentrated to 2 mL by ultrafiltration on a Centrplus-10 concentrator (Millipore, Bedford, MA). This material was applied to a TSK gel G3000SW column (10 mm × 60 cm) (Tosoh, Tokyo, Japan) equilibrated with 50 mM sodium acetate buffer (pH 6.7) containing 0.3 M NaCl, at a flow rate of 5 mL/min. The TTR peak was collected at an elution time of approximately 30 min. This sample was then concentrated to 2 mL by ultrafiltration on a Centricon-10 concentrator (Millipore), after which the concentrated sample was dialyzed against 0.15 M NaCl.

Analysis of the Purity by Sodium Dodecyl Sulfate–Polyacrylamide Gel Electrophoresis (SDS–PAGE) and High-Pressure Liquid Chromatography (HPLC). SDS–PAGE (TEFCO, Tokyo, Japan) was performed as previously described by Harmansen et al. (28). Gels were stained with Coomassie Brilliant Blue R-250. HPLC analysis (29) was performed on a TSK gel G3000SWXL column (4.6 mm × 30 cm) (Tosoh) equilibrated with 20 mM sodium phosphate buffer (pH 7.0) containing 150 mM NaCl, at a flow rate of 0.4 mL/min.

S-Nitrosylation of WT TTR and ATTR V30M. WT TTR and ATTR V30M were S-nitrosylated by using isoamyl nitrate (IAN) from Wako (Osaka, Japan) as described previously (30–32). Before S-nitrosylation of TTRs, each TTR (20 μM) was treated with 200 μM 1,4-dithiothreitol (DTT) in 100 mM sodium phosphate buffer (pH 7.4) with 5.0 mM ethylenediamine-*N,N,N',N'*-tetraacetic acid at 37 °C for 5 min, followed by Sephadex G-25 column chromatography to obtain reduced forms of TTRs. The contents of the free cysteine of TTRs were ~1 mol of the protein as determined by the 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB)

method (33). DTT-treated TTRs (20 μ M) were then reacted with 200 μ M IAN in 100 mM sodium phosphate buffer (pH 7.8) containing 0.5 mM diethylenetriamine pentaacetic acid at 37 °C for 1 h. *S*-Nitrosylated products were purified by Sephadex G-25 gel filtration and eluted with 100 mM sodium phosphate buffer (pH 7.4). TTRs that underwent the same procedure as the *S*-nitrosylated TTRs but without IAN (cTTRs) were also made in an effort to compare their reactivities with those of *S*-nitrosylated TTRs.

Determination of *S*-Nitrosothiols Levels in WT TTR and ATTR V30M. *S*-Nitroso compounds formed in the reaction mixture were analyzed by means of the HPLC flow reactor system to identify *S*-nitrosothiols. *S*-Nitroso compounds, including *S*-nitroso proteins eluted from the HPLC column, were decomposed with Hg^{2+} and detected as nitrite (NO_2^-) after the reaction with the Griess reagent (34). A deproteinization column was used in this system for determination of *S*-nitroso protein levels just before the reaction with the Griess reagent. The stability of *S*-nitrosothiols associated with TTRs (Figure 3) was analyzed by a colorimetric assay using the Griess reagent according to Saville's method (35). Briefly, 50 μ g of *S*-nitrosylated TTRs was incubated with a 10-fold molar excess of Hg^{2+} in 10 mM sodium acetate buffer (pH 5.5) at 37 °C for 30 min, and reacted with the Griess reagent at 40 °C for 10 min. The amount of NO_2^- released from each *S*-nitroso moiety was quantified with the Griess reagent reaction by measuring the absorbance at 540 nm. Simultaneously, 50 μ g of the *S*-nitrosylated TTRs solution without Hg^{2+} treatment was reacted with the Griess reagent. The quantities of *S*-nitroso moieties in *S*-nitrosylated TTRs were assessed by comparing the absorbance observed with Hg^{2+} -untreated *S*-nitrosylated TTRs to that of Hg^{2+} -treated *S*-nitrosylated TTRs.

Measurements of Circular Dichroism (CD) Spectra. CD spectra were obtained by using a JASCO (Tokyo, Japan) J-720 spectropolarimeter at 25 °C. A molecular mass of TTR of 14 kDa was used for calculation of the mean residue ellipticity. Far-UV and CD spectra were recorded at a protein concentration of 10 μ M, in 67 mM sodium phosphate (pH 7.4).

Tryptophan (Trp) Fluorescence Intensity. The fluorescence intensity of Trp was measured via a Hitachi (Tokyo, Japan) F-4500 spectrofluorimeter at 25 °C. All assays used excitation at 295 nm and emission at 340 nm. Excitation and emission slits were set at 5 nm. The fluorescence intensity was recorded at a protein concentration of 10 μ M, in 67 mM sodium phosphate (pH 7.4).

Amyloid Formation Induced by WT TTR and ATTR V30M. To evaluate amyloid fibril formation after *S*-nitrosylation, unmodified or *S*-nitrosylated WT TTR and ATTR V30M were diluted in 50 mM sodium acetate and 100 mM NaCl at the desired pH (3.0–6.5) in an Eppendorf tube to obtain a final TTR concentration of 15 μ M. The resulting stationary solutions were incubated at 37 °C for 5 days in the dark (36).

Thioflavin T-Based Fluorimetric Assays. To assess the amount of amyloid fibrils in vitro, the thioflavin T test has been widely used (37, 38). Fluorescence spectra were obtained by using a Hitachi F-4500 spectrofluorimeter with an assay volume of 1 mL. All assays used excitation at 450 nm and emission at 482 nm. Excitation and emission slits were set at 5 nm. The reaction mixture contained 5 μ M

thioflavin T and 50 mM Gly-NaOH buffer (pH 10.0) (37, 38). Ten microliters of a 15 μ M WT TTR or ATTR V30M solution was added to 1 mL of the reaction mixture. Spectra were recorded at 25 °C within minutes of the addition of the sample to the reaction mixture.

Amyloid Formation from TTRs in the Presence of Exogenous NO. To examine the direct effect of NO on the amyloidogenicity of TTRs, TTRs were reacted with various NO (NO^+)-releasing reagents, such as 1-hydroxy-2-oxo-3,3-bis(2-aminoethyl)-1-triazene (NOC18; Dojindo Laboratories, Kumamoto, Japan) and *S*-nitrosoglutathione (GSNO) (39). Samples of 15 μ M unmodified or *S*-nitrosylated WT TTR and ATTR V30M were incubated in 50 mM sodium acetate and 100 mM NaCl at pH 3.0 in the presence or absence of 50 μ M NOC18 and GSNO at 37 °C in the dark for 5 days. Amyloidogenicity was examined under the same conditions that are described above.

Nonboiled (Nonreducing) SDS-PAGE. Nonboiled SDS-PAGE was performed under nondenaturing conditions. One microgram of the TTR samples incubated for 37 °C for 5 days at pH 3.0 as described above was neutralized with PBS to obtain a final pH of >6.5. After neutralization, samples were mixed with 5% SDS sample buffer and loaded on 15% polyacrylamide gels, which were stained with Coomassie Brilliant Blue. Band intensities were evaluated by densitometric analysis using ATTO densito (ATTO, Tokyo, Japan).

Statistical Analysis. Statistical evaluation was performed by means of the paired *t* test. A *p* value of <0.05 was taken to be statistically significant.

RESULTS

Ratio of *S*-Nitrosylation of TTR Molecules. The ratio of *S*-nitrosylation of TTRs was determined by titrating *S*-nitrosothiol levels. The ratio of *S*-nitrosothiols for ATTR V30M was much higher than that for WT TTR (WT TTR, 0.39 mol/mol of protein; ATTR V30M, 0.67 mol/mol of protein). Dissociation of tetrameric forms to monomeric forms in TTRs was not recognized by nonreducing SDS-PAGE (data not shown).

Conformational Changes of *S*-Nitrosylated TTRs. The conformations of *S*-nitrosylated WT TTR and ATTR V30M were compared with those of unmodified TTRs by means of CD. Unmodified and *S*-nitrosylated WT TTR and ATTR V30M showed no significant difference in conformation in far-UV spectral analyses (Figure 1). However, *S*-nitrosylation of WT TTR induced a slight change in Trp fluorescence intensity at the 340 nm wavelength. Moreover, the Trp fluorescence intensity for *S*-nitrosylated ATTR V30M decreased to a greater degree compared with the change in Trp fluorescence intensity for WT TTR (Figure 2). CD spectra and the Trp fluorescence intensity of cTTRs did not significantly differ from that of unmodified TTRs (data not shown).

Stability of *S*-Nitrosothiol Associated with TTRs. The stability of *S*-nitrosothiol associated with WT TTR and ATTR V30M was measured at pH 3.0 (Figure 3A) and pH 7.0 (Figure 3B) in the dark. NO associated with TTR dissociated from the protein in a time-dependent manner: dissociation from TTR molecules started just after incubation began, and little NO remained associated with both TTRs after day 10 at pH 3.0 (Figure 3A). At pH 3.0, the half-life

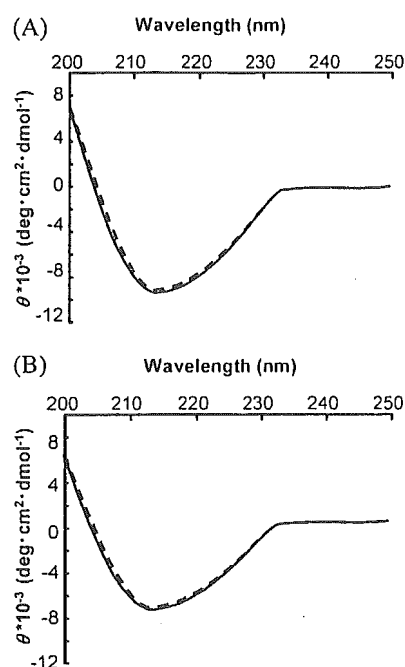


FIGURE 1: Far-UV CD spectral analysis. Samples (1.5 mL) of unmodified or *S*-nitrosylated WT TTR and ATTR V30M [$10 \mu\text{M}$ in 67 mM sodium phosphate (pH 7.4)] were applied to a JASCO J-720 spectropolarimeter as described in the text. (A) Unmodified WT TTR (—) and its *S*-nitrosylated form (---). (B) Unmodified ATTR V30M (—) and its *S*-nitrosylated form (---).

of *S*-nitrosylated TTRs was 6.6 days in WT TTR and 6.3 days in ATTR V30M. In contrast, at pH 7.0, more than 60% of NO associated with WT TTR and ATTR V30M remained unchanged 15 days after incubation started (Figure 3B). At pH 7.0, the half-life of *S*-nitrosylated TTRs was 22.05 days in WT TTR and 18.3 days in ATTR V30M. At both pH, *S*-nitrosothiols were more rapidly decayed in ATTR V30M than in WT TTR.

Effect of *S*-Nitrosylation on Amyloid Formation Induced by TTRs. The effect of *S*-nitrosylation on amyloid formation induced by WT TTR and ATTR V30M was evaluated by means of the thioflavin T-based test as described above. Both WT TTR and ATTR V30M exhibited increased fluorescence intensity as the pH of the incubation solution decreased (Figure 4). The fluorescence intensity of unmodified ATTR V30M was higher than that of WT TTR, and both *S*-nitrosylated TTRs had a higher intensity than unmodified TTRs. In addition, *S*-nitrosylated ATTR V30M exhibited greater fluorescence intensity than *S*-nitrosylated WT TTR. The fluorescence intensity of cTTRs did not significantly differ from that of unmodified TTRs (data not shown).

Amyloid Formation of TTRs in the Presence of Exogenous NO. To determine whether formation of *S*-nitrosothiols or direct inhalation of the proteins with NO affected amyloid fibril formation during incubation of *S*-nitrosylated TTRs, unmodified TTR and ATTR V30M were incubated with NOC18 or GSNO at pH 3.0 and 37 °C in the dark for 5 days, and amyloidogenicity was compared by means of the thioflavin T-based test. As demonstrated in Figure 5, TTRs incubated with NOC18 under acidic conditions showed no significant increase in the extent of amyloid formation, compared with unmodified TTRs. *S*-Nitrosylated WT TTR and *S*-nitrosylated ATTR V30M and ATTR V30M incubated with GSNO exhibited greater amyloidogenicity than unmodified TTR and

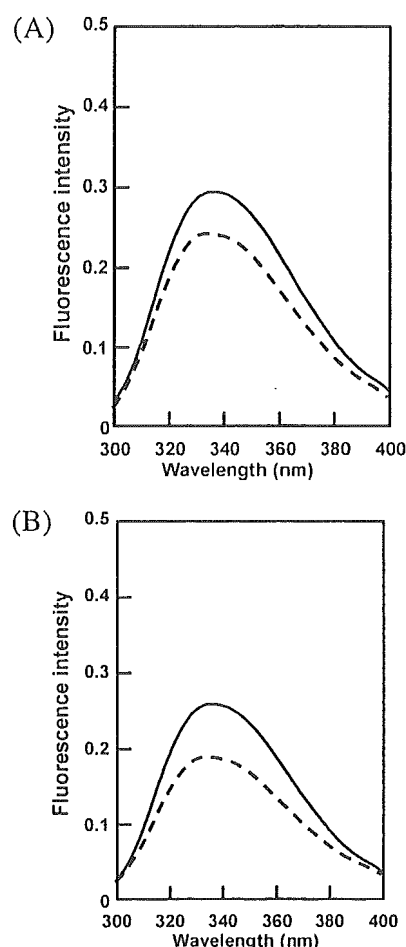


FIGURE 2: Trp fluorescence intensity of TTRs. The Trp fluorescence intensities of unmodified or *S*-nitrosylated WT TTR and ATTR V30M were monitored by using a Hitachi F-4500 spectropolarimeter. Aliquots of 2 mL of $10 \mu\text{M}$ WT TTR and ATTR V30M [in 67 mM sodium phosphate (pH 7.4)] were used in the experiments. (A) Unmodified WT TTR (—) and its *S*-nitrosylated form (---). (B) Unmodified ATTR V30M (—) and its *S*-nitrosylated form (---).

NOC18-treated TTR (unmodified ATTR V30M vs *S*-nitrosylated ATTR V30M, $p < 0.01$).

Effect of *S*-Nitrosylation on the Stability of the Tetrameric Form of TTRs. To examine the effect of *S*-nitrosylation on the stability of the tetrameric form of TTRs, nonboiled (nonreducing) SDS-PAGE was performed with unmodified and *S*-nitrosylated TTRs (Figure 6A). The ratio of the monomeric band to the tetrameric band was decreased in *S*-nitrosylated TTR, especially ATTR V30M. Figure 6B presents quantified results and shows that *S*-nitrosylated TTRs had greater proportions of monomers than did unmodified TTRs. The *S*-nitrosylated ATTR V30M exhibited the highest ratio of monomer to tetramer.

DISCUSSION

We demonstrated here that *S*-nitrosylated TTRs possessed increased amyloidogenicity in the amyloid-generating system *in vitro*, compared with unmodified TTRs, and that *S*-nitrosylated ATTR V30M induced more amyloid fibrils than did WT TTR. Conformational analyses have documented that Cys at position 10 in ATTR V30M is situated more to the outside of the molecule than in WT TTR (25). This finding suggested that the Cys of ATTR V30M may be more

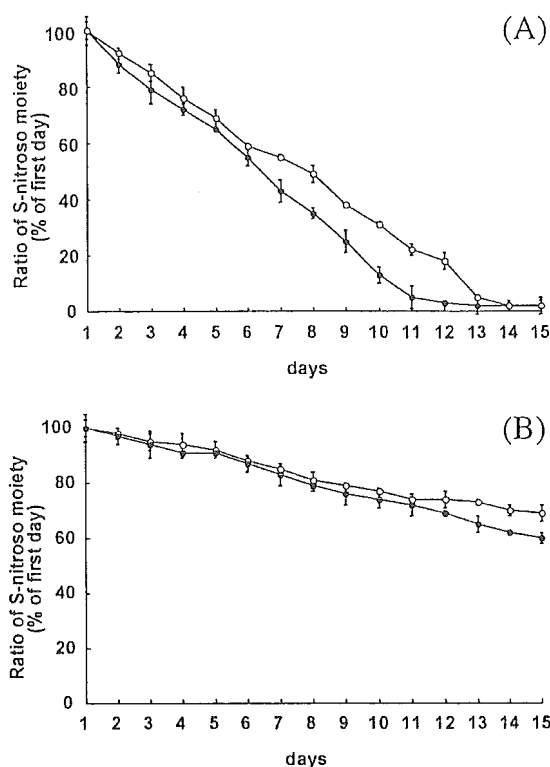


FIGURE 3: Stability of *S*-nitrosothiols associated with TTRs. This stability was examined until day 15 after the incubation started. *S*-Nitroso compounds formed in the reaction mixture were analyzed with the HPLC flow reactor system to identify *S*-nitrosothiols as described in the text. Samples of 50 μ g of unmodified or *S*-nitrosylated WT TTR and ATTR V30M were used in the incubations at pH 3.0 (A) and pH 7.0 (B) and 37 $^{\circ}$ C in the dark: (○) WT TTR and (●) ATTR V30M.

accessible to NO and thus susceptible to *S*-nitrosylation. In fact, the extent of *S*-nitrosylation of ATTR V30M was greater than that of WT TTR. Figure 4 indicates that *S*-nitrosylated ATTR V30M exhibited a greater thioflavin T fluorescence intensity than did WT TTR after incubation under acidic conditions. These findings demonstrate that the amount of amyloid fibrils formed increased as the ratio of *S*-nitrosylation increased.

Conformational study of *S*-nitrosylated TTRs revealed that, although far-UV spectral analyses indicated no significant difference between unmodified and *S*-nitrosylated proteins (Figure 1), the Trp fluorescence intensity at the 340 nm wavelength was significantly decreased in *S*-nitrosylated TTRs compared with unmodified TTRs, and *S*-nitrosylated ATTR V30M showed a greater reduction in intensity compared with wild-type TTR (Figure 2). These results suggest that, although *S*-nitrosylation of Cys at position 10 may not have a great influence on TTR structure, this modification may affect the microstructure around Trp at positions 41 and 79.

We also examined the possibility of dissociation of NO from TTRs during incubation at different pHs (Figure 3). Although *S*-nitrosothiols formed in proteins and peptides appear to be stable under acidic conditions (40), *S*-nitrosylated TTRs were found to be labile in an acidic solution (pH 3.0). Specifically, more than 60% of *S*-nitrosylated WT TTR and ATTR V30M remained unchanged at pH 7.0 15 days after incubation began. However, at pH 3.0, NO started to dissociate from the TTR molecule, and little NO remained

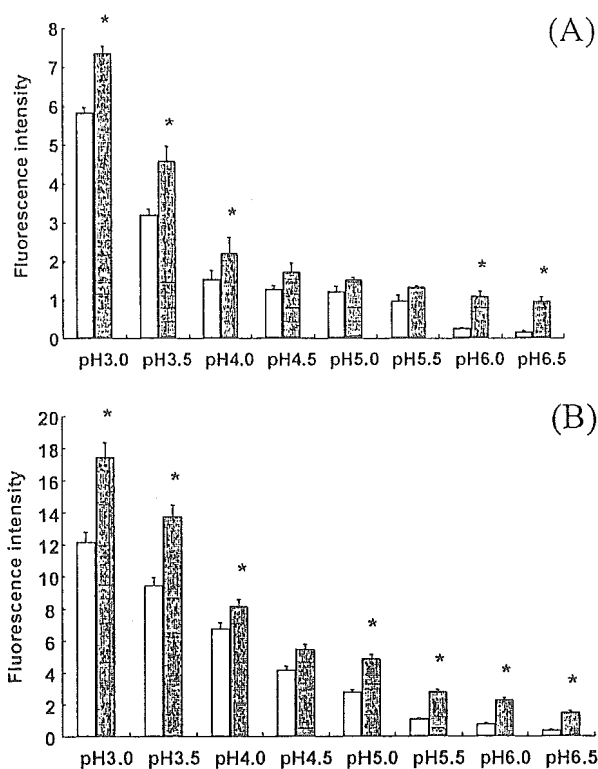


FIGURE 4: Effect of *S*-nitrosylation on amyloid formation induced by TTRs. Unmodified and *S*-nitrosylated WT TTR and ATTR V30M were diluted in 20 mM sodium acetate (pH 3.0–6.5) and 100 mM NaCl at the desired pH in an Eppendorf tube to obtain a final TTR concentration of 15 μ M. The resulting stationary solutions were incubated at 37 $^{\circ}$ C for 5 days. The data were expressed as the increased values compared to those of the time just starting the incubation with (A) WT TTR and (B) ATTR V30M: (white columns) unmodified TTRs and (gray columns) *S*-nitrosylated TTRs. In panel A, an asterisk indicates that $p < 0.01$, for unmodified WT TTR vs *S*-nitrosylated WT TTR. In panel B, an asterisk indicates that $p < 0.01$, for unmodified ATTR V30M vs *S*-nitrosylated ATTR V30M.

associated with the TTRs on day 15 at this pH value. At both pHs, *S*-nitrosothiols were more rapidly decayed in ATTR than in WT TTR. This NO dissociation may be induced by conformational changes of TTRs occurring at low pH, which facilitates dissociation of TTR tetramers to monomers. As demonstrated in Figure 6, in *S*-nitrosylated ATTR V30M, the ratio of monomeric forms to tetramers was much greater than in *S*-nitrosylated WT TTR.

In amyloid deposits in FAP as well as other amyloidoses, accumulation of human neutrophil elastase and cathepsin G derived from polymorphonuclear leukocytes has been documented (41). This result suggests that free radical species, including NO, may play a role in the amyloid formation mechanism in FAP. Therefore, it is quite important to elucidate whether the increased amyloidogenicity depends solely on the effect of *S*-nitrosylation of TTRs. We incubated both unmodified WT TTR and ATTR V30M with NOC18 or GSNO to generate NO or nitrosonium cation (NO^+) (39). The thioflavin T fluorescence intensity of TTRs did not significantly increase with NOC18 treatment. Incubation with GSNO, however, produced a significant increase in the thioflavin T fluorescence intensity of *S*-nitrosylated TTRs compared with unmodified and NOC18-incubated TTRs. Although NO generated from NOC18 cannot react with TTRs under acidic conditions, NO^+ generated from GSNO

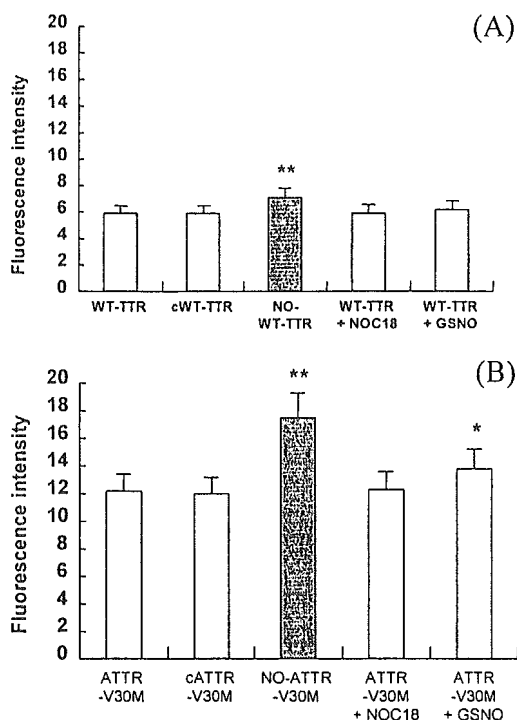


FIGURE 5: Amyloid formation of TTRs in the presence of exogenous NO. Samples of 15 μ M unmodified or S-nitrosylated WT TTR and ATTR V30M were incubated in 50 mM sodium acetate and 100 mM KCl at pH 3.0 in the presence or absence of 50 μ M NOC18 and GSNO. Amyloidogenicity was examined under the same conditions that are described in the legend of Figure 4: (A) WT TTR and (B) ATTR V30M. cTTRs are TTRs that underwent the same procedure as the S-nitrosylated TTRs but without IAN. Two asterisks indicate that $p < 0.01$, for unmodified TTRs vs S-nitrosylated TTRs at pH 3.0. One asterisk indicates that $p < 0.05$, for unmodified ATTR V30M vs TTR incubated with GSNO.

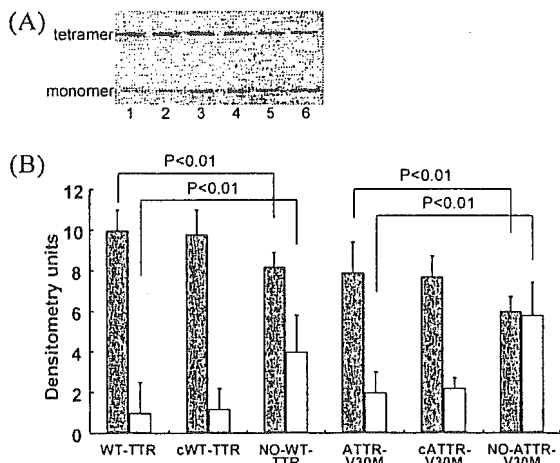


FIGURE 6: Effect of S-nitrosylation on the stability of the tetrameric forms of TTRs. Samples of 15 μ M unmodified or S-nitrosylated WT TTR and ATTR V30M were incubated at 37 $^{\circ}$ C for 5 days in acetate buffer (pH 3.0). (A) Samples were analyzed via nonboiled (nonreducing) SDS-PAGE as described in the text: lane 1, WT TTR; lane 2, WT cTTR; lane 3, NO-bound WT TTR; lane 4, ATTR V30M; lane 5, cATTR V30M; and lane 6, NO-bound ATTR V30M. (B) Intensities of the bands were evaluated by densitometric analysis (ATTO densito, ATTO, Tokyo, Japan): (white columns) TTR monomers and (gray columns) TTR tetramers.

induces transnitrosylation, which occurs even at acidic pH, and in those solutions, TTRs may undergo S-nitrosylation (19). These results suggest that NO dissociated from TTR may not directly participate in amyloid fibril formation.

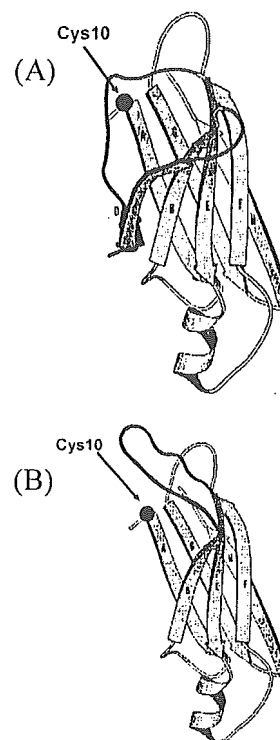


FIGURE 7: Conformation of WT TTR and ATTR V30M. A change in TTR structure caused by an amino acid substitution may induce amyloid formation by facilitating dissociation of TTR tetramers to monomers, which leads to amyloid fibril formation via misfolding C-D strands of the TTR molecule. S-Nitrosylation in TTRs may be one of the factors facilitating dissociation of TTR tetramers to monomers.

It is widely accepted that a change in TTR structure caused by an amino acid mutation may induce amyloid formation by facilitating dissociation of TTR tetramers to monomers (42) (Figure 7). To elucidate the mechanism of the increased amyloidogenicity in S-nitrosylated TTRs, we used nonboiled (nonreducing) SDS-PAGE to examine the ratio of tetramers to monomers for S-nitrosylated WT TTR and ATTR V30M. As Figure 6 shows, S-nitrosylated TTRs had higher ratios of monomeric to tetrameric forms than did unmodified TTRs, and the proportion of the monomeric form of S-nitrosylated ATTR V30M was greater than that of S-nitrosylated WT TTR. These results indicate that the increase in the extent of monomer formation occurs depending on S-nitrosylation of TTRs. This analysis and other conformational studies thus suggest that S-nitrosylation of TTRs induced a change in TTR structure, leading to reduced tetrameric stability and enhancing the amyloidogenicity of TTRs.

Unmodified Cys residues of circulating plasma proteins such as albumin and α_1 -protease inhibitor undergo S-nitrosylation, and these S-nitrosylated proteins protect against tissue injury (43). However, as Figure 5 demonstrates, S-nitrosylated TTR would function as a harmful protein for amyloid fibril formation, whereas S-nitrosylated TTR in the bloodstream may work as a beneficial molecule for tissue injury, like S-nitrosylated albumin and α_1 -protease inhibitor (18, 44, 45). Further study is needed.

In all types of TTR-related FAP, perivascular sites are known to show the greatest level of amyloid deposition, although the reason for this finding has not been well explained. These lesions are locations of NO reactions, where endothelial cells of vessels and smooth muscles generate NO.

In fact, nitrotyrosine antibody exhibited perivascular immunoreactivity at sites of abundant amyloid deposition in transgenic mice with ATTR V30M (26).

In conclusion, ATTR V30M was more prone to undergoing S-nitrosylation than WT TTR. S-Nitrosylated TTRs, especially ATTR V30M, induced formation of amyloid fibrils after a change in conformation. FAP, whose precursor protein is primarily ATTR V30M, usually starts to produce clinical manifestations when patients are ~35–45 years old; SSA, whose precursor protein is WT TTR, usually starts to produce clinical cardiac symptoms when patients are more than 80 years old (10). ATTR V30M is more prone than WT TTR to undergoing post-translational modification via the Cys residue at position 10. This distinction may contribute to the difference in the time of onset of these two types of systemic amyloidoses.

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The structure and function of oxidized albumin in hemodialysis patients: Its role in elevated oxidative stress via neutrophil burst

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Abstract

Oxidized albumin is a reliable marker of oxidative stress in hemodialysis (HD) patients. However, oxidized albumin in vivo and its possible clinical significance has been rarely investigated. In the present study, the qualitative modification of albumin in HD patients ($n = 20$) was examined and their results were compared with healthy age-matched controls ($n = 10$). The increase in plasma protein carbonyl levels in HD patients was largely due to an increase in oxidized albumin. Human serum albumin (HSA) of HD patients, HSA of HD patients (HD-HSA) and normal subjects (Normal-HSA) were purified on a blue Sepharose CL-6B column. Spectroscopic analysis confirmed that the HD-HSA samples contained higher levels of carbonyls than Normal-HSA. An HPLC analysis also suggested that the state of the purified HSA used throughout the experiments accurately reflects the redox state of albumin in blood. HD-HSA was found to have a decreased the antioxidant activity, and was able to trigger the oxidative burst of human neutrophils, compared to Normal-HSA. HD-HSA was conformationally altered, with its hydrophobic regions more exposed and to have a negative charge. In binding experiments, HD-HSA showed impaired Site II-ligand binding capabilities. Collectively, the oxidation of plasma proteins, especially HSA, might enhance oxidative stress in HD patients.

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The oxidative modification of proteins and lipids has been implicated in the etiology of numerous disorders and diseases [1,2]. Oxidatively modified plasma proteins can serve as important in vivo biomarkers of oxidative stress. Proteins are better candidates than plasma lipids for use in detecting specific pathways of oxidative stress, due to the accessibility of plasma protein for sampling, their relatively long half-lives, and their well-defined biochemical pathways.

Since extracellular fluids contain only small amounts of antioxidant enzymes, it has been proposed that the

major extracellular antioxidants are proteins and circulating albumin is a major antioxidant in plasma [3]. The in vitro oxidation of amino acid residues leads to protein degradation, aggregation, and cross-linking. In contrast, significant evidence for the presence of ROS-mediated protein damage in vivo and its possible clinical significance is not currently available. Witko-Sarsat et al. [4] reported the presence of elevated levels of oxidized protein products, termed advanced oxidation protein products (AOPP) such as oxidized albumin, in the plasma of dialysis patients. It has been well documented that human serum albumin (HSA) is quite vulnerable to reactive oxygen species (ROS) [5]. Therefore, HSA is continuously exposed to oxidative stress, so that alterations of the conformation and function of HSA could

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