

MINIPLEX STR AMPLIFICATION WORKSHEET

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Date: _____ Project Name: _____

Reaction Mix:	200 μ L dNTPs	}	Vortex mixture and spin down
	250 μ L 10X Buffer		
	50 μ L ddH ₂ O		

Prepare Primers:	5 μ L of 100 μ M Forward Primer	}	Prepare a separate primer mix for each individual loci ***If 200 μ M Primers, use 2.5 μ L of primers and adjust ddH ₂ O to keep final volume at 50 μ L.***
	5 μ L of 100 μ M Reverse Primer		
	40 μ L of ddH ₂ O		

Master Mix:			Mini 4			Big Mini		
<i>Mini 2</i>	<i>Per Sample (μL)</i>	<i>Samples</i>	<i>Per Sample (μL)</i>	<i>Samples</i>	<i>Per Sample (μL)</i>	<i>Samples</i>		
D5	1.0		vWA	1.0		TH01	0.4	
D8	1.0		D18	1.0		CSF	0.4	
D16	0.5		D13	1.4		TPOX	0.5	
Reaction Mix	5.0		Reaction Mix	5.0		FGA	0.6	
BSA	1.0		BSA	1.0		D21	0.6	
Taq	0.4		Taq	0.4		D7	0.8	
ddH ₂ O	15.1		ddH ₂ O	14.2		Reaction Mix	5.0	
Total	24.0 μL		Total	24.0 μL		BSA	1.0	
						Taq	0.4	
						ddH ₂ O	14.3	
						Total	24.0 μL	

Remember to add 2 to the number of samples for pipetting error

~Pipet 24 μ L of each master mix into prepared 0.2 μ L tubes for each sample.
~Add 1 μ L of sample DNA to each sample tube (add water to reagent blank and DNA standard to positive control).

~PCR Cycling Conditions:

Step 1:	10 minutes at 94oC warm-up	Step 2:	1 minute at 94oC 1 minute at 55oC 1 minute at 72oC 33 cycles	Step 3:	45 minutes at 60oC Forever at 25oC
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