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Brief report

First case of imported Zika virus infection in Spain



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ABSTRACT

We report a case of Zika virus (ZIKV) infection in a patient with diarrhea, fever, synovitis, non-purulent conjunctivitis, and with discreet retro-orbital pain, after returning from Colombia in January 2016. The patient referred several mosquito bites. Presence of ZIKV was detected by PCR (polymerase chain reaction) in plasma. Rapid microbiological diagnosis of ZIKV infection is needed in European countries with circulation of its vector, in order to avoid autochthonous circulation. The recent association of ZIKV infection with abortion and microcephaly, and a Guillain-Barré syndrome highlights the need for laboratory differentiation of ZIKV from other virus infection. Women with potential risk for Zika virus infection who are pregnant or planning to become pregnant must mention that fact during prenatal visits in order to be evaluated and properly monitored.

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Primer caso importado de infección por virus Zika en España

RESUMEN

Palabras clave:

Virus Zika

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Enfermedades transmitidas por vectores

Presentamos un caso de infección por virus Zika (ZIKV) en un paciente con diarrea, fiebre, sinovitis, conjuntivitis no purulenta, con discreto dolor retroorbital, después de regresar de Colombia en enero de 2016. El paciente refería múltiples picaduras de mosquito. La presencia de ZIKV fue detectada en plasma por reacción en cadena de la polimerasa (PCR). En los países europeos en los que existe circulación de su vector es necesario un diagnóstico microbiológico rápido de la infección por ZIKV para evitar transmisión autóctona. La asociación reciente de la infección por ZIKV con abortos y microcefalia y síndrome de Guillén-Barré pone de relieve la necesidad de la diferenciación de ZIKV de otras infecciones por virus. Las mujeres con riesgo potencial de infección por el virus Zika que están embarazadas o que planean quedar embarazadas deben mencionar esa circunstancia durante las visitas prenatales con el fin de ser evaluadas y monitorizadas adecuadamente.

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Case report

A 49 years-old man is being followed at our Hospital in Spain for a clinical history of irritable bowel syndrome, hypertension and

dyslipidemia and he is under treatment telmisartan and atorvastatin.

The patient traveled to Colombia where he stayed between December, 2015 and January, 2016. The day of his return to Spain, he began with diarrhea (up to 8 watery stools without blood, mucus or pus) and temperature of 37.3 °C. On January the patient was attended at our Hospital for signs of swelling in the right hand, arthralgia in fingers, right knee, ankle and elbow, and conjunctival

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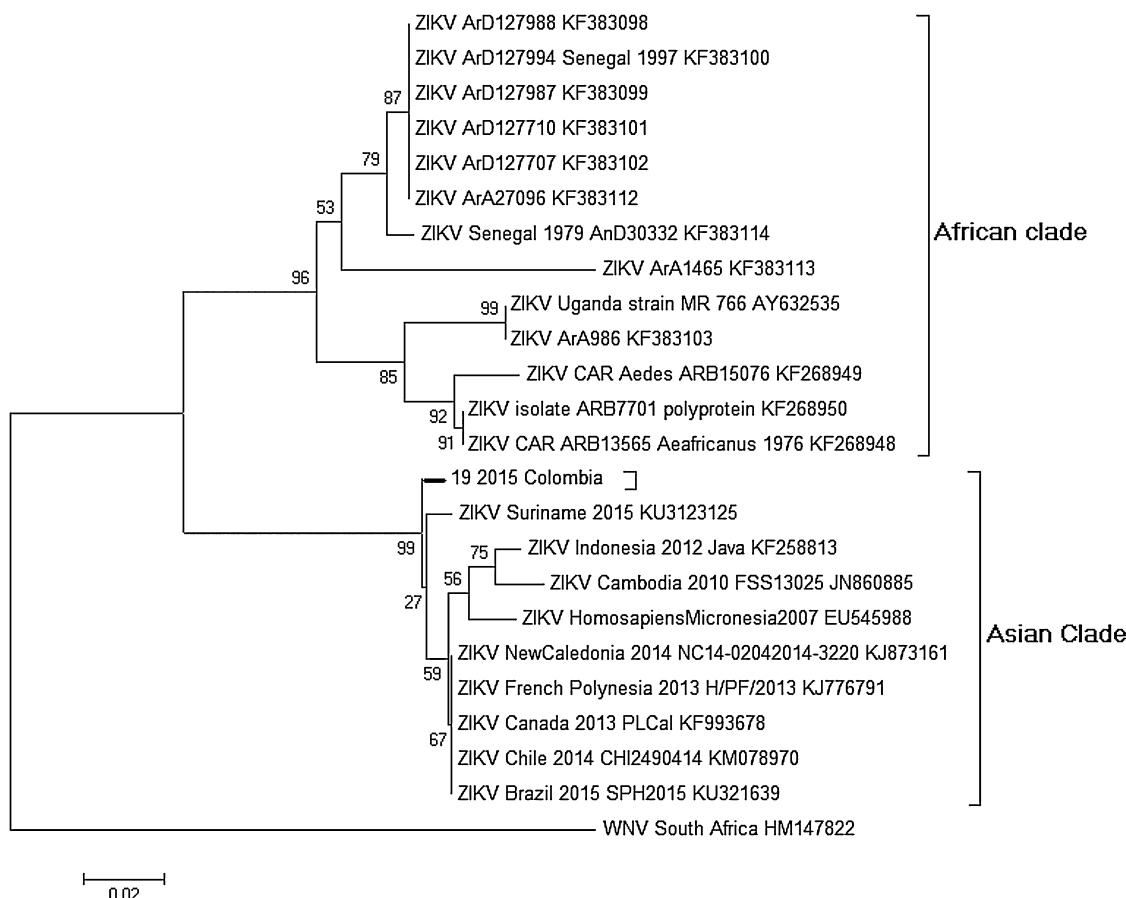


Fig. 1. Phylogenetic analysis from partial NS5 gene of ZIKV. Neighbor-joining phylogenetic tree constructed by Mega 5 software using p-distance Method with 1000 bootstrap replicates. The tree was rooted with West Nile virus. The sequence obtained in this study is marked with the branch bold.

injection with mild retro-orbital pain. He did not report headache, rash, myalgias, or urinary symptoms. The patient referred mosquito bites in Colombia.

Cutaneous lesions were not observed, but there was evidence of synovitis at the proximal interphalangeal joint of the first finger of his right hand with no other significant exploratory signs.

Having into consideration the epidemiological history, diagnosis of infection by Dengue (DENV), Chikungunya (CHIKV) and Zika (ZIKV) viruses was requested. A 198 bp amplicon was obtained from a sample of plasma taken 3 days after onset by using a RT-PCR specific for ZIKV (modified from Balm MN et al.).¹ The product was sequenced and ZIKV was identified after BLAST analysis. A phylogenetic analysis with this fragment was carried out using the Mega 5 software, and the phylogenetic tree show that the sequence of ZIKV detected in the patient clusters within the Asian clade (Fig. 1). The sample was also used to detect DENV and CHIKV by PCR with negative results. A serum sample obtained the same day was analyzed by IFA against Zika virus (Arboviral Fever Mosaic IgG and IgM, Euroimmun, Germany). Positive and undetermined results for IgG and IgM, respectively, were obtained. An urine sample taken 14 days after the onset was also positive by PCR. It is remarkable that urine samples can be used for direct diagnostic and that virus can be detected for a longer period than in blood or serum samples.² A case is confirmed when a positive result by PCR is obtained. This patient was confirmed by positive results in two different samples. The results of IFA are compatible with an acute Zika infection still without a clear IgM response in a patient with a previous exposition to another flavivirus such as DENV.

Table 1
Comparison of symptoms and signs between Dengue, Chikungunya and Zika.

Symptoms	Dengue	Chikungunya	Zika
Fever	++++	+++	++/++
Arthralgia/myalgia	+++	++++	++
Edema of extremities	0	0	++
Rash	++	++	+++
Retroocular pain	++	+	++
Conjunctivitis	0	+	+++
Leukopenia/thrombocytopenia	+++	+++	0
Hemorrhage	+	0	0
Lymphadenopathy	++	++	+
Hepatomegaly	0	+++	0

Other laboratory tests were unremarkable: leukocytes 5500 cells/ μ L (neutrophils 67%, lymphocytes 20%), hemoglobin 13.5 g/dL, platelets 208,000 cells/ μ L, ESR 12 mm/h; urinalysis with traces of proteinuria; AST 29 U/L, ALT 31 U/L. Research of fecal parasites and stool showed no significant findings.

Background

ZIKV is an arbo-virus (*Flavivirus* genus, *Flaviviridae* family). It was first identified in 1947 in the Zika forest (Uganda) from the blood of a Rhesus monkey. In 1948 it was isolated from a pool of *Ae. africanus* mosquitoes in the same forest.³ Thereafter, serological and entomological studies indicated human and animal ZIKV infections in the African continent. In 2007 an epidemic developed in Malaysia and Micronesia, including Yap Islands.⁴

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