Heterogeneity of the fibre sequence in subgenus C adenoviruses

A K Adhikary, U Banik, J Numaga, E Suzuki, T Inada, N Okabe

Aims: To determine the nucleotide sequences of adenovirus (Ad) types 1 and 6 fibre genes; to clarify the molecular basis of the distinct haemagglutination properties of subgenus CAds and their phylogenetic relations.

Methods: Human Ad1 and Ad6 fibre genes were sequenced from genomic DNA by direct sequencing. Primer selection was based on alignment of the fibre gene of human Ad serotypes Ad2 and Ad5. Fibre based subgenus C specific polymerase chain reaction (PCR) was performed to check for deletions in field isolates of Ad6, as revealed by sequence analysis of the Ad6 prototype. A phylogenetic tree was constructed from the predicted amino acid (AA) sequences of the fibre gene of important Ads.

Results: Ad1 and Ad6 comprise 1746 and 1584 nucleotides, encoding 582 and 528 AA, respectively. Ad6 showed deletions in motifs 15–17 (51 AA) of the shaft when compared with Ad1, Ad2, and Ad5. Subgenus C specific PCR with both prototype and field isolates also showed deletions in Ad6. In the shaft and knob, AA homology was 58.82–72.91% and 68.89–74.59%, respectively. The tail was 100% conserved. Phylogenetically, Ad1 and Ad6, including Ad2 and Ad5, formed a subgenus specific cluster, like other serotypes.

Conclusions: The fibre gene (including the knob region) of subgenus C Ads is heterogeneous, providing the molecular basis for lack of crossreactivity in the haemagglutination inhibition test. This heterogeneity could be helpful in fibre based genotyping of subgenus C field isolates. Phylogeny might be useful for subgenus specific identification of important field strains.

In our study, the full length fibre genes of both Ad1 and Ad6 prototypes were sequenced for the first time with the following aims: (a) to find possible variations in the Ad1 and Ad6 fibre genes compared with the other serotypes of subgenus C, and to confirm any variations by fibre based polymerase chain reaction (PCR); (b) to correlate the variations of subgenus C Ads in the HAI test with heterogeneity of the fibre protein in the knob region; and (c) to clarify any evolutionary relations between Ad1 and Ad6 using a phylogenetic method.

MATERIALS AND METHODS

Viruses and DNA
Ad prototype strains Ad1 (adenovirus 71), Ad2 (adenovirus 6), Ad5 (adenovirus 75), and Ad6 (tonsil 99) and serotypes of other subgenus Ads were obtained from the American Type Culture Collection (ATCC; Rockville, Maryland, USA). Seventeen geographically and temporally diverse field isolates (Ad1:4, Ad2:4, Ad5:4, and Ad6:5) previously typed by the SN were used in our study.

Abbreviations: AA, amino acids; Ad, adenovirus; CAR, coxsackie virus and adenovirus receptor; HAI, haemagglutination inhibition; nt, nucleotides; PCR, polymerase chain reaction; SN, serum neutralisation
Extraction of viral DNA
The viruses were passaged in A549 cells and DNA was extracted from the infected cells by the method described previously. In short, when a greater than 75% cytopathic effect was seen, the cells were dislodged and pelleted by low speed centrifugation, washed three times with phosphate buffered saline, and resuspended in lysis buffer (10mM Tris/HCl, pH 7.4, 10mM EDTA, 1% sodium dodecyl sulfate). Next, the suspension was incubated with proteinase K (Sigma Chemical, St Louis, Missouri, USA). Cellular DNA was precipitated with 5M NaCl, and pelleted by centrifugation. After incubation with RNase A (Sigma Chemical) the supernatant was extracted twice in phenol/chloroform and precipitated in two volumes of absolute ethanol. After drying, DNA was suspended in 50 μl of double distilled water.

DNA sequencing
The genomic DNA was used as a template for DNA sequencing. The initial selection of primers was based on alignment of the Ad2 and Ad5 fibre genes (GenBank accession numbers j01907 (Ad2) and m18369 (Ad5)), and subsequently, because of the heterogeneous nucleotide sequences, the primer walking method was used. DNA sequencing was performed on the sense and antisense strands with overlapping primers. The cycle sequence reaction was carried out with an ABI prism big dye terminator cycle sequencing ready reaction kit (Applied Biosystems, Foster City, California, USA). The sequences were read by a Genetic analyser model 310 (Applied Biosystems). The nucleotide sequences of the fibre gene were translated into AA sequences, and compared with the published AA sequences of Ad2 and Ad5 using the GENETYX-MAC program (Software Development Co, Tokyo, Japan).

Nucleotide sequence accession numbers
The nucleotide sequence data reported in this paper will appear in the DDBJ/EMBL/GenBank nucleotide sequence database with the accession numbers ab125750 (Ad1) and ab125751 (Ad6).

Ad subgenus C specific primers
Subgenus C specific PCR was performed with prototypes and field isolates to check the deletion in the shaft region of Ad6 field isolates noted by sequence analysis of the Ad6 prototypes. The design of the subgenus C specific primers (AdnCF, 5'-TGC TTG CGC THA AAA TGG GC-3' (654–674) and AdnCR, 5'-CGA TTC TTT ATT CYT GGG CAA TGT-3' (2228–2205)) was based on the alignment of the fibre gene sequences of Ad1 and Ad6 conducted in our laboratory (GenBank accession numbers Ad1 (ab125750), and Ad6 (ab125751)) and Ad2 and Ad5 obtained from GenBank (accession numbers Ad2 (j01917) and Ad5 (m18369)). The nucleotide positions of the primers refer to the fibre sequence of Ad2.

DNA amplification and detection
PCR amplification was carried out in a 50 μl reaction mixture containing 1 μl aliquots of DNA, 5 μl of 10× concentrated buffer, 0.5 μM of each of the primers, 200 μM of each deoxynucleoside triphosphate (dNTP), and 1.25 U of Taq polymerase (Boehringer-Mannheim, Mannheim, Germany). Sterile distilled water was used as a negative control. For field isolates, Ad2 DNA was used as a positive control. The assay was performed in a thermal cycler (model 9600-R; Perkin Elmer, Foster City, California, USA). Thermal cycling consisted of preliminary denaturation for three minutes at 94°C, followed by 35 cycles of denaturation at 94°C for one minute, annealing at 47°C for one minute, and extension at 72°C for two minutes, with a final extension at 72°C for seven minutes. The reaction products (5 μl aliquots) were electrophoresed on a 1.2% horizontal agarose gel. After staining with ethidium bromide the gel was photographed with a CCD camera. The specificity of the test was evaluated with pooled DNA from the A, B, D, E, and F subgenera of Ad and non-adenoviral DNA.

Phylogenetic analysis
A phylogenetic tree was constructed according to the full length predicted AA sequences of the fibre genes obtained from the GenBank. Sequence alignment was performed with Clustal X (http://www-igbmc.u-strasbg.fr/BioInfo/) with parameters provided in Clustal X. Evolutionary distances were estimated by Kimura’s two parameter method. A phylogenetic tree with 1000 bootstrap replicates was generated by the neighbour joining method with Clustal W and plotted by Tree View (http://taxonomy.zoology.gla.ac.uk/rod/treeview.html).

RESULTS
Sequence analysis
The Ad1 and Ad6 fibre genes were 1746 and 1584 nucleotides (nt) long (table 1), corresponding to 582 and 528 AA, respectively. The predicted AA sequences were consistent with the structural domains described for Ad2, which possesses three distinct regions: an N-terminal tail (44 AA); a central shaft of 357 residues consisting of 22 pseudorepeats, usually 15 residues; and a C-terminal knob of 181 residues. The fibre tail of Ad1 and Ad6 was 132 nt long, corresponding to 44 AA, and showed 100% conservation among the members of subgenus C. However, at the nucleotide level the homology varied from 89.39% to 93.18%. The two conserved sequences—2-KR&R (where λ is A in all the subgenus C serotypes) and FNPVYPYD—are present in subgenus C, similar to the other Ad subgenera (fig 1).

The shaft region of Ad1 and Ad6 displayed a 15 AA basic structural unit (motif), containing hydrophobic glycine (G) or proline (P) residues at position 8 of the β sheet/β turn model (fig 1). Ad1 comprises 357 polypeptides and displayed 22 repeats of the 15 AA motifs, similar to Ad2 and Ad5. The shaft of Ad6 was relatively shorter, with 306 AA (19 motifs) as a result of the deletion of motifs 15 to 17 (fig 2). The third motifs were 19 AA long except for in Ad6, which showed deletion of a residue (T) in position 8. In the shaft region, AA homology among the serotypes varied from 58.82% to 72.91%. The Ad1 and Ad6 fibre knobs were 543 and 534 nt long, corresponding to 181 and 178 AA, respectively. Their AA homology varied between 68.89% and 74.59% (table 2). Ad1 and Ad6 showed an overall homology of 71.95% at the DNA level and 70.1% at the AA level. The overall AA homology among the members of subgenus C varied from 65.06% to 75.47%, and is considered to be heterogeneous.

Table 1  Length of the fibre genes (number of nucleotides) and proteins (number of amino acids) among the members of subgenus C adenoviruses

<table>
<thead>
<tr>
<th></th>
<th>Tail</th>
<th>Shaft</th>
<th>Knob</th>
<th>Overall</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ad1</strong></td>
<td>DNA</td>
<td>132</td>
<td>1071</td>
<td>543</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>44</td>
<td>357</td>
<td>181</td>
</tr>
<tr>
<td><strong>Ad2</strong></td>
<td>DNA</td>
<td>132</td>
<td>1071</td>
<td>543</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>44</td>
<td>357</td>
<td>181</td>
</tr>
<tr>
<td><strong>Ad5</strong></td>
<td>DNA</td>
<td>132</td>
<td>1071</td>
<td>540</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>44</td>
<td>357</td>
<td>180</td>
</tr>
<tr>
<td><strong>Ad6</strong></td>
<td>DNA</td>
<td>132</td>
<td>918</td>
<td>534</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>44</td>
<td>306</td>
<td>178</td>
</tr>
</tbody>
</table>

Ad, adenovirus.
Polymerase chain reaction

The primer pair AdnCF/AdnCR amplified approximately 1575 bp, 1575 bp, and 1572 bp products from Ad1, Ad2, Ad5 prototypes, respectively, whereas the Ad6 prototype yielded a 1413 bp product (fig 2A). Seventeen field isolates (table 3) also yielded clearly visible products of the expected length (fig 2B). Ad6 could be distinguished from the other members of subgenus C by the length of the amplicon (1413 bp).
PCR using DNA from other subgenera of adenovirus and non-adenoviral DNA yielded no amplified products, indicating a high specificity of the test.

**Phylogenetic analysis**

Ad1 and Ad6 were included in a cluster formed by subgenus C adenoviruses. Ad6 was found to be more closely related to Ad1 than either Ad2 or Ad5. Other serotypes included in phylogeny also formed subgenus specific clusters. However, the cluster formed by subgenus F was divided into two subclusters because of the presence of two different fibres (long fibre and short fibres; fig 3).

**DISCUSSION**

Subgenus C Ads differ from other subgenera in that they have a highly variable genomic organisation, they are in widespread use as a vector for gene delivery, and they are responsible for approximately 60% of human infections caused by the different Ad serotypes. Our study revealed two important findings about the subgenus C Ad fibre protein. First, the structure of the knob is heterogeneous, which could explain the serological characteristics of subgenus C Ads and, second, the fibre shaft of Ad6 is relatively short, and this may be related to its lower rate of infection.

Subgenus C Ads are unique because of the lack of crossreactivity in both the SN and HAI tests. The HAI test in a large number of geographically diverse isolates showed that the fibre epitopes are relatively conserved, although the exact location and size of these epitopes are yet to be determined.

---

**Table 2**

<table>
<thead>
<tr>
<th>Fibre</th>
<th>Ad1/2</th>
<th>Ad1/5</th>
<th>Ad1/6</th>
<th>Ad6/2</th>
<th>Ad6/5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tail</td>
<td>DNA</td>
<td>90.15</td>
<td>90.91</td>
<td>93.18</td>
<td>90.91</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>100.0</td>
<td>100.0</td>
<td>100.0</td>
<td>100.0</td>
</tr>
<tr>
<td>Shaft</td>
<td>DNA</td>
<td>72.71</td>
<td>74.91</td>
<td>69.16</td>
<td>64.52</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>70.56</td>
<td>72.91</td>
<td>64.71</td>
<td>62.18</td>
</tr>
<tr>
<td>Knob</td>
<td>DNA</td>
<td>68.56</td>
<td>72.56</td>
<td>72.24</td>
<td>73.85</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>70.88</td>
<td>74.59</td>
<td>73.48</td>
<td>74.03</td>
</tr>
<tr>
<td>Overall</td>
<td>DNA</td>
<td>72.73</td>
<td>75.39</td>
<td>71.95</td>
<td>69.42</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>72.87</td>
<td>75.47</td>
<td>70.10</td>
<td>68.73</td>
</tr>
</tbody>
</table>

DNA and amino acid homologies were obtained using GENETYX-MAC software (Software Development Co, Tokyo, Japan). Ad, adenovirus.
and Ad37 show 97.7% and 95.5% homology in the knob region, respectively, and show major crossreactivity in the HAI test. Therefore, higher homology (> 80%) in the knob region is thought to be the basis of serological crossreactivity as a result of epitope sharing.19 20 22 Although not an absolute rule, lower homology in the knob region may imply the presence of distinct epitopes, as demonstrated by the absence of major crossreactivity in HAI. Thus, subgenus C Ads have no major crossreactivity in HAI with other subgenera or among themselves probably because they have low homology in the fibre knob region, between 68.89% and 74.59.

"Adenovirus 1 (Ad1) is more closely related to Ad6 than to Ad2 or Ad5"

To date, studies with Ad2 and Ad5 have shown that the subgenus C fibre knob mediates binding to the cowpox virus and Ad receptor (CAR), and is thus a major determinant of viral tropism.21 Mounting evidence suggests that at least two factors determine viral infectivity and tropism, namely: (a) the fibre knob–primary receptor interaction and (b) the length and flexibility of the fibre shaft.16 17 The length of the fibre shaft is variable because of deletion or insertion of certain numbers of repeats. The formation of a recombinant fibre shaft with decreasing length and increasing rigidity significantly reduced infectivity and attachment of Ad5.18 The major differences between the Ad6 fibre shaft and that of the other subgenus C Ads are the deletion of the 15–17th motifs (51 residues) of the shaft and a single residue (T) deletion in the eighth position of motif 3. The deletion was also detected in PCR products of the field isolates of Ad6. It may be possible that deletion of 51 AA residues in the shaft decreases its flexibility and reduces the mobility of the knob.22 The third motif of subgenus C is 19 AA long, except in Ad6. In all Ads, the third motif carries a hydrophilic residue in the second hydrophobic position of the β strand (position 15 in subgenus C). The fibre bends above the penton base at a position corresponding to this third motif of the shaft, which is also a flexing point to allow fibre movement. In Ad6, the third motif is 18 AA long (rather than 19 AA as in other subgenus C Ads), as a result of the deletion of a single residue (T) at position 8, and this may also slightly decrease the flexibility of the fibre shaft.23 Both these factors present in Ad6 might act together to decrease its efficiency of binding to the cellular receptor. Further experiments with recombinant Ad6 fibre proteins might be useful to investigate the relation between the length of the Ad6 fibre shaft and the rate of infection. It is also important to investigate whether Ad6 uses CAR or another cellular receptor.

In the phylogram, Ad1 and Ad6, together with Ad2 and Ad5, formed a subgenus C specific cluster, indicating that these Ads have the same ancestral origin. Ad1 is more closely related to Ad6 than to Ad2 or Ad5. Subgenus specific clusters were also formed by other serotypes as a result of high homology (60–80%) among the members of each subgenus.26 Subgenus specific clusters of the fibre gene that included important Ads could be useful in subgrouping and understanding the evolution of Ads.

Sequence heterogeneity in the fibre knob provides the molecular basis of the distinct serological properties of subgenus C Ads in the HAI test. The inclusion of serotypes Ad1, Ad2, Ad5, and Ad6 in a single cluster defines the evolutionary relations between the subgenus C Ads. Furthermore, the sequence data of Ad1 and Ad6 reported in our study will be useful in the identification of subgenus C field strains, either by fibre based type specific PCR or by comparison of sequences.

ACKNOWLEDGEMENTS

Supported by grant-in-aid for Scientific Research from the Japan Society for the Promotion of Science (JSPS), Japan.

Authors’ affiliations

A K Adhikary, U Banik, T Inada, N Okabe, Infectious Disease Surveillance Centre, National Institute of Infectious Diseases, 1-23-1 Toyama, Shinjuku-ku, Tokyo 162-8640

J Numaga, Department of Ophthalmology, Graduate School of Medicine, The University of Tokyo, 7-3-1 Hongo, Bunkyo-ku, Tokyo 113-0033, Japan

E Suzuki, Department of Developmental Medical Sciences, Graduate School of Medicine, The University of Tokyo, 7-3-1 Hongo, Bunkyo-ku, Tokyo 113-0033, Japan

REFERENCES


Heterogeneous fibre of subgenus C adenoviruses

3. Adam E, Nasz I, Lengyel A. Antigenic homogeneity among the members of the adenovirus hexon types of subgenus C. Arch Virol 1995;140:1297–301.