Cell-autonomous targeting of arabinogalactan by host immune factors inhibits mycobacterial growth

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Abstract

Deeper understanding of the crosstalk between host cells and *Mycobacterium tuberculosis* (Mtb) provides crucial guidelines for the rational design of novel intervention strategies against tuberculosis (TB). Mycobacteria possess a unique complex cell wall with arabinogalactan (AG) as critical component. AG has been identified as a virulence factor of Mtb which is recognized by host galectin-9. Here we demonstrate that galectin-9 directly inhibited mycobacterial growth through AG-binding property of carbohydrate-recognition domain 2. Furthermore, IgG antibodies with AG specificity were detected in serum of TB patients. Based on the interaction between galectin-9 and AG, we developed monoclonal antibody (mAb) screening assay and identified AG-specific mAbs which profoundly inhibit Mtb growth. Mechanistically, proteomic profiling and morphological characterizations revealed that AG-specific mAbs regulate AG biosynthesis, thereby inducing cell wall swelling. Thus, direct AG-binding by galectin-9 or antibodies contributes to protection against TB. Our findings pave the way for the rational design of novel immunotherapeutic strategies for TB control.

Keywords

Mycobacterium tuberculosis, tuberculosis, arabinogalactan, galectin-9, antibody

Introduction

Tuberculosis (TB), caused by *Mycobacterium tuberculosi*s (Mtb) remains a considerable threat to human health. In 2021, 10.6 million people fell ill with TB, 1.6 million people died from the disease, and 450,000 new TB cases suffered from rifampicin-resistant or multidrug-resistant TB on this globe (Bagcchi, 2023). Although TB treatment measures are in place in many parts of the world, cure rates are insufficient due to the increasing emergence of drug resistance. Therefore, novel intervention strategies are urgently needed. Rational design of novel TB therapeutics depends on better understanding of the crosstalk between Mtb and host cells.

Following inhalation of aerosols carrying Mtb, innate immune responses are initiated which constitute a first line of defense. Firstly, Mtb are engulfed by mononuclear phagocytes into the phagosome which matures and then fuses with lysosomes (Chandra et al., 2022). The pathogen-associated molecular patterns (PAMPs) from Mtb are recognized by a variety of pattern recognition receptors (PRRs), such as Toll-like receptors (TLRs), Nod-like receptors (NLRs), C-type lectins receptors (CLRs) and cyclic GMP-AMP synthase (cGAS), resulting in production of inflammatory cytokines, chemokines and anti-bacterial peptides to restrict bacterial growth (Fremond et al., 2004; Reiling et al., 2002; Watson et al., 2015; Wilson et al., 2015). Mtb can escape from the phagosome into the cytosol and can be recaptured in autophagosomes, through a process termed xenophagy, to form a degradative autolysosome (Lopez et al., 2018; Wang and Li, 2020). Cell-autonomous defense mechanisms also include production of reactive oxygen and nitrogen intermediates,

70 hypoxia, mild acidity and nutrition deprivation (Lupoli et al., 2018; Nathan and
71 Shiloh, 2000). These mechanisms help to limit growth and spread of the bacteria
72 within the cell, and contribute to the initiation of an adaptive immune response.
73 However, our knowledge about host immune factors that target Mtb components and
74 directly inhibit replication is limited.

75 Mtb can evade and resist immune defense by entering a state of dormancy which
76 can last for years. This is due to the capacity of Mtb to synthesize a sturdy cell wall,
77 slow down metabolism to promote growth arrest and implement the so-called
78 stringent response (Batt et al., 2020; Hauryliuk et al., 2015). These mechanisms
79 provide the basis for long-term persistence of Mtb until immune control weakens.
80 Once immune control deteriorates, Mtb acquires a metabolic active stage and induces
81 progression to active TB.

82 The complex cell wall of Mtb provides a barrier not only for host defense, but also
83 for antibiotics. Accordingly, components of the cell wall are well-established drug
84 targets. The essential core cell wall structure is composed of three distinct layers: a)
85 the cross-linked network of peptidoglycan (PGN), b) the highly branched
86 arabinogalactan (AG) polysaccharide, and c) the characteristic long-chain mycolic
87 acids (Jankute et al., 2015). Among these, AG has been an important target for
88 anti-TB drugs, though understanding of its biological functions is limited. E.g.,
89 ethambutol, one of the front-line anti-TB drugs, targets the arabinosyltransferases
90 EmbA, EmbB and EmbC, which are critical for AG synthesis (Escuyer et al., 2001;
91 Goude et al., 2009; Zhang et al., 2020). However, it remains unclear whether AG is

directly targeted by natural host immune factors in TB.

Recently, we identified AG as a virulence factor of Mtb that is recognized by galectin-9, a member of the β-galactoside binding gene family. Upon AG binding, galectin-9 initiates the downstream TAK1-ERK-MMP signaling cascade leading to pathologic impairment of the lung (Wu et al., 2021). Galectin-4 and galectin-8 directly kill *E.coli* by recognizing blood group antigens of bacteria (Stowell et al., 2010). This raises the question whether galectin-9 inhibits mycobacterial growth via targeting AG. Here, we demonstrate a novel cell-autonomous mechanism by which galectin-9 impedes mycobacterial growth via its AG-binding property in a carbohydrate recognition domain (CRD) 2-dependent mode. Moreover, in sera of TB patients, we identified anti-AG IgG antibodies which were supposed to defend against TB. Employing a monoclonal antibody (mAb) screening array, we identified anti-AG mAbs, CL010746 and CL046999, which were capable of restraining Mtb growth by regulating AG biosynthesis. Thus, AG possesses characteristic features of protective antigens. In sum, our work identified a previously unknown role of galectin-9 and anti-AG antibodies in TB control. Hence, our findings provide the basis for rational design of mAb-based immunotherapy of TB as a novel approach towards host-directed therapy of one of the deadliest infectious diseases globally.

Results

Galectin-9 inhibits mycobacterial growth

Our previous work demonstrated that galectin-9 directly interacts with AG and AG-containing bacteria (Wu et al., 2021). Given the capacity of galectin-4 and galectin-8 to kill *E.coli* (Stowell et al., 2010), we interrogated whether galectin-9 directly interferes with mycobacterial replication. Real-time monitoring of *in vitro* cultures revealed that recombinant galectin-9 protein inhibits growth of Mtb at a concentration as low as 10 ng/mL **(Figure 1A)**. Native structural conformation of galectin-9 was required for its bacteriostatic effect since heat inactivation at 95 °C for 5 min abrogated this activity **(Figure 1B)**. CFU assays further validated that galectin-9 inhibited Mtb growth **(Figure 1C)**. Galectin-9 also impaired replication of the fast-growing *Mycobacterium smegmatis* in a dose-dependent manner **(Figure 1D)**. Consistently, ELISA revealed profoundly higher abundance of galectin-9 in serum from TB patients than that from Tuberculin skin test (TST) negative (TST-) heathy donors, implying that galectin-9 contributes to resistance against Mtb infection **(Figure 1E)**. Of note, average concentration of galectin-9 in sera of TST positive (TST+) healthy donors was 3.748 ng/mL **(Figure 1E)**. We speculate that this high abundance of galectin-9 contributes to maintenance of latent TB infection by restricting Mtb spreading from granuloma, where the pathogen is contained. In sum, we conclude that galectin-9 directly inhibits mycobacterial growth.

Macrophages are part of the first line defense against invading mycobacteria. When human monocytic THP-1 cells were infected with *Mycobacterium bovis* BCG

fused with DsRed, immunofluorescence assays revealed recruitment of galectin-9 to mycobacteria in a time-dependent manner **(Figure 1F and 1G)**. In line with this observation, robust accumulation of galectin-9 around invading Mtb H37Rv-GFP was also observed in THP-1 cells post infection **(Figure 1H and 1I)**. Though galectin-9 has been reported to be critical for initiation of mTOR signaling and induction of autophagy (Bell et al., 2021; Jia et al., 2018; Jia et al., 2020), our work revealed a novel cell-autonomous mechanism whereby galectin-9 recruitment restricts mycobacterial growth in an autophagy-independent manner.

Carbohydrate recognition is essential for galectin-9-mediated inhibition of mycobacterial growth

Given that galectin-9 binds to β-galactoside, we interrogated whether carbohydrate recognition by galectin-9 is essential for inhibition of mycobacterial growth. Addition of lactose rich in β-galactoside (generally used for neutralization of carbohydrate binding of galectin-9) partially enhanced Mtb growth *in vitro*, and completely reversed mycobacterial growth inhibition by galectin-9 **(Figure 2A)**. In contrast, addition of glucose had no such effect **(Figure 2B)**. These results indicate that the β-galactose binding property of galectin-9 is involved in mycobacterial activity. Moreover, excess AG not only reversed growth inhibition by galectin-9, but also markedly promoted Mtb growth **(Figure 2C)**, indicating that the AG-binding property of galectin-9 is involved in anti-mycobacterial activity. Our previous work had demonstrated that CRD2, but not CRD1, of galectin-9 mediated its interaction with

AG (Wu et al., 2021). As expected, addition of purified CRD2, but not of CRD1, to some extent hindered Mtb growth, emphasizing that AG binding to galectin-9 was sufficient for anti-mycobacterial effects **(Figure 2D)**. Taken together, carbohydrate recognition is essential for galectin-9-mediated inhibition of mycobacterial growth.

Identification of anti-AG antibodies from TB patients

Given the higher abundance of galectin-9 in serum from active TB patients, we next interrogated whether anti-AG antibodies are present in serum of TB patients. An ELISA assay was developed for identification of anti-AG antibodies by coating AG on plates **(Figure 3A)**. An adequate window of serum dilutions allowed a linear correlation between OD_{450} and dilution over an appropriate range **(Figure 3B)**. Subsequently, we determined the relative abundance of anti-AG serum antibodies in 17 healthy donors and 25 active TB patients, all of whom had received BCG vaccination within 24 hours after birth. Our findings revealed a significant increase in the abundance of anti-AG antibodies among TB patients when compared to healthy donors **(Figure 3C)**. We speculate that during Mtb infection, anti-AG IgG antibodies are induced which contribute to protection against TB by directly inhibiting Mtb replication albeit apparently in vain. Whether anti-AG antibody levels at site of Mtb growth in patients are too low or whether growth inhibition is nullified by excessive replication of Mtb remains to be solved in the future.

Generation of anti-AG mAb

Based on the finding that AG binding of galectin-9 inhibits mycobacterial growth, we

embarked on the development of anti-AG mAbs with blocking acivity. Given the high affinity between galectin-9 and AG, we developed an antibody chip comprising 62,208 mAbs to screen for anti-AG activity (Wu et al., 2021). Briefly, the antibody chip was incubated with AG, and bound AG was subsequently detected using galectin-9 in conjunction with FITC-labeled anti-galectin-9 monoclonal antibody **(Figure 4A)**. We filtered out 12 candidate mAbs exhibiting binding affinity to AG **(Figure 4A, B)**. Subsequently, we validated their AG-binding capacity using ELISA. AG was coated on plates, and mAbs were added in 2-fold serial dilutions **(Figure 4C)**. The ELISA assay revealed a robust AG-binding curve for CL010746 (referred to as mAb1) and CL046999 (referred to as mAb2) **(Figure 4D)**. Furthermore, both mAbs exhibited specific binding to Mtb H37Rv-GFP as demonstrated by immunofluorescent assay **(Figure 4E**). Therefore, we have successfully developed anti-AG mAbs that bind Mtb directly.

Anti-AG antibody inhibits Mtb growth

For functional characterization, we monitored the *in vitro* mycobacterial growth in the presence or absence of anti-AG mAbs. Both mAb1 and mAb2 demonstrated inhibition of Mtb growth **(Figure 5A)**. This finding was further confirmed through CFU determination **(Figure 5B)**. Likewise, both mAbs markedly inhibited growth of *Mycobacterium smegmatis* as evidenced by real-time OD monitoring **(Figure 5C)** and CFU assay **(Figure 5D)**. In conclusion, the newly identified anti-AG mAbs demonstrated direct blockade of mycobacterial growth through binding to AG. Direct

inhibitory effects on Mtb growth by anti-AG antibodies emphasize that AG expresses features of protective antigens.

Proteomics profiling of the mycobacterial response to anti-AG mAb

To elucidate the molecular mechanisms underlying inhibition of mycobacterial growth by anti-AG antibodies, we conducted proteomic profiling of Mtb in response to anti-AG mAb1 treatment. Gene ontology (GO) enrichment analysis revealed significant enrichment of numerous cellular and metabolic processes, primarily related to biosynthesis of outer membrane, upon treatment with anti-AG mAb1. These processes include cell periphery, external encapsulating structure, organic substance metabolic process, cellular metabolic process, primary metabolic process, nitrogen compound metabolic process, and biosynthetic process (**Figure 6A**). Moreover, the formamidopyrimidine-DNA glycosylase N-terminal domain was enriched based on the analysis of functional enrichment and protein domain of differentially expressed antigens **(Figure 6B, C)**. Additionally, the KEGG pathway analysis demonstrated a significant enrichment of lipoarabinomannan (LAM) biosynthesis pathways **(Figure 6D)**. Consistently, the upregulated proteins Rv0236.1 and Rv3806c were involved in the biosynthesis of the mycobacterial cell wall arabinan **(Figure 6E)**. In the re-annotated genome sequence of Mtb, Rv0236.1 consists of Rv0236c and Rv0236A. Rv0236c is predicted to be a cognate of the GT-C superfamily of glycosyltransferases and likely acts as arabinofuranosyltransferase involved in AG synthesis (Skovierová et al., 2009). On the other hand, Rv0236A is a small secreted protein involved in cell

wall and cell processes (Marmiesse et al., 2004). Additionally, Rv3806c is a decaprenylphosphoryl-5-phosphoribose (DPPR) synthase involved in AG synthesis (He et al., 2015). These data provide compelling evidence to suggest that anti-AG mAbs regulate AG biosynthesis.

Targeting AG by mAbs modulates the cell wall of Mtb

To verify the impact of anti-AG mAbs on the biosynthesis of the mycobacterial cell wall, we characterized the morphological changes of Mtb treated with or without mAbs. Intriguingly, mAb1 and mAb2 treatment both led to a dispersed distribution of Mtb in cultures (**Figure 7A**). Acid fast staining further revealed the formation of a cord-like structure in Mtb treated with mAb1 or mAb2, which was not observed following ethambutol (EMB) treatment **(Figure 7B)**. Moreover, electron microscopy demonstrated that anti-AG mAbs treatment markedly increased the thickness of the Mtb cell wall **(Figure 7C)**. Based on these findings, we conclude that targeting AG by specific antibodies, and likely by galectin-9 as well, impairs growth of Mtb and other mycobacteria by modulating cell wall structure.

Discussion

Mtb, the etiologic agent of TB, is one of the leading causes of death worldwide, further aggravated by increasing incidences of antibiotic resistance (Miotto et al., 2018; Singh and Chibale, 2021). Hence, TB remains a major contributor to the global disease burden. Host-directed therapy is increasingly recognized as an alternative or

243 adjunct to antibiotic therapy (Kaufmann et al., 2018). Therefore, deeper insights into
244 the interactions between Mtb and the host immune system are warranted. We
245 previously demonstrated that mycobacterial AG binds to the galactoside-binding
246 protein galectin-9, causing pathologic impairments in the lung via the
247 TAK1-ERK-MMP signaling pathway (Wu et al., 2021). Here, we demonstrate that
248 galectin-9 directly impedes mycobacterial growth through its AG-binding property.
249 Further, we identified natural anti-AG antibodies in sera of TB patients, which are
250 predicted to inhibit Mtb growth. Based on these findings, we generated mAbs capable
251 of binding AG and hindering Mtb replication. Proteomics profiling of Mtb revealed
252 that the binding of anti-AG antibodies regulates AG biosynthesis which leads to
253 swelling of the cell wall, as validated by morphological characterization. We conclude
254 that galectin-9 and anti-AG antibodies serve as immune factors that restrain bacterial
255 growth by targeting AG in the cell wall. Increasing evidence suggests a role for
256 antibodies in protection against TB (Irvine et al., 2021; Lu et al., 2019). It is generally
257 assumed that the role of antibodies in TB is based on their interactions with
258 macrophages, which promote anti-mycobacterial activities such as phago-lysosome
259 fusion and production of reactive oxygen and nitrogen intermediates (Chandra et al.,
260 2022; Nathan and Shiloh, 2000). In striking contrast, our findings demonstrate that
261 anti-AG antibodies directly impair Mtb growth and thus emphasize that AG comprises
262 features of protective antigens.

263 Galectins are a highly conserved class of molecules that play critical roles in
264 multiple biological processes. Fifteen different types of galectins are known in

265 humans, which can be classified based on their structure, subcellular localization, and
266 function. For instance, galectin1 participates in regulating cell proliferation, apoptosis,
267 and immune responses through interactions with specific glycosylated receptors on
268 the cell surface, such as integrins and CD45 (Cedeno-Laurent et al., 2012; Ge et al.,
269 2016; Perillo et al., 1995). More recently, it was shown that galectin-4 disrupts
270 bacterial membranes and kills *E.coli* through interactions with lipopolysaccharides on
271 the bacterial outer membrane (Stowell et al., 2010). Here, we demonstrate that
272 galectin-9 significantly inhibits replication of Mtb by interacting with AG in Mtb via
273 its CRD2 domain. Similar to galectin-9, galectin-4, galectin-6 and galectin-8 also
274 comprise 2 CRDs in tandem connected by a linker sequence(Leffler et al., 2002). It
275 remains to be explored whether and how these galectins exert anti-mycobacterial
276 activities via the CRD2 domain, thereby providing general insights into the role of
277 galectin family cognates in immunity to TB.

278 After phagocytosis by pulmonary macrophages of the newly infected host, Mtb
279 ends up inside phagosomes, where it downregulates its metabolism and enters a
280 non-replicating persistent (NRP) state, termed dormancy, in response to host stress
281 (Gengenbacher and Kaufmann, 2012; Russell, 2001). Once the immune response
282 "breaks down", Mtb transits into a metabolically active and replicative state which
283 ultimately results in progression to active TB disease (van der Wel et al., 2007).We
284 demonstrated that galectin-9 accumulates around invading mycobacteria in host cells.
285 However, whether Mtb recruits galectin-9 during dormancy, its active stage, or during
286 both stages, has not been investigated. Of note, AG is hidden by mycolic acids in the

287 outer layer. We speculate that during Mtb replication, cell wall synthesis is elevated
288 and AG becomes exposed, thereby facilitating its binding to galectin-9 and leading to
289 Mtb growth arrest.

290 Major drugs in clinical use for TB treatment inhibit Mtb growth by targeting
291 different essential components and processes. For instance, Isoniazid inhibits mycolic
292 acid synthesis by targeting InhA enzyme (Quémard et al., 1995), Ethambutol blocks
293 AG biosynthesis by targeting EmbCAB complex (Telenti et al., 1997), and
294 Pyrazinamide disrupts the pH balance within the bacterial cell, thereby impairing
295 mycobacterial growth (Zhang et al., 1999). Although the mycobacterial cell wall has
296 been widely exploited as antibiotic target, to date, drugs that directly bind AG and
297 inhibit Mtb growth have not been reported. Here, we identified host galectin-9 and
298 anti-AG antibodies (both serum antibodies from patients and mAbs) which recognize
299 AG and thus inhibit Mtb replication. Hence, anti-AG mAbs can be harnessed for
300 design of novel biologics which address the challenge of drug resistance in TB.

301 Mechanisms underlying inhibition of mycobacterial growth induced by galectin-9
302 or anti-AG mAbs remain elusive. We propose they interfere with the activity of
303 enzymes involved in AG biosynthesis and/or modify the physical properties of the cell
304 wall, leading to disruption of AG side chain extension, thereby increasing Mtb
305 vulnerability to host immunity. They may also function through the two-component
306 system, that is commonly found in bacteria and allows bacteria to sense and respond
307 to changes in the environment, such as nutrient availability or stress (Glover et al.,
308 2007; James et al., 2012; Majumdar et al., 2012). Interactions between galectin-9 and

309 AG in the cell wall may alter membrane permeability, which restrains nutrient uptake
310 and activates sensor proteins, causing bacterial growth arrest.

311 Aside from their direct anti-Mtb activity, anti-AG antibodies in serum of TB
312 patients probably also opsonize Mtb, thereby promoting phagocytosis by mononuclear
313 phagocytes (Chen et al., 2016; Lu et al., 2019). However, the mechanisms by which
314 galectin-9 or antibodies inhibit mycobacterial growth depend on the details of the
315 molecular interactions and require further investigation.

316 Our knowledge about antibodies which target glycans is scarce, not the least due
317 to technical challenges. Glycan antigens have been identified on the surface of
318 numerous microorganisms and are also expressed by certain cancer cells. Antibodies
319 that recognize and bind these glycan antigens, therefore, are promising candidates for
320 therapy and diagnosis of infectious and malignant diseases. For instance, the mAb
321 2G12 neutralizes human immunodeficiency virus-1 (HIV-1) by recognizing
322 oligomannose-type N-glycans on the HIV-1 gp120 envelope protein, and the mAb
323 FH6 specifically binds the Sialyl Lewis X (SLeX) antigen on the surface of various
324 cancer cells (Fukushi et al., 1984; Kannagi et al., 1986; Trkola et al., 1996). In this
325 study, we not only discovered anti-AG mAbs which directly impair Mtb growth, but
326 also developed an efficient high-throughput screening for identifying mAbs with
327 specificity for glycans.

328 In conclusion, we (i) discovered a novel cell-autonomous mechanism by which
329 galectin-9 protects against TB via targeting AG in the cell wall of Mtb, (ii) identified
330 neutralizing antibodies against AG in serum of TB patients, (iii) selected anti-AG

331 mAbs for passive immunization against TB by means of a mAb screening array, (iv)
332 characterized inhibition of Mtb replication by induction of cell wall swelling as
333 critical mechanism of protection through AG targeting (**Figure 7D**). Our findings,
334 thus, not only provide deeper insights into humoral immune mechanisms involved in
335 protection against TB, but also serve as basis for new intervention strategies against
336 TB in adjunct to chemotherapy.
337

Materials and Methods

Bacteria

Mtb H37Rv, H37Rv-GFP, *Mycobacterium bovis* BCG, and *Mycobacterium smegmatis* mc^{2}155, were from Shanghai Key Laboratory of Tuberculosis and grown in Middlebrook 7H9 (Becton Dickinson, Cockeysville, MD) liquid medium supplemented with 0.25% glycerol, 10% oleic acid–albumin-dextrose-catalase (OADC) (Becton Dickinson, Sparks, MD) and 0.05% Tween-80.

In vitro growth of mycobacteria

Mycobacteria were harvested at mid-log phase and diluted to a calculated starting OD600 of 0.25 and added to 96-well culture plates containing Middlebrook 7H9 liquid medium together with antibody or galectin-9. OD600 of each time point of each strain was tested in real time by Bioscreen C microplate incubator (FP1100-C, Labsystems, USA) at 37℃.

CFU assay

Mycobacteria were harvested at mid-log phase and diluted to a calculated starting OD600 of 0.25 and incubated in Middlebrook 7H9 liquid medium with or without antibody or galectin-9 for 30 h at 37 ℃. Appropriate dilutions were plated on 7H10 agar plates for enumeration of CFU.

Immunofluorescence assay

For colocalization of galectin-9 and mycobacteria, immunofluorescence assays were performed as described previously (Liu et al., 2018). Briefly, THP-1 cells (ATCC Cat# TIB-202 RRID: CVCL_0006) were infected with bacteria for 2 h, fixed with 4%

formaldehyde for 30 min at R.T., permeabilized with 0.1% Triton X-100 in PBS for 5 min, and blocked with 5% BSA in PBS for 60 min at R.T.. Cells were stained with the anti-galectin-9 antibody (Cell Signaling Technology, Cat#54330, RRID:AB_2799456), antibodies at a dilution of 1:200 in 5% BSA in PBS overnight at 4 °C and then incubated with Alexa Fluor 488 or 555 conjugated secondary antibodies (Thermo Fisher Scientific, Cat# A-11008; RRID: AB_143165; Cat# A32732, RRID:AB_2633281) at a dilution of 1:1000 for 2 h at R.T.. Nuclei were stained with DAPI.

For binding of anti-AG mAb with Mtb, H37Rv-GFP strains were harvested at mid-log phase. Subsequently, 2×10^7 H37Rv-GFP/100 μL FACS buffer was incubated with anti-AG mAb (mAb1 or mAb2, 20 μg/mL) at R.T. for 1h, and washed three times with PBST (PBS containing 0.05 % Tween 20, pH 7.4) by centrifugation (12,000g, 5 min). The resulting sediment was resuspended in 10 μL of ddH2O and smeared on microscopic slides.

Images were acquired using a Leica TCS SP8 confocal laser microscopy system (Leica Microsystems) at ×63 magnification.

Validation of anti-AG antibodies by ELISA

AG antigen (10 μg in 100 μL 0.1 mol/L $NaHCO_3$ buffer, pH 9.4) was added to wells of microwell plates and incubated overnight at 4°C. After four rinses with PBST (PBS containing 0.05 % Tween 20, pH 7.4), the wells were saturated with blocking buffer. After four additional PBST rinses, serum or candidate anti-AG mAbs were added to each well and incubated for 1 h at 37°C. The wells were rinsed three times with PBS

containing 0.05 % Tween 20 and horseradish-peroxidase-labeled rabbit anti-human or anti mouse IgG (100 μL/well; Sigma-Aldrich, Germany) was added to each well and incubated for 1 h at 37°C. Finally, the TMB substrate was added and the absorbance was measured with the Thermo Fisher Scientific Multiskan FC microplate photometer. Clinical serum specimens were collected from 17 healthy volunteers and 25 pulmonary TB patients before undergoing treatment at Shanghai Pulmonary Hospital (Shanghai, PR China), all of whom had received BCG vaccination within 24 h after birth. The donors are between the ages of 50 and 65, ethnic Han, with an equal representation of males and females.

High throughput Screening of anti-AG antibody

Antibody chip harboring 62208 mAbs was incubated with 10 μg AG in 10 mL incubation buffer (1x PBS buffer containing 10% BSA) for 1 h, followed by incubation with 10 μg galectin-9 protein in 10 mL incubation buffer for 1 h. The chip was incubated with rabbit anti-galetin-9 antibody (1:5000 diluted in 10 mL incubation buffer; ab227046, Abcam, UK) followed by staining with FITC-labelled anti-rabbit IgG (1:5000 diluted in 10 mL incubation buffer; ab6717, Abcam, UK). Then the chip was scanned by GenePix 4200A Microarray Scanner (Molecular Devices LLC) and analyzed by GenePix Pro 6.0 software.

Proteomics analysis

LC-MS/MS

Mycobacteria at mid-log phase were diluted to a calculated starting OD600 of 0.25 and incubated in Middlebrook 7H9 liquid medium with or without mAb1 for 30 h at

37 ℃. The bacterial pellets were collected followed by three washes with sterile saline. The bacterial pellets were resuspended in lysis buffer (8 M urea, 1% Protease Inhibitor Cocktail) and inactivated for 10 min at 100 ℃. The lysate was sonicated three times on ice using a high intensity ultrasonic processor (Scientz) and centrifuged. The protein concentration of the lysate was determined with BCA kit according to the manufacturer's instructions.

After trypsin digestion, peptides were dissolved in 0.1% formic acid and separated with nanoElute UHPLC system (Bruker, Germany) and subjected to Capillary source followed by the timsTOF Pro mass spectrometry. The resulting MS/MS data were processed using Maxquant search engine (v1.6.6.0). Tandem mass spectra were searched against Mtb strain ATCC 25618 83332 PR 20191210 database (3993 entries) concatenated with reverse decoy database.

Enrichment of Gene Ontology analysis

GO annotations of proteins are divided into three broad categories: Biological Process, Cellular Component, and Molecular Function. For each category, a two-tailed Fisher's exact test was employed to test the enrichment of the differentially expressed protein against all identified proteins. The GO with a corrected p-value < 0.05 was considered significant.

Enrichment of pathway analysis

Kyoto Encyclopedia of Genes and Genomes (KEGG) database was used to identify enriched pathways by a two-tailed Fisher's exact test to test the enrichment of the differentially expressed protein against all identified proteins. The pathway with a

corrected p-value < 0.05 was considered significant. These pathways were classified into hierarchical categories according to the KEGG website.

Enrichment of protein domain analysis

For each category proteins, InterPro database (a resource that provides functional analysis of protein sequences by classifying them into families and predicting the presence of domains and important sites) was researched and a two-tailed Fisher's exact test was employed to test the enrichment of the differentially expressed protein against all identified proteins. Protein domains with a corrected p-value < 0.05 were considered significant.

Morphologic characterization of Mtb

Mtb strains were cultured in 96 U well culture plates containing Middlebrook 7H9 liquid medium with anti-AG antibody or without anti-AG antibody at 37 ℃, and were grown for 10-14 days until the formation of colonies. Morphologic characterization of tested strains in liquid medium were observed by 2 x magnifier. At the same time, 10 μl of culture were spread onto a glass slide. Smears on glass slides were fixed under ultraviolet light overnight. Glass slides were stained with Ziehl–Neelsen stain using a TB Stain Kit (Baso DIAGNOTICS TAIWAN, Zhuhai, China). Morphological characteristics or cell lengths of tested strains were observed using a Leica DM2500 microscope using the 100× objective.

For ultrastructural characteristics, strains at mid-log phase were collected, and analyzed by Tecnai transmission electron microscopy (TEM) with 160 kV according the procedures of the manufacturer (GOODBIO, Wuhan, China). A 10^7-10^8 bacterial

suspension was used for TEM examination.

Statistical analysis

The statistical significance of comparisons was analyzed with two tailed unpaired Student's t test or Mann-Whitney U test in GraphPad Prism version 8.0.1.. $P < 0.05$ was considered statistically significant. All data are showed as mean ± SD of two or more independent experiments performed in triplicate. Detailed statistical information on each experiment is provided in the respective figure legends.

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Disclosure Statement

The authors report there are no competing interests to declare.

Author Contributions

H.L. conceived the study and designed the experiments.

L.Q., J.X., J.C., and S.W. analyzed the data and wrote the original draft.

H.L., W.S., L.Z., and X.M. supervised the project and edited the manuscript.

R.Z, Z.C., Z.L., Y.W., J.W., and X.H. performed most of the experiments.

Z.W., M.W., and R.P. performed the bioinformatics analysis.

W.S. provided clinical samples.

S.H.E.K provided helpful discussions and edited the manuscript.

All authors commented on the paper.

477

478 **Materials availability statement**

479 Bacterial strains and monoantibodies generated in this study are available from the
480 Lead Contact upon reasonable request.

481

482 **Data availability statement**

483 The authors confirm that the data supporting the findings of this study are available
484 within the article.

485

486 **Statement of Ethics**

487 This work received approval from the Clinical Ethics Committee of the Shanghai
488 Pulmonary Hospital, which is affiliated with the Tongji University School of
489 Medicine, in accordance with governmental guidelines and institutional policies.
490 Patients providing the pulmonary samples gave informed consent. The ethics
491 committee approved this consent procedure. All protocols were approved by the local
492 ethics committee of Tongji University School of Medicine (permit number:
493 K21-114Y).

494

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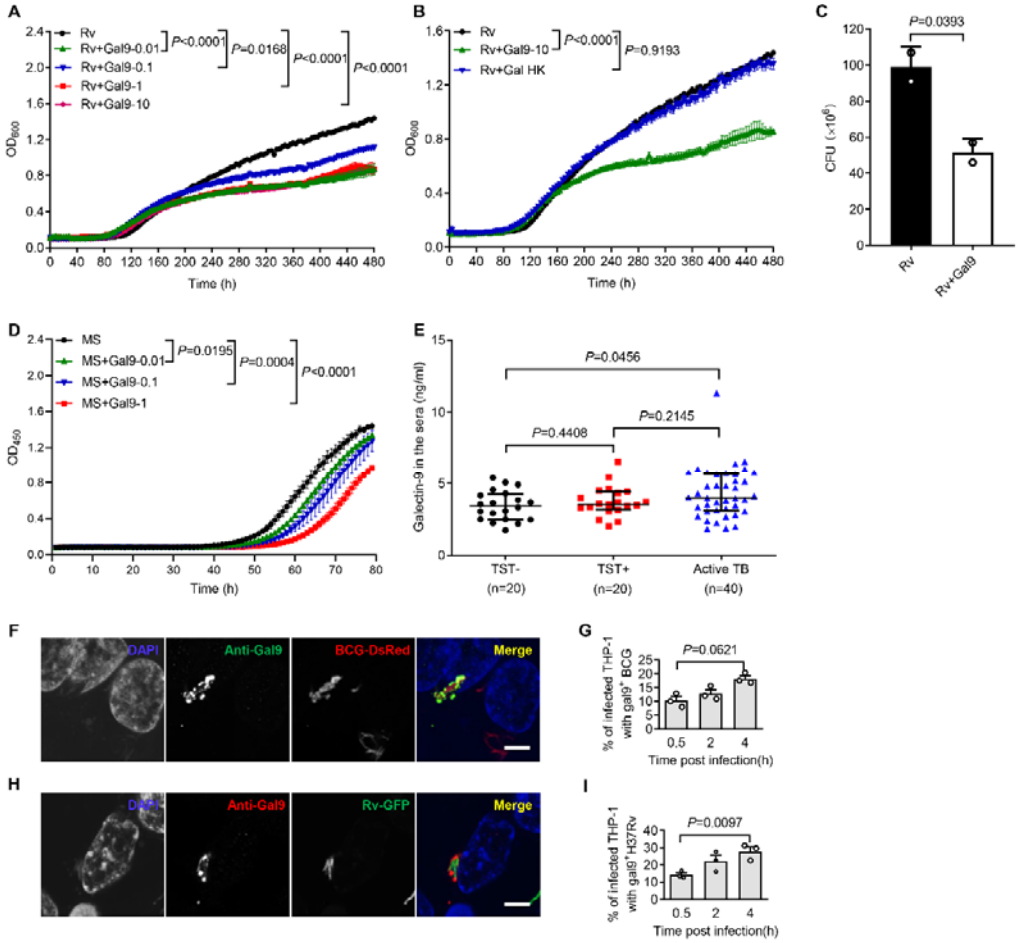
655 **Figures**



656 **Figure 1. Galectin-9 inhibits mycobacterial growth directly.**

657 A. Profile of Mtb H37Rv (Rv) grown at 37°C in Middlebrook 7H9 liquid medium

658 with different concentration of Galectin-9 (Gal9, 0, 0.01, 0.1, 1, 10 μg/mL).

659 Growth curve was measured using a Bioscreen Growth Curve Instrument. Optical

660 density was measured at absorbance at 600 nm every 2 h.

661 B. Growth profile of Mtb H37Rv (Rv) in Middlebrook 7H9 liquid medium with10

662 μg/mL galectin-9 (Gal9) or inactivated galectin-9 (Gal9 HK, heat killed at 95 °C

663 for 5 min).

664 C. CFU of Mtb H37Rv (Rv) on Middlebrook 7H10 solid medium with or without 10

μg/mL Galectin-9 (Gal9). Cultures were grown at 37°C for 4-8 weeks.

D. Growth profile of *Mycobacterium smegmatis* (MS) in Middlebrook 7H9 liquid medium with different concentrations of Galectin-9 (Gal9, 0, 0.01, 0.1, 1 μg/mL).

E. Concentrations of galectin-9 in sera of tuberculin skin test negative healthy donors (TST-, n = 20), tuberculin skin test positive healthy donors (TST+, n = 20) and active TB patients (Active TB, n = 40).

F. Confocal microscopy of *M. bovis* BCG-DsRed (BCG-DsRed, red) and Galectin-9 (Anti-Gal9, green) in THP-1 cells. Nuclei was stained with DAPI (blue).

G. Percent of cells with galectin-9 positive ($Gal9^+$) BCG in total infected THP-1 cells. Symbols indicate colocalization ratio of at least 12 fields in each experiment.

H. Confocal microscopy of Mtb H37Rv-GFP (Rv-GFP, green) and Galectin-9 (Anti-Gal9, red) in THP-1 cells. Nuclei were stained with DAPI (blue).

I. Percent of cells with galectin9 positive ($Gal9^+$) Mtb H37Rv in total infected THP-1 cells. Symbols indicate colocalization ratio of at least 12 fields in each experiment.

Data are shown as mean ± SD, n = 3 biologically independent experiments performed in triplicate (A-D). Data are representative of three independent experiments with similar results (F and H). Two-tailed unpaired Student's t test (A-D, G, and I) or Mann-Whitney U test (E). $P < 0.05$ was considered statistically significant.

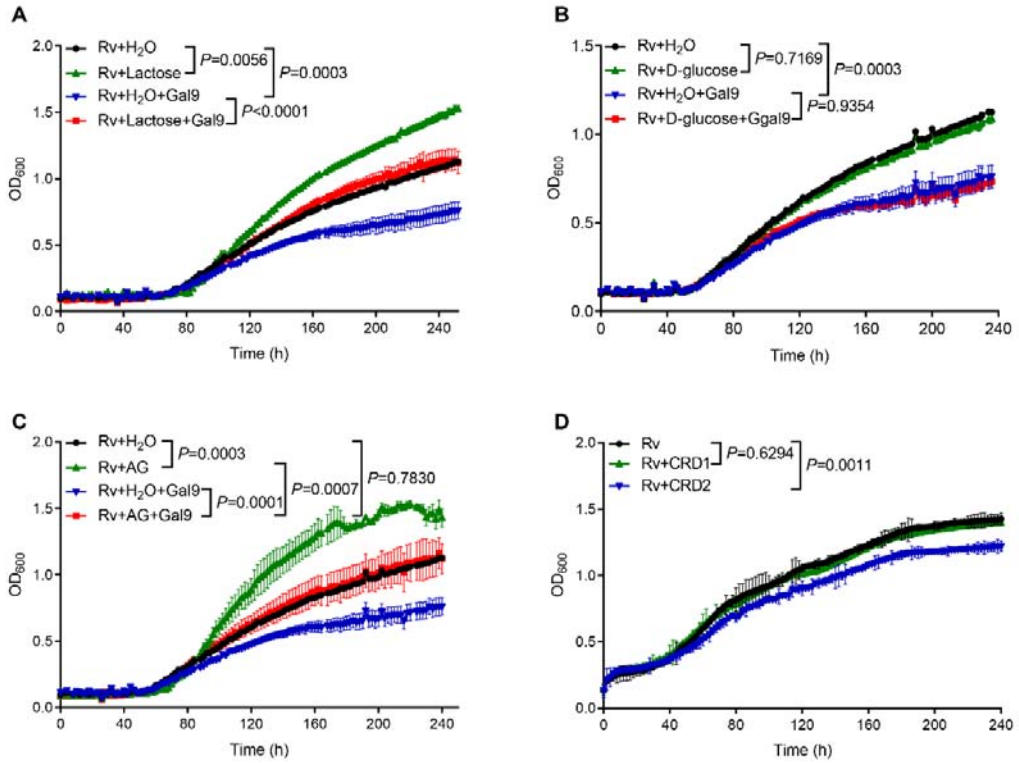


Figure 2. Carbohydrate recognition is essential for galectin-9-mediated inhibition of Mtb growth.

A. Growth profile of Mtb H37Rv (Rv) in Middlebrook 7H9 liquid medium with or without galectin-9 (Gal9, 10 μg/mL) and lactose (10 μg/mL).

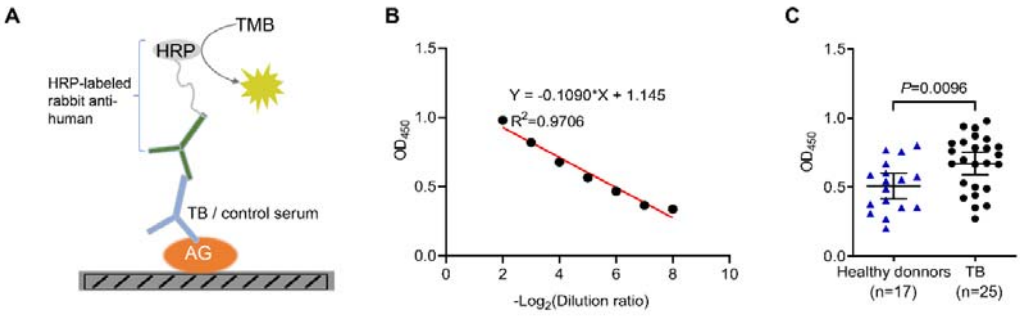
B. Growth profile of Mtb H37Rv (Rv) in Middlebrook 7H9 liquid medium with or without galectin-9 (Gal9, 10 μg/mL) and D-glucose (10 μg/mL).

C. Growth profile of Mtb H37Rv (Rv) in Middlebrook 7H9 liquid medium with or without galectin-9 (Gal9, 10 μg/mL) and AG (10 μg/mL).

D. Growth profile of Mtb H37Rv (Rv) in Middlebrook 7H9 liquid medium with 1μg/mL CRD1 or CRD2 of galectin-9.

Data are shown as mean ± SD, n = 3 biologically independent experiments performed in triplicate (A-D). Two-tailed unpaired Student's t test (A-D). $P < 0.05$ was

696 considered statistically significant.



697 **Figure 3. Identification of anti-AG antibodies from TB patients.**

698 A. Schematic presentation of ELISA assay for detecting anti-AG IgG antibodies in the

699 serum of TB patients.

700 B. Linear correlation between OD and serum dilution ratio determined by ELISA

701 assay.

702 C. Anti-AG IgG antibodies levels in TB patients (n = 25) and healthy

703 BCG-immunized controls (n = 17) determined via ELISA.

704 Data are representative of three independent experiments with similar results (B).

705 Mann-Whitney U test (C). $P < 0.05$ was considered statistically significant.

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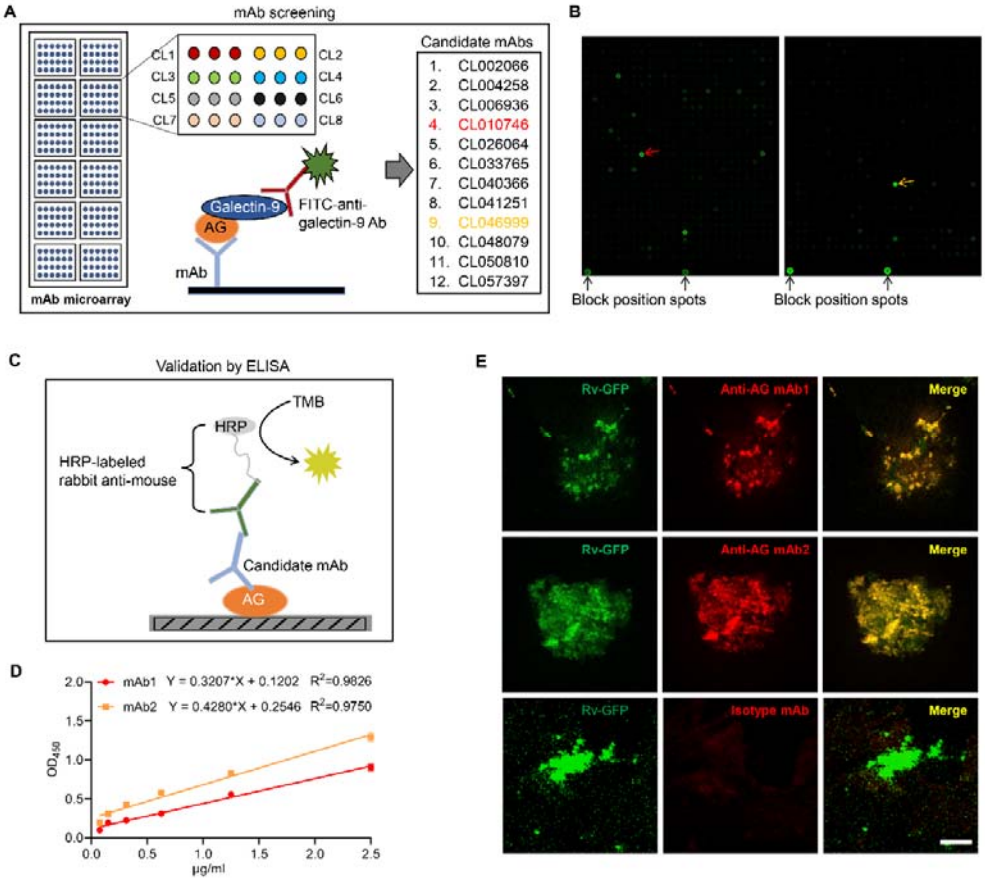


Figure 4. Development of anti-AG mAbs.

A. Schematic presentation of mAb screening for AG specificity.

B. Representative image of chip hybridization for mAb screening. Bright spots in the bottom mark the end line of each array block. Other spots represent AG binding to mAbs. CL010746 (mAb1) and CL046999 (mAb2) were labeled with red arrow and yellow arrow, respectively.

C. Schematic presentation of candidate anti-AG mAbs validation by ELISA.

D. Binding curve of mAb1 and mAb2 to AG determined by ELISA assay.

E. Confocal microscopy of Mtb H37Rv-GFP (Rv-GFP, green) and anti-AG mAbs (red).

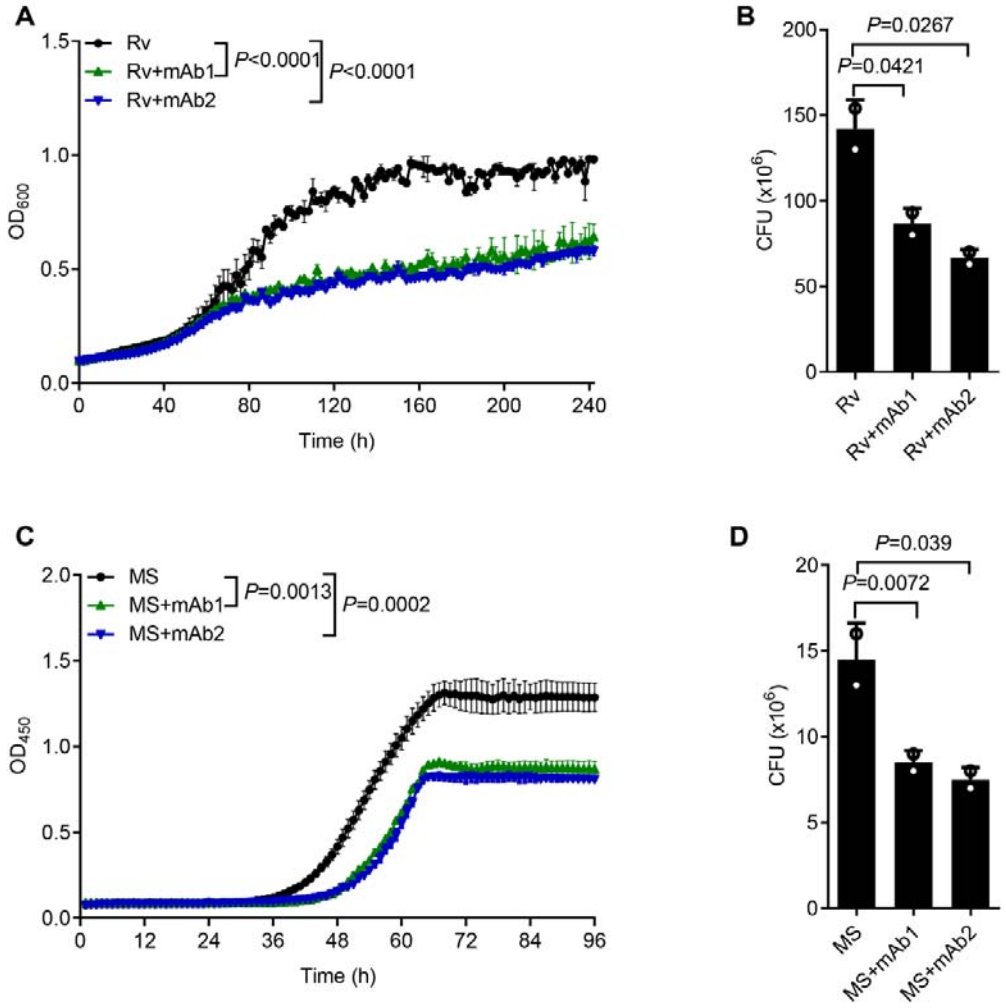


Figure 5. anti-AG antibody inhibits mycobacterial growth.

A. Growth profile of Mtb H37Rv (Rv) in Middlebrook 7H9 liquid medium with or without mAb1/mAb2 (1 μg/mL).

B. CFU of Mtb H37Rv (Rv) on Middlebrook 7H10 solid medium with or without mAb1/mAb2 (1 μg/mL). Cultures were grown at 37°C for 4-8 weeks.

C. Growth profile of *Mycobacterium smegmatis* (MS) in Middlebrook 7H9 liquid medium with or without mAb1/mAb2 (1 μg/mL).

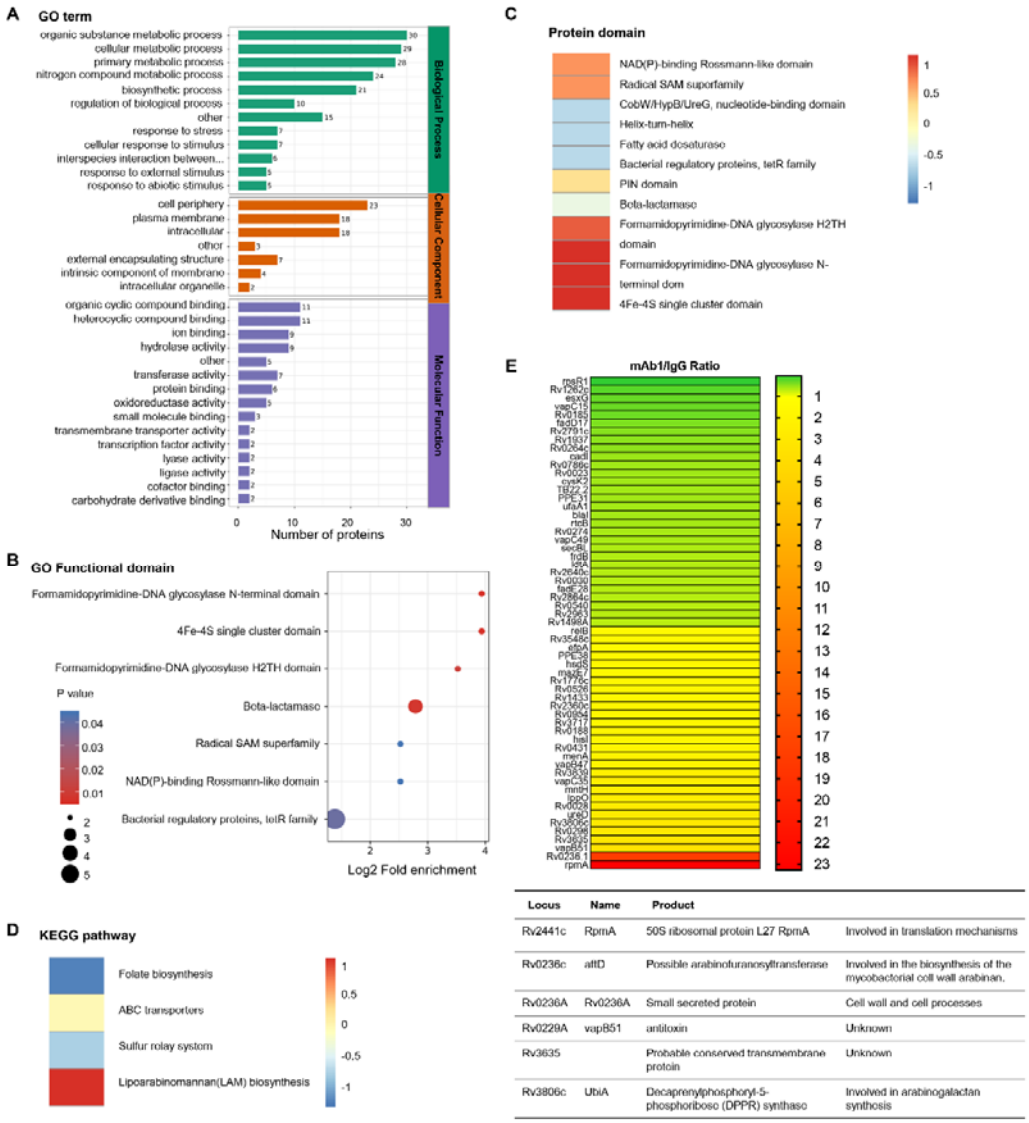
D. CFU of *Mycobacterium smegmatis* (MS) on Middlebrook 7H10 solid medium with or without mAb1/mAb2 (1 μg/mL). Cultures were grown at 37°C for 5-10 days.

727 Data are shown as mean ± SD, n = 3 (A, C) and n = 2 biologically independent

728 experiments performed in triplicate (B, D). Two-tailed unpaired Student's t test (A-D).

729 $P < 0.05$ was considered statistically significant.

730



Locus	Name	Product	
Rv2441c	RpmA	50S ribosomal protein L27 RpmA	Involved in translation mechanisms
Rv0236c	aftD	Possible arabinofuranosyltransferase	Involved in the biosynthesis of the mycobacterial cell wall arabinan.
Rv0236A	Rv0236A	Small secreted protein	Cell wall and cell processes
Rv0229A	vapB51	antitoxin	Unknown
Rv3635		Probable conserved transmembrane protein	Unknown
Rv3806c	UbiA	Decaprenylphosphoryl-5-phosphoribose (DPPR) synthase	Involved in arabinogalactan synthesis

Figure 6. Proteomics profiling of the response of Mtb to anti-AG antibody.

A. GO class of differentially expressed proteins in Mtb H37Rv treated with mAb1 (1 μg/mL) for 30 h followed by proteomics analysis. IgG was set as control.

B. Functional enrichment of differentially expressed proteins in Mtb H37Rv in (A).

C. Protein domain of differentially expressed proteins in Mtb H37Rv in (A).

D. KEGG class of differentially expressed proteins in Mtb H37Rv in (A).

E. Upregulation or downregulation genes in Mtb H37Rv in (A).

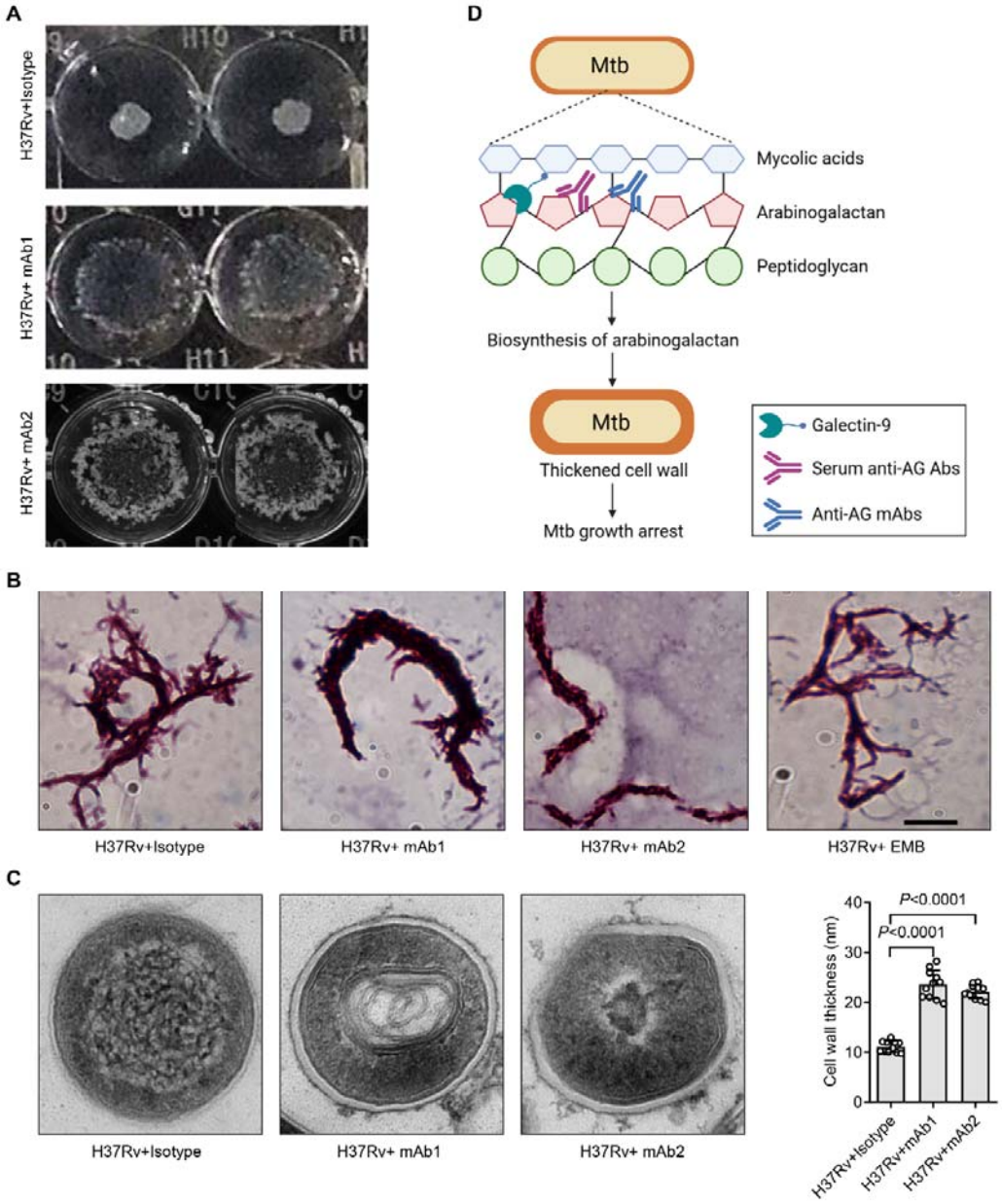


Figure 7. Mtb cell wall modulation by anti-AG antibodies.

A. Morphologic characteristics for Mtb H37Rv strain grown in liquid culture with or without anti-AG mAbs (1 μg/mL) observed by × 2 magnifier.

B. Bacterial shape of Mtb H37Rv strain treated as in (A) observed by acid fast staining under a Leica DM2500 microscope using the 100× oil microscopy. EMB,

744 Ethambutol. Scale bar, 20μm.

745 C. Ultrastructural morphology of Mtb H37Rv treated as in (A) analyzed by

746 transmission electron microscopy (TEM).

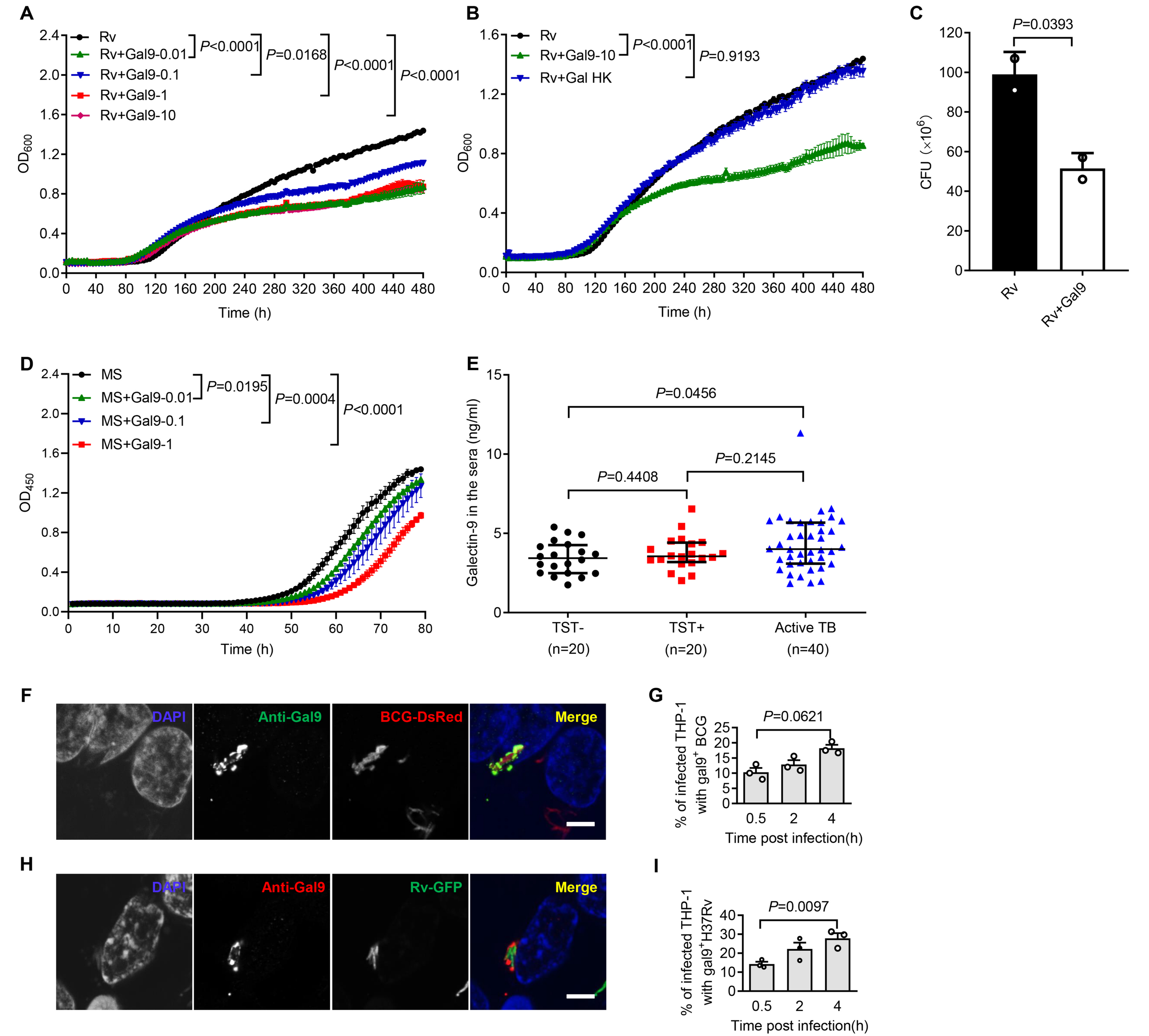
747 D. Schematic presentation of Mtb growth arrest by Galectin-9 or anti-AG antibodies.

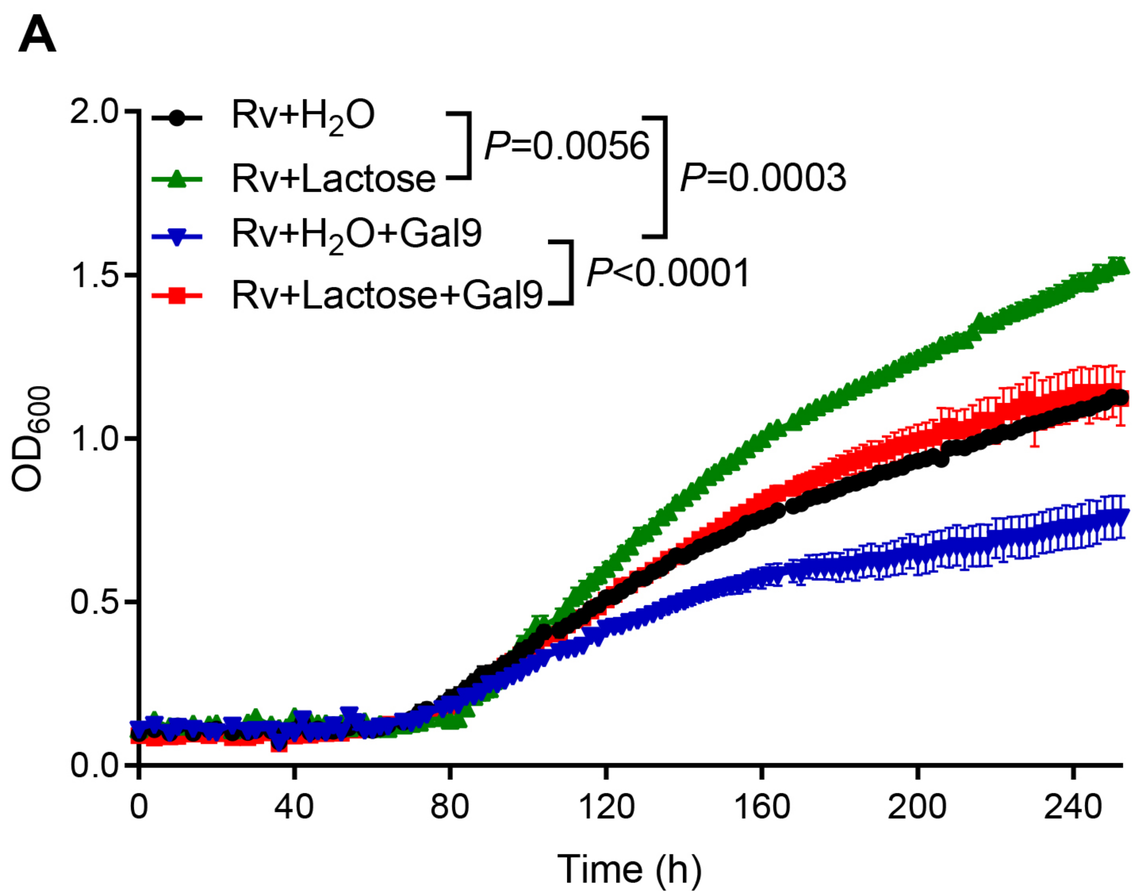
748 Data are representative of three independent experiments with similar results (A, B

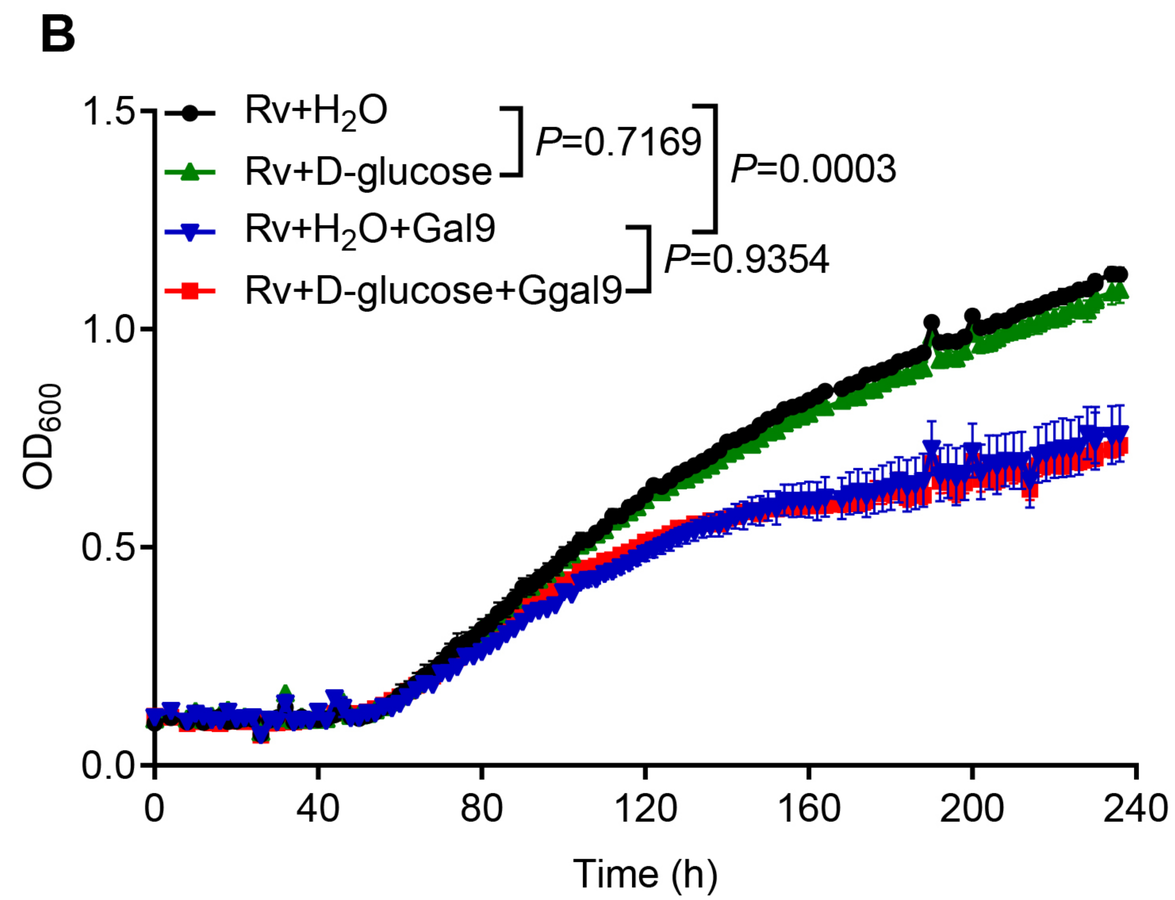
749 and C, left panel). Data are means ± SD of five bacteria, representatives of three

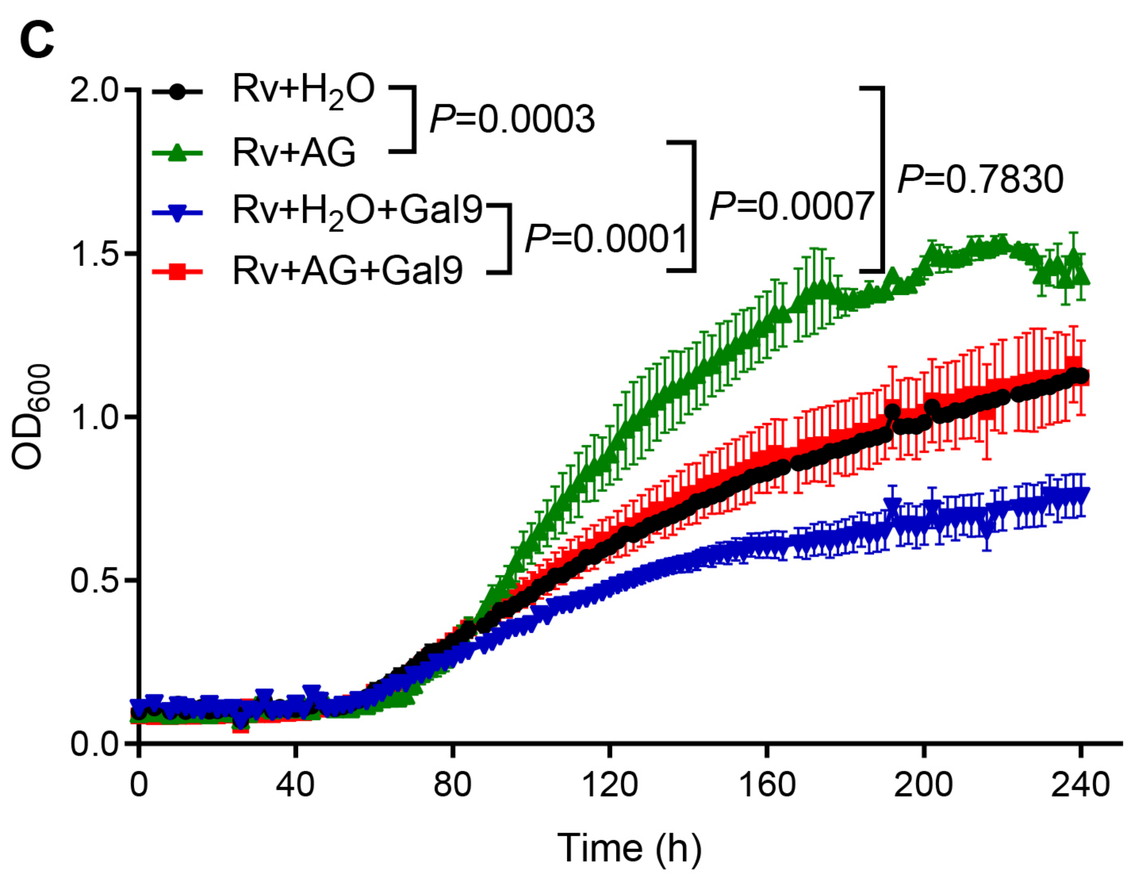
750 independent experiments (C, right panel). Two-tailed unpaired Student's t test (C,

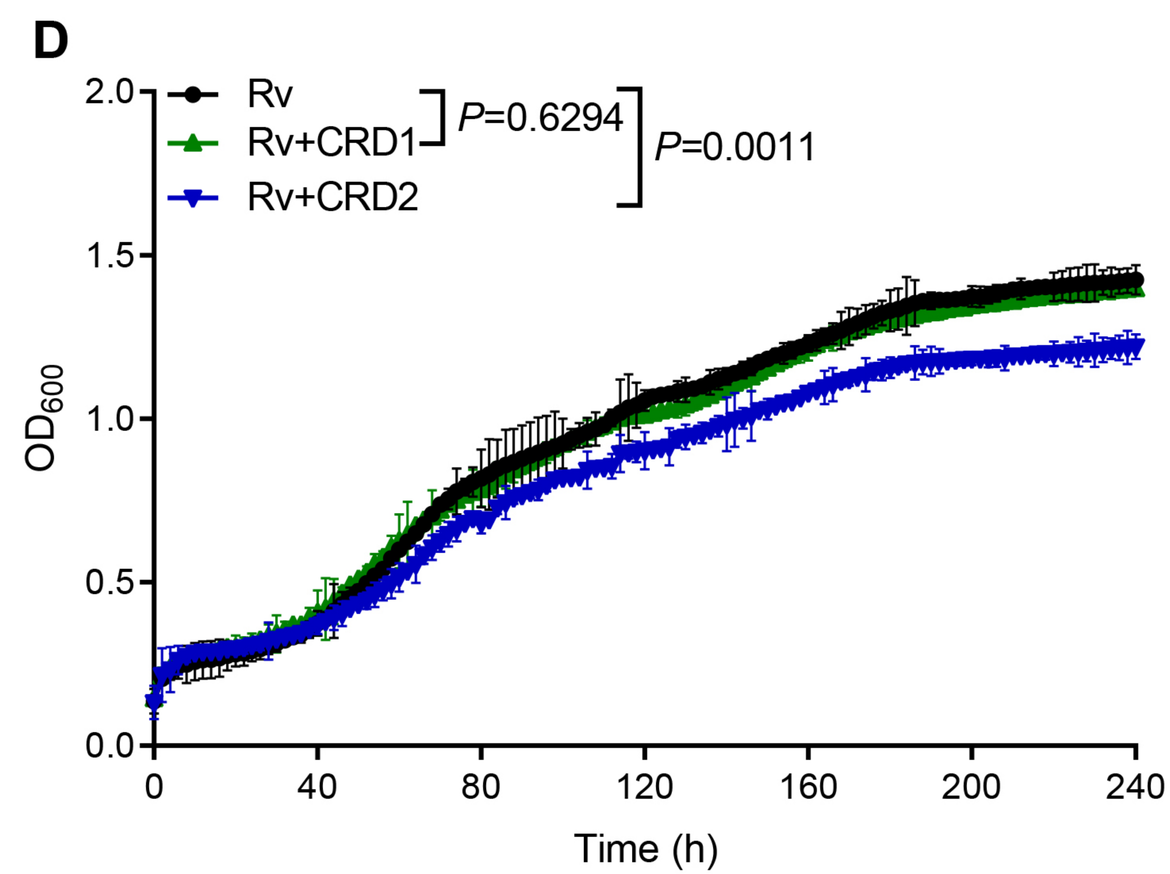
751 right panel). $P < 0.05$ was considered statistically significant.

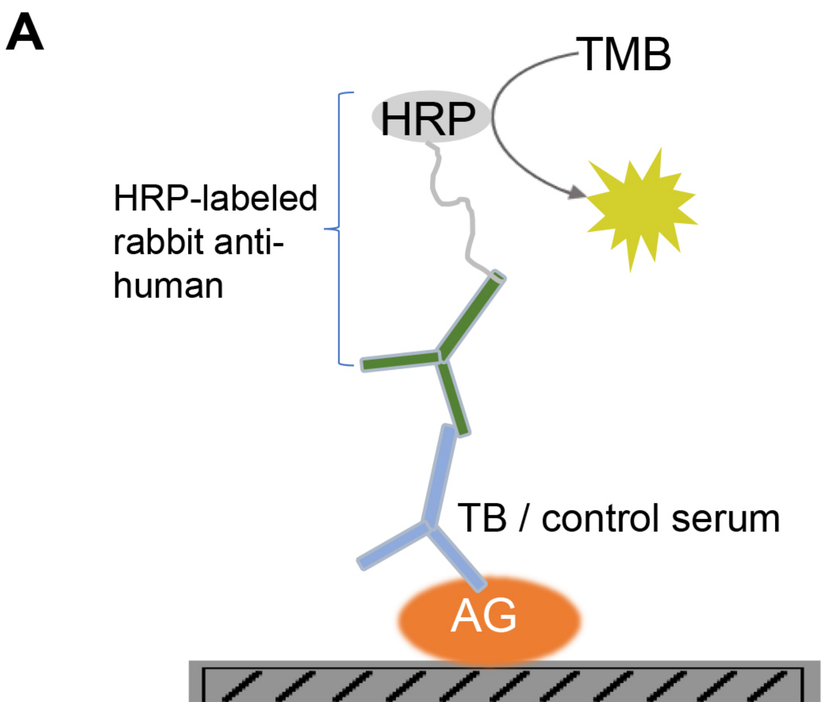

A
Rv
Rv+Gal9-0.01
Rv+Gal9-0.1
Rv+Gal9-1
Rv+Gal9-10
P<0.0001
P=0.0168
P<0.0001
P<0.0001
OD600
Time (h)
B
Rv
Rv+Gal9-10
Rv+Gal HK
P<0.0001
P=0.9193
OD600
Time (h)
C
P=0.0393
CFU (×10^6)
Rv
Rv+Gal9
D
MS
MS+Gal9-0.01
MS+Gal9-0.1
MS+Gal9-1
P=0.0195
P=0.0004
P<0.0001
OD450
Time (h)
E
P=0.0456
P=0.2145
P=0.4408
Galectin-9 in the sera (ng/ml)
TST-
(n=20)
TST+
(n=20)
Active TB
(n=40)
F
DAPI
Anti-Gal9
BCG-DsRed
Merge
G
P=0.0621
% of infected THP-1 with gal9+ BCG
Time post infection(h)
H
DAPI
Anti-Gal9
Rv-GFP
Merge
I
P=0.0097
% of infected THP-1 with gal9+H37Rv
Time post infection(h)

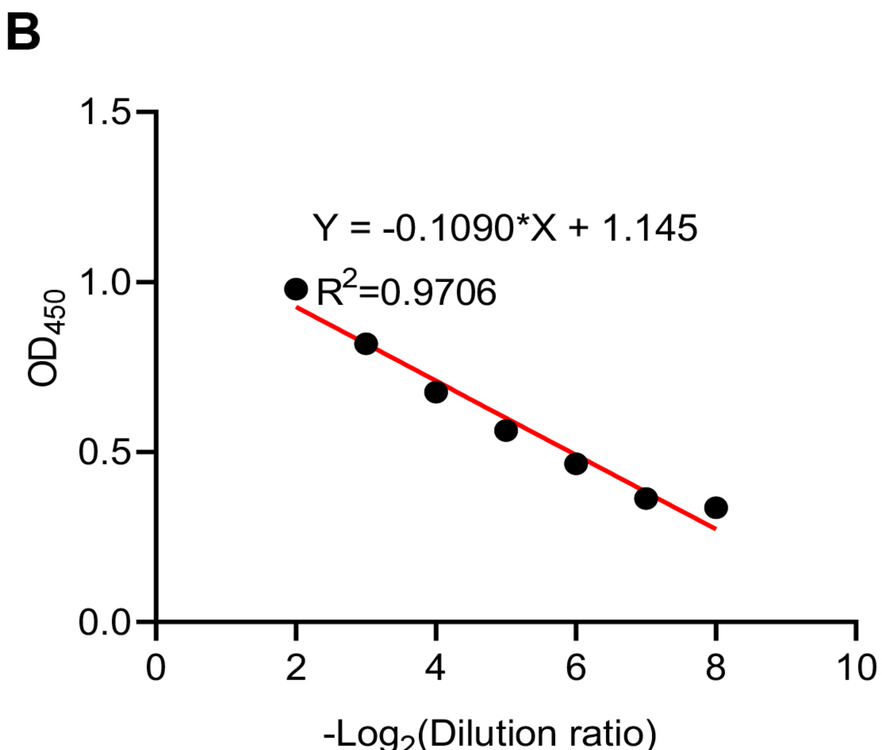

A
Rv+H2O
Rv+Lactose
Rv+H2O+Gal9
Rv+Lactose+Gal9
P=0.0056
P=0.0003
P<0.0001
OD600
Time (h)

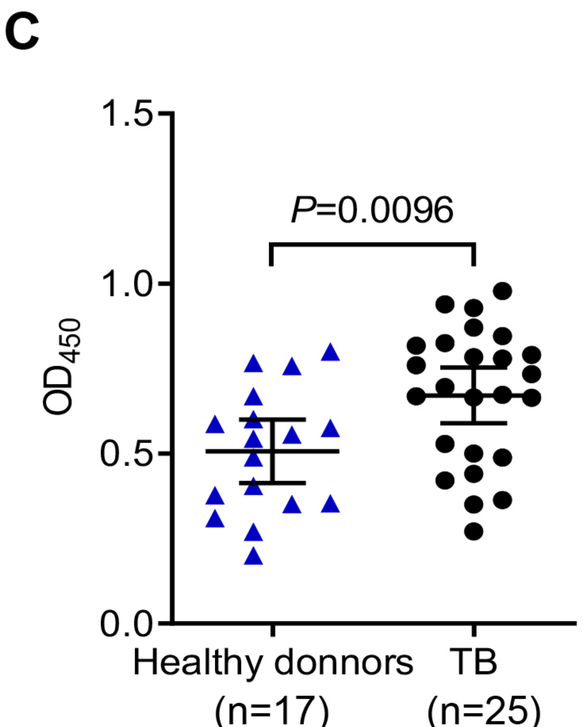

B
Rv+H2O
Rv+D-glucose
Rv+H2O+Gal9
Rv+D-glucose+Ggal9
P=0.7169
P=0.0003
P=0.9354
OD600
Time (h)

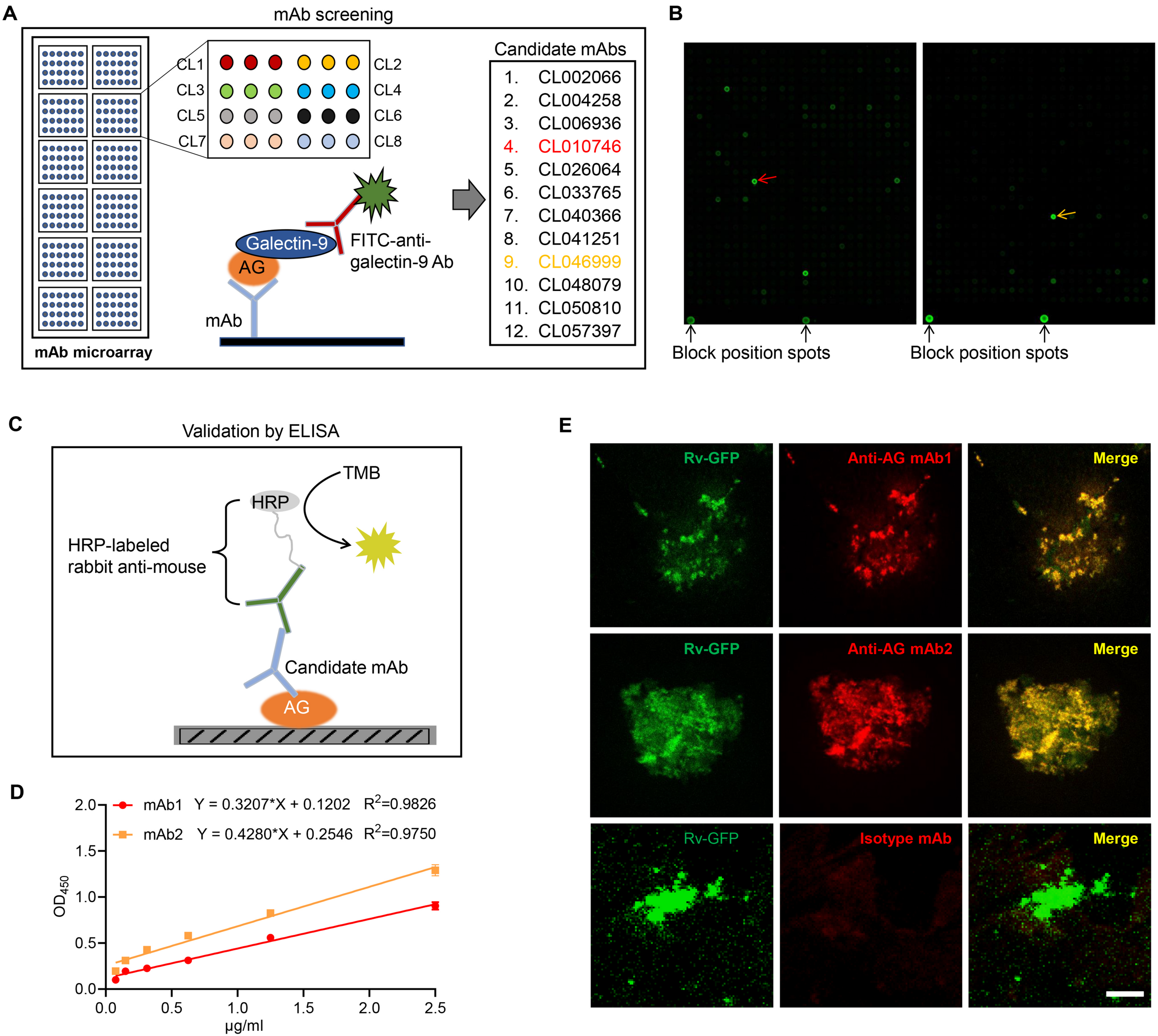

C
Rv+H2O
Rv+AG
Rv+H2O+Gal9
Rv+AG+Gal9
P=0.0003
P=0.0001
P=0.0007
P=0.7830
OD600
Time (h)

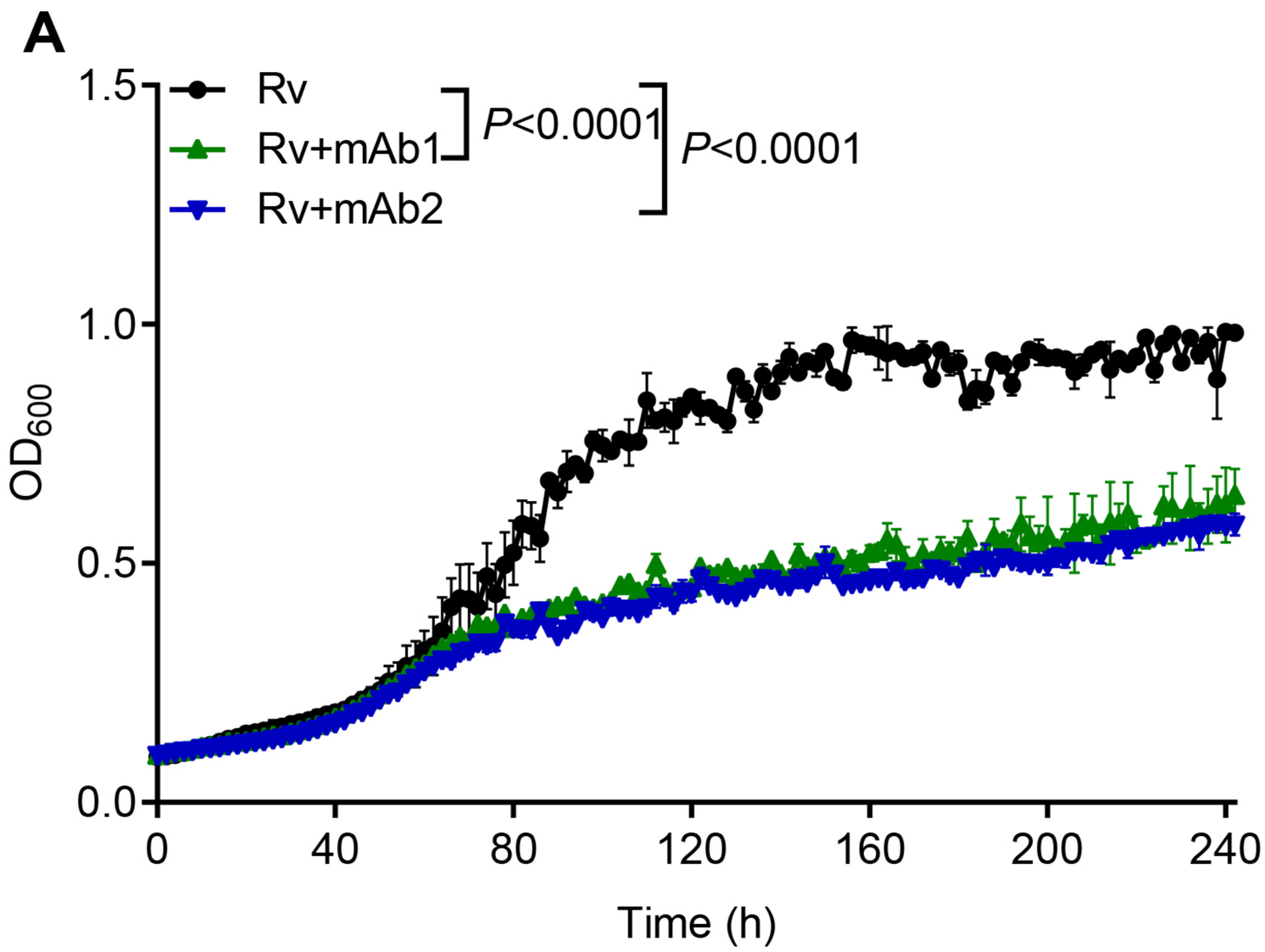

D
Rv
Rv+CRD1
Rv+CRD2
P=0.6294
P=0.0011
OD600
Time (h)

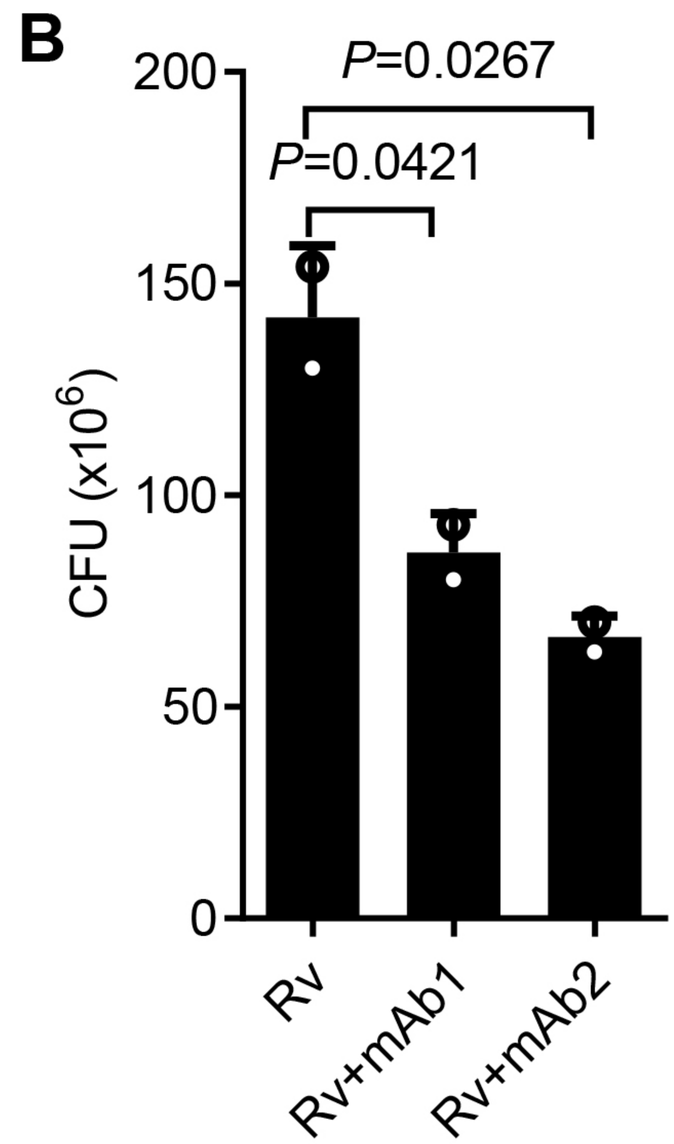

A
TMB
HRP
HRP-labeled rabbit anti-human
TB / control serum
AG

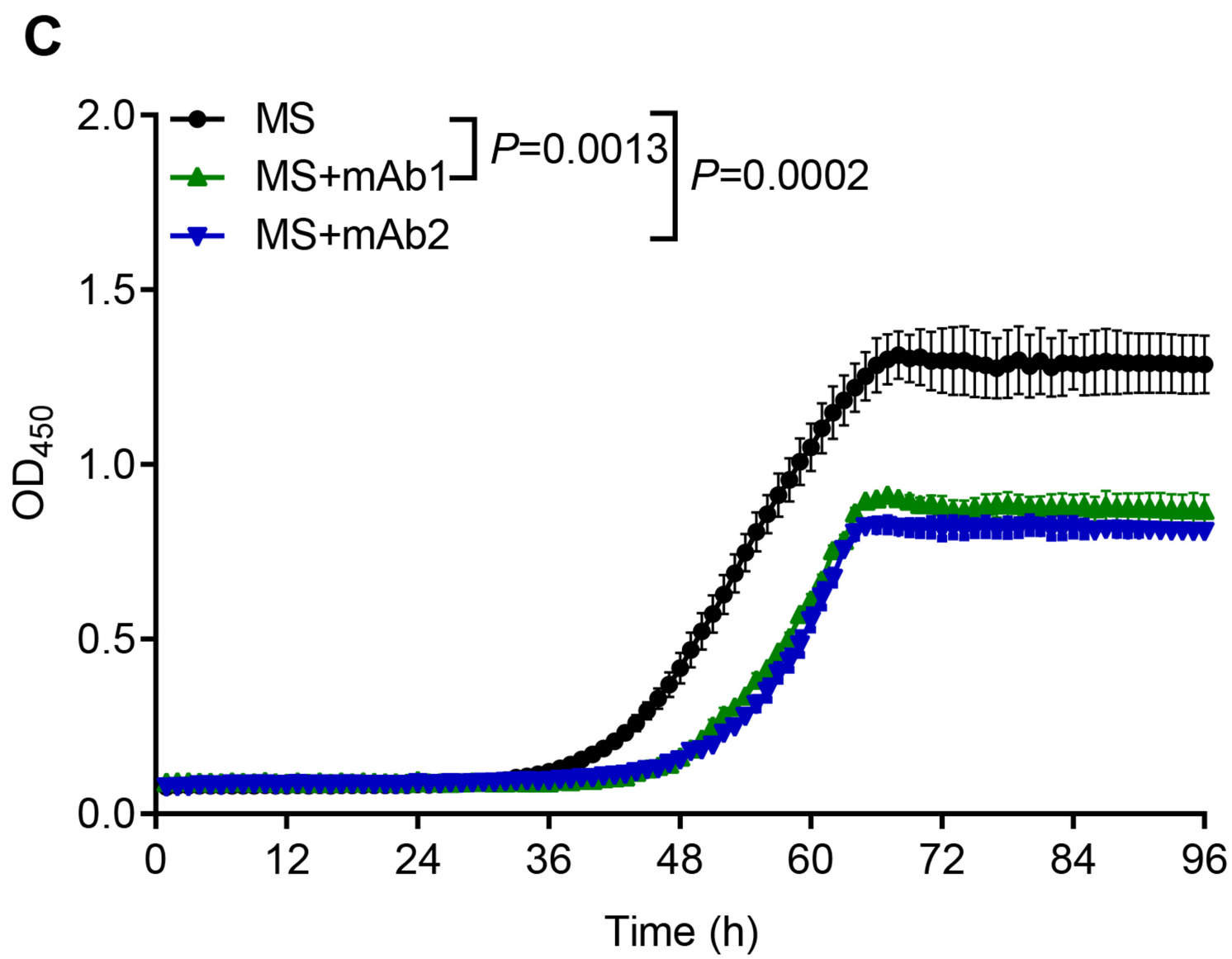

B
Y = -0.1090*X + 1.145
R2=0.9706
OD450
1.5
1.0
0.5
0.0
0
2
4
6
8
10
-Log2(Dilution ratio)

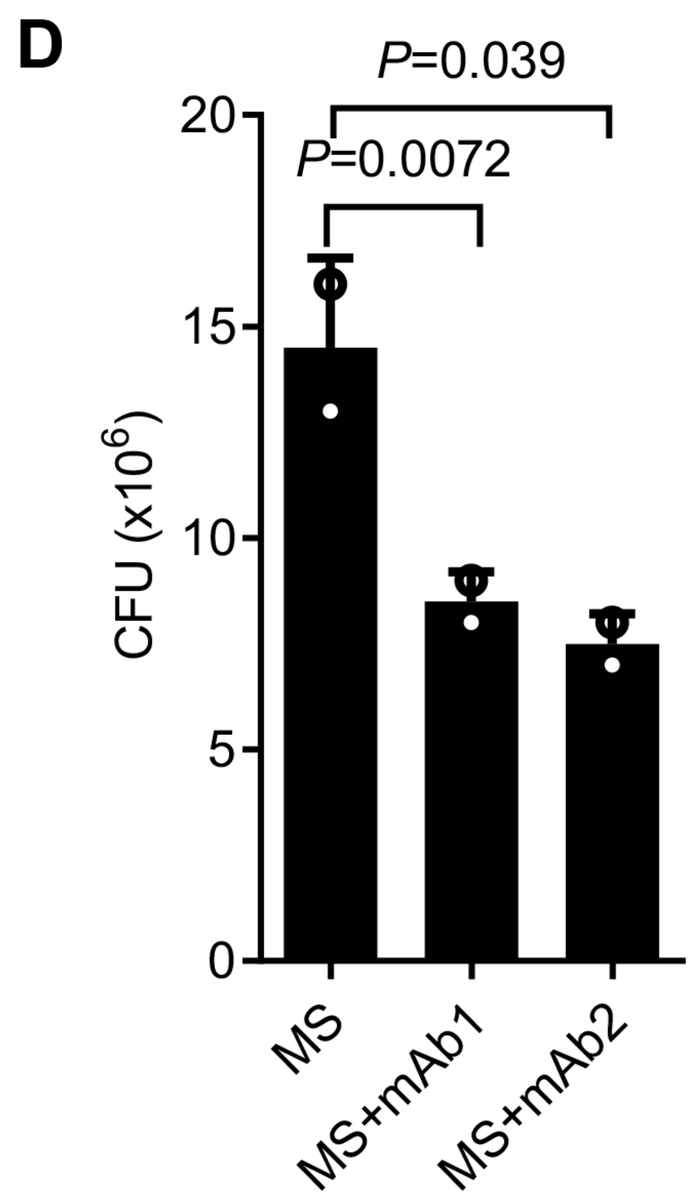

C
P=0.0096
OD450
1.5
1.0
0.5
0.0
Healthy donors (n=17)
TB (n=25)

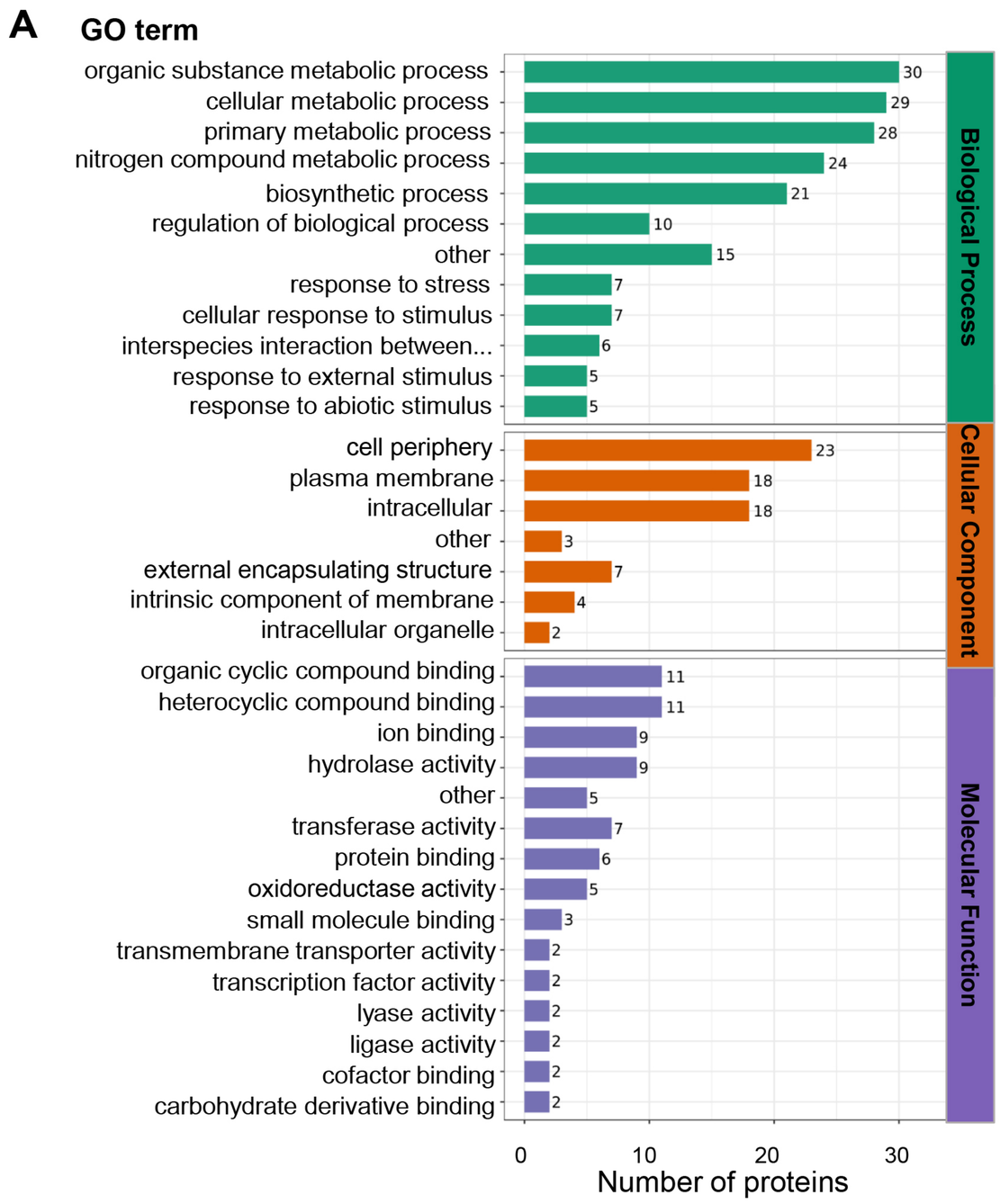

A
mAb screening
CL1
CL2
CL3
CL4
CL5
CL6
CL7
CL8
mAb microarray
Galectin-9
AG
mAb
FITC-anti-galectin-9 Ab
Candidate mAbs
1. CL002066
2. CL004258
3. CL006936
4. CL010746
5. CL026064
6. CL033765
7. CL040366
8. CL041251
9. CL046999
10. CL048079
11. CL050810
12. CL057397
B
Block position spots
Block position spots
C
Validation by ELISA
TMB
HRP
HRP-labeled rabbit anti-mouse
Candidate mAb
AG
D
mAb1 Y = 0.3207*X + 0.1202 R2=0.9826
mAb2 Y = 0.4280*X + 0.2546 R2=0.9750
OD450
2.0
1.5
1.0
0.5
0.0
0.0
0.5
1.0
1.5
2.0
2.5
μg/ml
E
Rv-GFP
Anti-AG mAb1
Merge
Rv-GFP
Anti-AG mAb2
Merge
Rv-GFP
Isotype mAb
Merge

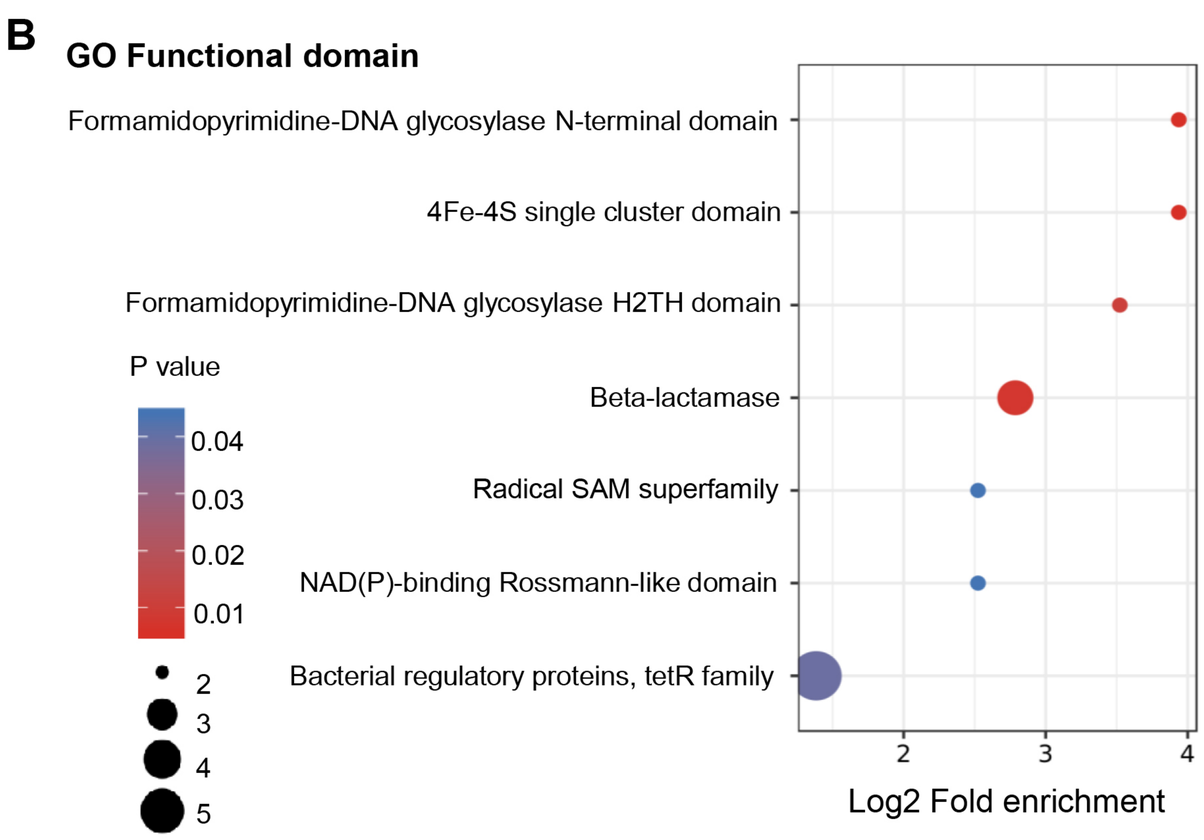

A
Rv
Rv+mAb1
Rv+mAb2
P<0.0001
P<0.0001
OD600
Time (h)
0.0
0.5
1.0
1.5
0
40
80
120
160
200
240

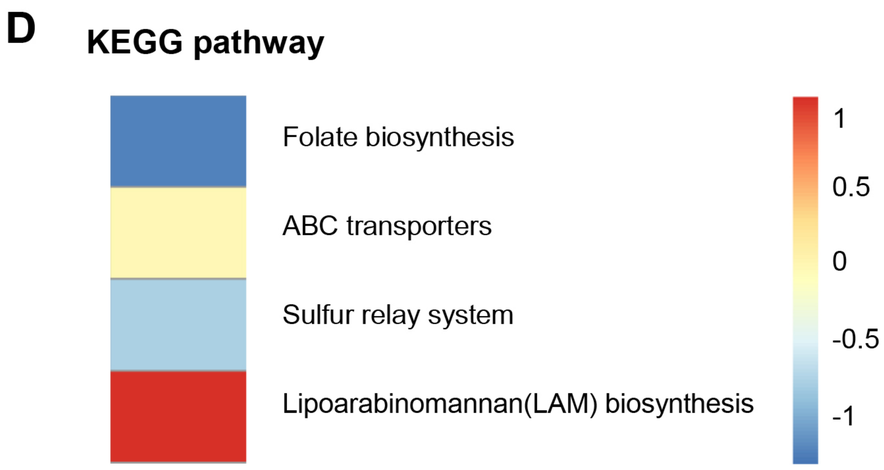

B
P=0.0267
P=0.0421
CFU (x10^6)
0
50
100
150
200
Rv
Rv+mAb1
Rv+mAb2

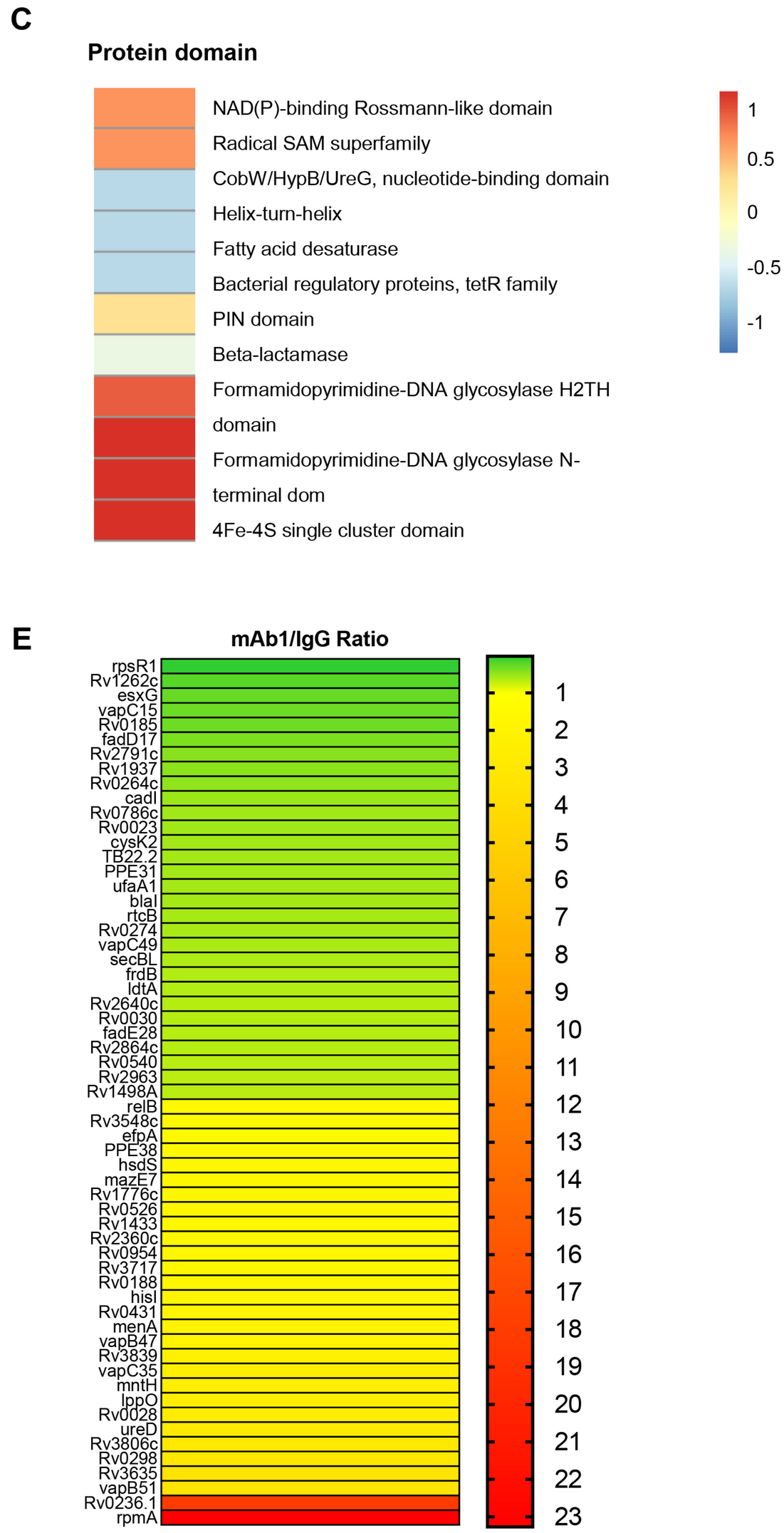

C
MS
MS+mAb1
MS+mAb2
P=0.0013
P=0.0002
OD450
Time (h)
0.0
0.5
1.0
1.5
2.0
0
12
24
36
48
60
72
84
96


D
P=0.039
P=0.0072
CFU (x10^6)
0
5
10
15
20
MS
MS+mAb1
MS+mAb2



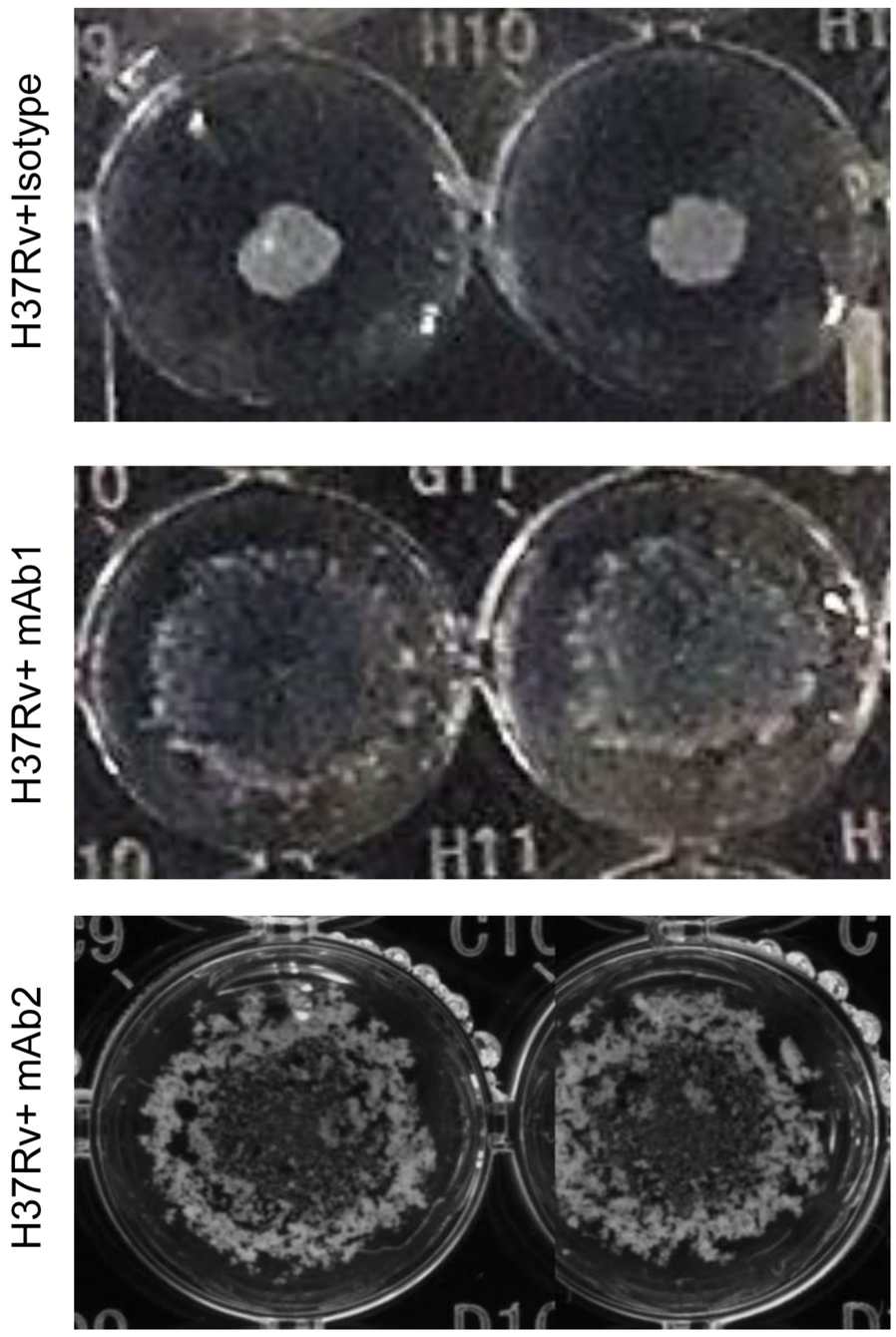




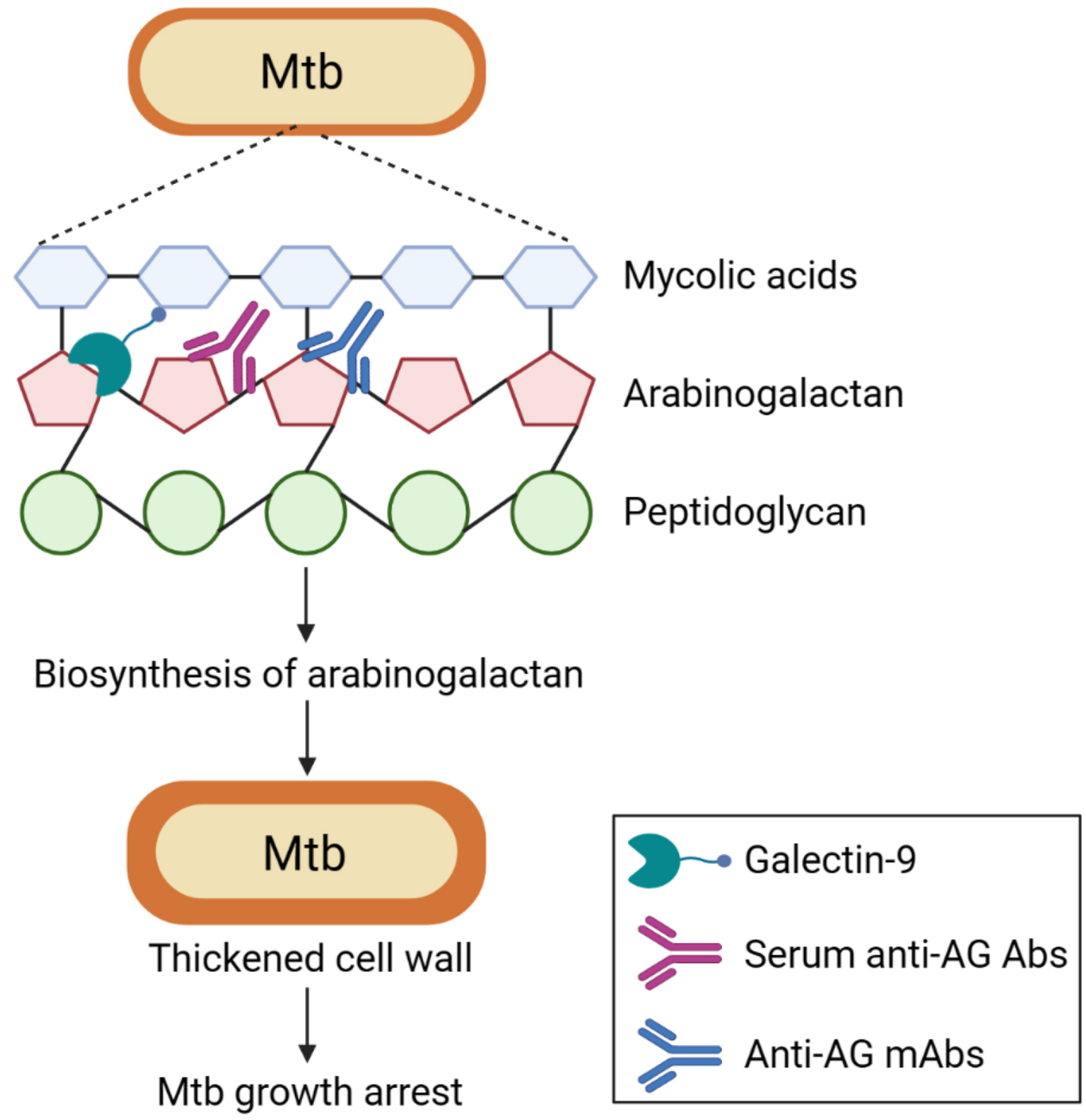


Locus	Name	Product	
Rv2441c	RpmA	50S ribosomal protein L27 RpmA	Involved in translation mechanisms
Rv0236c	aftD	Possible arabinofuranosyltransferase	Involved in the biosynthesis of the mycobacterial cell wall arabinan.
Rv0236A	Rv0236A	Small secreted protein	Cell wall and cell processes
Rv0229A	vapB51	antitoxin	Unknown
Rv3635		Probable conserved transmembrane protein	Unknown
Rv3806c	UbiA	Decaprenylphosphoryl-5-phosphoribose (DPPR) synthase	Involved in arabinogalactan synthesis

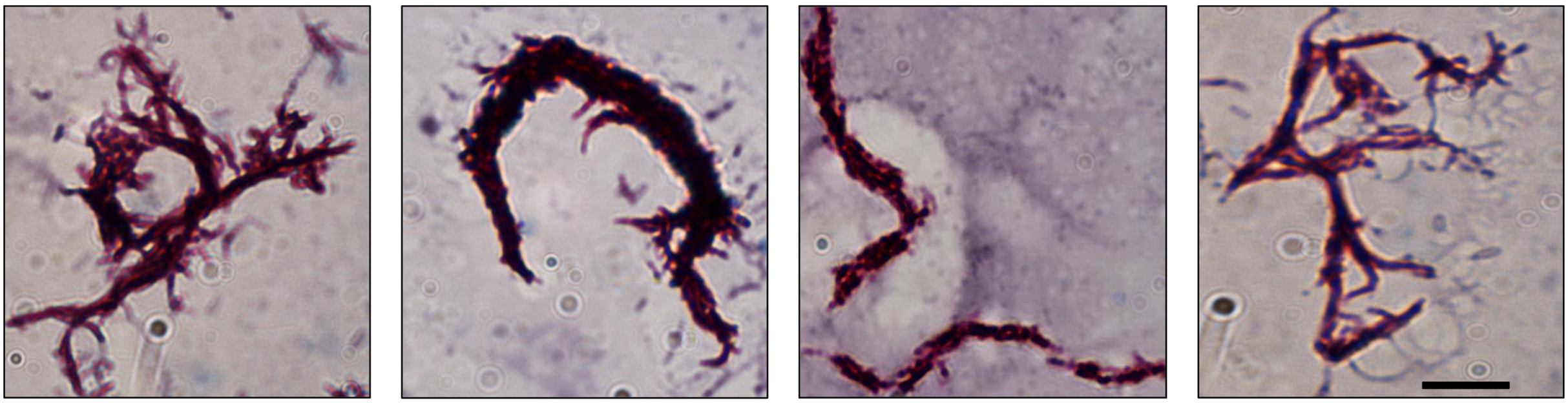
A



D

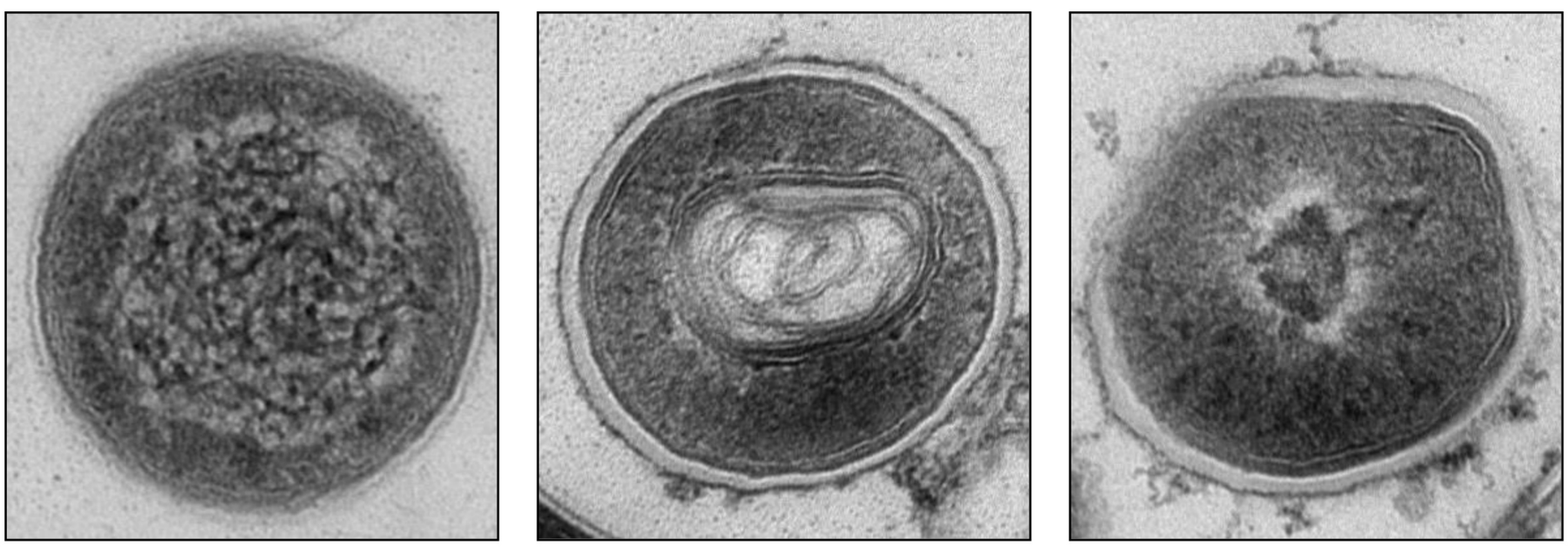
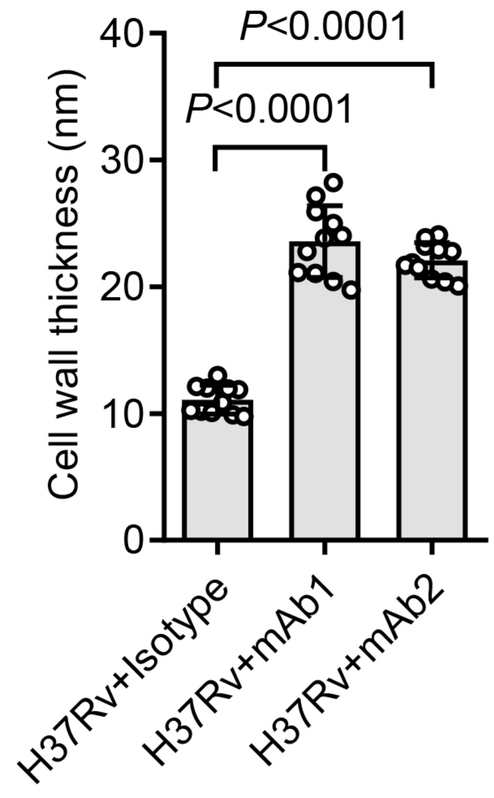


B



H37Rv+Isotype | H37Rv+ mAb1 | H37Rv+ mAb2 | H37Rv+ EMB

C



H37Rv+Isotype | H37Rv+ mAb1 | H37Rv+ mAb2