

1 **Supplementary Information**

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13 **1.1 Summary of latent class analysis output**

Assay result	Assay +ve / Group N	Risk	Risk Ratio
<b>GP ELISA High Titre</b>			
Group A	11 / 20	0.550	17.4
Group B	3 / 95	0.032	(5.3 – 56.8)
<b>WB GP positive</b>			
Group A	10 / 20	0.500	23.7
Group B	2 / 95	0.021	(5.6 – 100.2)
<b>WB NP positive</b>			
Group A	17 / 20	0.850	5.8
Group B	14 / 95	0.147	(3.4 – 9.7)
<b>WB VP40 positive</b>			
Group A	4 / 20	0.042	14.2
Group B	12 / 95	0.600	(5.1 – 39.7)
<b>WB negative</b>			
Group A	0 / 20	0.789	-
Group B	75 / 95	0	

14 **Supplementary table 1:** Results of latent class analysis from GP-ELISA

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18 **1.2 Consort diagram of sample processing pipeline and selection process**

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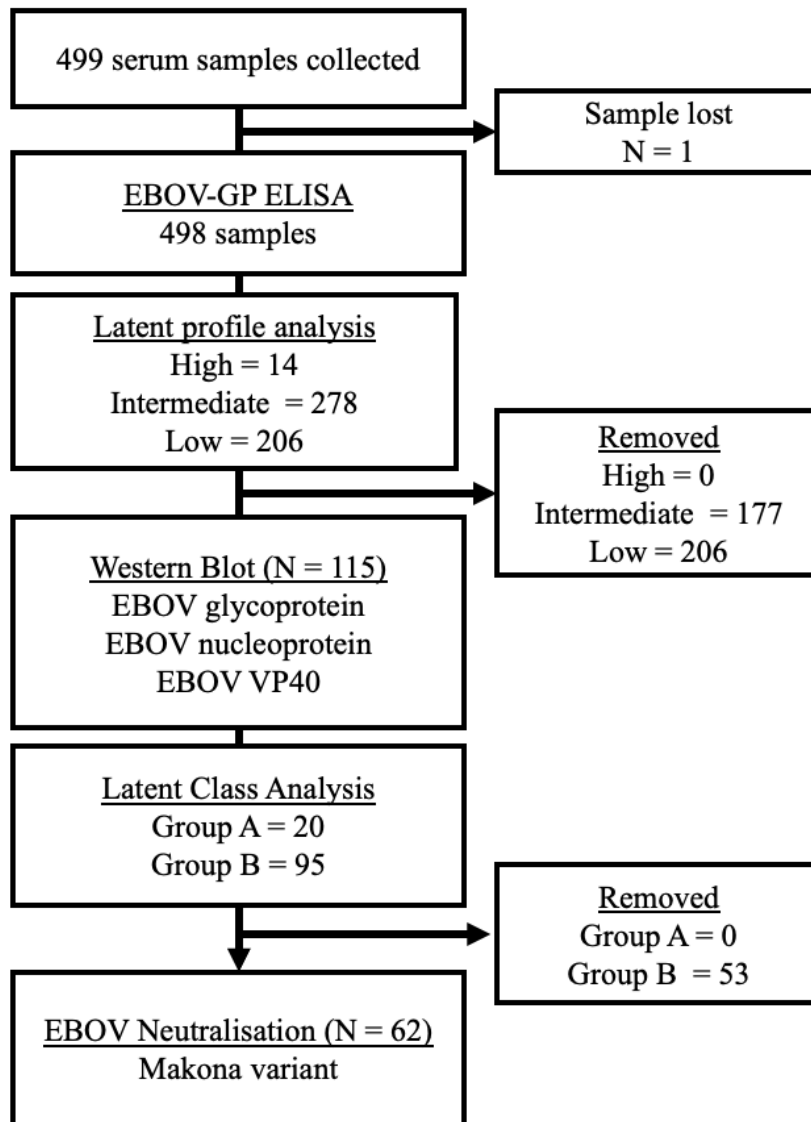
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36 **Supplementary figure 1:** Consort diagram of serum sample processing.

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43 **1.3 Outcomes of serological analysis stratified by village status**

	<b>Affected</b> (N=194)	<b>Unaffected</b> (N=304)	<b>Overall</b> (N=498)
<b>Anti-GP ELISA</b>			
High	9 (4.6%)	5 (1.6%)	14 (2.8%)
Intermediate	111 (57.2%)	167 (54.9%)	278 (55.8%)
Low	74 (38.1%)	132 (43.4%)	206 (41.4%)
<b>Latent class group</b>			
Group A	8 (16.3%)	12 (18.2%)	20 (17.4%)
Group B	41 (83.7%)	54 (81.8%)	95 (82.6%)
Not tested	145	238	383
<b>Neutralisation data</b>			
High	3 (10.3%)	2 (6.1%)	5 (1.0%)
Low	3 (10.3%)	2 (6.1%)	5 (1.0%)
Negative	23 (79.3%)	29 (6.1%)	52 (10.4%)
Not tested	165	271	436

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45 **Supplementary table 2:** Serological outcome data stratified by village status. Villages were

46 classified as affected or unaffected by 2013-2016 EBOV outbreak (see methods).

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60 **1.4 Ecological associations with EBOV immunological outcomes: Sensitivity analysis**

Predictors	Odds ratio	95% CI	p-value
<b>Outcome</b>			
LCA group A	20 / 498		
<b>Village status</b>			
Affected	Reference		0.86
Unaffected	1.09	0.40 – 2.94	
<b>Age</b>			
	1.03	0.99 – 1.06	0.12
<b>Closed forest</b>			
Shape index (500m)	0.28	0.08 – 0.98	0.02
<b>Vegetation</b>			
Perimeter area ratio (20,000m)	0.35	0.08 – 0.98	0.01
<b>Random Effects</b>			
ICC	0.02		
N village	38		

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62 **Supplementary table 3:** Multivariable generalised linear mixed effects model (binomial  
63 family) of immunological group defined by latent class analysis of ELISA and Western Blot  
64 analysis (Group A vs Group B). Success defined as Group A. Variables were selected using a  
65 forward, stepwise approach using AIC. P-values estimated by likelihood ratio test. Mixed  
66 effect models not used due to singular fit from village-level random intercepts. Two-sided test.

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Predictors	Estimate	95% CI	p-value
<b>Age</b>			
18 – 30	Reference		
31 – 50	0.19	-0.15 – 0.53	0.55
51 – 90	0.15	-0.25 – 0.55	
<b>Closed canopy cover</b>			
Perimeter area ratio (500m)	-0.63	-1.24 – -0.02	0.05
<b>Random Effects</b>			
ICC	0.14		
N <sub>village</sub>	24		

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77 **Supplementary table 4:** Multivariable mixed-effects linear regression of log<sub>2</sub> anti-EBOV-GP  
78 total antibody titre excluding all participants from villages with confirmed EBOV cases during  
79 2013-2016 outbreak (195/498; 39.2%). Variables were selected using a forward, stepwise  
80 approach using AIC. P-values estimated by likelihood ratio test. Two-sided test.

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Predictors	Odds ratio	95% CI	p-value
<b>Outcome</b>			
High titre GP-ELISA	14/498		
<b>Village status</b>			
Affected	Reference		0.24
Unaffected	0.35	0.14 – 1.56	
<b>Age</b>			
	1.02	0.98 – 1.06	0.35
<b>Vegetation</b>			
Perimeter area ratio (20,000m)	0.37	0.12 – 1.04	0.06

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96 **Supplementary table 5:** Multivariable generalised linear model (binomial family) of log<sub>2</sub> anti-  
97 EBOV-GP total antibody titre classified by finite mixture models (high titre individuals vs.  
98 intermediate and low titre individuals combined; see figure 1). Success defined as high titre  
99 individual. Variables were selected using a forward, stepwise approach using AIC. P-values  
100 estimated by likelihood ratio test. Two-sided test.

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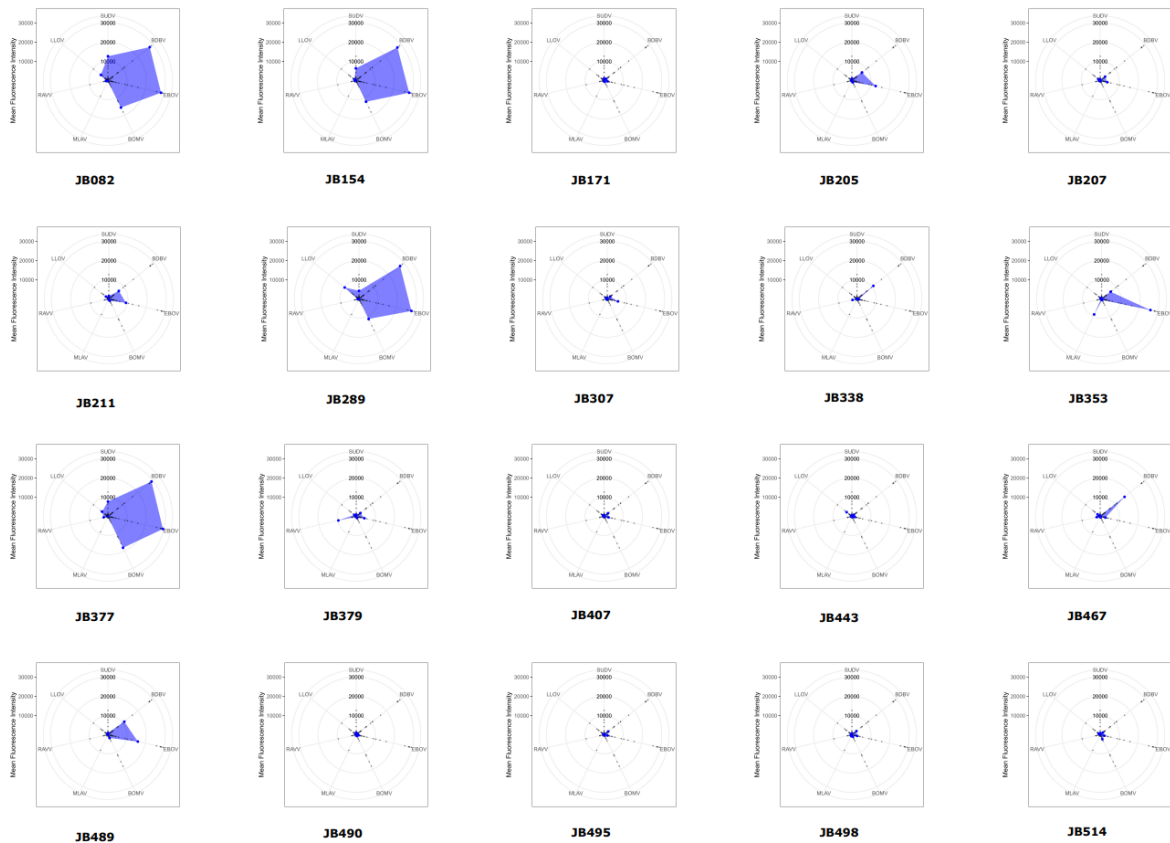
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114 **Supplementary Figure 2**

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117 Individual serological profile of participants within group A (n=20). Shows antigen-specific

118 total binding IgG antibody response (median fluorescence intensity) against a multiplexed

119 panel of filovirus antigens. Detected by Luminex-based multiplexed microsphere binding

120 immunoassay