

1

Create Sample Sheet

Date/Time: \_\_\_\_\_

Sample DNA Plate ID: \_\_\_\_\_

Operator: \_\_\_\_\_

Kit PN: \_\_\_\_\_

2

Make ARX

Date/Time: \_\_\_\_\_

ARX Plate ID: \_\_\_\_\_

Operator: \_\_\_\_\_

NaOH Batch: \_\_\_\_\_

- Set centrifuge to 15° to 25°C, if refrigerated
- Preheat incubating microplate shaker (68°C)
- Preheat heat block (95°C)
- Attach pool guide label to VBP and label plate
- Add 5 µl DNA to 3 DNA sample well locations
- Vortex 0.1N NaOH (5 s)
- Add 5 µl 0.1N NaOH (pipette up and down)
- Vortex MTR4A, MTR4B, MTR4C (5 s)
- Centrifuge MTR4A, MTR4B, MTR4C (briefly)
- Add 10 µl MTR4A (red cap) to columns 1-4
- Add 10 µl MTR4B (yellow cap) to columns 5-8
- Add 10 µl MTR4C (blue cap) to columns 9-12
- Seal and centrifuge ARX plate (1,000 xg, 1 m)
- Incubate ARX plate (program step 2)
- Place ARX plate on heat block (95°C, 1 m):
- Place ARX plate on bench (RT, 3 m):
- Incubate ARX plate (program step 3)
- Seal and centrifuge ARX plate (1,000 xg, 1 m)
- Add 5 µl NaOH
- Seal and incubate ARX plate (program step 4)
- Vortex AB1 (5 s)
- Add 30 µl AB1
- Seal and incubate ARX plate (program step 5)
- Centrifuge ARX plate (1,000 xg, 1 m)
- Place ARX plate on magnetic plate (1 m)
- Remove supernatant
- Vortex UB3 (5 s)
- Add 40 µl UB3
- Seal and incubate ARX plate (program step 6)
- Place ARX plate on magnetic plate (1 m)

Kit Lot Number: \_\_\_\_\_

MTR4A Lot Number: \_\_\_\_\_

MTR4B Lot Number: \_\_\_\_\_

MTR4C Lot Number: \_\_\_\_\_

AB1 Lot Number: \_\_\_\_\_

UB3 Lot Number: \_\_\_\_\_

AOP0 Lot Number: \_\_\_\_\_

AOP4A Lot Number: \_\_\_\_\_

AOP4B Lot Number: \_\_\_\_\_

AOP4C Lot Number: \_\_\_\_\_

Incubating Microplate Shaker ID: \_\_\_\_\_

Heat Block ID: \_\_\_\_\_

Pipette ID: \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

Project: \_\_\_\_\_ Batch: \_\_\_\_\_ Date: \_\_\_\_\_

- Vortex AOP0 (5 s)
- Add 1,200 µl AOP0 to AOP4A, AOP4B, AOP4C
- Vortex AOP4A, AOP4B, AOP4C (5 s)
- Centrifuge AOP4A, AOP4B, AOP4C (briefly)
- Remove supernatant
- Add 40 µl AOP4A (red cap) to columns 1-4
- Add 40 µl AOP4B (yellow cap) to columns 5-8
- Add 40 µl AOP4C (blue cap) to columns 9-12
- Seal and incubate ARX plate (program step 7)

## 3

## Add ELM2

Date/Time: \_\_\_\_\_

AE1 Lot Number: \_\_\_\_\_

Operator: \_\_\_\_\_

UB3 Lot Number: \_\_\_\_\_

- Centrifuge ARX plate (1,000 xg, 1 m)
- Place ARX plate on magnetic plate (1 m)
- Remove supernatant
- Vortex AE1 (5 s)
- Add 40 µl AE1
- Seal and incubate ARX plate (program step 8)
- Place ARX plate on magnetic plate (1 m)
- Remove supernatant
- Add 40 µl AE1
- Seal and incubate ARX plate (program step 9)
- Place ARX plate on magnetic plate (1 m)
- Remove supernatant
- Vortex UB3 (5 s)
- Add 40 µl UB3
- Seal and incubate ARX plate (program step 10)
- Place ARX plate on magnetic plate (1 m)
- Vortex ELM2 (5 s)
- Remove supernatant
- Add 40 µl ELM2
- Seal and incubate ARX plate (program step 11)

ELM2 Lot Number: \_\_\_\_\_

**4** Add MAM1

Date/Time: \_\_\_\_\_ Titanium Taq DNA Polymerase Lot Number: \_\_\_\_\_

Operator: \_\_\_\_\_

 Add 48 µl Titanium Taq DNA Polymerase to MAM1 MAM1 Lot Number: \_\_\_\_\_ Vortex MAM1 **with Taq** (5 s) Centrifuge ARX plate (1,000 xg, 1 m) Place ARX plate on magnetic plate (1 m) Remove supernatant Add 40 µl MAM1 **with Taq** Seal and incubate ARX plate (program step 12)**Cycle PCR:** Check Cycler Program Cycler ID: \_\_\_\_\_ Cycle ARX plate: Start: \_\_\_\_\_ Stop: : \_\_\_\_\_ Cycler Program ID: \_\_\_\_\_ Check Cycler Program Complete**5** Make MSS

Date/Time: \_\_\_\_\_ VBP Plate ID: \_\_\_\_\_

Operator: \_\_\_\_\_ Incubator ID: \_\_\_\_\_

 Preheat shaking incubator (47°C) MSS Lot Number: \_\_\_\_\_ Centrifuge ARX plate (680 xg, 1 m) Place ARX plate on magnetic plate (1 m) Centrifuge VBP (680 xg, 1 m) Add 40 µl MSS to VBP Transfer 45 µl from ARX plate to VBP**6** Hyb VeraCode Bead Plate

Date/Time: \_\_\_\_\_ VW2 Lot Number: \_\_\_\_\_

Operator: \_\_\_\_\_

 Vortex VBP in shaking incubator (1,200 rpm, 2.5 hours, 47°C):

Start: \_\_\_\_\_ Stop: \_\_\_\_\_

 Centrifuge VBP (680 xg, 1 m) Add 200 µl VW2 Swirl, wait (1 m), remove supernatant Add 200 µl VW2 Swirl, wait (1 m), remove supernatant

## 7 Prepare BeadXpress® Reader

Date/Time: \_\_\_\_\_

Operator: \_\_\_\_\_

Preventive maintenance:  Up to date

Fluidics bottles:  Checked and filled

## 8 Scan VeraCode Bead Plate

Date/Time: \_\_\_\_\_

VBP Plate ID: \_\_\_\_\_

Operator: \_\_\_\_\_

BeadXpress Reader ID: \_\_\_\_\_

Select ADME Kit

Enter VBP ID

Select sample sheet

Insert VBP in correct plate orientation

Scan

Generate report