Kazu Buffer Feedback Form

Name: Shuiyi Thames

Lab: Eveland Lab

Material: Maize leaf

Primer set: IDDP1(F/R1) and JSR01

Amplification size (kb): 0.268kb

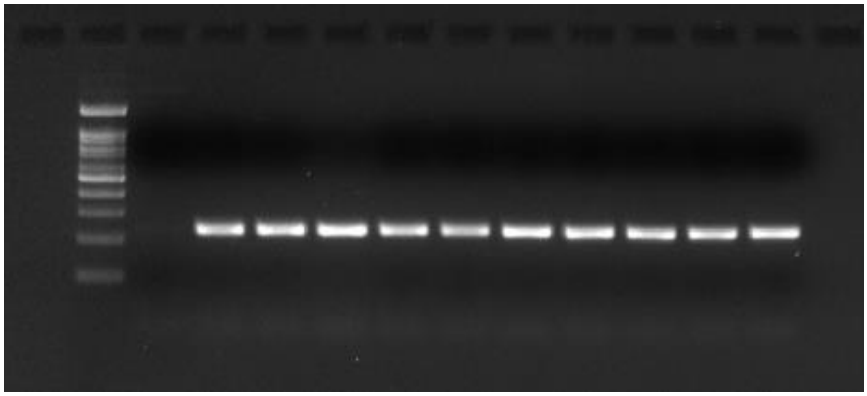
PCR taq: Invitrogen Platinum Taq DNA Polymerase

PCR amplification:

Reagent	Volume
Kazu DNA extract	2 µl
Taq	0.1 µl
Forward primer	1 µl
Reverse primer	1 µl
	µl
	µl
H_2O	15.9 µl
Total volume	25 µl

Step	Temperature	Time	Cycles
Initial Denaturation	94℃	2 min	N/A
Denaturation	94℃	30s	x34
Annealing	57℃	30s	
Extension	72℃	1min	
Final Extension	72℃	5min	N/A
Hold	12℃	Infinite hold	N/A

PCR Gel Picture:



Would you prefer using Kazu Buffer over your current lab DNA extraction protocol? ☐ **Yes** ☐ **No**

Why or why not?

It is so easy and convenient. Kazu's method saves me a lot of time of extraction DNA!

Other Comments:

I am looking forward to the new buffer for extracting RNA ☺