The identification of an integral membrane, cytochrome c urate oxidase completes the catalytic repertoire of a therapeutic enzyme

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In living organisms, the conversion of urate into allantoin requires three consecutive enzymes. The pathway was lost in hominid, predisposing humans to hyperuricemia and gout. Among other species, the genomic distribution of the two last enzymes of the pathway is wider than that of urate oxidase (Uox), suggesting the presence of unknown genes encoding Uox. Here we combine gene network analysis with association rule learning to identify the missing urate oxidase. In contrast with the known soluble Uox, the identified gene (puuD) encodes a membrane protein with a C-terminal cytochrome c. The 8-helix transmembrane domain corresponds to DUF989, a family without similarity to known proteins. Gene deletion in a PuuD-encoding organism (Agrobacterium fabrum) abolished urate degradation capacity; the phenotype was fully restored by complementation with a cytosolic Uox from zebrafish. Consistent with H2O2 production by zfUox, urate oxidation in the complemented strain caused a four-fold increase of catalase. No increase was observed in the wild-type, suggesting that urate oxidation by PuuD proceeds through cytochrome c-mediated electron transfer. These findings identify a missing link in purine catabolism, assign a biochemical activity to a domain of unknown function (DUF989), and complete the catalytic repertoire of an enzyme useful for human therapy.

As a drug used to treat hyperuricemia associated with gout, tumor lysis syndrome, and the Lesch-Nyhan disease, urate oxidase (Uox, also called uricase) is an enzyme of considerable biomedical interest1–3. The intravenous administration of urate oxidase is a particular example of enzyme-replacement therapy, in which the enzyme is not just missing in the individual but in the entire species, resulting from progressive mutations of the uox coding sequence in hominid ancestors4. In consequence of this evolutionary inactivation, urate is the end product of purine catabolism in human and apes, whereas in other mammals the end product is the more soluble allantoin.

In Bacteria, Archaea, and Eukaryotes, where present, the oxidative conversion of urate into allantoin proceeds through a three-step enzymatic pathway5. In the first step, uric acid (urate at neutral pH) is converted to 5-hydroxyisourate (HIU) in an oxygen-dependent reaction6. In the second step, HIU is hydrolysed to 2-oxo-4-hydroxy-4-carboxy-5-ureidoimidazoline (OHCU), which is decarboxylated to give dextrorotatory allantoin in the third step (Fig. 1a). As HIU and OHCU are metastable compounds with a half-life of 7.2 and 9.6 minutes at physiological conditions, racemic allantoin is obtained in vitro as a final product of the Uox reaction. However, in nature the presence of Uox is almost invariably associated with both HIU hydrolase (Urah) and OHCU decarboxylase (Urad). Besides affecting the reaction...
stereochemistry\textsuperscript{7}, the presence of these enzymes appears to be important for the rapid elimination of the metastable intermediates of urate oxidation\textsuperscript{8}.

The functional coupling between Uox, Urnah, and Urad is reflected by the evolutionary link of the corresponding genes, which are usually present or absent together in a given genome. This link has been key to the identification of urah and urad as genes associated with uox\textsuperscript{5}; in prokaryotes, however, cases were also noticed in which both urah and urad were present in the absence of uox genes. This observation was followed in recent years by the identification of alternative urate oxidases—non homologous enzymes involved in the same reaction step. The uox gene encodes the first identified urate oxidase\textsuperscript{9}, a

Figure 1. Identification of COG3748 as urate oxidase. (a) Pathway for the conversion of urate into allantoin. (b) String association network obtained with COG3648, COG2351, and COG3195. Nodes represent gene families according to the COG classification. Edges represent predicted functional associations; stronger associations are shown as thicker lines. The node identified as a candidate urate oxidase (COG3748) is indicated with an arrow. (c) Map of urate oxidation capacity in complete genomes. The tree represents 431 distinct species possessing either the uox, hpxO, hpyO, or COG3748 genes or both the urah and urad genes. The presence (red) or the absence (blue) of genes in complete genomes is shown alongside the organism tree. Main taxonomic groups and organisms discussed in the text are indicated.
cofactor-less enzyme with a T-fold domain\(^\text{10}\) that is found in all tree domains of life. The existence of an alternative urate oxidase gene, named \(hpxO\) (hypoxanthine-xanthine utilization O), has been demonstrated in \(Klebsiella\) spp. possessing \(urah\) and \(urad\) but not \(uox\)\(^\text{1,12}\). The \(HpxO\) enzyme has been biochemically and structurally characterized as a flavoprotein belonging to a large family of hydroxylases\(^\text{13,14}\). An isofunctional non homologous urate oxidase has been identified in \(Xanthomonas\) spp. and named \(HpyO\)\(^\text{15}\); although \(HpyO\) belongs to a distinct family of hydroxylases, similarly to \(HpxO\), it uses \(NAD(P)H\) as a cosubstrate and \(FAD\) as a cofactor\(^\text{15}\).

From a mechanistic standpoint, \(Uox\) and \(HpxO/HpyO\) represent distinct solutions to the same biochemical problem. Each of these solutions has advantages and disadvantages. \(Uox\) does not depend on cofactors for activity, but its two-electron reduction of dioxygen generates \(H_2O_2\). This potent oxidant is readily detoxified by peroxisomes, explaining the organellar localization of the pathway in eukaryotes\(^\text{16–18}\). However, the formation of hydrogen peroxide by the \(Uox\) reaction can pose problems with organisms lacking peroxisomes, and is a matter of concern for the therapeutic administration of the enzyme\(^\text{19}\). On the other hand, the four-electron dioxygen reduction catalysed by \(HpxO/HpyO\) generates \(H_2O\), but requires a labile cofactor and a cosubstrate that is consumed in the reaction (Fig. 1a).

Surprisingly, the enzymatic variety above described does not fully account for urate oxidation in living organisms. It has been pointed out that homologs of \(uox, hpxO,\) or \(HpyO\) genes are not identified in completely sequenced organisms known to perform urate oxidation\(^\text{20}\), and our preliminary search identified numerous species (mostly proteobacteria) possessing \(urah\) and \(urad\) but not recognizable uricase genes. As \(Urah\) and \(Urad\) act on metastable products of urate oxidation, the presence of the corresponding genes could be justified only by the presence of a gene capable of urate oxidation. However, no candidate oxidoreductases can be identified by homology among the genes associated with purine catabolism. To identify this missing gene and possibly a different urate oxidase mechanism, we used here gene network analysis and a data mining technique able to discover association rules between items of an item set\(^\text{20,21}\). The application of this technique identified with high confidence genes belonging to \(COG3748\) as the missing urate oxidase. In striking contrast with the known genes involved in urate oxidation, the candidate gene (\(puad\)) was found to encode an integral membrane domain (\(DUF989\)) fused with a cytochrome \(c\). Experiments with deleted and complemented bacterial strains provided evidence for the functional assignment and revealed a novel mechanistic solution for urate oxidation.

**Results**

**Identification of \(COG3748\) as urate oxidase.** To find candidate genes encoding enzymes with urate oxidase activity we interrogated a database of predicted gene/protein associations (String)\(^\text{22}\) using as input Cluster of Orthologous Groups (COGs)\(^\text{23}\) corresponding to the \(Uox, Urah,\) and \(Urad\) proteins involved in the enzymatic pathway (Fig. 1a,b). Among the numerous associated COGs reported, the \(COG3748\) ‘predicted membrane protein’ ranked in fifth position in the String list (Supplementary Fig. S1) was considered a possible candidate as a domain with unknown function exhibiting the expected relation for a protein with urate oxidase activity, that is a connection with \(Urah\) and \(Urad\) and not with \(Uox\) in the association network (Fig. 1b). Indeed, genes involved in consecutive reactions of a pathway are expected to be correlated (e.g. they are found in the same genomes or in the same operons), while no correlation or anti-correlation is expected for different genes involved in the same reaction. Visual inspection of various operons containing \(COG3748\) genes confirmed the presence of other genes involved in purine degradation and the absence of known genes encoding urate oxidase (Supplementary Fig. S2).

To evaluate in detail the association between \(COG3748\) and the other genes of urate oxidation we examined gene distributions in 1689 complete genomes from different species. Using a semi-automated method for gene classification we found 431 species possessing both the \(urah\) and \(urad\) genes or either \(COG3748\) or a known urate oxidase gene (\(uox, hpxO, hpyO\)). This distribution clearly shows that the different genes involved in the first step of urate oxidation occur alternatively in different species (Fig. 1c). The cofactor-independent urate oxidase (\(Uox\)) occurs in eukaryotes, in some bacteria (particularly Actinomycetes), and in some archaea. Alternative oxidases (\(HpxO\) and \(HpyO\)) occur in a limited number of bacteria, while \(COG3748\) occurs in several bacterial species (mostly proteobacteria) possessing both \(urah\) and \(urad\) but not known urate oxidase genes. Among these species are organisms (e.g. \(Paracoccus denitrificans, Pseudomonas aeruginosa\)) that were previously reported to be capable of oxidative urate degradation\(^\text{24,25}\).

Bioinformatics methods make it possible to quantitatively measure correlated\(^\text{26}\) or anti-correlated\(^\text{27}\) gene distributions. Such methods were devised for pairwise associations. The generalization introduced with the analysis of logic implications\(^\text{28}\), however, allows one to evaluate comparisons involving any number of gene families. The gene distribution of Fig. 1c shows that the relations among urate oxidation genes are not well described by pairwise comparisons. For instance, \(COG3748\) is positively correlated with the presence of \(Urad\) (\(Urad\) implies \(COG3748\) ) and negatively correlated with \(Uox\) (not \(Uox\) implies \(COG3748\)). Although significant, these “association rules”\(^\text{20,21}\) have low confidence (Table 1), reflecting the fact that in many cases \(urad\) is present in the absence of \(COG3748\), and the absence of \(uox\) does not imply the presence of \(COG3748\) (see Fig. 1c and Supplementary Fig. S3). When more genes were included in the comparisons, association rules with increasing confidence and significance were obtained (Table 1). Interestingly, these associations are consistent with the supposed role of \(COG3748\), and the best association rule obtained (93% confidence; \(P = 1 \times 10^{-196}\)) was exactly that expected \(a\) priori by
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Accordingly, we observed growth of

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in a minimal medium supplemented with urate as the

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biochemical reasoning: that is, the presence of both Urah and Urad in the absence of known urate oxi-

dases implies the presence of COG3748.

Based on the above analysis, we predicted a urate oxidase activity for COG3748. The presence of an

integral membrane domain in these proteins (confirmed by sequence analysis, see below) was initially

surprising, as the known urate oxidases are soluble proteins. However, early experiments reported that

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Urate degradation without soluble uricase activity in organisms encoding COG3748
genes. For the anticipated difficulties in the recombinant expression and purification of the integral

membrane COG3748 proteins, we sought to validate the bioinformatics prediction using a genetic

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| Association rule | TT | TF | FT | FF | Confidence | P       |
|------------------|----|----|----|----|------------|---------|
| Urah Urad \( \rightarrow \) zfUox \( \rightarrow \) HpxO \( \rightarrow \) HpyO \( \rightarrow \) COG3748 | 154 | 12 | 5 | 1519 | 0.928 | \( 1 \times 10^{-10} \) |
| Urad \( \rightarrow \) HpxO \( \rightarrow \) HpyO \( \rightarrow \) COG3748 | 155 | 26 | 4 | 1505 | 0.856 | \( 1 \times 10^{-7} \) |
| Urad \( \rightarrow \) COG3748 | 155 | 44 | 4 | 1487 | 0.779 | \( 2 \times 10^{-7} \) |
| zfUox \( \rightarrow \) COG3748 | 157 | 95 | 2 | 1436 | 0.623 | \( 6 \times 10^{-6} \) |
| COG3748 | 158 | 272 | 1 | 1259 | 0.367 | \( 1 \times 10^{-10} \) |
| zfUox \( \rightarrow \) COG3748 | 158 | 1339 | 1 | 192 | 0.106 | \( 1 \times 10^{-8} \) |

Table 1. Association rules implicating COG3748 as consequent. “Best pairwise and higher-order
association rules involving genes for urate oxidation. Genes are connected by logical ‘and’ operators;
exclamation marks indicate the logical ‘not’ operator. The truth table reports the number of cases in which
the antecedent and the consequent of a rule are true (T) or false (F).
**PuuD urate oxidase is an integral membrane protein with a cytochrome c domain.** Pfam analysis of the *A. frarium* PuuD and homologous proteins revealed a bi-domain organization with an N-terminal (aa 3–300) integral membrane domain (DUF898; E = 6 × 10^{-135}) and a C-terminal (aa 330–405) cytochrome c domain (Cyt_c; E = 3 × 10^{-8}). This domain organization was confirmed by the more sensitive HHpred analysis, although this search identified a more significant similarity with another c-type cytochrome domain in Pfam (Haem_bd; E = 6 × 10^{-13}). Besides the significant similarity to Hidden Markov Model (HMM) descriptions of cytochrome c, the presence of this domain is supported by the strict conservation of the canonical CXXCH motif in all PuuD sequences. Sequences representative of the phylogenetic diversity of the PuuD family (Supplementary Fig. S6), where selected to analyse the sequence and structure conservation. The multiple alignment (Fig. 3a and Supplementary Fig. S7) shows high conservation in the N-terminal DUF989 domain, lower conservation in the Cyt_c domain and high variability in the linker region. In the N-terminal domain alignment, eight ungapped hydrophobic blocks are observed. Accordingly, the predictors of transmembrane (TM) domains identified the presence of eight strong TM helices. Consistent results were obtained using single sequences or multiple alignments (Supplementary Fig. S8a,b) and different prediction methods (Supplementary Fig. S8c). Similarly, consistent predictions were obtained for a N_out-C_out membrane topology. With this organization (Fig. 3b), the longer extra-membrane loops (specifically the loops connecting helix I to helix II and helix V to VI) are located in the cytoplasm, whereas the Cyt_c domain is located outside the plasma membrane. Cleaveable signal peptides are not identified in PuuD sequences, so in these proteins helix I is assumed to function as a signal anchor for the targeting to the plasma membrane. Within the DUF989 domain, the stronger conservation is observed in helices I, II, V, VI and in the cytosolic loops I-II and V-VI. Within the Cyt_c domain, conservation pertains especially to the residues that interact with the heme cofactor. Invariant residues in all PuuD sequences are two cysteines of the CXXCH motif (Cys340 and Cys343) that covalently bind the vinyl side-chains of heme through thioether bonds plus the histidine (His344) that provides the fifth heme iron ligand. Also, invariant is a methionine residue (Met385) located about 40 residues further on towards the C-terminus, which in c-type cytochromes provides the sixth iron ligand.

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**Figure 2.** Experimental evidence for the urate oxidase activity of Atu2314 (COG3748). Growth curves of wild-type and engineered *A. frarium* C58 strains in M9 minimal medium supplemented with (a) urate or (b) ammonia as nitrogen source. (c) Urate utilization by concentrated cell cultures. Error bars represent standard deviations obtained from three independent experiments. (d) Uox activity of 250 μg of cell-free extracts as monitored by the decrease in absorbance at 293 nm; extracts were added (arrows) to 0.1 ml solutions containing 0.11 mM urate.
A search with the PuuD C-terminal domain in the Swissmodel server identified more than 50 distinct cytochrome proteins suitable as templates for 3D homology model reconstruction. The best model was obtained using as template the crystal structure of cytochrome c\textsubscript{1} from *Hyphomicrobium denitrificans* (PDB 2D0W), a soluble cytochrome of the periplasmic space acting as the physiological electron acceptor of the methanol dehydrogenase quinoprotein through transient protein-protein interaction\textsuperscript{38}. Only some parts of the structure can be modelled with high confidence, reflecting the low local similarity (22\%) between the template and target sequences (Supplementary Fig. S9a). The positions of residues Cys340, Cys343, His344, and Met385 in the structural model is that expected for heme coordination (Supplementary Fig. S9b), while another invariant residue in the PuuD alignment, Arg398, is directed
towards the protein surface and could be involved in protein-protein interactions. Electrostatic bonds are typically involved in the interaction of c-type cytochromes with partner redox proteins.

**Role of cytochrome c in PuuD urate oxidation.** As a cofactor-free oxidase, the zfUox protein used in our experiments to complement the activity of PuuD deletion mutant transfers two electrons to oxygen and generates hydrogen peroxide as a by-product of the reaction. On the other hand, the structure of the PuuD protein and the presence of an electron transfer cytochrome c domain suggest a completely different mechanism of urate oxidation. When the catalase activity was measured in protein extracts of ΔpuuD/zfUox cells, a four-fold increase of the catalase activity (p < 0.001) was observed for cells grown in urate with respect to cells grown in ammonia (Fig. 4a). Conversely, no differences in the catalase activity were observed in the parental strain in the two growth conditions. Also, no differences were observed when the parental strain was transformed with the empty vector (Fig. 4a). These results can be explained with production of H2O2 in the urate oxidation reaction catalysed by zfUox but not in the reaction catalysed by PuuD. The slight increase in the catalase activity observed in the complemented strain grown in ammonia with respect to the other strains (p < 0.1) could be explained by the degradation of endogenously produced urate. Consistent with induction of catalase by the Uox activity, we found that urate-grown ΔpuuD/zfOux strain was more resistant than the wild-type to oxidative stress, as observed by plating cells on agar medium supplemented with increasing concentrations of H2O2 (Fig. 4b).

Finally, given the presence of a cytochrome c domain in the PuuD protein we observed the effect of an inhibitor of cellular respiration (azide) on urate degradation. Azide inhibited urate utilization by *A. fabrum* C58 in a dose-dependent manner, with a 50% inhibition obtained with about 0.2 mM azide. However, a similar effect was observed in the ΔpuuD/zfOux mutant (Supplementary Fig. S10). As the purified zfOux protein was inhibited by azide at much higher concentrations in vitro (IC50 ~20 mM), we
concluded that the inhibition observed \textit{in vivo} was due to a general effect on cellular metabolism rather than a specific effect on the enzymes of urate oxidation.

\textbf{Discussion}

We have described the identification of a protein family of unknown function (COG3478) as an integral membrane, cytochrome \textit{c} urate oxidase (PuuD). This identification was initially suggested by the analysis of the gene association network (Fig. 1b) and then confirmed by the analysis of gene distribution (Fig. 1c), as quantitatively evaluated through the use of association rules (Table 1). This method, inspired by the logic extension\cite{28} of the widely used correlated and anti-correlated phylogenetic profiles\cite{26,27}, can be useful in the presence of functional associations not adequately described by pairwise relations, as exemplified here. The same analysis that enabled the functional assignment of COG3748, also provides evidence that the identification of PuuD completes the genetic repertoire of enzymes involved in urate oxidation. There is a very small number of cases in which any of the four urate oxidation genes is found in the absence of \textit{urah} and \textit{urad}, suggesting that there are no alternative genes encoding these enzymatic activities; on the other hand, among the 1689 different species considered there are only twelve cases in which the presence of \textit{urah} and \textit{urad} is not explained by the presence of \textit{uox}, \textit{hpxO}, \textit{hpyO}, or \textit{puuD} genes (Table 1). Most of these exceptions can be explained by errors of the gene identification procedure (Supplementary Table S1). Genes encoding PuuD urate oxidase have a peculiar organism distribution, being found only in aerobic bacteria with two cell membranes -diderms, approximately corresponding to gram-negative in the traditional classification\cite{40}. In these organisms, PuuD is the prevalent form of urate oxidase, while Uox is prevalent in monoderm (−gram-positive) prokaryotes (Fig. 5).

A genetic approach was selected for the experimental validation of the activity of PuuD proteins predicted by bioinformatics. The expression and purification of integral membrane proteins is notoriously challenging, and we did not observe protein overexpression when the gene was cloned in a heterologous (\textit{E. coli}) expression system (Supplementary Fig. S11). Furthermore, the presence of a cytochrome \textit{c} domain suggested that the enzymatic activity of this protein could depend on other components of an electron transfer chain. Ultimately, the results obtained with deleted and complemented strains along with bioinformatics provide conclusive evidence for the assignment of a urate oxidase activity to COG3478.

PuuD proteins are characterized by the presence of DUF989, an integral membrane domain with 8 TM helices. Although this organization can be reminiscent of transporter or channel proteins, the evidence presented here indicates that the domain is involved in urate oxidation and not in urate transport. Elements of evidence are the frequent genomic association of PuuD protein with known urate transporters (see e.g. Supplementary Fig. S1) and the complete rescue of the phenotype obtained through complementation of the PuuD deletion mutant with a cytosolic Uox. The analysis of gene context and gene distribution (see Fig. 1c) indicates that all full-length DUF989 proteins have a role in purine degradation; these proteins additionally contain a cytochrome \textit{c} domain. Included in the list of DUF989 members in Pfam is a group of shorter proteins characterized by a partial match to the domain HMM and the absence of the C-terminal cytochrome. These shorter DUF989 proteins contain 4 TM helices corresponding to helices I, II, V, and VI of PuuD (Supplementary Fig. S12). This suggests that helices I, II, V, and VI - the most conserved elements in the PuuD alignment (see Fig. 3) - constitute the core of the DUF989 fold. Genes encoding the shorter DUF989 variant have a genetic context completely different from that of \textit{puuD} genes (Supplementary Fig. S13). Searches based on HMM-HMM comparisons

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure5.png}
\caption{Distribution of urate oxidation genes in monoderm and diderm prokaryotes. The occurrence of the different genes for urate oxidation in 341 prokaryotic species (see Fig. 1c) classified based on the presence of a single (monoderm) or a double (diderm) cell membrane\cite{40}.}
\end{figure}
PuuD proteins are also characterized by the presence of the heme-binding cytochrome c domain. Cytochrome c is an extremely widespread electron transfer domain that mostly occurs in single-domain proteins. However, as in the case of PuuD, it is also often found fused with other protein domains. The association with DUF989 is the third most common multi-domain architecture involving cytochrome c and another Pfam domain. The most common association is with the cytochrome CBB5 domain as is found in ubiquinol-cytochrome c reductases. Other common associations are with the heme binding CCP_MauG domain as is found in di-heme cytochrome c peroxidases, and with the COX2 domain as is found in cytochrome c oxidases. All the cited examples are integral membrane or periplasmic oxidoreductases in which the electron transfer activity of cytochrome c is an essential component of the enzyme catalysis. The presence of a cytochrome c domain in PuuD urate oxidases explains the otherwise puzzling observation that in Sinorhizobium meliloti—a PuuD-encoding organism (see Fig. 1c)—mutations of the ccmC gene coding for an integral membrane heme exporter impair the ability to use purines as nitrogen source. As a protein with a signal anchor, PuuD will be directed to the membrane in an unfolded state, implying that the heme must be assembled with the apo-Cyt_c domain in the extra-cytoplasmic space. Together with the notion that cytochrome c proteins typically function outside the plasma membrane in prokaryotes, the purine degradation deficient phenotype of ccmC mutants provides additional support for the N\textsubscript{out}-C\textsubscript{out} topology of PuuD proteins (see Fig. 3b). According to this topology, conserved extra-membrane regions are exposed towards the cytosol, suggesting that urate oxidation by PuuD takes place on the cytosolic side of the membrane. The electrons removed from the urate molecule could be transferred to the extra-cellular (periplasmic) Cyt_c domain through physical interaction with the transmembrane DUF989 domain. The hypothesis of an oxidation reaction catalysed by DUF908 on the cytosolic side is also consistent with the cytosolic localization of the enzymes (Urah and Urad) acting on purines hypoxanthine and xanthine, by xanthine oxidase (XO), can produce superoxide according to the reaction. If the PuuD reaction mechanism imitates that of the preceding enzyme in the pathway, the presence of the cytochrome c domain would be justified by the scavenging of a reactive oxygen species produce by urate oxidation. In this hypothesis, however, the cytochrome c domain would have an accessory function, while the evidence of the invariant presence of this domain in PuuD proteins and the phenotype of the ccmC mutant rather suggest that the cytochrome c is an essential component of the enzyme.

With four protein families and three different reaction mechanisms, urate oxidase provides an illustrative example of the variety of solutions that can be found in nature for the same biochemical problem. Among these independent inventions, the cofactor-less Uox, which is found in all three domains of life (see Fig. 1c), is probably the most ancient solution, but not necessarily the best one. It appears an ideal solution for an enzyme localized in a single-membrane compartment specialized in the detoxification of reactive oxygen species. PuuD, with its more restricted organism distribution, was apparently invented later in a diderm ancestor (possibly a proteobacteria) through a gene fusion of a preexisting integral membrane domain with a cytochrome c-type heme suggests an unsuspected link between purine catabolism and bacterial respiration. However, we did not observe specific inhibition of the PuuD activity in the presence of a classical inhibitor of the electron transport chain. Azide was not expected to inhibit urate utilization through direct binding to the PuuD protein, but through binding to the cytochrome c oxidase, the terminal acceptor of the respiratory electron chain. The interpretation of the results obtained with the azide inhibitor is complicated by the fact that bacteria have branched respiratory pathway with multiple terminal oxidases exhibiting different sensitivity to the typical respiratory inhibitors. Another complication arises from the known ability of azide to bind the heme cofactor of catalase, which is expected also to affect the activity of the H\textsubscript{2}O\textsubscript{2}-producing zUox. An alternative hypothesis to the electron transfer can be put forward by considering the known ability of cytochrome c to act as a scavenger of the superoxide ion (O\textsubscript{2}−)\textsuperscript{−}. The oxidation of the oxopurines hypoxanthine and xanthine, by xanthine oxidase (XO), can produce superoxide according to the reaction. If the PuuD reaction mechanism imitates that of the preceding enzyme in the pathway, the presence of the cytochrome c domain would be justified by the scavenging of a reactive oxygen species produce by urate oxidation. In this hypothesis, however, the cytochrome c domain would have an accessory function, while the evidence of the invariant presence of this domain in PuuD proteins and the phenotype of the ccmC mutant rather suggest that the cytochrome c is an essential component of the enzyme.

Methods

Analysis of gene associations. The predicted association network for genes involved in urate oxidation was obtained by interrogating the String database (http://string-db.org) with a list of Cluster of Orthologous Groups (COG3648, COG2351, and COG3195) corresponding, respectively, to the uox, urah, and urad gene families. The “gene neighborhood”, “gene fusion” and “gene co-occurrence” evidence were used to predict functional associations. To analyse the distribution of genes involved in urate oxidation in complete proteomes, homology searches where performed using Hidden Markov Models of the corresponding protein families in Pfam (see Supplementary Table S1) and the Hmmsearch program. The results for each family were clustered using Blast best reciprocal hits (BRH) and the MCL program with an inflation index of 3. Genes clusters comprising reference genes involved in purine degradation (see Supplementary Table S1) were considered. The entire procedure was automated using Perl scripts. In the case of hpxO and hpyO, genes belonging to large families containing many paralogs,
the clusters were further refined through phylogenetic analysis (Supplementary Fig. S14). The representation of gene distributions along the species phylogeny (see Fig. 1c) was obtained with the R software using the Ape library\(^\text{34}\) and the heatmap.phylo function (Johan Renaudie; http://stackoverflow.com/questions/15153202).

Association rules\(^\text{30}\) based on gene distributions were determined using the Apriori program\(^\text{31}\) using as input vectors of a gene distribution and its logical negation. Rules implicating COG3748 as consequent (see Table 1) were ranked according to their confidence\(^\text{30}\), defined as the number of cases in which the rule is correct relative to the number of cases in which it is applicable. The significance of the dependency of the antecedent and consequent of a rule was computed through the Fisher's Exact Test. Note that significance is not an indication of the correctness of the logic implication, as the opposite of a rule may be true.

**Sequence and structure analysis.** The ClustalW program was used for multiple sequence alignments and tree reconstruction (based on neighbor-joining clustering of Kimura-corrected distances). Domain analysis was conducted using Pfam\(^\text{36}\) and HHpred\(^\text{37}\). Figures of multiple alignments decorated with structural elements were obtained with the ESPript web server\(^\text{35}\). Transmembrane topology was determined with Phobius\(^\text{46}\) and TopCons\(^\text{37}\). The representation of membrane topology of PudD proteins were obtained with the Protter web server\(^\text{56}\). Homology modelling was performed using Swiss Model (http://swissmodel.expasy.org/) and the Pymol program was used to analyse structural models.

**Deleted and complemented strains.** The deletion of Atu2314 gene of *A. fabrum* C58 was obtained with a described procedure\(^\text{46}\). Briefly, the upstream (1800 bp.) and the downstream (1748 bp.) sequences will receive the same p-value. That significance is not an indication of the correctness of the logic implication, as the opposite of a rule may be true.

**Bacterial growth.** *A. fabrum* C58 strains were first grown on LB medium at 28 °C. After 20 h, bacteria were spun down, washed with M9 minimal medium without any nitrogen and carbon sources and diluted to an optical density at 600 nm (OD\(_{600}\)) of 0.05 in M9 minimal medium with NH\(_4\)Cl (0.02%) and arabinose (0.2%) as sole nitrogen and carbon sources. The cultures were grown for another 20 h, spun down, washed twice with M9 minimal medium without any nitrogen or carbon sources and then diluted to an OD\(_{600}\) of about 0.1 in M9 minimal medium containing arabinose (0.2%) plus NH\(_4\)Cl (0.02%) or uric acid (0.02%), with the addition of IPTG 1 mM where no differently specified. Cultures were incubated with shaking at 28 °C. At intervals, OD\(_{600}\) measurements were taken (Ultraspex 2000, Pharmacia Biotech).

**Urate utilization assays.** Bacteria inoculated at OD\(_{600}\) 0.05 were grown in M9 medium supplemented with arabinose and NH\(_4\)Cl as described. After 20 h, cells were spun down, washed twice with M9 medium without nitrogen or carbon sources and resuspended to an OD\(_{600}\) ~1 in M9 medium containing arabinose and without any nitrogen sources. After 90 minutes, urate (0.02%) was added to the cultures, together with IPTG 1 mM where no differently specified. At intervals, 1 ml of culture was spun down and 0.1 ml of supernatant was diluted with 0.9 ml H\(_2\)O to measure the absorbance at 293 nm (Cary 50, Varian). Urate utilization was measured in the absence and in the presence of different concentrations of sodium azide.

To measure soluble urate oxidase activity, an aliquot of each culture was taken after 5 hours after the addition of urate and IPTG, and the cells were resuspended in 50 mM NaP pH 7.0 containing 1 mM PMSF and lysed by sonication. After centrifugation (7500 rpm for 15 min), the protein concentration of the soluble fraction was determined using the Bradford colorimetric assay, with bovine serum albumin as standard. Cell-free extracts (250 μg) were added to 0.1 ml solutions containing 0.11 mM urate and the reaction was monitored by following the decrease in absorbance at 293 nm.

**Catalase activity and oxidative stress assay.** Cells were grown starting from OD\(_{600}\) ~0.1 in M9 medium containing arabinose (0.02%) and NH\(_4\)Cl (0.02%) or urate (0.02%) with the addition of IPTG 1 mM where no differently specified. After 12 hours, cells were spun down and the cell-free extracts were obtained as described above. The catalase activity of each sample was measured spectrophotometrically by following the decrease of absorbance at 240 nm\(^\text{59}\). One unit of enzyme was defined as the amount of enzyme catalysing the turnover of 1 μmol of substrate per minute under the assay conditions. To measure the resistance of different strains to H\(_2\)O\(_2\), cells grown in urate for 12 h were spun down, washed once with 50 mM sodium phosphate buffer pH 7.0 and diluted in the same buffer to an OD\(_{600}\) of 0.1. Progressive dilutions (1:5) of the cells were spotted on LB agar plates with different H\(_2\)O\(_2\) concentration and incubated overnight at 28 °C.
PuuD cloning and expression. A PuuD expression vector was constructed by cloning the Atu2314 (PuuD) coding sequence into the pET11b vector. The Atu2314 coding sequence was amplified from A. fabrum strain C58 genomic DNA using Taq DNA polymerase with primer forward 5′-CATATGAGCATACTGCCATTG and primer reverse 5′-GGATCCACGAGCAAGAACG in standard reaction conditions. The PCR product was cloned in pGEM-T easy vector (Promega), which was then digested with NdeI and BamHI (Thermo Scientific) and ligated to a linearized pET11b vector DNA with compatible ends. The correct insertion of the PuuD coding sequence was verified by sequencing (Supplementary Fig. S11). Expression attempts were conducted in transformed BL21 cells (Novagen). Different induction conditions with IPTG 1 mM were tested, from 4 to 16 hours and from 20°C to 37°C.

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Acknowledgements
We thank Romina Corsini, Ileana Ramazzina, and Giulia di Punzio for practical help and suggestions. The kind gift of the pSRKGm plasmid by Stephen K. Farrand is gratefully acknowledged. This work was funded by Telethon Italy (grant GGP13149).

Author Contributions
N.D. and E.M. performed the experiments, R.P. carried out the bioinformatics analysis, A.D.P. and A.M. assisted with the data analysis, and R.P. and N.D. wrote the manuscript. All authors reviewed the manuscript.

Additional Information
Supplementary information accompanies this paper at http://www.nature.com/srep
Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Doniselli, N. et al. The identification of an integral membrane, cytochrome c urate oxidase completes the catalytic repertoire of a therapeutic enzyme. Sci. Rep. 5, 13798; doi: 10.1038/srep13798 (2015).

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