

# **A Novel Method that allows SNP Discrimination with 160:1 ratio for Biosensors Based on DNA-DNA hybridization**

Satish Balasaheb Nimse,<sup>a†</sup> Keum-Soo Song,<sup>b†</sup> Shrikant Dashrath Warkad,<sup>b</sup> Taisun Kim<sup>a\*</sup>

<sup>a</sup>Institute for Applied Chemistry and Department of Chemistry, Hallym University, Chuncheon, 200-702, Korea, Fax: +82-33-256-3421, E-mail: [tskim@hallym.ac.kr](mailto:tskim@hallym.ac.kr)

<sup>b</sup>Biometrix Technology, Inc. 202 BioVenture Plaza, Chuncheon, 200-161, Korea

## **Supporting Information**

**Table S1.** Sequences of the probes used for derivation of SWAT

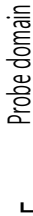
Probes	Sequence	T <sub>m</sub> (°C)	
		[4(G+C)+2(A+T)-5]	Nearest neighbor method
<b>Probe1</b>	5'-9G-vertical spacer-TAC CGA CCC AC <sup>G</sup> CGG GC-3'	55	55.96
<b>Probe2</b>	5'-9G-vertical spacer-TAC CG <sup>G</sup> CCC AC <sup>G</sup> CGG GC-3'	57	58.45
<b>Probe3</b>	5'-9G-vertical spacer-TAC CG <sup>G</sup> CCC ACC CGG GC-3'	57	56.30
<b>Probe4</b>	5'-9G-vertical spacer-TAC CGA CCC AC <sup>G</sup> TGG GC-3'	53	52.37
<b>Probe5</b>	5'-9G-vertical spacer-TAC CG <sup>G</sup> CCT AC <sup>G</sup> CGG GC-3'	55	56.78
<b>Probe6</b>	5'-9G-vertical spacer-TAC CG <sup>G</sup> CCC ACC TGG GC-3'	55	54.3

9G- nine consecutive guanines (GGG GGG GGG) used for the immobilization of the probes on the AMCA DNACChip; Vertical spacer- nine consecutive thymidine's (TTT TTT TTT); T<sub>M</sub>: melting temperature

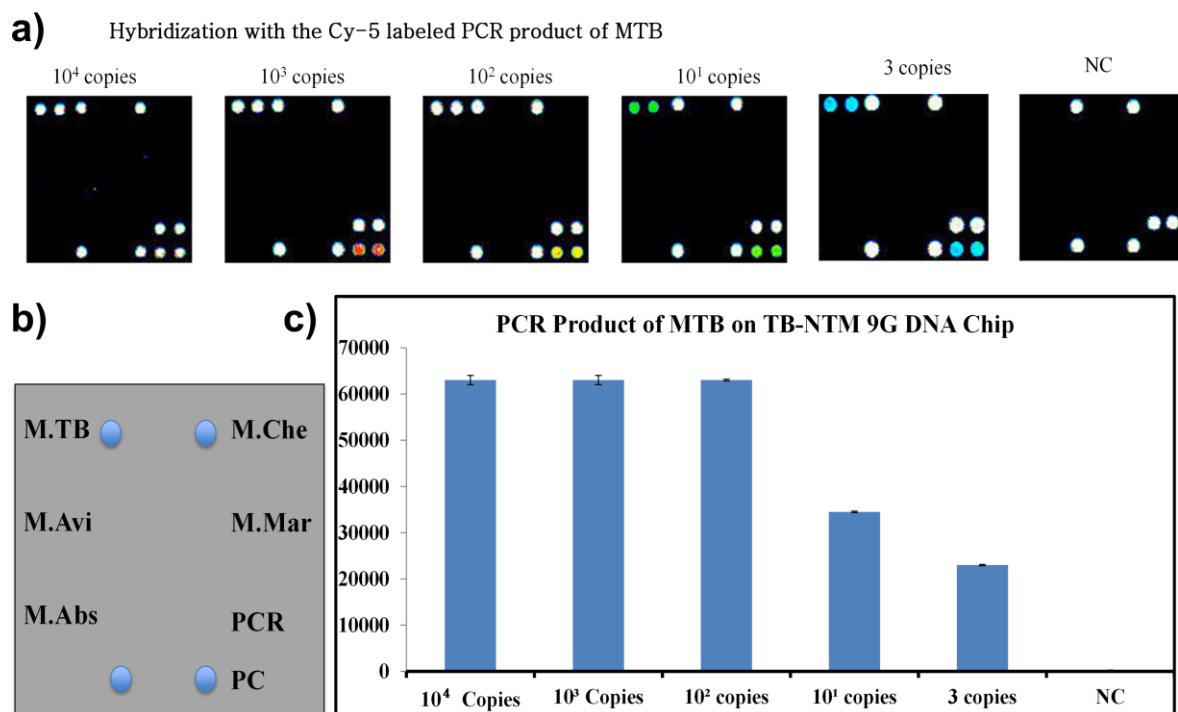
**Table S2.** Sequences of probes used for the detection and discrimination of MTB strains and NTM Strains

Probes	Probe Description	Sequence	T <sub>m</sub> (°C)
<b>Probe7</b>	<i>MTB</i>	5'-9G-vertical spacer-TAG CGA CCG AC <sup>G</sup> TGG GC-3'	53
<b>Probe8</b>	<i>Chelonae</i>	5'-9G-vertical spacer-TAC CGG CCT AC <sup>G</sup> CGG GC-3'	55
<b>Probe9</b>	<i>Avium</i>	5'-9G-vertical spacer-TAG CGG CCC ACC TGG GC-3'	55
<b>Probe10</b>	<i>Marinum</i>	5'-9G-vertical spacer-TAC CGG CCC ACC TGG GC-3'	55
<b>Probe11</b>	<i>Abscessus</i>	5'-9G-vertical spacer-TAC CGA CCC AC <sup>G</sup> TGG GC-3'	53
<b>Probe12</b>	<i>kansasii</i>	5'-9G-vertical spacer-TAT CGG CCG ACATGG GC-3'	53
<b>Probe13</b>	PCR	5'-9G-vertical spacer-CGA CCS ACV CGS GCC AGG TC -3'	61
<b>Probe14</b>	NC	5'-9G-vertical spacer-GCC AGG TCG TAG CGC TTC TC-3'	61
<b>Probe15</b>	HC	5'-9G-vertical spacer-CCT AGT GGC TCT ATG GTA AC-3'	55
<b>HC-Cy5-T1</b>	HC Target DNA	3'- GGA TCA CCG AGA TAC CAT TG GAG ACT GCG -Cy5-5'	

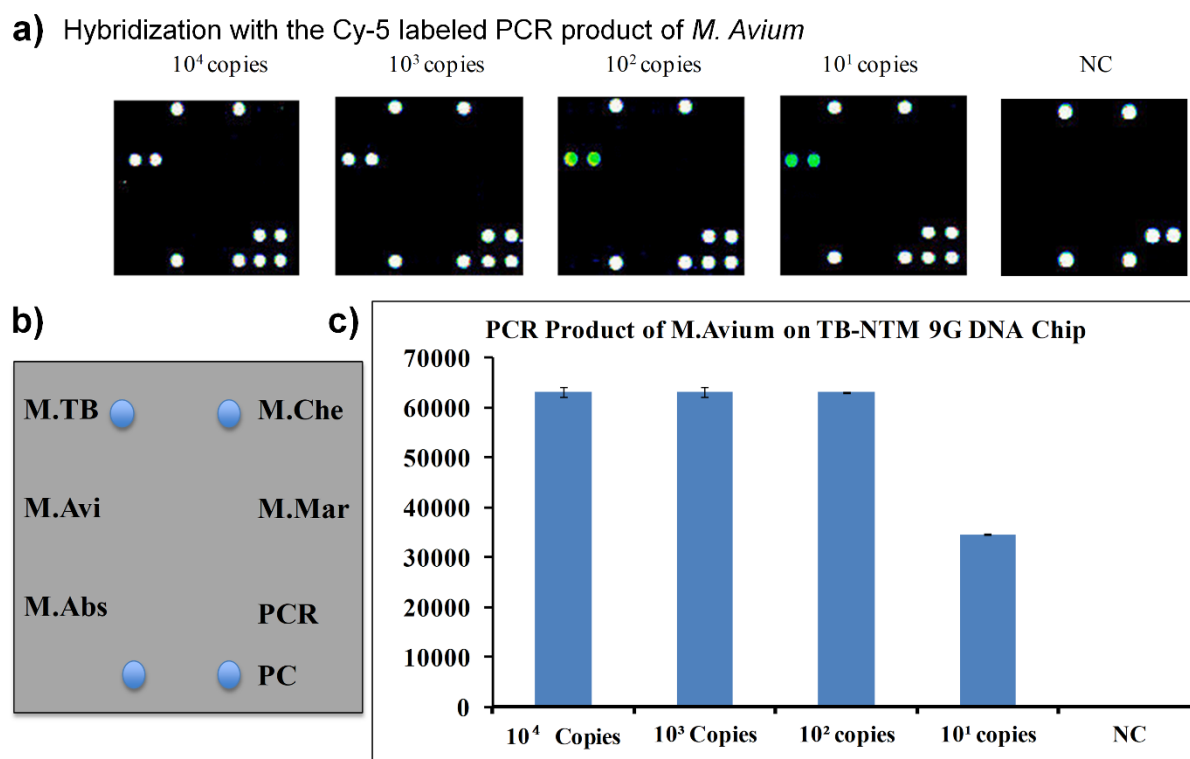
HC – probe for the Hybridization control; HC-Cy5-T1 – Target oligonucleotide for HC probe9G- nine consecutive guanines (GGG GGG GGG) used for the immobilization of the probes on the AMCA DNACChip; Vertical spacer- nine consecutive thymidine's (TTT TTT TTT); T<sub>M</sub>: melting temperature



**S3**

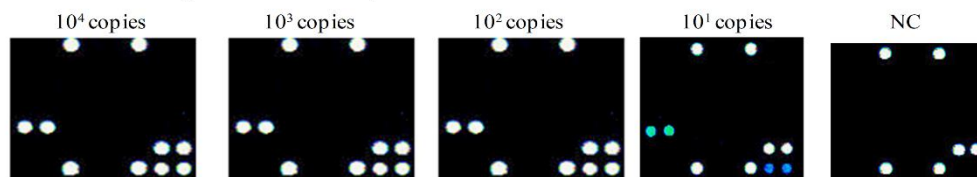


**Figure S2:** Limit of detection for the identification of MTB.

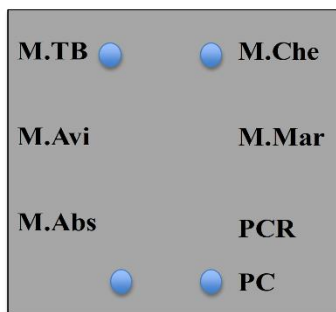


**Figure S3:** Limit of detection for the identification of *Mycobacterium Avium*.

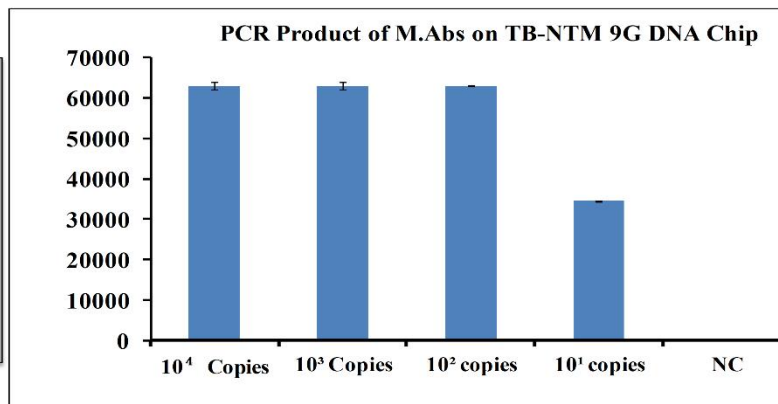
a) Hybridization with the Cy-5 labeled PCR product of *M. Abscessus*



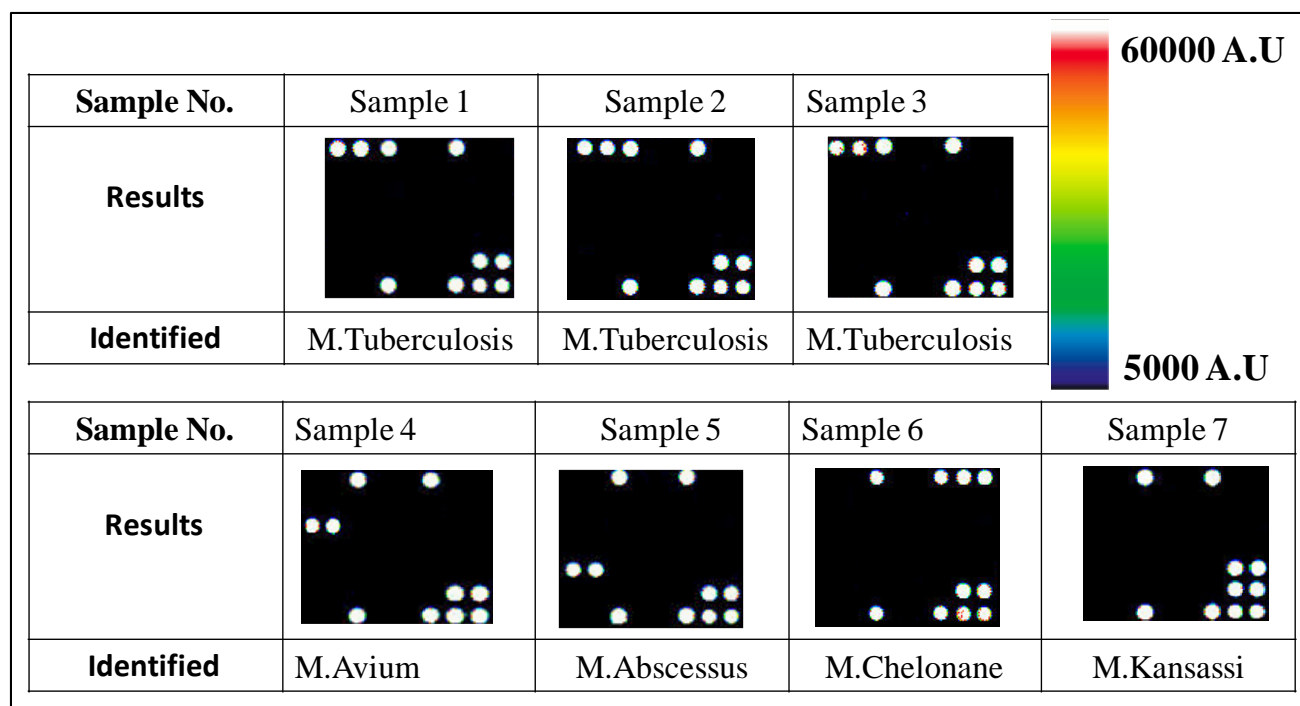
b)



c)



**Figure S4:** Limit of detection for the identification of *Mycobacterium Abscessus*.



**Figure S5:** The detection and discrimination of the MTB and NTM strain in the clinical samples.