Genomic and antigenic characterization of the newly emerging Chinese duck egg-drop syndrome flavivirus: genomic comparison with Tembusu and Sitiawan viruses

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Duck egg-drop syndrome virus (DEDSV) is a newly emerging pathogenic flavivirus causing avian diseases in China. The infection occurs in laying ducks characterized by a severe drop in egg production with a fatality rate of 5–15%. The virus was found to be most closely related to Tembusu virus (TMUV), an isolate from mosquitoes in South-east Asia. Here, we have sequenced and characterized the full-length genomes of seven DEDSV strains, including the 5' and 3'-non-coding regions (NCRs). We also report for the first time the ORF sequences of TMUV and Sitiawan virus (STWV), another closely related flavivirus isolated from diseased chickens. We analysed the phylogenetic and antigenic relationships of DEDSV in relation to the Asian viruses TMUV and STWV, and other representative flaviviruses. Our results confirm the close relationship between DEDSV and TMUV/STWV and we discuss their probable evolutionary origins. We have also characterized the cleavage sites, potential glycosylation sites and unique motifs/modules of these viruses. Additionally, conserved sequences in both 5' and 3'-NCRs were identified and the predicted secondary structures of the terminal sequences were studied. Antigenic cross-reactivity comparisons of DEDSV with related pathogenic flaviviruses identified a surprisingly close relationship with dengue virus (DENV) and raised the question of whether or not DEDSV may have a potential infectious threat to man. Importantly, DEDSV can be efficiently recognized by a broadly cross-reactive flavivirus mAb, 2A10G6, derived against DENV. The significance of these studies is discussed in the context of the emergence, evolution, epidemiology, antigenicity and pathogenicity of the newly emergent DEDSV.

These authors contributed equally to this work/paper.

The GenBank/EMBL/DDBJ accession numbers for the genome sequences of seven strains of DEDSV are JQ920420–JQ920426. The GenBank/EMBL/DDBJ accession numbers for the ORF sequences of TMUV and STWV are JX477685 and JX477686, respectively.

Supplementary material is available with the online version of this paper.
INTRODUCTION

Since April 2010, a severe duck disease has spread throughout the major duck-producing regions in China (Hu et al., 2011; Su et al., 2011). The infected ducks display anorexia accompanied by a heavy drop in egg production. As the disease progresses, some ducks exhibit an uncoordinated gait and loss of balance. Total mortality can range from 5 to 15%. Consequently, this disease is a major concern for the Chinese poultry industry (Hu et al., 2011). The aetiological agent was previously identified as a newly emerged flavivirus and designated Baiyangdian virus (BYD virus) (Su et al., 2011). However, since we now recognize that a plant virus, barley yellow dwarf virus, already has the designation BYDV, we propose the alternative name duck egg-drop syndrome virus (DEDSV) for this important avian pathogen. Analysis of partial sequences of the envelope (E) and non-structural (NS) NS5 gene, demonstrated its close relationship with the Ntaya virus (NTAV) group in the genus Flavivirus, family Flaviviridae. Additionally, DEDSV was shown to be particularly closely related to Tembusu virus (TMUV) and Sitiawan virus (STWV) (Hu et al., 2011; Su et al., 2011). Other research groups have performed similar investigations and defined their newly isolated viruses as strains of TMUV or Tembusu-like virus (Cao et al., 2011; Yan et al., 2011).

Members of the genus Flavivirus in the family Flaviviridae are positive-sense, ssRNA viruses that infect humans and other vertebrates (Gould & Solomon, 2008). By taking into account the vector, vertebrate hosts, antigenic phylogenetic and biogeographical characteristics, the flaviviruses were divided into three groups: mosquito-borne viruses, tick-borne viruses and viruses with no known arthropod vector (Porterfield, 1980). For the mosquito-borne flaviviruses, they have been further subdivided into seven subgroups including the NTAV group (refer to the ICTV database – ICTV 2010, http://talk.ictvonline.org/files/ictv_documents/m/msl/1231.aspx). The NTAV group currently comprises the following viruses: NTAV, Bagaza virus (BAGV), TMUV, Ilheus virus (ILHV) and Israel turkey meningoencephalomyelitis virus (ITV). Two other viruses, namely STWV and Rocío virus (ROCV), are considered to be subtypes of TMUV and ILHV, respectively. The NTAV group causes disease in several different hosts; ITV infects turkeys producing non-suppurative meningoencephalitis and myocardial necrosis (Guy & Malkinson, 2008). ILHV is not associated with epidemic disease but has been isolated sporadically from patients in Brazil, Trinidad, Panama, Argentina, Colombia and Ecuador, with acute febrile illness presenting as headache, myalgia and malaise (Shope, 2003). ROCV is the only member known to have caused at least two human epidemics of severe encephalitis between 1973 and 1990 in the south-eastern region of Brazil (Shope, 2003). STWV infects chickens causing encephalitis and growth retardation (Kono et al., 2000). BAGV causes both human illness (Gould & Solomon, 2008) and avian infections in Africa and Spain (Agüero et al., 2011; Digoutte, 1978; Traore-Lamizana et al., 1994). TMUV and NTAV were isolated from mosquitoes. No known diseases in either human or animals have been recorded (Weissenböck et al., 2010). To date, whole genomic sequences are available for only three members of the NTAV group, namely BAGV, ILHV and ROCV (Kuno & Chang, 2005, 2007; Medeiros et al., 2007). Thus, their evolutionary characteristics and genetic relationships are currently inadequately defined.

The flavivirus genomic RNA contains a unique ORF, flanked by a type 1 capped 5’-terminal non-coding region (NCR) and a 3’-terminal NCR that, together, form specific secondary stem–loop structures required for RNA translation, replication and/or expression of biological traits most likely including pathogenetic determinants (Lindenbach & Rice, 2003; Rice et al., 1985). The ORF encodes the viral proteins as a large polyprotein precursor that is co- and post-translationally processed by cellular and viral proteases into at least 10 separate products. The N-terminal region encodes three structural proteins: capsid (C), pre-membrane (prM) (which is post-translationally cleaved to produce pr and M protein) and E, followed by the non-structural proteins NS1 (soluble complement-fixing antigen), NS2A, NS2B, NS3 (serine protease/RNA helicase), NS4A, NS4B and NS5 (RNA-dependent RNA polymerase/methyltransferase) (Chambers et al., 1990).

In this study, we have characterized the antigenicity of DEDSV in terms of its relationship with other mosquito-borne flaviviruses and sequenced the full-length genomes of seven strains of DEDSV (including their 5’- and 3’-NCR) and the ORF sequences of both TMUV and STWV. The close relationship between DEDSV and TMUV/STWV was confirmed. In detailed antigenic studies with representative human pathogenic flaviviruses, we demonstrate significant antigenic cross-reactivity with the implication that these DEDSV shared epitopes may indicate the potential for infection of human beings by DEDSV. At the same time, we also found that the broadly neutralizing mAb 2A10G6, which was originally raised against dengue virus (DENV) (Deng et al., 2011), binds to a very high titre with DEDSV. We therefore propose DEDSV should be considered to be a new member of the NTAV group of flaviviruses, forming a subgroup together with TMUV and STWV.

RESULTS

Full-genome sequencing and genome structures of DEDSV, TMUV and STWV

We have isolated seven strains of DEDSV (Table 1), all of which are pathogenic. The laying ducks infected with isolated DEDSV-JXSP showed analogous clinical manifestations, like a sudden decline of food uptake (Fig. S1a, available in JGV Online), severe egg drop (Fig. S1b) and haemorrhage of spleen and ovaries (Fig. S2), with those...
infected with DEDSV-byd1 we have performed previously (Su et al., 2011). Seven strains of DEDSV and one strain each of TMUV and STWV were sequenced (Table 1). The whole genome of DEDSV, including the 5'- and 3'-NCRs, was 10990 nt in length with an ORF of 10278 nt flanked by 5'- and 3'-NCRs of 94 and 618 nt, respectively. The ORF encoded a polyprotein of 3425 aa. The ORFs for both TMUV and STWV were also 10278 nt, encoding 3425 aa. The 10 recognized flaviviral proteins (defined above) were identified. The lengths of these proteins and the specific positions of relevant genes are shown in Fig. 1(a).

The similarity between these DEDSV isolates was at least 99.7 %, confirming they are all strains of the same virus. As shown in Fig. 1(b), a phylogenetic tree was constructed based on the ORF nucleotide sequences. The DEDSV-JX2 lineage roots to the DEDSV major cluster. Notably, DEDSV-goose and DEDSV-pigeon share a clade with DEDSV-JS and DEDSV-JXSP. Moreover, the duck ‘Tembusu virus’ strains YY5, ZJ-407 and GH-2 recently identified by another group (Yun et al., 2012), all of which were isolated from the same province in China, segregated into one cluster. All of the other strains including duck flavivirus strain TA (Liu et al., 2012) formed another cluster and they were isolated from neighbouring provinces in China (Table 1).

TMUV and STWV showed 88.8 and 87.4 % nt sequence identity and 97 and 96.5 % aa sequence identity with DEDSV, respectively. The close relationship between TMUV and DEDSV strongly implies that DEDSV, like STWV, can be designated a subtype of TMUV.

**Comparison with other flaviviruses**

In order to study the phylogenetic relationship between DEDSV/TMUV/STWV and other members of the genus

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**Table 1. Viruses studied in this report**

<table>
<thead>
<tr>
<th>Name or ORF available</th>
<th>Source of isolation</th>
<th>Location of isolation</th>
<th>Sequence available</th>
</tr>
</thead>
<tbody>
<tr>
<td>DEDSV-byd1</td>
<td>Egg-laying duck</td>
<td>Hebei Province, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>DEDSV-JXSP</td>
<td>Meat-type duck</td>
<td>Beijing Autonomous City, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>DEDSV-Duan</td>
<td>Egg-laying duck</td>
<td>Beijing Autonomous City, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>DEDSV-JX2</td>
<td>Egg-laying duck</td>
<td>Jiangxi Province, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>DEDSV-JS</td>
<td>Egg-laying duck</td>
<td>Jiangsu Province, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>DEDSV-goose</td>
<td>Goose</td>
<td>Beijing Autonomous City, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>DEDSV-pigeon</td>
<td>Pigeon</td>
<td>Beijing Autonomous City, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>TMUV MM1775</td>
<td>Culex tritaeniorhynchus</td>
<td>Kuala Lumpur, Malaysia</td>
<td>ORF</td>
</tr>
<tr>
<td>STWV</td>
<td>Broiler chicken</td>
<td>Perak state, Malaysia</td>
<td>ORF</td>
</tr>
<tr>
<td>TMUV ZJ-407</td>
<td>Breeder duck</td>
<td>Zhejiang Province, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>TMUV YY5</td>
<td>Shaoxing duck</td>
<td>Zhejiang Province, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>TMUV GH-2</td>
<td>Goose</td>
<td>Zhejiang Province, PR China</td>
<td>Full sequence</td>
</tr>
<tr>
<td>Duck flavivirus TA</td>
<td>Duck</td>
<td>Shandong Province, PR China</td>
<td>Full sequence</td>
</tr>
</tbody>
</table>

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**Fig. 1.** DEDSV genome structures and the phylogenetic analysis of seven strains of DEDSV. (a) Genome structure, RNA elements and polyprotein constitutions. Structural proteins are highlighted in grey. (b) Bayesian phylogeny of the ORF nucleotide dataset of seven strains of DEDSV, TMUV strain MM1775 and STWV were used as outgroups. Posterior probabilities are shown. The tree is midpoint-rooted. Bar, 0.01 substitutions per site.
GenBank (NCBI accession no. NC_015843.1). This strain isolated from a DEDS outbreak in China was deposited in genome sequence of DEDSV-like virus (strain TMUV-JS804) (Gaunt et al., 1980). On the basis of their phylogenetic, ecological and virus–vector relationships, the mosquito-borne flaviviruses were subdivided into *Culex* spp.-associated and *Stegomyia* spp.-associated groups (Gaunt et al., 2001; Porterfield, 1980). The number of potential N-linked glycosylation sites (N-LGlyS) in the prM, E and NS1 proteins of DEDSV-byd1, TMUV and STWV were 2–2–3, 2–2–2 and 2–1–2, respectively, whereas BAGV has 1–1–3. No glycosylation site was predicted to exist in the hydrophobic domains in the viruses studied. Disulfide bonds formed by cysteine residues are important for the stabilization and conformation of the proteins. For prM, the conserved six cysteine residues clustered in the pr domain. This is similar to other flaviviruses. The pattern of 12 cysteine residues found in all mosquito-borne flaviviruses in the E and NS1 genes is also conserved in the three flaviviruses.

**Functional and structural motifs/modules**

Many motifs/modules conserved in the flaviviruses (Gao et al., 1993) are also found in the DEDSV-byd1, TMUV and STWV. In each E monomer of flaviviruses, three structural domains are discernible. Domain II is involved in dimerization (Modis et al., 2004). It contains a highly conserved fusion peptide at its tip, i.e. aa residues 98–110, which binds to the membrane of the host cell during fusion (Bakonyi et al., 2004). This fusion loop is also shared by DEDSV-byd1, TMUV and STWV. Domain III has been implicated in receptor binding. As shown in Table 3, a putative integrin-binding motif (a tripeptide sequence of aa 388–390) is encoded by RGD in JEV, Usutu virus, Murray Valley encephalitis virus and Alfuy virus (Bakonyi et al., 2004; Hurrelbrink et al., 1999; May et al., 2006; Sumiyoshi et al., 1987), RGE in West Nile virus (WNV) (Castle et al., 1985) and Kunjin virus (Coia et al., 1988) and RGP in St. Louis encephalitis virus (SLEV) (Ciotà et al., 2007). However, the corresponding sequence is TGE in BAGV (Kuno & Chang, 2007), TGP in ROCV (Medeiros et al., 2007) and QEN in ILHV (Kuno & Chang, 2005). We identified the sequence as SGK in DEDSV-byd1, TMUV and STWV. In the N-terminal third of the NS3, the catalytic triad (H47, D75 and S135) and the proposed substrate-binding pocket of the trypsin-like serine protease (Gly133, Ser135, Gly136, Gly148, Leu149 and Gly153) (Valle & Falgout, 1998) were also found to be conserved in the three viruses, as well as the RNA helicase motif D285-E286-A287 in the C-terminal third of the protein (Gorbalenya et al., 1989). Similar to other flaviviruses, a homologous G667-D668-D669 motif common to RNA-dependent RNA polymerase (Rice et al., 1985) was also found in all DEDSV-byd1, TMUV and STWV in the C-terminal third of the NS5 protein.

**Secondary structures of the NCRs**

The MFOLD program was used to predict the secondary structure (SS) of the viral terminal region of the DEDSV
Fig. 2. Bayesian phylogeny of representative flaviviruses with DEDSV, TMUV and STWV. Posterior probabilities are shown for all the clades. (a) Phylogenetic tree was based on the ORF sequence dataset. Those flavivirus sequences that are published for the first time in the current study, namely DEDSV strain byd1, TMUV strain MM1775 and STWV, are highlighted in bold. All horizontal branch lengths are drawn to scale; bar, 0.3 substitutions per site. (b) Phylogenetic tree was based on the complete genome sequences. DEDSV is highlighted in grey. All horizontal branch lengths are drawn to scale; bar, 0.5 substitutions per site. NKV, No known vector flaviviruses; SS, secondary loss NKV flaviviruses. See Table S1 for virus abbreviations and GenBank accession numbers.
Table 2. Comparison of the genes or genomic regions of the four flaviviruses in the NTAV group

Numbers in parentheses represent the percentage amino acid or nucleotide identity with DEDSV, respectively.

<table>
<thead>
<tr>
<th>Gene or genomic region</th>
<th>DEDSV</th>
<th>BAGV</th>
<th>TMUV</th>
<th>STWV</th>
</tr>
</thead>
<tbody>
<tr>
<td>5'-NCR</td>
<td>94 nt</td>
<td>94 nt (74.7)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>C</td>
<td>120 aa</td>
<td>122 aa (67.2)</td>
<td>120 aa (95.8)</td>
<td>120 aa (93.3)</td>
</tr>
<tr>
<td>prM</td>
<td>167 aa</td>
<td>167 aa (81.4)</td>
<td>167 aa (97.6)</td>
<td>167 aa (95.2)</td>
</tr>
<tr>
<td>E</td>
<td>501 aa</td>
<td>501 aa (83.6)</td>
<td>501 aa (97.0)</td>
<td>501 aa (96.8)</td>
</tr>
<tr>
<td>NS1</td>
<td>352 aa</td>
<td>352 aa (78.4)</td>
<td>352 aa (94.0)</td>
<td>352 aa (93.2)</td>
</tr>
<tr>
<td>NS2A</td>
<td>227 aa</td>
<td>226 aa (63.9)</td>
<td>227 aa (92.5)</td>
<td>227 aa (92.1)</td>
</tr>
<tr>
<td>NS2B</td>
<td>131 aa</td>
<td>132 aa (95.6)</td>
<td>131 aa (97.0)</td>
<td>131 aa (97.0)</td>
</tr>
<tr>
<td>NS3</td>
<td>619 aa</td>
<td>619 aa (85.6)</td>
<td>619 aa (98.7)</td>
<td>619 aa (98.2)</td>
</tr>
<tr>
<td>NS4A</td>
<td>126 aa</td>
<td>126 aa (82.5)</td>
<td>126 aa (96.8)</td>
<td>126 aa (95.2)</td>
</tr>
<tr>
<td>2K</td>
<td>23 aa</td>
<td>23 aa (87.0)</td>
<td>23 aa (95.6)</td>
<td>23 aa (95.6)</td>
</tr>
<tr>
<td>NS4B</td>
<td>254 aa</td>
<td>253 aa (78.7)</td>
<td>254 aa (98.0)</td>
<td>254 aa (98.0)</td>
</tr>
<tr>
<td>NS5</td>
<td>905 aa</td>
<td>905 aa (85.9)</td>
<td>905 aa (97.8)</td>
<td>905 aa (98.0)</td>
</tr>
<tr>
<td>Polypeptide</td>
<td>3425 aa</td>
<td>3426 aa (81.3)</td>
<td>3425 aa (97.0)</td>
<td>3425 aa (96.5)</td>
</tr>
<tr>
<td>ORF</td>
<td>10278 nt</td>
<td>10281 nt (71.8)</td>
<td>10278 nt (88.8)</td>
<td>10278 nt (87.4)</td>
</tr>
<tr>
<td>3'-NCR</td>
<td>618 nt</td>
<td>566 nt (64.5)</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Total length of genome</td>
<td>10990 nt</td>
<td>10941 nt (71.4)</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

Genomic RNA. First, we predicted the SSs of 5'-terminal sequence and 3'-terminal sequence separately (Fig. 3a). Stem–loop A (SLA), stem–loop B (SLB) and the capsid hairpin (cHP) were as defined for other flaviviruses (Gritsun et al., 2006). Both in vivo and in vitro data supported the idea that SLA, present in the 5'-UTR, functions as the promoter element for the viral polymerase (Villordo & Gamarnik, 2009). SLA is essential for the viral NS5 protein binding to the viral RNA, while the SLB and cHP are dispensable (Dong et al., 2007). The 3'-terminal sequence has a conserved long stem–loop (3'-SL) structure, which is predicted to be conserved in all flavivirus genomes. The structure has a conserved pentanucleotide CACAG in the anthropod-borne flaviviruses (Brinton et al., 1986; Hahn et al., 1987; Wengler & Castle, 1986), which is critically important for viral RNA replication, it is located in the bulge at the top of the 3'-SL in DEDSV. The conserved sequence (CS)1 is located just upstream of the 3'-SL. The 5'-CS is 12 nt long in the C gene. As already known, cyclization of the viral genome is crucial for viral genome replication. The predicted SS of the cyclized viral RNA is shown in Fig. 3(b). As previously reported in other flaviviruses, the 5'-CS and 3'-CS are involved in genome cyclization and form double-stranded structures. The complementarity rather than the sequence per se is essential for RNA synthesis. Regarding the upstream AUG region (UAR) found conserved in other mosquito-borne flaviviruses, a functionally homologous sequence for DEDSV could not be found, this is also the case for BAGV (Kuno & Chang, 2007).

CS sequences in 3'-NCR play a role in viral RNA replication (Alvarez et al., 2005). The organization, names of the structures and/or domains used in this study follow the nomenclatures established earlier (Chambers et al., 1990; Hahn et al., 1987; Proutski et al., 1997). Besides the CS1, some other conserved sequences in the 3'-NCR were also identified (Table 4). Five CSs were found with the pattern ImRCS3-CS3-RCS2-CS2-CS1, where the ImCS3 describes the imperfectness of the conserved sequence differing from the corresponding consensus sequences in three or more bases. However, the CS organization of BAGV is CS1-CS2-CS3-RCS3 (Kuno & Chang, 2007). The CS pattern of DEDSV is different from BAGV but is the same as the JEV group.

Antigenic cross-reactivity with representative flaviviruses

Indirect ELISAs were carried out to investigate the cross-reactivity of immune sera from DEDSV-infected ducks, with other representative flaviviruses. Except for TMUV and BAGV, all of the viruses were standardized at 1000 p.f.u. ml⁻¹ in the ELISA experiments. After coating plates with these viruses, the antiserum against DEDSV-byd1 was tested at a 1:200 dilution. The duck sera exhibited high levels of cross-reactivity with representative flaviviruses although the level of cross-reactivity was lower than the homologous test with DEDSV, as the positive control antigen (Fig. 4a). Whilst TMUV and BAGV appear to show lower readings in the ELISA experiments. After coating plates with these viruses, the antiserum against DEDSV-byd1 was tested at a 1:200 dilution. The duck sera exhibited high levels of cross-reactivity with representative flaviviruses although the level of cross-reactivity was lower than the homologous test with DEDSV, as the positive control antigen (Fig. 4a).
observation that the broadly cross-reactive DENV-2-derived mAb 2A10G6, which recognizes an epitope within the \textsuperscript{98}DRXW\textsuperscript{103} motif localized in the fusion loop of the E protein (Deng et al., 2011), also exhibited high binding activity to DEDSV-byd1. Compared with DENV-2, DEDSV-byd1 shows similar reactivity with the mAb. The comparative sequence alignment data confirmed that the conserved motif does exist in DEDSV-byd1 (Fig. 4g).

Neutralization profiles of 2A10G6 against representative flaviviruses

To investigate the neutralization potential of mAb 2A10G6 to the DEDSV, the neutralization activities of 2A10G6 against various flaviviruses including DEDSV were assessed using a standard plaque reduction neutralization assay. The results (Fig. 4i) demonstrated that 100 µg 2A10G6 ml\textsuperscript{-1} potently neutralized DENV-1–4, yellow fever virus (YFV) and WNV on BHK-21 cells with different neutralization potential. However, no neutralization activity of 2A10G6 was observed for JEV, TBEV and DEDSV.

DEDSV replication and growth were also tested in the mosquito cell line C6/36 and the results showed that the virus can grow in this cell line with high titre, but failed to produce any obvious cytopathic effect (CPE).

DISCUSSION

In 2010, a highly contagious duck disease spread widely and quickly in the duck-farming regions of China. The disease caused severe decreases in egg production and deaths in many birds, leading to huge economic losses (Hu et al., 2011; Su et al., 2011). Previously, we identified the pathogen as a new TMUV-related flavivirus, initially named as BYDV (Su et al., 2011), which we have now renamed DEDSV. Entire genome sequencing, phylogenetic analysis and antigenic cross-reactivity studies were undertaken to characterize DEDSV in this study. In the light of its close relationship with TMUV and STWV, we have also carried out sequencing of the ORFs of TMUV and STWV for comparative analysis.

Seven strains of DEDSV were sequenced, including both the 5' and 3'-NCRs, and their comparative genome alignments revealed high levels of identity. Phylogenetic analyses based on the whole genomes of 39 flaviviruses and the polyproteins of 42 viruses demonstrated the close relationships between DEDSV and members of the NTAV group viruses, but a particularly close relationship with TMUV.

In conclusion, analysis of the phylogenetic data (Fig. 2) provides us with a plausible explanation for the evolutionary origins and dispersal of DEDSV. First, the phylogenetic cluster that contains DEDSV exclusively contains Old World viruses of either African or Asian origin, all of which diverged from a sister group of New World viruses represented by ILHV and ROCV. In turn, the JEV-related viruses diverged from this DEDSV-related cluster which comprised Old World viruses and, in this tree, a single New
World virus (SLEV), the lineage of which clearly diverged from the Old World viruses. Thus, the most likely evolutionary scenario for DEDSV and indeed for TMUV is that they represent Old World lineages, probably of African origin (cf BAGV) that were dispersed into Asia via migratory birds and ornithophilic Culex spp. mosquitoes.

The genome strategy of DEDSV is the same as that of all the other mosquito-borne flaviviruses. Likewise, the distribution of cysteine residues in C, prM and E are identical to the other flaviviruses. However, the presence of the predicted N-LGlyS of DEDSV is different from that of the other NTAV group viruses. Currently, we do not know the function of these sites for DEDSV; nevertheless, in the absence of experimental data, there have been controversial results. Winkler et al. (1987) reported that non-glycosylated viruses were antigenically indistinguishable from glycosylated viruses. In contrast, other groups reported that glycosylation was essential for the virus, as viral replication, virulence, maturation or release of viral-like particles were perturbed in non-glycosylated variants.
(Beasley et al., 2005; Crabtree et al., 2005; Goto et al., 2005; Li et al., 2006). Further studies will be needed to resolve these issues.

DEDSV shares many conserved motifs with other flaviviruses, including the catalytic triad motif, the substrate-binding motif and the RNA helicase motif in the NS3 and NS5 proteins. However, with regard to the RGX (D, E or T) motif in the E protein, which has been reported to be conserved among the JEV group, the sequence is SGK in DEDSV, TMUV and STWV. Notably, the corresponding sequences of BAGV, ILHV and ROCV are TGE, QEN and TGP, respectively. The motif has been postulated to be involved in receptor binding when viruses interact with the host cells. Therefore, this observation presumably has a significant impact on host specificity of these viruses and may partly explain why DEDSV has emerged recently to cause egg-drop syndrome in China.

At least one base-pairing (rather than the sequences per se) between the 5'-3'-terminal sequences is essential for viral RNA cyclization (Villordo & Gamarnik, 2009), which is critical for genome replication and translation (Khromykh et al., 2001). DEDSV has one such conserved base-pairing as is the case for the other viruses in the NTAV group. For the CS in the 3'-NCR, we know that CS1 is necessary for the replication of the genome, but CS2 and CS3 are dispensable. DEDSV has an ImRCS3-CS3-RCS2-CS2-CS1 pattern, differing from the other viruses in the NTAV group, but is the same as that of the JEV group viruses. In this respect, DEDSV is more closely related to the JEV group viruses than to the other NTAV group viruses. Clearly, the secondary structures of the CSs require further studies to elucidate their functions.

The cross-reactivity experiments illustrated significant shared antigenic relationships between DEDSV and JEV, WNV, DENV, YFV and TBEV. All of the representative flaviviruses that were included in this analysis are pathogenic for human beings, consequently the potential threat of DEDSV to human health cannot be overlooked, especially when taking into consideration the similarities with the JEV group analysed above. The broad neutralization mAb 2A10G6 can react with DEDSV at a high degree, but has no neutralization activity.

In conclusion, we have characterized an important newly emerging avian arboviral pathogen. We have assigned a new name to this virus to avoid potential confusion with the plant virus BYDV. DEDSV is a new and highly pathogenic member of the NTAV group of mosquito-borne flaviviruses and most-closely related to TMUV and STWV. From the evidence presented above, we propose that these three viruses, isolated from distinct and distant localities, should be grouped together to form a new subgroup in the NTAV group. On the basis of phylogenetic, phylogeographic and ecological data, these three viruses are most probably

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**Table 4.** Conserved sequences (CSs) in the 3'-NCRs of DEDSV compared with BAGV

<table>
<thead>
<tr>
<th>Region</th>
<th>Consensus sequence*</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>CS1</td>
<td>ASCATATGGACRCWGGAAGAC</td>
<td>DEDSV AGCATATGGACACCTGGGA–TAGAC</td>
</tr>
<tr>
<td></td>
<td>AGCATATGGACACCTGGGA–TAGAC</td>
<td>BAGV AGCATATGGACACCTGGGA–GAGAC</td>
</tr>
<tr>
<td>CS2</td>
<td>GGWCTAGAGGGTAWGGAGACCC</td>
<td>DEDSV GGACTAGAGGTTAGGAGGACCC</td>
</tr>
<tr>
<td></td>
<td>GGACTAGAGGTTAGGAGGACCC</td>
<td>BAGV GGACTAGAGGTTAGGAGGACCC</td>
</tr>
<tr>
<td>RCS2</td>
<td>GGWCTAGAGGGTAWGGAGACCC</td>
<td>DEDSV GGACTAGAGGTTAGGAGGACCC</td>
</tr>
<tr>
<td></td>
<td>GGACTAGAGGTTAGGAGGACCC</td>
<td>BAGV GGACTAGAGGTTAGGAGGACCC</td>
</tr>
<tr>
<td>CS3</td>
<td>YCCCAAGWGGAGACTGGGTADAMCAAAKSBR</td>
<td>DEDSV CCCCAGGGAGGACTGGGTTAACAAATCG</td>
</tr>
<tr>
<td></td>
<td>CCCCAGGGAGGACTGGGTTAACAAATCG</td>
<td>BAGV CCCCAGGGAGGACTGGGTTAACAAATCG</td>
</tr>
<tr>
<td>RCS3</td>
<td>YCCCAAGWGGAGACTGGGTADAMCAAAKSBR</td>
<td>DEDSV CCCCAGGGAGGACTGGGTTAACAAATCG</td>
</tr>
<tr>
<td></td>
<td>CCCCAGGGAGGACTGGGTTAACAAATCG</td>
<td>BAGV CCCCAGGGAGGACTGGGTTAACAAATCG</td>
</tr>
</tbody>
</table>

*Consensus sequences of CS1, CS2 and CS3 are based on the sequences of YFV, DENV complex viruses and JEV complex viruses. Solid line indicates a gap created artificially for alignment purposes. The following single letter codes are used: B, C, G or T; D, A, G or T; K, G or T; M, A or C; R, A or G; S, C or G; W, A or T; Y, C or T.

†The 3'-NCR of BAGV does not contain this sequence.

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descendant lineages of an African virus, which is transmitted between avian species via Culex spp. mosquitoes.

**METHODS**

**Virus isolation and identification.** Five DEDSV strains were isolated from the brains and ovaries of ducks presenting with DEDS (Su et al., 2011). Two additional strains were isolated from the brain tissue of a pigeon and a goose. The RNA of all of these DEDSV strains had been identified by RT-PCR assay as described previously (Su et al., 2011). TMUV MM1775 was originally identified in 1955 (Berge, 1975; Institute for Medical Research Federation of Malaya, 1957). The virus had been isolated from Culex tritaeniorhynchus collected on the outskirts of Kuala Lumpur, Malaysia by inoculation of suckling mouse brains. STWV was obtained as inactivated RNA through the courtesy of Dr Yuji Kono (Kono et al., 2000).

**Cell culture and virus propagation.** DEDSV, at the third passage in embryonated duck eggs, was stored at -20 °C. For cell culture passage, BHK-21 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM; Invitrogen) supplemented with 10% FCS, 100 IU penicillin ml⁻¹ and 100 mg streptomycin ml⁻¹. Cell monolayers were inoculated with stored infectious allantoic fluid, diluted in tenfold steps. The cultures were incubated at 37 °C in a 5% CO₂ atmosphere and checked daily for CPE. When approximately 75% of the cells showed CPE, the cultures were harvested, diluted and used to infect fresh cells. Virus from the third passage was used for virus genome amplification. The records show that TMUV MM1775 had been cultivated in suckling mouse brains for at least 11 sequential passages.

**The infectivity assay of DEDSV to the C6/36 cell line.** DEDSV-bvd1 (1 × 10⁵ p.f.u. ml⁻¹) was first added into C6/36 cell monolayers (1 × 10⁶ cells per well) in six-well plates, then incubated at 33 °C for 1 h. The supernatants were removed and RPMI 1640 cultures with
Preparation of virus RNA, RT-PCR and sequencing. Viral RNA was extracted directly from the third passage of cell cultures for DEDSV or from mouse brain for TMUV and STWV using the QiAamp Viral RNA Mini kit (Qiagen) following the manufacturer’s protocol. cDNA was synthesized using the Reverse Transcription System for RT-PCR (Promega). After reverse transcription, PCR was performed using the Phusion High-fidelity DNA Polymerase kit (NEB). Primers used for amplifying and sequencing the ORF regions (about 10 400 bp long) were described previously (Su et al., 2011). Temperature cycles used were one cycle of heat denaturation at 94 °C, 5 min; 30 cycles of three temperature changes (94 °C for 1 min, 53 °C for 1 min and 72 °C for 1 min) for amplification and one cycle for final extension at 72 °C for 10 min. The amplicons were electrophoresed on a 1 % agarose gel. 5'- and 3'-ends of the genome were amplified using 5'-RACE and 3'-RACE, respectively. 5'-RACE was conducted according to the methods described previously (Freeman et al., 2009). 3'-Ends were first poly(A) tailed and then amplified using 3'-RACE kits, according to the manufacturer’s instructions (Invitrogen). The amplicons were sequenced using dideoxynucleotide cycle sequencing and products were run on an automated sequencer (Sunbiotech). For TMUV MM1775 and STWV, sequencing procedures were carried out as described previously (Grard et al., 2007).

Multiple sequence alignment and phylogenetic analysis. Multiple sequence alignment and phylogenetic analysis were performed as described previously (Cook et al., 2012). Briefly, alignments were conducted via MUSCLE embedded in MEGA5 (Tamura et al., 2011). For the analysis of 40 representative flaviviral complete genome sequences and 42 representative flaviviral ORF nucleotide sequences, alignment was submitted to the GBLOCKS program, which objectively eliminates poorly aligned positions and divergent regions (Edgar, 2004). The best-fit models of nucleotide substitution were selected by using jModelTest (Posada, 2008). Phylogenetic analyses were conducted using these selected models under the Bayesian Markov chain Monte Carlo method implemented in MrBayes v3.2.0 (Huelsenbeck & Ronquist, 2001), with a minimum of 10 million generations and a burn-in of 10%.

Determination of cleavage sites, glycosylation sites, cysteine residues and conserved motifs. For the polypeptides of DEDSV, TMUV and STWV, potential cleavage sites were determined by using the SignaP-NN program (http://www.cbs.dtu.dk/services) as described previously. Glycosylation sites were predicted by the NetNGlyc program (http://www.cbs.dtu.dk/services), and the cysteine residues were found by Protran [v. 7.1.0.4(44)] included in the LaserGene Program (DNA Star). Conserved motifs were localized by alignment using MEGA software (Tamura et al., 2011).

Prediction of the genome secondary structure. The secondary structures in the 5'- and 3'-terminal sequences and cyclization between 5'- and 3'-terminal regions were implemented with the Mfold program (Zuker, 2003). The first 150 nt of the genome including the entire 5'-NCR and the 5'-CS in the C gene and the last 110 nt of the 3'-NCR were used for the analysis, because the domains for cyclization of the genome were included (Villordo & Gamarnik, 2009).

Serology (indirect ELISA). In order to determine whether or not serum from DEDSV-infected ducks can cross-react antigenically with other representative flaviviruses, including TMUV, BAGV, JEV, WNV, DENV, YFV and TBEV, an indirect ELISA was performed as described previously (Deng et al., 2011). All experiments involving the use of live TBEV and WNV were performed under the BSL-3 containment facilities. The strain names of the representative flaviviruses are listed as follows: JEV strain Beijing-01 (Hashimoto et al., 1988), WNV strain Chin-01 (Li et al., 2010), DENV-2 strain D2-43 (Liu et al., 2010), YFV strain 17D (Galler et al., 1998) and TBEV strain Senzhang (Si et al., 2011).

Briefly, all of the tested flaviviruses except TMUV and BAGV were standardized at 1000 p.f.u. ml⁻¹. Flavivirus antigens were prepared from culture supernatants of the infected mosquito C6/36 cells or infected suckling mouse brain suspensions and were inactivated for 30 min at 56 °C. Subsequently, 96-well microtitre plates were coated overnight at 4 °C with these heat-inactivated antigens (1 : 100 dilution) in a pH 9.6 carbonate buffer. Plates were washed five times with PBST and blocked for 2 h at 37 °C with PBST containing 5 % skimmed milk. Plates were rinsed five times in PBST and then incubated with serum from DEDSV-infected ducks. Duck serum from non-infected birds was used as the control (in triplicate) for 1 h at 37 °C. Plates were washed five times and then incubated with peroxidase-conjugated goat anti-duck IgG (1 : 500) (KPL) for 1 h at 37 °C. Plates were washed five times and then incubated with TMB substrate (Promega). The reaction was stopped by the addition of 1 M H₂SO₄, and emission (450 nm) was read using a microplate reader (Beckman). Independently, cross-reactivities between DEDSV and different flavivirus-specific antibodies or the flavivirus cross-reactive mAb 2A10G6 were determined by ELISA.

2A10G6 (100 µg ml⁻¹) was pre-incubated (1 h, 37 °C) with approximately 200 p.f.u. of various flavivirus strains and the mixtures were added to BHK-21 cell monolayers in a 12-well plate and incubated at 37 °C for 1 h. Then mixtures were removed and 1 ml of 1.0 % (w/v) LMP agarose (Promega) in 2 x DMEM plus 4 % (v/v) FBS was layered onto BHK-21 cells. After further incubation at 37 °C for 4–5 days, the wells were stained with 1 % (w/v) crystal violet dissolved in 4 % (v/v) formaldehyde to visualize the plaques. The percentage of plaque reduction was calculated as described previously (Deng et al., 2011).

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REFERENCES


Characterization of DEDS flavivirus


