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SDI FINAL EVALUATION FORM 1.1

PART 1:

Journal Name:	Asian Journal of Research in Biochemistry	
Manuscript Number:	Ms_AJRB_43243	
Title of the Manuscript:	High-level expression and purification of DNA and DNase free Taq DNA polymerase	
Type of Article:	Original Research Article	

PART 2:

FINAL EVALUATOR'S comments on revised paper (if any)	Authors' response to final evaluator's comments
A revised version of the manuscript addresses the most points raised during the review	
process. However, I do think that the author(s) need(s) to work a little more on the	
manuscript. The pLoxGentrc vector encoding Taq DNA polymerase was used to produce	
the recombinant enzyme. However, a construction of this plasmid has not been described	
in the Methods section. It is an important part of work and can not be omitted. Protein gel staining methods were also omitted in the methodology section.	
Protein ger staining methods were also omitted in the methodology section.	
Information on clinical samples used in the study could be provided in section 2.7 and not	
at the end of the manuscript.	
Fig. 5a DNA size marker is not marked.	
I would like to emphasize that the scientific name of the species is always written in italics.	
Therefore it should be <i>Tubercle bacilli</i> or <i>Tubercle bacillus</i> (lane 120) instead of tubercle	
bacilli. Symbols for genes are also italicized, so <i>rpoB</i> gene (lanes 118, 284), <i>chuA</i> gene	
(lanes 119, 284) instead of rpoB gene, chuA gene.	
I think that one name of the bacterium <i>Tubercle bacillus</i> or <i>M. tuberculosis</i> (lane 284)	
should be used throughout the manuscript.	
The manuscript still requires editing of English language and style. Some examples:	
- lanes 344 and 345: in my opinion it should be: samples were obtained from/sample	
was obtained from instead of samples were got from/sample is got from;	
- lane 99: it should be DNase activity instead of DNaase activity;	
- lanes 217,221,226, 319: it should be with specific primers instead of with specific primer;	
- lane 325: it should be with all types of samples instead of with all types of sample.- Fig, 5b: Unit of tag enzyme?	
- Fig. 3b. Officer tag enzyme? - I am afraid that Tag DNA polymerase can not be used as a template (lanes 218, 222).	
- Lane 124-125: text reads: "The recombinant plasmid containing Tag DNA polymerase	
gene was confirmed by PCR with gene specific primers "In my opinion, the presence of	
Taq DNA polymerase gene within the recombinant vector/plasmid was confirmed by PCR	
and not the recombinant plasmid.	
and not the recombinant placeman	

Reviewer Details:

Name:	Sabina Kędzierska-Mieszkowska
Department, University & Country	Faculty of Biology, University of Gdańsk, Poland

Created by: EA Checked by: ME Approved by: CEO Version: 1.5 (4th August, 2012)