Non-coding RNA and its potential role in *Mycobacterium tuberculosis* pathogenesis

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**Abbreviations:** sRNA, small RNA; RNA-seq, RNA sequencing; MTS, *M. tuberculosis* sRNA; RLM-RACE, RNA ligase-mediated RACE; PE, proline-glutamate; PPE, proline-proline-glutamate; UTR, untranslated region; CRISPR, clustered regularly interspaced short palindromic repeat

It is estimated that one third of the human population is infected with *Mycobacterium tuberculosis*. Efforts to understand the molecular basis of its gene regulation have been focused on identification of protein encoding genes and regulons implicated in pathogenesis. Recently, a number of studies have described the identification of several non-coding RNAs that are likely to contribute significantly to the regulatory networks responsible for adaptation and virulence in *M. tuberculosis*. We have reviewed emerging information on the presence and abundance of different types of non-coding RNA in *M. tuberculosis* and consider their potential contribution to the adaptive responses that underlie disease pathogenesis.

**Introduction**

*Mycobacterium tuberculosis* is regarded as one of the most successful human pathogens. It is responsible for more deaths than any other microorganism, and moreover, one third of the global population is showing immunological evidence of past or current infection. This occurs predominantly in the form of an asymptomatic latent infection, resembling carriage of a commensal organism, and only a minority of individuals develop the progressive lung disease that is necessary for maintenance of the transmission cycle. In contrast to well-characterized enteric pathogens that rely on the acquisition and expression of specialized virulence factors, pathogenesis of *M. tuberculosis* depends on an ability to adjust to changing environments within the host and, under appropriate circumstance, to transmit signals that subvert the immune response in order to cause localized immunopathology.

Efforts to understand the molecular basis of mycobacterial pathogenesis, with the aim of identifying interventions that prevent or resolve disease, focus on understanding of the ability of the bacteria to modulate their metabolism and replication in response to different environmental signals. The 4.4 megabase genome of *M. tuberculosis* encodes 13 sigma factors and more than 100 annotated transcriptional regulators, and an extensive literature documents transcriptional changes in response to environmental stresses imposed in vitro, during macrophage infection, and in human disease (reviewed in ref. 3–5).

In this article, we will review emerging information on the presence of non-coding RNA in *M. tuberculosis* and consider its potential contribution to the adaptive responses that underlie disease pathogenesis and persistence.

**Different Types of Non-coding RNA in Bacteria**

Application of high-density tiling arrays and RNA sequencing technologies (RNA-seq) have uncovered an extensive and previously unknown repertoire of non-coding RNA in bacteria, including 5’ and 3’ untranslated regions (UTRs), antisense transcripts and intergenic small RNAs (sRNAs). This is also the case in *M. tuberculosis*.8

The 5’ UTR, refers to the region between the transcriptional and the translational start sites. This part of the transcript can in some cases change its conformation in response to external stimuli such as changes in temperature or the availability of certain metabolites, in which case the RNA element is referred to as a riboswitch.10 The change in conformation in turn leads to a change in the expression of the downstream gene either by blocking/unblocking of the ribosome binding site or by transcriptional termination/antitermination.11 An intriguing finding is that the riboswitch moiety of the SAM-IV riboswitch from *Listeria monocytogenes* can act in trans as an sRNA thereby adding further complexity to the regulation by riboswitches.12 Whether this applies to other riboswitches or perhaps attenuated transcripts in general remains to be determined.

sRNAs are, as the name indicates small transcripts, mostly in the range of 50 to 250 nucleotides, and they can be encoded opposite open reading frames (antisense or cis-encoded) or between open reading frames (intergenic or trans-encoded).

The majority of sRNAs regulate gene expression by base-pairing to one or more target mRNAs thereby modifying translation efficiency and/or mRNA stability.13,14 One widespread exception is 6S RNA, which binds to RNA polymerase thereby specifically downregulating the usage of σ70 promoters.15 Other examples of protein binding sRNAs include CsrB and CsrC, which both
bind to the carbon storage regulatory protein, CsrA (reviewed in ref. 16).

Although sRNAs are generally referred to as regulatory and/or non-coding RNAs, a few have been found to encode small peptides as well as acting on RNA level. Other small intergenic transcripts appear to be acting purely as templates for the synthesis of small peptides.18,19

The RNA Chaperone Hfq

In Gram negative bacteria, including many pathogens the RNA chaperone Hfq is required to facilitate the interaction between trans-encoded sRNAs and their targets.

Deletion of the hfq gene often leads to loss of virulence, which may in part be due to loss of proper sRNA function.20,21 However, the role of Hfq in the Gram positive Staphylococcus aureus is more controversial,22,23 and the protein is completely absent in several pathogens including M. tuberculosis.20 This lack of Hfq represents a limitation in extrapolation of experimental approaches as well as sRNA mechanisms from other bacterial systems both in terms of sRNA-mRNA interaction, but also in terms of the subsequent fate of transcripts via engagement with the degradosome, which is mediated by Hfq.24

The obvious question is, whether M. tuberculosis sRNAs can and do interact with their targets unaided or whether in M. tuberculosis there is one or more alternative, hitherto unidentified chaperones. The high GC content (67%) as well as the structure of intrinsic terminators in M. tuberculosis (see below), limits the frequency of AU-rich stretches implicated in conventional Hfq interactions.25 Nevertheless, an alternative chaperone could be considered an appropriate Hfq analog if it were to display a broad, albeit different sequence specificity toward the majority of sRNAs and their targets. One candidate for this type of chaperone is Rv2367. This protein is a homolog of the Sinorhizobium meliloti YbeY protein that has been shown to display certain Hfq functions,26 although the low level expression of Rv2367 contrasts with the abundance of Hfq in Escherichia coli.8,27

Alternatively (or in parallel) one could imagine a range of regulon-specific chaperones, each of which could interact with a subset of sRNAs and their targets in a manner similar to the FbpABC proteins in Bacillus subtilis.28 Finally there is the possibility that sRNA function in M. tuberculosis is independent of chaperones. But what are the pros and cons of riboregulation without RNA chaperones? On one hand the nucleotide composition of M. tuberculosis sRNAs will in all likelihood lead to structures that have a significantly higher degree of intramolecular stability, which are more difficult to disrupt and these would therefore impair the interaction with target RNAs. On the other hand it is also possible that a higher proportion of G and C nucleotides in the seed sequence will facilitate the interaction between sRNA and target. Supporting this notion is the fact that some M. tuberculosis sRNAs are predicted to harbour unpaired C-rich stretches that would make for strong first interactions.29

Future investigations, in which tagged sRNAs are used as bait for the isolation of RNA-binding proteins and hence putative chaperones30 will hopefully shed more light on these questions in near future.

sRNAs and Pathogenesis

While functional characterization of intergenic sRNAs has only recently been initiated in M. tuberculosis, numerous sRNAs have been characterized in other pathogens. Many of these sRNAs are induced by stress and hence associated with host adaptation and virulence.6,21,31 In some cases sRNAs enhance their effect by regulating regulators such as the Qrr1-4 sRNAs, which regulate HapR in Vibrio cholerae,32 or ArcZ, which regulates the expression of RpoS in both E. coli and Salmonella.33,34 Other sRNAs that are intimately related to pathogenesis and survival within the host are RybB and MicA, which upon acid stress promote the degradation of several outer membrane protein (OMP) mRNAs in Salmonella. This leads to a rapid rearrangement of the outer membrane, and hence the host-pathogen interface until a new homeostasis has been achieved.35 In spite of the close relationship between stress, host adaptation and pathogenesis, only a fraction of sRNAs has been shown to directly affect virulence in different infection models.21 These include the S. aureus Sprd,36 L. monocytogenes Rli38,37 Legionella pneumophilia srs (encoding 6S RNA),38 and the three Salmonella sRNAs IstM,39 IstR and SroA;40 all of these sRNAs result in impaired virulence upon deletion. Deletion of L. monocytogenes RliB and V. cholerae VrrA lead to increased colonization of spleen and intestines, respectively.37,41

Coding vs. Non-coding RNA in M. tuberculosis

The identification of putative regulatory sRNAs in M. tuberculosis is so far limited to three studies.8,25,62 Figure 1 presents an overview of some of the putative RNA-based regulatory networks in M. tuberculosis.

The most recent study is a detailed quantitative analysis of M. tuberculosis transcriptomes from exponential and stationary phases using RNA-seq.8 After removal of signals from rRNAs, 17% of the reads identified from exponentially growing M. tuberculosis H37Rv mapped to intergenic regions, with a further 12% mapping in antisense orientation to coding sequences. Non-coding RNA increased to 58% of total (non-rRNA) reads in stationary phase cultures, largely due to accumulation of a single, highly abundant sRNA transcript, MTS2823 (see below). It is likely that these numbers represent an underestimate of the total non-coding transcriptome, since the protocol used in this study was not optimised for detection of shorter transcripts (< 200 nucleotides), which was reflected by the relatively low ratio of 5S to the larger 16S and 23S rRNA species.8

Not all intergenic transcripts will be non-coding and the distinction between coding and non-coding transcripts is contingent on the methods employed to predict protein-coding transcripts; the outcome varies according to different algorithms...
that have been used to generate multiple annotations of *M. tuberculosis* genomes. Identical short open reading frames are more frequently annotated as coding sequences in *M. tuberculosis* CDC1551 \(^\text{43,44}\) than in the closely related *M. tuberculosis* H37Rv.\(^2\) For example, transcription of the sequence annotated as an intergenic region between Rv2395 and Rv2396 in H37Rv has been described as an sRNA (mcr7),\(^\text{42}\) but it is annotated as two hypothetical proteins, MT2466 and MT2467 in the CDC1551 genome, and was recently reported to encode two small proteins involved in the mycobacterial response to low pH.\(^\text{49}\) Conversely, an abundant transcript encoded between Rv3661 and Rv3662c has been alternatively annotated as sRNA MTS2823,\(^8\) and as hypothetical protein MT3762.\(^\text{44}\) In this case, the inverse orientation of the transcript with respect to the predicted protein strongly favors the sRNA annotation. Moreover, it is likely that some transcripts will have dual function as both sRNA and mRNA as it has been found in other bacteria.\(^7\) Initial predictions of translation start sites are also likely to be subject to revision in light of experimental data, and this will result in reassignment of coding and non-coding portions of mRNA transcripts. Application of mass spectrometry technology for proteome analysis, e.g., reference 46 and 47, will play an important role in resolving these ambiguities.

**Antisense RNAs**

Bacterial gene regulation by short antisense transcripts that base-pair with 5’ regions of mRNAs has been well characterized for a subset of bacterial genes, including several involved in transposition of foreign genetic elements.\(^\text{48}\) A surprising finding from several studies using RNA-seq is the extent and heterologous nature of antisense transcriptomes. Thus, an antisense component has been identified for up to 75% of all protein-coding genes, with transcripts ranging in size from less than 50 nucleotides to several kilobases and mapping to sequences throughout the length of cognate mRNAs.\(^\text{49}\) A similar pattern is observed for *M. tuberculosis*, where 65% of genes have an antisense component corresponding to ≥ 10% of the coding transcript during exponential growth; a number that increases to > 90% in stationary phase.\(^8\) Antisense transcripts have been shown to regulate transcription and translation of coding transcripts by a variety of mechanisms.\(^\text{48}\) Recently, RNase III-mediated digestion of double stranded sequences has been proposed as an attractive model by which antisense transcription could play a pervasive genome-wide role in fine-tuning of expression patterns.\(^\text{49}\)

During exponential growth in *M. tuberculosis*, all gene classes have some level of antisense RNA, but the gene classes with the lowest occurrence of antisense, are also the ones that are the most highly expressed, i.e., energy metabolism and synthesis of macromolecules.\(^8\) Several *M. tuberculosis* antisense transcripts have been identified by cloning and sequencing from low molecular weight RNA fractions or by RNA-seq, and their sizes as well as locations relative to their cognate open reading frames (ORFs) vary significantly; some are encoded at the 5’ end of the ORF, some in the center and some at the 3’ end, and a few cover an entire ORF or more.\(^\text{8,29,42}\) Most of the *M. tuberculosis* antisense RNAs appear to be independent transcripts, but a few are derived from long, overlapping 3' UTRs (see below).\(^8\)

PhoPR is a pathogenesis-associated two-component system that may be subject to riboregulation. There is a prominent internal antisense transcript toward the 5’ end of the *phoP* gene,

**Figure 1.** Overview of the potential roles of regulatory RNA in *M. tuberculosis*. The figure illustrates how different external signals, shown as orange arrows, can regulate either riboswitches, antisense or intergenic RNAs (italics). Known protein regulators, i.e., DosR and SigF as well as selected sRNAs have been indicated with names. Blue lines indicate that the resulting regulation can be either activation or inhibition; full lines indicate known activation/inhibition; dashed arrows indicate possible consequence.

**Figure 2.** Expression of the *ino1* antisense RNA is significantly downregulated in stationary phase. The figure illustrates RNA-seq data visualized with the Artemis genome browser; reads mapping to the plus strand, i.e., antisense to *ino1*, are shown in red, and reads mapping to the minus strand, i.e., *ino1* itself, are shown in blue. Expression levels in the two phases have been adjusted for comparative reasons.
and the 3' UTR of Rv0795c generates an antisense transcript that extends over the 3' end of the converging phoR and the 3' UTR of Rv0795c generates an antisense transcript mRNA, DesA2 (Rv1094). The second example, ASpks, is for base-pairing with the mRNA of a second homologous target and has the potential to the increased ASpks levels. During oxidative stress, and to a lesser extent at low pH, an extended 200-nucleotide ASpks transcript is induced. A subset of M. tuberculosis mRNAs have well-defined 3' ends associated with a recently identified mycobacterial terminator, TRIT, but the general paucity of L-shaped intrinsic terminators—i.e., a stem-loop followed by a stretch of U residues—frustrates bioinformatics approaches to the prediction of 3' ends in M. tuberculosis.

Riboswitches

A number of riboswitches have been identified in M. tuberculosis by sequence homology; one of these, the cobalamin riboswitch has been subject to further investigation. This type of riboswitch represses the expression of the downstream gene in the presence of their cognate mRNA partner as well as coordinated transcriptional regulation. 8

Figure 3. M. tuberculosis riboswitches and PE-PPE genes; the figure illustrates how riboswitches in M. tuberculosis are sometimes separated from their predicted cognate mRNAs (shown in blue), by the insertion of one or more PE-PPE genes (shown in red). Due to the sequence conservation inherent in these riboswitches, the regions have been annotated as 'conserved hypotheticals' (shown in green). Other genes are shown in gray. The top part shows how the B12 riboswitch is separated from cobQ1 and cobU genes by a PPE2 insertion. The bottom part shows how the Mbox has been separated from mgtC by several PE-PPE genes.
of cobalamin (vitamin B12) and is often found upstream of genes involved in the synthesis and/or transport of cobalamin. There are two copies of this motif in M. tuberculosis; one is upstream of the metE gene, encoding a B12-independent methionine synthase, and the second is upstream of an operon comprising PPE2, cobQ1 and cobU (Fig. 3). PPE2 belongs to an extensive family of proteins sharing proline-glutamate (PE) or proline-proline-glutamate (PPE) N-terminal motifs that were identified from the M. tuberculosis genome sequence but for the most part lack any obvious biological function. The genomic location of PPE2, together with the presence of several transmembrane domains has led to the suggestion that it may be a coaptor transporter. Another riboswitch motif that is represented twice in the genome of M. tuberculosis but which has not been functionally characterized, is the ykok leader or Mbox, which is usually involved in the synthesis and/or transport of cobalamin. There are two copies of this motif in M. tuberculosis; one is upstream of the metE gene, encoding a B12-independent methionine synthase, and the second is upstream of an operon comprising PPE2, cobQ1 and cobU (Fig. 3). PPE2 belongs to an extensive family of proteins sharing proline-glutamate (PE) or proline-proline-glutamate (PPE) N-terminal motifs that were identified from the M. tuberculosis genome sequence but for the most part lack any obvious biological function. The genomic location of PPE2, together with the presence of several transmembrane domains has led to the suggestion that it may be a coaptor transporter. Another riboswitch motif that is represented twice in the genome of M. tuberculosis but which has not been functionally characterized, is the ykok leader or Mbox, which is usually found in the context of magnesium transporters. The Bacillus subtilis Mbox has been shown to attenuate transcription upon binding Mg2+, thereby ensuring magnesium homeostasis. In M. tuberculosis one highly expressed Mbox is found upstream of Rv1535, a conserved hypothetical protein that is induced during Mg-starvation. Rv1535 is followed by and co-transcribed with a downstream Tbox (a riboswitch that binds tRNAs) that forms the 5’ UTR of the essential isoleucine tRNA synthase mRNA (IleS, Rv1536), (Arnvig KB, Young DB unpublished). The second Mbox is located upstream of a putative operon encoding four PE-PPE genes, a conserved hypothetical protein and a predicted magnesium transporter, MgtC (Fig. 3), all of which are induced by magnesium starvation. The fact that the genes downstream of the two Mboxes in M. tuberculosis are induced upon magnesium starvation, strongly suggest that these are in fact functional magnesium-responsive RNA elements, which are activated by the low concentration of magnesium found in macrophages.

Table 1. Selected intergenic sRNAs, verified by northern blotting

| sRNA     | Alternative name | Strand | Left CDS       | Right CDS      | Size* | Regulation**          | Ref. |
|----------|------------------|--------|----------------|----------------|-------|------------------------|-----|
| MTS0194  | F6               | F      | Rv0243         | Rv0244c        | 55, 110 | SigF, starvation, H2O2, low pH | 29  |
| MTS0479  | B55              | F      | Rv0609A        | Rv0610c        | 60    | H2O2                   | 29  |
| MTS0823  | mpr5             | F      | Rv1051c        | Rv1052         | 120   | -                       | 42  |
| MTS0858  |                 | R      | Rv1092c        | Rv1093         | 100   | -                       | 8   |
| MTS0997  | mcr11            | R      | Rv1264         | Rv1265         | 120   | CRP, stationary phase, infection | 8, 42 |
| MTS1082  | Rv1373           | F      | Rv1375         | 130            | -     | -                       | 8   |
| MTS1310  | G2               | R      | Rv1689         | Rv1690         | 65    | Exponential phase       | 29  |
| MTS1338  | F173c            | F      | Rv1734c        | 120            | -     | DosR, hypoxia, infection | 8   |
| MTS2774  | mpr17            | F      | Rv3596c        | Rv3597c        | 80, 110 | -                       | 42  |
| MTS2822  | B11              | R      | Rv3660c        | Rv3661         | 95    | H2O2                   | 28  |
| MTS2823  | mpr4             | F      | Rv3661         | Rv3662c        | 305   | stationary phase, infection, low pH | 8, 42 |
| MTS2975  |                 | F      | Rv3843c        | Rv3844         | 100   | Exponential phase       | 8   |

Notes: *Approximate size (nucleotides) of the dominant transcripts as determined by northern blotting; **Regulation refers to relevant factors and conditions where increased expression is observed.

Another riboswitch motif that is represented twice in the genome of M. tuberculosis but which has not been functionally characterized, is the ykok leader or Mbox, which is usually found in the context of magnesium transporters. The Bacillus subtilis Mbox has been shown to attenuate transcription upon binding Mg2+, thereby ensuring magnesium homeostasis. In M. tuberculosis one highly expressed Mbox is found upstream of Rv1535, a conserved hypothetical protein that is induced during Mg-starvation. Rv1535 is followed by and co-transcribed with a downstream Tbox (a riboswitch that binds tRNAs) that forms the 5’ UTR of the essential isoleucine tRNA synthase mRNA (IleS, Rv1536), (Arnvig KB, Young DB unpublished). The second Mbox is located upstream of a putative operon encoding four PE-PPE genes, a conserved hypothetical protein and a predicted magnesium transporter, MgtC (Fig. 3), all of which are induced by magnesium starvation. The fact that the genes downstream of the two Mboxes in M. tuberculosis are induced upon magnesium starvation, strongly suggest that these are in fact functional magnesium-responsive RNA elements, which are activated by the low concentration of magnesium found in macrophages.

Inspection of M. tuberculosis RNA-seq profiles identifies numerous additional candidate riboswitches expressed as truncated 5’ transcripts (Arnvig KB, et al. unpublished). One of these is located in the 5’ UTR of Rv0282, the first gene in the ESX3 operon that is essential for virulence and regulated at the level of transcription initiation by iron and zinc. In addition to its transcriptional attenuation profile the 5’ UTR contains a highly conserved region, which could constitute a ligand-binding aptamer domain. It is intriguing to note that once again this putative riboswitch is located upstream of an operon containing a PE-PPE gene pair, extending a pattern of PE-PPE gene insertion between riboswitches and their predicted regulated targets. A YdaO riboswitch (candidate) is present upstream of rpfA (Rv0867c) in M. tuberculosis. No ligand has been identified for the YdaO riboswitch though it is often associated with genes involved in processes related to cell wall degradation.

These examples suggest that riboswitches have a widespread role in regulation of M. tuberculosis gene expression and growth.

M. tuberculosis Intergenic sRNAs and Stress

Around 20 M. tuberculosis intergenic sRNAs have been identified and verified by northern blotting. They display varying degrees of conservation; some are restricted to closely related members of the M. tuberculosis complex, others are present in multiple pathogenic mycobacteria with Mycobacterium leprae being the most distant relative, and finally some are conserved in all mycobacteria and even other actinomycetes. The M. tuberculosis sRNAs range in size from ~50 to > 300 nucleotides, they are often differentially expressed in response to changing conditions.

Notes: *Approximate size (nucleotides) of the dominant transcripts as determined by northern blotting; **Regulation refers to relevant factors and conditions where increased expression is observed.
environments, and a couple have been shown to be lethal upon overexpression. Selected examples are listed in Table 1 using an MTS (Mycobacterium tuberculosis sRNA) designation followed by a number corresponding to the intergenic region i.e., the TIGR website annotation of intergenic regions (cmr.jcvi.org/tigr-scripts/CMR/CmrHomePage.cgi).

The differential expression of sRNAs upon changing growth conditions may give some clues about function in terms of under what conditions a given sRNA may be required, although it will not reveal the entire story.

MTS194, MTS479 and MTS2822. Oxidative stress is a potent inducer of stress responses in M. tuberculosis. Three of the sRNAs listed in Table 1, namely MTS194 (“F6”29), MTS479 (“B55”29) and MTS2822 (“B11”29) show increased expression in response to H2O2, which mimics the oxidative stress encountered inside the host macrophage, and hence these sRNAs could be associated with intracellular survival during the early stages of infection. MTS194 is encoded in the intergenic region between two genes involved in lipid degradation, Rv0243 and Rv0244c. It is transcribed by the starvation associated alternative sigma factor, SigF and the MTS194 promoter has recently been shown to represent the highest occupancy of this sigma factor suggesting that expression of this sRNA has high priority under certain conditions such as starvation. Overexpression of MTS194 leads to reduced growth in M. tuberculosis, but although an MTS194 homolog is present in the distantly related, non-pathogenic Mycobacterium smegmatis, overexpression of MTS194 in M. smegmatis has no obvious phenotype.29

MTS2822. MTS2822 is ~95 nucleotides in size and located in the intergenic region between Rv3660c and Rv3661, both predicted to be involved in cellular differentiation. Downstream of Rv3661 is another sRNA, MTS2823 (see below), and the entire locus is highly conserved among mycobacteria, with the notable exception of M. leprae. MTS2822 contains a so-called 6C motif, consisting of two C rich loops found to be widespread between Actinobacteria.77 The structure of MTS2822 has led to the suggestion that this sRNA may be a structural/protein binding RNA rather than a conventional regulatory sRNA.77 M. smegmatis harbours an sRNA that is ~90% identical to MTS2822 and while overexpression of MTS2822 in M. tuberculosis is lethal, overexpression of MTS2822 in M. smegmatis results in a dramatic phenotype; growth is very poor on solid and in liquid media, and the cells display aberrant morphology including irregular and elongated shape, suggesting association with cell wall synthesis and/or cell division.29

MTS997, MTS1338 and MTS2823. Three other sRNAs, MTS997, MTS1338 and MTS2823 accumulate to high levels during the transition from exponential to stationary phase, (representing a mixture of stresses), and to even higher levels during infection, suggesting a role in pathogenesis. Generally, the roles of individual sRNAs within a cell’s regulatory networks vary substantially. Well-characterized sRNAs can be, and in the case of Salmonella and E. coli have been, assigned to expressing this sRNA may be in a different metabolic state than the remaining population and therefore they could represent putative persister cells. Moreover, the induction of MTS1338 seen in stationary phase is almost eliminated by deletion of dormancy regulator, DosR.8 DosR interacts with DosS or DosT to form a phosphorylation-dependent two-component regulatory system activated during growth arrest induced by hypoxia or exposure to nitric oxide.79 The DosR regulon includes genes required for remodeling of protein, lipid and energy metabolism together with members of the universal stress protein family and multiple proteins of unknown function.3 The DosR genes are strongly expressed during infection in mice and in sputum samples from patients, and include several prominent antigens, and their expression has been linked to the generation of drug tolerant persister cells.3,80,81 It remains to be determined if MTS1338 is directly involved in the generation of persister cells or simply a marker for this subpopulation.

The most in-depth functional analysis has been performed on the relatively large (>300-nucleotide) MTS2823 sRNA, which is encoded on the plus strand between Rv3661 and Rv3662c. It is the most highly expressed non-rRNA in exponentially growing cultures, is induced a further 10-fold in stationary phase, and accumulates to a level approaching 1:1 stoichiometry with rRNA in tissues from infected mice. MTS2823 is conserved in a broad range of mycobacteria (and other actinomycetes).

Constitutive overexpression of MTS2823 in exponential phase cultures results in a slightly reduced growth rate and in a widespread downregulation of energy metabolism genes, analogous to that observed during transition from exponential growth to stationary phase. The most marked reduction is seen for genes associated with the methyl citrate cycle in particular prpC (methyl citrate synthase, >100 fold) and pepD (methyl citrate dehydratase, >60 fold). This metabolic pathway contributes to detoxification of metabolites generated during catabolism of odd-numbered fatty acids and cholesterol, which has been shown to be an essential carbon source during intracellular growth.82,83

Downregulation of active growth functions during stationary phase transition in bacterial cultures is commonly associated with increased expression of the structurally conserved 6S RNA molecule, that binds to RNA polymerase and inhibits transcription of genes controlled by the principal sigma factor.15,84,85 A bioinformatics search based on matching for energetically sub-optimal structures recently identified the M. smegmatis homolog of MTS2823 as a potential 6S RNA candidate, but lack of binding to RNA polymerase led the authors to conclude that it is not a genuine 6S RNA,86 and so far no 6S RNA homolog has been identified in mycobacteria. It remains to be determined if MTS2823 is a conventional sRNA that acts via base-pairing with specific mRNA targets or with protein or whether its mode of action differs entirely from other sRNAs.

M. tuberculosis sRNA Networks?
different types of regulatory circuits depending on the interplay between regulators, sRNAs and targets. However, in the case of *M. tuberculosis* this requires a significant increase in the amount of data about individual sRNAs that we are only just beginning to collect. Once the different regulators and targets of *M. tuberculosis* sRNA have been identified, monitoring the fate of target mRNAs upon deletion or overexpression of sRNAs will help to distinguish between different mechanistic models.

**CRISPR**

The Clustered Regularly Interspaced Short Palindromic Repeat (CRISPR) locus provides a specialized source for generation of *M. tuberculosis* sRNAs. The CRISPR locus incorporates sequences from phage and other invading genetic elements and repackages them as sRNA defense molecules that confer resistance to re-infection. CRISPRs are found in only a subset of mycobacteria, and the presence of the CRISPR in *Mycobacterium canetti* suggests acquisition by horizontal transfer prior to final branching of the *M. tuberculosis* complex. Sporadic deletion of integrated foreign sequence elements from the CRISPR domain has been exploited in a widely used system for differentiation of *M. tuberculosis* strains referred to as spoligotyping. Phylogenetically, the *M. tuberculosis* CRISPR system belongs to the Type III-A family, and is related to a well-characterized CRISPR system in *Staphylococcus epidermidis*. The Type III-A system is characterized by the presence of Cas10 (Rv2823c) and Csm6 (Rv2818) in the CRISPR-associated complex for antiviral defense (CASCADE), which in *M. tuberculosis* is likely to target foreign DNA rather than RNA. Inspection of RNA-seq profiles shows that the CRISPR domain is transcribed and processed into a series of 50–60 nucleotide sRNAs that are presumably incorporated into a CASCADE effector complex (Arnvig KB, et al. unpublished). The transcription start site driving CRISPR sRNA expression overlaps with a divergent start site that generates a long antisense transcript covering Rv2816c and Rv2817c encoding Cas1 and Cas2 proteins involved in incorporation of novel sequences into the CRISPR locus (Arnvig KB, et al. unpublished). This profile is consistent with silencing of the genes required to capture novel CRISPR sRNAs, but functional expression of Cas6 and Cas10 enzymes (i.e., Rv2824c and Rv2823c) required for processing of pre-existing sRNAs. An exception is the sub-lineage of *M. tuberculosis* known as the Beijing family, which is undergoing a current global expansion in marked association with multiple drug resistance. A genomic deletion has removed most of the CRISPR locus and many of the associated Cas genes, eliminating expression of CRISPR sRNAs in these strains. A similar deletion has been found in an unrelated sub-lineage referred to as “pseudo-Beijing.” It is open to speculation whether the presence of a functional CRISPR system in the majority of *M. tuberculosis* strains has current biological relevance or reflects fortuitous retention of a defense system appropriate to an ancestral lifestyle.

**Implications for Pathogenesis and Disease Control**

Progression from documentation of the non-coding transcriptome to elucidation of its biological function will require an extensive program of experimental research. Given the limitations of working with a slow-growing pathogen under strict containment condition—the timeframe of biologically relevant experiments using *M. tuberculosis* routinely extends from 6 to 12 mo—it may be important to explore fundamental mechanisms using more tractable organisms and to focus experimental work with *M. tuberculosis* on aspects of particular relevance to pathogenesis.

It can be envisaged that research on non-coding RNA in *M. tuberculosis* will contribute in three areas: improved understanding of the regulation of pathways already implicated in pathogenesis, characterization of *M. tuberculosis* phenotypes relevant to human infection, and hence, although more long-term, improved interventions for disease control.

**Improved understanding.** Over more than a decade since the sequencing of the *M. tuberculosis* genome, research in many laboratories has focused on identification of protein encoding genes and regulons implicated in pathogenesis; typically by generation of recombinant strains and subsequent testing of their phenotype in a mouse model of infection. Analysis by genome-wide transposon mutagenesis identified almost 200 protein encoding genes that are dispensable for growth in laboratory culture but are required for infection, and individual studies have characterized multiple genetic loci involved in control of defense against host antimicrobial systems, adaptation to changing nutritional and physicochemical constraints, and production of macromolecules that modulate interactions with host cells. RNA-seq profiling identifies non-coding RNA elements associated with many of these loci.

**Characterization of *M. tuberculosis* phenotypes.** While mouse infection provides a convenient model to analyze *M. tuberculosis* growth dynamics in the presence of host innate and adaptive immune cells, it does not recapitulate the more complex biological processes underlying the disease transmission cycle. In humans and non-human primates, infection with *M. tuberculosis* induces formation of a heterogeneous and dynamic range of highly structured granulomatous lesions that provide a multiplicity of bacterial microenvironments. Successful pathogenesis involves an ability to tolerate the resulting unfavorable environments, and to exploit environments that are favorable for bacterial multiplication, access to airways and onward transmission. It is likely that the mouse model interrogates only a portion of the phenotypic repertoire involved in human disease, and characterization of phenotypic states relevant to different stages of human pathology presents a major research challenge. Accumulation of sRNAs linked to different stress regulons may act as sensitive markers of the physiological status of the small numbers of bacteria present in different parts of human and non-human lesions, providing crucial information to support rational targeting of improved therapies.

**Improved interventions.** The success of treatment-based strategies for tuberculosis control is limited by the requirement for
therapeutic regimens extending over 6 months or more. Prolonged treatment is needed to clear a residual population of bacteria that persist in a phenotypic condition in which they are tolerant to antibacterial drugs. Identification of drugs that have a cidal effect against persister populations is a key research goal. Phenotypic tolerance can be reproduced in vitro by driving bacteria into a non-replicating state by limitation of oxygen or nutrients. In vitro persister models share the characteristic of a widespread downregulation of genes required for active growth, and it is likely that MTS2823 makes a contribution to this process. An alternative to the environment-induced model is that the persister phenotype is generated spontaneously in a few individuals within an apparently homogeneous bacterial population as a result of stochastic variations in gene expression. Mathematical modeling suggests that post-transcriptional regulation could provide an important source of such stochastic variation, and measurement of uninduced levels of an sRNA such as MTS1338 does indeed suggest occurrence at less than one copy per cell.

In summary the abundance of non-coding RNA in M. tuberculosis and its association with genes implicated in pathogenesis suggests that post-transcriptional regulation plays a significant role in the intracellular survival of this bacterium as it is the case for many other pathogens. Moreover, it is possible that riboregulation may also be involved in the generation of persister cells that are phenotypically tolerant to conventional antibacterial drugs. Future investigations will hopefully shed more light on this aspect of M. tuberculosis biology.

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