**ABSTRACT**

Introduction: Cholera is one of the most common diarrhoeal diseases in Nepal. Etiological agent of cholera is *Vibrio cholerae* which removes essential body fluids, salts and vital nutrients, which are necessary for life causing dehydration and malnutrition. Emerging antimicrobial resistant is common. The aim of the present study was to determine the antibiotic susceptibility pattern of cholera patients in Nepal.

Methods: All the laboratory works were conducted in the bacteriology section of National Public Health Laboratory, Teku, Kathmandu, from March to September 2005. During this period a total of 340 stool samples from diarrhoeal patients were collected and processed according to the standard laboratory methods. Each patient suffering from diarrhoea was directly interviewed for his or her clinical history during sample collection.

Results: A total of 340 stool samples were processed and studied from both sex including all ages of patients. Among the processed sample 53 *Vibrio cholerae* cases were found. All isolated *Vibrio cholerae* O1 were El Tor, Inaba. All isolated (100%) *Vibrio cholerae* O1 were sensitive to Ampicillin, Ciprofloxacin, Erythromycin and Tetracycline whereas all were resistant to Nalidixic acid and Cotrimoxazole. Only 15.1% cases were sensitive to Furazolidone whereas 84.9% were resistant.

Conclusion: All *V. cholerae* strains isolated in this study were found resistant to Multi Drug Resistant (resistant to at least two antibiotics of different group). Ampicillin, Ciprofloxacin, Erythromycin and Tetracycline were found still more potent antibiotics against *Vibrio cholerae* isolated during the study.

**Keywords:** antibiotics, susceptibility, *Vibrio cholerae*

**INTRODUCTION**

Cholera is a disease caused by *Vibrio cholerae*. Since 1817, six pandemics have swept over the world, and the seventh one is in progress. Seventh pandemic was started from an Indonesian island which was caused by the E1 Tor biotype and it spread across Asia and Africa. In Nepal, owing to the low socioeconomic status and poor hygienic condition of the people, Cholera constitute an important cause of morbidity and mortality. The case fatality rate in Nepal from diarrhoea is highest (2%) in the age group of 1-4 and is lowest (1%) among the infants.

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There are several serogroups of *V. cholera*, the O1 serogroup being the cause of endemic and epidemic cholera. Using specific antisera to detect different fractions of the O antigen, the O1 serogroup is divided into Inaba (AC), Ogawa (AB) and Hikojima (ABC) serotypes. Drug resistant *V. cholera* have been reported since 1977 and some of these strains were mult drug resistant.6,7 Due to the development of these resistant strains, it is now important to know the local antimicrobial susceptibility patterns of *V. cholerae*.

**METHODS**

This observational study was conducted in the bacteriology section of National Public Health Laboratory (NPHL), Teku, Kathmandu from March to September 2005, following standard microbiological techniques. Each patient suffering from diarrhoea was directly interviewed for his or her demographic profile (name, age, sex, address and hospital treated) and significant clinical history (sign, symptoms, stool passage frequency, stool consistency, presence of blood and mucus in the stool) during sample collection. Data was entered on a standard excel worksheet. Data analysis was performed by using Microsoft Excel 2007 and statistical package of social sciences (SPSS).

Stool sample was primarily inoculated on the plate of MacConkey Agar (MA) medium and Thiosulphate Citrate Bile Salt Sucrose (TCBS) medium, at the same time a loopful sample was inoculated in Alkaline Peptone Water (APW) for enrichment and incubated for 4-6 hours. From APW, sample was subcultured on second plate of MA and TCBS. All inoculated plates were incubated aerobically at 37ºC for 24 hours. *Vibrio* suspected non-lactose fermenting colonies on MA and sucrose fermenting yellow colonies on TCBS were subcultured on Nutrient Agar (NA) medium and incubated for 24 hours at 37ºC. Further processing was performed from the colonies on NA plate.

Identification was done by colony characteristics, Gram’s stain, different biochemical tests (Different sugar utilization, Oxidase, Catalase, Indole, Methyl Red, Voges-Proskaur, Citrate Utilization, Oxidative-Fermentative, Motility, Urea Hydrolysis, Salt Concentration Tolerance and String test), serotyping (O-antigen groups were serologically identified by using commercially available polyvalent O, polyvalent O1, monospecific Ogawa-Inaba antisera and with specific anti-O 139 antisera, Denka Seiken Co., Japan), and biotyping of *Vibrio cholerae* O1 (Haemagglutination of Chick RBC, Haemolysis of Sheep RBC, Susceptibility to Polymyxin B and VP Reaction). Antibiotic susceptibility testing was done on Mueller-Hinton Agar plate by Kirby-Bauer disc diffusion method.

**RESULTS**

Out of 340 patients, 82.94% were from Shukraraj Tropical Hospital and 17.05% were from different locality of Kathmandu valley attending NPHL by requesting stool culture. There were total of 53 (15.58%) strains of *V. cholerae* and all of them were El Tor, Inaba. Association of isolation between male and female was not statistically significant. Majority of the patients were belonged to age group 21-30 years (Table 1). Isolation rate was found highest in August (45.28%) (Figure 1). Vomiting, dehydration, nausea and passage of rice watery stool with more than 10 times per day were reported as the main clinical features of the cholera (Table 2 & 3). All isolates were sensitive to Ciprofl oxacin, Ampicillin, Erythromycin and Tetracycline. Only 15.1% isolates were sensitive towards Furazolidon but all isolates were resistant to Nalidixic acid and Cotrimoxazole (Table 4). Hundred percent *V. cholerae* isolates were Multi Drug resistant.

**Table 1. Age and gender wise distribution of samples and *V. cholerae* cases**

<table>
<thead>
<tr>
<th>Age groups</th>
<th>Male No. of patient</th>
<th>V. cholera cases</th>
<th>Female No. of patient</th>
<th>V. cholera cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-10yr</td>
<td>19</td>
<td>1</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>11-20yr</td>
<td>38</td>
<td>9</td>
<td>19</td>
<td>3</td>
</tr>
<tr>
<td>21-30yr</td>
<td>54</td>
<td>10</td>
<td>42</td>
<td>9</td>
</tr>
<tr>
<td>31-40yr</td>
<td>37</td>
<td>5</td>
<td>25</td>
<td>6</td>
</tr>
<tr>
<td>41-50yr</td>
<td>24</td>
<td>3</td>
<td>14</td>
<td>2</td>
</tr>
<tr>
<td>51-60yr</td>
<td>18</td>
<td>2</td>
<td>16</td>
<td>2</td>
</tr>
<tr>
<td>61-70yr</td>
<td>6</td>
<td>0</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>71-80yr</td>
<td>3</td>
<td>1</td>
<td>5</td>
<td>0</td>
</tr>
<tr>
<td>&gt; 80yr</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Total no. (%)</td>
<td>199(58.52%)</td>
<td>31(58.49%)</td>
<td>141(41.47%)</td>
<td>22(41.75%)</td>
</tr>
</tbody>
</table>
Table 2. Clinical symptoms exhibited by cholera patients

<table>
<thead>
<tr>
<th>No. of V. cholerae isolates</th>
<th>Abdominal pain/cramp</th>
<th>Fever</th>
<th>Vomiting</th>
<th>Dehydration</th>
<th>Nausea</th>
</tr>
</thead>
<tbody>
<tr>
<td>53</td>
<td>6</td>
<td>2</td>
<td>51</td>
<td>47</td>
<td>31</td>
</tr>
<tr>
<td></td>
<td>(11.3%)</td>
<td>(3.77%)</td>
<td>(96.2%)</td>
<td>(88.69%)</td>
<td>(58.4%)</td>
</tr>
</tbody>
</table>

Table 3. Physical characteristics of stool exhibited by cholera patients

<table>
<thead>
<tr>
<th>Stool passage frequency Per day (%)</th>
<th>Stool texture (Consistency)</th>
<th>Presence of blood or mucus in the stool</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥3</td>
<td>Loose</td>
<td>Rice watery</td>
</tr>
<tr>
<td>≥10</td>
<td>Watery</td>
<td>Bloody</td>
</tr>
<tr>
<td>6(11.32%)</td>
<td>0</td>
<td>13(24.52%)</td>
</tr>
<tr>
<td>47(88.67%)</td>
<td>40(75.47%)</td>
<td>11(20.7%)</td>
</tr>
</tbody>
</table>

Figure 1. Seasonal distribution of V. cholerae cases

DISCUSSION

Cholera is an important cause of acute diarrheal diseases in Nepal. Poor environmental hygiene, poor water sanitation and improper management of sewage wastes are the predominant factors that V. cholerae are established in Kathmandu valley. V. cholerae is one of the most important enteropathogens in all countries. The sixth pandemic of cholera and, presumably, the earlier pandemics were caused by the classical biotype of Vibrio cholerae O1, which was progressively replaced by the El Tor biotype representing the seventh cholera pandemic. Eighth pandemic in cholera history, could be due to Vibrio cholerae O1 El Tor, Inaba serotype, infection. Our result also support this view.

It was reported that for the last six years V. cholerae El Tor O1 Ogawa was responsible for the cholera endemic in Kathmandu, but in 1990 it was V. cholerae El Tor O1 Inaba, so it can be said that after 15 years later V. cholerae O1 Inaba emerged hugely in Kathmandu. Some previous outbreaks of cholera in Kathmandu valley like, in 1994 and in 2004 were due to V. cholerae, El Tor, O1 Ogawa. Although Ogawa was predominant, frequently Hikojima and to less often Inaba serotypes was also isolated in Nepal.

Vibrio cholerae O1 strains with a periodic shift of different serotypes (Inaba and Ogawa) may be a consequence of genetic reversion which is known to take place in vitro and vivo. The only changes observed between the Ogawa and Inaba serotypes are related to a mutation in the wbeT region on a 21.6-kb SacI fragment of DNA. The epidemic strains had undergone a serotype conversion, possibly because of immune pressure in the population; or due to seletive pressure of anti-Ogawa antibodies and the emergence of anti-Inaba antibodies.

In the present study, the highest cholera cases were seen among the age group between 21-30 years and were predominant among male patients. Although children are the primary victims of cholera, the present result may be due to higher frequency of adult patients visiting NPHL. The incidence of cholera are frequently reported in August and September. Vomiting, dehydration, nausea with rice watery stool and stool passage frequency ≥10 were found the important clinical manifestations exhibited by cholera patients which are the universal clinical features of cholera.

Antibiotics shorten the course and diminish the severity of the illness, but they are not as important as rehydration, which is the cornerstone of therapy for cholera. The susceptibility pattern of the isolates, against seven different antibiotics, belonging to different groups were studied and found all with multiple antibiotic resistant capacity. It appears that antibiotic-resistant strains are
increasingly being found worldwide.\textsuperscript{20,21} Since, Tetracycline is the drug of choice for cholera because it is more effective, non-toxic, easily available and comparatively cheap. In our study and in some other studies, all isolates were sensitive to Tetracycline and Cotrimoxazole.\textsuperscript{22} But Tetracycline, Cotrimoxazole and Furazolidon resistant \textit{V. cholerae} strains were also reported.\textsuperscript{11,23} Resistance to Cotrimoxazole, Furazolidone and Streptomycin suggests the possibility of the presence of the SXT element in Inaba strains.\textsuperscript{24} Genes and their mechanisms of transfer may be important in the maintenance and transfer of resistance among \textit{V. cholerae} and other enteric pathogens. In 1989, Erythromycin was reported to be effective for the treatment of Tetracycline resistant strains.\textsuperscript{25} All isolates in this study were found sensitive towards Erythromycin.

This study can help health authorities to review the cholera treatment guideline. Health education campaigns about prevention of diarrhoeal diseases and proper disposal of sewage, good sanitation and hygiene may be effective practice to control the cholera in Kathmandu Valley.

CONCLUSIONS

In Kathmandu Valley there is an incidence of cholera in the rainy season almost every year. The marked ability of \textit{V. cholerae} isolates to resist the inhibitory action of antibiotics has become a global problem. Increasing antibiotic resistance against commonly prescribed drugs signify that treatment options have become difficult in cases of cholera. The results indicate a continuing need for resistance surveillance and rational use of antimicrobial agents to reduce the multi-drug resistant strains of \textit{Vibrio}.

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