

Target-triggered exponential signal amplification reaction for RNA detection



Seoyoung Lee^a and Hyun Gyu Park^{a*}

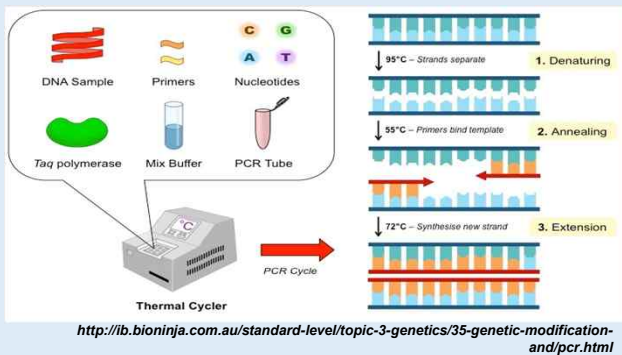
^a Department of Chemical and Biomolecular Engineering, KAIST, 291 Daehak-ro, Yuseong-gu, Daejeon 305-701, Republic of Korea

* hgpark1@kaist.ac.kr

A. Introduction

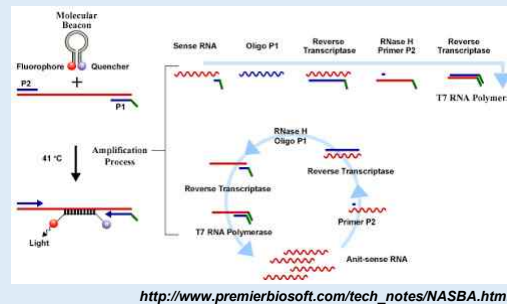
Polymerase chain reaction (PCR)

Isothermal nucleic acid amplification

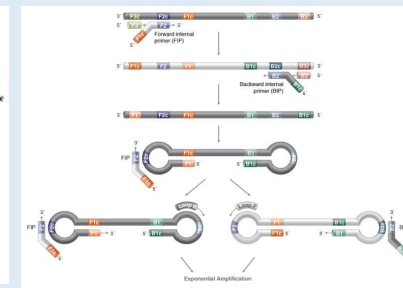


Disadvantages

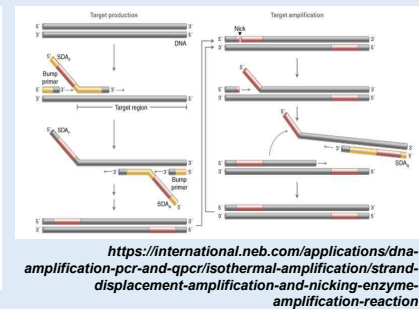
- Massive & expensive thermal cycler
- Not applicable to Point-of-care testing technology



Nucleic-acid sequence-based amplification (NASBA)



Loop-mediated isothermal amplification (LAMP)



Strand displacement amplification (SDA)

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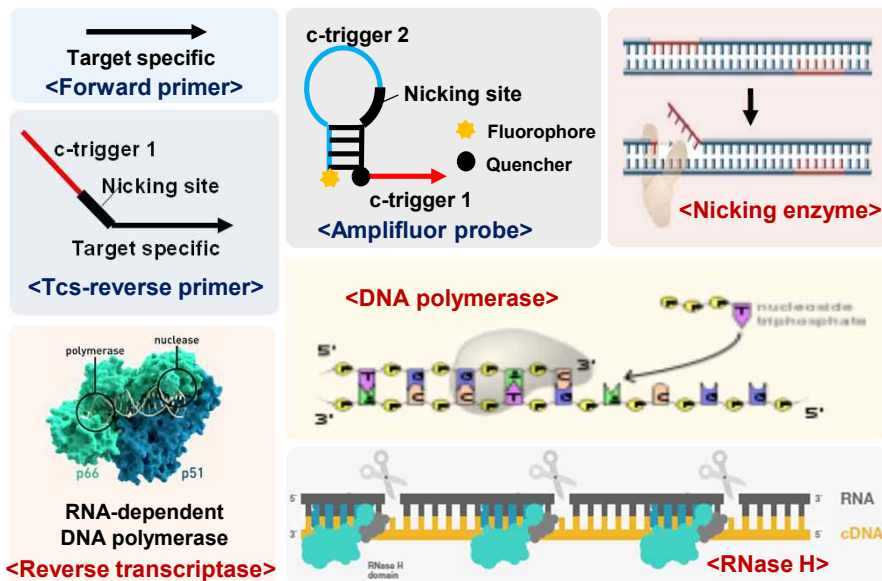
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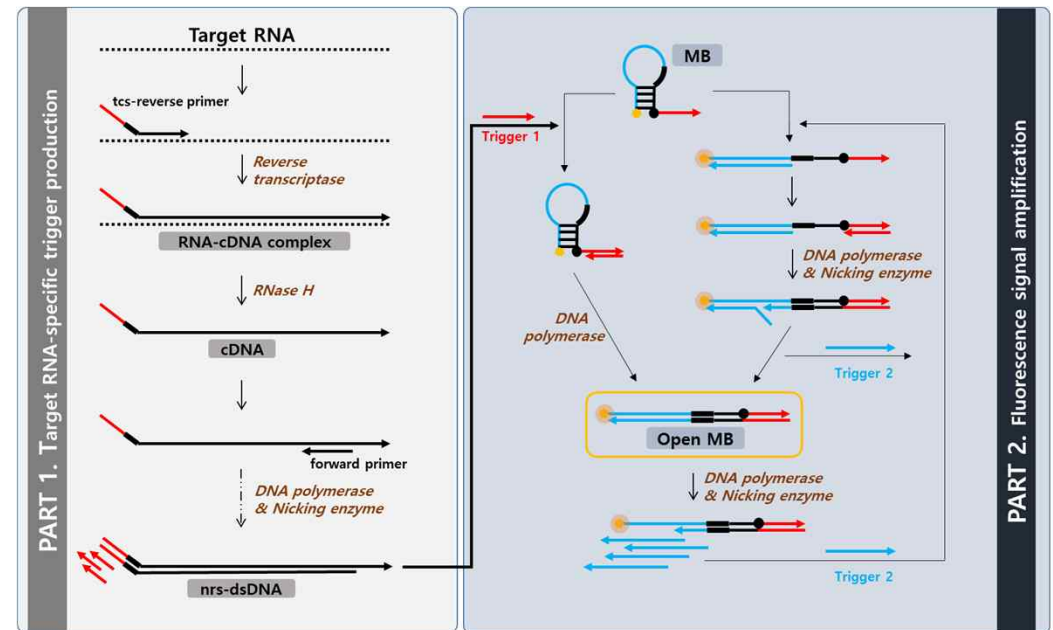
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B. Schematic representation

Key components



Target RNA-specific trigger induced exponential signal amplification reaction



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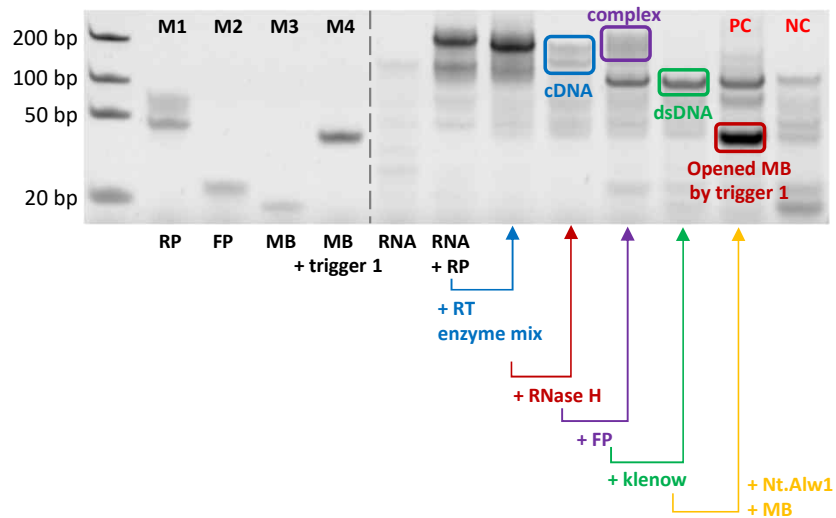
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C. Results & Discussion

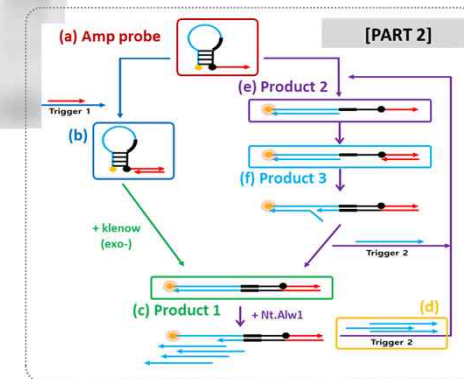
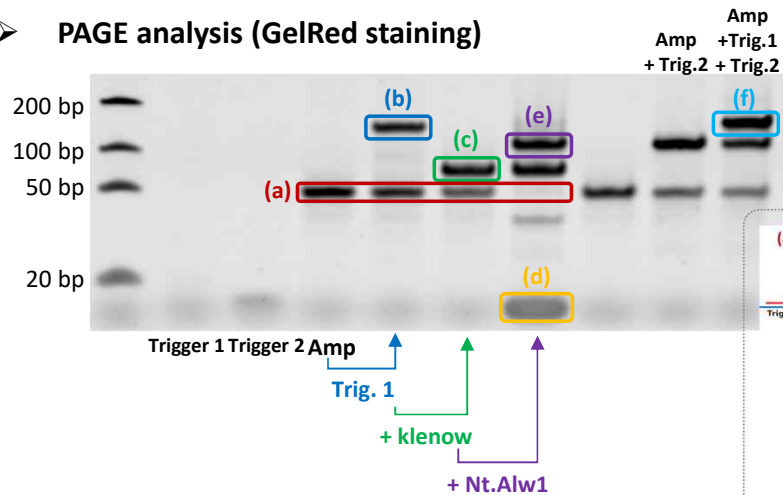
PART 1 feasibility test

PAGE analysis (GelRed staining)



PART 2 feasibility test

PAGE analysis (GelRed staining)



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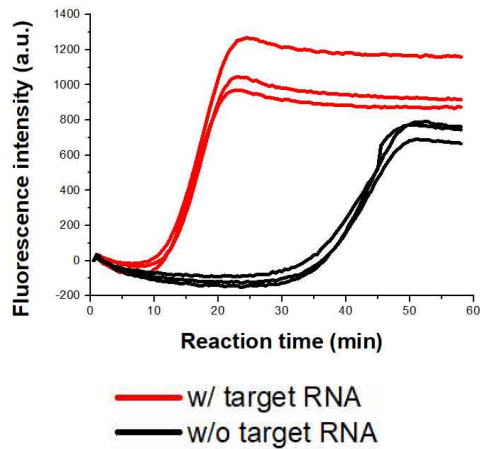
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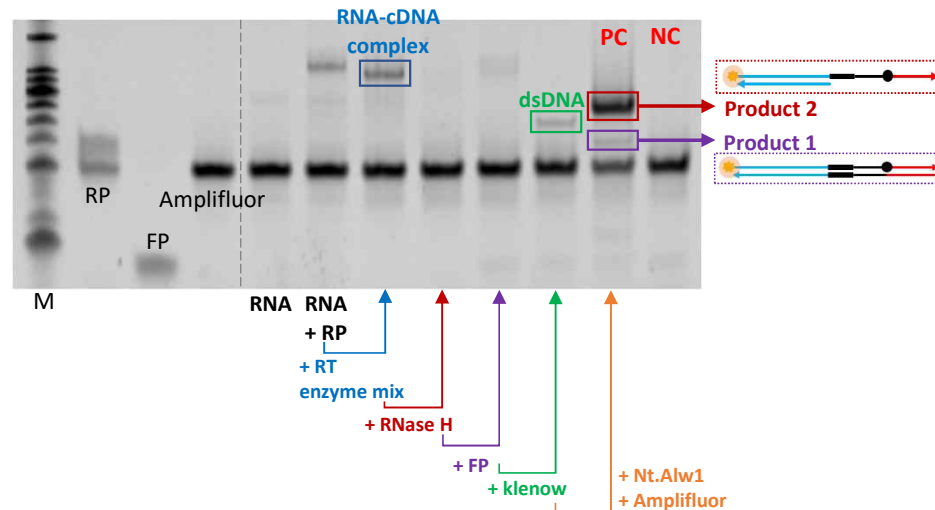
C. Results & Discussion

Feasibility test of whole reaction

Real-time fluorescence analysis

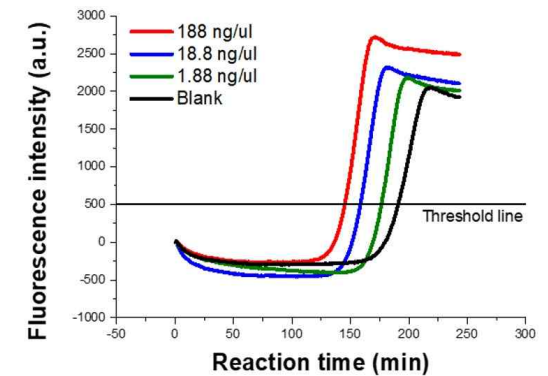


PAGE analysis (GelRed staining)



Sensitivity test

Target: total RNA of *Brachionus rotundiformis*



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D. Conclusion

Target-specific trigger-induced exponential signal amplification reaction

Development of novel real-time RNA detection method

Great insight for the development of self-operative isothermal amplifying system enabling target RNA detection

Applicable to the development of isothermal amplification system for detection of several virus (e.g. SARS-CoV-2, MERS-CoV), pathogens, or cancer cells

A powerful platform for the multiplex isothermal detection of several RNA target by employing multiple trigger-specific Amplifluors