

Zika Virus Persistence and Higher Viral Loads in Cutaneous Capillaries Than in Venous Blood

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We collected venous and capillary serum samples from 21 Zika virus–infected patients on multiple days after symptom onset and found RNA load was higher and median duration of virus detection significantly longer in capillary than in venous blood. These findings raise questions about the role of the capillary compartment in virus transmission dynamics.

Zika virus, belonging to the family *Flaviviridae* and genus *Flavivirus*, is transmitted to humans by mosquito bites but can also be contracted through sexual and vertical transmission (1). Zika virus was first detected in Brazil in 2015 and has since spread with the same speed as chikungunya virus through South and Central America and the Caribbean Islands, despite a shorter and lower viremia in humans and a longer replication period in the vector (2,3). As with other arboviruses, Zika virus viremia is commonly measured in venous blood, even though mosquitoes introduce virus into cutaneous capillary blood. We conducted a prospective descriptive study with Zika virus patients during the Zika virus epidemic in French Guiana to evaluate the kinetics of Zika virus RNA load in serum samples collected sequentially from venous and skin capillary blood.

The study population comprised 21 symptomatic and consenting Zika virus patients infected during March–September 2016. We confirmed Zika virus infection by real-time reverse transcription PCR (RT-PCR) of serum and urine samples provided by patients during the first few days after symptom onset. We obtained serum samples from the venous and cutaneous capillary blood (collected from the fingertip) sequentially 1–18 days after the onset of symptoms. The median age of the population was 40 (range 28–63) years and the sex ratio (male:female) was 1.6. We observed no co-morbidities, and all participants were found

to be free of dengue and chikungunya virus infections by methods previously described (4,5).

We extracted RNA from 150- μ L samples using the QIAamp Viral RNA kit (QIAGEN, Hilden, Germany) and performed Zika virus RNA amplification using the RealStar Zika Virus RT-PCR Kit 1.0 (Altona Diagnostics GmbH, Hamburg, Germany) according to the manufacturer's instructions. We performed Zika virus RNA quantification using a reference strain provided by the European Zika Virus Archive (SKU no. 001N-01648) and estimated the Zika virus RNA load as \log_{10} copies per milliliter (online Technical Appendix Table, <https://wwwnc.cdc.gov/EID/article/23/11/17-0337-Techapp1.pdf>).

Zika virus RNA loads in capillary blood correlated with those in venous blood (Spearman correlation test, $r = 0.54$, $p < 0.0001$) but were significantly higher in the capillary samples (Wilcoxon signed rank test, $p = 0.0003$), except for 3 patients (nos. 2, 13, and 21; online Technical Appendix Table). The median duration of Zika virus detection after symptom onset was significantly greater in capillary blood than in venous blood ($p = 0.005$ by log rank test; hazard ratio = 2.99, 95% CI 1.39–6.43), even though the duration of detection in capillary blood was underestimated; RNA was still detectable in the last capillary blood samples taken from 8 patients (nos. 1, 5, 6, 8, 11, 15, 16, and 19; online Technical Appendix Table). The duration of Zika virus RNA detection was greater in capillary than in venous blood for 12 (57%, 95% CI 34%–78%) of the 21 patients (nos. 1, 3, 5, 6, 7, 8, 11, 14, 15, 16, 19, and 20). The maximum duration of RNA detection in capillary blood samples was 18 days after the onset of symptoms (seen with patient no. 19) with a load of 1.9 \log_{10} copies/mL. The duration of detection was equal between the 2 compartments for 7 (33%, 95% CI 15%–57%) of 21 patients (nos. 4, 9, 10, 12, 13, 17, and 18) and longer in venous blood for 2 (10%, 95% CI 1%–30%) of 21 patients (nos. 2 and 21) (online Technical Appendix Table).

These data raise questions about the consequences of longer persistence and higher loads of Zika virus RNA in the cutaneous capillary blood compartment, although we did not test Zika virus replication capacity. The higher load and longer detection of Zika virus RNA in this compartment might be attributable to Zika virus replication in permissive cells of the skin (e.g., human dermal fibroblasts and epidermal keratinocytes); capillaries; or both (6). If the Zika virus RNA observed in the serum samples taken from the capillary compartment reflects the presence of infectious virus particles, symptomatic Zika virus–infected patients would need to be shielded from mosquitoes for a longer period than is currently practiced to limit potential vectorborne transmission.

This study comparing the kinetics of Zika virus RNA load between the venous and capillary compartments

highlights the need to further investigate the infectivity and pathophysiology of the virus located in the often neglected capillary compartment. These findings provide new information on this biologic compartment, which plays a key role in vectorborne transmission and transmission dynamics. Moreover, these observations, if validated with more patients and extended to other vectorborne infections, will be vital for preventing and controlling the transmission of Zika virus and other arboviruses.

Institutional review board approval was granted by the Comité de Protection des Personnes Sud-Méditerranée I corresponding to the following study “Etude descriptive prospective de la maladie à virus Zika au sein de la communauté de défense des Forces Armées en Guyane” and was registered February 2016 under the number RCB: 2016-A00394-47. Written informed consent was obtained from each patient as required by the Comité de Protection des Personnes Sud-Méditerranée I.

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Dr. Matheus is a research assistant at the Institute Pasteur de la Guyane, French Guiana, with research interests in the diagnosis and pathophysiology of arboviruses. She is currently studying viral emergence, particularly that of a hantavirus in French Guiana.

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Detection of Spotted Fever Group *Rickettsia* DNA by Deep Sequencing

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After conventional molecular and serologic testing failed to diagnose the cause of illness, deep sequencing identified spotted fever group *Rickettsia* DNA in a patient’s blood sample. Sequences belonged to *R. honei*, the causative agent of Flinders Island spotted fever. Next-generation sequencing is proving to be a useful tool for clinical diagnostics.

When conventional laboratory tests cannot identify an etiologic agent, unbiased deep sequencing performed directly on a clinical sample has the potential to identify a probable cause of disease. We used deep sequencing to detect spotted fever group (SFG) *Rickettsia* DNA in the blood of a patient for whom diagnosis was not possible through conventional molecular and serologic testing.

In late 2016, a middle-aged woman was admitted to a regional hospital in Queensland, Australia, after 2 weeks of mild cough, myalgia, fever, and lethargy. The day before admission, she experienced a blanching rash and pains in her feet, after which her condition deteriorated and a definite petechial rash appeared. Chest radiographs showed atelectasis on 1 side. Meningococcal septicemia was suspected, and the patient was transferred to intensive care with septic shock. Despite treatment with inotropes and several antimicrobial drugs (including ceftriaxone, vancomycin, meropenem, doxycycline), the patient died the next morning.

Clinical testing did not identify an infectious disease agent in the patient’s blood; serologic test results

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Technical Appendix

Technical Appendix Table. Kinetics of Zika virus load in venous and capillary blood samples from 21 patients, French Guiana, 2016*

Patient no.	Age, y	Sex	Day after symptom onset	Cycle threshold		Zika virus load, log ₁₀ copies/mL	
				Venous blood	Urine	Venous blood	Capillary blood
1	40	M	3	33.26	32.77	3.8	NA
			8			<1.8	2.8†
2	45	M	3	33.26	32.77	2.8	<1.8†
			6			2.3	<1.8†
			7			<1.8	<1.8
3	59	F	2	36.75	34.02	2.6	3.9†
			3			<1.8	3.1†
			5			<1.8	<1.8
			10			<1.8	<1.8
4	34	M	2	38.01	28.41	2.2	3.7†
			4			<1.8	<1.8
			11			<1.8	<1.8
5	37	M	1	30.27	34.08	4.3	4.2
			3			<1.8	2.4
			7			<1.8	2.6
6	28	F	4	31.00	NA	4.6	NA
			5			4.1	4.2
			7			<1.8	2.8
			8			<1.8	<1.8†
7	42	M	2	34.29	36.88	3.6	4.8†
			3			<1.8	3.5
			5			<1.8	2.7
			8			<1.8	<1.8†
			10			<1.8	<1.8
8	41	F	3	Negative	29.00	<1.8	3.7
			7	<1.8	<1.8†		
9	34	F	2	36.92	35.00	2.9	3.1
			5			<1.8	<1.8†
			10			<1.8	<1.8
10	41	M	3	35.35	33.63	3.3	3.4†
			5			<1.8	<1.8
			8			<1.8	<1.8
			10			<1.8	<1.8
11	35	M	2	36.64	30.17	2.6	2.9
			3			<1.8	2.6
			5			<1.8	2.5
			8			<1.8	2.4
			10			<1.8	<1.8†
12	39	M	3	34.16	32.9	2.7	4.0†
			6			<1.8	<1.8†
			10			<1.8	<1.8†
13	39	M	3	36.97	27.63	2.5	2.4
			5			<1.8	<1.8
			10			<1.8	<1.8†
14	45	M	2	33.48	36.45	3.8	NA
			3			<1.8	3.3†
			6			<1.8	<1.8†
			8			<1.8	<1.8†
15	46	M	1	33.96	29.27	3.9	3.9†
			3			3.1	3.5
			5			NA	3.8†
			7			2.3	4.2
16	63	F	3	31.37	34.44	3.9	3.4
			8			<1.8	<1.8
			10			<1.8	2.5

Patient no.	Age, y	Sex	Day after symptom onset	Cycle threshold		Zika virus load, log ₁₀ copies/mL	
				Venous blood	Urine	Venous blood	Capillary blood
17	44	M	2	37.38	34.17	2.5	5.7†
			3			<1.8	<1.8†
			5			<1.8	<1.8†
			7			<1.8	<1.8†
18	32	F	1	30.92	34.70	4.3	NA
			3			3.0	3.1†
			7			<1.8	<1.8†
			15			<1.8	<1.8
19	35	M	4	Negative	36.56	<1.8	3.8†
			6			<1.8	3.2†
			8			<1.8	3.7
			18			NA	1.9
20	41	F	2	31.99	37.00	4.1	4.4†
			4			<1.8	3.7†
			7			<1.8	3.6
			14			<1.8	<1.8
21	35	F	1	34.91	NA	3.4	<1.8†
			3			<1.8	<1.8†
			6			<1.8	<1.8
			13			<1.8	<1.8†

*NA, not available; RT-PCR, reverse transcription PCR. Quantification was performed by real-time RT-PCR (RealStar Zika Virus RT-PCR Kit 1-0 CE; Altona Diagnostics GmbH, Hamburg, Germany) with a detection threshold of 1.8 log₁₀ copies/mL.
†Volume of sample used for extraction was <50 µL.