

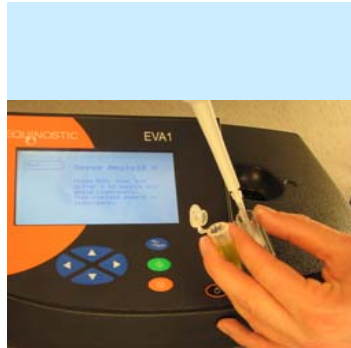


**1**

Use the **blue pipette** to dispense **100 µl** Buffer 1 into the cuvette.

### Important!

- 1) Start EVA1
  - 2) Press F2 to accept diagnostic report
  - 3) Press F2 to enter Equinostic methods
  - 4) Press F1 for SAA measurement
- Avoid bubbles in tip



**2**

Followed by **2 µl** sample, taken by the **read pipette**.

### Important!

Avoid Serum/plasma on surface of the tip. Can gently be wiped off by filter paper

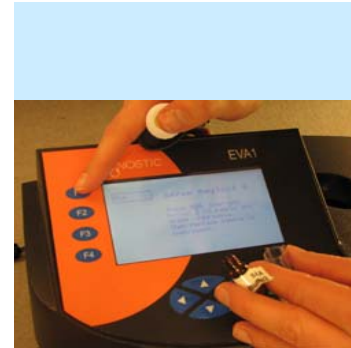


**3**

Shake the cuvette with vortex mixer and place the cuvette in cuvetteholder of EVA1, alternatively in a 37 C incubator, for 10 minutes

### Important!

Buffer 1 and sample can be placed in incubator for 1 hour, without effecting result.

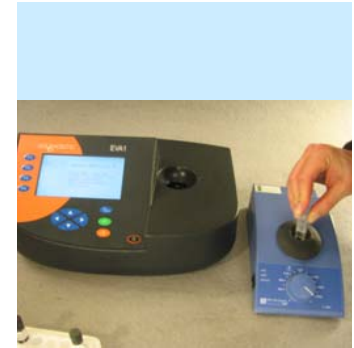


**4**

Press **F1(run)** then immediately add **100 µl** Buffer2, by the **blue pipette**.

### Important!

Aspirate **100 µl** Buffer 2 to tip before pressing **F1**.



**5**

Immediately vortex vigorously for 10 sec – see countdown on screen. Replace in cuvetteholder, arrow against arrow. Readout result after 2.45min – see countdown on screen

### Important!

Before replacement of cuvette, hit the cuvette gently against a hard surface to assure enough liquid is in the measuring area at the bottom