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## APPENDIX A

# Preparation of reagents and buffer

# 10% stool suspension

1.	Phosphate buffered saline (PBS), pH 7.4	
	NaCl	8.0 g
	KH <sub>2</sub> PO <sub>4</sub>	0.2 g
	Na <sub>2</sub> HPO <sub>4</sub> .12H <sub>2</sub> O	2.9 g
	KCI	0.2 g
	Distilled water was added up to 1000 ml	
2.	0.1% DEPC	
	DEPC	100.0 μ1
	Dissolved in 100 ml distilled water	
Agarose gel electrophoresis		
1.	TAE Buffer (50X) Tris-base	
	Tris-base UNI	242.0 g
	Glacial acetic acid	57.1 ml
	0.5 M EDTA (pH 8.0)	100.0 ml
	Distilled water was added up to 1000 ml	1110
	The working solution (1X) contains 20 ml TAE (50X) and 980 m	l distilled
	water	e d
2.	Agarose gel (1.5%)	
	TAE Buffer (1X)	100.0 ml
	Agarose gel powder	1.5 g

## 3. Working 1:5 RedSafe<sup>TM</sup> Nucleic Acid Staining Solution

RedSafe solution 20.0 μl
Distilled water 80.0 μl

### 4. DNA marker preparation

0.5  $\mu$ g/ $\mu$ l DNA ladder (Thermo Scientific, USA) 100.0  $\mu$ l 6X Loading dye 100.0  $\mu$ l Distilled water 400.0  $\mu$ l



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