Incidence and Molecular Analysis of *Vibrio cholerae* among Some Primitive Tribes in Odisha, India

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**Abstract**

Cholera has been reported in the state of Odisha, India since more than two decades. The present study was carried out during the year 2000 to 2004 to find out the incidence and molecular analysis of *V. cholerae* causing diarrhoea among some primitive tribes namely Bondo, Didayi, Juanga and Kondha residing in Odisha, India. Bacteriological analysis of 1066 rectal swabs collected from hospitalized diarrhoea patients in tribal areas revealed 9.8% were positive for *V. cholerae* O1 Ogawa and 3.8% for *V. cholerae* O139 serogroup. The major antibiogram profile of O1 and O139 revealed resistance to ACoFzNNaS and AFzNSNa respectively. The emergence of fluoroquinolone resistant *V. cholerae* O1 and O139 and 100% nalidixic acid resistant O139 were observed. Cholera toxin production of *V. cholerae* O1 isolated from the tribal areas exhibited similar to the strains prevail in the endemic coastal districts of Odisha. PCR assay revealed that all the isolates of *V. cholerae* O1 and O139 were positive for ctxA, tcpA (El Tor), ace and zot genes. Except some minor variation in RAPD profiles, both O1 and O139 strains exhibited similar pattern of PFGE and RAPD among each serogroup compared with the strains of Kolkata and other parts of the state. For the first time, this study revealed that *V. cholerae* was one of the important aetiopathologic agents responsible for causing diarrhoea in the remote primitive tribal areas in the state, the possible origin and spread of which could be due to the population movement from the endemic areas of the state to the remote tribal areas.

**Keywords**  
Molecular Analysis of *Vibrio cholerae*, Primitive tribes, *V. cholerae* O1 Ogawa, *V. cholerae* O139

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**Introduction**

Epidemic and endemic cholera is a major public health problem and continues to be an important cause of morbidity and mortality in developing world. The causative agent *V. cholerae* O1 and since 1992 *V. cholerae* O139 is capable of pandemic spread (Levine *et al.*, 1993). Seven distinct pandemics of cholera have been recorded since 1817 (Kaper *et al.*, 1995). The eighth pandemic was suspected when a new serotype O139 was emerged as an epidemic strain and spread from Chenai to other parts of the country and Bangladesh (Kaper *et al.*, 1995; Ramamurthy *et al.*, 1993). The causative agent of the sixth and perhaps the fifth pandemic belonged to the classical biotype of *V. cholera* (Kaper *et al.*, 1995). The seventh pandemic was caused by *V. cholera* O1 El Tor biotype. After its emergence in the Island of Sulawesi of Indonesia, it spread to Indian subcontinent and reached other countries including Latin America through different routes. Cholera due
to *V. cholera* O1 El Tor reemerged in Americas in 1991 after 100 years absence in the form of explosive outbreak in Peru and spread rapidly to other South American countries at the rate of one country per month approximately (Tauxe and Blake, 1992). Between 1991 and 1994 more than one million cases affected by cholera were reported in Latin American countries (Koo *et al.*, 1999). The explosive outbreak among Rwandan refugees in Zaire in July 1994 which affected 70,000 people and killed 12000 is another example of its potentiality for rapid spread (Goma, 1994; Sidique *et al.*, 1995) particularly among marginalized or secluded populations. The invasion of cholera in Nichobar island (India) among tribal population in 2002 is another startling event in the history of cholera where *V. cholerae* gained access to this hitherto unaffected population (Sugunan *et al.*, 2004). Cholera has been reported in coastal districts of Odisha, since two decades (Niyogi *et al.*, 1994; Pal *et al.*, 2000; Chhotray *et al.*, 2002; Samal *et al.*, 2001). However its etiology was not reported among primitive tribes living in the hill regions of Northern plateau and Eastern Ghats, 500-600 kilometer away from the Bay of Bengal.

Odisha an eastern peninsular state of Indian subcontinent occupies a unique place in the tribal map including primitive 13 tribes. Amongst these 13 primitive tribes, 4 primitive tribes (Bondo, Didayi, Juanga and Kondha) belonging to the Mundari group of Proto-Astroid family are very poor in health status due to their isolation, remoteness, illiteracy, poverty, faulty feeding habits and being largely unaffected by the developmental processes going in the country. Diarrhoeal disorder is one of the major public health problems among primitive tribes with increased morbidity and higher mortality. *V. cholerae* being one of the important potential etiological agents of diarrhoeal disease, the knowledge of its incidence was still lacked among the primitive tribes in Odisha. This is the first study to report the incidence, origin and spread of *V. cholerae* causing diarrhoea amongst the four primitive tribes.

**Materials and Method**

**Study population**

A cross sectional study was undertaken from 2000 to 2004 amongst 4 primitive tribes (Bondo, Didayi, Kondha and Juanga) living in four different geographical regions of the state (Fig-1). The study was conducted in four phases in 17 villages from Bondo, 15 from Didayi, 25 from Kondha and 35 from Juanga, which were selected by PPS (population proportion size) sampling procedure. The sample size was calculated taking into account the prevalence rate of diarrhoeal diseases from the government of Odisha, health department (statistics). The study population includes 1012 Bondo, 1009 Didayi, 1298 Kondho and 1137 Juanga tribes. All the diarrhea patients admitted to hospitals of that locality were recruited during the study period. Informed consent was obtained from patients and/or close relatives. The study was reviewed and approved by the ethical committee of Regional Medical Research Center (RMRC) under Indian Council of Medical Research (ICMR).

**Bacteriology and Serogroup**

Rectal swabs were collected from freshly admitted cases in the hospitals in tribal areas having acute diarrhoeal symptoms before any treatment was initiated by the government health authorities. Rectal swabs in Cary Blair transport medium (CBT) were transported to the Regional Medical Research Center (RMRC) laboratory for bacteriological analysis to isolate *V. cholerae*
Diarrhea is defined as three or more watery or loose stools in a 24-hour period prior to admission to the hospital. A multi-test medium was used for presumptive identification of V. cholerae (WHO, 1987; Kaper, 1979). All the strains were subsequently subjected for oxidase reaction. Serology was done using slide agglutination with polyvalent and mono-valent antisera specific for Ogawa and Inaba (Difco, USA) and O139 antisera prepared and supplied by National Institute of cholera and Enteric Diseases (NICED), Kolkata, India. Strains of V. cholerae from each serogroup of O1 and O139 were selected for molecular typing based on antibiograms and geographical locations of tribal and non-tribals for comparison.

Antimicrobial susceptibility

Antimicrobial sensitivity pattern was tested for isolated V. cholerae strains following the Kirby-Bauer technique (16) with the antibiotics (Hi-media Laboratories, Bombay, India) ampicillin (A, 10μg), chloramphenicol (C, 30μg), co-trimoxazole (Co, 25μg), ciprofloxacin (Cf, 5μg), furazolidone (fz, 100μg), gentamicin (G, 10μg), neomycin (N, 30g), nalidixic acid (Na, 30μg), norfloxacin (Nx, 10μg), streptomycin (S, 10μg) and tetracycline (T, 30μg). Characterization of strains as susceptible, intermediate or resistant was determined based on the size of the inhibition zone around each disc according to manufacturer’s instruction. Strains showing an intermediate zone of inhibition were interpreted as resistant to that drug on the basis of previous minimum inhibition of concentration (MIC) studies conducted with V. cholerae (Yamamoto et al., 1995).

Cholera toxin assay

A monosialoganglioside (GM1) enzyme-linked immunosorbent assay (ELISA) was used to examine cholera toxin production synthesized by isolated V. cholerae O1 and O139 including two control strains (VC20, El Tor Biotype, O1 and SG24, O139) by the method Svennerholm and Holmgren (Svennerholm and Holmgren, 1978).

Molecular analysis

Polymerase Chain Reaction (PCR) assay

The virulence factors of V. cholerae encoding ctxA (A-sub unit of cholera toxin gene), tcpA (major structural subunit of toxin co-regulated pilus), zot (zonnula occluden toxin) and ace (accessory cholera entero toxin) were detected by PCR technique using specific primers as described elsewhere (Chhotray et al., 2002; Shi et al., 1998). Randomly amplified polymorphic DNA (RAPD) PCR assay was performed selected V. cholerae strains with 1281 and 1283 primers to check the clonality as in previous study (Chhotray et al., 2002). The fingerprinting pattern obtained from RAPD (1281) of tribal and coastal strains were compared to ascertain the phylogenetic relationship between the two groups by using Quantity-1 software and dendrogram was generated using SAS (SAS Institute, 1989).

Pulsed Field Gel Electrophoresis (PFGE)

Genomic DNA of strains of V. cholerae was prepared in agarose plugs as described previously by (Sinha et al., 2001) to carryout PFGE. PFGE was performed using the counter clamped homogenous electric field method on a CHEF Mapper system (BioRad, California, USA) with 1% PFGE grade agarose in 0.5X TBE buffer for 26 hour. Run conditions were generated by the autoalgorithm mode of CHEF Mapper PFGE system using a size range of 20-300kb. The gels were stained for 30 min with solution of 20-10μg/ml ethidium bromide, destained in water for 15 min and photographed under UV
Results and Discussion

A total of 1066 rectal swabs collected from patients with secretory diarrhoea were bacteriologically analyzed during the period under study. Of the total samples *V. cholerae* O1 Ogawa was 9.8% and *V. cholerae* O139 3.8% serogroup, Table-1.

The antibiogram profile of strains of *V. cholerae* O1 & O139 against commonly used antibiotics, revealed interesting shifts in the patterns of resistance during the study period. The dominant drug resistance profile of *V. cholerae* O1 and O139 in 2000 and 2001 were ACoFzNNaS and AFzNSNa respectively. The incidence of fluoroquinolone resistant isolates of O1 & O139 was observed in variable proportion. The striking feature is the emergence of complete (100%) nalidixic acid resistant strains of *V. cholerae* O139 for the first time during the study. Thus the results obtained demonstrate that *V. cholerae* O1 & O139 isolated during the study period had high rates of resistant to various antibiotics tested.

All the strains of *V. cholerae* O1 & O139 were analyzed using specific primers to detect ctxA, tcpA, and two other genes zot and ace which putatively encode additional toxins. Our results revealed the presence of the core toxin region among the isolates of O1 and O139 and confirmed the co-occurrence of tcpA (El Tor), ctxA, zot and ace in all O1 and O139 strains.

RAPD PCR fingerprinting carried out on the five representative isolates from each of the two serogroups generated identical arrays of DNA fragments with one primer, 1283 (Fig 2a, b). Interestingly, the other primer (1281) (Fig3a,b) exhibited a slightly varied RAPD PCR profile for the O1 and O139 strains of Primitive tribes from those of the other part of the state as well as of the country. The relatedness between the genomic DNA fingerprinting obtained from RAPD band profile of the O1 and O139 strains of coastal and tribal district were estimated and presented as a dendrogram (Fig 4a, b) using average linkage determined by the CLUSTER and TREE procedure SAS.

PFGE profiles of strains of both the serogroups from the study area and from the other geographical locations of the state is shown in Fig-5. All the 4 strains of O1 Ogawa examined, showed identical banding pattern to that exhibited by CO 840, the new clone of O1, which appeared in Calcutta in 1994. Similarly, the banding pattern of four O139 strains of both tribal and coastal districts matched with that of O139 reference strain ATCC 51394, the Madras (India) outbreak strain in 1992. Thus, each serogroup of O1 and O139 of tribal and coastal districts of widely distanced geographical locations exhibited similar PFGE profile as prevailing in other parts of the country.

In the present study the outbreak of cholera was reported in 2000 due to *V. cholerae* O1 and O139 among the four primitive tribes in different geographical regions, O1 being the predominant serogroup. All the O1 strains were biotype El Tor, Ogawa, similar to the trend documented in other part of the state as well as in India. Very interestingly although *V. cholerae* O139 after its emergence in Madras in 1992 was spread to Kolkata in 1992 and Visakhapatanum in 1993 (Nair et al., 1994), its emergence was not reported till Sept 1995 (Pal et al., 2000) in Odisha which is situated midway between these two places. After emergence in 1995 (Pal et al., 2000) and reemergence in June 1999 in coastal district (Unpublished data), O139 gained it’s entry to the tribal land during November 2000 for the first time. In the present study the incidence and isolation of cholera was on a
higher side during the interval of first year of study. There was a gradual decrease in the isolation rate of *V. cholerae* in the second and third year and there was not a single case of cholera reported during fourth year in whole of the study area. This may be due to the institution of effective intervention measures in the form of immediate treatment, health awareness and intense IEC activities or the spread might be temporary at high hilltop region.

The results of antibiogram pattern demonstrate that *V. cholerae* isolated during 2000 in the study area had resistance to various commonly used antibiotics. The emergence of fluoroquinolone resistant *V. cholerae* strains, which is being reported during the first year of the study, bear an important testimony in the therapeutic use of antibiotics during the outbreaks. Again emergence of nalidixic acid resistant strains of *V. cholerae* O139 in 2000 represent another shift and is noteworthy because it was sensitive to nalidixic acid since its’ emergence at Kolkata and elsewhere (Nair *et al.*, 1994). Emergence of fluoroquinolone resistant *V. cholerae* O1 and O139 have appeared as a great threat in therapeutic practice as ciprofloxacin and norfloxacin are broad-spectrum fluoroquinolone and possess excellent activity against *V. cholerae* O1 and O139 serogroup (Yamamoto *et al.*, 1995). These drugs are effective in treatment of cholera among adults and children (Bhattacharya *et al.*, 1990). Possibly selective pressure exerted by nalidixic acid, disproportionate use of fluoroquinolone and increase in the incidence of nalidixic acid resistant strains of O1 and O139 (due to single mutation in gyrA and other related gene) might be the leading cause of appearance of fluoroquinolone resistance (with two or more mutation at gyrA).

The phenotypic character of synthesis of CT exhibited by both O1 and O139 strains had the similar feature producing almost equal quantity. Cholera toxin is the major cause of profuse watery diarrhoea caused by *V. cholerae* that is the hallmark of cholera (Kaper *et al.*, 1994).

| Year     | 2000-2001 | 2001-2002 | 2002-2003 | 2003-2004 | Total |
|----------|-----------|-----------|-----------|-----------|-------|
| Rectal swabs | 222       | 332       | 276       | 236       | 1066  |
| B-70     | 32         | 70        | 41        | 50        | 105   |
| D-41     | 41         | 73        | 64        | 52        | 230   |
| K-67     | 67         | 93        | 82        | 54        | 296   |
| J-44     | 44         | 74        | 52        | 54        | 226   |

|          | B-70       | 32         | 67         | 52        | 165   |
|-----------|------------|------------|------------|-----------|-------|
| V. cholerae O1 (%) | 48 (36.3) | 14 (6.7)  | 5 (3.0)   | 00        | 61(9.8)|
| V. cholerae O139 (%) | 18 (13.6) | 6 (2.8)   | 00         | 00        | 24(3.8)|

Table 1 Year-wise distribution of *V. cholerae* from diarrhoea patients among Primitive tribes, Odisha during 2000-2004
Fig. 1 Map of Orissa showing the study area of four primitive tribes, Coastal districts & Regional Medical Research Center (RMRC)
**Fig. 2** Comparison of the RAPD profiles of *V. cholerae* O1 (a) and O139 (b) isolated from primitive tribes of Orissa using primer 1283. (a) and (b) Lane 1 through 5: POCY 130, *V. cholerae* O1 from coastal area, JP1, JP23, JP3, & JP4, *V. cholerae* O1 from tribal districts. (b) ) Lane 1 through 5: POCY194, *V. cholerae* O139 of coastal district; JP6, JP2, KH4, *V. cholerae* O139 from tribal districts.

**Fig. 3** RAPD profiles of the *V. cholerae* O1 (a) and O139 (b) isolated from primitive tribes of Orissa using primer 1281 (a) Lane1: O395, *V. cholerae* O1 classical; Lane2: *V. cholerae* biotype El Tor 20; Lane3 through 8 (JP1, JP23, JP3, JP4, JP29), *V. cholerae* O1 from tribal district and POCY 130, *V. cholerae* O1 from coastal area. (b) Lane1:100 bp ladder, Lane2: POCY194, *V. cholerae* O139 of coastal district; Lane3 through 7 (JP6, JP2, KH4, JP11 & KH3) *V. cholerae* O139 from tribal districts.
**Fig. 4a** UPGMA (unweighted pair-group method using arithmetic averages) dendrogram for *V.cholerae* O1 of coastal and tribal districts of Orissa based on average linkage cluster analysis using RAPD marker (1281)

![Dendrogram for V.cholerae O1](image)

**Fig. 4b** UPGMA (unweighted pair-group method using arithmetic averages) dendrogram for *V.cholerae* O139 of coastal and tribal districts of Orissa based on average linkage cluster analysis using RAPD marker (1281)

![Dendrogram for V.cholerae O139](image)
**Fig.5** PFGE profile generated with Not1 digested genomic DNA of *V. cholerae* O1, El Tor Ogawa, biotype and O139 strains. Lane1: Bacteriophage λ- ladder; Lane2: 569B, O1 Classical, Inaba; Lane3: CO840, El Tor O1, Ogawa; Lane4 O139 reference strain ATCC 51394; Lane5: PU16 (O1, Ogawa); Lane6: JP23 (O1, Ogawa); Lane7: JP3 (O1, Ogawa); Lane8: JP4 (O1, Ogawa); Lane9: POCY194 (O139); Lane10: KH3 (O139); Lane11: JP2 (O139); Lane12: KH4 (O139).

*In vitro* comparative analysis of CT during the present study revealed almost equal amount of CT synthesis by both the serogroup indicating O139 strains are also equally invasive as O1 contrary to the earlier hypothesis (Sinha *et al.*, 2000; Nair *et al.*, 1994).

Production of equal quantity of CT corroborates the similar spectrum of severity of the disease infected by both the serogroups. Again production of almost equal quantity of cholera toxin by *V. cholerae* O1 and O139 isolates bear another homologous character between the strains of two geographical regions of tribal and coastal districts.

Phenotypic characterization is less discriminatory in identifying the clonal nature of bacterial strains within a serogroup. To overcome this many molecular techniques have been used recently to distinguish clones of enteric pathogens for epidemiological investigation. The data obtained from RAPD fingerprinting analysis, substantiates the existence of a single clone in each serogroup within the location of 4 primitive tribes except some minor variations similar to the clones present in other part of the state which corroborate our earlier reports (Chhotray *et al.*, 2002). The variations in the RAPD PCR profile exhibited by the O1& O139 serogroup isolated from the geographical regions of primitive tribes may be due to the interplay of genetic changes and natural selection caused by unidentified environmental factors or due to the consequences of rapid changes in drug resistance pattern.

However it requires further investigation by characterization of the varied region of the chromosome towards a better understanding of the parameters for the microbial evolution. The relatedness between the RAPD band pattern of the tribal and coastal strains of O1 and O139 were estimated and represented as dendogram (Fig 4a & 4b). In the cluster
analysis we found that tribal strains were clonal in nature with the coastal strains.

The PFGE patterns of *V. cholerae* O1 and O139 strains isolated from primitive tribal population were not different from the strains of coastal endemic regions Previous studies have identified the clones of each serogroup present in the state is similar to the clones present in the Indian subcontinent (Chhotray *et al.*, 2002; Sinha *et al.*, 2000). From the present study it is evident that the phenotypic and genotypic expression of *V. cholerae* strains of each serogroup isolated in the primitive tribal region are similar to the strains of coastal region of Orissa which in turn is similar to the clone prevails in India.

The outbreaks of cholera among primitive tribes may be correlated with the rapid spread from the outbreak areas of coastal districts, which occurred after a period of six months gap of super cyclonic cholera due to population migration through different routes to tribal areas where developmental activities are on progress. It is worth to mention that due to the super cyclone in November 1999, a large outbreak of cholera was occurred in the 9 coastal districts of Orissa due to *V. cholerae* O1 and O139 (Chhotray *et al.*, 2002). This requires a continuous surveillance in these difficult and remote tribal areas of the state.

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