Temporal–Spatial Distribution of *Vibrio cholerae* in Cuba: July 1997–December 2019

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**ABSTRACT**

**INTRODUCTION** *Vibrio cholerae* is a microorganism that causes acute diarrheal diseases and cholera, one of the leading causes of global morbidity and mortality, especially in children under five years old. It is present in many regions and has been isolated from diverse sources such as water, soil and food. Surveillance of this microorganism in Cuba from 1985 through June 1997 showed circulation of non-epidemic non-O1/non-O139 serogroups, but surveillance continued to identify distribution of *V. cholerae* serotypes and serogroups in the different geographical regions of the country during the following years, due to the risk of introducing cholera-causing serogroups that provoked cholera epidemics in other countries of the region.

**OBJECTIVE** Describe the temporal–spatial distribution of serogroups and serotypes of *V. cholerae* in Cuba.

**METHODS** A cross-sectional study was conducted that included isolates from passive surveillance of *V. cholerae* in 16 hygiene and epidemiology centers throughout Cuba from July 1997 through December 2019, submitted to the National Reference Laboratory for Acute Diarrheal Diseases of the Pedro Kourí Tropical Medicine Institute in Havana, Cuba. The timeline was subdivided into three five-year periods and one eight-year period. The centers submitting isolates were grouped into three geographical regions: western, central and eastern Cuba. A total of 1060 *V. cholerae* isolates were studied, from the 1438 samples sent from 15 Provincial Hygiene, Epidemiology and Microbiology Centers and the Municipal Hygiene, Epidemiology and Microbiology Center of the Isle of Youth Special Municipality. Genus, species and serotype of all specimens were studied and reviewed in the context of the outbreaks of acute diarrheal diseases reported in the country.

**RESULTS** All 1060 isolates were confirmed as *V. cholerae*. In the distribution by time period and region, the highest percentage occurred in the 2012–2019 period, and the eastern region contributed the most isolates in all periods. Approximately 63.9% (677/1060) were from outbreaks, and in the 2012–2019 period, the most epidemic-causing isolates came from the western region. Approximately 52.8% (560/1060) were identified as non-O1/non-O139 *V. cholerae*, and 47.2% (500/1060) as O1 *V. cholerae*; of these, 96.4% (482/500) corresponded to Ogawa serotype and 3.6% (18/500) to Inaba. Circulation of non-O1/non-O139 *V. cholerae* occurred throughout the entire period. The O1 serogroup began to circulate in 2012 and continued through 2016; however, since 2017, it has not been identified again. In the western region, there were smaller percentages of isolates of non-O1/non-O139 *V. cholerae* in all periods, except 2012–2019. In that period, *V. cholerae* O1 was identified to a lesser degree in the central region.

**CONCLUSIONS** *Vibrio cholerae* circulated in all three Cuban regions during the years studied, with a higher percentage of isolates of the non-O1/non-O139 serogroup, which caused outbreaks or sporadic cases of diarrhea in the eastern region, with the exception of the 2012–2019 period, when epidemic outbreaks of the O1 serogroup (which causes cholera) occurred in all three regions, with higher percentages in the western region.

**KEYWORDS** *Vibrio cholerae*; *Vibrio cholerae* O1; *Vibrio cholerae* non-O1; *Vibrio cholerae* O139; dysentery; cholera; epidemiological monitoring; infectious diarrheal disease; disease transmission, infectious; gastrointestinal diseases; Cuba

**INTRODUCTION**

Acute diarrheal diseases—including cholera—are one of the main causes of global morbidity and mortality, especially in children under five years old.[1] Causal factors include various bacteria; among these, *Vibrio cholerae* is one of the most important.[2,3]

Morphologically, the microorganism has a curved structure with a polar flagellum.[4] Based on the O antigen of the capsular lipo-polysaccharide, *V. cholerae* is classified in more than 200 biochemically identical serogroups, including O1 and O139, the two epidemic serogroups that cause cholera.[5] Non-O1/non-O139 serogroups are not considered epidemic, although they can cause outbreaks or sporadic cases of diarrhea with clinical symptoms and signs different from cholera.[3]

*V. cholerae* is present in several regions of the world and can be isolated from diverse sources such as water, soil and food. Humans are an accidental and transitory host, yet they are the ones who disseminate it in the environment.[6,7] Intestinal infections are the most clinically relevant, particularly cholera.[8] But extra-intestinal infections can occur as well, such as fasciitis, bacteremia, meningitis, otitis and those that can develop in surgical wounds.[9]

Since 1991, in the region of the Americas, various cholera outbreaks have been reported[10] as well as sporadic cases of gastroenteritis...
After incubating for 24 hours, three or more colonies were selected, based on their characteristics in the corresponding media: i) convex, with regular borders and sucrose fermenters in TCBS agar; ii) translucent, convex, with regular borders in MacConkey agar; and iii) hemolytic or not, convex, with regular borders in blood agar.

All colonies were inoculated by puncture and streaked in the primary differentiated media (Kligler’s iron agar with double sugar fermentation and lysine iron agar). Both were aerobically incubated from 18–24 hours at 37 °C.[18]

Once incubation was complete, cultures that exhibited the following characteristics were selected: a) they did not oxidize or ferment lactose, they did oxidize and ferment glucose, they did not produce gas or hydrogen sulfide in Kligler’s iron agar iron with double sugar fermentation; and b) they decarboxylated L-lysine in iron and lysine agar.[18]

In all isolates, presence of cytochrome oxidase enzyme was confirmed, in accordance with the Kovacs method. Those that were oxidase positive were submitted to a complementary physiological study for confirmation in the Vibrio genus and V. cholerae species.[18]

To identify the genus, the Möeller method was utilized for biochemical tests measuring use of amino acids and carbohydrates. [18] To identify the species, isolates were submitted to tolerance tests for sodium chloride (NaCl) and use of sucrose. Serological agglutination tests were performed on slides using V. cholerae O1 and V. cholerae O139 polyvalent antiserum to identify these serogroups.[18]

Analysis Descriptive statistics were used, such as frequency and percentages, to analyze and present results.

RESULTS
The 1060 isolates were confirmed as V. cholerae as they were Gram-negative bacilli, facultative anaerobic, positive oxidase, with positive reaction to lysine and ornithine decarboxylase tests and negative for arginine dihydrolase; they grew in tryptone soy broth with 6% NaCl concentration and used sucrose and mannitol, not inositol.

The highest percentage of V. cholerae isolates occurred in the 2012–2019 period and in the eastern region in all periods (Table 1). Approximately 63.9% (677/1060) came from outbreaks. Figure 1 shows the proportion calculated for each period, according to geographic region. In the 1997–2001 period, the lowest percentage of isolates coming from outbreaks was reported in the western region (5.0%; 34/677), but in the 2012–2019 period, that percentage of isolates coming from outbreaks was reported in the central region (95.0%; 643/677). In the 2012–2019 period and in the eastern region in all periods (Table 1) or bacteremia caused by non-O1/non-O139 V. cholerae.[8,11,12] In 2010, acute diarrheal diseases caused by V. cholerae spread rapidly in Haiti,[13] a new scenario that could lead to reemergence and dissemination of cholera in countries such as the Dominican Republic, Cuba and Mexico, where positive cases were reported.[14,15] Considering circulation of this pathogen in the region, the National Reference Laboratory for Acute Diarrheal Diseases of the Pedro Kouri Tropical Medicine Institute (IPK) established in 1995 a surveillance system to identify outbreaks and sporadic cases of diarrheal illnesses that could be caused by V. cholerae.[16] This surveillance documented evidence of the circulation of non-O1/non-O139 strains from 1985 through June 1997, but surveillance of sporadic outbreaks of diarrheal diseases and epidemiological events possibly attributable to V. cholerae has continued, in order to provide relevant information to health authorities. The purpose of this study was to describe the temporal–spatial distribution of V. cholerae serogroups and serotypes in Cuba from July 1997 through December 2019.

METHODS
Design and samples A cross-sectional study was conducted from July 1997 through December 2019. It was subdivided into four periods: P1: 1997–2001, P2: 2002–2006, P3: 2007–2011 and P4: 2012–2019 (three five-year periods and one eight-year period, the latter in order to include the most recent results).

Of the 1483 V. cholerae isolates from passive surveillance in the culture collection of the IPK’s National Reference Laboratory for Acute Diarrheal Diseases, 1060 viable isolates (microorganisms with the possibility of multiplying) were studied; 423 resulted non-viable or were contaminated. The isolates were originally collected in 15 of Cuba’s Provincial Hygiene, Epidemiology and Microbiology Centers (CPHEM) and the Municipal Hygiene, Epidemiology and Microbiology Center (CMHEM) of the Isle of Youth Special Municipality.

The centers submitting isolates were grouped into three regions: western, central and eastern Cuba. The western region included CPHEMs of Pinar del Rio, Artemisa, Mayabeque, Havana, and Matanzas provinces, as well as the CMHEM of the Isle of Youth Special Municipality; the central region included CPHEMs of Cienfuegos, Ciego de Ávila, Villa Clara, Santís Spiritus and Camagüey provinces; and the eastern region included CPHEMs of Las Tunas, Holguín, Granma, Santiago de Cuba and Guantánamo provinces.

Isolates were classified according to genus, species, serogroup and serotype, and those related to outbreaks were identified. Cases were considered to be an outbreak when the unit submitting the V. cholerae isolates reported that they belonged to a single event in which two or more patients had been in contact with one another. The moment of onset of symptoms, the place where the episodes occurred, and the individuals’ characteristics were all recorded.[17]

Procedures Isolates were preserved in Pasteur’s conservation medium for Enterobacteriaceae; they were inoculated in brain-heart infusion broth and anaerobically incubated for 18–24 hours at 37 °C. After incubation, a loopful of the broth culture was seeded by colony depletion plating on plates with selective medium for Vibrio thiosulfate-citrate-bile salts-sucrose (TCBS) agar (Biofite, Italy), MacConkey agar (Biofite, Italy) and blood agar (5% sheep blood) and incubated under recommended conditions.[18]
O1/non-O139 were in continuous circulation all those years. The O1 serogroup began circulating in 2012; the number of O1 isolates increased in 2013 and 2014, decreased in 2015, and since 2017, the O1 serogroup has not been reported at all.

Table 1: Distribution of confirmed isolates of Vibrio cholerae by geographic region. Cuba, July 1997‒December 2019

<table>
<thead>
<tr>
<th></th>
<th>Western</th>
<th>Central</th>
<th>Eastern</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>1997‒2001</td>
<td>20</td>
<td>105</td>
<td>100</td>
<td>225</td>
</tr>
<tr>
<td>2002‒2006</td>
<td>14</td>
<td>80</td>
<td>103</td>
<td>197</td>
</tr>
<tr>
<td>2007‒2011</td>
<td>8</td>
<td>14</td>
<td>36</td>
<td>58</td>
</tr>
<tr>
<td>2012‒2019</td>
<td>187</td>
<td>157</td>
<td>236</td>
<td>580</td>
</tr>
<tr>
<td>Total</td>
<td>229</td>
<td>356</td>
<td>475</td>
<td>1060</td>
</tr>
</tbody>
</table>

Figure 1 presents the distribution (in percentages) of serogroup strains according to geographic region. The western region presented the lowest percentage of non-O1/non-O139 V. cholerae isolates in all periods except 2012‒2019. In that same period, V. cholerae O1 was identified in the central region, but exhibiting the lowest percentages.

**DISCUSSION**

Worldwide, Vibrio spp. are widely distributed in aquatic environments.[18] It has been suggested that as a result of climate change, increases in ocean temperatures may be responsible for outbreaks of Vibrio infections in countries such as Israel, Denmark, Spain, Chile and the United States.[19,20]

According to data published by the Global Burden of Diarrheal Diseases Collaborators (GBDDC), acute diarrheal diseases have persisted in Latin America as a significant public health problem.

[21] Levels during the last three decades have been relatively stable and mortality in several countries has dropped, which Herrera-Benavente attributes to the success of WHO-sponsored control programs (under auspices of PAHO).[1]

In a study prior to implementation of Cuba’s surveillance system, Bravo[16] found that in the three Cuban regions, non-O1/non-O139 V. cholerae circulated from 1985 through 1997 in patients with acute diarrheal diseases, most commonly in the eastern region. The author suggested that an ecological niche in that region might favor persistence of diarrheas caused by this agent.

In June 2012, V. cholerae O1 was isolated in Cuba, when the Ministry of Public Health reported a cholera outbreak in Granma province.[22] Although no events caused by this strain had been observed in previous years, it is not unusual for such events to occur. The cholera-causing agent cannot be eliminated from the environment since it is a native species and is associated with marine habitats such as algae and crustaceans. In addition, changes in climate patterns favor the growth of V. cholerae in aquatic ecosystems and contribute to the occurrence of outbreaks and epidemics.[23,24]

In Cuba, more outbreaks and cases of diarrhea occur in the summer months, in part due to high temperatures and summer rains.[25] However, no studies have been conducted on Cuban climate variability and its effect on the temporal–spatial behavior of V. cholerae.

The epidemic O1 and O139 V. cholerae serogroups produce cholera, particularly in developing countries due to faulty infrastructure affecting basic sanitation and drinking water, as well as poor access to quality medical services.[2,26,27] while non-O1/non-O139 strains are often isolated in the environment and are associated with sporadic cases of gastroenteritis and extraintes-
From 2012 to 2016, there were various outbreaks in Cuba associated with the circulation of *V. cholerae* O1.[22,29] In the present study, an increase in isolates was observed from 2012 to 2019 from outbreaks in the country's western region, which could be related to the joint circulation of the epidemic serotypes that can cause outbreaks of greater magnitude.[27]

In a 2012 study in Cuba by Romero-Placeres to investigate whether shortcomings in the quality of drinking water and sanitation influenced the burden of acute diarrheal diseases, it was found that more municipalities in the west faced high risk of a diarrheal outbreak due to both these problems. In addition, Havana is located in western Cuba, and is the province with the largest population and most dealings (through business, tourism, travel) with the rest of the island and also with other countries.[30] Another important factor is that 96.8% of households in the capital receive water via aqueduct; the supply service is inconsistent and in some places has low pressure, which along with the current poor condition of distribution pipes, can lead to water contamination.[31]

According to the 2012 census by Cuba’s National Statistics Bureau,[31] despite the fact that the country enjoys extensive coverage of water and sanitation services, it still faces deficiencies from deteriorating infrastructure due to aging distribution systems, with high risk of microbiological contamination, a conclusion coinciding with observations by Romero-Placeres.[30]

The 2012 and 2013 cholera outbreaks in Havana were associated with food markets, where several asymptomatic carriers worked and manipulated foods.[25] The O1 serogroup contains three serotypes; the two most common—Ogawa and Inaba—are capable of producing cholera outbreaks. The difference between the two lies in methylation of the capsular lipopolysaccharide by the enzyme methyltransferase, codified by the *webT* gene. When an isolate presents a methylated lipopolysaccharide, it belongs to the Ogawa serotype, and when it does not, it is Inaba.[32]

In endemic areas, the change from one serotype to another is not a random process, rather, it depends on several factors.[33] Genetic studies suggest that the mobile elements present in the genome of *V. cholerae*, such as those related to virulence (CTXØ), antibiotic resistance (SXT ICE) and phages and their resulting clonal expansion, can contribute to a preponderance of a particular serotype during a specific time period. Another important factor in the change of serotype is pressure generated by the immune system, because when a specific serotype circulates in a population over a long time, the host will present immunity to this serotype. The effect of environmental factors on the viability of these two serotypes is not well known.[34]

In stool samples of patients with diarrhea obtained from 2006 through 2016 in sentinel centers in Nepal, *V. cholerae* O1 was isolated, mostly the Ogawa serotype,[35] and in 2015 and 2016 this strain was also isolated in stool samples of patients with acute diarrheal disease and identified in samples of water used for drinking and domestic chores in Ghana, coexisting with non-O1/non-O139 serogroup.[36]

During cholera outbreaks in Cuba, circulating together with *V. cholerae* O1 were the non-epidemic serogroups, as reported in Thailand, Iraq and Japan.[37] The coexistence of toxigenic and nontoxigenic serogroups represents a global health emergency, due to the horizontal transfer of genes that contribute virulence and antimicrobial resistance among serogroups of the same species, which can lead to the appearance of more virulent strains.[38]

The results of the present study on circulation of serogroups and serotypes coincided with the above-cited works and with cocirculation of *V. cholerae* O1 and non-O1/non-O139 from 2012 to 2016, underscoring the importance of epidemiological surveillance of this bacterium in Cuba.

In January 1991, a cholera epidemic broke out on the coast of Peru, attributed to the O1 serogroup, which quickly spread to almost all Latin American countries, although Cuba was not affected.[25] In October 2010, cholera was introduced into Haiti, and from then through 2018, persistent circulation and outbreaks of cholera were reported, associated with poor sanitary conditions and lack of access to safe drinking water in that country.[13] Subsequent outbreaks in Cuba were related to the Haitian epidemic, according to Cuban

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**Figure 3:** Distribution of *Vibrio cholerae* serogroup O1 and *Vibrio cholerae* non-O1/non-O139 responsible for diarrheal diseases (n = 1060) according to geographic region. Cuba, July 1997–December 2019
Our temporal–spatial heterogeneity study confirmed that *V. cholerae* circulated for 23 years in all three regions of Cuba. Additionally, non-O1/non-O139 serogroups that caused outbreaks or sporadic cases of diarrhea were isolated in all time periods and regions, most often in the eastern region, with the exception of the last period, when epidemic outbreaks of the cholera-causing O1 strain occurred in all three regions, most commonly in the western region. This study shows that gastrointestinal infections from *V. cholerae* constitute a health problem in Cuba that needs to be addressed and suggests that the results should be used to design focused strategies, in accordance with the microorganism’s temporal-spatial distribution pattern.

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23. 190, 372. Spanish.


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