

Caseation of human tuberculosis granulomas correlates with elevated host lipid metabolism

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The progression of human tuberculosis (TB) to active disease and transmission involves the development of a caseous granuloma that cavitates and releases infectious *Mycobacterium tuberculosis* bacilli. In the current study, we exploited genome-wide microarray analysis to determine that genes for lipid sequestration and metabolism were highly expressed in caseous TB granulomas. Immunohistological analysis of these granulomas confirmed the disproportionate abundance of the proteins involved in lipid metabolism in cells surrounding the caseum; namely, adipophilin, acyl-CoA synthetase long-chain family member 1 and saposin C. Biochemical analysis of the lipid species within the caseum identified cholesterol, cholesteryl esters, triacylglycerols and lactosylceramide, which implicated low-density lipoprotein-derived lipids as the most likely source. *M. tuberculosis* infection *in vitro* induced lipid droplet formation in murine and human macrophages. Furthermore, the *M. tuberculosis* cell wall lipid, trehalose dimycolate, induced a strong granulomatous response in mice, which was accompanied by foam cell formation. These results provide molecular and biochemical evidence that the development of the human TB granuloma to caseation correlates with pathogen-mediated dysregulation of host lipid metabolism.

INTRODUCTION

Despite advances in modern medicine, tuberculosis (TB) still remains one of the most deadly infectious diseases with about 8 million infected individuals progressing to active disease and about 2 million people dying of TB annually (Dye et al, 2008). The synergy between HIV and *Mycobacterium tuberculosis* (Mtb)

infections is a cause for considerable concern, and TB has emerged as the diagnostic infection for the progression to acquired immunity deficiency syndrome (AIDS) in sub-Saharan Africa.

Shortly after Mtb infection, the tissue site becomes organized into a granuloma, which comprises of a core of infected macrophages surrounded by foamy and epithelioid macrophages, monocytes and multinucleated giant cells (MGCs) (Russell, 2007).

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The periphery of the granuloma is marked by fibroblasts that lay down a fibrous capsule around the macrophage-rich centre. Once the capsule has formed, lymphocytes are scarce within the granuloma centre and are most abundant at the periphery of granuloma. In this state, the granuloma represents a stable structure that maintains a fairly constant bacterial load but does not cause any overt signs of disease in the infected individual. Progression of the latent TB towards active disease correlates with the increased accumulation of caseum in granulomas, which is thought to result from the death of macrophages in the granuloma centre (Dannenberg, 1994; Dannenberg & Sugimoto, 1976; Russell et al, 2009). The granuloma becomes increasingly necrotic until the centre liquefies and ruptures into the lung airway. At this late stage, many of the Mtb bacilli are extracellular and free bacilli are released into the airways to be exhaled into the atmosphere, thus facilitating spread of the infection.

Progression of disease is determined locally at the level of the granuloma; however, the factors that lead to granuloma development remain to be determined. Progression to active disease is almost universally regarded as a 'failure' of the host immune system to control the infection. However, this is a 'host-centered' view that minimizes Mtb's ability to manipulate the host response locally and to induce the damage that is required for the pathogen to complete its life cycle. Recent data on granuloma model systems have revealed how Mtb is extremely adept at inducing responses that lead to extensive tissue remodelling and pathology (Actor et al, 2001; Geisel et al, 2005; Karakousis et al, 2004; Puissegur et al, 2004; Rhoades et al, 2005). These data indicate that Mtb actively modulates the localized tissue response throughout the course of infection. Despite the critical importance of understanding the changes in tissue metabolism within TB granuloma, few studies have looked beyond immune mechanisms to probe the underlying pathology required to generate active disease.

In the current study, we exploited laser capture microdissection (LCM) and genome-wide microarray analysis to investigate human TB granulomas. Amongst several physiological changes revealed by transcriptional profiling, we noted that host genes for lipid sequestration and metabolism, namely, adipophilin (*ADFP*), acyl Co-A synthase long chain family member 1 (*ACSL1*) and prosaposin (*PSAP*), were upregulated in caseous human pulmonary TB granulomas. Immunohistological analysis of human TB granulomas confirmed the disproportionate abundance of ADFP, ACSL1 and the PSAP product, saposin C (SapC), in cells surrounding the caseum. Biochemical analysis of the major lipid species within the caseum revealed an abundance of cholesterol ester (CE), cholesterol (CHO), and triacylglycerol (TAG) and also a high level of lactosylceramide (LacCer), which is known to influence both CHO sequestration and cell death (Chatterjee et al, 1997; Garner et al, 2002). Mtb bacilli induced similar lipid sequestration and foam cell formation in murine and human macrophages *in vitro*. Significantly, using a murine granuloma model we developed previously (Rhoades et al, 2005), we observed that the Mtb cell wall lipid trehalose dimycolate (TDM) injected with Matrigel into the subcutis of mice induced a strong granulomatous response and extensive foam cell formation.

The data presented here provide compelling evidence that the development of the lipid-rich caseum in the human TB granuloma correlates with a re-alignment in host lipid metabolism within the granuloma. The induction of a similar response by the bacterial lipid TDM implies that this is a pathogen-driven response, which results in a metabolic shift that induces the pathology necessary for the pathogen to complete its life cycle.

RESULTS

Transcriptional profiling of caseous human pulmonary TB granulomas

Previous studies of the human response to Mtb infection at the molecular level have examined the gene expression profiles of bronchoalveolar lavage fluid (BALF) samples, peripheral blood mononuclear cells (PBMCs), human cell lines or human TB lesions by using quantitative real-time RT-PCR and/or microarray for a subset of genes (Grassi et al, 2006; Kim et al, 2006; Ragno et al, 2001; Ulrichs & Kaufmann, 2006; Volpe et al, 2006; Zhu et al, 2003). Although these studies have provided valuable information, we feel that characterization of global gene expression within human pulmonary TB granulomas is required to further our appreciation of the disease process. In the current study, lung tissues were surgically excised from TB patients with extensive lung cavitation and tissue degeneration, and LCM was employed to dissect out granulomas, excluding uninvolved lung tissue. We specifically selected caseous granulomas for this study because accumulation of caseum is a histologically defined process that is correlative of granuloma liquefaction and cavitation, which leads ultimately to transmission (Fig S1) (Dannenberg, 1994; Dannenberg & Sugimoto, 1976). Total ribonucleic acids (RNAs) were extracted from LCM samples, and messenger RNAs (mRNAs) were amplified, biotin-labelled and hybridized on to genome-wide Human X3P array that is specifically designed based on 200 base pairs towards 3' end of mRNA.

We analyzed the array data in two different ways. First, selection of an appropriate control is problematic because the TB granuloma is strikingly different from uninfected lung parenchyma and contains a markedly different set of cell lineages; therefore the comparison between infected and uninfected tissue is of debatable value. In addition, there is no justification to biopsy healthy lung tissue, so we are limited to uninvolved lung tissue from TB patients. Nonetheless, we did generate an array profile from 'normal', uninvolved lung parenchyma to include as a control and used this to produce a relative list of transcripts from caseous granulomas compared to the normal lung tissue sample (GEO Accession# GSE20050, Table SII). Second, we generated a hierarchical list of absolute transcript abundance from caseous granulomas (GEO Accession# GSE20050, Table SI), in case the comparative list generated misleading results. Fortunately, both the comparative and hierarchical array lists showed very similar rankings of transcript abundance.

Preliminary examination of granuloma-associated abundant transcripts identified gene subsets involved in a number of

destructive tissue pathologies such as invasive cancers and metabolic diseases, suggesting overlap in pathological mechanisms (Bild et al, 2006; Ma et al, 2009; Tuomisto & Yla-Herttuala, 2005). However, we focused on lipid metabolism as previous reports have documented the abundance of lipids in Mtb-infected host cells in mice and human (D'Avila et al, 2006; Hunter et al, 2007; Kondo & Kanai, 1976; Kondo & Murohashi, 1971; Kondo et al, 1970; Peyron et al, 2008; Russell et al, 2009). We noted that many of the genes encoding enzymes involved in lipid catabolism and synthesis were markedly upregulated in the transcriptome of the TB granuloma (Table 1) and generated an Ingenuity[®] metabolism plot that was centered around this extensive array of lipid-processing enzymes (Fig 1). The upregulated genes involved in lipid metabolism are interconnected and many of them are unified around the pro-inflammatory cytokine tumour necrosis factor- α (TNF- α), indicating that this response is consistent with a sustained pro-inflammatory insult.

Localization of proteins associated with lipid synthesis and sequestration in human TB granulomas

We decided to characterize in greater depth the proteins encoded by three highly abundant transcripts relevant to three different aspects of the lipid sequestration and metabolism themes revealed in Fig 1. The selected genes encode ADFP, ACSL1 and PSAP, the precursor of SapC (Table 1). These three proteins play different, but key roles in the intracellular processing and sequestration of lipids. ADFP is known to be required for lipid droplet synthesis and remains strongly associated with these droplets following their formation (Robenek et al 2005). Upregulation of ADFP expression increases sequestration of CE, increases synthesis of long-chain fatty acid (LCFA) and TAG, and inhibits catalysis of lipid through the β -oxidation pathway (Bickel et al, 2009; Chang & Chan, 2007; Larigauderie et al, 2004). ACSL1 can also lead to the lipid droplet formation through the *de novo* synthesis of LCFAs that are incorporated into TAG (Parkes et al, 2006; Soupené &

Table 1. Genes encoding enzymes involved in lipid metabolism

Gene	Description	Hierarchical signal intensity ^a	Fold change ^b	Function
<i>ABHD5</i>	Abhydrolase domain containing 5	9.6	34.3	Triacylglycerol metabolism
<i>ACLY</i>	ATP citrate lyase	12.4	22.7	Acetyl-CoA synthesis
<i>ACSL1</i>	Acyl-CoA synthetase long chain fatty acid family member 1	12.7	39.4	Fatty acyl-CoA ester synthesis
<i>ACSL3</i>	Acyl-CoA synthetase long chain fatty acid family member 3	11.1	71.2	Fatty acyl-CoA ester synthesis
<i>ACSL4</i>	Acyl-CoA synthetase long chain fatty acid family member 4	10.5	261.5	Fatty acyl-CoA ester synthesis
<i>ACSL5</i>	Acyl-CoA synthetase long chain fatty acid family member 5	11.7	75.5	Fatty acyl-CoA ester synthesis
<i>ADFP</i>	Adipose differentiation-related protein	13.1	19.1	Lipid accumulation
<i>CRAT</i>	Carnitine <i>O</i> -acetyltransferase	10.6	14.8	Acyl-CoA/CoA regulation
<i>CYP11B1</i>	Cytochrome P450, family 1, subfamily B, polypeptide 1	13.6	36.5	Sterol synthesis
<i>CYP27A1</i>	Cytochrome P450, family 27, subfamily A, polypeptide 1	12.6	48.0	Sterol synthesis
<i>DEGS1</i>	Degenerative spermatocyte homologue 1, lipid desaturase	11.8	180.2	Unsaturated fatty acid synthesis
<i>DHCR7</i>	7-Dehydrocholesterol reductase	10.0	8.8	Cholesterol synthesis
<i>EBP</i>	Emopamil binding protein	10.4	18.3	Cholesterol synthesis
<i>ELOVL5</i>	Elovl family member 5, elongation of long chain fatty acids	12.4	36.8	Fatty acid synthesis
<i>FADS1</i>	Fatty acid desaturase 1	12.2	45.0	Fatty acid synthesis
<i>FDPS</i>	Farnesyl diphosphate synthase	11.2	8.6	Cholesterol synthesis
<i>GBA</i>	Glucosidase, beta, acid	11.6	32.0	Sphingolipid metabolism
<i>GLA</i>	Galactosidase, alpha	13.0	152.5	Sphingolipid metabolism
<i>GLB1</i>	Galactosidase, beta 1	11.4	19.9	Sphingolipid metabolism
<i>GPD2</i>	Glycerol-3-phosphate dehydrogenase 2	10.2	60.9	Glycerophospholipid metabolism
<i>HADHA</i>	Hydroxyacyl-Coenzyme A dehydrogenase/3-ketoacyl-Coenzyme A thiolase/enoyl-Coenzyme A hydratase, alpha subunit	11.0	16.7	Fatty acid metabolism
<i>HMGR</i>	3-Hydroxy-3-methylglutaryl-Coenzyme A reductase	8.9	38.4	Cholesterol synthesis
<i>IDI1</i>	Isopentenyl-diphosphate delta isomerase 1	10.9	7.3	Cholesterol synthesis
<i>LIPA</i>	Lipase A, lysosomal acid, cholesterol esterase	12.6	209.6	Sterol metabolism
<i>LSS</i>	Lanosterol synthase	11.0	8.1	Cholesterol synthesis
<i>PLSCR1</i>	Phospholipid scramblase 1	11.8	137.1	Regulation of lipid accumulation
<i>PSAP</i>	Prosaposin (precursor of sapC)	14.6	30.0	Sphingolipid metabolism
<i>SCD</i>	Stearoyl-CoA desaturase	13.1	76.8	Fatty acid synthesis
<i>SC5DL</i>	Sterol-C5-desaturase	10.1	18.7	Cholesterol synthesis
<i>SOAT1</i>	Sterol <i>O</i> -acyltransferase 1	10.3	47.2	Cholesterol esterification
<i>SPHK2</i>	Sphingosine kinase 2	11.6	32.1	Sphingolipid metabolism
<i>TPI1</i>	Triosephosphate isomerase 1	12.4	10.9	Fatty acid synthesis

^aGene expression profiles from three independent caseous human TB granulomas were averaged, and the hierarchy of the absolute expression levels was generated.

^bGene expression profiles from three independent caseous human TB granulomas were averaged, and then compared to that of normal lung tissue sample. The upregulation of selected genes were statistically significant ($P < 0.05$).

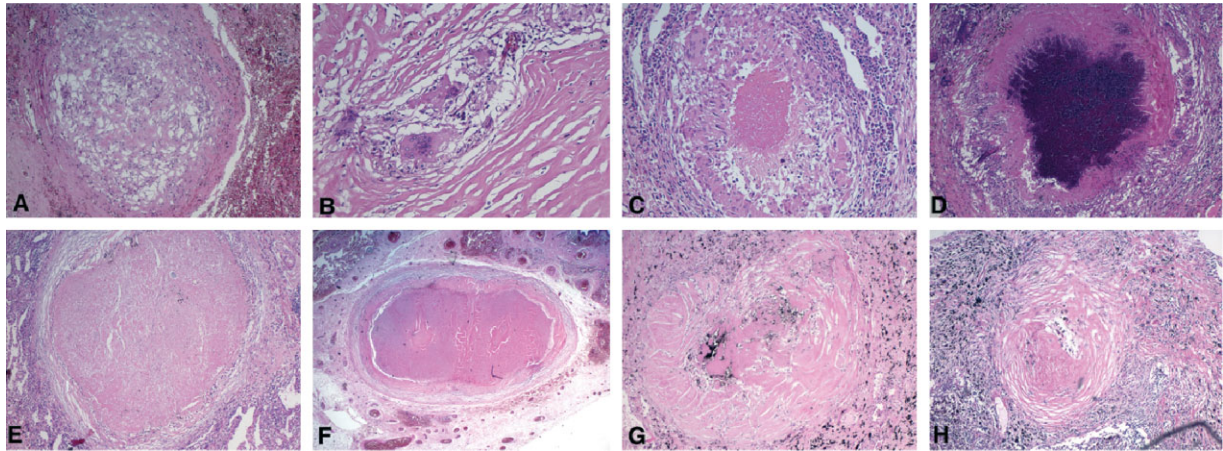


Figure 2. Definition of the Stages of Human TB lung Granulomas. Human TB lung granulomas were categorized into stages as illustrated by the following examples.

- A, B. nascent granuloma (A) ($\times 40$) and nascent granuloma (B) ($\times 100$).
- C, D. caseous granuloma (C) ($\times 100$) and caseous granuloma (D) ($\times 40$).
- E, F. fibrocaseous granulomas ($\times 40$), and
- G, H. resolved granulomas ($\times 40$).

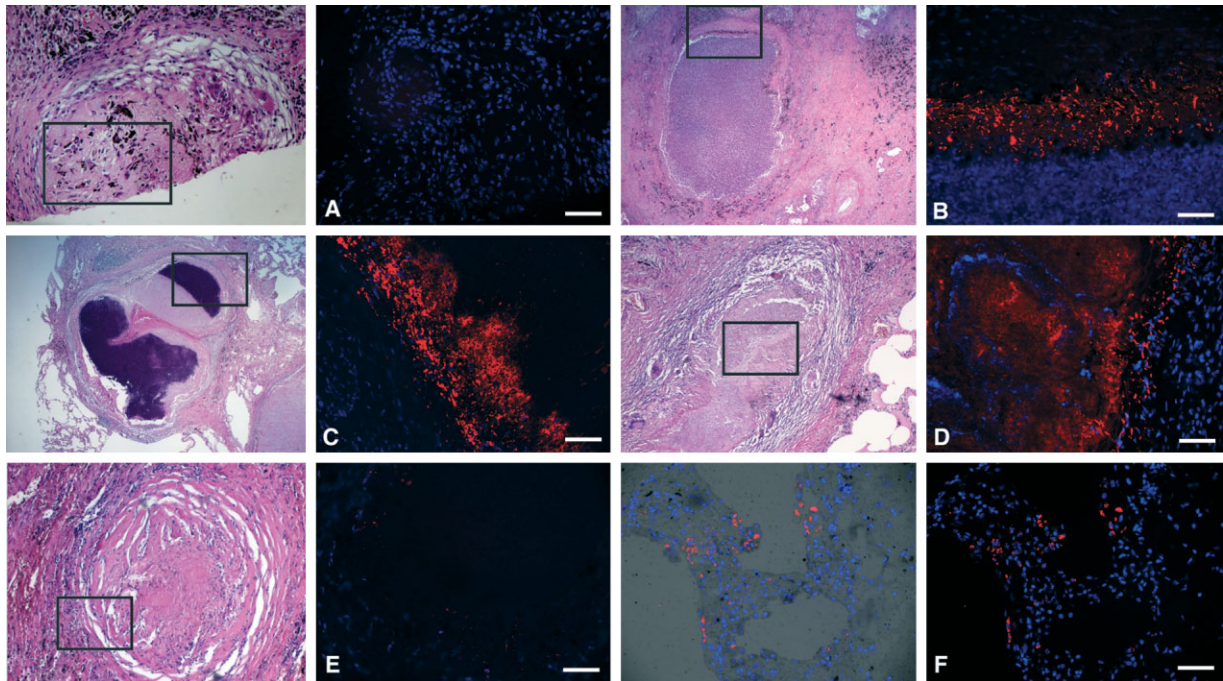


Figure 3. ADFP expression in human TB granulomas. Immunofluorescence signals were obtained for each granuloma, and a representative image (right) and the corresponding region from an H&E stained slide (left, boxed) are shown. Nuclei are seen in blue and antigens in red.

- A. Nascent granulomas exhibit weak ADFP expression.
- B. Caseous granulomas stain strongly for ADFP.
- C. Fibrocaseous granulomas also label strongly for ADFP expression.
- D. The caseous center also has intense ADFP expression, together with nuclear debris.
- E. Resolved granulomas exhibit low levels of ADFP labeling.
- F. Control normal lung parenchyma shows ADFP expression in pneumocytes and alveolar macrophages (the left image is a merged image with bright field). Scale bar is 50 μm .

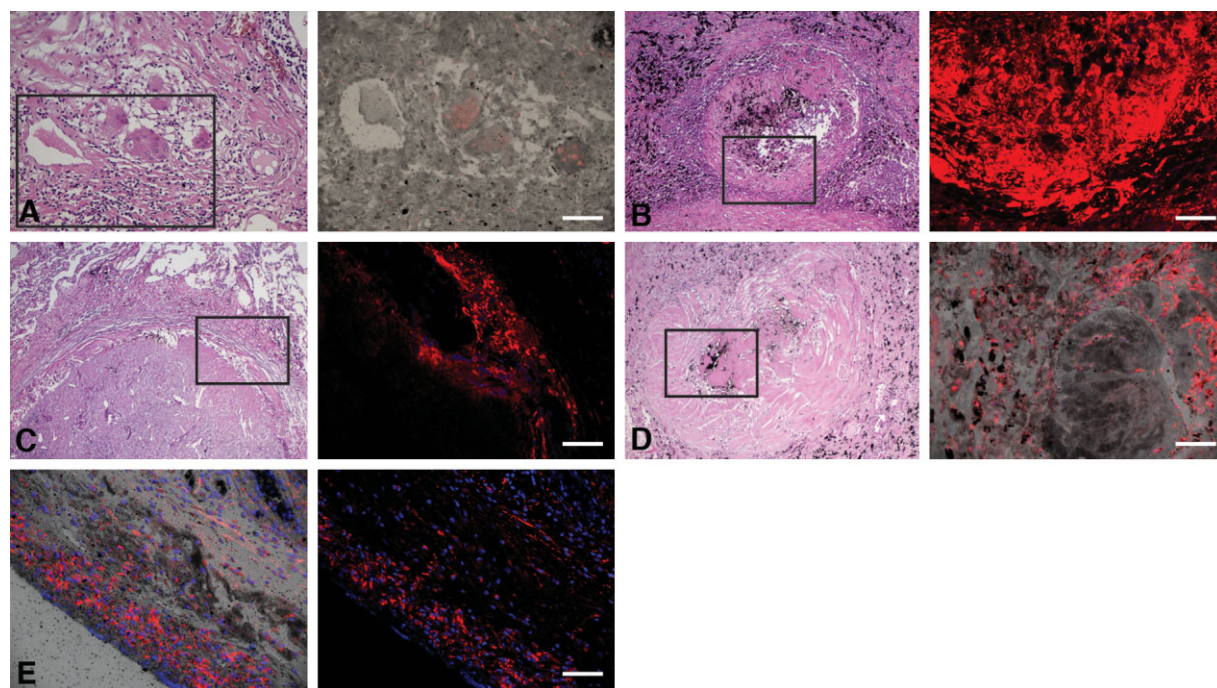


Figure 4. ACSL1 expression in human TB granulomas. Immunofluorescence signals were obtained for each granuloma, and a representative image (right) and the corresponding area from an H&E stained slide (left, boxed) are shown. Nuclei are seen in blue and antigens in red.

A, D. ACSL1 expression is weak in nascent granulomas (**A**) and in resolved granulomas (**D**).

B, C. ACSL1 is strongly expressed in caseous (**B**) and fibrocaceous granulomas (**C**).

E. Normal lung parenchyma shows ACSL1 expression (the left is a merged image with bright field). Scale bar is 50 μ m.

exception of a few cells (Fig 4D). ACSL1 is important for normal cell functions; thereby it was detectable in normal lung parenchyma (Fig 4E). SapC expression differed slightly from ADFP and ACSL1 in that it was strongly expressed in macro-

phages and MGCs even in nascent granulomas (Fig 5A). In caseous and fibrocaceous granulomas, SapC was expressed along the capsule that contained many macrophages and MGCs (Fig 5B and C). Similarly to ADFP and ACSL1, SapC expression

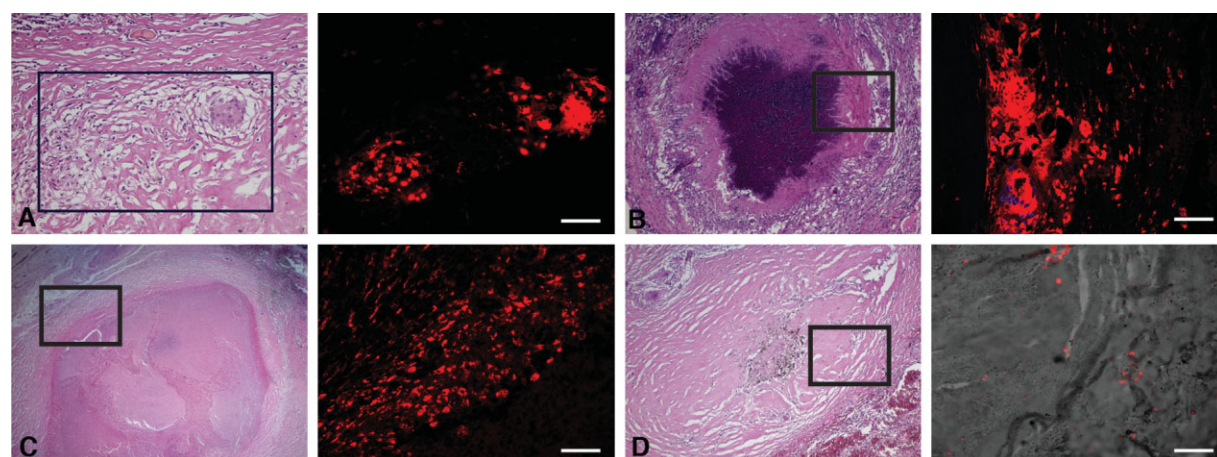


Figure 5. SapC expression in human TB granulomas. Immunofluorescence signals were obtained for each granuloma, and a representative image (right) and the corresponding area from an H&E stained slide (left, boxed) are shown. Nuclei are seen in blue and antigens in red.

A, B, C. SapC is expressed in nascent granuloma (**A**), caseous granulomas (**B**), and fibrocaceous granulomas (**C**).

D. Resolved granulomas show weak expression. Scale bar is 50 μ m.

was very weak or not detected in resolved granulomas (Fig 5D). There was no detectable level of SapC expression in normal lung parenchyma (data not shown).

The enhanced expression of these proteins, most notably ADFP, in the macrophage-rich centre of the granuloma is indicative of lipid sequestration and foam cell formation. We examined cryosections of human pulmonary TB tissues by staining neutral lipids with Oil Red O and observed lipid-laden foam cells in both respiratory bronchioles and caseous granulomas (Fig S2A–D).

The immunohistological analysis of human TB granulomas enables us to form a more dynamic interpretation of granuloma biology. Both ADFP and ACSL1 are indicative of lipid accumulation through either sequestration or *de novo* synthesis. The paucity of ADFP and ACSL1 expression in nascent granulomas, coupled with their increased expression upon acquisition of the lipid-laden caseum in caseous and fibrocaceous granulomas, suggests a functional linkage.

Analysis of caseum: neutral lipids and lactosylceramide

The lipid accumulation in *Mtb*-infected tissues has been reported in studies dating back to early last century (Caldwell,

1919; Kondo & Murohashi, 1971; Kondo et al, 1970; Pagel & Pagel, 1925). Moreover, our data indicate that it is a re-alignment in host metabolism that is the most likely explanation for lipid accumulation. To test this hypothesis, we sought to determine the identity of the major lipid species present in the caseum from human pulmonary TB granulomas.

We extracted total lipids from the caseum of caseous or fibrocaceous human pulmonary TB granulomas and from uninvolved lung parenchyma and analyzed them by thin-layer chromatography (TLC). There were noticeable differences in lipid profiles; the caseum had markedly increased amounts of lipids that co-migrated with the CE, CHO and TAG standards, when compared to lipids isolated from normal lung tissues (Fig 6A). The identities of CE, CHO and TAG were confirmed by mass spectrometric analysis. Electron impact gas chromatography/mass spectrometry (EI-MS) and total ion current (TIC) chromatogram generated a peak at 12.23 min and an EI mass spectrum that confirmed the presence of CHO (Fig S3A and B). The tandem mass spectrum contained the major $[M + Na]^+$ ions at m/z 671.5 and m/z 673.8, corresponding to CE (Fig S3C). The dominating $[M + NH_4]^+$ ions were at m/z 876.8 corresponding to TAG possessing a total of C_{52} with the presence of two total

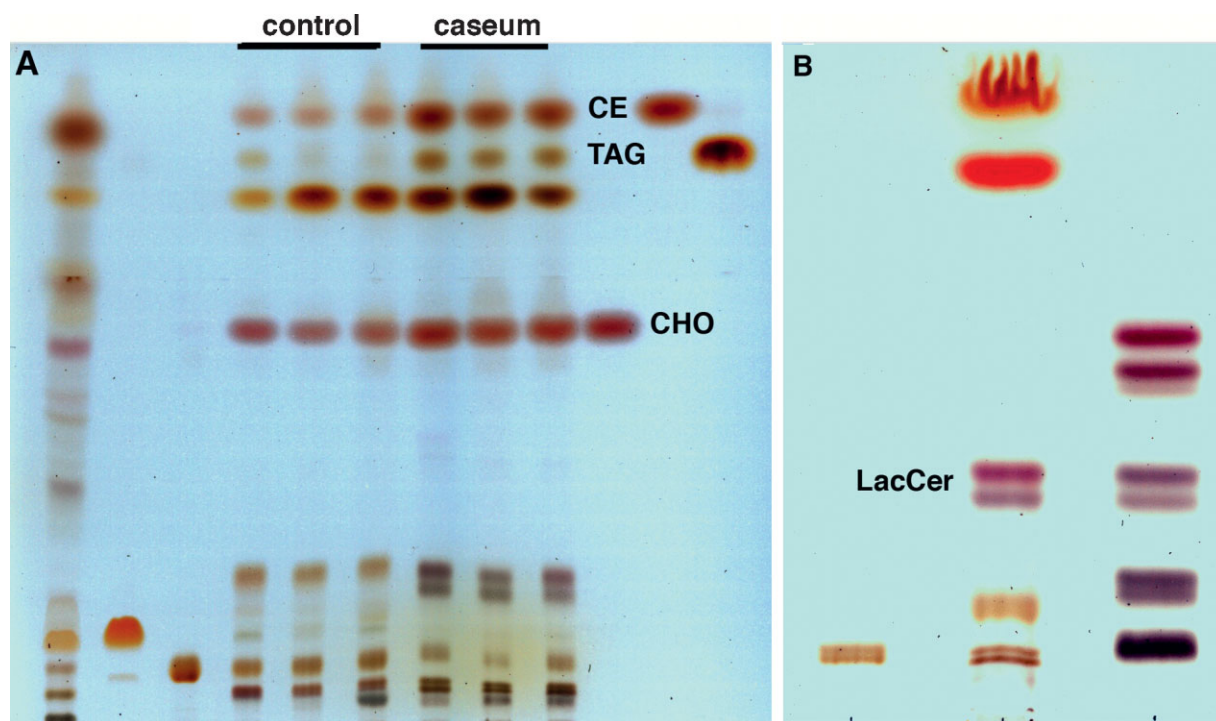


Figure 6. Thin-layer chromatographic analysis of lipids from the caseum and normal lung tissues.

- A.** Dual solvent separation of granuloma lipids by TLC. Total lipids were extracted from normal lung tissues (lane 4–6, 120 μ g) and caseous TB lung granulomas (lane 7–9, 120 μ g), and then equivalent amounts were run in parallel with total lipids from *Mtb* CDC1551 (lane 1, 400 μ g), and standard lipids, cardiolipin (lane 2, 50 μ g), phosphatidylcholine (lane 3, 50 μ g), CHOL (lane 10, 20 μ g), CE (lane 11, 25 μ g), TAG (lane 12, 20 μ g). The lipids from the TB caseum exhibit an abundance of CHOL, CE, and TAG.
- B.** Single solvent separation of lipid species with similar migration to the sphingolipids, illustrating the enrichment of LacCer in the caseum extract. Total lipids from the caseum (lane 2, 120 μ g) were run together with sphingomyelin standard (lane 1, 5 μ g) and neutral glycosphingolipids standard (lane 3, 30 μ g). Caseous and fibrocaceous granuloma samples were derived from seven independent tissue samples, and normal lung tissue samples.

unsaturated bonds situated on the fatty acyl chains (52:2) (Fig S3D).

In addition to these neutral lipids, we observed lipids near the origin that migrated similarly to glycosphingolipids (Fig 6B). Among glycosphingolipids, we detected a unique double band that was identified as LacCer, and ESI-MS analysis revealed $[M+Na]^+$ ions at m/z 884.6 and m/z 994.8, corresponding to d18:1/16:0-LacCer and d18:1/24:1-LacCer, respectively (Fig S3E).

These data support the hypothesis that there is a functional correlation between the major lipid species in the caseum and the upregulated transcripts and proteins (Fig 1). Foamy macrophages acquire much of their lipids from low-density lipoproteins (LDLs), which contains CHO, TAG and glycosphingolipids. The CHO becomes esterified to CE during incorporation into lipid droplets, which are formed through the activity of ADFP. Therefore, the presence of CE implicates passage through a macrophage prior to delivery to the caseum.

Mtb-infected macrophages exhibit upregulation of lipid synthesis and sequestration pathways

It has been reported previously that Mtb infection induces lipid droplet formation *in vitro* (Kondo & Kanai, 1976; Peyron et al, 2008). To determine if this process was associated with increased expression of *ADFP*, *ACSL1* and *PSAP* (SapC), we infected murine bone marrow-derived macrophages (BMM Φ) and human PBMC-derived macrophages with Mtb. Quantitative real-time RT-PCR (Table 2) analysis revealed increased expression of *ADFP*, *ACSL1* and *PSAP* (SapC) in infected-macrophages. Mtb-infected cells demonstrated lipid droplet formation concomitant with the expression of *ADFP*, *ACSL1* and *SapC*, revealed by confocal microscopy (Fig 7A–F). The upregulation of these genes was modest in the human PBMC-derived macrophages, however we observed lipid droplet formation even in uninfected human cells (Fig 7G) suggesting that they were already pre-disposed to accumulate lipids.

TDM, a strong inducer of foam cells in mice

Recently it was reported that mycolic acids from the cell wall of Mtb were strong inducers of foamy macrophage formation *in vitro* (Korf et al, 2005; Peyron et al, 2008). We had shown previously that Mtb cell wall lipids are released in copious

amounts and traffic out of infected macrophages (Beatty et al, 2000, 2001). We had also demonstrated that amongst these lipids, TDM was the most potent inducer of a granulomatous response and would be an ideal candidate for triggering the pathological response at the infection site (Geisel et al, 2005; Hunter et al, 2006a,b).

To determine whether or not this granulomatous response was accompanied by lipid sequestration, we used the granuloma model we had developed previously (Geisel et al, 2005; Rhoades et al, 2005). Polystyrene beads coated with TDM from Mtb were suspended in Matrigel and inoculated into mice subcutaneously. Murine cells invade the matrix and remodel the site under the influence of the bacterial lipid. Histology of the tissue site revealed an abundance of ADFP-expressing foam cells clustering around the inoculated beads (Fig 8A and B). Cryosections of TDM-induced granuloma also showed an abundance of lipid droplets in foam cells (Fig 8C). Control lipid phosphatidylglycerol (PG)-coated beads recruited very few cells and did not induce foam cell formation (Fig 8D and E). *Acs11* was also detectable in the cells recruited to the TDM-induced granuloma; however, we did not observe detectable levels of *SapC* expression (data not shown). These data indicate, not surprisingly, that the Mtb lipid TDM is capable of inducing some, but not all, of the features found in the human TB granuloma.

DISCUSSION

The human TB granuloma is a complex structure that has evolved to enable both the host and the pathogen to survive and replicate. However, unlike many chronic infectious agents, Mtb actually promotes the development of a robust, systemic immune response, which it subverts at the localized tissue-level to facilitate its prolonged survival, and, ultimately facilitate its transmission (Russell, 2007). The TB granuloma plays a pivotal role in TB; yet, our knowledge of what determines containment versus active disease is superficial.

In the present study, we sought to elucidate tissue changes that accompany granuloma progression through LCM and microarray analysis of caseous granulomas from human TB patients. Because of the caveats associated with the use of uninfected lung tissue as a control, we generated both hierarchical and comparative lists of transcript abundance; the overall transcriptional profiles from both methods were strikingly similar. One of the interesting themes to emerge from the microarray data was the apparent re-alignment of lipid metabolism. We chose to focus on the proteins ADFP, *ACSL1* and *SapC* as three key representatives of the lipid-modulating pathways that are impacted by the proinflammatory cytokine TNF- α (Fig 1). ADFP is a member of PAT (perilipin, ADRP and tail-interacting protein of 47 kDa) domain proteins and is present on the surface and the core of intracellular lipid droplets formed in foam cells. Its mRNA level is increased by excess LCFAs, fasting, modified LDL, peroxisome proliferator-activated receptor (PPAR) agonists or transcription factors that play a role in lipid homeostasis (Dalen et al, 2006; Edvardsson et al, 2006;

Table 2. Expressions of *ADFP*, *ACSL1* and *PSAP* (*SapC*) are induced by Mtb infection

Gene	Fold induction	
	Murine BMM Φ	Human PBMM
<i>ADFP</i>	2.95	1.22
<i>ACSL1</i>	32.67	1.86
<i>SapC</i>	2.28	1.37

Murine and human macrophages were infected with Mtb H37Rv and mRNA levels of *ADFP*, *ACSL1* and *PSAP* (*SapC*) were determined by quantitative real-time RT-PCR. Three technical replicates were included for all conditions and genes. The fold change was calculated using the formula, $2^{-\Delta\Delta C_t}$, and the average of replicates is shown here.

Gao & Serrero, 1999; Larigauderie et al, 2004; Wang et al, 1999; Wei et al, 2005). In the current study, ADFP was detected in the cells subtending the caseum in caseous and fibrocaseous human pulmonary TB granulomas. ADFP was also observed in newly caseating granulomas, mingled with nuclear debris, implying that the protein was associated with lipid droplets that were released by cells undergoing necrotic or apoptotic death (Fig 3D). ACSL1 and SapC were also strongly expressed in caseous and fibrocaseous human pulmonary TB granulomas. ACSL1 mediates the formation of fatty acyl-CoA esters from fatty acids (preferably chain length of 12–20 carbons), which is a key step for lipid biosynthesis and fatty acid degradation, and is also found associated with lipid droplets (Soupene & Kuypers, 2008).

SapC aids the degradation of glycosphingolipids into ceramide, and sphingosine in acidic compartments, which is critical in maintaining the turn-over of cellular components (Hannun & Obeid, 1995; Kolter & Sandhoff, 2005; Qi et al, 1994; Schuette et al, 2001). In addition to glycosphingolipid metabolism, SapC is also known to transfer mycobacterial lipid antigen from intralysosomal membrane to CD1b, thereby activating the antigen specific T cells (Winau et al, 2004). Our hypothesis that the TB granuloma undergoes a shift in lipid metabolism and accumulates host-derived lipids was supported further by the analysis of the lipid constituents of the caseous material isolated from human pulmonary TB granulomas. Compared to normal lung tissues, the caseum samples were enriched for CE, CHO,

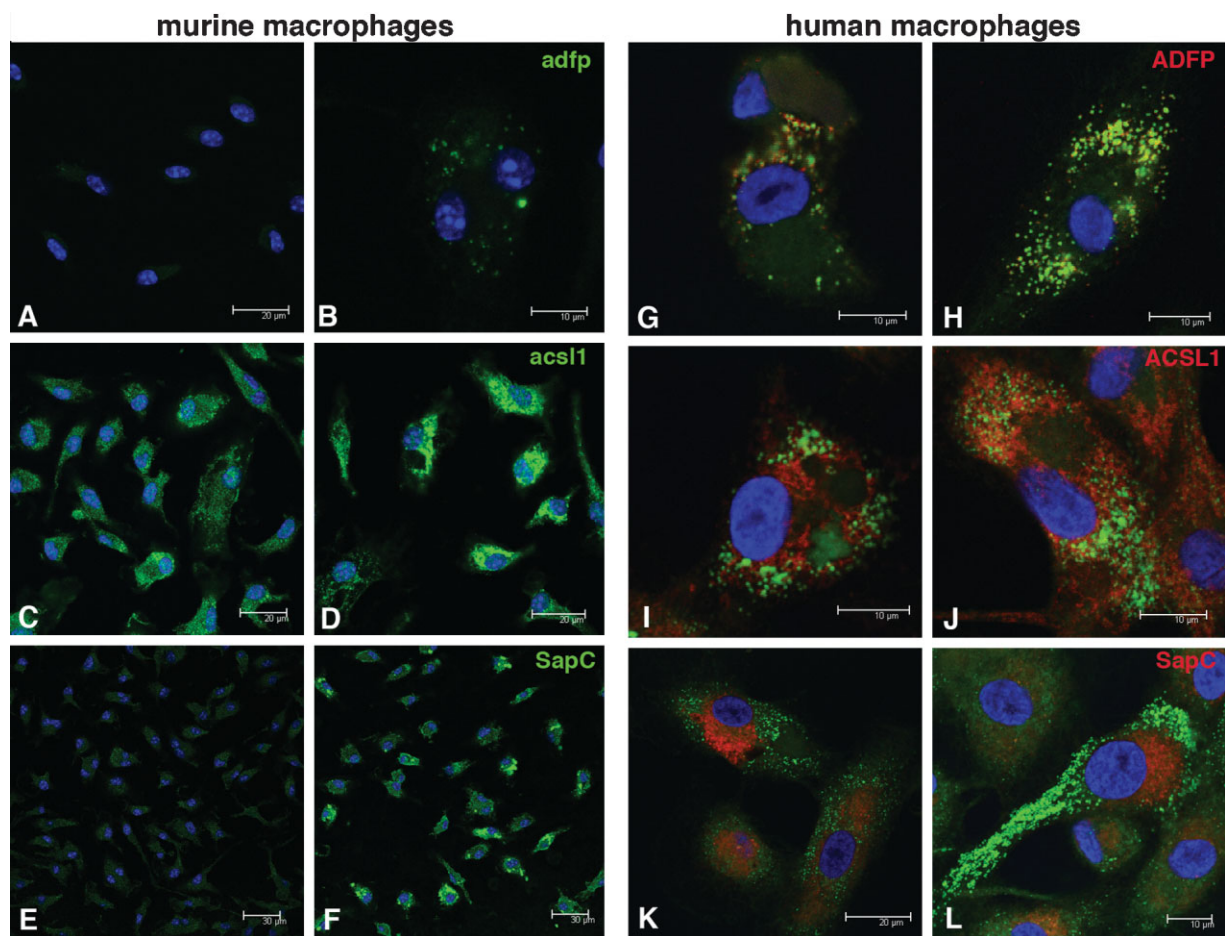


Figure 7. ADFP, ACSL1, and SapC in Mtb-infected murine and human macrophages. Murine bone marrow-derived macrophages infected with Mtb H37Rv for 24hr and examined by confocal microscopy.

A, B. Expression of Adfp in (A) control cells versus (B) Mtb-infected cells.

C, D. Expression of Acs11 in (C) control and (D) Mtb-infected macrophages. Blue, nuclei, Green: antigens.

E, F. Expression of SapC in (E) control and (F) Mtb-infected macrophages. Blue, nuclei, Green: antigens.

G, H. ADFP is expressed and is associated tightly with lipid droplets (G) control and (H) Mtb-infected macrophages. Lipid droplets were observed in both uninfected and infected human PBMC-derived macrophages.

I, J. ACSL1 is also detected in association with lipid droplets, (I) control and (J) Mtb-infected macrophages.

K, L. SapC is expressed and localizes throughout the cytoplasm (K) control and (L) Mtb-infected macrophages. Blue: nuclei, red: antigens, green: lipid droplets.

TAG and LacCer (Fig 6A and B). This profile of lipid enrichment is most consistent with the lipids coming from the lipid droplets formed in macrophages that have acquired their lipids from LDL.

The TB literature is rich with reports of lipid accumulation in Mtb infections such as lipid-laden foam cells in human TB lung tissues (Hunter et al, 2006b, 2007; Pagel & Pagel, 1925), the accumulation of CE in Mtb infected murine lungs during persistent infection (Kondo & Murohashi, 1971; Kondo et al, 1970) and the development of lipid droplets in murine or human macrophages infected with virulent Mtb or BCG (D'Avila et al, 2006, 2008; Kondo & Kanai, 1976; Peyron et al, 2008). Lipid droplets are attracting increased attention as dynamic organelles that participate in numerous biological functions linked to metabolic syndromes and diseases including atherosclerosis, diabetes, obesity, heart attack, sepsis, Chagas' disease and AIDS (Bozza et al, 2009; Desruisseaux et al, 2007; Ducharme & Bickel, 2008; Murphy, 2001). The data presented in this study provide a new hypothesis for the alterations in tissue physiology that accompany the progression of the human TB granuloma to caseation.

Cholesterol has long been linked to TB; however, data on role(s) of the sterol in disease progression or control are contradictory. The administration of CHO-rich diets to TB patients reduced the number of bacilli in the sputum (Deniz et al, 2007; Kozarevic et al, 1981; Perez-Guzman et al, 2005; Taylor & Bamgboye, 1979). However, hypercholesterolemic mice through deletion of *ApoE* showed increased Mtb growth and exacerbated lung pathology (Martens et al, 2008). This is consistent with the observations of Pandey (Pandey & Sasseti, 2008) that CHO supports the growth of persistent Mtb, and the findings in this current study that the local tissue response to Mtb infection is tightly associated with lipid accumulation. Garton *et al* reported that Mtb bacilli isolated from sputum samples had TAG-rich intracellular lipophilic inclusions (ILIs) (Garton et al, 2002, 2008). Prokaryotes have the ability to accumulate lipids (Alvarez & Steinbuchel, 2002; Waltermann et al, 2005), and Mtb is known to synthesize and accumulate TAG under stress, *e.g.* starvation and hypoxia (Daniel et al, 2004; Deb et al, 2009; Sirakova et al, 2006), and encode enzymes to utilize the TAG (Deb et al, 2006; Mishra et al, 2008). Therefore, the accumulated CHO and TAG in caseum of human granulomas may represent a convenient carbon source for either intracellular or extracellular bacilli on the cavitary surface of TB granulomas. LacCer is an extremely interesting glycosphingolipid to be found in abundance in the caseum with respect to its biological properties in glycosphingolipid metabolism. It is normally present in trace amounts in mammalian cells because it is an intermediate in the glycosphingolipids metabolism (Kolter & Sandhoff, 2005, 2006). Its accumulation in cells could be the result of either increased ceramide synthase activity or the increased catabolism of complex ceramides (Chatterjee & Pandey, 2008). The upregulation of SapC within the granulomas favors the explanation that the tissue may be experiencing LacCer excess and is trying to degrade the lipid through promoting the activities of β -galactosidase and β -glucosidase, the expression of which were upregulated 20-fold and 32-fold,

respectively, in the TB granuloma (Table 1). In addition to the endogenous generation of LacCer, treatment of cells with LacCer is known to reduce CHO efflux and enhances foam cell formation (Glaros et al, 2005). Further studies need to be done to fully verify the origin(s) of CE, CHO, TAG and LacCer that accumulate in human TB granuloma caseum.

Lesions of *Mycobacterium leprae*-infected lepromatous leprosy patients show significant accumulation of oxidized phospholipids (Cruz et al, 2008). Intriguingly, several of the genes reported as upregulated in our study overlap with those observed in lepromatous leprosy lesions, including *PSAP* and *ADFP* (Cruz et al, 2008). Another study also demonstrated the accumulation of neutral lipids including CE and the localization of ADFP protein in leprosy lesions (Mattos et al, 2010). These data indicate that there are many parallels to the metabolic reprogramming induced by both Mtb and *M. leprae*. The point at which the two infection sites diverge could hinge on the formation of the fibrous capsule in the TB granuloma, which may focus and constrain the inflammatory response leading to the accumulation of the sequestered lipids as caseum. This pathology is unique to Mtb infection and precedes the liquefaction and cavitation of the lesion. In mice, this late-stage pathology was absent from experimental Mtb infections, which has led many researchers to be concerned about the validity of the mouse as an experimental model. Recently, however, Hunter and colleagues reported caseum formation in reactivation disease in antibiotic-treated mice (Hunter et al, 2007). Significantly, these murine caseous lesions exhibited a highly developed fibrous capsule comparable to that observed in fibrocaveous human granulomas.

We also have shown that Mtb infection of macrophages *in vitro* induces foam cell formation, and that inoculation of beads coated with the bacterial cell wall lipid TDM into mice caused the clustering of Adfp-expressing foam cells around the particles. These data argue strongly that lipid sequestration is a pathogen-induced response. We have demonstrated previously that mycobacterial cell wall lipid-containing vesicles, which contain TDM, are released from infected macrophages and spread the influence of the bacterium to neighbouring cells (Beatty et al, 2000; Rhoades et al, 2003; Russell et al, 2002). TDM activates macrophages by signalling through TLR2, which forms a functional complex with the scavenger receptor MARCO to bind the mycobacterial lipid (Bowdish et al, 2009). This amplification of the response from infected cells to uninfected bystander cells could explain the extreme tissue reaction observed in human TB granulomas when the bacterial density is known to be relatively low (data not shown) (Ulrichs et al, 2005).

The current study provides a new framework for appreciating the physiological responses to Mtb infection that accompany the progression of TB granulomas to caseation and cavitation, and ultimately disease transmission. We hypothesize that bacterial components, such as TDM, stimulate the host's innate immune response to enhance synthesis and/or sequestration of host lipids in the form of lipid droplets. The foamy macrophages accumulate in the macrophage-rich centre of the granuloma, where ultimately they die through either apoptosis or necrosis. This leads to the buildup of lipids as caseum, and, finally to

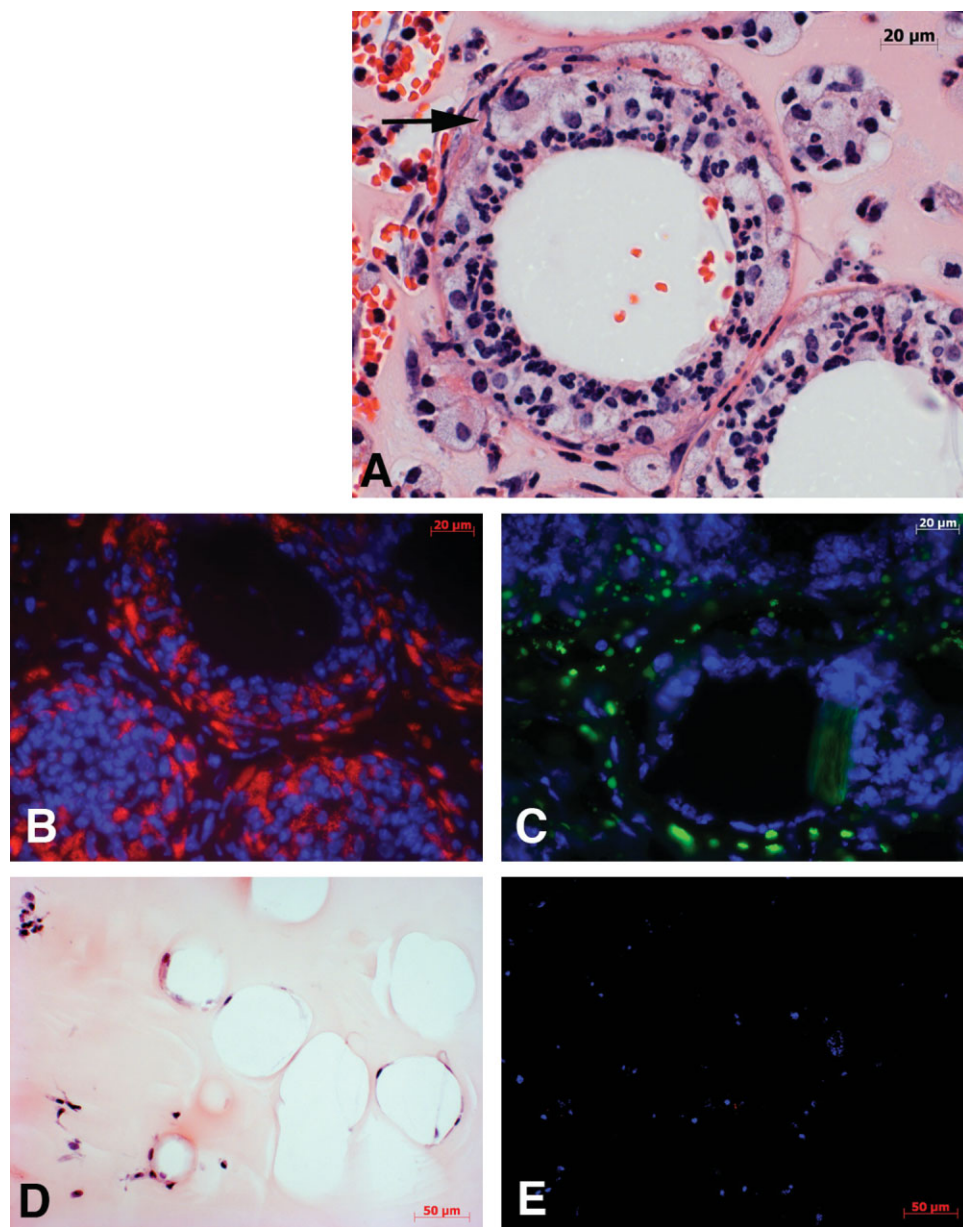


Figure 8. TDM-coated beads induce foam cell formation in mice in an experimental granuloma model. TDM-coated beads resuspended in Matrigel were subcutaneously injected to wild-type mice and the granulomas were excised at day6 post-inoculation.

- A.** Histological examination demonstrated the extensive recruitment of cells to TDM-coated beads, many of which had the appearance of lipid droplet-filled foam cells (A representative cell is marked with an arrow).
- B.** Immunohistology on the consecutive section of the same tissue block revealed a strong expression of lipid droplet-associated protein, Adfp (nuclei in blue and Adfp in red).
- C.** Lipid droplets were detected by staining neutral lipids with Nile Red on cryosections from TDM-induced granuloma (nuclei in blue and neutral lipids in green).
- D.** In contrast, control lipid PG-coated beads demonstrated considerably reduced recruitment of cells and no induction of foam cell formation (H&E).
- E.** Moreover, probing with anti-Adfp antibody on the consecutive section cut from the same tissue block as (D) did not label any of the cells surrounding the PG-coated beads (nuclei in blue and Adfp in red). The experiments were repeated three times, and three mice were used for PG- or TDM-Matrigel mixture injection.

cavitation and release of infectious Mtb bacilli (Fig 9). Our data argue that the development of novel therapeutics should encompass manipulation of the host’s local response and not be restricted to antimicrobials. Targeting fatty acid metabolism,

lipid deposition and lipid-mediated cell death in host cells are unlikely to impact on bacterial growth directly, however, they may influence the tissue response to Mtb infection and thereby modulate TB disease progression.

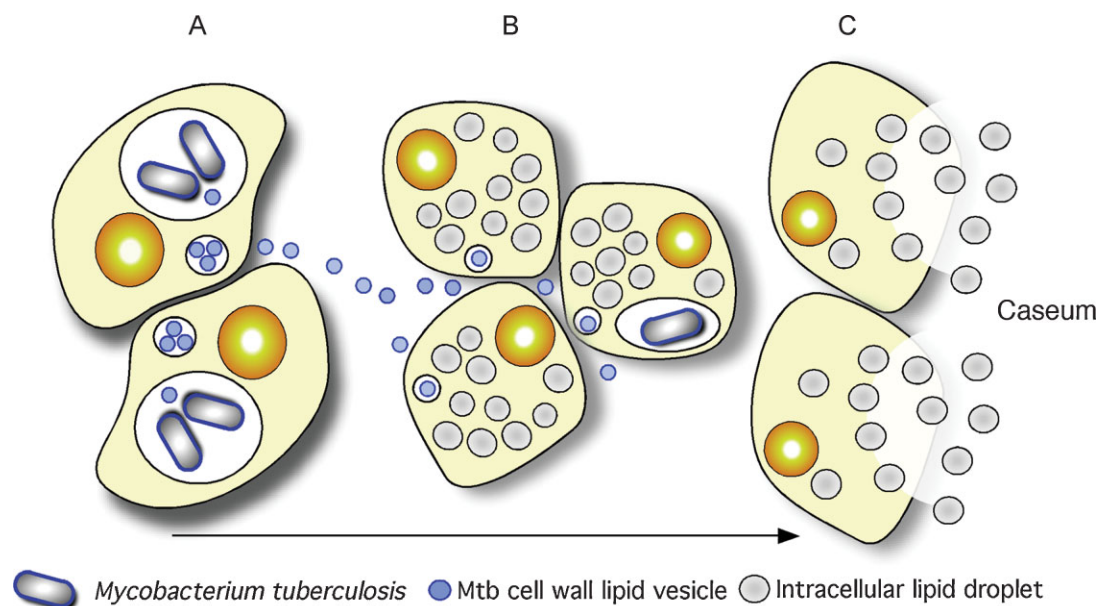


Figure 9. A model illustrating the linkages between Mtb-infection, foam cell formation and accumulation of caseum in the human TB granuloma.

- A.** Intracellular Mtb bacilli synthesize and release cell wall components inside their host cells. We have demonstrated previously that these lipids accumulate in the internal vesicles in multi-vesicular bodies, which are exocytosed from the cell in vesicular form.
- B.** Because of the release of these vesicles, both infected and uninfected macrophages are exposed to cell wall mycolates and induced to form foamy macrophages, as illustrated in Fig 7. The foamy macrophages have been shown to support the maintenance and growth of persistent bacteria.
- C.** We now propose that these cells die via an inflammatory, necrotic process and release their lipid droplets into the extracellular milieu within the granuloma. As a result of the fibrotic capsule, the human granuloma is an enclosed, isolated structure with minimal vasculature. The enclosed nature of the human granuloma leads to accumulation of necrotic debris as caseum. In this model, this process is an integral part of the pathology that leads to active disease and transmission.

MATERIALS AND METHODS

Human subjects

Human subjects were included in this study according to protocols approved by Institutional Review Boards at University of Cape Town, South Africa, the Public Health Research Institute, Newark, NJ and Cornell University, Ithaca, NY. Informed consent was obtained from all of patients.

Animals

C57BL/6J mice were obtained from the Jackson Laboratory and housed under specific-pathogen-free conditions. All experiments were approved by Cornell University Institutional Animal Care and Use Committee.

Tissue specimens

Human TB lung tissues were surgically excised from TB patients who had extensive lung cavitation and tissue degeneration, that showed little response to the antibiotics. Snap-frozen tissues were transported from Groote Schuur Hospital of Cape Town to our biosafety level-3 (BL3) laboratory. Frozen samples were embedded in OCT compound (Tissue-Tek[®]), cut into 10 μ m-thick sections at 25°C on a Cryocut 1800 cryostat (Leica Microsystems) and attached onto PET-membrane frame slides (Leica Microsystems) for LCM or Superfrost Plus Micro slides (Erie Scientific) for histological examination.

Histology of human tissues

Human TB lung tissues embedded in paraffin were cut at 3 μ m thickness and stained with hematoxylin & eosin (H&E) for histological examination. Cryosections of human TB lung tissues were fixed in 4% paraformaldehyde in PBS for 30 min and stained with H&E, and then paired to PET-membrane slides prepared for LCM. For the detection of neutral lipids, fixed cryosections were stained with Oil Red O (Alfa Aesar) and counterstained with hematoxylin.

Laser capture microdissection

Tissue sections mounted onto PET-membrane slides were fixed in graded ethanol (70, 75, 96 and 100%) containing 0.5% sodium azide for 30 s per each step. Slides were air dried briefly, and areas of interest were dissected by using Leica AS LMD system (Leica Microsystems).

Microarray

Total RNA was isolated from LCM-derived materials using PicoPure[™] RNA Isolation Kit (Arcturus) and treated with DNase using TURBO DNase-free[™] (Ambion). Purity, quality and quantity of RNA were assessed by the Agilent NanoChip Bioanalyzer assay, spectrophotometer and reverse-transcription PCR (RT-PCR) of β -actin (234 bp), glyceraldehyde-3-phosphate dehydrogenase (GAPDH, 189 bp), 18S rRNA (236 bp) and ubiquitin (198 bp). To obtain sufficient quantities of RNA for microarray, two-rounds of linear amplification of RNA were conducted using RiboAmp HS[™] Amplification Kit (Arcturus) and BioArray High Yield[™]

RNA Transcript Labeling Kit (T7) (Enzo Life Sciences, Inc.). Biotinylated, amplified antisense RNA was fragmented and hybridized onto GeneChip[®] Human X3P Array (Affymetrix), and the chip was scanned by GeneArray 300 scanner (Affymetrix). We performed three independent arrays on one LCM-derived granuloma sample, which confirmed the reliability of the technique. Thereafter, two other granulomas were further analyzed by microarray for biological replicates.

Microarray data analysis

For the generation of a hierarchical list of absolute gene expressions in caseous human TB lung granulomas, raw CEL files were analyzed at the probe level using R and Bioconductor. Only perfect match (PM) probes were included in the estimation for expression level of probe sets. Raw PM intensities from each array were subjected to local background correction using MAS5 method, \log_2 -transformed and median-centered. Informative PM probes from each probe set were selected as having the highest mean across all arrays. In this study, the mean signal of top 5 informative PM probes of each probe set was used to represent its expression level. For a list of genes differentially expressed between caseous human pulmonary TB granulomas and normal lung parenchyma, raw CHP files were summarized by MAS5 method and scaled to median of all samples, and the baseline was transformed to the median of all samples, by using GeneSpring GX 10.0.2 program (Agilent Technologies). The networks were generated through the use of Ingenuity Pathway Analysis (Ingenuity[®] Systems, www.ingenuity.com).

Statistical analysis

The statistical significance of differences in gene expressions between normal human lung tissues and caseous human TB lung granulomas was calculated using an unpaired t-test with Benjamini–Hochberg correction. Results with a *P*-value less than 0.05 were considered significant.

Culture and *in vitro* infection of macrophages

Bone marrow cells were isolated from femurs and tibias of 6- to 8-week-old female C57BL/6 mice and cultured in Dulbecco's Modified Eagle Medium (DMEM, Gibco) supplemented with 20% L-929 cell-conditioned medium, 10% heat-inactivated foetal calf serum (HyClone), 2 mM L-glutamine (Gibco), 1 mM sodium pyruvate (Gibco) and 100 U/ml penicillin-100 µg/ml streptomycin (Gibco) at 37°C, 5% CO₂ and 90% humidity for 5 days. BMMΦ were incubated with Mtb H37Rv at a multiplicity of infection 20:1 (bacteria per macrophage) for 2 h, washed with PBS and cultured DMEM medium without antibiotics.

Monocytes were isolated from human blood (Zen-Bio Inc.) by gradient separation. Briefly, blood was mixed with an equal volume of PBS, and 25 ml of this mixture was laid over 12 ml of Histopaque[®]-10771 (Sigma, density = 1.0771 g/ml). The leukocyte-containing layer was harvested, washed and cultured in DMEM medium as in BMMΦ culture, but without L-929 cell-conditioned medium. Differentiated human PBMC-derived macrophages were incubated with Mtb H37Rv at a MOI 10:1 (bacteria per MΦ) for 2 h, washed with PBS and cultured in DMEM medium without antibiotics.

Quantitative real-time RT-PCR

Infected and uninfected macrophages were lysed with TRIZOL[®] Reagent (Invitrogen) and homogenized by passing cell lysate through

a pipette. The homogenized samples were mixed with 0.2 volume of chloroform per 1 volume of TRIZOL[®] Reagent, and phases were separated by centrifugation. The RNA-containing aqueous phase was column-purified using RNEasy mini kit (Qiagen). RNA samples were treated with DNase using TURBO DNA-free[™] (Ambion). The cDNA was synthesized using iScript[™] cDNA Synthesis Kit (BIO-RAD) and used for real-time PCR using iTaq[™] SYBR[®] Green Supermix with ROX (BIO-RAD) in 7500 Fast Real-Time PCR System (Applied Biosystems). Each PCR was done in triplicate, and PCR conditions are as follows: initial activation at 95°C for 2 min and 40 cycles of denaturation at 95°C for 30 s and annealing and extension at 53°C for 45 s. Forward and reverse primers designed by using Primer3 are shown in Table SIII (Rozen & Skaletsky, 2000). The calculated threshold cycle (*C_t*) value for each gene was normalized to *C_t* of house-keeping gene 18S rRNA, and then the fold induction of gene of interest between control MΦ and Mtb-infected MΦ was calculated and presented by using the formula, $2^{-\Delta\Delta C_t}$.

Confocal microscopy of macrophages

Cells were fixed in 4% paraformaldehyde in PBS for 30 min, washed and blocked with 5% bovine serum albumin (Sigma), 1.5% glycine (VWR International) and 0.1% saponin in PBS at room temperature for 1 h. Cells were incubated with guinea pig polyclonal antibody against murine Adfp or mouse monoclonal antibody against human ADFP (1:100, RDI division of Fitzgerald Industries Intl.), rabbit polyclonal antibody against human ACSL1 (1:200, GenWay Biotech) or rabbit polyclonal antibody against human SapC (1:50, Santa Cruz Biotechnology) at 37°C for 1 h. After washing with PBS, cells were incubated with Texas Red[®]-conjugated antibodies against guinea pig, mouse or rabbit (1:250, Jackson ImmunoResearch) and BODIPY[®] 493/503 (1 µg/ml, Invitrogen) for lipid droplets, at room temperature for 1 h. After wash with PBS, the nuclei were stained with 5 µg/ml of bisbenzimidazole (Sigma) prior to mounting the cells in ProLong Gold Antifade Reagent (Molecular Probes[®]). Cells without primary antibodies were included as a control. Cells were examined by Leica TCS SP5 confocal laser scanning microscope (Leica Microsystems) using a 63× oil immersion lens.

Immunofluorescence microscopy of human TB lung tissues

Paraffin-embedded sections of 3 µm thickness were heat-fixed, deparaffinized and rehydrated. Epitopes were heat-retrieved in a pressure cooker with 1 mM EDTA buffer pH 8.0, and tissue sections were blocked with 0.2% gelatin in PBS with 0.05% Tween-20 (PBS-T) at room temperature for 1 h. Primary antibodies, mouse monoclonal antibody against human ADFP (1:100, RDI division of Fitzgerald Industrial Intl.), rabbit polyclonal antibody against human ACSL1 (1:150, GenWay Biotech) or rabbit polyclonal antibody against human SapC (1:50, Santa Cruz Biotechnology) were applied to tissue sections at 37°C for 1 h. After washing with PBS-T, sections were incubated with 1% Sudan Black B (Sigma) in 7% ethanol at room temperature for 10 min. Sections were washed with PBS-T and incubated with anti-Cy3[™]-goat antibody against mouse or rabbit (1:250, Jackson ImmunoResearch) at room temperature for 45 min. After PBS-T wash, nuclei were stained with 1 µg/ml of DAPI (Invitrogen) and the tissue sections were mounted in ProLong Gold Antifade Reagent, prior to examination. Images were captured by Zeiss Axioskop 40 epifluorescence microscope with AxioVision 4.3 software (Carl Zeiss MicroImaging). Further image analysis was done by AxioVision LE software (Carl Zeiss MicroImaging).

The paper explained

PROBLEM:

Tuberculosis has a serious impact on global health with a disproportionate impact on low-income countries. It is estimated that one-third of the planet is infected with its causative agent, *M. tuberculosis*, yet only 5–15% of these individuals will go on to develop disease. The balance between containment of the infection and the progression to active disease is determined at the site of infection, the granuloma. Despite the significance of this issue, little is known about the factors that regulate this process.

RESULTS:

We use human whole genome array analysis of granulomas from TB patients to demonstrate that progression of the infection in

the granuloma is accompanied by dysregulation of host lipid metabolism. Furthermore, *M. tuberculosis* induces lipid sequestration and the formation of foamy macrophages (loaded with lipid droplets) both *in vitro* and *in vivo*. The accumulation of the caseum appears to result from the death of these foamy macrophages. The expansion of the necrotic, caseous centre of the granuloma is significant both for the progression of the disease and as a potential carbon source for bacterial growth.

IMPACT:

These findings enhance our understanding of the factors, both host- and pathogen derived, that impact progression of the infection from containment to active disease.

Lipid analysis

Frozen human TB lung tissues or normal lung tissues were thawed, and caseum or normal tissue segments were isolated and transferred into glass vials (Wheaton Science) and then mixed with chloroform:methanol (2:1, v/v). After sonication at 50°C for 1 h, the extracts were filtered through 0.2 µm-pore size PTFE membrane (GD/X[®] syringe filter, Whatman) and dried down using nitrogen. TLC plates (aluminium-backed Silica gel 60 plates, EMD) were pre-run in chloroform:methanol (90:10, v/v) to remove contaminants from silica gel followed by heat-activation at 110°C for 30 min. Total lipids were developed either in a dual solvent system of chloroform:methanol:water (65:25:4, v/v/v) and petroleum ether:ethyl ether:acetic acid (70:30:2, v/v/v), or in a single solvent system of chloroform:methanol:water (65:25:4, v/v/v). Plates were sprayed with 50% sulfuric acid in ethanol and charred. The lipid standards used were cholesteryl oleate, TAG, cardiolipin and phosphatidylcholine (Sigma), CHO (MP Biomedicals), sphingomyelin (Avanti Polar Lipids) and neutral glycosphingolipid qualmix (Matreya). Caseous and fibrocaseous granuloma samples were derived from seven independent tissue samples, and normal lung tissues were from the unaffected lung area of seven independent tissue samples.

Mass-spectrometry

Structural analysis of lipids including SM, LacCer, CE and TAG were carried out by a Finnigan TSQ-7000 triple stage quadrupole mass spectrometer (Thermo Electron Corporation, CA) equipped with a Finnigan ESI source and controlled by Finnigan ICIS software operated on a DEC alpha workstation. Lipid extract dissolved in methanol/chloroform (2:1, v/v) were continuously infused into the ESI source with a Harvard syringe pump at a flow rate of 1 µl/min. The skimmer was at ground potential, and the electrospray needle was at 4.5 kV. The heated capillary temperature was 250°C. The collisionally activated dissociation (CAD) tandem mass spectra were obtained under collision energy from 30 to 45 eV, and argon (2.3 mTorr) was used as target gas. The product ion spectra were acquired in the profile mode at the scan rate of one scan per 3 s.

Structural characterization of CH was conducted by Agilent 5973 MSD GC/MS system operated by ChemStation software. Lipid extract in methanol/chloroform (2:1, v/v) was injected into a Hewlett-Packard 6890 GC in a splitless mode and analyzed with a 15 m DB-1701 column (0.25 mm i.d., 0.25 µm film thickness; J & W Scientific). The initial temperature of GC was set at 100°C for 1 min, increased to 220°C at a rate of 50°C/min, then to 290°C at a rate of 10°C/min and maintained at 290°C for another 10 min. The temperatures of the injector, transfer line of the GC column and of the ion-source were set at 280, 280 and 230°C, respectively. The full scan mass spectra (50–500 Da) were acquired at a rate of 1 scan/0.25 s.

In vivo murine granuloma model

Mycobacterial lipid-bearing beads were prepared as described previously, with slight modifications (Geisel et al, 2005; Rhoades et al, 2005). Briefly, 9.25×10^3 of 80 µm polystyrene microspheres (Duke Scientific) were coated with 15 µg of Mtb-derived TDM (Sigma) or control lipid PG (Sigma) by alternative cycles of vortexing and sonication. The TDM- or PG-coated beads were resuspended in 300 µl of ice-cold growth-factor reduced Matrigel (BD). The mixture was injected sub-dermally into the skin fold at the scruff of the neck. Granulomas were excised at 6 days post-inoculation. Three or four mice were used for the injection of TDM- or PG-Matrigel mixture. The experiment was repeated more than three times.

Histology and immunohistology of murine granulomas

Granulomas were fixed in 4% paraformaldehyde in PBS at 4°C for over 24 h and transferred to 70% ethanol. Sections were stained with H&E for histological examination. For immunostaining, paraffin-embedded sections were heat-fixed, deparaffinized and rehydrated. Epitopes were heat-retrieved in 1 mM EDTA buffer pH 8.0, and tissue sections were blocked with 5% BSA in PBS at room temperature for 1 h. Guinea pig polyclonal antibody against murine Adfp (1:200, RDI division of Fitzgerald Industrial Intl.) or rabbit polyclonal antibody against human ACSL1 (1:150, GenWay Biotech) was applied to

sections at 37°C for 1 h. After washing with PBS three times, sections were incubated with 1% Sudan Black B (Sigma) in 70% ethanol at room temperature for 10 min. Sections were washed with PBS three times and incubated with Texas Red[®]-donkey antibody against guinea pig IgG (H + L) (1:250, Jackson ImmunoResearch) or Texas Red[®]-goat antibody against rabbit IgG (H + L) (1:250, Jackson ImmunoResearch) at room temperature for 45 min. After three washes with PBS, nuclei were stained with 5 µg/ml of bisbenzimidazole (Sigma) in PBS at room temperature for 5 min. After thorough wash with PBS, sections were mounted in ProLong Gold Antifade Reagent (Molecular Probes[®]). Sections without primary antibody were included as a negative control. Images were captured by Zeiss Axio Imager with AxioVision LE software (Carl Zeiss MicroImaging).

Cryosections were cut at 10 µm thickness and fixed in 4% paraformaldehyde in PBS for 20 min. The sections were stained with Nile Red (1 µg/ml, Sigma) for neutral lipids and with bisbenzimidazole for nuclei. Mounted sections were visualized by Zeiss Axio Imager with AxioVision LE software (Carl Zeiss MicroImaging).

Author Contributions

M-JK, GK and DGR planned the experiments. M-JK, GK and DGR devised the concept. M-JK, F-FH and UW performed the experiments. GBW performed the surgery. HCW, ML, L-GB, GBW, CD, AV and LT generated and processed samples and performed pathology on the tissues. M-JK and WW performed microarray data analysis. M-JK, GK and DGR wrote and edited the manuscript.

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Supporting information is available at EMBO Molecular Medicine online.

The authors declare that they have no conflict of interest.

For more information

GEO Database; Human TB granuloma array data:

<http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?token=nns1pmyesqieqbm&acc=GSE20050>

OMIM Online Mendelian Inheritance in Man: Adipophilin (PLIN2)

<http://www.ncbi.nlm.nih.gov/omim/103195>

ACSL1

<http://www.ncbi.nlm.nih.gov/omim/152425>

PSAP

<http://www.ncbi.nlm.nih.gov/omim/176801>

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