Characterization of group B coxsackieviruses isolated from non-polio acute flaccid paralysis patients in Pakistan: vital assessment before polio eradication

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Summary
Pakistan is at the verge of polio eradication but isolation of non-polio enteroviruses (NPEVs) from acute flaccid paralysis (AFP) cases may result in serious or even fatal outcome. Many enteroviruses share similar symptoms and epidemiology as is the case with poliovirus and coxsackievirus (CV). The present study was designed to genetically characterize coxsackievirus B (CV-B) serotypes isolated from non-polio acute flaccid paralytic children, as well as to understand their probable role in paralysis. A total of 63 (20·1%) out of 313 stool samples during 2013 were found positive for NPEVs in rhabdomyosarcoma cells. Only 24 (38·0%) NPEVs were typed as CV-B by microneutralization assay and were further characterized by sequencing of the viral protein 1 (VP1) gene. Molecular phylogenetic analyses classified the study strains into six coxsackievirus B serotypes (coxsackievirus B1 to B6) with their respective prototype strains with evidence of epidemiological linkage and distinct clusters. Moreover, four major differences were found within the amino acid sequences of BC-loop in VP1 of CV-B strains. In conclusion, this study presented the molecular evolutionary genetic overview and distinct phylogenetic pattern of CV-B isolates from AFP cases in Pakistan, and explored the possible link between CV-B infections and AFP cases. Furthermore, our data reveal that these viruses might contribute to the incidence of paralysis in population and there is need of time to establish an enterovirus surveillance system for better understanding of epidemiological and virological characteristics of NPEV infections associated with AFP cases in the country.

Key words: Acute flaccid paralysis, coxsackievirus B, enterovirus, molecular epidemiology, Pakistan.

INTRODUCTION
Coxsackievirus (CV) is a member of species Enterovirus B of genus Enterovirus in the family Picornaviridae [1]. Originally, these viruses were discovered in 1948 by Gilbert Dalldorf from the fecal specimens of children with poliomyelitis like illness, residing in town ‘Coxsackie’ situated on the Hudson River in New York [2]. These are non-enveloped, single-stranded, positive sense RNA viruses of 7–8 kb pairs long and were divided into two subgroups called A and B based on characteristic disease induced in suckling mice inoculated intracerebrally.
Coxsackievirus A (CV-A) cause a flaccid paralysis by infecting skeletal muscle, whereas coxsackievirus B (CV-B) cause a spastic paralysis due to focal muscle injury and degeneration of neuronal tissue [3]. Currently, 23 CV-A and six CV-B serotypes are recognized [4].

CV-B have worldwide distribution with different geographic patterns of circulation. These viruses have long been recognized as significant pathogen of infants and children and are also known as pediatric pathogen [5]. CV-B infections can result in various diseases, including aseptic meningitis, acute myocarditis and pericarditis, but most commonly recognized serious manifestation of CV-B infection is associated with paralysis alone or in combination with encephalitis. The presentation may mimic that of poliovirus related paralysis with fever at the time of onset of paralysis and a rapid progression to flaccid paralysis [6]. Nearly all serotypes of CV-B have been associated with paralytic infections [7]. These viruses can be isolated from CSF (cerebrospinal fluid), stool, rectal swabs, throat swabs and nasal secretions, but the isolation from stool is most sensitive due to the presence of virus in higher titer and longer than in any other specimen [8].

The application of molecular biology techniques to clinical virology has begun to significantly change approaches to enterovirus diagnostics. The sequence analysis of viral protein 1 (VPI; one of the main exposed regions of the viral capsid) has been successfully implemented for enterovirus serotype identification [9]. Currently, the evolutionary distances and phylogenetic investigations based on VPI gene are used for the identification and classification of CV-B [9]. In addition, phylogenetic analyses based on sequencing data of VPI region is considered to be the standard method of molecular analysis for epidemiological purpose or clustering or genotyping or to identify emerging new variants [10].

The objective of this study was to understand the molecular epidemiology of group B CVs isolated from the stool of non-polio acute flaccid paralytic children and to identify their probable role in causing paralysis from the perspective of poliomyelitis eradication.

METHODS

Ethics statement

This study was approved by Institutional Internal Review Committee of the National Institute of Health, Pakistan. Written informed consent was obtained from all the individuals participated. A check box was included in data form to document the consent taking procedure.

Sample collection

Stool samples (n = 313) were collected during 2013 from patients with acute flaccid paralysis (AFP) under the age of 15 years. All stool samples collected according to World Health Organization (WHO) guidelines within 14 days from the date of onset and were delivered to WHO Regional Reference Laboratory for Polio Eradication Initiative, Virology Department, National Institute of Health Islamabad, Pakistan and stored at −20 °C.

Virus isolation and microneutralization

All these samples were processed as per WHO standard procedures [11]. Briefly, these samples were processed and inoculated into RD (human rhabdomyosarcoma) and L20B (recombinant murine cell lines expressing human poliovirus receptor) cells and were observed daily for the development of a characteristic enterovirus cytopathic effect (CPE). Isolates from RD cells were re-passaged to L20B cells, and were designated as non-polio enterovirus (NPEV) if no CPE was observed. According to WHO standard protocols, typing of the NPEV isolates was carried out by microneutralization using typing sera kit [provided by National Institute for Public Health and the Environment (RIVM), Bilthoven] containing the polyclonal antisera to identify 23 enterovirus serotypes [11].

VPI amplification

RNA was extracted from 140 µl of the infected cell culture using QIAamp viral RNA extraction kit (Qiagen, Valencia, CA, USA). NPEVs were confirmed by real-time reverse transcription polymerase chain reaction (qRT–PCR) targeting the 5′ untranslated region [12]. Reverse transcription–polymerase chain reaction (RT–PCR) was performed to amplify the VPI gene of NPEVs. Initially, complementary DNA (cDNA) was synthesized in a 20 µl reaction mixture comprising of 11 µl of each viral RNA, 4 µl of 5× transcriptase buffer, 2 µl of 10 mM dithiothreitol (DTT), 0·5 µl of 25 mM each deoxynucleotide (dNTP) solution mix (Roche), 1 µl of 40 U of ribonuclease Inhibitor, 1 µl of 20 U Avian Myeloblastosis Virus (AMV) reverse transcriptase (Thermo Scientific) and 0·5 µl of nuclease free
Water at 42°C for 45 min. For amplification of VP1 region RCR was performed by using 2 µl of cDNA, 5 µl of 10×Taq buffer, 3.5 µl of MgCl₂ (50 mM), 1× µl of 12.5 mM dNTPs, 0.5 µl of Taq DNA polymerase (5 units/µl), 37 µl of water and 0.5 µl of each primer (10 µM) 224 and 222 targeting VP3 (nt 2204–2223) and VP1 (nt 2969–2951) region of virus genome, respectively [13]. Thermocycler parameters for RT–PCR was 95°C for 3 min followed by 40 cycles of 94°C for 45 s, 42°C for 45 s and 72°C for 1 min. Specific RT–PCR products were purified using the Qiaquick PCR purification kit (Qiagen) and stored at −20°C till further procedure.

Nucleotide sequencing and phylogenetic analyses

The purified products were sequenced using the same forward and reverse primers as used in RT–PCR with the BigDye Terminator v3.1 cycle sequencing kit (Applied Biosystems, Foster City, CA), and sequencing was performed with ABI 3130 genetic analyzer (Applied Biosystems). The obtained sequences were curated and aligned using Sequencher version 4.9 software (GeneCodes) (http://www.gene codes.com). Molecular typing based on VP1 sequences was performed using automated genotyping tool for enterovirus version 0.1 [14]. VP1 sequences were also subjected to online BLAST analyses and the sequence of CV-B prototype and all the closest relative strains of study isolates retrieved from GenBank (http://www.ncbi.nlm.nih.gov/BLAST). These sequences were aligned using ClustalW. The genetic distances were calculated by using Kimura-2 parameter model and Jones–Thornton–Taylor (JTT) model for nucleotide and deduced amino acid sequences, respectively. Phylogenetic tree was constructed using the neighbor-joining method with 1000 bootstrap replicates in MEGA software (version 5.0) [15]. Geneious Package (version 10.1.3; http://www.geneious.com) was used for VP1 deduced amino acid sequence analysis and to generate sequence logos, with the column height signifies conservation of the sequence at that position, while the height of the amino acids within the column shows relative frequency [16, 17].

The sequence data described in this study have been submitted to GenBank data library under accession numbers KY593460–KY593483.

RESULTS

A total of 313 stool samples from AFP patients were processed and inoculated on RD and L20B cell lines. Overall on cell culture, 63 (20.1%) stool samples were found positive for NPEV, 110 (35.1%) for poliovirus and 140 (44.7%) remained negative. Microneutralization assay successfully typed 56 (88.8%) NPEV isolates into echovirus (n = 25; 44.7%), numbered enterovirus (n = 7; 12.5%) and CV-B (n = 24; 42.8%), while seven (11.1%) isolates remained untypeable. All CV-B isolates were subjected to RT–PCR and characterized by VP1 gene sequencing. The partial (479 bp) VP1 nucleotide sequences of these isolates were compared with the CV-B sequences selected among those available in GenBank. The prototype sequence of each CV-B serotype was also included. BLAST analysis of the sequences indicated homology of the study strains to CV-B1 (n = 3), CV-B2 (n = 1), CV-B3 (n = 2), CV-B4 (n = 1), CV-B5 (n = 12) and CV-B6 (n = 5). The clinical and demographic details of all CV-B strains were summarized in Table 1.

Phylogenetic analysis

A phylogenetic tree was constructed based on partial (479 bp) VP1 gene sequences of CV-B strains (n = 24) of the present study to determine the phylogenetic clustering among isolates and to analyze the genetic diversity and molecular aspects of CV-B epidemiology. The VP1 sequence of CV-A1 was included in the phylogenetic tree as an out group. All isolates clustered into six distinct CV-B serotypes (CV-B1 to CV-B6) with their respective prototype strains (Fig. 1). The genetic relationships of the study strains within and between CV-B serotypes were further analyzed.

The analysis revealed that three Pakistan (PAK) strains, detected in 2013 from Balochistan (BN) and Khyber Pakhtunkhwa (KP) Provinces were clustered into the CV-B1 and showed 19.2% (17.2–20.9%) mean nucleotide divergence with their reference strain (accession no. M16560). In addition, two genetic clusters were identified in this serotype: one contained two study strains (PAK-MEV3757B and -3688B) from BN, and the other contained the one strain (PAK-MEV5726A) from KP. The nucleotide divergence of PAK-MEV5726A strain with PAK-MEV3757B and PAK-MEV3688B strains were 15.7% and 12.7%, respectively. The closest relative strain of PAK-MEV5726A was an Indian strain A210 (accession no. JX513569) having 6.7% nucleotide divergence. CV-B2 serotype contained only one study strain (PAK-MEV2058A) from Sindh (SD) province and had 16.4% nucleotide divergence with its prototype strain (Ohio-1; accession no. AF081312). Similarly, two strains (PAK-MEV4565A
and PAK-MEV1452A) isolated from Sindh clustered in CV-B3 and showed 21.3% nucleotide divergence with each other. In addition, CV-B4 serotype had only one study strain (PAK-MEV076B) isolated from KP province and their closest relatives were the strains (PK1Pak92, accession no. AF160019 and P234Pak92, accession no. AF160018) earlier isolated from Karachi, Pakistan in 1992 having 11.2% nucleotide divergence. Furthermore, phylogenetic analysis revealed the existence of two genetic clusters (I – II) in CV-B5 serotype. Each cluster differed with each other by at least 11.2% (11.2 – 14.2%) mean nucleotide divergence. Cluster I contained three strains (PAK-MEV1620B, -1177A, -467B) isolated from KP and Punjab (PB) patients, while cluster II contained nine strains (PAK-MEV5777A, -5659B, -3586B, -5926A, -3271A, -5959A, -2005B, -3903A and -5810B) from PB, SD and BN isolated in 2013. The closest relative of both clusters was strain (accession no. GU323568) isolated from Sweden and had mean nucleotide divergence 9.2% (9.0 – 9.7%) and 9.18% (8.2 – 11.9%), respectively. CV-B6 serotype is composed of five study strains collected from KP and SD provinces having 20.4% (20.1 – 20.9%) mean nucleotide divergence with their reference strain (Schmitt, accession no. AF081313). The closest relative of the CV-B6 study strains was isolate N-165 (accession no. JN203635) isolated from Indian patient having 15.2% (12.7 – 16.4%) mean nucleotide divergence.

**Genetic divergence among CV-B study isolates**

Six distinct CV-B serotypes could be individualized and appear more clearly in Figure 1. Pairwise sequence comparisons among CV-B study isolates are summarized in Table 2. Isolate of CV-B2 serotype was found the most divergent study strain and differed in nucleotide sequence from other CV-B isolates by 20.1 to 31.3% (11.4 – 20.5%) amino acid sequence difference while it showed 11.2 – 18.4% (2.0 – 4.0% amino acid) nucleotide divergence within the serotype. The more study isolates (n = 12) were classified into CV-B5 serotype. The strains within CV-B5 differed from one another by up to mean 8.0% (3.0 – 14.2%)

### Table 1. Clinical and demographic data of acute flaccid paralytic patients with CV-B infection characterized in this study

<table>
<thead>
<tr>
<th>Serotype identified in this study</th>
<th>Isolate ID</th>
<th>Sex</th>
<th>Age (months)</th>
<th>Fever</th>
<th>Asymmetric paralysis</th>
<th>Progression</th>
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</thead>
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<tr>
<td>CV-B1</td>
<td>PAK-MEV3757B</td>
<td>F</td>
<td>30</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
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<tr>
<td>CV-B1</td>
<td>PAK-MEV3688B</td>
<td>M</td>
<td>86</td>
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<td>Yes</td>
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<tr>
<td>CV-B1</td>
<td>PAK-MEV5726A</td>
<td>M</td>
<td>21</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B2</td>
<td>PAK-MEV2058A</td>
<td>F</td>
<td>24</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B3</td>
<td>PAK-MEV1452A</td>
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<td>36</td>
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<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B3</td>
<td>PAK-MEV4565A</td>
<td>M</td>
<td>24</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B4</td>
<td>PAK-MEV076B</td>
<td>M</td>
<td>36</td>
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<tr>
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<td>PAK-MEV1620B</td>
<td>M</td>
<td>4</td>
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<td>Yes</td>
<td>Yes</td>
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<tr>
<td>CV-B5</td>
<td>PAK-MEV1177A</td>
<td>M</td>
<td>16</td>
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<td>Yes</td>
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<tr>
<td>CV-B5</td>
<td>PAK-MEV467B</td>
<td>M</td>
<td>18</td>
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<td>PAK-MEV5777A</td>
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<td>36</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
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<tr>
<td>CV-B5</td>
<td>PAK-MEV5959A</td>
<td>M</td>
<td>36</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
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<tr>
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<td>PAK-MEV5926A</td>
<td>M</td>
<td>49</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B5</td>
<td>PAK-MEV3271A</td>
<td>M</td>
<td>4</td>
<td>Yes</td>
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<td>Yes</td>
</tr>
<tr>
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<td>PAK-MEV5959A</td>
<td>M</td>
<td>36</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B5</td>
<td>PAK-MEV2005B</td>
<td>M</td>
<td>23</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B5</td>
<td>PAK-MEV3903A</td>
<td>F</td>
<td>72</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B5</td>
<td>PAK-MEV5810B</td>
<td>M</td>
<td>36</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B6</td>
<td>PAK-MEV674B</td>
<td>F</td>
<td>18</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B6</td>
<td>PAK-MEV2475A</td>
<td>M</td>
<td>24</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B6</td>
<td>PAK-MEV4600A</td>
<td>F</td>
<td>18</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B6</td>
<td>PAK-MEV627A</td>
<td>M</td>
<td>18</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>CV-B6</td>
<td>PAK-MEV4553A</td>
<td>F</td>
<td>36</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
</tbody>
</table>

M, male; F, female; CV, coxsackievirus.
Fig. 1. A Phylogenetic tree based on partial VP1 coding sequence of 24 coxsackievirus B strains including CV-B1 to B6 and representative sequences of coxsackievirus serotypes retrieved from GenBank (accession numbers are included in the virus names). Neighbor-joining (NJ) method in MEGA 5.0 was used and evaluated with 1000 bootstrap pseudo-replicates. Bootstrap values greater than 50 are indicated at the respective nodes and the scale bar represents the evolutionary distance. Clusters are marked on the tree. The isolates from this study and prototype strains are represented by ‘●’ and ‘▲’ taxon markers, respectively.
nucleotide sequence divergence (up to 0·3% amino acid sequence difference). Whereas within CV-B1, CV-B3 and CV-B6 serotypes, nucleotide sequences varied by up to mean 12·2% (8·2–15·7%), 18·7% and 9·9% (0–13·4%), respectively (up to 0·4%, 9·1%, 2·7% amino acid sequence difference, respectively).

**VP1 deduced amino acid sequence analysis**

A comparison of the deduced amino acid sequences of the BC-loop regions of the study strains with their respective reference prototype strains was conducted to investigate the possible amino acid changes in the region. The BC-loop is generally located on the viral surface (VP1) and is associated with enteroviral infectious properties. Therefore, it has been suggested that unique sequence variations in the BC-loop might result in conformational changes that can alter antigenicity in these viruses. Figure 2 shows the alignment of 30 amino acids in the VP1 gene corresponding with amino acid 74–113 of the VP1 from all strains studied in this work. An analysis of the amino acid sequences of the BC-loop region (amino acid 84–93) of all strains showed that the strains of CV-B2 and CV-B4 had no amino acid change, while strains belonging to CV-B1, CV-B3, CV-B5 and CV-B6 had some changes in this region. All study strains from CV-B1 had serine (S) to threonine (T) substitution at position 86 (S86T), while CV-B3 strains had substitution of the amino acid, alanine (A) to serine (S) at position 87 (A87S). Similarly, the substitution of the 88th and 89th amino acids; aspartic acid (D) to asparagine (N) (D88N) for CV-B5 and lysine (K) to threonine (T) (K89T) for CV-B6 strains were found, respectively.

**DISCUSSION**

The Global Polio Eradication Initiative (GPEI) has succeeded in reducing the number of polio cases worldwide, but NPEVs (Echoviruses, CV and enterovirus 68, 70, 71, 77) have been reported frequently in association with AFP worldwide [18–21]. Similar to other NPEVs, the neurological complications of CV infections include AFP, encephalitis, meningoencephalitis and aseptic meningitis [22–24]. CV-B destroy motor neurons leading to the paralysis [25]. Though, mouse models for CV infection of the central nervous system are available, but much remains to be learned to understand the pathogenesis of this disease [26, 27]. Most research on CV has focused on infection of heart and pancreas. In Pakistan, the role of CVs in causing paralysis is not very clear due to lack of enterovirus surveillance system in the country. The CVs reported earlier in the country were identified in the context of AFP surveillance because poliovirus is the pathogen of interest and detailed testing for identification is not required for NPEVs. Additionally, to the best of our knowledge there is no study on CV-B serotypes in Pakistan that explored their molecular epidemiology, genetic diversity and possible link with AFP. Therefore, in the current study, all these questions were addressed.

The coxsackievirus B serotypes (CV-B1 to B6) have been identified for more than 50 years causing a wide spectrum of diseases and are repeatedly represented in listings of the most commonly isolated enteroviruses [28]. Their genetic analysis has improved our understanding of evolution, epidemiology and pathogenesis [29, 30]. In the current study, all six serotypes of CVs were identified with high degree of genetic diversity, which is particularly important for a country with no enterovirus surveillance system and a relatively short observation period, i.e. 1 year. Our study also identified distinct grouping pattern of CV-B1 and CV-B5. The phylogenetic analysis of CV-B5 and CV-B6 strains that had circulated in different provinces of the country showed that they can be divided into two clusters. The nucleotide divergences between

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</tr>
</thead>
<tbody>
<tr>
<td>CV-B1</td>
<td>36·4</td>
<td>42·9</td>
<td>31·3</td>
<td>29·7</td>
<td>32·1</td>
<td>36·3</td>
<td>13·1</td>
<td>9·1</td>
<td>6·5</td>
<td>12·1</td>
</tr>
<tr>
<td>CV-B2</td>
<td>13·1</td>
<td>9·1</td>
<td>6·5</td>
<td>12·1</td>
<td>17·0</td>
<td>14·7</td>
<td>20·7</td>
<td>11·1</td>
<td>15·8</td>
<td>18·2</td>
</tr>
<tr>
<td>CV-B3</td>
<td>39·8</td>
<td>35·9</td>
<td>39·8</td>
<td>35·9</td>
<td>39·8</td>
<td>39·8</td>
<td>33·7</td>
<td>33·7</td>
<td>33·7</td>
<td>33·7</td>
</tr>
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</table>

*Table 2. Summary of pairwise nucleotide and amino acid sequence comparisons among coxsackievirus B isolates*
study strains and the corresponding virus serotype strains reported in other countries were relatively high. In addition, the presence of multiple distinct genetic clusters is very important observation because the emergence of new genetic clusters of enterovirus in a particular area is often associated with large-scale outbreaks [31]. Furthermore, temporal and geographical comparisons of the study strains with previously characterized CV-B were limited by the availability of comparison sequences in the country. Till to date, there are only three CV-B4 nucleotide sequences in GenBank from Pakistan under accession numbers AF160018, AF160019 and AF160047. Therefore, the information on the range of CV-B serotypes circulating in Pakistan and their nucleotide sequences from the present study contribute significantly to the database of circulating enterovirus strains. These baseline data can be used in the future for determining temporal and geographical relationships of CV-B viruses, which is particularly useful for outbreak investigations and for detecting the emergence of new genetic clusters of enteroviruses.

The BC-loop is one of the regions associated with viral antigenicity, and substitutions resulting in conformational changes in this region are believed to play a role in host adaptation of picornaviruses [32]. In this study, the data analysis of deduced amino acid sequences in BC-loop region in the VP1 gene revealed four major substitutions at positions 86, 87, 88 and 89 in strains, on comparison with the
representative prototype strains from four different serotypes of coxsackieviruses (CV-B1, CV-B3, CV-B5 and CV-B6), respectively (Fig. 2). It is possible that these changes in the BC loop region might have played some role in conformational changes and associated with viral antigenicity. However, our findings do not confirm this hypothesis, but more detailed investigations by sequencing of all capsid proteins and conformational analyses may give the better answer.

This study found a different detection frequency pattern of CV-B serotypes. Overall CV-B5 (50%) strains were isolated more as compared with other serotypes of CV-B. These findings are in line with data reported earlier as CV-B5 consistently appears in the top five commonly identified enterovirus types in the USA and France [5, 33] and has exhibited the highest annual prevalence in a number of countries. Similarly, it has been found an emerging serotype that correlated with some outbreaks of meningitis and other diseases in the last 10 years [34]. In our study, the low frequency of other CV-B types might indicate low circulation of these strains in Pakistani population. It is important to note that the sensitivity of RD cells for CV-B is reported to be low in different reports, but the present study was able to isolate and detect at least one representative of each of the six CV-B serotypes using the RD cell. However, it is possible that some CV-B strains that cannot be propagated in RD and may have been undetected. Therefore, additional cell lines (e.g., MRC-5, Vero, RMK, BGMK and A-549) are recommended for isolation of such viruses in culture to elucidate their actual patterns of circulation.

There are many NPEVs that have been reported as the probable etiologic agents of paralytic illness clinically resembling poliomyelitis, including CV-B [35]. In earlier studies, five coxsackievirus B serotypes (CV-B2 to B6) have been repeatedly isolated from the stool of clinical paralytic poliomyelitis patients with slight to moderate paralysis and all these patients had no evidence of poliovirus infection [36]. CV-B2 and CV-B5 were isolated from stool of patients having spinal paralytic illness in which poliovirus was excluded [37] while CV-B2 to B4 were recovered from brain and spinal cord of patients with spinal paralytic disease [35]. In another study conducted in South Africa, Yui and Gledhil isolated CV-B from the stool of patients presenting with limb paralysis and hospitalized in Kalafong Hospital [38]. CV-B1, B2, B4 and B5 were also isolated from stool of AFP patients in Hong Kong during 2012 [39]. Similarly, in our study, the isolation of CV-B serotypes from AFP patients demonstrated their possible association with paralysis. Additionally, in all these patients, at the time of onset of disease, fever, asymmetrical paralysis and rapid progression were found (Table 1). Most patients had residual weakness on 60 days follow up investigation. Consequently, all these observations propose CV-B as the possible causative agent of paralysis. Furthermore, the different animal studies exhibit their neurotropic properties in experimentally infected animals that provide convincing evidence that these viruses have the capacity to produce neurological disease, which may clinically simulate poliomyelitis [27].

This data reveal the isolation of diverse CV-B from wide geographic regions of the country from the stool of paralytic patients. As we are approaching toward global eradication of poliomyelitis, consideration toward NPEV infections causing AFP is an equal cause of concern. Therefore, there is need to monitor such viruses by either the polio eradication program or the larger state health care system for better characterization and epidemiological investigation.

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DECLARATION OF INTEREST
None.

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