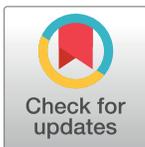


CORRECTION

Correction: CD4 is expressed on a heterogeneous subset of hematopoietic progenitors, which persistently harbor CXCR4 and CCR5-tropic HIV proviral genomes in vivo

The *PLOS Pathogens* Staff

There are two errors in [Table 3](#). The purity values (98, 98) should be listed below donor ID 409000. In addition, in the first column of the 409000 donor ID row, the third line should read 1*. The publisher apologizes for the errors.



 OPEN ACCESS

Citation: The *PLOS Pathogens* Staff (2017) Correction: CD4 is expressed on a heterogeneous subset of hematopoietic progenitors, which persistently harbor CXCR4 and CCR5-tropic HIV proviral genomes in vivo. *PLoS Pathog* 13(9): e1006617. <https://doi.org/10.1371/journal.ppat.1006617>

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Table 3. Analysis of *env* amplicons isolated from HIV+ donors.

Donor ID (% purity) (S1, S2)	HSPC Sort 1 (S1) <i>env</i> amplicons				HSPC Sort 2 (S2) <i>env</i> amplicons				PBMC <i>env</i> amplicons		
	#	FPR	Geno ^b	Pheno ^c	#	FPR	Geno	Pheno	#	FPR	Geno
409000 (98, 98)	1**** _w	61	R5		ND				17	38–61	R5
	1**** _w	0.7	X4/Dual						11	0.7	X4/Dual
	1*	0.7	X4/Dual								
413402 (94, 91)	1*	1.7	X4/Dual	Dual	ND				6	1.7	X4/Dual
414000 (95, 85)	1*	57	R5	R5	ND				5	57	R5
415000 (95, NA)	1**	71	R5	R5	NA				20	43–83	R5
419000 (95, NA)	1***	89	R5		NA				21	22–99	R5
420000 (99, 92)	1****	19	R5	R5	ND				11	14–73	R5
	1**** _w	73	R5						7	0.5–1.7	X4/Dual
	1****	0.7	X4/Dual	Dual							
	1**** _w	8.1	X4/Dual								
421000 (99, 96)	1**** _w	7.8	X4/Dual		1** _w	7.8	X4/Dual		4	24–94	R5
	2***	7.4,7.8	X4/Dual	X4					5	4.7–9.6	X4/Dual
426000 (97, 92)	ND				4****	29–60	R5		10	20–38	R5
					2**** _w	20–31	R5		1	3.2	X4/Dual
					1**** _w	4.8	X4/Dual				
428408 (95, 86) (91, 91)	2****	83,84	R5						22	30–100	R5
	1****	1.3	X4/Dual		1** _w	1.3	X4/Dual		16	0.7–6.8	X4/Dual
431000 (93, 89)	2****	75	R5	R5	ND				8	38–90	R5
	1****	79	R5	R5							
432000 (98, 99)	1****	17	R5						1	49	R5
									8	3.4–8.5	X4/Dual
	1****	2.8	X4/Dual	X4	1****	2.8	X4/Dual	NF	11	74	R5
					1****	3.4	X4/Dual		4	2.8–4.7	X4/Dual
									4	3.4–8.5	X4/Dual
434423 (92, 89) (94, 84)	1**	38	R5		1**	46	R5		18	38–82	R5
435412406 (99, 99) (98, 83) (95, 94)	1**	83	R5	R5							
	2****	87	R5	R5							
	1**	89	R5	R5	1**	83	R5	R5	103	42–99	R5
	1** _w	45	R5								
436000 (93, 85)	ND				2**	31	R5	R5	8	31–85	R5
437000 (94, 92)	1**	55	R5				ND		8	12–55	R5
449000 (90, 95)	1*	100	R5		1**	81	R5		5	59–99	R5
453000 (96, 83)	ND				1***	74	R5	R5	6	41–86	R5
454304 (92,93)	ND				1*	31	R5		24	11–97	R5
					1* _w	9.6	X4/Dual		1	6.8	X4/Dual

(Continued)

Table 3. (Continued)

Donor ID (% purity) (S1, S2)	HSPC Sort 1 (S1) <i>env</i> amplicons				HSPC Sort 2 (S2) <i>env</i> amplicons				PBMC <i>env</i> amplicons		
	#	FPR	Geno ^b	Pheno ^c	#	FPR	Geno	Pheno	#	FPR	Geno
456000 (93,90)	ND				3***	13–39	R5		10 2	10.5–39 7.4, 9.2	R5 X4/Dual

indicates the number of amplicons of that type isolated from each donor.

^bgenotypic prediction of co-receptor usage by Geno2pheno using a false positive rate (FPR) cutoff of $\geq 10\%$ [36, 37].

^cphenotypic analysis of co-receptor usage (see Table 4).

“w” indicates amplicons from whole genome PCR and black text indicates amplicons from multiplex PCR reaction.

Asterisks reflect the likelihood that CD3+ T cells contamination was responsible for the amplicon

* $p < 0.05$ by (1) only

** $p < .05$ by (1) and (2)

*** $p < .01$ by (1) and (2)

**** $p < .001$ by (1) and (2) where (2) is using the more conservative assessment (Fig 2) [10].

Abbreviations: HSPC, hematopoietic stem and progenitor cells; PBMC, peripheral blood mononuclear cells; NA, not analyzed due to purity concerns; R5, CCR5; X4, CXCR4; ND, not detectable; NF, non-functional.

Each set of three non-zero numbers in the donor name represents an independent donation.

<https://doi.org/10.1371/journal.ppat.1006617.t001>

Reference

1. Sebastian NT, Zaikos TD, Terry V, Taschuk F, McNamara LA, Onafuwa-Nuga A, et al. (2017) CD4 is expressed on a heterogeneous subset of hematopoietic progenitors, which persistently harbor CXCR4 and CCR5-tropic HIV proviral genomes in vivo. *PLoS Pathog.* 13(7): e1006509. doi: [10.1371/journal.ppat.1006509](https://doi.org/10.1371/journal.ppat.1006509)