# A novel family of plasmid-transferred anti-sense ncRNAs

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The genome of Xanthomonas campestris pv. vesicatoria encodes a constitutively expressed small RNA, which we designate PtaRNA1, "Plasmid transferred anti-sense RNA". It exhibits all hallmarks of a novel RNA antitoxin that proliferates by frequent horizontal transfer. It shows an erratic phylogenetic distribution with occurrences on chromosomes in a few individual strains distributed across both beta- and gammaproteobacteria. Moreover, a homologous gene located on plasmid pMATVIM-7 of Pseudomonas aeruginosa is found. All ptaRNA1 homologs are located anti-sense to a putative toxin, which in turn is never encountered without the small RNA. The secondary structure of PtaRNA1, furthermore, is very similar to that of the FinP anti-sense RNA found on F-like plasmids in Escherichia coli.

### Introduction

Several toxin-antitoxin systems of type 1, in which the toxin is a short protein and the antitoxin an anti-sense RNA and of type 2, where both elements are proteins, are frequently found in both prokaryotic chromosomes and plasmids.<sup>1-4</sup> The paradigmatic example for type 1 is the plasmid encoded hok/sok system in Escherichia coli and its close relatives. The toxinencoding stable mRNA encodes a protein that rapidly leads to cell-death unless its translation is suppressed by a short-lived small RNA. The plasmid encoded module prevents the growth of plasmid-free offsprings thus ensuring the persistence of the plasmid in the population: After

cell division, plasmid-free cells still contain the stable toxin mRNA, while the comparably unstable antitoxin is quickly depleted. It is poorly understood how the chromosomally encoded systems function. Interestingly, the SOS-induced genes *tisB* and *symE* are expressed under very specific stress conditions. The corresponding antitoxins (SymR and Sib) are constitutively expressed.

Although distinct toxin-antitoxin systems have been found in widely separated bacterial groups (e.g., *hok/sok* in *E. coli* and *txpA/ratA* in *Bacillus subtilis*<sup>5</sup>), each of the known examples exhibits a very narrow phylogenetic distribution.

In this contribution we characterize by computational means a small RNA that has all the hallmarks of the known type 1 toxin-antitoxin systems but shows a rather wide spread erratic phylogenetic distribution that hints at frequent horizontal gene transfers.

### Results

The founding member, PtaRNA1 ("Plasmid transferred anti-sense RNA"), of the family was detected in a library of pyrosequencing data of Xanthomonas campestris pv. vesicatoria strain 85-10 (Xcv) that was prepared and analyzed for unrelated purposes. The superposition of the individual reads revealed a small RNA encoded adjacent to the trbL gene. Expression and approximate size of the small RNA was verified by northern blot (Fig. 1). These analyses revealed a constitutive expression with respect to the tested growth phases. Interestingly,



**Figure 1.** Expression of PtaRNA1 and 5S RNA in exponential and stationary growth phase of *Xcv* analysed by northern blot. The size of corresponding marker bands is indicated on the left.

two bands which indicate procession of the full length PtaRNA1 are detected in the exponential but not in the stationary growth phase.

Chromosomally encoded homologs of *ptaRNA1* were found in beta-proteobacteria (Nitrosomonas eutropha C91, Azoarcus sp. EbN1, Verminephrobacter eiseniae EF01-2, Burkholderia cenocepacia J2315, B. pseudomallei K96243, B. pseudomallei 9, B. pseudomallei 91, and Acidovorax JS42) as well as gamma-proteobacteria (X. campestris pv. vasculorum NCPPB702, Shigella flexneri 2a 2457T, Acinetobacter baumannii ATCC 17978, Marinobacter aquaeolei VT8, and Pseudomonas aeruginosa UCBPP-PA14). Two ptaRNA1 copies were found in N. eutropha C91 and are named *ptaRNA1*-a and *ptaRNA1*-b. This observation is in accordance with various reported insertion, duplication

and rearrangement events within this species.<sup>6</sup>

Conspicuously, ptaRNA1 was not found in other closely related genomes, e.g., other strains of Burkholderia, Pseudomonas or Xanthomonas. This distinguishes PtaRNA1 from most other bacterial small RNAs, such as the cyanobacteria-specific Yfr RNAs.7 In addition to the chromosomal loci listed above, we found a *ptaRNA1* homolog in the P. aeruginosa plasmid pMATVIM-7, adjacent to the transfer region, a gene cluster that encodes proteins of unknown function and the plasmid stabilization protein ParE. Figure 2A depicts the alignment and the resulting consensus secondary structure of all detected PtaRNA1 homologs.

Phylogenetic analysis of the PtaRNA1 sequences (Fig. 3) shows that the



**Figure 2.** (A) Consensus secondary structure model of PtaRNA1 based on the depicted seed alignment. The structure is highly stable (minimum free energy -37 kcal/mol) and supported by various compensatory mutations within the stem on the right-hand side. Marked in blue is the region complementary to the putative Shine-Dalgarno sequence of the *XCV2162* mRNA. (B) Amino acid alignment of XCV2162 homologs. The alignment shows various totally (indicated by '\*') and by substitutions (indicated by ':' and '.') supported and therefore conserved columns. The protein topology of a trans-membrane domain, predicted by MEMSAT3,<sup>8</sup> is indicated as well. '+' marks inside loop, '~' outside loop, 'O' outside helix cap, 'X' central trans-membrane helix segment and 'l' inside cap. The truncated Verminephrobacter sequence was not used for the calculation of the conservation track.



**Figure 3.** Phylogenetic tree based on PtaRNA1 alignment (similar for XCV2162 alignment, data not shown). Class of the "host" species is shown by the symbols on the right hand side. Numbers indicate bootstrap values of the inner nodes.

phylogeny of the PtaRNA1 sequences is not congruent with the phylogeny of their "host" species. This indicates that the proliferation of *ptaRNA1* depends on frequent horizontal transfer, presumably by means of plasmids.

The *ptaRNA1* gene in *Xcv* is located anti-sense to a so far uncharacterized protein coding gene (*XCV2162*). The gene is adjacent to *trbL*, encoding a type IV secretion system protein. The small overlap of both genes strongly suggests that PtaRNA1 is an anti-sense regulator of *XCV2162* (Fig. 4). We therefore searched the complete set of eubacterial genomes for homologs of *XCV2162* and found that *ptaRNA1* and *XCV2162* co-occur in all cases, indicating their functional linkage. This is in particular also the case in the *P. aeruginosa* plasmid pMATVIM-7, whose uncharacterized gene p07-406.22is an ortholog of *XCV2162*, as in *Xcv* adjacent to *trbL* (Fig. 4). According to MEMSAT3<sup>8</sup> prediction, XCV2162 contains a trans-membrane domain (Fig. 2B), as it is also the case for many reported toxic proteins.<sup>3,4</sup>

Furthermore, the gene phylogeny of the XCV2162 proteins (not shown) is congruent with the phylogeny of PtaRNA1 sequences, indicating that they are transferred together. We observed a frequent co-occurrence of *ptaRNA1/XCV2162* and *trbL* homologs, albeit *trbL* was detected in at least 155 eubacterial genomes, suggesting that *trbL* might have a role in the frequent chromosomal insertions of the *ptaRNA1/XCV2162* system. In *V. eiseniae* EF01-2 we found a truncated *XCV2162* homolog. Verminephrobacter is also the only *ptaRNA1* encoding species in which no *trbL* homolog was found.

Analysis of the putative *ptaRNA1* promoter regions revealed the existence of two highly conserved sequence motifs of eight nucleotides. The first one starts between 42 and 36 nt upstream and the other 13/12 nt upstream of the transcription start corresponding to the -35 and -10 box, respectively (Fig. 4). In the upstream region of *XCV2162*, an ultra conserved AG-rich motif was found (Fig. 4), probably representing the Shine-Dalgarno sequence of the mRNA. This motif is entirely covered by the complementary *ptaRNA1* sequence (Fig. 2A).

The consensus secondary structure of PtaRNA1 consists of a 5'-stem loop and a long 3' stem which presumably acts as terminator hairpin, (Fig. 2A). This structure is very similar to that of FinP found in *E. coli* (data not shown). Interestingly, FinP is the anti-sense regulator of TraJ, a transcriptional activator required for expression of various conjungative protein components.<sup>9</sup> Thus, FinP is only one component in a complex network of several interacting molecules.

# Discussion

All evidence available for PtaRNA1 suggests that XCV2162/ptaRNA1 is a novel toxin-antitoxin system: both genes are only found as combined cluster and in fact do not appear as single genes; XCV2162 encodes a relatively short protein that shows the typical topology of toxins with a trans-membrane domain; the presence on a plasmid in combination with the erratic phylogenetic distribution of the system indicates a frequent horizontal gene transfer. It is known that type 2 systems, where both molecules are proteins, show an erratic phylogenetic distribution.<sup>4</sup> We assume that this might also be the case for type 1 systems, such as the one presented here.

Furthermore, the phylogenetic distribution of the *XCV2162/ptaRNA1* pair indicates a very rapid loss of the chromosomal homologs: the erratic distribution suggests that we only see very recent chromosomal insertions. The homolog in Verminephrobacter with its truncated *XCV2162* coding sequence might



**Figure 4.** Surrounding genomic location of the *ptaRNA1* gene in *Xcv*. On the plus strand the coding sequences of *trbL* and *XCV2162* are indicated. In front of *XCV2162* an ultra conserved AG rich motif, the putative Shine-Dalgarno sequence is shown. *ptaRNA1* is encoded on the minus strand. A conserved -10 as well as -35 box (sequence logos) was found directly upstream of this gene.

represent the first step towards the complete loss of the system.

This apparent evolutionary instability further supports the hypothesis that *XCV2162/ptaRNA1* is a toxin-antitoxin pair. Only chromosomal integration of the toxin-antitoxin pair makes the plasmid dispensable. Thus, cell-death is prevented by chromosomal integration of the system. Plasmid-loss and subsequent destruction of the toxin *XCV2162* then leaves the chromosomal copy of the antitoxin PtaRNA1 without function, so it is also rapidly removed from the genome.

## **Materials and Methods**

For northern blot analysis Xanthomonas campestris pv. vesicatoria strain 85-10 was cultivated at 30°C in nutrient-yeast-glycerol medium.10 Cells were harvested at exponential and stationary growth phase at OD of 0.6 and 1.5, respectively. RNA was extracted as described in.11 Northern blots were done according to12 with the following modifications: 20 µg RNA were separated on 8.3 M urea-6% polyacrylamide gels. For detection of PtaRNA1 and 5S rRNA membranes were incubated for 1 h at 42°C with Rapid-hyb<sup>TM</sup> Buffer (GE Healthcare) containing <sup>32</sup>P 5' endlabeled oligodeoxyribonucleotides NB39 (5'-ATG GAG AGG TGA ATC ATG GC-3') and NB5S (5'-ATG ACC TAC TCT CGC ATG GC-3'), respectively.

Homology searches were based on scans of the bacterial NCBI genome database (ftp://ftp.ncbi.nih.gov/genomes/ Bacteria/downloaded 06/12/2009) as well as the plasmid genome database (http:// www.genomics.ceh.ac.uk/plasmiddb/ downloaded 06/12/2009). Homologs of protein coding genes were searched using tblastn of the Blast package.<sup>13</sup> Since noncoding RNAs may vary in sequence but still fold into the same secondary structure a semi-global alignment implementation, GotohScan,<sup>14</sup> was used to scan for *ptaRNA1* homologs. The microbial web Blast (http://www.ncbi.nlm.nih. gov/sutils/genom\_table.cgi) was used to search for additional homologs especially in unfinished genome projects.

Alignments were calculated with ClustalW<sup>15</sup> and locARNATE<sup>16</sup> for sequence structure alignments, respectively. The consensus structure model was calculated with RNAalifold.<sup>17</sup>

Using, MEME<sup>18</sup> we analyzed the 100 nt upstream region of all homologous *ptaRNA1* loci. MEME searches for similarities among the given sequences and calculates descriptors for these motifs. To search for known regulatory sites within the 100 nt upstream region the PRODORIC database was queried using the Virtual Footprint v3.0 web tool.<sup>19</sup>

Submitted data. This manuscript documents the seed alignment ptaRNA1. seed.stk. A short summary is available as Wikipedia Entry at http://en.wikipedia. org/wiki/User:SveFinBioInf/ptaRNA1.

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Note

Supplementary materials can be found at:www.landesbioscience.com/supplement/FindeissRNA7-2-Sup.txt

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