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# Functional characterization of *Arabidopsis thaliana* transthyretin-like protein

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## Abstract

**Background:** *Arabidopsis thaliana* transthyretin-like (TTL) protein is a potential substrate in the brassinosteroid signalling cascade, having a role that moderates plant growth. Moreover, sequence homology revealed two sequence domains similar to 2-oxo-4-hydroxy-4-carboxy-5-ureidoimidazole (OHCU) decarboxylase (N-terminal domain) and 5-hydroxyisourate (5-HIU) hydrolase (C-terminal domain). TTL is a member of the transthyretin-related protein family (TRP), which comprises a number of proteins with sequence homology to transthyretin (TTR) and the characteristic C-terminal sequence motif Tyr-Arg-Gly-Ser. TRPs are single domain proteins that form tetrameric structures with 5-HIU hydrolase activity. Experimental evidence is fundamental for knowing if TTL is a tetrameric protein, formed by the association of the 5-HIU hydrolase domains and, in this case, if the structural arrangement allows for OHCU decarboxylase activity. This work reports about the biochemical and functional characterization of TTL.

**Results:** The TTL gene was cloned and the protein expressed and purified for biochemical and functional characterization. The results show that TTL is composed of four subunits, with a moderately elongated shape. We also found evidence for 5-HIU hydrolase and OHCU decarboxylase activities *in vitro*, in the full-length protein.

**Conclusions:** The *Arabidopsis thaliana* transthyretin-like (TTL) protein is a tetrameric bifunctional enzyme, since it has 5-HIU hydrolase and OHCU decarboxylase activities, which were simultaneously observed *in vitro*.

## Background

The *Arabidopsis thaliana* transthyretin-like protein (TTL) was first identified as a potential substrate of Brassinosteroid-Insensitive 1 (BRI1), the principal brassinosteroid (BR) receptor, playing a negative role in BR-mediated plant growth [1].

Sequence analysis shows that TTL displays an N-terminal domain corresponding to 2-oxo-4-hydroxy-4-carboxy-5-ureidoimidazole (OHCU) decarboxylase, and a C-terminal domain that has approximately 42% sequence identity with transthyretin (TTR), a vertebrate-specific transport protein [1,2]. TTR is a plasma protein which transports the thyroid hormones 3,5,3',5'-tetraiodo-L-thyronin (T<sub>4</sub>) and 3,5,3-triiodo-L-thyronin (T<sub>3</sub>) as well as retinol by association with the retinol-binding protein [3]. TTRs are homotetrameric proteins, each monomer containing two four-stranded  $\beta$ -sheets and

one short  $\alpha$ -helix. They have a channel that runs through the dimer-dimer interface, where two identical thyroid hormone binding sites are located [4,5]. Proteins with sequence homology to TTR and with the characteristic C-terminal sequence motif Tyr-Arg-Gly-Ser, which is absent in TTRs, are named transthyretin-related proteins (TRPs) and are found in bacteria, fungi, animals (both vertebrate and invertebrate) and plants [2,6,7]. TRPs are thought to be TTR ancestors [2,7].

Three TTL splice variants were reported; two of them are cytoplasmic (311- and 286-residues each) and the other one is peroxisomal (324-residues) [8]. In all three variants the N-terminal domain, OHCU decarboxylase, is composed of 180 amino acids, whereas the C-terminal domain varies in length, having 144, 131 and 106 amino acids for the 324-, 311- and 286-residues isoforms, respectively [2,8]. The 324-residues isoform contains a type-2 peroxisomal targeting sequence (PTS-2), which is deleted in the other two isoforms [2,8]. In most eukaryotes, 5-HIU hydrolases contain a PTS-2 and OHCU

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decarboxylases contain a type-1 peroxisomal targeting sequence (PTS-1), except in plants [9]. In *A. thaliana* these two enzymes are fused into a single polypeptide chain, containing a PTS-2 sequence, between the two domains, in the 324-residues isoform.

In a previous study, the C-terminal domain of TTL was expressed and studied separately. This single-domain was named transthyretin-like protein (TLP) [2] and is distinct from TTL, which contains both this C-terminal domain, with 5-HIU hydrolase activity, and a N-terminal domain, with OHCU decarboxylase activity, fused into the same polypeptide chain [8]. TLP was described as a tetramer [2]. The three-dimensional structure of TRPs from *Salmonella Dublin*, *Escherichia coli*, zebrafish and the homologous domain from *Bacillus subtilis* were determined [10-13]. The overall structures of these TRPs and also the model predicted for TLP [2] are very similar to those of TTRs, as expected due to sequence homology, although functionally they are different proteins. TRPs are hydrolases with a role in uric acid degradation [7,14] and they do not bind thyroid hormones [2,6]. TTL is the *A. thaliana* TRP member, which contains an extra domain of 180 amino acids as compared to other TRPs. Until now, only TRPs from *Magnetospirillum magnetotacticum*, *Bradyrhizobium japonicum*, *Bacillus subtilis* and plant species have been described as composed of two sequence domains, one related to TTR and the other

with features common to OHCU decarboxylases [2,7,9].

TTL was predicted by sequence homology as being a bifunctional enzyme with 5-hydroxyisourate (5-HIU) hydrolase and OHCU decarboxylase activities, catalysing the two final steps in the uric acid degradation pathway (Fig. 1) [8]. In most organisms, uric acid, the end product of purine degradation, is catabolised to allantoin. The process is initiated by uricase, which oxidizes uric acid into 5-HIU; this intermediate compound is then hydrolysed by 5-HIU hydrolase, leading to OHCU. Finally, a third enzyme, OHCU decarboxylase, catalyses the decarboxylation of OHCU producing (S)-allantoin [9]. The spontaneous degradation of uric acid follows the same pathway, resulting into 5-HIU and OHCU intermediates, which can be further oxidized, producing potentially harmful reactive species. However, many organisms have an enzymatic pathway, possibly to convert these intermediates more rapidly into allantoin, a much less reactive species. The spontaneous pathway produces racemic allantoin, whereas the enzymatic pathway produces exclusively (S)-allantoin [9].

So far, two different functions have been assigned to TTL: i) a signalling role to the cytoplasmic isoforms and ii) an enzymatic role to the peroxisomal isoform. While the signalling role has already been studied [1], it is not known if the two enzymatic sequence domains that are in this protein fused into a long polypeptide chain do

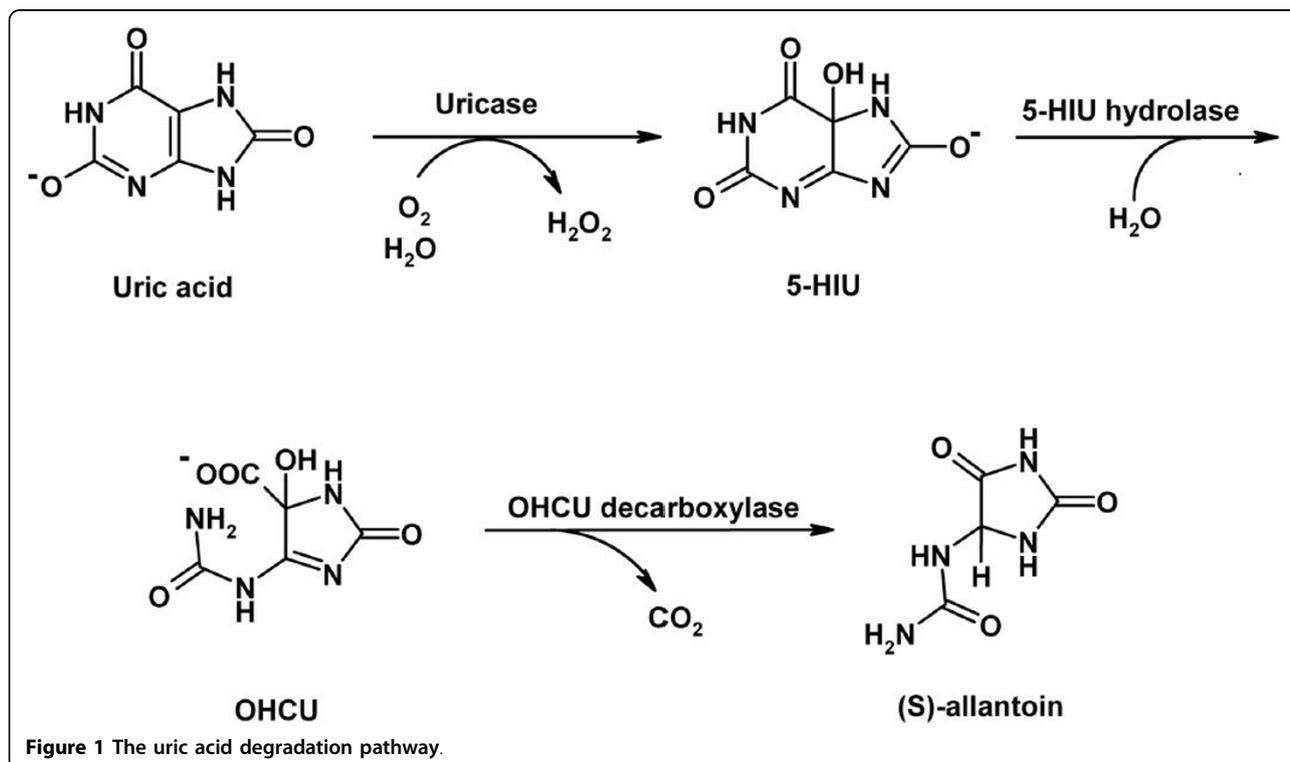


Figure 1 The uric acid degradation pathway.

not compromise the role of each domain in an isolated form. In this work we studied the 311-residue cytoplasmic isoform, referred as TTL throughout the manuscript, to address the protein enzymatic role.

## Results

### TTL oligomerization state

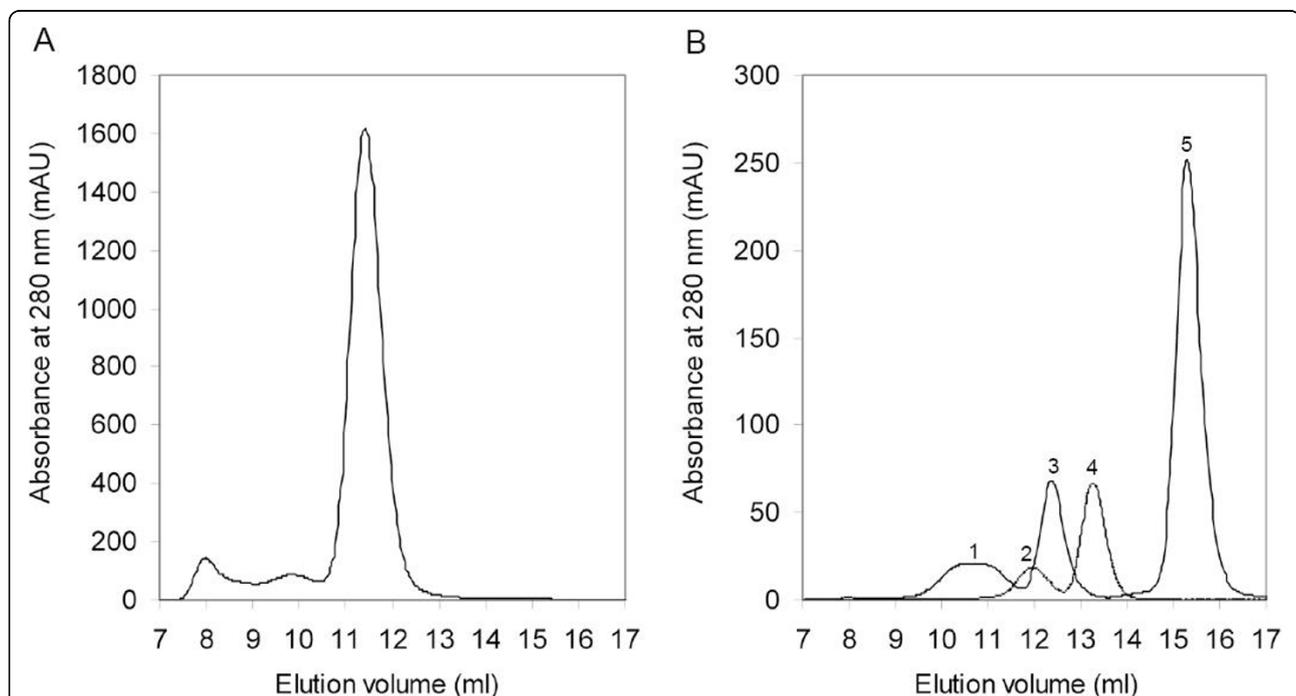
Recombinant TTL was purified and used to investigate the protein oligomerization state. We decided to use the method developed by Siegel and Monty in 1966 [15], because it allows calculation of the molecular mass of a protein independently of its shape, using a combination of the Stokes radius ( $a$ ) derived from gel filtration chromatography and the sedimentation coefficient ( $S$ ) obtained from density gradient centrifugation. The gel filtration chromatograms are presented in Fig. 2, showing in Fig. 2A a major peak of oligomeric TTL and in Fig. 2B superposed chromatograms of standard Stokes radius markers. Using the equations described in Methods, section on "Determination of the protein oligomeric state",  $a$  and  $S$  were estimated to be 5.3 nm and 6.3 S, respectively, and with these values the native molecular mass for TTL was estimated to be 137.3 kDa. Since each TTL subunit has 36.6 kDa, we concluded that TTL is a tetramer, which is in agreement with the fact that TTR, as well as *A.thaliana* TLP, also have a tetrameric

structure [2]. Moreover, the calculated frictional ratio, 1.5, is consistent with a moderately elongated non-globular protein [15].

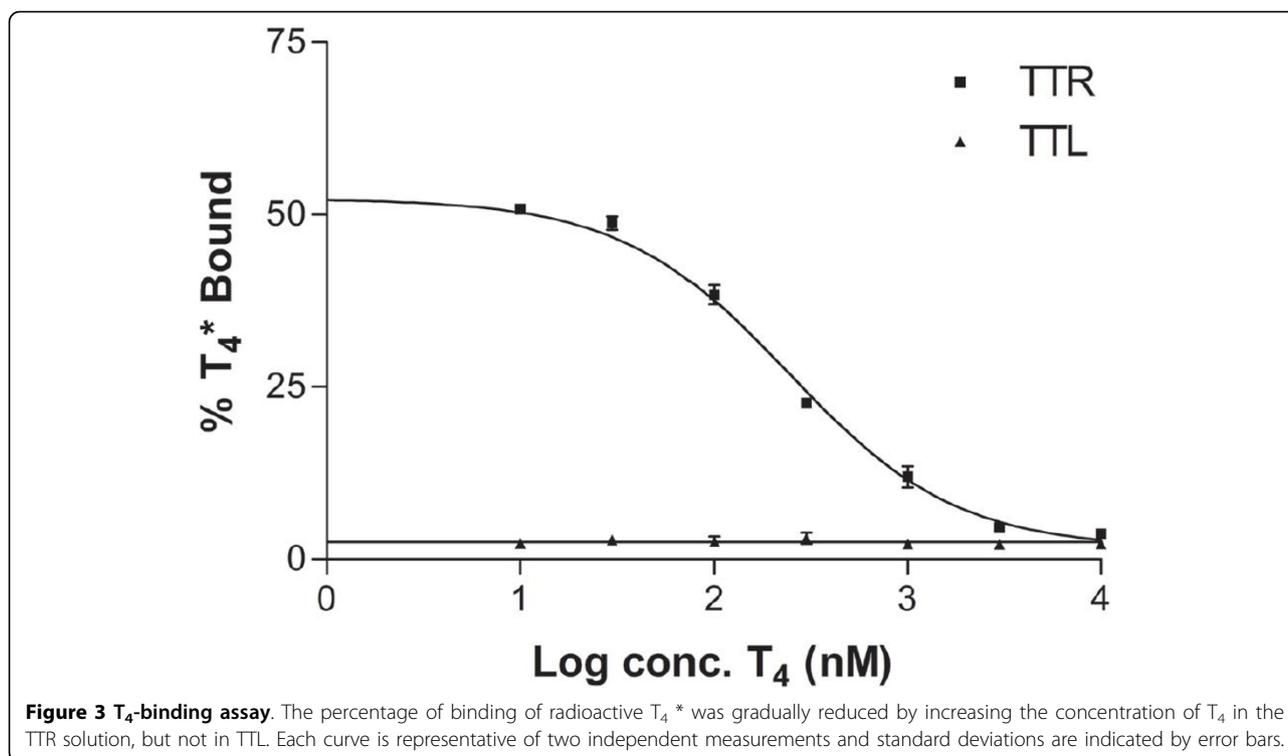
### Thyroxin ( $T_4$ )-binding assays

Since TTR is a tetramer with two equivalent  $T_4$ -binding sites, we decided to study the binding of  $T_4$  to TTL, using radiolabeled thyroxin and TTR as the sample control (Fig. 3). The TTR control has the typical behaviour of a specific  $T_4$ -binding protein, in which the increase of cold  $T_4$  concentration causes the displacement of specifically bound radiolabeled  $T_4^*$ . On the contrary,  $T_4^*$ -binding to TTL, besides being very low, was constant at different  $T_4$  concentrations, showing a non-specific binding. This is a very different behaviour as compared to TTR, for which non-specific binding was only detected at the highest  $T_4$  concentration. We concluded that the binding of  $T_4$  to TTL is not significant and clearly non-specific, since it is not altered in the presence of a competitor.

Sequence alignment of human TTR, TTL and TRPs from *Salmonella dublin*, *B. subtilis* and mouse are presented in Fig. 4. As mentioned previously, only the C-terminal domain of TTL has homology with TTR or TRPs. Several amino acids were reported as important for the binding of  $T_4$  to TTR, namely Lys15, Glu54,



**Figure 2 Analytical gel filtration chromatograms for determination of TTL Stokes radius.** (A) shows a chromatogram of recombinant TTL (elution volume: 11.4 ml). In (B), two chromatograms of standard Stokes radius markers are superposed, where 1 refers to ferritin (Stokes radius: 6.1 nm; elution volume: 10.7 ml), 2 to aldolase (4.81 nm; 12.0 ml), 3 to albumin (3.55 nm; 12.4 ml), 4 to ovalbumin (3.05 nm; 13.3 ml) and 5 to chymotrypsinogen A (2.09 nm; 15.3 ml).



Thr106 and Thr119 [5]. Interestingly, they are not conserved in TTL, being replaced by His196, Arg245, His295 and Tyr308, respectively (Fig. 4), which are amino acids conserved in TRPs, and were reported to be important for 5-HIU hydrolase catalytic activity [10,13,16]. According to our sequence alignment data (Fig. 4) and the three-dimensional structure of several TTR and TRP proteins [5,10-13], these substitutions will probably have two main effects: i) create a narrower region at the inner part of the binding site due to the replacement of Thr for Tyr, which has a larger side chain and ii) alter the surface topography and introduce charge differences at the entrance of the channel, due to substitutions Lys-His, Glu-Arg and Thr-His. These modifications will probably exclude the binding of T<sub>4</sub> to TRPs and TTL.

#### TTL enzymatic activities

Since TRPs are functionally associated to 5-HIU hydrolases, we decided to study the predicted 5-HIU hydrolase and OHCU decarboxylase activities for TTL *in vitro*. The experiments were started by testing how TTL would affect the activity of uricase over uric acid by monitoring the differences in absorbance at 292 nm in the absence or presence of TTL (Fig. 5). Although uricase alone was able to decompose uric acid, its activity was significantly accelerated in the presence of TTL. By contrast, TTR did not influence this reaction. This result is consistent with the predicted 5-HIU hydrolase and

OHCU decarboxylase activities. The rapid consumption of 5-HIU and OHCU should favour an equilibrium shift in the reaction catalysed by uricase.

5-HIU hydrolase activity was demonstrated *in vitro* for single-domain TRPs present in *S. dublin*, *B. subtilis* and mouse [10,14,16]. The authors reported that uric acid was converted into 5-HIU by uricase and both the formation and subsequent degradation of this compound was monitored at 312 nm, since OHCU does not absorb at this wavelength [17]. In our assay conditions, within 60 seconds incubation of uric acid with the *Candida sp.* uricase, the production of 5-HIU reached the maximum level (Fig. 6). At this point, TTL was added to the reaction solution resulting in a rapid decrease in absorbance at 312 nm (Fig. 6). As expected, addition of TTR, which was used as the negative control, failed to accelerate 5-HIU degradation. This result shows that TTL is an enzyme that facilitates the hydrolysis of 5-HIU and also indicates that most probably four protein C-terminal domains associate forming a tetrameric structure, since the 5-HIU hydrolase catalytic site are located at the dimer-dimer interface in all structurally and functionally characterised TRP members [10,13].

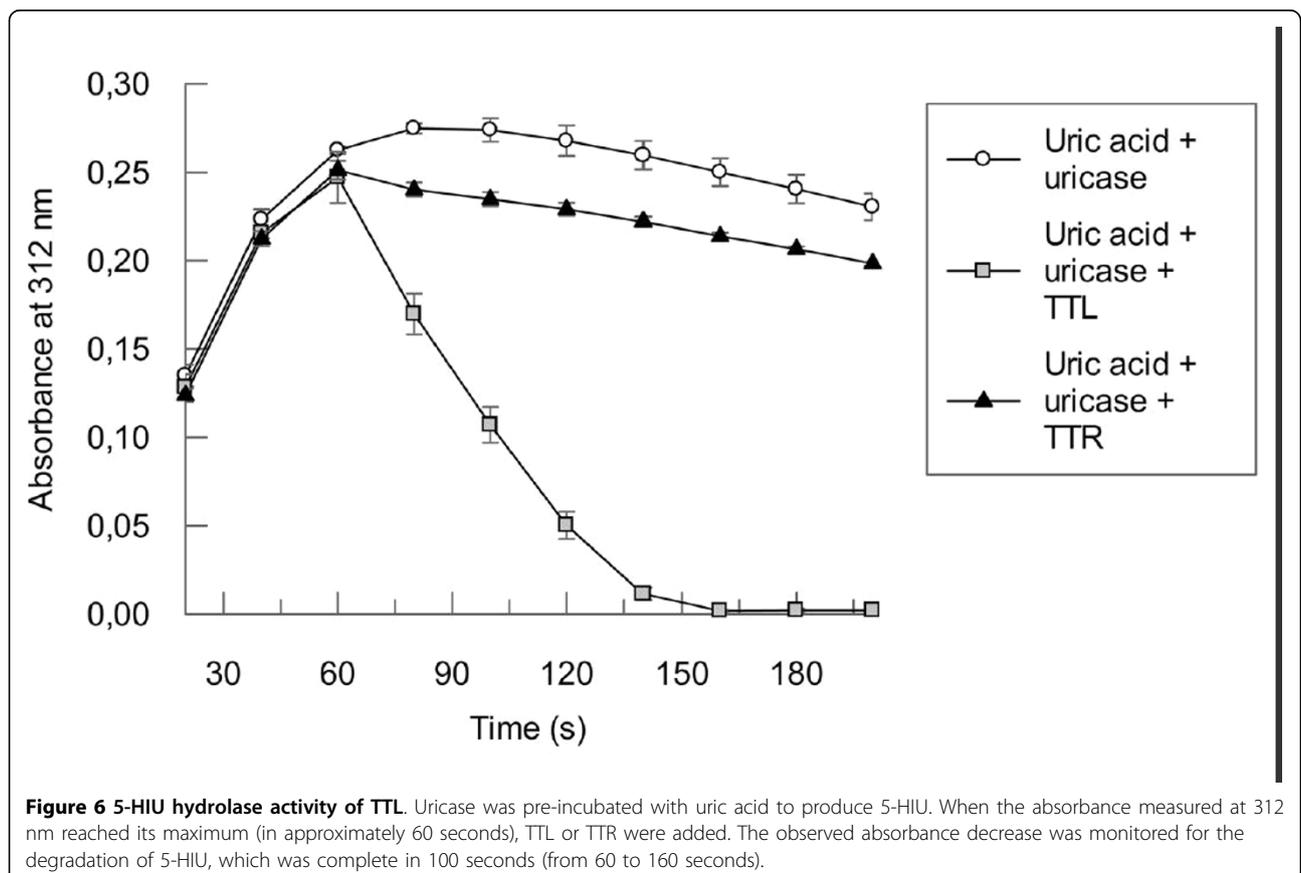
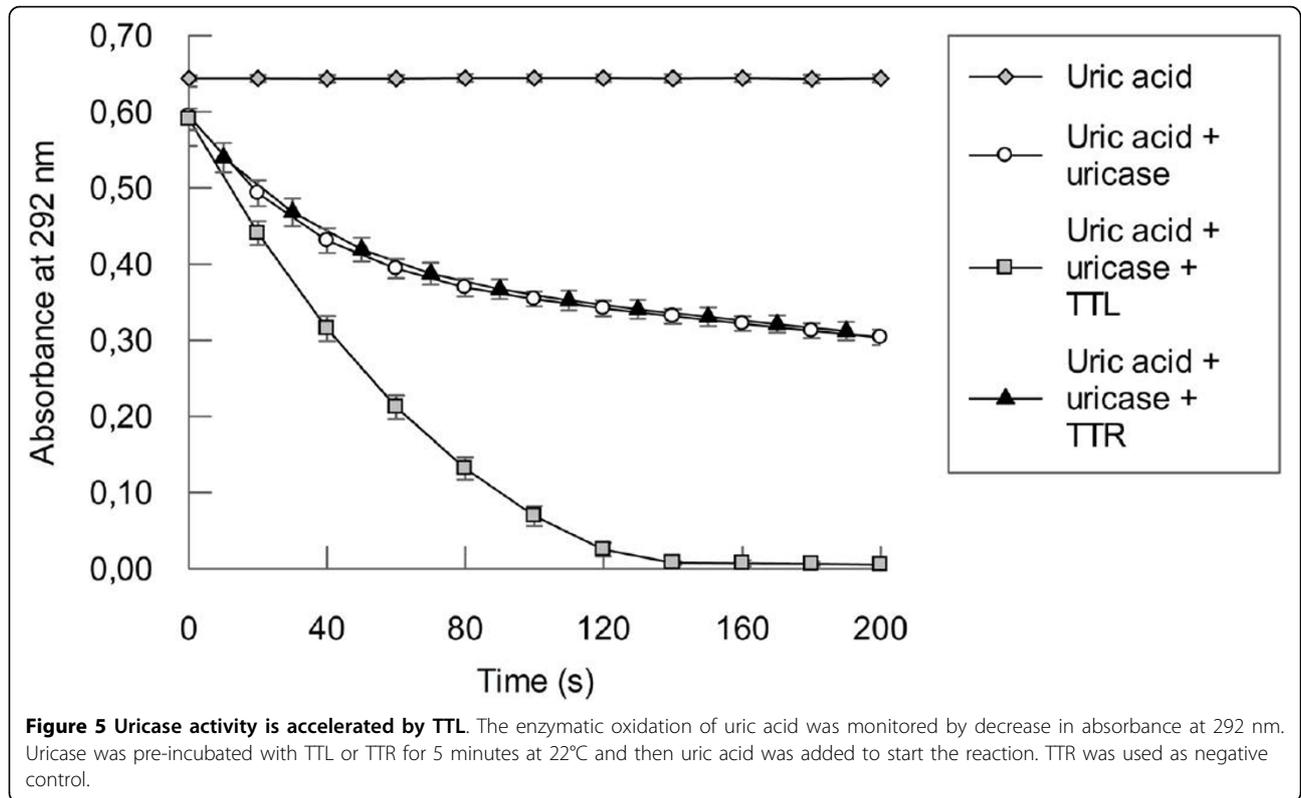
Although the crystal structure of the N-terminal OHCU domain of TTL from *A. thaliana*, which contains conserved residues critical for the OHCU decarboxylase activity, was recently reported [18], its predicted enzyme activity was not experimentally demonstrated. In fact, it was only observed *in vitro* in

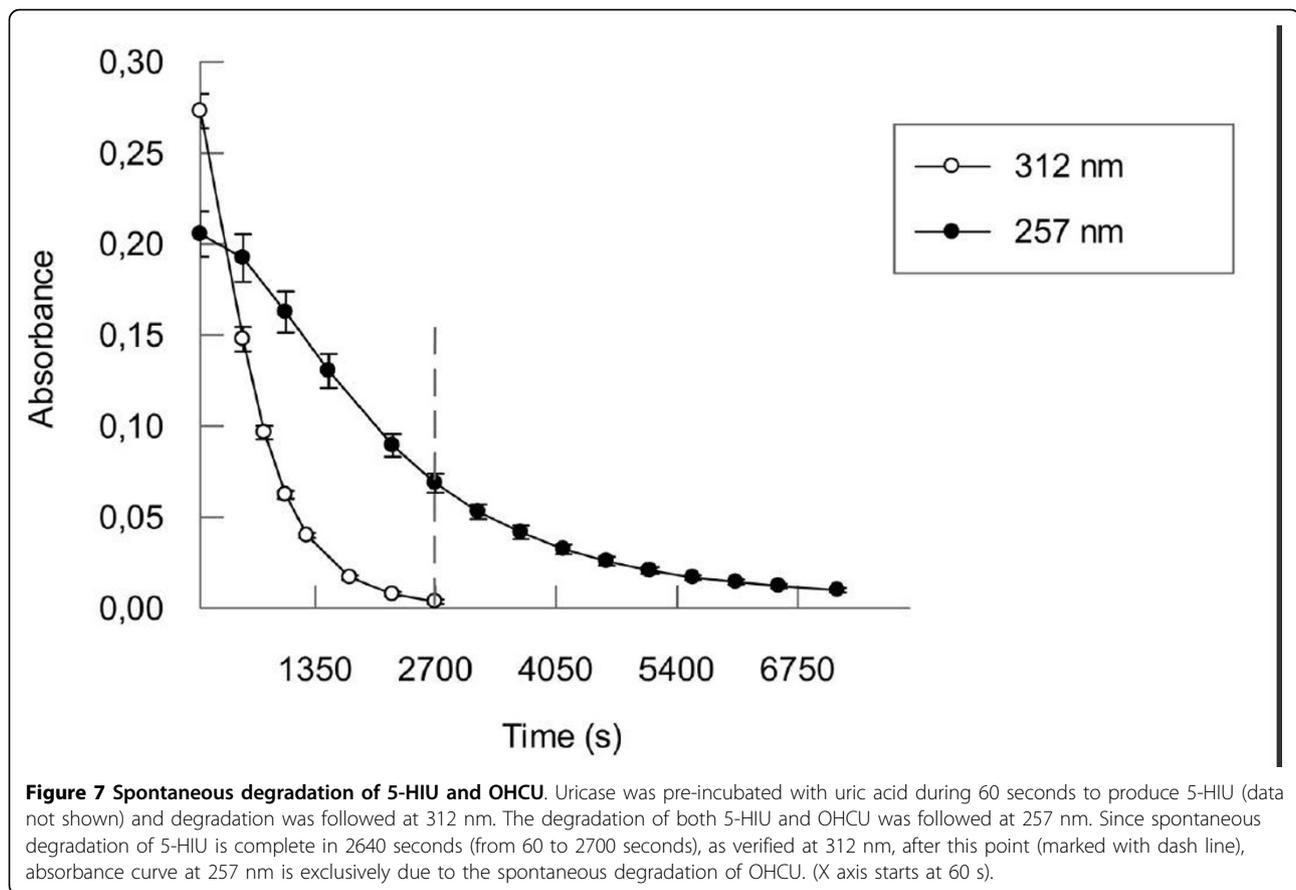
TTR	-----	
TTL	MAMEIGEDEWKVCCGSSEFAKQMS TSGPLTSQEAIYTARDI WFNQVNVTDWLEAFSAHPQ	60
mTRP	-----	
SdTRP	-----	
BsTRP	-----	
TTR	-----	
TTL	IGNTPSPSINSDFARRSVSEQSTAFATT SASALQELAEWNVLYKKKFGFI F I ICASGRTH	120
mTRP	-----	
SdTRP	-----	
BsTRP	-----	
TTR	-----GPTGTGES-----	8
TTL	AEMLHALKERYENRP IVELEIAAMEQMKITELRMAKLFSDKAKVISETDSSSPVSTKPQ	180
mTRP	-----MATES S-----	6
SdTRP	-----N-----	1
BsTRP	-----MSEPESLMG-----	9
TTR	-----KCPLMVKVLDAVRGSPA INVAVHVFRKAAD-----DTWEPFASGKT	49
TTL	AEAKAPKRSRPPI TTHVLDVSRGAPAAGVEVHLEVWSGTTGPSFVHGGGGVWSSVGTSAT	240
mTRP	-----PLTTEHVLDTASGLPAQGLCLRSLRLEAPCQQ-----WMELRTSYT	46
SdTRP	-----ILSVHI LDQQTGKPA PGVEVVLEQKKD-----NGWTQLNTGHT	39
BsTRP	-----KLTTTHI LDLTGKPAANVKIGLKR-----LGESIMKEVYT	44
TTR	SESGELHG-LTTEEEFVEGIYKVEIDTKSYWKALG----ISPFHEHAEVVFTANDSGP-R	103
TTL	DRDGRSGPLMDLVDALNPGTYRISFDTAKYSPG-----CFFPYVSIVFQVTESQKWE	292
mTRP	NLDGRC PGLLTPS-QIKPGTYKLF DTERYWKER----GQESFYYPYVEVFTITKET--Q	99
SdTRP	DQDGR IKA-LWPEKAAAPGDYRVI FKTCQYFESKKLDT----FFPEI PVEFHISKTN--E	92
BsTRP	NNDGRVDVPLLAGEELMSG EYVMEFHAGDYFASKNMNAADQPFLTIVTVRFQLADPD--A	102
TTR	RYTHAALLSPYSYSTTAVVTNPKE	127
TTL	HFEVPLLLAPFSFSTYRGS-----	311
mTRP	KFEVPLLLSPWSYTTYRGS-----	118
SdTRP	HYEVPLLLSQYGYSTYRGS-----	111
BsTRP	HYHIPLLLSPFGYQVYRGS-----	121

**Figure 4** Sequence alignment of human TTR, TTL and TRPs from mouse (mTRP), *S. dublin* (SdTRP) and *B. subtilis* (BsTRP). Amino acids involved in the binding of T<sub>4</sub> to TTR and their substitutes in TTL and TRPs are presented in black. Sequence alignment was obtained using ClustalW [24].

an isolated domain of a bifunctional enzyme from *B. subtilis* and also in zebrafish OHC decarboxylase [18,19]. Monitoring the decarboxylase activity of TTL by spectrophotometry is not straightforward since OHCU absorbs below 300 nm and its absorption spectrum overlaps with that of 5-HIU [17]. Therefore, measurements at 257 nm are likely to be a sum of contributions by 5-HIU and OHCU. However, the spontaneous degradation observed at this wavelength is clearly distinct

from what is observed at 312 nm (Fig. 7). The spontaneous degradation followed at 312 nm shows that 5-HIU is completely degraded in 2640 seconds (from 60 s to 2700 s Fig. 7), thus the spontaneous absorbance decay measured after this point at 257 nm (marked with dash line in Fig. 7) is exclusively due to the spontaneous degradation of OHCU. Therefore this degradation was used as control in the experiments testing the effect of TTL on degradation of OHCU.





In presence of TTL the degradation of 5-HIU followed at 312 nm was completed in 100 seconds (from 60 s to 160 s in Fig. 6) therefore after this point the absorbance decay measured at 257 nm (marked with a dash line in Fig. 8) corresponds to the degradation of OHCU affected by TTL. The rate of this degradation is excessively high as compared to the control spontaneous degradation of OHCU (Fig. 8). Addition of TTL accelerated significantly the degradation of OHCU, supporting the decarboxylase activity of the protein.

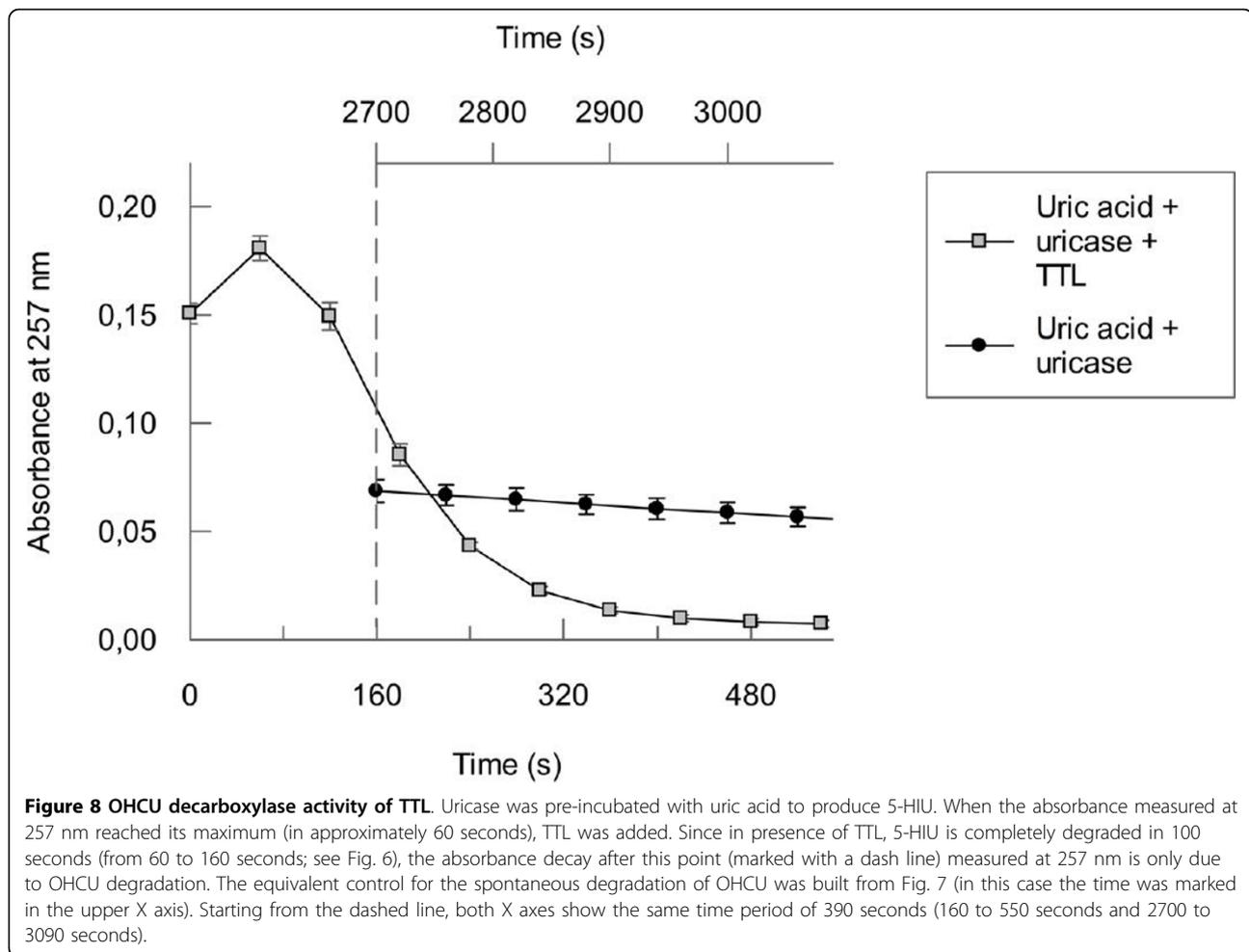
Taken together, our results demonstrated that TTL has 5-HIU hydrolase and OHCU decarboxylase activities *in vitro*, and they are both performed at the same time.

## Discussion

It seems that the TTR domain evolved from an enzyme present in bacteria, fungi, plants and animals to a hormone transporter present in vertebrates [2,7,12]. Its function changed completely, although there are only minor differences in sequence. In the present study, we showed that TTL from *A. thaliana* is a tetramer with a moderately elongated shape. The tetrameric structure is in agreement with the observation that the protein has 5-HIU hydrolase activity. We expect that four C-terminal domains associate in a tetramer forming two active

sites, as reported for TRPs, which contain only one domain. Moreover, the elongated shape is possibly due to the N-terminal OHCU decarboxylase domain associated to each of the four C-terminal sequences forming the tetramer. The molecular model of this domain was determined in the presence of allantoin [18]; it is an alpha-domain that associates as dimers in its crystal form. However, a careful inspection of the available molecular models for TRPs and OHCU decarboxylase domain indicates that allantoin binding site is not in the region of the interaction between the two subunits that form the dimer [18] and this kind of association most probably will not be present in TTL, since the TTL tetramer will be formed through contacts among the C-terminal domains.

We also demonstrated that TTL does not bind  $T_4$  *in vitro*, corroborating the observation that the ligand-binding site has characteristics which are different from those present in TTR. These results are in agreement with previous observations, since the functionally characterized TRPs do not bind thyroid hormones [2,6] and are 5-HIU hydrolases with active sites located in the regions corresponding to the  $T_4$ -binding sites in TTR [7,10,12,13]. The OHCU decarboxylase activity was also detected in the full-length TTL, indicating that the



N-terminal domains do not inhibit the active sites formed by the C-terminal regions; the reverse is also valid, thus supporting the idea of a bifunctional enzyme.

TTL was initially characterized as a specific BRI1-interactor that regulates BR-mediated plant growth [1]. Here we show that the same protein is also enzymatically active. Several proteins with at least two totally different functions have been reported and named moonlighting proteins; the switch between functions is due to changes in cellular localization, cell type, oligomeric state or cellular concentration of a ligand [20,21]. We think that TTL is a moonlighting protein that switches its function according to its subcellular location. In eukaryotes, peroxisomes are the major organelles where oxidative reactions with molecular oxygen consumption are carried out. In these organisms, uricase is located in peroxisomes, oxidizing uric acid to 5-HIU [9]. As a consequence, 5-HIU accumulates in peroxisomes, providing the initial TTL substrate. Therefore, in peroxisomes, where uric acid degradation occurs, TTL, through its 324-residues isoform, functions as a bifunctional enzyme

with 5-HIU hydrolase and OHCU decarboxylase activities. In the cytoplasm, 5-HIU and OHCU are absent, since they are produced in peroxisomes. Consequently, the TTL 311-residues isoform most probably will not have these enzymatic functions. Moreover, it was shown to interact with BRI1, mediating BR-regulated plant growth. BRI1 is an intrinsic membrane protein, located in the plasma membrane and containing a cytoplasmic kinase domain that interacts with TTL [1]. The main difference between the 324- and 311-residues isoforms is the presence or absence of PTS-2. The subcellular location of three TTL isoforms is probably fundamental to define their role *in vivo*. TTL 324-residues isoform location into peroxisomes was already shown by fluorescence microscopy, and provided conclusive results regarding its import [8]. It was also observed that its import efficiency was low and it was hypothesized that could result from a low accessibility of the internal targeting signal to the peroxisomal import machinery. Internal PTS-2 signal sequences are thought to be rare in nature and to occur preferentially in bifunctional enzymes resulting from gene

fusion [8]. The 311-residues isoform will probably remain in the cytoplasm *in vivo*, since it does not contain the PTS-2 sequence [1,8]. Therefore in the absence of uric acid degradation intermediates, it interacts with the plasma membrane-located BRI1, mediating BR-regulated plant growth [1].

Attempts for the three-dimensional structure determination of TTL are in progress and should represent a fundamental step in order to clarify TTL function, namely its signalling role in BR-mediated responses.

## Conclusions

The amino acid sequence of TTL from *A. thaliana* reveals two domains similar to 5-HIU hydrolase and OHCU decarboxylase. As a consequence, TTL was predicted to be a bifunctional enzyme. Here we reported the first experimental data showing that in fact the protein has both enzymatic activities *in vitro*. Moreover, we confirmed that TTL is in fact a bifunctional enzyme, by showing that it performs both activities simultaneously.

## Methods

### Cloning, expression and purification of *A. thaliana* TTL

TTL ORF was amplified by PCR and cloned into pET-28a(+) (Novagen). Recombinant TTL was overexpressed in *E. coli* BL21 (DE3) codon plus and purified as follows: cells were grown in LB at 37°C to OD<sub>600</sub> = 0.8 and expression induced with 1 mM IPTG for 18 hours at 20°C. Cells were harvested (10500 g, 20 minutes), resuspended in 140 mM NaCl, 2.7 mM KCl, 1.0 mM KH<sub>2</sub>PO<sub>4</sub>, 50 µg/ml lysozyme, 1 mM phenylmethylsulfonyl fluoride (PMSF), 1 µg/ml DNase I, 100 µM MgCl<sub>2</sub>, 10 mM Na<sub>2</sub>HPO<sub>4</sub>, pH 7.4 and lysed by sonication. Lysates were centrifuged (27200 g, 20 minutes), supernatants filtered and loaded onto a 5 ml HisTrap HP column (GE Healthcare) previously equilibrated with buffer A (20 mM imidazole, 500 mM NaCl, 20 mM NaH<sub>2</sub>PO<sub>4</sub>/Na<sub>2</sub>HPO<sub>4</sub>, pH 7.4). After a washing step with 20% buffer B (500 mM imidazole, 500 mM NaCl, 20 mM NaH<sub>2</sub>PO<sub>4</sub>/Na<sub>2</sub>HPO<sub>4</sub>, pH 7.4), TTL was eluted with 50% buffer B. Fractions containing His-tagged TTL were dialyzed against buffer A followed by buffer C (150 mM NaCl, 1 M EDTA, 20 mM Tris-HCl, pH 8.0), concentrated and loaded onto a HiPrep 26/60 Sephacryl S-300 HR column (GE Healthcare). After an isocratic elution in buffer C, TTL containing fractions were pooled and concentrated to 7 mg/ml. Protein concentration was determined by absorption at 280 nm using a theoretical 49430 M<sup>-1</sup>cm<sup>-1</sup> molar extinction coefficient [22].

### Determination of the protein oligomeric state

For analytical size-exclusion chromatography a Superose 12 10/300 column (GE Healthcare) was equilibrated with buffer C at 0.5 ml/min and calibrated with protein

standards of known Stokes radius. The Stokes radius (*a*) for the experimental data was calculated using:  $(-\log K_{av})^{1/2} = f(a)$ . For isokinetic rate zonal ultracentrifugation, two continuous gradients of 5 to 20% sucrose (Merck) were prepared in buffer C, calibrated with standard proteins of known sedimentation coefficient and ran for 25 hours and 15 minutes at 260800 g in a Beckman SW41 Ti rotor. TTL native molecular mass (*M*) and frictional ratio (*f/f*<sub>0</sub>) were calculated according to Siegel and Monty [15], using the following equations:

$$M = \frac{6\pi\eta N a s}{1 - v\rho}$$
$$f / f_0 = \frac{a}{((3 v M)/(4\pi N))^{1/3}}$$

where *M* represents the molecular mass, *a* the Stokes radius, *s* the sedimentation coefficient, *v* the partial specific volume, *f/f*<sub>0</sub> the frictional ratio, *η* the viscosity of the medium, *ρ* the density of the medium and *N* Avogadro's constant. Partial specific volume for TTL was calculated from the amino acid sequence of the protein using the program SEDNTERP v1.08 <http://www.jphilo.mailway.com/default.htm>.

### Thyroxin-binding assays

The assays were performed as previously described [23], using 100 nM TTR or TTL, 5-10 × 10<sup>4</sup> counts per minute (cpm) radiolabeled thyroxin, <sup>125</sup>I-T<sub>4</sub> (T<sub>4</sub>\*) (Perkin Elmer) and different cold T<sub>4</sub> (Sigma) concentrations. Counts were measured in a Wallac 1470 wizard™ automatic gamma counter.

### Enzyme assays

The enzymatic assays were based on previously described procedures for enzymes with sequence similarity to the TTL hydrolase domain [10,14]. Briefly, 500 µl assay mixtures contained 0.05 units/ml uricase from *Candida sp.* (Sigma) in 50 mM potassium phosphate buffer pH 7.8. When 50 µM uric acid (Sigma) were added, 5-HIU and OHCU were generated *in situ* and their degradation was followed at 312 nm and 257 nm, in the presence and absence of 0.003 µM TTL or TTR. TTL effect on uricase activity was monitored at 292 nm in the same conditions. Triplicate measurements were done aerobically at 22°C.

### Abbreviations

5-HIU: 5-hydroxyisourate; BR: Brassinosteroid; BRI1: Brassinosteroid-Insensitive 1; OHCU: 2-oxo-4-hydroxy-4-carboxy-5-ureidoimidazole; PTS-2: type-2 peroxisomal targeting sequence; T<sub>4</sub>: 3,5,3',5'-tetraiodo-L-thyronin, thyroxin; T<sub>3</sub>: 3,5,3-triiodo-L-thyronin; T<sub>4</sub>\*: radiolabeled thyroxin; TLP: transthyretin-like protein; TTL: transthyretin-like protein; TRP: transthyretin-related protein; TTR: transthyretin.

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#### Authors' contributions

JP carried out the biochemical and functional analysis work under ZS and FFS's supervision; SM collaborated on the protein expression work; MRA supervised the T<sub>4</sub> binding studies; JL supplied the TTL-coding vector and discussed the results; AMD coordinated the study. All authors gave ideas, revised, read and approved the final manuscript.

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