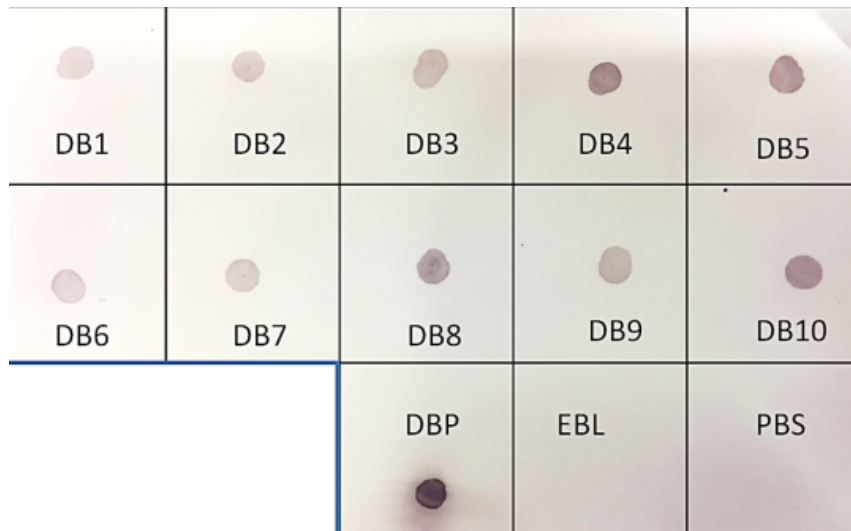


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Structural basis for inhibition of *Plasmodium vivax* invasion by a broadly neutralizing vaccine-induced human antibody

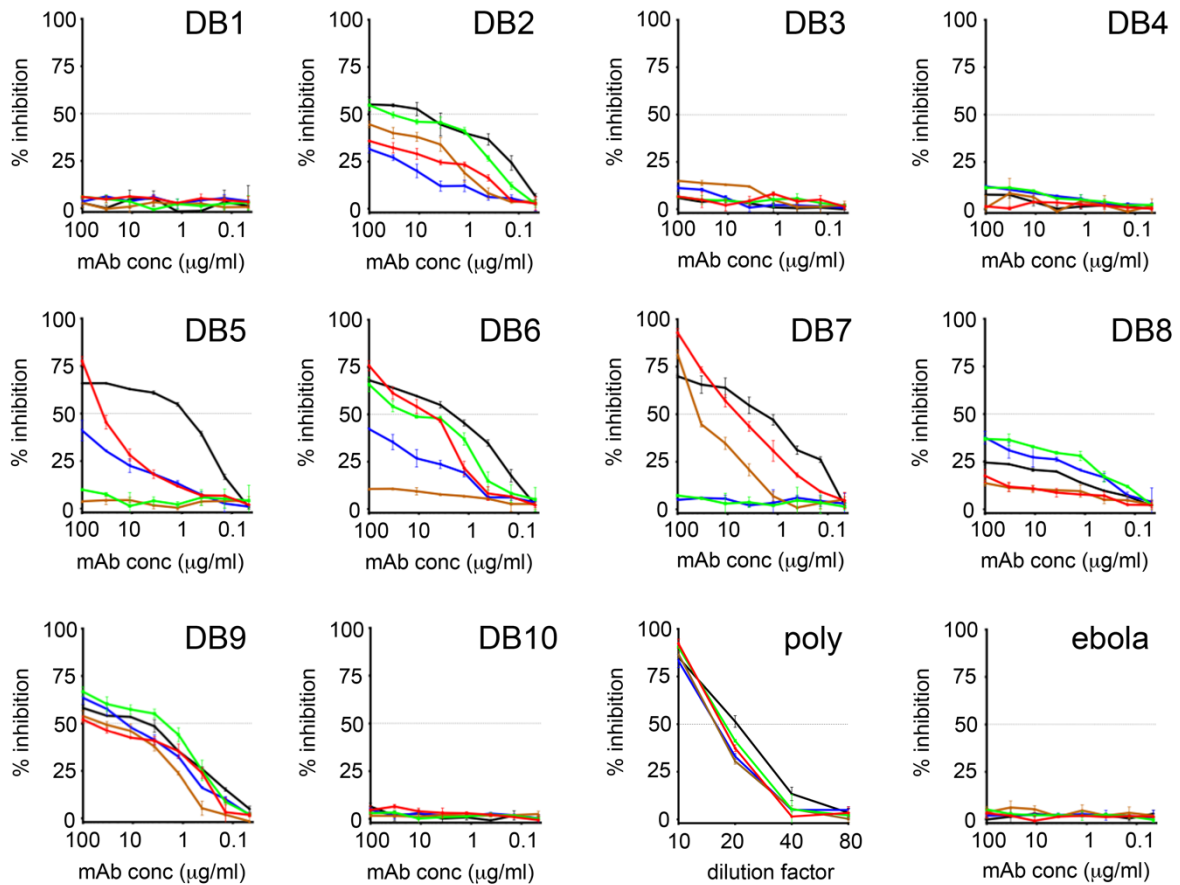
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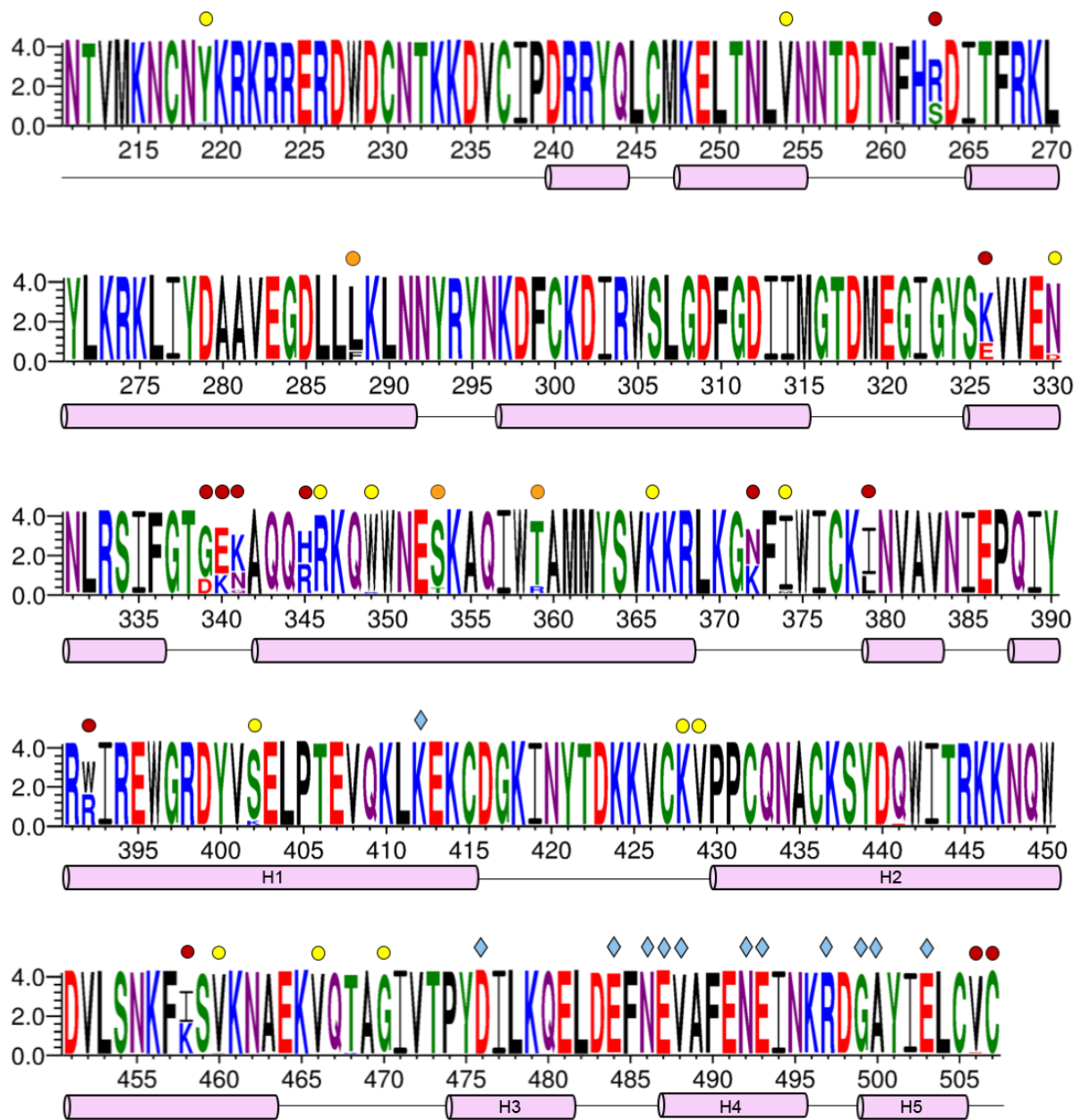
Supplementary Figure 1: Recognition by *Pv*DBP from culture supernatants by dot blot

Dot blot assay showing binding of the ten human anti-*Pv*DBP mAbs to native *Pv*DBP secreted in the supernatant of a short-term *in vitro* culture of *Plasmodium vivax*. The positive control was recombinant *Pv*DBP (DBP) and the two negative controls were a human anti- *Ebolavirus* GP IgG1 mAb (EBL) and PBS. This assay was conducted once due to limited quantities of culture supernatant.



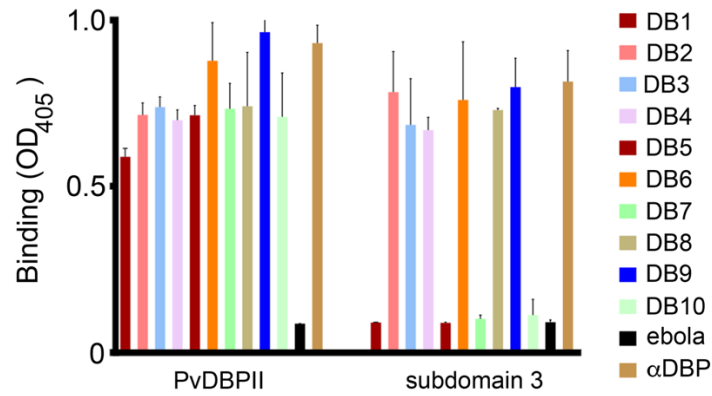
Supplementary Figure 2: Anti-*PvDBP2* mAb inhibition of the binding of recombinant *PvDBP2* variants to the recombinant N-terminal 60 amino acid DARC ectodomain

The binding of five naturally occurring variants of *PvDBP2* to the DARC ectodomain in the presence of increasing concentrations of each of DB1-DB10. The variants are Sall (red), AH (green), O (brown), P (blue) and HMP013 (black). “poly” is polyclonal human anti-*PvDBP2* serum from the VAC051 clinical trial¹⁹, while “ebola” is an anti-*Ebolavirus* recombinant human IgG1 mAb included as a negative control. Data points represent the mean of three technical replicates, while the error bars represent the standard deviation.



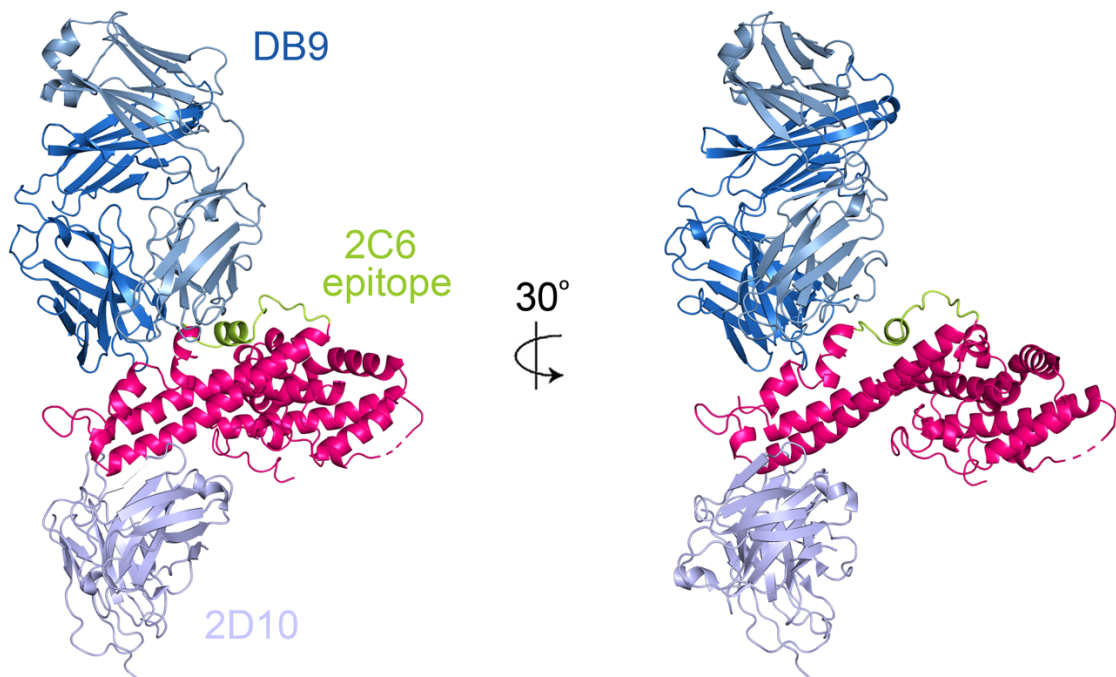
Supplementary Figure 3: Sequence logo representing sequence conservation across PvDBPII

Sequence logo derived from 383 sequences of PvDBPII from *Plasmodium vivax* isolates. Underneath the logo is the residue number from the Sall PvDBPII variant. Cylinders represent the location of helices while lines represent loops. Above the sequence, blue kites indicate residues which directly contact DB9. Yellow, orange and red circles represent residues with sequence entropies of 0.15-0.3, 0.3-0.45 and >0.45 respectively.



Supplementary Figure 4: Analysis of the binding of monoclonal antibodies to *Pv*DBPII subdomain 3

An ELISA of the ten anti-*Pv*DBPII mAbs (at 10 µg/mL) binding to recombinant *Pv*DBPII (left) and subdomain 3 (right). Columns represent the mean of three technical replicates, while the error bars represent the standard deviation and the assay was performed twice with similar results. Human polyclonal anti-*Pv*DBPII serum (αDBP) from the VAC051 clinical trial¹⁹ at 1:100 dilution and a human anti-*Ebolavirus* IgG1 mAb (ebola) at 10 µg/mL were used as positive and negative controls, respectively.



Supplementary Figure 5: Comparison of the epitope for DB9 with those of previously identified mouse monoclonal antibodies

*Pv*DBPII is shown in pink and DB9 in dark blue. The ScFv fragment of mouse antibody 2D10²⁶ is shown in lilac while the epitope of 2C6 in *Pv*DBPII, identified by hydrogen-deuterium exchange mass spectrometry is shown in green.

Supplementary Table 1: Genetic lineage of heavy and light chain variable regions from PvDBPII-specific mAbs

The allele usage, amino acid sequences and percentage of nucleotide substitutions relative to germline are shown.

mAb	Chain	Fv allele usage and amino acid sequence	Germline change
DB1	heavy	IGHV5-51*03 IGHD3-22*01 IGHJ3*02 EVQLVQSGAEVKKPGESLKISCKGSGYSFTDYWIGWVRQMPGKG LEWMGIIYAGDSDTRYSPSFQGQVTISADKSISTASLQWSSLKA SDTAMYYCARLAYDSSGYYYAFDIWGQGTMTVTVSS	1.7% (5/294)
	light	IGKV1-39*01,IGKV1D-39*01 IGKJ5*01 DIVMTQSPSSLSASVGDRVTITCRASQTISSYLNWYQQKPGKAP KLLIYAASSLQSGVPSRFSGSGSGTDFTLTISLQPEDFATYYC QQSYSTPLITFGQGRLEIKRTV	1.7% (5/286)
DB2	heavy	IGHV4-59*08 IGHD6-13*01 IGHJ3*02 QVQLQESGPGLVKPSSETLSLTCTVSGGSISSYYWSWIRQPPGKG LEWIGYISYTGSTNYPNPSLKSRVTISVDTSKNQFSLKLSSVTAA DTAVYSCARHFHSSSTAFAFDIWGQGTMTVTVSS	1.4% (4/293)
	light	IGKV3-20*01, IGKV3D-20*01, IGKJ1*01 EIVLTQSPGTLSSLSPGEGATLSCRASQSVSN SYLAWYQQKPGQA PRLLIYGASIRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYY CQQYGRSPRT	2.1% (6/289)
DB3	heavy	IGHV4-31*03 IGHD2-2*01,IGHD2-2*02,IGHD2-2*03 IGHJ4*02 QLQLVESGPGLVKPSQTLSTCTVSGGSTSSGGYYWNWIRQHGP KGLEWIGYIHNSGSTYYNPSLKSRIISVDTSKHQFSLRLRSVT AADTAEYYCARSQGYCSSSSCLLPRGYFDYWGPGLVTVTVSS	6.4% (19/297)
	light	IGLV1-51*01 IGLJ2*01,IGLJ3*01 QSALTQPPSVSAAPGQKVTISCSGSSSNIGNNFVSWYQLFPGTA PKLLIYDNNRPSGIPDRFSGSRSGTSATLGITGLQTGDEADYY CGTWDSSLSAVVFGGGTKLTVLGQP	3.7% (11/295)
DB4	heavy	IGHV1-69*06 IGHD6-6*01 IGHJ4*02 QLVQSGAEVKKPGSSVKVCKASGDTSSSYAISWVRQAPGQGLE WMGGIIPFGTANYAQKFQGRFTITAHKSTSTAYMELSSLRSD TAVYYCARDGGHHGQLVFDYWGGTGLVTVTVSS	2.7% (8/295)

	light	IGKV1-16*02 IGKJ4*01 DIQLTQSPSSLASVGDRTITCRASQVISNYLAWFQQKPGKAP KSLIYAASSLQSGVPSKFSGSGSGTDFTLTISSLQPEDFATYYC QQYNSYPLTFGGGTKVEIRRTV	1.4% (4/284)
DB5	heavy	IGHV1-46*03 IGHD1-26*01,IGHD4-11*01,IGHD4-4*01 IGHJ6*02 QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYMHWVRQAPGQG LEWMGIINPSGGSTSYAQKFQGRVTMTRDTSTSTVYMELSSLRS EDTAVYFCARDNSEGAAYSYYYYYGMDVWGQGTTVTVSS	0.7% (2/296)
	light	IGLV1-44*01 IGLJ3*02 QSALTQPPSASGTPGQRTISCSGSSSNIGSNTVNWYQQVPGTA PKLLIYSNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYY CAAWDDSLNGPRFGGGTKLTVLGQP	1% (3/296)
DB6	heavy	IGHV1-2*02 IGHD3-3*02,IGHD5-12*01,IGHD6-13*01 IGHJ6*02 QLVQSGAEVKKPGASVKVSCKASGYSTGYFLHWVRQAPGQGLE WMGWINPNSGGTKYAQKFQGRVTMTRDTSISTAYMELSLRSDD TAVYYCARGLRYSIAWYSDYGLDVWGQGTTVTVSS	2.4% (7/294)
	light	IGKV3-20*01 IGKJ4*01 EIVLTQSPGTLSLSPGERATLSCRASQSVTSTYLAWYQQKPGQA PRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYY CQQFGSSLTFGGGTKVEIKRT	1.4% (4/286)
DB7	heavy	IGHV4-39*01 IGHD3-10*01,IGHD3-10*02,IGHD5-18*01 IGHJ5*02 QVQLQESGPGLVKPSSETLSLTCTVSGGSIXSISYFWGWIRQPPG KGLEWIGSIYYSGSTYYNPSLKSRTVSVSDTSKNQFSLKLSSVT AADTAVYFCARRSLGYFFGPWGQGTTLVTVSS	2.7% (8/298)
	light	IGLV3-21*02 IGLJ2*01,IGLJ3*01 SYELTQPPSVSVAPGQTARITCGGNNIGSKRVHWYQQKPGQAPV LVVYDDSDRPSGIPERFSGSNSGNTATLTIXWVEAGDEADYYCQ LWDTSSDHPVFVGGGTKLTVLGX	2.4% (7/290)
DB8	heavy	IGHV1-3*01 IGHD6-19*01 IGHJ4*02 QVQLVQSGAEVKKPGASVKVSCKASGYTFSSYAMHWVRQAPGQR LEWMGWINAGNGNTKYSQKFQDRVTITRDTSASTAYMELSSLSS EDTAVYYCARSYRSSIGWFWMFYWGQGTTLVTVSS	2% (6/294)
	light	IGKV4-1*01 IGKJ2*01 DIVMTQSPDSLGVSLGERATINCKSSQSVLYSSNNKNYLAWYQQ	3% (9/302)

		KPGQPPKLLIYWASTRESGVPDRFSGSGSGTDFTLTITGLQAED VAVYYCLQYYSIPYTFGQGTKVEIKRTV	
DB9	heavy	IGHV4-39*02 IGHD1-1*01,IGHD1-14*01,IGHD1-20*01 IGHJ2*01 EVQLQESGPGLVKPSSETLSLTCTVSGGSVSSSTYYWGWRQPPG KGLEWIGSIYYSGSTYYNPSLKSRTVISVDTSKNQFSLKLSSVT AADTAVYYCARDGTGALDLWGRGTLTVSS	2.3% (7/298)
	light	IGKV1-5*03 IGKJ1*01 DIVMTQSPSTLSASVGDRVTITCRASQSISSWLAWYQQRPGKAP RLLIYKASSLLSGVPSRFGGSGSGTDFTLTISLQPDDEFATYHC QHYNTPWTFGQGTKVEIKRTV	4.4% (12/274)
DB10	heavy	IGHV3-21*01 IGHD3-10*01,IGHD3-10*02 IGHJ6*02 QVQLVESGGGLVKPGGSLRLSCAASGFTFSTYSMNWVRQAPGKG LEWVSSITSSSSYMDYADSVKGRFTISRDNANKSLYLQMTSLRA EDTAVYYCARDVAGPFYFYAMDVWGQGTTVTVSS	2.4% (7/295)
	light	IGLV2-14*01 IGLJ2*01,IGLJ3*01 NFVLTQPASVSGSPGQSITISCTGTSSDVGGYNFVSWYQQHPGK APKLMIEVSDRPSGVSNRFGSGSKSGNTASLTISGLQAED EADY YCSSYTSSSTVVFGGGTKLTVLGQP	0.7% (2/284)

Supplementary Table 2: Kinetic parameters for binding of antibodies to *Pv*DBP-II

mAb	k_{on} ($M^{-1}s^{-1}$)	k_{off} (s^{-1})	K_D (pM)
DB1	1.88E+07	6.46E-03	344
DB2	9.02E+05	6.05E-04	671
DB3	2.98E+06	4.27E-04	143
DB4	1.30E+06	4.39E-04	337
DB5	2.33E+06	4.33E-04	186
DB6	8.65E+06	2.53E-04	29.2
DB7	1.72E+07	1.13E-04	6.55
DB8	1.73E+06	3.92E-04	227
DB9	1.79E+07	3.57E-04	20.0
DB10	1.06E+07	5.05E-04	47.8

Supplementary Table 3: Data collection and refinement statistics

PvDBPII:DB9	
Data collection	
Space group	P6 ₂ 22
Cell dimensions	
<i>a</i> , <i>b</i> , <i>c</i> (Å)	173.58, 173.58, 169.88
α , β , γ (°)	90.0, 90.0, 120.0
Resolution (Å)	84.94 – 3.04 (3.09 – 3.04)
<i>R</i> _{PIM}	4.0 (46.9)
<i>I</i> / σI	12.8 (1.2)
Completeness (%)	100.0 (100.0)
Redundancy	19.6 (20.1)
Refinement	
Resolution (Å)	3.04
No. reflections	30703
<i>R</i> _{work} / <i>R</i> _{free}	20.0 / 23.8
No. atoms	
Protein	5754
Ligand/ion	0
Water	0
<i>B</i> -factors	
Protein	115.3
R.m.s. deviations	
Bond lengths (Å)	0.01
Bond angles (°)	1.24

All structures were determined from one crystal.

Values in parentheses are for highest-resolution shell.

Supplementary Table 4: List of contacts between PvDBPII and DB9

PvDBPII residue	Group	Fab Chain	Residue	Group	Interaction
K412	Side Chain	Heavy Chain	Y74	Side Chain	Hydrogen bond
D476	Side Chain	Light Chain	S75	Side Chain	Hydrogen bond
E484	Side Chain	Heavy Chain	E20	Main Chain	Hydrogen bond
E484	Side Chain	Heavy Chain	G45	Main Chain	Hydrogen bond
N486	Side Chain	Heavy Chain	R118	Side Chain	Hydrogen bond
E487	Side Chain	Light Chain	Y68	Side Chain	Hydrogen bond
E487	Side Chain	Light Chain	S75	Main Chain	Hydrogen bond
V488	Side Chain	Light Chain	Y68	Side Chain	Hydrophobic
V488	Side Chain	Light Chain	L74	Side Chain	Hydrophobic
N492	Side Chain	Heavy Chain	T121	Main Chain	Hydrogen bond
E493	Side Chain	Heavy Chain	T52	Side Chain	Hydrogen bond
R497	Side Chain	Light Chain	W51	Side Chain	Cation-pi
R497	Main Chain	Heavy Chain	T121	Side Chain	Hydrogen bond
R497	Side Chain	Heavy Chain	T121	Main Chain	Hydrogen bond
G499	Main Chain	Heavy Chain	T52	Main Chain	Hydrogen bond
A500	Main Chain	Heavy Chain	S51	Main Chain	Hydrogen bond
E503	Side Chain	Heavy Chain	Y54	Side Chain	Hydrogen bond
E503	Side Chain	Heavy Chain	Y79	Side Chain	Hydrogen bond

Supplementary Table 5: Summary of the properties of the monoclonal antibodies

	k_{on} ($M^{-1} s^{-1}$)	k_{off} (s^{-1})	K_D (pM)	Inhibitory in protein- based binding assay	Inhibitory in transgenic <i>P. knowlesi</i> assay	Inhibitory against homologous <i>P. vivax</i> isolates	% of <i>P.</i> <i>vivax</i> strains where invasion inhibited	Binds to PvDBPII subdomain 3
DB1	1.88×10^7	6.46×10^{-3}	344	no	potent	potent	50	no
DB2	9.02×10^5	6.05×10^{-4}	671	yes	low	low	0	yes
DB3	2.98×10^6	4.27×10^{-4}	143	no	intermediate	intermediate	0	yes
DB4	1.30×10^6	4.39×10^{-4}	337	no	low	low	0	yes
DB5	2.33×10^6	4.33×10^{-4}	186	yes	intermediate	n/a	66	no
DB6	8.65×10^6	2.53×10^{-4}	29.2	yes	intermediate	n/a	33	yes
DB7	1.72×10^7	1.13×10^{-4}	6.55	yes	intermediate	intermediate	50	no
DB8	1.73×10^6	3.92×10^{-4}	227	no	low	low	0	yes
DB9	1.79×10^7	3.57×10^{-4}	20.0	yes	potent	potent	91	yes
DB10	1.06×10^7	5.05×10^{-4}	47.8	no	potent	potent	40	no

Supplementary Table 6: Primers.

Primer number	Primer sequence (5' to 3')
1	ACAGGTGCCCACTCCCAGGTGCAG
2	AAGGTGTCCAGTGTGARGTGCAG
3	CCCAGATGGGTCCTGTCCCAGGTGCAG
4	CAAGGAGTCTGTTCCGAGGTGCAG
5	GGAAGGTGTGCACGCCGCTGGTC
6	ATGAGGSTCCCYGCTCAGCTGCTGG
7	CTCTTCCTCCTGCTACTCTGGCTCCCAG
8	ATTTCTCTGTTGCTCTGGATCTCTG
9	GTTTCTCGTAGTCTGCTTTGCTCA
10	GGTCCTGGGCCAGTCTGTGCTG
11	GGTCCTGGGCCAGTCTGCCCTG
12	GCTCTGTGACCTCCTATGAGCTG
13	GGTCTCTCTCSCAGCYTGTGCTG
14	GTTCTTGGGCCAATTTTATGCTG
15	GGTCCAATTCYAGGCTGTGGTG
16	GAGTGGATTCTCAGACTGTGGTG
17	CACCAGTGTGGCCTTGTTGGCTTG
18	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTGCAGCTGGTGCAG
19	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCTGAGGTGCAGCTGGTGGAG
20	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTGCAGCTGCAGGAG
21	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCTGAGGTGCAGCTGTTGGAG
22	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTGCAGCTACAGCAGTG
23	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTTCAGCTGGTGCAG
24	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTCCAGCTGGTACAG
25	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCTGAAGTGCAGCTGGTGGAG
26	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTACAGCTGCAGCAG
27	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCCGAGGTGCAGCTGCAGGAG
28	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCTCAGGTGCAGCTGGTGGAG
29	GATGGGCCCTTGGTCGACGCTGAGGAGACGGTGACCAG
30	GATGGGCCCTTGGTCGACGCTGAAGAGACGGTGACCATTG
31	GATGGGCCCTTGGTCGACGCTGAGGAGACGGTGACCGTG
32	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCTGACATCCAGATGACCCAGTC
33	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTCTGACATCCAGTTGACCCAGTCT
34	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTGTGCCATCCGGATGACCCAGTC
35	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATGGGGATATTGTGATGACCCAGAC
36	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATGGGGATATTGTGATGACTCAGTC
37	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTGAGAAATTGTGTTGACACAGTC
38	CTTTTTCTAGTAGCAACTGCAACCGGTGTACATTGAGAAATAGTGATGACGCAGTC

39	CTTTTCTAGTAGCAACTGCAACCGGTGTACATTCAGAAATTGTGTTGACGCAGTCT
40	CTTTTCTAGTAGCAACTGCAACCGGTGTACATTCGGACATCGTGATGACCCAGTC
41	ATGGTGCAGCCACCGTACGTTTGATYTCCACCTTGGTC
42	ATGGTGCAGCCACCGTACGTTTGATATCCACTTTGGTC
43	ATGGTGCAGCCACCGTACGTTTAATCTCCAGTCGTGTC
44	ATGGTGCAGCCACCGTACGTCTGATTTCCACCTTGGTC
45	CTTTTCTAGTAGCAACTGCAACCGGTTCTGGGCCCAGTCTGTGCTGACKCAG
46	CTTTTCTAGTAGCAACTGCAACCGGTTCTGGGCCCAGTCTGCCCTGACTCAG
47	CTTTTCTAGTAGCAACTGCAACCGGTTCTGTGACCTCCTATGAGCTGACWCAG
48	CTTTTCTAGTAGCAACTGCAACCGGTTCTCTCSCAGCYTGCTGACTCA
49	CTTTTCTAGTAGCAACTGCAACCGGTTCTGGGCCAATTTATGCTGACTCAG
50	CTTTTCTAGTAGCAACTGCAACCGGTTCCAATTCYCAGRCTGTGGTGACYCAG
51	GGCTTGAAGCTCCTCACTCGAGGGYGGGAACAGAGTG