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Mycobacterium Tuberculosis Signaling via c-AMP

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1. Introduction

1.1 Cyclic Adenosine Monophosphate (cAMP) metabolism in mycobacteria

1.1.1 General aspects of Adenylyl Cyclases and their presence in *M. tuberculosis* genomes

Adelynate cyclases (ACs), which catalyze synthesis of cAMP from ATP and yield pyrophosphate as a by-product, can be classified into four different classes according to their common features: Class I cyclases, related to enterobacterial adenylate cyclases; Class II, toxic adenylate cyclases isolated from bacterial pathogens; Class III, a large and probably ancient class that comprises cyclases from both eukaryotes and prokaryotes and is strongly related to guanylate cyclases; and Class IV, with mainly one example that differs entirely from all other classes (McCue *et al.*, 2000).

In class I ACs (the enterobacterial type) no long stretch of hydrophobic amino acid residues is present to explain the membrane-bound localization of the adenylate cyclases. In all cases, the proteins are very rich in cysteine residues, an uncommon feature for proteins located in the cytoplasm or at the cytoplasmic border of the membrane. They are also rich in histidine residues, which could indicate that metal ions take part in the folding and/or activity of the polypeptide chain (Mock *et al.*, 1991).

Class II ACs (the calmodulin-activated toxic class) is represented by *Bordetella pertussis* adenylate cyclase. It is synthesized as a large bifunctional polypeptide chain of 1706 amino acid residues. The N-terminal segment of the protein (400 residues) alone displays calmodulin-activated adenylate cyclase activity, whereas the rest of the molecule is responsible for hemolytic activity and for transporting the toxin. After attempts to isolate other members of this class, several examples of similar proteins have now been discovered in *Bacillus anthracis*, *Pseudomonas aeruginosa*, and in *Yersinia* species. Comparison of the catalytic regions of the *B. pertussis* and *B. anthracis* adenylate cyclases identified four conserved regions that are involved in catalysis, calmodulin binding and activation. The first region comprises a sequence, Gly-XXXX-Gly(Ala)-Lys-Ser, similar to the nucleotide-binding motif found in many ATP- or GTP-binding proteins. Analysis of the region conserved between the *B. anthracis* and *B. pertussis* enzymes, indicates that these proteins

may form a catalytic center from the cooperation of two halves. The function of calmodulin may be to trigger the appropriate conformational change necessary to form an active catalytic center (Drum *et al.*, 2002).

Class III ACs (the “universal” class) form a very diverse collection of enzymes in eubacteria. They comprise two domains: the catalytic domain is carboxy-terminal and the regulatory domain is likely an ion transporter in one case and the phosphorylated moiety of a two-component regulatory system in another. Most mammalian ACs are monomeric integral membrane proteins that are catalytically active as pseudoheterodimers (Sunahara *et al.*, 1996), while prokaryotes and lower eukaryotes produce both soluble and membrane-bound nucleotidyl cyclases of variant domain compositions functioning as homodimers (Guo *et al.*, 2001). In general, class III ACs are the most widespread class of cAMP-generating enzymes, and they are further subdivided into four subclasses: IIIa-III d. Dimerization is required for all class III ACs in order to be active, given the substrate-binding sites are formed at the dimer interface (Abdel Motaal *et al.*, 2006). Class III adenylyl and guanylyl cyclases are proteins with a central four stranded anti-parallel β -sheet structurally similar to the palm domain of DNA polymerases, and α -helices on either side (Shenoy & Visweswariah, 2006b). So far, all Cyclase Homology Domain (CHDs) proteins operate as dimers with mostly two catalytic centers positioned at the dimer interface, where catalysis is based on six highly conserved residues. Two aspartate residues coordinate two metal cofactors (Mg^{2+} or Mn^{2+}), an asparagine and an arginine stabilize the transition-state and a lysine-aspartate couple specifies ATP as a substrate. Several mycobacterial ACs gene products were early annotated as putative cyclases, but have now been characterized biochemically and structurally.

Class IV ACs was assigned to *Aeromonas hydrophila*, which synthesizes a very small cyclase of 193 residues. This class IV cyclase has an optimal temperature for activity of 65°C and is at least ten times more active than the class I adenylyl cyclase in the same organism (Sismeiro *et al.*, 1998). No function has yet been discovered for this protein. Currently, it has been found only in various isolates of *A. hydrophila* and in *Y. pestis*.

Most prokaryotes contain a single adenylyl cyclase (AC, e.g. *E. coli*) and some contain none (e.g., *Bacillus* species); however, 16 or 17 genes have been identified as ACs in mycobacteria, and in particular *M. tuberculosis*. AC enzymes require the presence of conserved metal-, substrate-binding (either ATP or GTP) amino acid residues and transition-state stabilizing amino acid residues in their catalytic site (Shenoy & Visweswariah, 2006b).

The genome of *M. tuberculosis* H37Rv encodes 16 class III cyclases while the CDC1551 strain has 17 cyclases. Using the 16 H37Rv class III ACs genes as query, we found they possess different numbers of orthologous genes in the up-to-date sequenced mycobacterial genomes available at MycoDB (<http://xbase.bham.ac.uk/mycodb/>, Table 1). This suggests that some of these cyclases have more conserved roles than others. Several mycobacterial gene products initially classified as cyclases have now been biochemically and structurally characterized as ACs. In mycobacteria, they retain similar catalytic properties to the mammalian enzymes, in terms of the requirement for divalent cations, and dimerization as a prerequisite to generate the catalytic site.

Bioinformatics analysis has predicted subcellular localization of mycobacterial ACs (Shenoy *et al.*, 2004). Nevertheless, experimental studies have identified proteins like Rv0386 and Rv1358 in mycobacterial cell wall and membranes, despite the absence of

hydrophobic, predicted transmembrane domains (Agarwal, 2009). This suggests that a necessary improvement to prediction software must emphasize these particularities of mycobacteria.

Gene name	Orthologous (No.)
<i>Rv0386</i>	7
<i>Rv0805</i>	7
<i>RV0891c</i>	4
<i>Rv1264</i>	13
<i>Rv1318c</i>	14
<i>Rv1319c</i>	14
<i>Rv1320c</i>	14
<i>Rv1358</i>	7
<i>Rv1359</i>	3
<i>Rv1625c</i>	11
<i>Rv1647</i>	16
<i>Rv1900c</i>	10
<i>Rv2212</i>	6
<i>Rv2564</i>	3
<i>Rv2565</i>	3
<i>Rv3645</i>	16

Table 1. Orthologous to H37Rv AC-domain containing genes present in Mycobacteria

As mycobacteria lack G-proteins, it is unknown how bacterial adenylate cyclase AC activities are modulated. However, polyphosphates isolated from *M. bovis* BCG were potent inhibitors of *Rv1625c*, *Rv1264*, and *Rv3645*, and the mechanism of inhibition proposed is that polyphosphates possibly obstructs the catalytic fold of ACs (Guo et al., 2001).

1.2 cAMP producing (Adenylyl cyclases) and degrading (Phosphodiesterase) mycobacterial enzymes

Six *M. tuberculosis* AC-domain containing proteins (*Rv0891c*, *Rv1264*, *Rv1359*, *Rv1647*, *Rv2212* and *Rv1625c*) contain just a cyclase domain. *Rv1647* and *Rv1625c* are phylogenetically distant and biochemically distinct AC-domain containing proteins (Shenoy & Visweswariah, 2006a). The remaining AC-domain containing proteins have additional domains that presumably allow them to respond to multiple signals, regulate their activity in response to environmental conditions, and/or expand their repertoire with effector function capability. Five of these multidomain AC-domain containing proteins (*Rv1318c*, *Rv1319c*, *Rv1320c*, *Rv2435c* and *Rv3645c*) are membrane-associated and contain HAMP (histidine kinases, adenylyl cyclases, methyl-accepting chemotaxis proteins and phosphatases) domains. HAMP domains are often associated with two-component signal transduction pathways and connect the sensing of extracellular environmental signals with

responding intracellular signalling domains. Mycobacterial ACs with HAMP domains have six transmembrane regions, followed by the HAMP domain and a C-terminal AC domain. Modulators of *M. tuberculosis* AC activity include pH and fatty acids, which were seen to enhance the pH sensitivity of the holoenzyme, and CO₂ levels, and could be signals present during *M. tuberculosis* host infection (Abdel Motaal et al., 2006, Bai et al., 2011, Barba et al., 2010).

An N-terminal autoregulatory domain in Rv1264 is a pH-response element that inhibits cyclase activity above pH 6.0. Additional AC-domain containing proteins (Rv0386, Rv1358 and Rv2488c) possess both ATPase and helix–turn–helix (HTH) domains. Rv1900c contains an $\alpha\beta$ -hydrolase domain (Barba et al., 2010, Bai et al., 2011). Rv0386 is the first AC-domain containing protein as a representative of the family of putative DNA-binding domain-containing cyclases in mycobacteria (Castro et al., 2005). Rv0386 has guanylyl cyclase activity that is 20% of its adenylyl cyclase activity.

Rv1625c adenylyl cyclase is particularly unusual, in that it is catalytically active when expressed in mammalian cells or in *E. coli*. The Rv1264 holoenzyme shows higher AC activity at acidic pH (pH 6.0). Similarly, the Rv1264 AC is activated only at low pH (pH 5.5) whereas Rv1647 is active only at high pH (pH 8.5). The localization of Rv1647 in the cell wall and membrane fractions of *M. tuberculosis* might enable the bacteria to sense and respond to extracellular pH shifts. Because *M. tuberculosis* actively avoids phagosomal acidification, sensing of pH and bicarbonate and/or CO₂ could be crucial signaling events in *M. tuberculosis* pathogenesis (Shenoy & Visweswariah, 2006a). In fact, cAMP production in pathogenic mycobacteria increased when the pH of their growth medium was shifted from pH 6.7 to pH 5.5 (Gazdik et al., 2009), thus suggesting activation of the acid-responsive ACs.

In *M. tuberculosis*, Rv0998 was recently shown to regulate protein lysine acetylation in a cAMP-responsive manner (Nambi et al., 2010), where acetylation is enhanced in the presence of either cAMP or cGMP. Recently, it was shown that a cAMP-dependent protein acetyltransferase inactivates ACs through acetylation of a single, specific lysine residue, and Rv1151c is a NAD⁺-dependent deacetylase that reactivates ACs. This acetylation/deacetylation system in mycobacteria is likely to sense the extracellular environment through cAMP levels and also key intracellular metabolites, including NAD⁺ and AcCoA, since cAMP, AcCoA, and NAD⁺ are required for acetylation and deacetylation, respectively (Xu et al., 2011).

Adenylyl cyclase Rv2212 from *M. tuberculosis* has a domain composition identical to the pH-sensing isoform Rv1264, an N-terminal regulatory domain and a C-terminal catalytic domain. The maximal velocity of Rv2212 was the highest of all 10 mycobacterial cyclases investigated to date. Unsaturated fatty acids strongly stimulated Rv2212c activity by increasing substrate affinity (Findeisen et al., 2007). In addition, fatty acids greatly enhanced the pH sensitivity of the holoenzyme, thus converting Rv2212c to a pH sensor adenylyl cyclase (Abdel Motaal et al., 2006).

To date, the only identifiable cAMP phosphodiesterase (PDE) in the genome of *M. tuberculosis* H37Rv is the one encoded by the *Rv0805* gene. This enzyme is a dimeric Fe(3⁺)-Mn(2⁺) binuclear PDE, where metals coordinated at the catalytic site contribute to dimerization and thus play an additional structural role apart from their involvement in

catalysis (Shenoy *et al.*, 2007). As this gene is found only in pathogenic mycobacteria, Rv0805 may therefore play a key role in the pathogenicity of mycobacteria, not only by hydrolyzing bacterial cAMP, but also by indicating as a protein that can alter cell wall functioning (Podobnik *et al.*, 2009).

2. Participation of cyclic adenosine 3',5'-monophosphate (cAMP) in mycobacterial gene regulation

Cyclic adenosine 3',5'-monophosphate (cAMP) is one of the most important second messengers used in bacteria and it has been characterized principally in *Escherichia coli*. However its signaling role in *M. tuberculosis* is beginning to emerge. A large number of AC genes are present in *Mycobacterium tuberculosis* in comparison with other microorganisms (Cha *et al.*, 2010, Klengel *et al.*, 2005, Shenoy *et al.*, 2004), and also that *Mycobacterium* just have one class III cNMP phosphodiesterase (Rv0805) that has been identified in *M. tuberculosis* (McCue *et al.*, 2000, Shenoy *et al.*, 2007) with a modest ability to efficiently hydrolyse 3',5'-cAMP and that responds to H₂O₂ *in vitro* (Bai *et al.*, 2011, Barba *et al.*, 2010). cAMP may serve as both an extra and intracellular signaling molecule in mycobacteria (Agarwal, 2009). cAMP levels are modulated by stress conditions in *M. smegmatis*, where it has been found in concentrations between 10 µM and 1 mM, suggesting that cAMP participates in signalling events within the bacterial cell, mediating its action by downstream effectors (Dass *et al.*, 2008).

It appears that Mycobacteria need to maintain a steady level of cytoplasmic cAMP in many conditions. For example, it has been shown that cytoplasmic cAMP levels in *E. coli* are reduced three- to fourfold when the carbon source is ~0.2% glucose rather than glycerol (Bai *et al.*, 2011). In contrast, a recent study showed no significant change in the cytoplasmic cAMP levels of *M. bovis* BCG incubated with 0.2% glucose (Bai *et al.*, 2009), or carbon-starved bacteria (Dass *et al.*, 2008). cAMP levels decrease in both fast- and slow-growing mycobacteria in response to very high levels of glucose (2%) (Bai *et al.*, 2011). cAMP levels have been suggested to be high in *Mycobacterium* cells, exceeding up to 100-fold levels found in other bacteria (Nambi *et al.*, 2010, Stapleton *et al.*, 2010, Shenoy & Visweswariah, 2006b, Rickman *et al.*, 2005). However, it is difficult to make a comparison between studies, due to cAMP variations presents in the conditions tested, as well as the different normalization and reporting methods used.

2.1 CRP and cNMP binding proteins

In silico studies predict 10 cNMP binding proteins that encompass a wide range of potential effector functions, suggesting a more complex role for cAMP signalling, probably important during host infection. From these 10 cNMP binding proteins, 7 (Rv0073, Rv0104, Rv2434c, Rv2564, Rv2565, Rv3239c and Rv3728) contain an assortment of putative functional domains, including those associated with transport functions and esterase activities (McCue *et al.*, 2000, Shenoy & Visweswariah, 2006a). Only three cNMP binding proteins have been functionally characterized to date (Bai *et al.*, 2011). Two of these proteins, referred to as CRP (Rv3676, for Catabolite Represor Protein) and Cmr (Rv1675c, for cAMP and macrophage regulator), contain a HTH DNA binding domains, and belong to the CRP-FNR family of transcription factors (McCue *et al.*, 2000). The third protein from this group, is encoded by

Rv0998 in *M. tuberculosis*, and regulates lysine acetylation in mycobacterial proteins in a cAMP-responsive manner (Nambi et al., 2010). The biological effects of this acetylation are not yet defined.

CRP (cAMP receptor protein) in *Escherichia coli*, is one of the best-studied prokaryotic transcription factor. Currently, a total of 378 target promoters on the *Escherichia coli* genome are proposed to be under the control of cAMP-bound CRP, using a SELEX approach (Shimada et al., 2011). Some of the CRP regulon genes include those encoding the transporters and the catabolic enzymes of glucose (Perrenoud & Sauer, 2005) and non-glucose sugars (Wickstrum et al., 2010), virulence genes (Espert et al., 2011), motility genes (Hollands et al., 2010), GMP synthesis (Husnain et al., 2009), anaerobic growth and nitrate reductases genes (Stewart et al., 2009). To date, in *E. coli*, 2 different ways in which CRP leads the sigma 70 promoter transcription have been described: in the Class I promoters, CRP binds upstream of the promoter -35 element, at a site centered at position -61.5 or further upstream, and an activating region (AR1) in the downstream subunit of the CRP dimer makes contact with the C-terminal domain of one of the two RNA polymerase α subunits (α CTD). In class II promoters, CRP binds at a target that overlaps the promoter -35 element and is usually centered at position -41.5. AR1 in the upstream subunit of the CRP dimer interacts with α CTD, while a second activating region (AR2) in the downstream subunit interacts with the N-terminal domain of one of the two RNA polymerase α subunits (α NTD) (Hollands et al., 2010).

In a similar manner to the CRP protein from *E. coli*, the *M. tuberculosis* CRP (Rickman et al., 2005) is the best-studied example of a protein implicated in cAMP-mediated signaling. In *M. tuberculosis* CRP is encoded by the gene *Rv3676* and it is homologous to *E. coli* CRP. In *E. coli*, it has been shown that CRP regulates expression of genes required to control metabolism, as well as growth under hypoxic and nutrient-deprived conditions. Like *E. coli* CRP, *M. tuberculosis* *Rv3676* possesses an N-terminal cAMP-binding domain and a C-terminal DNA-binding domain. The crystal structures of *M. tuberculosis* CRP at 2.2 and 2.0 Å resolution of cAMP-bound (Reddy et al., 2009) and the apo-form (Gallagher et al., 2009), have been reported. Conformational changes required for DNA-binding do not take place in the absence of the second messenger (Reddy et al., 2009). In fact, as opposed to *E. coli* CRP where cAMP binding follows a cooperative mechanism, cAMP binding sites are independent, and DNA-binding activity is not as enhanced with *M. tuberculosis* *Rv3676* (Stapleton et al., 2009). The CRP orthologous in *M. bovis* BCG (CRP_{BCG}) is a fully functional transcription factor, since CRP_{BCG} overcame the virulence deficiency of an *M. tuberculosis* *crp* mutant (Hunt et al., 2008). Although both of them have a similar ability to bind cAMP and DNA, CRP_{BCG}'s DNA binding affinity is approximately twice that of *Mycobacterium tuberculosis* CRP (CRP_{Mt}) (Bai et al., 2007, Hunt et al., 2008). Interestingly, CRP_{BCG} differs from CRP_{Mt} in just two amino acid residues (L47P and E178K).

The other predicted cAMP-dependent transcriptional regulator, Cmr, was found to negatively regulate the expression of five proteins (GroEL2, Rv2971, PE_PGRS6a, Mdh and Rv1265) (Gazdik & McDonough, 2005). The upstream regions of three of these genes (*mdh*, *groEL2* and *Rv1265*) bound specifically to Cmr in electrophoretic mobility shift assays, consistent with direct regulation of these genes by Cmr. Expression of three of these genes was found to be regulated within macrophages, and this regulation was mediated by Cmr in both *M. tuberculosis* and *M. bovis* BCG (Gazdik et al., 2009). Despite

the importance of Cmr for the cAMP- dependent regulation of these genes, Cmr has not been shown to directly bind cAMP *in vitro*, and cAMP did not affect Cmr's binding to any of their promoter sequences (Gazdik et al., 2009). The mechanism by which Cmr responds to cAMP levels has yet to be discovered, and it is possible that a second factor plays a facilitating role.

2.2 CRP_{Mt} regulon

The cAMP binding transcriptional regulator encoded by *Rv3676* in *M. tuberculosis* has been strongly associated with growth during mycobacterial life cycle. Its deletion resulted in impaired growth in macrophage cell lines and in mice (Hunt et al., 2008, Rickman et al., 2005). Using an exponential enrichment (SELEX) approach, Bai et al. defined the CRP_{Mt}'s palindromic binding motif (C/TGTGANNNNNT CACG/A) based on 58 predicted binding sites from the *M. tuberculosis* genome, using a combination of *E. coli* CRP binding sites and *M. tuberculosis* DNA sequences recovered by affinity capture using CRP_{Mt} to seed the computational analyses (Bai et al., 2005). Akhter et al. used the positional Shannon relative entropy method to predict 19 new putative binding sites for *M. tuberculosis* (Rv3676) CRP, in addition to the 73 sites previously predicted by Bai et al. (Akhter et al., 2008, Bai et al., 2005). These additional sites resulted from a difference where Akhter et al. used only the information available from the *M. tuberculosis* CRP-regulon instead of adding up the one available from the *E.coli* CRP-regulon. According to Akhter et al., the *M. tuberculosis* CRP-regulon comprises genes required for critical functions like: (i) cell-wall biogenesis, (ii) central metabolism pathways, (iii) putative regulatory elements controlling cAMP signaling, and (iv) antibiotic resistance (Akhter et al., 2008) (Table 2).

Experimental validation of these predictions has only been completed for Bai et al. model (Bai et al., 2005). Their research showed that mutation of nucleotides G2 or C15 from the palindromic binding motif, abolished CRP_{Mt}'s binding. Both positions are conserved in all predicted binding sites. They also evaluated CRP_{Mt} and CRP_{BCG} binding to seven putative CRP_{Mt} sites, located within intergenic regions, where 6 out of these 7 were found to be functional *in vitro* and *in vivo* (Bai et al., 2007). This increases confidence in the prediction algorithm that was used to identify CRP_{Mt} binding sites, although at this point it is possible to suggest that CRP_{Mt} regulon requires further refinement. On the other hand, it has been shown that *rpfA* is directly activated by CRP_{Mt} (Rickman et al., 2005). The resuscitation-promoting factor (Rpf) is a growth factor that stimulates the growth of aged *M. tuberculosis* cultures, and members of this family are thought to play a role in reactivation of dormant *M. tuberculosis* (Mukamolova et al., 2002). Regulation of *rpfA* by CRP_{Mt} suggests that CRP_{Mt} plays a role in persistence and/or reactivation of tuberculosis, but this is only one of many biological functions that may be regulated by CRP_{Mt} in *M. tuberculosis*.

Expression of *M. tuberculosis whiB1*, a member of the Wbl (WhiB-like) family, is controlled positively and negatively by CRP_{Mt} (Stapleton et al., 2010; Agarwal et al., 2006). A CRP_{Mt} binding site (CRP_{Mt}1) was detected in *whiB1*'s upstream regulatory region (Smith et al., 2010) (Rickman et al., 2005). Reporter assays with native and mutated promoter sequences indicated that transcription from the native, but not the mutant, promoter was affected by cAMP levels via the direct binding with CRP_{Mt} (Agarwal et al., 2006). Other evidences indicate that a second CRP_{Mt} binding site (CRP_{Mt}2) in the *whiB1* promoter altered CRP_{Mt}'s

Associated function	Gene	Predicted function
Cell-wall biogenesis	<i>Rv0993</i>	<i>galU</i> , uridine diphosphate-glucose pyrophosphorylase
	<i>Rv3031</i>	Members of the family of enzymes transferring activated sugars
	<i>Rv3032</i>	
	<i>Rv0643c</i>	<i>mma3</i> , methoxy mycolic acid synthase
	<i>Rv0904c</i>	AccD3, a putative acetyl CoA carboxylase carboxyl transferase, which catalyzes the initial steps of fatty acid and mycolic acid biosynthesis.
Central metabolism pathways	<i>Rv2918</i>	<i>glnD</i> , uridyl transferase
	<i>Rv0992c</i>	CHP with putative 5-formyltetrahydrofolate cyclo-ligase
	<i>Rv0520</i>	Methyl transferase believed to be involved in ubiquinone pathway
	<i>Rv3113</i>	Phosphatase
	<i>Rv3114</i>	Nucleoside deaminase
	<i>Rv3505</i>	FadE27, protein possibly involved in regulating probable acyl-CoA dehydrogenase
	<i>Rv3617</i>	EphA, putative epoxide hydrolase
Putative regulatory elements controlling cAMP signaling	<i>Rv0104</i>	Hypothetical protein probably implied in cAMP mediated signaling in <i>M. tuberculosis</i>
	<i>Rv0103c</i>	Probable cation transporter
	<i>Rv3645</i>	Membrane linked adenyl cyclase
Antibiotic resistance	<i>Rv0906</i>	Hypothetical proteins belonging to the β -lactamase family
	<i>Rv0907</i>	
	<i>Rv0905</i>	<i>echA6</i> , enoyl-CoA hydratase
	<i>Rv0908</i>	CtpE, methyl-accepting chemotaxis protein

Table 2. *Mycobacterium* genes belonging to CRP regulon and their associated function

regulatory effect inhibiting *whiB1* expression (Stapleton et al., 2010). Interestingly, only CRP_{Mt}1 enhances *whiB1* expression, while CRP_{Mt}2 alone represses *whiB1* expression. Dnase I footprinting assays allowed to determine the presence at -58.5 of an activating CRP_{Mt} binding site (CRP_{Mt}1), which matches in seven of the eight nucleotides from the proposed CRP_{Mt} consensus (NGTGNNANNNNCACA), and also it overlapped with the repressing CRP_{Mt} binding site (CRP_{Mt}2) centred at -37.5. This second site has a poorer match to the consensus (six of the eight defined bases are matched)(Stapleton et al., 2010, Rickman et al., 2005). CRP_{Mt}1 site was occupied before the CRP_{Mt}2 site by a titration test done with increasing concentrations of CRP_{Mt}. It is worth noting that although it was shown that cAMP enhanced binding of recombinant CRP_{Mt} to target DNA, this enhancement was not equivalent to that observed for *E. coli* CRP, where DNA-binding affinity is enhanced by several orders of magnitude in the presence of 0.1 mM cAMP, allowing specific DNA binding at nanomolar concentrations. For CRP_{Mt} a much less significant enhancement of DNA binding was observed, and higher concentrations of cAMP compared with *E.coli* CRP were required (Stapleton et al., 2010).

On the other hand, out of 92 genes found and predicted by bioinformatics to be regulated by CRP in *M. tuberculosis*, only 18 [including *echA6*, *ctpE*, *accD3*, *sucC*, *sucD*, *glnD*, *fadE9*, *fbpC1* (Ag85C) and *galU*] have conserved orthologous in *M. leprae*, *M. avium* subsp. *paratuberculosis*, and *M. smegmatis* (Akhter et al., 2008). Incorporation of a plasmid harboring and expressing *M. tuberculosis* Rv3676 in an *M. tuberculosis* strain in which Rv3676 was absent, induced differential expression of 27 genes when compared to the same mutant harboring the BCG gene orthologous to Rv3676 (Hunt et al., 2008). This constitutes evidence that differences in gene regulatory sequences or the regulators among species exist, and it could explain the number of genes that do not respond in a similar manner. Further characterization of the *M. tuberculosis* CRP regulon as well as analysis of what physiological conditions regulate activation/inactivation are required.

3. Participation of cAMP in mycobacterial pathogenic processes

Little is known about the role of cAMP in mycobacteria, although it is found in both pathogenic and non-pathogenic species. Ingestion of live *M. microti* or *M. bovis* BCG (but not *M. lepraemurium*) increased macrophage intracellular cAMP levels, whereas no change occurred in cells engulfing dead bacilli, latex beads or colloidal gold (Lowrie et al., 1979). The rise in cAMP levels appears directly related to mycobacterial capacity to interfere with phagolysosome formation, evidence suggesting that these microorganisms modify cAMP-dependent signalling pathways as a manner to control virulence and infection (Lowrie et al., 1975, Lowrie et al., 1979). Elevated cAMP levels were correlated with reduced phagolysosome fusion during mycobacterial infection of macrophages (Kalamidas et al., 2006). Increased cAMP levels inside phagocytes were shown to negatively modulate actin-dependent processes, including chemotactic movement and phagocytosis. Macrophage passage was found to have a stimulatory effect on cAMP production by mycobacteria, as cAMP levels within macrophage-passaged mycobacteria were ~50-fold higher than cAMP levels within bacteria incubated in the tissue culture medium alone (Bai et al., 2009).

Evidences exist that cAMP regulates gene expression in mycobacteria during bacterial growth *in vitro* (Stapleton et al., 2010, Dass et al., 2008), and during macrophage infection (Rickman et al., 2005) where some studies identified Cmr as a transcription factor that regulates cAIGs (cAMP-induced genes) within macrophages, and suggest that multiple factors affect cAMP-associated gene regulation in tuberculosis-complex mycobacteria (Gazdik et al., 2009). Even during phagocytosis, expression may be down regulated in response to high cAMP or NO levels inside the macrophage environment, providing a mechanism to integrate the transcriptional response to two important signals associated with infection (Smith et al., 2010). Generally, increases in cAMP levels compromise the bactericidal activity of the host immune system.

It is likely that each cyclase is associated with a distinct signaling pathway. It is expected that specific cyclases are activated to modify cAMP levels in response to different physiological conditions as for example hypoxia, intramacrophage environment or pH changes. cAMP receptor protein (CRP) Rv3676 was found to exist as dimer and exhibited cAMP binding in a concentration-dependent manner and could bind specifically to the putative CRP/FNR nucleotide sequence elements in response to hypoxia (Akhter et al., 2008). The protein itself is composed of three distinct regions of the polypeptide: a large N-terminal domain that binds cAMP, a long α -helix (termed the C-helix) that mediates most of

Gene	Induction	Repression
<i>Rv0386</i>	Palmitic acid Tetracyclin	7H9 medium shaking Streptomycin Purified surfactant lipids
<i>Rv0891c</i>	Oleic acid	DETA/NO
<i>Rv1120c</i>	Pulmonary surfactant	Non-replicative persistence
<i>Rv1264</i>	Carbonyl cyanide chlorophenylhydrazone	Palmitic acid DETA/NO
<i>Rv1318c</i>	Oligopeptide permease (<i>Rv3662c-Rv3665c</i>) mutant Palmitic acid	Pulmonary surfactant protein A (human) Acetate
<i>Rv1319c</i>	Hypoxia	Arachidonic acid Hydrogen peroxide Thioridazine Nicotinamide
<i>Rv1320c</i>	Acetate	
<i>Rv1359</i>	Econazole Macrophages Iron	Arachidonic acid Palmitic acid
<i>Rv1625c</i>	Clofazimine + S-nitrosoglutathione Arachidonic acid Thioridazine	Palmitic acid Oleic acid
<i>Rv1647</i>	Linoleic acid Palmitic acid Oleic acid	Streptomycin Arachidonic acid DETA/NO
<i>Rv1900c</i>	Non-replicative persistence PA 824 Palmitic acid	Tetracycline S-nitrosoglutathione + Chlorpromazine Acetate
<i>Rv2212</i>	Tetracycline	5-chloro-pyrazinamide
<i>Rv2345</i>	Starvation	Thioridazine DETA/NO
<i>Rv2488c</i>	Arachidonic acid Acetate	Tetracycline Palmitic acid Hydrogen peroxide
<i>Rv3645</i>	Oleic acid	Oligopeptide permease (<i>Rv3662c-Rv3665c</i>) mutant Acetate Ceramide

Table 3. Growth conditions affecting expression of mycobacterial cyclase domain containing genes

the inter monomer interactions, and a small C-terminal DNA-binding domain. It is also capable of binding to two, three, or four cAMP molecules, but the specificity of recognition sequentially diminishes beyond two cAMP molecules bound to CRP, although the physiological importance of these molecular interactions are not yet known (Stapleton et al., 2010). Data available at Tuberculosis Database (<http://www.tbdb.org/>, Table 3) might help suggesting particular conditions where each AC gene may be required.

Currently, few studies have addressed the participation of cAMP signalling in *M. tuberculosis* pathogenesis *in vivo*. Rickman et al. (Rickman et al., 2005) found that a mutant in Rv3676 (CRP) had diminished bacterial burden in lung and spleen after intravenous infection of BALB/c mice, compared to wild type bacteria. On the other hand, Agarwal et al. (Agarwal et al., 2009) found that a Rv0386 *M. tuberculosis* mutant was affected in its capacity to replicate in BALB/c or C57BL/6 mice lungs, following aerosol infection. Neither publication mentioned the capacity of either strain to kill mice.

In addition to determining bacterial burden in infected lungs Agarwal et al. (Agarwal et al., 2009) demonstrated that cAMP produced by *M. tuberculosis* Rv0386 during J774.16 macrophage infection was a substrate for protein kinase A, in order to control the amount of phosphorylated CREB (CREB-P), by using specific chemical inhibitors of selected signalling transduction pathways. They found CREB-P induced TNF- α production, and led to an unregulated host inflammatory response, which favoured bacterial survival. To date, this is the first indication of how adenylyl cyclase action helps sustaining an infection by pathogenic mycobacteria.

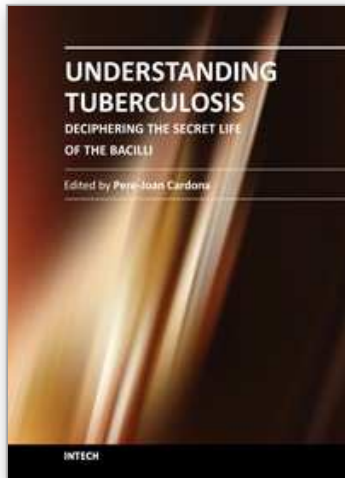
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Mycobacterium tuberculosis, as recent investigations demonstrate, has a complex signaling expression, which allows its close interaction with the environment and one of its most renowned properties: the ability to persist for long periods of time under a non-replicative status. Although this skill is well characterized in other bacteria, the intrinsically very slow growth rate of *Mycobium tuberculosis*, together with a very thick and complex cell wall, makes this pathogen specially adapted to the stress that could be generated by the host against them. In this book, different aspects of these properties are displayed by specialists in the field.

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