CHR24



Supplementary Figure 1 Map of AGM (*Chlorocebus sabaeus*) immunoglobulin heavy chain locus on chromosome (CHR) 24. Representation of the identified functional and potentially functional, open reading frames, and pseudogenes of AGM V_H , D_H and J_H germline segments along with the Ig constant region genes of Ig M, D, G, A and E located on CHR24. Four additional functional AGM V_H gene segments that were identified from the unassembled contigs are not showed here (see Supplementary Data 1).

	50	100	
	I	1	
HuIgG1	${\tt ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSSKSTSGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSSKSTSGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTING {\tt ASTKGPSVFPLAPSS}{\tt ASTKGPSVFPLAPSSSLGTQTING {\tt ASTKGPSVFPLAPSS}{\tt ASTKGPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKGPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPSSCSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTFFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPS}{\tt ASTKFPSVFPLAPSSSSVFPLAPS$ {\tt ASTKFPLAPSVFPLAPSVFPLAPSSSVFPLAPS{\tt ASTKFPAPSVFPLA	IYICNVNHKPSNTKVDKKV <u>E</u> -	
RhIgG1	RES		
AGMIgG1	RES		
HuIgG2	C.RES	TD	
RhIgG2	SC.RQSQ.	VV.ET.GL	
AGMIgG2			
HuIgG3	C.R	T	
HuIgG4	C.RES		
RhIgG3	SC.RQS	VV.ERF	
RhIgG4	SRES	.VV.ERF	

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21	υ	υ	

	Hinge region	ł
HuIgG1	<u>PKSCDKTHTCPPCPAPELLGGP</u> SVFLFPPKPKDTLMISRTPEVTCVVVDVSHED)
RhIgG1	KTCG	•
AGMIgG1	<u>KTCG</u> N	•
HuIgG2	PVA	
RhIgG2	PCRS	S
AGMIgG2	Q.E	S
HuIgG3	KTPLGDTTHTCPRCPEPKSCDTPPPCPRCPEPKSCDTPPPCPRCPETPPPR	•
HuIgG4	YGPP.SF	•
RhIgG3	TPPCGDTT-PP	
RhIgG4	трР	13

					250				300
					1				1
HuIgG1	PEVKFNWYVDGVE	VHNAKTKPI	REEQYNSTYRV	SVLTVLHQDWLN	GKEYKCKVSN	KALPAPIEKTI	SKAKGQPRE	PQVYTLPPSRDELTKNQVS	LTCLVK
RhIgG1	.DN.A.	H.Q	т	T	T	Q	D	E	
AGMIgG1	Q	Q	т	R	T	Q	D	AE	
HuIgG2	Q		FF	v		.G	T	E.M	
RhIgG2	.D	Q	<mark>.</mark> .F	T	T	RQV	T	P.E	
AGMIgG2	Q	Q	N	T	T	RQV	7 T	P.E	I.
HuIgG3	Q.K		F				T	E.M	
HuIgG4	Q		F			.GSS		QE.M	
RhIgG3	QA.	H.Q		T	T	.G		IPQE	T
RhIgG4	0	0		T	T	.G		IPOE	T

350 |

HuIgG1	GFYPSDIAVEWESNGQPENNYKTTPP	/LDSDGSFFLYSKLTVDK	SRWQQGNVFSCSVMHEA	LHNHYTQKSLSLSPGK
RhIgG1	VST	Y		
AGMIgG1	VT	¥	T	. v
HuIgG2		4		
RhIgG2	VAT	¥	T	
AGMIgG2	VRT	¥	T	. v
HuIgG3	N	4	I	RF
HuIgG4			E	L
RhIgG3	T	¥	T	
RhIgG4	T	N.	PI.T	

)	5 0	100
HulgAl	${\tt SPTSPKVFPLSLCSTQPDGNVVIACLVQGFFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACLVQGFPQEPLSVTWSESGQGVTARNFPPSQDASGDLYTTSSQLTLPATQCLAGKSVTCHVIACUVACUVACUVACUVACUVACUVACUVACUVACUVACUV$	KHYTNPSQDVTVPC
HuIgA2	D.PQVN	
RMIgA_AI	KEGS.~VKNKAVIRGAP.SE	ERA
RMIgA_AII	REGS.~VNK.AVIRGAP.SEI	ERG
RMIgA_BI	RNS.~V	E
RMIgA_BII	RS.~	E
RMIgA_CI	RNS.~VNKADVIGAP.SEI	E
RMIgA_CII	REGS.~V	EARA
RMIgA_DI	REGS.~VNK.AVIRGAP.SE	ERG
RMIgA DII	REGS.~VNKAVI	EARA
RMIGA_RJZ7	KGSVNNK.AVIGP.TE	E
RMIGA RUA8		E
AGMIGA	EGSV	EA

		15 0		20 0
	Hinge region			1
HuIgA1	PV <u>PSTPPTPSPSTPPTPSPS</u> CCHPRLSLHRPALEDLLLGSEANLTCT	LTGLRDASGVTFTWTPSSG	KSAVQGPPERDLCGCYSVSSVLPGCAEPW	NHGKT
HuIgA2	P.P		Q	E.
RMIGA_AI	R~VPP.KCLK.DKR	PA	.NQSH.P	. NRV .
RMIgA_AII	R~VPP.NCRL.DKR	PAA	.NQSH.P	.NRV.
RMIgA_BI	VP.ICER	K.PA	.NQS.KP	.NRE.
RMIgA_BII	V~.QPK.CLDKR	PA	.NQSH.P	.NRV.
RMIgA_CI	VP.ICER	PA	.NQS.KP	.NRE.
RMIgA_CII	V~SQ.K.CLDER	PA	.NQS.KP	.NRV.
RMIgA DI	R~VPP.NCRL.DKR	PAA	.NQSH.P	.NRV.
RMIgA_DII	V~SQ.K.CLDER	PA	.NQS.KP	.NRV.
RMIGA_RJZ7	RD.QPK.CLDK.QR.	PAA	.NQSH.P	.NRE.
RMIGA RUA8	VP.ICER	K.PA	.NQS.KP	.NRE.
AGMIGA	V~PKTCFDER	PA	.NQSH.P	.N.V.

25 0	
1	

0

		1
HuIgAl	FTCTAAYPESKTPLTATLSKSGNTFRPEVHLLPPPSEELALNELVTLTCLARGFSPKD	CDVLVRWLQGSQELPREKYLTWASRQEPSQGTTTFAVTSILRVA
HuIgA2	HLNIT	
RMIgA_AI	EHLE.QI	EK.TEQDEK
RMIgA_AII	.NNHLIE.	EK.TEQVDEK
RMIgA_BI	NHLEIE.	EK.TEQDEK
RMIgA_BII	EH.QLI	EK.TEQDEK
RMIgA_CI	NHLEIE.	EK.TEQDEK
RMIgA_CII	K.EHLEIE.	EK.TEQDEK
RMIgA_DI	.NNHLIE.	EK.TEQVDEK
RMIgA DII	K.EHLEIE.	EK.TEQDEK
RMIGA RJZ7	NHLIE.	SK.TEQDEK
RMIGA_RUA8	NHLEIE.	EK.TEQDE
AGMIGA	NHLE.QITE.	EK.TEQDEKP

	35 0	
HuIgA1	AEDWKKGDTFSCMVGHEALPLAFTQKTIDRLAGKPTHVNVSVVMAEVDGTCY	
HuIgA2		
RMIgA_AI	······	
RMIgA_AII		
RMIgA_BI	······	
RMIgA_BII		
RMIgA_CI	······	
RMIgA_CII		
RMIgA_DI	······	
RMIgA_DII		
RMIGA_RJZ7	······	
RMIGA_RUA8	······	
AGMIGA		

С

	5 0	100
		1
HuIgD	APTKAPDVFPIISGCRHPKDNSPVVLACLITGYHPTSVTVTWYMGTQSQPQRTFPEIQRRDSYYMTSSQLSTPLQQWRQGEYKCVVQHTA:	SKSKKE-I
RhIgD	AA.QLN.KHLI.N.IMTE.EGTAPTPS.QHT.KP	.DTSKT
AGMIgD	.ST.A.ELV-TPS.RHT.KP	.NTNKT
	150	200
Hu TaD		CERACCOL
BhIgD		SF VVGSDL
AGMIqD	.HYPSG.SRSNRE.EDEKQ.GRIPLF	
2	-	
	25 0	30 0
	1	
HuIgD	KDAHLTWEVAGKVPTGGVEEGLLERHSNGSQSQHSRLTLPRSLWNAGTSVTCTLNHPSLPPQRLMALREPAAQAPVKLSLNLLASSDPPEAA:	SWLLCEVS
RhigD	QS	
AGMIGD	QS	
	25 0	
HulgD	${\tt GFSPPNILL} MWLEDQREVNTSGFAPARPPPQPRSTTFWAWSVLRVPAPPSPQPATYTCVVSHEDSRTLLNASRSLEVSYVTDHGPMK$	
RhIgD	DNWTTH.TG.MG.T	
AGMIgD	D	
d	50	100
		100
HulgM	GSASAPTLFPLVSCENSPSDTSSVAVGCLAQDFLPDSITLSWKYKNNSDISS-TRGFPSVLRGGKYAATSQVLLPSKDVMQGTDEHVVCKVQHPN	GNKEK
RhIgM		Q
AGMIgM		Q
	150	200
HuTaM	I NVPLPVTARLPPKVSVFVPPRDGFFGNPRKSKLTCOATGFSPROTOVSWLREGKOVGSGVTTDOVOAFAKESGPTTYKVTSTLTTKESDWLGOSM	FTCRV
RhIgM	L.R.N	
AGMIgM	VQNIVEEEIEFVS.RSV	· · · · ·
	250	300
HutoM		WNSCE
RhIaM	K.V.V.G.NP. G.	WNSGE
AGMIgM		
-		
	35 0	40 0
HulgM	RFTCTVTHTDLPSPLKQTISRPKGVALHRPDVYLLPPAREQLNLRESATITCLVTGFSPADVFVQWMQRGQPLSPEKYVTSAPMPEPQAPGRYFA	HSILT
AGMICM	W R DM T	
Ashigh		
	43 0	
HulgM	VSEEEWNTGETYTC-VAHEALPNRVTERTVDKST	
RhIgM	DVV	

AGMIgMD......V......V.....

	50	100
	1	1
HulgE	${\tt ASTQSPSVFPLTRCCKNIPSNATSVTLGCLATGYFPEPVMVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTCDTGSLNGTTMTLPATTLTLSGHYATISLLTVSGAWAKQMFTCRVTGTGSLNGTTMTLPATTLTLSGHYATISLTVSGAWAKQMFTCRVTGTGSLNGTTMTLPATTLTLSGHYATISLTVSGAWAKQMFTCRVTGTGSLNGTTMTLPATTTTTTLSGAWAKQMFTCRVTGTGSLNGTTMTTGTGSLNGTTMTTGTGSLNGTTMTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT$	AHTPSSTDWVDNK
RhIgE	.IFIPH.AETH.	VA.KEV
AGMIgE	IFIPH.A.DETH.	VA.KEV
	15 0	200
	I	1
HulgE	${\tt TFSVCSRDFTPPTVKILQSSCDGGGHFPPTIQLLCLVSGYTPGTINITWLEDGQVMDVDLSTASTTQEGELASTQSELTLSQKHWLSSCDGGGHFPPTIQLLCLVSGYTPGTINITWLEDGQVMDVDLSTASTTQEGELASTQSELTLSQKHWLSSCDGGHFPPTIQLLCLVSGYTPGTINITWLEDGQVMDVDLSTASTTQEGELASTQSELTLSQKHWLSSCDGGHFPPTIQLLCLVSGYTPGTINITWLEDGQVMDVDLSTASTTQEGELASTQSELTLSQKHWLSSCDGGHFPTIQLLSQNGVDVDJSTASTTQEGELASTQSELTLSQKHWLSSCDGGYTNGTWLEDGQVMDVDJSTASTTQEGELASTQSELTLSQKHWLSSCDGGYTNGTWLEDGQVMDVDJSTASTTQEGELASTQSELTLSQKHWLSSCDGFFFTIQJLJSQHWLSTASTQSEJQSGYTPGTINITWLEDGQVMDVDJSTASTTQEGELASTQSELTLSQKHWLSSCDGFFFTTQFFFTTQFFFFTTQFFFTTQFFFTTQFFTTQFF$	3DRTYTCQVTYQGH
RhIgE	.GN	. T
AGMIgE		G
	25 0	300
	1	
HulgE	${\tt TFEDSTKKCadsnprgvsaylsrpspfdlfirksptitclvvdlapskgtvnltwsrasgkpvnhstrkeekqrngtltvtstlpvgrasgkpresservers$	TRDWIEGETYQCR
RhIgE	.YNP.IPTT.K	7.Q
AGMIgE	.Y	7.Q
	35 0	40 0
	1	
HulgE	$\tt vth phipralmrsttktsgpraapevy afat pewpgsrdkrtlaclign fmpedisvgw lhnev glpdarhsttgprktkgsgff vission of the standard $	SRLEVTRAEWEQK
RhIgE	VMVVKLE	K
AGMIgE	VMVVKLE	ĸ
	43 0	
HulgE	DEFICRAVHEAASPSQTVQRAVSVNPGK-	
RhIgE	X	
AGMIgE	WIQL.X	

Supplementary Figure 2 Sequence alignment of AGM, RM, and human Ig constant regions. Panel a shows sequence alignment of IgGs from the three species. The underlined text shows the hinge region of IgGs of three species. Sequences highlighted in yellow represent the identical residues in the hinge region between AGM IgG1 and RM IgG1. Sequences in red show the shared unique mutations in the C_{H3} domain of AGM IgG2 and RM IgG2. Panel b shows sequence alignment of IgAs of three species. The hinge region of human IgA1 is highlighted in red. Ten RM intra-species heterogenetic IgA alleles were used in the alignment. The sequence alignments of IgD (c), IgM (d) and IgE (e) of three species are also represented.

е

						50					100
						1		μ Ε 1		μ E 5	
AGM Igh Eu	CATCATAAATC	AAGITIAT		ATICACICAA	CICCA	GOCCAICA	IGAGIIAGAGI	AAGATOGCTO	CATCOCCET	CITIGOGCACCC	ACA
Rhesus IgH Eu									.G	c	G
Human IqH Eu					A.	A	A			œ	
						1 50					20 0
	шE2	шA	щEЗ	ш	в	1					1
AGM TOTH FU	GCAGGIGGCAG	TAACTAC		GICIATUTA			CAACTIGGTAGA		IGGIIGA	GAGAIGGITT	m
Rhesus IaH Eu	G					C	.G.T A.	G			
Human Toff Eu			с. С								т.
						•••••					
											30 0
					uF4	Octamor	motif				1
ACM TO HE	സ്റ്റാനസംവ							ممصحمص	22000222220		י_חידע_י
Phoenic Toff Fu	cidebuung	licidine	C	<u></u>	1001001			7	GLUGIN		, 11 m
Riesus_ign_in	~~~~					·····	•••••		•••••	۰۰۰۰۰ ۳	1
Hullen_Igh_Eh	.CA	• • • • • • • • •	• • • • • • • • • • • •		• • • • • • •	.A	•••••		•••••	A	
						050					
						35 0		_			
			CACA BO	x			TATA	Box			
AGM_IgH_Eµ	TITTTAACIZ	AICOGAAI'I	TCATTICATI	CITAGCITAI	CAACIGO	TAGITIGI	GCAAACAGG <mark>AT</mark>	AICAACIICIA	AACIGCAIT	ATTT	
Rhesus_IgH_Eµ		A	C		•••••	• • • • • • • • •			C.		
Human_IgH_Eµ		T					C				

Supplementary Figure 3 Comparison of IgH intronic enhancer core region ($cE\mu$) of AGM, **RM and human.** The $cE\mu$ elements, Octamer motif, CACA box, and TATA box were highlighted and underlined. RM $cE\mu$ intronic enhancer region was identified from the assembly on Oct. 2010 (BGI CR_1.0/rheMac3).



Supplementary Figure 4 Comparison of the centromeric region of V_H locus in AGM and human genomes. Mapping of AGM VH gene segments in reference of human genome indicated that a large DNA fragment is potentially missing in the cerntromeric side of AGM V_H locus, in which VH6, VH7 and VH2 might be located by comparing human V_H locus.



Supplementary Figure 5 AGM, RM and human immunoglobulin kappa chain variable gene (V_{κ}) homology. AGM V_{κ} genes are genetically more similar to those of RM than human. Panel a shows numbers of functional V_{κ} genes from each species. Panel b shows phylogenetic comparison of V_{κ} families of AGM, RM and human. AGM V_{κ} 3-13 is located out of V_{κ} 3 family branch and has a human homologue gene as IGKV3-NL5*01 F. Panel c shows joint radial phylogenic tree of AGM (red lines), RM (green lines), and human (blue lines) V_{κ} genes; the AGM V_{κ} genes that are genetically more similar to RM than human V_{κ} homologs are indicated with star (*). Panel d indicates the number of AGM V_{κ} genes that are genetically more similar to RM than human V_{κ} homology of AGM V_{κ} genes to those of RM and human. Percent homology between V_{κ} germline genes between AGMs, humans and RMs were compared (p<0.0001) using an Exact Wilcoxon rank sum test. The data in Panel e shows means±sd.



Supplementary Figure 6 AGM, RM and human immunoglobulin lambda chain variable gene (V_{λ}) homology. AGM V_{λ} genes are genetically more similar to those of RM than human. Panel a shows number of functional V_{λ} genes in each species. Panel b shows phylogenetic comparison of AGM, RM and human V_{λ} families. AGMLV1-87 is located out of $V_{\lambda}1$ family branch, and has the human homologue gene as IGLV1-62*01 P. Panel c shows joint radial phylogenic tree of AGM (red lines), RM (green lines), and human (blue lines) V_L genes; the AGM V_{λ} genes that are genetically more similar to RM than human V_{λ} homologs are indicated with star (*). Panel d shows number of AGM V_{λ} genes genetically more similar to RM (grey box) or human (white box) than AGM. Panel e shows percent homology of AGM V_{λ} genes to those of RM and human. Percent homology between V_{λ} germline genes between AGMs, humans and RMs were compared (p<0.0001) using an Exact Wilcoxon rank sum test. The data in Panel e shows means±sd.

CHR14

Centromeric



Supplementary Figure 7 Map of AGM (*Chlorocebus sabaeus*) immunoglobulin κ locus on chromosome (CHR) 14. Shown are the identified functional, potentially functional, open reading frames, and pseudogenes of AGM V_{κ} and J_{κ} germline segments along with the Ig kappa chain constant region gene located on CHR14.

Supplementary Figure 8 Map of AGM (*Chlorocebus sabaeus*) immunoglobulin λ locus on chromosome (CHR) 19. Shown are the identified functional, potentially functional, open reading frames, and pseudogenes of AGM V_{λ} and J_{λ} germline segments along with the Ig lambda chain constant region gene located on CHR19.

Supplementary Figure 9 Gating strategy for total memory B cell populations in blood and milk of SIV/HIV-infected, AGM, RM, and human. Total memory B cells of SIV-infected AGM and RM were gated as CD3⁻/CD20⁺/IgD⁻ (top two rows). Total memory B cells from blood and milk of HIV-infected human were gated as CD3⁻/CD19⁺/IgD⁻ (bottom row).

Supplementary Figure 10 Genetic distribution of D_H and J_H of AGM Env-specific and non Env-specific mAbs. The distribution of D_H gene families (Panel a) and J_H gene families (Panel b) isolated from AGM blood and milk samples.

Supplementary Figure 11 The phylogenetic relationship of 10 purified SIV Env-reactive mAbs to AGM heavy and light chain variable germline genes. The phylogenetic relationship of V_H (Panel a), V_{κ} (Panel b) and V_{λ} (Panel c) genes of 10 purified SIV Env-reactive mAbs to AGM V_H germline genes.

Supplementary Figure 12 Schematic diagram for the generation of linear full-length Ig heavy- and light-chain gene constructs. (a) Schematic representation of cloning rhesus immunoglobulin heavy and light chain variable genes by semi-nest PCR. (b) The Ig H/L PCR products were sequenced and the functional genes were then assembled into full-length linear expression cassettes with CMV promoter and BGH Poly A by overlapping PCR.

Supplementary Table 1 Demographic, virologic, and immunologic characterization of

		Number of Antibodies Isolated				Patient Characteristics					
#	Patient ID	Blood	Milk	Total	Age (yrs)	Sex	Country of Origin	Specimen Draw Date	Viral Load (RNA copies/mL)	CD4 count (cells/mm ¹)	ARV
1	711-12-071-8	93	0	93	21	Μ	USA	24-Jun-10	<40	528	Yes
2	700-12-051-1	68	0	68	59	Μ	USA	9-Oct-09	16,800	957	No
3	714-90-060-8	22	0	22	41	F	Malawi	3-Dec-07	<400	275	Yes
4	715-90-060-5	0	3	3	28	F	Malawi	2-Apr-08	32,164	519	No
5	714-09-360-1	0	31	31	18	F	Malawi	10-Mar-08	60,335	452	No
6	715-90-030-1	6	5	11	32	F	Malawi	26-Mar-08	75,934	406	No
7	715-90-120-9	14	2	16	25	F	Malawi	18-Jun-08	174,976	153	No
			Blood	203							
Antibodies Isolated		ated	Milk	41							
			Total	244	-						

HIV-1-infected individuals used for antibody isolation from blood and breast milk

¹ VL and CD4 count performed at the time of PBMC/milk cell isolation

Supplementary Table 2 Amino acid sequences of the upper, core and lower hinge of IGG

	Upper Hinge	Core	Lower Hinge
HU_IgG1	EPKSCDKTHT	CPPC	PAPELLGGP
RM_IgG1	EIKTCGGGSKPPT	CPPC	PAPELLGGP
AGM_IgG1	EIKTCGGGGKPPT	CPPC	PAPELLGGP
HU_IgG2	ERK	CCVECPPC	PAPPVAGP
RM_IgG2	GLP	CRSTCPPC	PAELLGGP
AGM_IgG2	not available	not available	ELLGGP (partial)
HU_IgG3	ELKTPLGDTTHT	CPRCP EPKSCDTPPPCPRCP EPKSCDTPPPCPRCP EPKSCDTPPPCPRCP	APELLGGP
RM_IgG3	EFTPPCST	CPPC	PAPELLGGP
HU_IgG4	ESKYGPP	CPSC	PAPEFLGGP
RM_IgG4	EFTPP	CPPC	PAPELLGGP

molecules from AGM (Chlorocebus Sabaeus), RM, and human.

Primers	Sequences	Tm
RT primers		
G_RT	TCTTGTCCACCTTGGTGTTG	51.8
M-RT	TGAATTTCCAGGAGAAAGTGATG	51.7
A_RT	TGTTTCCRGATTTTGAGATGGTG	53.5
D_RT	AAGGTCTTCTCCTTGCTTGTATC	53.5
E_RT	TTGGAATCTGCACACTTCTTG	50.5
CK_RT	TCTCTGGGATAGAAGTTATTCAG	51.7
CL_RT	CTTGTTGTTGCTCTGTTTGGAG	53
PCR1 primers		
IgH primers		
VH1+7_EXT1	CACCATGGACTGGACCTGGAGGMTCCTC	67.9
VH1+7_EXT2	CACCATGGACCTGACCCGGAGGATCCTTTTC	72.4
VH2_EXT	CACCATGGACACGCTTTGCTCCACRCTC	68
VH3_EXT1	CATGGAGTTgGGGCTGAGCTGGGTYTTCC	71.3
VH3_EXT2	CATGGAGTTgGGGCTGAGYTGGGTTTTCC	69.7
VH3_EXT3	CATGGAGTTTGGGCTGAGCTGGR	61.1
VH3_EXT4	CATGGAGTTTGGGCTGAGCTKGGTTTTYC	66.8
VH4_EXT1	ACCATGAAGCACCTGTGGTTCTBCCTCCTCC	69.7
VH4_EXT2	CACCATGAAGCACCTGKGGTTCTTY	60.1
VH5_EXT	CACCATGGGGTCAACTGCCMTCCTC	65.3
VH6_EXT	CCATGTCTGTCTCCTTCCTCATCGTCC	63.8
A_EXT	GAAGAAGCCCTGGACCAGGCAGGC	66.6
G-EXT	AAGGTGTGCACGCCGCTGGTCAG	66.9
M_EXT	GTCGGGAAGGAAGTCCTGTGCGAGG	66.5
D_EXT	TCCCCAGGTGCCAGGTGACAGTCAC	66.8
E_EXT	ACGGTCAGCAAGCTGATGGTGGCA	66.3

Supplementary Table 3 Primer list for amplifying both AGM and rhesus macaque Ig pairs

IgK primers		
VK1_EXT1	CACCATGGACATGAGGGYCCC	58.4
VK1_EXT2	CACCATGGACATGAGGGTCCCCAGTC	66.9
VK1_EXT3	CACCATGGACATGAGGGTCCCCGGTtAtC	70.2
VK1_EXT4	CACCATGGACATGAGGGTCYCCGGTCAG	69.5
VK1_EXT5	CACCATGGACATGAGGGTCCCCGGTCAGCTYC	75.9
VK2_EXT1	CACCATGAGGCTCCCWGCTCAG	60.6
VK2_EXT2	CACCATGAGGCTCCCTGCTCAGCTYCTGGGGC	76.8
VK3_EXT1	CACCATGGAAGCCCCAGCTCRGCTTCTC	69.7
VK3_EXT2	CACCATGGAAGCCCCAGCACAGCTTCTC	70.4
VK4_EXT	CACCATGGTGTCACAGACCCAAGWCTTC	64.4
VK5_EXT	CACCATGGGATCCCAGGTTCACCTCCTCAG	71.4
VK6_EXT1	CACCATGGTGTCCCCATTGCAACTCCTG	69.9
VK6_EXT2	CACCATGtTGTCtCCATCACAACTCATtG	63.2
VK7_EXT	CACCATGGGGTCCTGGGCTCCTTTCCTG	72.2
CK_EXT	ACCTGATCCTCAGATGGCGGGAAGATG	66.7
IgL primers		
VL1_EXT1	CACCATGGCCTGGTCTCCTCTCSTCCTCAC	69.7
VL1_EXT2	CACCATGGCCTGGTCTCCTCTCCTtCTC	67.4
VL2_EXT	CACCATGGCCTGGGCTCTGSTCCTC	67.2
VL3_EXT1	CACCATGGCCGGGACCCYTCTCCTCCTC	73.1
VL3_EXT2	CACCATGGCCTGGACCCCTGTTCTGCTC	71.8
VL3_EXT3	CACCATGGCCtGGACCCcTCcCCTRCTC	73.1
VL4_EXT	CACCATGGCCTGGACCCCACTCCTCCTC	72.6
VL5_EXT	CACCATGGCCTGGACTCYTCTC	57.3
VL6_EXT	CACCATGGCCTGGGCTCCACTCCTCCTC	72.7
VL7_EXT	CACCATGGCCTGGACTCTGCTCCTCCTCC	72
VL8_EXT	CACCATGGCCTGGATGATGCTTCTCCTCG	71.6
VL11_EXT	CACCATGGCCCTGACTCCTCCTCCTCCTC	69.1
CL_EXT	TGCCATCTGCCTTCCAGGCCACTT	66.3
Primers	Sequences	Tm

PCR2 primers

IgH primers		
VH1+7_INT1	CCAAGCTGGCTAGCACCATGGACTGGACCTGGAGGMTCCTC	80.1
VH1+7_INT2	CCAAGCTGGCTAGCACCATGGACCTSAcccGGAGSATCCTTTTC	78.4
VH2_INT	CCAAGCTGGCTAGCACCATGGACACGCTTTGCTCCAC	78.9
VH3_INT1	CCAAGCTGGCTAGCACCATGGAGTTGGGGGCTGAGYTG	78.2
VH3_INT2	CCAAGCTGGCTAGCACCATGGAGTTTGGGCTGAGCTKG	78.5
VH4_INT	CCAAGCTGGCTAGCACCATGAAGCACCTGKGGTTC	75.3
VH5_INT	CCAAGCTGGCTAGCACCATGGGGTCAACTGCCMTCCTC	79
VH6_INT	CCAAGCTGGCTAGCACCATGTCTGTCTCCTCCTCATCGTC	77.8
A_ACD_Int	CAGGGCCGCTGTGCTCTCGGAGGTGCTCCTGCCCTCGAGG CTCAGCGGGAAGAC	76.3
A_BC_Int	CAGGGCCGCTGTGCTCTCGGAGGTGCTCCTGCTGCAGAGG YTCAGCGGGAAGAC	75.5
G_Int	CAGGGCCGCTGTGCTCTCGGAGGTGCTCCTGGAG	80.9
M_Int	CAGGGCCGCTGTGCTCTCGGAGGTGCTCCTACAGGAGACG AGGGGGAAAAGGG	61.2
D_Int	CAGGGCCGCTGTGCTCTCGGAGGTGCTCCTCGCTGATATG ATGGGGAACACATCC	62.1
IgK primers		
VK1_INT1	CCAAGCTGGCTAGCACCATGGACATGAGGGYCCC	76.6
VK1_INT2	CCAAGCTGGCTAGCACCATGGACATGAGGGTCYCCG	78
VK2_INT1	CCAAGCTGGCTAGCACCATGAGGCTCCCWGCTC	75.8
VK3_INT1	CCAAGCTGGCTAGCACCATGGAAGCCCCAGCWC	76.7
VK4_INT	CCAAGCTGGCTAGCACCATGGTGTCACAGACCCAAG	76.4
VK5_INT	CCAAGCTGGCTAGCACCATGGGATCCCAGGTTCACCTCC	80.3
VK6_INT1	CCAAGCTGGCTAGCACCATGGTGTCCCCATTGCAACTC	79
VK6_INT2	CCAAGCTGGCTAGCACCATGTTGTCtCCATCACAACTC	74.9
VK7_INT	CCAAGCTGGCTAGCACCATGGGGTCCTGGGCTCCTTTCC	81.5
CK_INT	TGGCGGGAAGATGAAGACAGATGGTG	66.2
IgL primers		
VL1_INT	CCAAGCTGGCTAGCACCATGGCCTGGTCTCCTCTC	76.7
VL2_INT	CCAAGCTGGCTAGCACCATGGCCTGGGCTCTGSTCC	79.5
VL3_INT1	CCAAGCTGGCTAGCACCATGGCCGGGACCCYTC	78.8

VL3_INT2	CCAAGCTGGCTAGCACCATGGCCtGGACCCCTS	77.4			
VL4_INT	CCAAGCTGGCTAGCACCATGGCCTGGACCCCACTCC				
VL5_INT	CCAAGCTGGCTAGCACCATGGCCTGGACTCYTCTC	75.4			
VL6_INT	CCAAGCTGGCTAGCACCATGGCCTGGGCTCCACTC	79.4			
VL7_INT	CCAAGCTGGCTAGCACCATGGCCTGGACTCTGCTCCTC	79.5			
VL8_INT	CCAAGCTGGCTAGCACCATGGCCTGGATGATGCTTCTC	78.6			
VL11_INT	CCAAGCTGGCTAGCACCATGGCCCTGACTCCTCTCCTC	78.9			
CL_INT	GTCACTGATCAGACACACTAGTGTGG	54.4			
Sequencing primers					
VHKLSEQ_F	CCAAGCTGGCTAGCACCATG	54.7			
VHSEQ_R	CTGTGCTCTCGGAGGTGCTCC	55.7			
VKSEQ_R	GGAAGATGAAGACAGATGGTG	48.4			
VLSEQ_R	TGATCAGACACACTAGTGTGG	45.1			
Primers for the construction of 1	inear cassettes				
Primers for P _{CMV} fragment ¹					
HV13220CMV-P-F262	AGTAATCAATTACGGGGTCATTAGTTCATAG	57.9			
CMV-P_R	CATGGTGCTAGCCAGCTTGGGTC	60.6			
Primers for Rh IgG1C fragment	2				
RhIGC_HF	GGAGCACCTCCGAGAGCACAGC	62			
BGH-R1235	TCCCCAGCATGCCTGCTATTGTC	61.9			
Primers for Rh IgKC fragment ³					
RhIGC_KF	CACCATCTGTCTTCATCTTCCCGCCA	65.7			
BGH-R1235	TCCCCAGCATGCCTGCTATTGTC	61.9			
Primers for Rh IgLC fragment ⁴					
RhIGC_LF	CCACACTAGTGTGTCTGATCAGTG	51.3			
BGH-R1235	TCCCCAGCATGCCTGCTATTGTC	61.9			
Primers for overlapping PCR					
HV13220CMV-P-F262	AGTAATCAATTACGGGGTCATTAGTTCATAG	57.9			
BGH-R1235	TCCCCAGCATGCCTGCTATTGTC	61.9			
 ¹ pcDNA3.1 plasmid as template ² Plasmid with rhesus whole IgG1 gene as template ³ Plasmid with rhesus whole IgK gene as template ⁴ Plasmid with rhesus whole IgL gene as template 					

Figure	Category	Test	Comparison	Variable	Compartment	Germline	Raw_P	fdr_p
Figure 2E	Homology	Wilcoxon	AGM:HU vs AGM:RM			V-Heavy	0.0000	0.0000
Figure 2E	Homology	Wilcoxon	AGM:HU vs AGM:RM			V-Kappa	0.0000	0.0000
Figure 2E	Homology	Wilcoxon	AGM:HU vs AGM:RM			V- Lambda	0.0000	0.0000
Figure 3B	Genomic	Fisher's Exact	HIV+/-: AGM vs HU		All		0.0000	0.0000
Figure 3B	Genomic	Fisher's Exact	HIV+/-: AGM vs RM		All		0.3095	0.5364
Figure 3B	Genomic	Fisher's Exact	HIV+/-: AGM vs HU		BM		0.0037	0.0113
Figure 3B	Genomic	Fisher's Exact	HIV+/-: AGM vs HU		PBMC		0.0000	0.0000
Figure 3B	Genomic	Fisher's Exact	HIV+/-: AGM vs RM		PBMC		1.0000	1.0000
Figure 3B	Genomic	Fisher's Exact	gp120+/-: AGM vs HU		All		1.0000	1.0000
Figure 3B	Genomic	Fisher's Exact	gp120+/-: AGM vs RM		All		0.0014	0.0052
Figure 3B	Genomic	Fisher's Exact	gp120+/-: AGM vs HU		BM		0.5403	0.7236
Figure 3B	Genomic	Fisher's Exact	gp120+/-: AGM vs HU		PBMC		0.4375	0.6319
Figure 3B	Genomic	Fisher's Exact	gp120+/-: AGM vs RM		PBMC		0.0004	0.0018
Figure 4	Genomic	Kolmogorov- Smirnov	ENV+ vs ENV-	HCDR3_Length			0.8996	1.0000
Figure 4	Genomic	Wilcoxon	ENV+ vs ENV-	H_Mutated			0.6547	0.7738
Figure 4	Genomic	Kolmogorov- Smirnov	gp120 vs Env cross- reactive	HCDR3_Length			0.3934	0.6016
Figure 4	Genomic	Exact Wilcoxon	gp120 vs Env cross- reactive	H_Mutated			0.5566	0.7236
Figure 4	Genomic	Kolmogorov- Smirnov	gp120 vs gp41	HCDR3_Length			0.9733	1.0000
Figure 4	Genomic	Exact Wilcoxon	gp120 vs gp41	H_Mutated			0.0401	0.0745
Figure 4	Genomic	Kolmogorov- Smirnov	gp41 vs Env cross- reactive	HCDR3_Length			0.6483	0.7738
Figure 4	Genomic	Exact Wilcoxon	gp41 vs Env cross- reactive	H_Mutated			0.3369	0.5475

Supplementary Table 4 Statistical analysis Raw p values and false discovery rate p values.

Supplementary Table 5 % blocking by AGM gp120-specific ADCC mediating antibodies

with undefined SIVsabgp120 fine-epitope specificity

% blocking				
	DH546	DH550	DH549	DH552
DH546-biotin	73.9	0	0	17
DH550-biotin	0	35.4	48.6	6.9
DH549-biotin	0	19.7	57.3	5.1
DH552-biotin	22.7	0	3.6	53.23

	animal ID	total B cell count (cells/µl) ¹	Mann- Whitney test	total memory B cell count $(cells/\mu l)^2$	Mann- Whitney test
AGM	013-89-10	220		118	
	013-90-10	21		8	
	013-91-10	36		11	
	013-92-10	116		59	
	013-93-10	12	m 0.07	4	m 0.26
	013-94-10	149	p = 0.07	83	$\mathbf{p}=0.20$
	206-96	289	-	109	
DM	257-97	95		40	
RM	402-98	662		376	
	403-98	251		66	

Supplementary Table 6 AGM and RM total B cell (CD3⁻CD20⁺) and memory B cell

(CD20⁺ IgD⁻ CD27⁺) count (cells/µl) at week 45 post infection

¹ denotes CD3⁻ CD20⁺ cells

² denotes CD20⁺ IgD⁻ CD27⁺