

Supplementary Material

Immunomagnetic separation improves the detection of mycobacteria by paper-based lateral and vertical flow immunochromatographic assays

Alejandra Ben Aissa¹, Barbara Araújo^{1,4}, Esther Julian³, Maria Valnice Boldrin Zanoni⁴ and María Isabel Pividori^{1,2,*}

- 1 Departament de Química, Universitat Autònoma de Barcelona, 08193, Bellaterra, Spain.
- 2 Institute of Biotechnology and Biomedicine, Universitat Autònoma de Barcelona, Bellaterra, Spain.
- 3 Departament de Genètica i de Microbiologia, Facultat de Biociències, Universitat Autònoma de Barcelona, Bellaterra, Spain
- 4 Department of Analytical Chemistry, Institute of Chemistry, UNESP-Univ Estadual Paulista, Araraquara, SP, Brazil

* Correspondence: Isabel.Pividori@uab.cat; Tel.: (+34 93 581 1976)

2. Materials and Methods

2.9. Double-tagging PCR for the amplification of Mycobacteria

Table S1. Thermal cycler conditions for the double-tagging PCR

	Initial step	DNA denaturation	Annealing	Extension	Last step
	1 cycle	40 cycles			1 cycle
Temperature (°C)	95	95	60	72	72
Time (s)	300	40	30	120	600

3. Results and Discussion

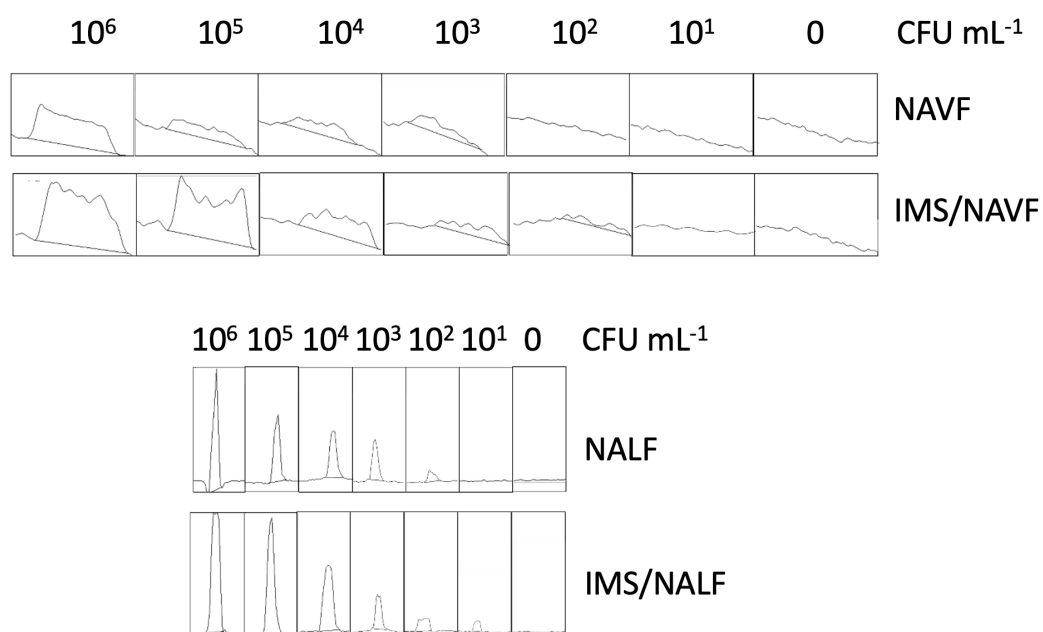


Figure S1. The relative areas obtained after processing the images, and which were plotted in Figure 6 and 7, are shown for both platforms and sample treatments. NAVF, Nucleic Acid Vertical Flow; IMS, immunomagnetic separation; NALF, Nucleic Acid Lateral flow.