

## DNA extraction from Meat

[Empty box for title or introduction]



**Reagents**  
 Mix the Cellease A, B and distilled water  
 (20 ul Cellease A, 20 ul CellEase B, 60 ul distilled water )



Add 100 ul of the mixture to the samples.



Incubate at 37°C for 6 minutes  
 Then incubate at 95°C for 3 minutes



Transfer 5-7ul of extracts to PCR reaction mixture and  
 amplify the target DNA fragment



Pork (about 5mg) in 0.2 ml micro test tube

PCR

- 5~7ul Test sample
- 5.0 ul × 10 buffer(+Mg<sup>2+</sup>)
- 5.0 ul dNTPs
- 1.0 ul Forward Primer (10pmol/ul)
- 1.0 ul Reverse Primer (10pmol/ul)
- 0.5 ul Ex Taq (5 U/ul)

Fill up to 50 ul by distilled water

PCR Cycle

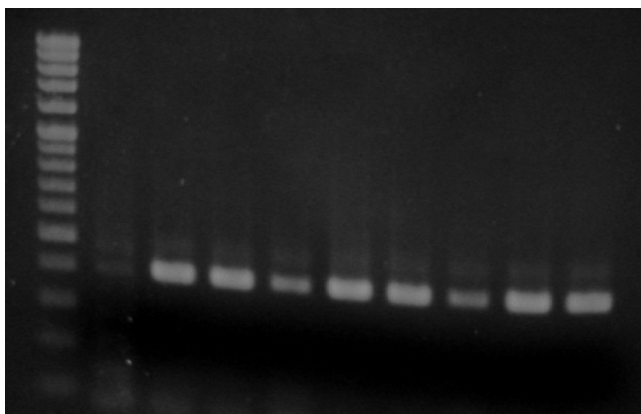
94°C	1min
94°C	30sec
55°C	30sec
72°C	60sec
72°C	4min

35 Cycles

## <Results>

### ①DNA extraction and detection from porch samples

M 1 2 3 4 5 6 7 8 9

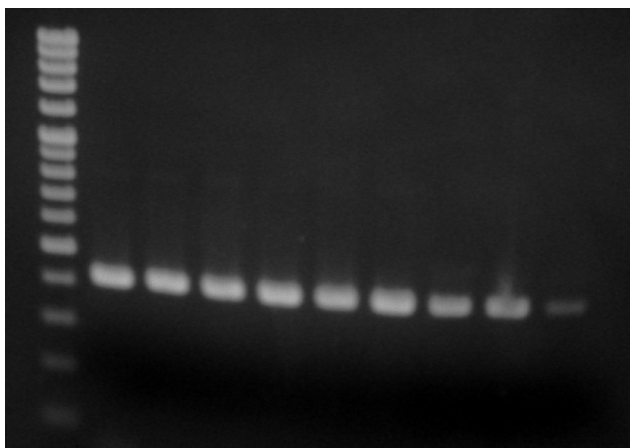


M Marker (100bp ladder)  
1 Sample 4mg, Add 5 $\mu$ l of DNA extract to PCR  
2 Add 6 $\mu$ l of DNA extract to PCR  
3 Add 7 $\mu$ l of DNA extract to PCR  
4 Sample 8mg, Add 5 $\mu$ l of DNA extract to PCR  
5 Add 6 $\mu$ l of DNA extract to PCR  
6 Add 7 $\mu$ l of DNA extract to PCR  
7 Sample 12mg, Add 5 $\mu$ l of DNA extract to PCR  
8 Add 6 $\mu$ l of DNA extract to PCR  
9 Add 7 $\mu$ l of DNA extract to PCR

As a results, 6~7 $\mu$ l of DNA extract was thought to be best for PCR.

### ②DNA extraction and detection from fish (Tuna) samples

M 1 2 3 4 5 6 7 8 9



M Marker (100bp ladder)  
1 Sample 6mg, Add 5 $\mu$ l of DNA extract to PCR  
2 Add 6 $\mu$ l of DNA extract to PCR  
3 Add 7 $\mu$ l of DNA extract to PCR  
4 Sample 9mg, Add 5 $\mu$ l of DNA extract to PCR  
5 Add 6 $\mu$ l of DNA extract to PCR  
6 Add 7 $\mu$ l of DNA extract to PCR  
7 Sample 18mg, Add 5 $\mu$ l of DNA extract to PCR  
8 Add 6 $\mu$ l of DNA extract to PCR  
9 Add 7 $\mu$ l of DNA extract to PCR

As a results, less than 10mg of test sample was thought to be best for PCR.