



**PROFICIENCY TESTING SERVICE**  
**AMERICAN ASSOCIATION OF BIOANALYSTS**  
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**PARTICIPANT STATISTICS**

**SECOND QUADRIMESTER 2018**

**HIV-1 Waived/Rapid**

**anti-HIV-1 Waived**

Method	Specimen 1			Specimen 2		
	Neg	Pos	Ind	Neg	Pos	Ind
Alere Determine	14					14
Chembio HIV 1/2 STAT-PAK	3					3
OraSure OraQuick ADVANCE, Wv	10					10
<b>Total Population</b>	<b>32</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>32</b>	<b>0</b>
<b>Flagging</b>		***	***	***		***

**p24 Antigen**

Method	Specimen 1			Specimen 2		
	Neg	Pos	Ind	Neg	Pos	Ind
Alere Determine		13		13		
<b>Total Population</b>	<b>0</b>	<b>13</b>	<b>0</b>	<b>13</b>	<b>0</b>	<b>0</b>
<b>Flagging</b>	***		***		***	***

\* Alere Determine users appear to correctly have identified the positive p24 Antigen, but have largely called the screen negative. Methods that report antigen and anti-HIV separately should have reported Negative for Specimen 2.

Correct responses are defined as those reflecting agreement among 80% or more of all participants or referees. Unacceptable responses are indicated by "\*\*\*\*\*" on the Flagging line of each specimen.