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## Open Isothermal Platform Protocal

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1 Works for me dx.doi.org/10.1

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- 1 Extract RNA as per dx.doi.org/10.17504/protocols.io.bkynkxve
- 2 Transfer 5ml of extracted RNA to pre-aliquoted LAMP mix. Mix by pipetting up and down 5 times.
- 3 Add 5ml of H<sub>2</sub>O in negative control well 7.
- 4 Add 5ml of positive control in well 8.
- 5 Cap all wells

- 6 Place strips in FABL-8
- 7 Run FABL-8 as per manual

## 8 LAMP reaction (can refer to N1-STOP-LAMP paper)

Single reaction setup:

Reagent	Volume (ml)
Optigene Master Mix	15
N1 primer	5
Extracted RNA	5
Total	25

To reduce pipetting error, Master Mix and N1 primer are premixed according to the number of reactions desired (allow for pipetting error by preparing enough for  $\sim$ 1.1x the number of reactions, rounded to the nearest whole number); 20ml of this mixture is then aliquoted per reaction.