Virulence and Evolution of West Nile Virus, Australia, 1960–2012


Worldwide, West Nile virus (WNV) causes encephalitis in humans, horses, and birds. The Kunjin strain of WNV (WNV$_{KUN}$) is endemic to northern Australia, but infections are usually asymptomatic. In 2011, an unprecedented outbreak of equine encephalitis occurred in southeastern Australia; most of the ≈900 reported cases were attributed to a newly emerged WNV$_{KUN}$ strain. To investigate the origins of this virus, we performed genetic analysis and in vitro and in vivo studies of 13 WNV$_{KUN}$ isolates collected from different regions of Australia during 1960–2012. Although no disease was recorded for 1984, 2000, or 2012, isolates collected during those years (from Victoria, Queensland, and New South Wales, respectively) exhibited levels of virulence in mice similar to that of the 2011 outbreak strain. Thus, virulent strains of WNV$_{KUN}$ have circulated in Australia for ≥30 years, and the first extensive outbreak of equine disease in Australia probably resulted from a combination of specific ecologic and epidemiologic conditions.

West Nile virus (WNV) is a mosquito-transmitted flavivirus that causes encephalitis. Outbreaks of potentially fatal neurologic syndromes have occurred in Europe and Africa (1); recently, however, strains of WNV have caused large outbreaks of encephalitis in humans and horses in the Americas and Australia (2,3). The Kunjin strain of WNV (WNV$_{KUN}$) is indigenous to Australia and historically has caused only relatively mild, nonfatal disease in humans and horses. However, in 2011, a large unprecedented outbreak of encephalitis in horses, involving ≈900 reported cases, occurred in southeastern Australia; a high proportion of cases were attributed to the emergence of a virulent strain of WNV$_{KUN}$ (3,4). WNV$_{KUN}$ has been shown to be enzootic to northern Australia and to have epidemic activity in southern regions thought to be associated with periods of heavy rainfall (5). However, the epidemiology of WNV$_{KUN}$ seems to have changed over the past decade; virus activity has been detected in the absence of prior flooding and in areas where it was previously not detected (4).

Studies comparing the virulence of various WNV strains in mouse models have identified several motifs, residing in both structural and nonstructural genes as well as in the 5′ and 3′ untranslated regions (UTRs). These motifs were associated with enhanced viral invasion of the central nervous system and onset of neurologic disease (5–11).

To identify potential markers of virulence of WNV$_{KUN}$ in Australia, we investigated evolutionary mechanisms behind the emergence of virulent strain(s) by using established mouse models to compare the neuroinvasive properties of WNV$_{KUN}$ isolates collected from different regions of Australia during 1960–2012. To investigate known markers of WNV virulence, we conducted comparative analyses of viral genome sequences.

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Methods

Cell Culture, Virus Production, and Titration
We used 13 WNV\textsubscript{KUN} strains isolated during 1960–2012 (Table 1) and African green monkey kidney (Vero) and \textit{Aedes albopictus} mosquito (C6/36) cells, cultured as previously described (3). The 1960 prototype WNV\textsubscript{KUN} strain (WNV\textsubscript{KUN1960}) was used as the attenuated reference virus and was either an unknown passage of the original isolate (WNV\textsubscript{MRM16}) or derived from an infectious clone of a plaque-purified virus (WNV\textsubscript{MRM61C}) (12, 13). These isolates were previously shown to be phenotypically identical (13, 14). WNV strain NY99-4132 was obtained from the US Centers for Disease Control and Prevention (Fort Collins, CO, USA) and used as a virulent control. Virus stocks and methods for determination of infectious titers have been described (3).

Antigenic Analysis
We compared reactivity of WNV\textsubscript{KUN} isolates with a panel of monoclonal antibodies (mAbs) with that of reference strains WNV\textsubscript{KUN1960} and WNV\textsubscript{NY99}. To do so, we used a fixed-cell ELISA, as previously described (3, 5, 15).

Virus Replication Kinetics
We performed growth kinetics analysis by infecting Vero and C6/36 cells at a multiplicity of infection of 1 at 37°C (Vero) or 28°C (C6/36) with WNV\textsubscript{KUN}. Culture supernatants were harvested at 0, 24, 48, and 72 h after infection (Vero) and 0, 24, 48, 72, 96, and 120 h after infection (C6/36) and titrated (3). Statistical significance from 3 independent experiments was determined by using 2-way analysis of variance following log transformation (16). Mean virus titers were compared between viruses by using the Tukey method for pairwise multiple comparisons (GraphPad Prism, version 6.0; GraphPad Software Inc., San Diego, CA, USA).

Virulence in Mice
Performance of all animal procedures was approved by The University of Queensland Animal Ethics Committee. To determine virus virulence in mice, we intraperitoneally inoculated 20 Swiss white outbred CD-1 mice (weanlings [18–19 days of age] and young adults [28 days of age]) with a range of doses (0.1–10,000 PFU) of each WNV strain (Table 1) (3). The significance of clinical differences between groups was calculated by Kaplan-Meier analysis and analyzed by log-rank test where noted (GraphPad Prism, version 6.0). A virus strain was designated as virulent if survival times for mice infected with this strain (both age groups) differed significantly from those of mice infected with the attenuated reference strain WNV\textsubscript{KUN1960}. All virus strains that did not meet this criterion for virulence were designated as attenuated.

Full-length Genome Sequencing
We sequenced 9 WNV\textsubscript{KUN} genomes (online Technical Appendix Table 1, http://wwwnc.cdc.gov/EID/article/22/8/15-1719-Techapp1.pdf) by using random primer sequencing on extracts of C6/36 cell cultures (17). Viral RNA was extracted by using a MagMAX-96 Viral RNA Isolation Kit (Ambion, Waltham, MA, USA) according to the manufacturer’s instructions. cDNA synthesis and random PCR amplification were conducted according to previously described methods (18), and resultant PCR amplicons were used for sequencing library preparation. DNA libraries were prepared by using a Nextera XT DNA Sample Preparation Kit (Illumina, San Diego, CA, USA) according to the manufacturer’s protocols. Paired-end sequencing of 150-bp fragments was performed by using a MiSeq Reagent Kit V2 (300 cycles) and MiSeq Sequencing System (Illumina). Sequencing data were analyzed by using CLC Genomics Workbench version 6.5.0 (http://www.clcbio.com). The sequence data were trimmed by

Table 1. WNV\textsubscript{KUN} strains used during study of virulence and evolution of WNV, Australia, 1960–2012*

<table>
<thead>
<tr>
<th>Isolate</th>
<th>Year collected</th>
<th>Location</th>
<th>Source</th>
<th>Passage history</th>
</tr>
</thead>
<tbody>
<tr>
<td>MRM16/MRM61C‡</td>
<td>1960</td>
<td>Mitchell River Mission, Queensland</td>
<td>Mosquito†</td>
<td>Unknown</td>
</tr>
<tr>
<td>Boort</td>
<td>1984</td>
<td>Victoria</td>
<td>Horse spinal cord</td>
<td>Unknown</td>
</tr>
<tr>
<td>K2499</td>
<td>1984</td>
<td>Kimberley region, Western Australia</td>
<td>Mosquito†</td>
<td>2× C6/36; 1× PSEK</td>
</tr>
<tr>
<td>Hu6774</td>
<td>1991</td>
<td>New South Wales</td>
<td>Human</td>
<td>Unknown</td>
</tr>
<tr>
<td>K6453</td>
<td>1991</td>
<td>Kimberley region, Western Australia</td>
<td>Mosquito†</td>
<td>2× C6/36; 1× PSEK</td>
</tr>
<tr>
<td>SH183</td>
<td>1991</td>
<td>Victoria</td>
<td>Chicken</td>
<td>Unknown</td>
</tr>
<tr>
<td>Gu0631</td>
<td>2000</td>
<td>Gulf of Carpentaria, Queensland</td>
<td>Mosquito†</td>
<td>3× C6/36</td>
</tr>
<tr>
<td>Gu1009</td>
<td>2000</td>
<td>Gulf of Carpentaria, Queensland</td>
<td>Mosquito†</td>
<td>3× C6/36</td>
</tr>
<tr>
<td>K68967</td>
<td>2009</td>
<td>Kimberley region, Western Australia</td>
<td>Mosquito†</td>
<td>3× C6/36</td>
</tr>
<tr>
<td>P9974</td>
<td>2009</td>
<td>Pilbara region, Western Australia</td>
<td>Mosquito†</td>
<td>3× C6/36</td>
</tr>
<tr>
<td>NSW2011</td>
<td>2011</td>
<td>New South Wales</td>
<td>Horse brain</td>
<td>2× C6/36; 1 Vero</td>
</tr>
<tr>
<td>K74015</td>
<td>2011</td>
<td>Kimberley region, Western Australia</td>
<td>Mosquito†</td>
<td>3× C6/36</td>
</tr>
<tr>
<td>NSW2012</td>
<td>2012</td>
<td>New South Wales</td>
<td>Mosquito†</td>
<td>3× C6/36</td>
</tr>
</tbody>
</table>

*C6/36, from Aedes albopictus mosquitoes; PSEK, porcine squamous equine kidney cells; WNV, West Nile virus; WNV\textsubscript{KUN}, Kunjin strain of WNV.
†Prototype strain.
‡Culex annulirostris.
using quality scores specified in CLC Genomics Workbench before performing read-mapping analysis. The genome sequences were assembled by read mapping against the reference WNV\textsubscript{KUN} strain genome (GenBank accession no. JX276662) with use of default parameters in the mapping algorithm. Where gaps in the genome sequence or low coverage (<10 reads/site) were observed, conventional Sanger sequencing was performed to complete or verify the sequence. Oligonucleotide primers sequences designed for these purposes are available upon request to A.A.K or R.A.H.

**Bioinformatics Analysis**

We used MUSCLE, as implemented in MEGA6 (19), to align complete open reading frame (ORF) (10,320 nt) and partial (402 nt) envelope (E) gene nucleotide sequences of the newly sequenced WNV\textsubscript{KUN} strains, together with those available for 6 other WNV\textsubscript{KUN} strains and selected WNV isolates, representing different lineages. We estimated maximum-likelihood phylogenetic trees by using PhyML version 3.0 (20) and by using substitution models and rates among sites selected with JModelTest version 2.1.5 (21). We tested reliability of the inferred trees by using the bootstrap method with 1,000 replicates. All trees were rooted with analogous ORF sequences from Murray Valley encephalitis virus (GenBank accession no. NC000943) and Japanese encephalitis virus (GenBank accession no. EF571853) and visualized by using FigTree version 1.4.0 (http://tree.bio.ed.ac.uk/software/figtree/). Pairwise distances were determined at the nucleotide and amino acid levels by using the p-distance model in MEGA6.

**Results**

**WNV\textsubscript{KUN} Strains**

A panel of previously characterized mAbs (5,15,22–24) was used to antigenically type 13 WNV\textsubscript{KUN} strains. The binding profiles of these mAbs confirmed that all WNV\textsubscript{KUN} isolates closely resembled the prototype WNV\textsubscript{KUN1960} strain, including strong recognition by mAb 10A1, known to be specific for WNV\textsubscript{KUN} strains (Table 2) (2,24). Only WNV\textsubscript{KUN} strains isolated before 2000 were recognized by mAb 5H1, which binds a linear epitope in the αA3 motif (residues 39–53) in the methyltransferase domain of nonstructural (NS) protein 5 (15). Lack of 5H1 binding was associated with a substitution at residue 49 (Ile-Val) in αA3 (Table 3); this finding was consistent with previous mutagenesis study findings that a substitution of Ile to Ala at this residue was associated with abolition of 5H1 binding (26).

**Glycosylated E Proteins**

All analyzed WNV\textsubscript{KUN} isolates collected after 1960 contain glycosylated E proteins. The sequence analysis of the E gene of WNV\textsubscript{KUN} isolates revealed the presence of a conserved potential N-linked glycosylation site at residue 154 in all but the prototype WNV\textsubscript{KUN1960} isolate (Table 3). To confirm that this site was indeed glycosylated on the viral E protein, each virus was assessed for recognition in ELISA by mAb 17D7 and 3.101C (Table 2), which specifically recognize glycosylated and unglycosylated WNV\textsubscript{KUN} E proteins, respectively (5,23). The results supported the predictions from our sequencing data; all WNV\textsubscript{KUN} strains except the prototype WNV\textsubscript{KUN1960} strain were recognized by mAb 17D7 but not by 3.101C.

**Table 2. Binding patterns of monoclonal antibodies to WNV strains in ELISA**

<table>
<thead>
<tr>
<th>Strain</th>
<th>Year of isolation</th>
<th>Pan-WNV\textsubscript{2012}, anti-env</th>
<th>Pan-WNV\textsubscript{2012}, 4G2, anti-env</th>
<th>Pan-WNV\textsubscript{2012}, 2B2, anti-env</th>
<th>10A1, anti-env</th>
<th>5D4, anti-NS5</th>
<th>5H1, anti-NS5</th>
<th>Glycosylated E, 17D7</th>
<th>Unglycosylated E, 3.101C</th>
<th>MVEV specific, 10C6</th>
</tr>
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<td>WNV\textsubscript{KUN}\textdagger</td>
<td>1960</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>KUN1960</td>
<td>1984</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
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<td>1991</td>
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<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
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<tr>
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<td>+</td>
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<tr>
<td>Gu1009</td>
<td>2000</td>
<td>+</td>
<td>+</td>
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<td>+</td>
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<tr>
<td>K89867</td>
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</tr>
<tr>
<td>P9974</td>
<td>2011</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
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</tr>
<tr>
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<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
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<td>+</td>
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<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
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</tbody>
</table>

*Binding of a monoclonal antibody was designated as positive if the optical density was at least double the optical density of the negative control (mock-infected C6/36 cells). E, envelope protein; MVEV, Murray Valley encephalitis virus; NS5, nonstructural protein 5; WNV, West Nile virus; WNV\textsubscript{KUN}, Kunjin strain of WNV; †WNV\textsubscript{KUN} strains collected during 1960–2012, Australia.

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Table 3. Amino acid sequences in the West Nile virus genome*

<table>
<thead>
<tr>
<th>WNV strain</th>
<th>Year of isolation</th>
<th>prM, residue</th>
<th>E protein, residues</th>
<th>NS3, residue</th>
<th>NS5, residue</th>
<th>3' UTR residues</th>
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<tbody>
<tr>
<td>NY99</td>
<td>1999</td>
<td>Val/Ser</td>
<td>154–156†</td>
<td>Pro</td>
<td>Phe</td>
<td>Val</td>
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<tr>
<td>KUN1960</td>
<td>1960</td>
<td>Ile/Leu</td>
<td>249§</td>
<td>Ala</td>
<td>Phe</td>
<td>Absent</td>
</tr>
<tr>
<td>Boort</td>
<td>1984</td>
<td>Ile/Leu</td>
<td>653¶</td>
<td>Ala</td>
<td>Ile</td>
<td>Absent</td>
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<td>1984</td>
<td>Ile/Leu</td>
<td>49</td>
<td>Val</td>
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<td>Absent</td>
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<td>K6453</td>
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<td>64–71</td>
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<tr>
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<td>2000</td>
<td>Ile/Leu</td>
<td>2011</td>
<td></td>
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<td>Gu1009</td>
<td>2009</td>
<td>Ile/Leu</td>
<td>2012</td>
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</tr>
<tr>
<td>K68967</td>
<td>2009</td>
<td>Ile/Leu</td>
<td>2013</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*E, envelope; NS, nonstructural; prM, premembrane; UTR, untranslated region.
†(17).
‡(10).
§(25).

Growth Kinetics and Plaque Morphology of WNV KUN Strains

Infection of Vero cells at 24 h postinfection demonstrated that WNV KUN1960, WNV K2499, WNV K6453, and WNV K68967 isolates yielded significantly lower titers than did WNV NY99 (p<0.05) (Figure 1, panel A). However, by 48 h postinfection, similar titers were reached for all WNV isolates except WNV KUN1960.

A similar trend was observed in C6/36 cells, in which viral titers of WNV KUN1960, WNV K2499, WNV K6453, and WNV K68967 and WNV Gu1009 were significantly lower than those for WNV NY99, WNV NSW2011, and WNV Boort at 48 h after infection (p<0.05) and titers for WNV K6453, WNV K74015, and WNV Gu0631 were intermediate. By 96 h after infection, the titers of all WNV isolates except WNV KUN1960 were similar.

In terms of plaque morphology of WNV KUN strains, in Vero cells, WNV NSW2011 and WNV K74015 produced large plaques (average size 4.3 ± 0.63 and 4.3 ± 0.77 mm, respectively), a size similar to those produced by the WNV NY99 strain (average size 4.8 ± 0.45 mm). The prototype virus, WNV KUN1960, produced very small plaques (average size 2.7 ± 0.47 mm), which differed significantly from those of all other viruses tested during this study (p<0.0001). The remaining isolates produced intermediate-sized plaques (average size 3.5–3.9 ± 0.45–0.84 mm) (Figure 2) (5). Plaques formed by WNV K6453, WNV K74015, WNV Gu1009, and WNV Gu0631 were less well defined than those formed by WNV NY99, WNV NSW2011, WNV K2499, and WNV Boort.

Virulence of WNV KUN Strains in Mice

We previously demonstrated that differentiation between virulent and attenuated strains of WNV can be detected in weanling and young adult mice (3,27). In this study, we found that in addition to WNV NSW2011, 3 other WNV isolates (WNV Boort†, WNV Gu0631‡, WNV NSW2012§) were neuroinvasive in both mouse models (Figure 3; online Technical Appendix Table 2). The WNV Boort strain, obtained from the spinal cord of a symptomatic horse during a small outbreak of equine disease in southeastern Australia in 1984, was neuroinvasive in young adult mice (40% mortality rate at 1,000 PFU); this finding did not statistically differ in this respect from that for WNV NSW2011 (p = 0.3218).
Two other isolates obtained from mosquitoes, 1 from the Gulf of Carpentaria in 2000 (WNV_Gu0631) and 1 from southeastern Australia in 2012 (WNV_NSW2012), also exhibited levels of neuroinvasive properties in young adult mice similar to those caused by WNV_NSW2011 (Figure 3). In weanling mice, the virulence of WNV_Boort, WNV_Gu0631, and WNV_NSW2012 was also similar to that of WNV_NSW2011; mortality rates, 50% lethal dose, or time to death did not differ significantly (Table 4). Of note, WNV_Gu1009 isolated at the same time and from the same region as WNV_GU0631 was significantly less virulent in young adult mice (Figure 3; online Technical Appendix Table 2). The remaining isolates were relatively attenuated in both young adult (Figure 3) and weanling (online Technical Appendix Table 2) mice and did not differ significantly from the attenuated prototype WNV_KUN1960 strain (p>0.05).

Sequence of Viral Genomes
We sequenced WNV_KUN isolates to analyze their relationship to the prototype WNV_KUN strains from 1960 (WNV_MRM16, WNV_MRM61C); the 2011 outbreak strains (WNV_NSW2011, WNV_SA2011, WNV_V11-07); and exotic strains of WNV known to be virulent in humans and horses (WNV_NY99) or representing different WNV lineages. Phylogenetic analysis of the ORF sequences demonstrated that the WNV_KUN strains form a single genetically homogeneous clade within lineage 1 (Figure 4), as previously recognized (24); nucleotide and amino acid identities between strains were 96.1%–99.4% and 98.5%–100%, respectively. As expected, the most recent isolates, including the 2011 outbreak strain, were the most divergent, and the early prototype strains (WNV_MRM61C and WNV_MRM16) occupied the basal lineage of this clade. Recent strains isolated in 2011 and 2012 from southeastern Australian states clustered together and shared high levels of nucleotide (98.6%–100%) and amino acid (99.6%–100%) identities, indicating transmission of a genetically homogeneous virus population during this period. These strains were either virulent for horses or shown in this study to be virulent in mice (Figure 3; online Technical Appendix Table 2). No other association between phylogenetic relationships and virulence was found; other virulent strains clustered closely and interspersed with the attenuated strains identified in this study. An expanded phylogenetic analysis that used 45 partial E gene sequences (402 nt) and a larger range of reference WNV_KUN strains showed a similar pattern of relationships (online Technical Appendix Figure).

Of note is the very close relationship (99.9% aa identity) between a virulent 2011 strain isolated from a horse and an isolate obtained from Culex annulirostris mosquitoes trapped in New South Wales, Australia, in 2012 (WNV_NSW2012). Only 3 nonconservative changes were identified between WNV_NSW2011 and WNV_NSW2012 located in NS1 (Lys33Arg), NS3 (Phe509Leu), and NS4A.
analyses of the complete ORF sequence of each WNV<sub>M</sub> isolate revealed that, in addition to the glycosylation site at residues 154–156 in the E protein, all strains isolated after 1960 contained a Phe residue at position 653 in the NS5 protein, which has previously been shown to play a role in resistance to antiviral activity of interferon-α/β (10) (Table 3). In contrast, WNV<sub>M1960</sub> contained a Ser residue at position 653 in NS5 (3,5,24,28). The Pro residue at position 249 in the NS3 protein, previously shown to be present in WNV strains and associated with increased virulence in birds of some species (25), was not present in any of the WNV<sub>M</sub> isolates, which all contained an Ala residue at this position (Table 3).

In addition to an Ile→Val substitution at position 49 in NS5 of WNV<sub>M</sub> isolates collected after 2009, analysis of more contemporary WNV<sub>M</sub> isolates also revealed a consistent 8-nt deletion in the 3′ UTR, just downstream of the ORF stop codon. This deletion was identical in WNV<sub>G1009</sub> and all isolates collected after 2000. In contrast, this deletion was not present in isolates collected before 2000 (Table 3) or in another isolate from Gulf of Carpentaria collected in 2000 (WNV<sub>G6063</sub>). We suggest that these 2 features (Ile→Val 49 residue in NS5 and an 8-nt deletion in the 3′ UTR) can be considered as potential evolutionary markers.

In addition to the genetic variability described above, sequence analysis between virulent and attenuated WNV<sub>M</sub> strains identified other nucleotide differences between isolates, located throughout the viral genome. These differences result in amino acid substitutions (Table 4) and may contribute to observed phenotypic differences.

We also sequenced WNV<sub>M</sub> viral RNA extracted directly from mosquito saliva expectorated onto sugar-soaked nucleic acid preservation cards placed in mosquito traps in Darwin, Northern Territory, in 2012 (WNV<sub>NT2012</sub>). When partial sequences from E, NS5, and the 3′ UTR from this RNA were aligned, we observed a high level (99.7%) of identity with the WNV<sub>M1960</sub> strains, indicating that viruses genetically homologous to the prototype virus are still circulating in some regions of Australia (online Technical Appendix Figure). Closer analysis revealed a lack of E glycosylation, similar to that found in the prototype strain. However, Phe was identified at position 653 of NS5, similar to that found in recent isolates.

**Discussion**

Historically, WNV<sub>M</sub> has been associated with only mild disease in humans and rare cases of disease in horses, consistent with data from mouse virulence studies that revealed a relatively attenuated phenotype (3,8,27). Thus, the emergence of an equid-virulent strain of WNV<sub>M</sub>, responsible for ≈900 cases of encephalitis in horses in southeastern Australia, was unprecedented.

Although most WNV<sub>M</sub> isolates examined in this study exhibited an attenuated phenotype, similar to that of the prototype WNV<sub>M1960</sub>, we identified an additional 3 strains with neuroinvasive properties in mice similar to those reported for WNV<sub>NSW2011</sub> (3). The first, WNV<sub>Boort</sub>, was isolated from the spinal cord of a horse with nonsuppurative encephalomyelitis and severe neurologic symptoms in northern Victoria in 1984 (31). At that time, 53 animals in the same area were clinically affected. However, a high incidence of Ross River virus–specific antibody in these animals implicated that virus rather than WNV<sub>M</sub> as the primary etiologic agent (37). Our results are also supported by another recent study showing virulence of WNV<sub>Boort</sub> in 18–19-day-old mice (32).

The second virulent strain identified in this study, WNV<sub>G6063</sub>, was isolated from *Cx. annulirostris* mosquitoes collected from Normanton, Gulf of Carpentaria, in April 2000. Of note, this virus was isolated in the absence of any reported disease outbreak, as part of a survey for the presence of Japanese encephalitis virus in northern Queensland.

Figure 3. Survival curves for young adult (28-day-old) Swiss outbred mice after intraperitoneal infection with 1,000 PFU of West Nile virus (WNV) strains isolated in Australia, 1960–2012. Groups of 10 mice were infected with each virus. The mice were monitored for 21 days after infection for signs of encephalitis and then euthanized. WNV<sub>M1960</sub> and WNV<sub>NSW2011</sub> with previously demonstrated virulence were included as controls. The significance of clinical differences between groups was calculated by Kaplan-Meier analysis and analyzed by log-rank test. Significantly increased virulence over that of WNV<sub>M1960</sub> is indicated by an asterisk (*): WNV<sub>Boort</sub> (p = 0.0285), WNV<sub>G6063</sub> (p = 0.0115), and WNV<sub>NSW2012</sub> (p = 0.0011). No significant differences were observed between WNV<sub>Boort</sub>, WNV<sub>G6063</sub>, and WNV<sub>NSW2012</sub> compared with WNV<sub>NSW2011</sub> (p > 0.05).
(33). The second Gulf of Carpentaria isolate, WNV
\textsubscript{Gu1009}, was also collected in April 2000, from the town of Karumba, which is \textasciitilde 30 km from Normanton. However, WNV\textsubscript{Gu1009} is genetically distinct and attenuated to the same degree as the prototype WNV\textsubscript{KUN1960} in 28-day-old mice (Figure 4). These observations demonstrated that virulent WNV\textsubscript{KUN} strains might co-circulate with attenuated strains in some regions of Australia. Furthermore, the circulation of neuroinvasive strains may often appear in the absence of disease outbreaks. This suggestion is consistent with our finding that WNV\textsubscript{NSW2011} was genetically almost identical to the WNV\textsubscript{NSW2011} and exhibited similar levels of neuroinvasiveness in mice. However, no cases of disease in equids were associated with WNV\textsubscript{KUN} infection during the 2012 season (34). This lack of cases suggests that the persistence of virulent strains in southeastern Australia is not the sole determinant for initiating disease outbreaks and that specific climatic and ecological conditions, perhaps influencing mosquito populations and viral transmission, are also required.

A similar scenario occurred in North America, where an unusually high number of cases in humans (5,387), most in Texas, USA, were reported in 2012. However, sequence analysis of WNV isolates from 2012 revealed that the strains circulating in Texas were virulent and attenuated, and no specific virulence determinants responsible for the increase in cases could be identified (35). Instead, other factors, including temperature and changes in mosquito or bird populations, were speculated to have contributed to the magnitude of the 2012 outbreak (36).

To identify a phylogenetic association with virulence and to identify potential virulence determinants encoded in the genome of WNV\textsubscript{KUN} strains, we also performed full-length sequencing of the ORF of several of the viruses studied. Although recent virulent strains were phylogenetically closely related, no other association between phylogenetic grouping and virulence phenotype was found (Figure 4; online Technical Appendix Figure). One notable change in the genome that was clearly associated with the temporal distribution of these viruses was a highly conserved 8-base deletion in the 3’ UTR, just downstream of the ORF stop codon. Isolates from samples collected after 2000, including the virulent WNV\textsubscript{NSW2011} and attenuated

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* C, capsid; E, envelope; NS, nonstructural; pM, premembrane.
strains, invariably contained this deletion. This finding suggests that the deletion is an evolutionary marker but is not directly associated with virulence. This finding is also consistent with our observation that the neuroinvasive 2000 Gulf of Carpentaria isolate, WNV\textsubscript{Gu0631}, did not have this deletion but that the co-circulating attenuated isolate, WNV\textsubscript{Gu1009}, collected from the same region at the same time, did have this deletion.

An additional evolutionary change was observed in the a-A3 motif of the methyltransferase domain of the NS5 protein. Isolates obtained before 2009, including the prototype WNV\textsubscript{KUN1960}, contained a conserved Ile residue at position 49. However, all isolates collected after 2009 displayed an Ile\rightarrow Val substitution at this position. Coincidentally, this substitution abolished the binding of a mAb (5H1) that recognizes a linear epitope comprising the a-A3 peptide (15).

Initial comparisons between the virulent isolate WNV\textsubscript{NSW2011} from a horse and the attenuated prototype WNV\textsubscript{KUN1960} revealed that several previously identified WNV virulence markers were detected in the former but not in the latter isolate (3). These markers included the conserved N-linked glycosylation of the E protein (7) and the Phe residue at position 653 in the NS5 protein, associated with resistance to antiviral activity of interferon a/b (8). Although these initial observations suggested the involvement of these motifs in the enhanced neuroinvasive properties of the isolate collected from a horse in 2011, our study revealed that, with the exception of WNV\textsubscript{KUN1960}, all strains examined contain both of these markers, regardless of virulence phenotype in mouse models. Thus, it seems that, although these motifs contribute to virulence in mice, they are not likely to be solely responsible for enhancing the neuroinvasive properties of some WNV\textsubscript{KUN} strains and, hence, not likely to be markers of evolving virulence in recent isolates of WNV\textsubscript{KUN}.

Additional markers of WNV virulence identified in WNV strains from North America were not detected in any of the WNV\textsubscript{KUN} isolates. This finding is consistent with our repeated observations that even the equid-virulent
WNV_{NSW2011} is substantially less neuroinvasive than WNV_{NY99} in young adult mice (3). These motifs may include Val 22 and Ser 72 residues in the premembrane, which enhance mouse neuroinvasiveness when introduced into the prototype WNV_{KUN1960} (11), and the Pro residue at position 249 in NS3, which is associated with enhanced virulence in birds (25). The absence of the latter motif in all WNV_{KUN} strains is also consistent with the perceived lack of illness and death among birds in Australia, notably during the 2011 outbreak among equids. Some isolates included in this study (including WNV_{KUN1960*} WNV_{SH183*} WNV_{BooRT*} and WNV_{Hu6774*}) have an unknown passage history. Extensive passage through cells is known to occasionally lead to passage-adapted mutations, and care should be taken when interpreting sequencing data from these virus strains.

WNV_{KUN} is thought to be endemic to the tropical areas of northern Australia, suggesting that virulent viruses emerging in southeastern Australia probably originate from northern Australia. However, WNV_{KUN} recently isolated from mosquitoes in northern Australia, including the 2011 Kimberley isolate WNV_{K74015}, were more attenuated than WNV_{NSW2011}. This finding suggests a different explanation for the evolution of virulent WNV_{KUN} viruses, which may be associated with the adaption of WNV_{KUN} to different hosts (avian and terrestrial) or different vector species in temperate regions. In this context, virulence in equids may be just a coincidental outcome of the constraints placed on virus fitness in different geographic locations (35–37).

Overall, our results show that virulent strains of WNV_{KUN} have been circulating in Australia for ≥30 years and that the first extensive outbreak of disease among horses in Australia in 2011 probably resulted from a combination of ecologic and epidemiologic conditions rather than the emergence of a novel, more virulent strain. Further studies evaluating viral fitness of West Nile virus quasispecies in terms of population-dependent host–virus interactions, are warranted.

Acknowledgments
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Dr. Prow is a postdoctoral research fellow at the QIMR Berghofer Medical Research Institute and is engaged in research to characterize the pathogenesis and virulence of WNV strains in murine models. She also has a special interest in understanding the neuroinvasive properties of neurotrophic flaviviruses.

References

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Virulence and Evolution of West Nile Virus, Australia

Technical Appendix

Technical Appendix Table 1. GenBank accession numbers and details of WNV strains used for phylogenetic analysis in study of virulence and evolution of WNV, Australia

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**Technical Appendix Table 2. Virulence of West Nile virus strains in 18–19-d-old mice after intraperitoneal infection**

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*Reference virus, data from (1).*
Technical Appendix Figure. Maximum-likelihood phylogenetic tree estimated using partial envelope gene sequences (402 nt) of WNV_{KUN} strains and reference WNV strains. The tree was estimated using a general time-reversible model of nucleotide substitution with a gamma distribution and invariant sites. Bootstrap values are shown on the nodes and are expressed as a percentage of 1,000 replicates; values >70% only are shown. Horizontal branch lengths indicate genetic distance proportional to the scale bar. The tree generated using the same reference sequences as for the open reading frame phylogeny and was rooted with Murray Valley encephalitis and Japanese encephalitis sequences. To improve resolution, all branches of reference sequences have been removed except for lineage 1, clade a. Strains sequenced as part of this study are indicated by a closed circle. Those that were assessed as having an attenuated virulence phenotype are indicated by a single asterisk, while virulent strains are indicated by a double asterisk. The state of origin for WNV_{KUN} strains is shown with the following abbreviations: NSW, New South Wales; QLD, Queensland; SA, South Australia; VIC, Victoria; WA, Western Australia. Virus source is indicated in parentheses next to its identity, as follows: Av, avian; Eq, equine; H, human; M, mosquito. WNV, West Nile virus.
Reference

http://dx.doi.org/10.3201/eid1805.111720