## TB diagnosis in children: metabolite biomarkers

Cresque				
19	LEBI	NAP Patients with industries some police trial green		
Patients with active TR	Latently M. Grindward individuals			
13-	31	.29		
11-	1	14		
(14-14)	(9-12)	(7-90)		
I CIPO	24 (86%)	7 (894)		
Remo	28 (14%)	12(00%)		
6036	4470	6.76		
66.7	41.8	10.1		
le.e	43.6	10.7		
12	8.0	9.0		
lat.	4.7	3.6		
0.9	6.9	8.6		
LANCOO	4,730,000	4,790,000		
13.1	13.3	10.0		
18.9	29.4	99.2		
18.ei	11.6	30.1		
544,700	304,600	307,500		
11.9	4/7	10.0		

Tuberculosis (TB) caused by the intracellular pathogen mycobacterium tuberculosis (M. tb) is one of the most contagious diseases. Up to a quarter of the world's population is estimated to be latently infected with M. tb (LTBI). Pediatric TB contributes significantly to worldwide TB infections but is difficult to diagnose due to insufficient detection methods (READ MORE).

Many biomarkers of pulmonary and extrapulmonary TB have already been identified with the help of mass spectroscopy since infection with *M. tb* causes significant changes in the energy and protein metabolism of the host. Druszczynska and colleagues now used targeted liquid chromatography-mass spectrometry (LC-MS/MS) to assess the levels of 30 small metabolites in the serum and *M. tb* antigen-stimulated blood cultures of active TB children (Figure 1). These were then compared with those of LTBI children, infectious non-mycobacterial pneumonia (NMP) children and controls without active infection.

Description	Groups				
	TB  Patients with active TB	LTBI Latently M.th-infected individuals	NMP Patients with infectious nonreycobacterial prevenonia	HC Healthy control	
Age					
Median years (IQR)	15	5	14	7	
	(11-16)	(5-12)	(7-16)	(9-11)	
Ses, a					
M (%)	5 (33%)	24 (46%)	7 (35%)	87 (58%)	
F (%)	10 (67%)	28 (34%)	13 (63%)	62 (42%)	
WEC, counts/mm*	9770	8150	8176	8962	
Neutrophils (%)	60.7	43.8	53.1	40.1	
Lymphocytes (%)	26.8	42.6	33.7	47.5	
Monocytes (%)	9.2	8.0	9.0	8.1	
Essinophils (%)	2.6	4.7	3.6	3.6	
Basophile (%)	0.5	0.5	0.6	0.6	
RBC, counts/mm <sup>2</sup>	4,630,000	4,730,000	4,720,000	4,780,000	
HGR, gVII	13.1	13.3	12.9	13.2	
HCT.%	38.9	39.4	39.2	39.2	
MCHC, g/dl	33.6	33.8	33.1	30.7	
PLT, counts/mm <sup>2</sup>	348,300	306,800	337,500	308,800	
CRE mgG	41.9	6.7	12.9	0.8	

Table 1: Demographic characteristics of study subjects. CRP C-reactive protein, F female, haematocrit, hGB haemoglobin, M male, MCHC men corpuscular haemoglobin concentration, M.tb Mycobacterium tuberculosis, n number, PLT platelets, RBC red blood cells, SD standard deviation, TB tuberculosis, WBC white blood cells (Druszcynska, et al., 2022).

Comparison of the levels of metabolites between the groups revealed that serum leucine levels were significantly higher in the TB and LTBI groups, whereas the abundance of citrulline was significantly lower in these patients compared to the healthy control group. Further metabolites with significant differences included hydroxyproline, proline, valine, tyrosine, leucine, kynurenine, valine and neopterine. Using receiver operator characteristic (ROC) curves, area under the curve (AUC) and elastic-net logistic regression analysis, they identified serum leucine as the best discriminator of TB + LTBI versus healthy control + NMP. Unfortunately, the small sample size of the TB group (n = 15) did not allow for a selection of the most informative metabolites to discriminate this group against the others. Nevertheless, the obtained

results are a proof of principle of the usefulness of metabolites in the diagnosis of TB in children.

Journal article: Druszczynska et al., 2022. <u>Targeted</u> <u>metabolomics analysis of serum and *Mycobacterium tuberculosis* antigen-stimulated blood cultures of pediatric patients with active and latent tuberculosis. *Scientific Reports*.</u>

Summary by Dr. Jasmin Knopf