CHRONIC WASTING DISEASE OF CAPTIVE MULE DEER: A SPONGIFORM ENCEPHALOPATHY

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CHRONIC WASTING DISEASE OF CAPTIVE MULE DEER: A SPONGIFORM ENCEPHALOPATHY

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Abstract: In the past 12 years (1967-79) a syndrome we identify as chronic wasting disease has been observed in 53 mule deer (Odocoileus hemionus hemionus) and one black-tailed deer (Odocoileus hemionus columbianus) held in captivity in several wildlife facilities in Colorado and more recently in Wyoming. Clinical signs were seen in adult deer and included behavioral alterations, progressive weight loss and death in 2 weeks to 8 months. Gross necropsy findings included emaciation and excess rumen fluid admixed with sand and gravel. Consistent histopathologic change was limited to the central nervous system and characterized by widespread spongiform transformation of the neuropil, single or multiple intracytoplasmic vacuoles in neuronal perikaryons and intense astrocytic hypertrophy and hyperplasia. Presented is a clinical characterization of chronic wasting disease and pathologic evidence supporting the conclusion that the disease is a specific spontaneously occurring form of spongiform encephalopathy.

INTRODUCTION

The purpose of this report is to characterize the clinical aspects and pathology of chronic wasting disease (CWD) a hitherto undescribed encephalopathic disease of mule deer (Odocoileus hemionus hemionus).

CWD has been observed in captive mule deer in a wildlife facility in Colorado for the past 12 years (1967-79), and more recently in a similar facility in Wyoming. The disease has a high morbidity in young adult animals which have been maintained in captivity for periods of 2.5 to 4 years. In the period 1974-79, CWD affected 53 of 67 mule deer and one black-tailed deer (Odocoileus hemionus columbianus) held captive for more than 2 years.

The wildlife facilities in which the disease has been observed subserve experimental nutritional, metabolic and disease studies of deer and other nondomesticated ruminants. The majority (approximately 90%) of mule deer were hand raised from infancy after they received colostrum and were separated from does at 1-5 days of age. Most were born to does trapped in the wild during pregnancy, brought to the facility, and released after parturition. The remainder were either born to resident does bred in captivity or orphan fawns captured in the wild. Fawns were bottle fed cow’s milk supplemented with vitamins and weaned at 70-90 days of age. Thereafter, groups of 2 to 10 deer occupied pens approximately 0.1 to 1.0 ha in size and received rations of alfalfa hay, commercially mixed grains supplemented with minerals and vitamins, and fresh water. Over the years a variety of grain mixes have been used with protein content ranging from 12 to 20%. From time to time a few young adult deer trapped in the wild have been added to the permanent captive populations.

Within the facilities, deer have had irregular and discontinuous contact with other wild ruminants[1] and with

[1] This study was supported in part by N.I.H. (BRSG) Grant 5807-BB-05458-17.
[2] Elk (Cervus canadensis), white-tailed deer (Odocoileus virginianus), pronghorn antelope (Antilocapra americana), Rocky Mountain bighorn sheep (Ovis canadensis), mouflon (Ovis musimon).
domestic cattle, goats and sheep. In addition, other feral mammalian species either reside within or traverse the facilities’ pens.

CLINICAL SIGNS

Clinical signs of CWD occurred only in mule deer maintained in captivity for a period of 2.5 to 4 years, whether raised from infancy by hand or captured in the wild as young adults. The disease affected males, females and castrates. Listlessness, progressive weight loss and depression with insidious onset and protracted course occurred over 2 weeks to 8 months, leading to emaciation (Fig. 1) and death. Onset occurred during all seasons. The majority of affected deer developed signs of polydipsia, polyuria, excessive salivation, grinding of the teeth, flaccid hypotonia of facial muscles, lowering of the head, drooping of the ears and terminal anorexia. In some affected animals esophageal hypotonia and dilatation, difficulty in swallowing, regurgitation of rumenal fluid and ingesta, and deprived appetite occurred. Behavioral changes included episodes of apparent lack of awareness. Affected deer often stood for several minutes with lowered head and a fixed stare, and then reverted to a more normal state of alertness. Decreased interactions with unaffected deer in the group, occasional abnormal response to restraint and hyperexcitability were noted. Although behavioral changes were a consistent feature of CWD, specific motor or sensory neurologic signs were not identified. Equivocal visual deficits occasionally were suspected. In many instances, the disease was interrupted or terminated by secondary complications,
notably pneumonia, or by euthanasia. A summary of clinical signs is presented in Table 1.

CLINICAL PATHOLOGY

Blood samples in EDTA were obtained from 11 affected deer for hematologic evaluation and serum from 10 deer for chemistry and electrolyte levels prior to death. Deer were restrained manually or sedated with xylazine hydrochloride during sample collection. Samples usually were analyzed within 24 h of collection. Erythrocyte and white blood cell counts were performed on a Coulter Counter. Hemoglobin content was determined using a Coulter hemoglobinometer. A Hycel-17 autoanalyzer was used for analysis of blood urea nitrogen (BUN), creatinine, cholesterol, bilirubin, sodium, inorganic phosphorus, chloride, potassium, creatinine phosphokinase (CPK), serum glutamic oxaloacetic transaminase (SGOT) and alkaline phosphatase. Data are shown in Tables 2 and 3. Alterations from normal values were noted only in deer in which secondary intercurrent disease existed, however consistent changes were not identified.

Cerebrospinal fluid (CSF) was collected from five affected deer immediately following electrocution. Cell counts, protein content, specific gravity, pH, and CPK levels (Biodynamics atomic spectrometry) were determined within 24 h (Table 4). CPK values ranged from 16-102 U/l which are elevated from normal levels; however, in these cases the elevation was felt to be due to the method of euthanasia. Urinalysis was performed on urine collected from the bladder of 6 deer at necropsy. Urine specific gravity ranged from 1.002 to 1.018 with 3 values below 1.006. These levels were lower than expected and considered significantly decreased in animals which were mildly to moderately dehydrated.

Sera from 1 to 5 affected deer were tested for antibody titers to Toxoplasma (indirect hemagglutination), Leptospira (slide agglutination), Brucella abortus (slide agglutination), infectious bovine rhinotracheitis virus (serum neutralization), bovine virus diarrhea virus (serum neutralization) and blue-tongue virus (immunodiffusion). All reactions were judged to be insignificant except for two deer with titers of 1:16 and 1:64 to bovine virus diarrhea virus.

Samples of liver and kidney from two deer were analyzed by atomic absorption spectrometry for selenium, lead, copper, mercury and molybdenum. Levels were considered to be within normal limits.

GROSS PATHOLOGY

Since 1974, 41 mule deer affected with CWD have been examined post-mortem. Gross pathologic findings consistently included emaciation and in 28 (68%) the rumen contained a marked excess of fluid admixed with sand or gravel. In 18 (44%) no gross lesion was found, while in 23 (56%) various other inconsistent gross lesions were observed and presumed to be the non-specific result of secondary disease processes. These included pneumonia, abscesses, enteritis, focal
TABLE 1. Summary of clinical features and gross pathologic findings in deer with histopathologically confirmed CWD.

<table>
<thead>
<tr>
<th>Sex</th>
<th>Duration of Clinical Signs (months)</th>
<th>Clinical Features</th>
<th>Gross Pathologic Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Emaciation</td>
<td>Behavioral Changes</td>
</tr>
<tr>
<td>Female</td>
<td>0.5-6</td>
<td>100%</td>
<td>100%</td>
</tr>
<tr>
<td>Male</td>
<td>1-8</td>
<td>100%</td>
<td>100%</td>
</tr>
<tr>
<td>17% (5/29)</td>
<td></td>
<td>(5/5)</td>
<td>(5/5)</td>
</tr>
<tr>
<td>Castrates</td>
<td>0.5-8</td>
<td>100%</td>
<td>100%</td>
</tr>
<tr>
<td>24% (7/29)</td>
<td></td>
<td>(7/7)</td>
<td>(7/7)</td>
</tr>
<tr>
<td>Total</td>
<td>0.5-8</td>
<td>100%</td>
<td>100%</td>
</tr>
</tbody>
</table>

aFractions refer to the number of deer showing a particular feature over the number of animals which were examined for that feature.
<table>
<thead>
<tr>
<th>Deer number</th>
<th>Date of Sample Collection</th>
<th>Age (years)</th>
<th>Sex</th>
<th>PCV%</th>
<th>Hb</th>
<th>Serum Protein</th>
<th>WBC's per mm$^3$</th>
<th>Neutrophils</th>
<th>Lymphocytes</th>
<th>Monocytes</th>
<th>Eosinophils</th>
<th>Basophils</th>
<th>Significant Intercurrent Pathology</th>
</tr>
</thead>
<tbody>
<tr>
<td>77W577</td>
<td>5/4/77</td>
<td>3.0</td>
<td>CM</td>
<td>38</td>
<td>13.6</td>
<td>6.5</td>
<td>2,300</td>
<td>70</td>
<td>24</td>
<td>4</td>
<td>1</td>
<td>1</td>
<td>None</td>
</tr>
<tr>
<td>78W23</td>
<td>2/18/78</td>
<td>3.7</td>
<td>F</td>
<td>37</td>
<td>12.6</td>
<td>6.0</td>
<td>3,200</td>
<td>69</td>
<td>26</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>Typhilitis</td>
</tr>
<tr>
<td>78W68</td>
<td>5/14/78</td>
<td>3.0</td>
<td>CM</td>
<td>53</td>
<td>17.6</td>
<td>8.4</td>
<td>7,000</td>
<td>70$^b$</td>
<td>24</td>
<td>6</td>
<td>0</td>
<td>0</td>
<td>Bronchopneumonia</td>
</tr>
<tr>
<td>78W180</td>
<td>10/12/78</td>
<td>3.3</td>
<td>CM</td>
<td>12$^c,d$</td>
<td>4.0$^c$</td>
<td>6.2</td>
<td>18,000</td>
<td>81</td>
<td>18</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>Suppurative arthritis</td>
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<tr>
<td>79W93</td>
<td>2/26/79</td>
<td>3.7</td>
<td>F</td>
<td>35</td>
<td>12.6</td>
<td>5.9</td>
<td>3,900</td>
<td>52</td>
<td>46</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>Aspiration pneumonia</td>
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<tr>
<td>79W193</td>
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<td>F</td>
<td>37</td>
<td>14.4</td>
<td>8.5</td>
<td>14,000</td>
<td>82</td>
<td>14</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>Multiple abscesses</td>
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<tr>
<td>79W200</td>
<td>3/9/79</td>
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<td>F</td>
<td>40</td>
<td>15.4</td>
<td>7.2</td>
<td>5,800</td>
<td>81</td>
<td>13</td>
<td>4</td>
<td>2</td>
<td>0</td>
<td>None</td>
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<tr>
<td>79W203</td>
<td>3/13/79</td>
<td>2.8</td>
<td>CM</td>
<td>59</td>
<td>22</td>
<td>7.0</td>
<td>8,000</td>
<td>83</td>
<td>17</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>Enteritis</td>
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<tr>
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<td>2.9</td>
<td>F</td>
<td>43</td>
<td>18</td>
<td>5.9</td>
<td>7,100</td>
<td>72</td>
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<td>0</td>
<td>0</td>
<td>0</td>
<td>None</td>
</tr>
<tr>
<td>79W258</td>
<td>5/16/79</td>
<td>4.0</td>
<td>F</td>
<td>37</td>
<td>12.6</td>
<td>5.7</td>
<td>40,600$^{c}$</td>
<td>73</td>
<td>26</td>
<td>1</td>
<td>0</td>
<td>0</td>
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<tr>
<td>79W314</td>
<td>8/17/79</td>
<td>3.1</td>
<td>F</td>
<td>31</td>
<td>10.8</td>
<td>5.1</td>
<td>3,400</td>
<td>70</td>
<td>28</td>
<td>2</td>
<td>0</td>
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<td>None</td>
</tr>
<tr>
<td>Mean</td>
<td></td>
<td>3.4</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td>0.2</td>
</tr>
<tr>
<td>Range</td>
<td></td>
<td>2.8-4.0</td>
<td></td>
<td>31-59</td>
<td>10.8-22</td>
<td>5.1-8.5</td>
<td>2,300-40,800</td>
<td>52-83</td>
<td>13-46</td>
<td>0.6</td>
<td>0.2</td>
<td>0.1</td>
<td></td>
</tr>
</tbody>
</table>

$^a$F=female, CM=castrated male  
$^b$23% immature neutrophils  
$^c$excluded from mean calculations  
$^d$regenerative anemia
TABLE 3. Serum chemistry and electrolyte profile of mule deer with histopathologically confirmed CWD.

<table>
<thead>
<tr>
<th>Deer number</th>
<th>BUN mg/dl</th>
<th>Globulin g/dl</th>
<th>Cholesterol mg/dl</th>
<th>Total Bilirubin mg/dl</th>
<th>Creatinine mg/dl</th>
<th>Alk. phos. (U/L)</th>
<th>SGOT (U/L)</th>
<th>CPK (U/L)</th>
<th>Inorganic Phosphorus mg/dl</th>
<th>Calcium mg/dl</th>
<th>Chloride mg/dl</th>
<th>Sodium mg/dl</th>
<th>Potassium mg/dl</th>
<th>Significant Intercurrent Pathology</th>
</tr>
</thead>
<tbody>
<tr>
<td>77W577</td>
<td>23</td>
<td>4.2</td>
<td>70</td>
<td>0.2</td>
<td>1.6</td>
<td>5</td>
<td>100</td>
<td>156</td>
<td>5.5</td>
<td>9.4</td>
<td>106</td>
<td>144</td>
<td>144</td>
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<tr>
<td>77W32</td>
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<td>3.4</td>
<td>57</td>
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<td>8</td>
<td>100</td>
<td>48</td>
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<td>9.4</td>
<td>100</td>
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<td>143</td>
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<tr>
<td>77W288</td>
<td>115</td>
<td>5.4</td>
<td>100</td>
<td>0.8</td>
<td>4.2</td>
<td>18</td>
<td>150</td>
<td>ND</td>
<td>11.0</td>
<td>10.0</td>
<td>100</td>
<td>100</td>
<td>146</td>
<td>Bronchopneumonia Suppurative arthritis</td>
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<td>50</td>
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<td>1.0</td>
<td>12</td>
<td>190</td>
<td>ND</td>
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<td>7.5</td>
<td>104</td>
<td>142</td>
<td>142</td>
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<tr>
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<td>12</td>
<td>130</td>
<td>44</td>
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<td>8.8</td>
<td>107</td>
<td>142</td>
<td>142</td>
<td>Multiple abscesses</td>
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<tr>
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<td>144</td>
<td>144</td>
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<td>38</td>
<td>1420</td>
<td>147</td>
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<tr>
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<td>5</td>
<td>150</td>
<td>280</td>
<td>9.2</td>
<td>8.2</td>
<td>103</td>
<td>143</td>
<td>4.6</td>
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<td>0.2</td>
<td>2.2</td>
<td>7</td>
<td>150</td>
<td>150</td>
<td>5.8</td>
<td>9.4</td>
<td>100</td>
<td>143</td>
<td>7.2</td>
<td>Abomasal ulcer</td>
</tr>
<tr>
<td>77W314</td>
<td>70</td>
<td>3.1</td>
<td>35</td>
<td>0.4</td>
<td>1.8</td>
<td>13</td>
<td>140</td>
<td>59</td>
<td>6.4</td>
<td>7.6</td>
<td>113</td>
<td>145</td>
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<td>None</td>
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<tr>
<td>Mean</td>
<td>44</td>
<td>4.0</td>
<td>55</td>
<td>0.4</td>
<td>2.0</td>
<td>12.9</td>
<td>261</td>
<td>114</td>
<td>7.0</td>
<td>9.0</td>
<td>106</td>
<td>144</td>
<td>5.5</td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>16-115</td>
<td>2.9-5.4</td>
<td>35-100</td>
<td>0.2-0.8</td>
<td>1.0-4.2</td>
<td>5.38</td>
<td>80-1420</td>
<td>34-280</td>
<td>2.7-11.0</td>
<td>7.5-10.6</td>
<td>100-116</td>
<td>142-147</td>
<td>3.7-9.0</td>
<td></td>
</tr>
</tbody>
</table>

*International units

\(\text{ND} = \text{Not determined}\)
alopecia and external or internal parasitism (Table 1).

HISTOPATHOLOGY
An extensive selection of formalin-fixed tissues examined by light microscopy of hematoxylin-eosin stained sections from 41 affected deer did not result in identification of consistent pathologic changes in extra-neural organs or systems. Brain or other tissues of the central nervous system (CNS) were removed from 29 of these cases and fixed in neutral buffered 10% formalin solution or in formalin-ammonium bromide (FAB). Tissue sections representing various regions of the CNS were processed for paraffin embedment, sectioned at 6-7 μm and stained with hematoxylin-eosin, periodic acid-Schiff, luxol fast blue-creosyl echt violet, oil red-0, Holzer's or Weil-Weigert methods. Selected FAB-fixed tissues were sectioned at 15-20 μm on a freezing microtome and stained by the Ramon Y Cajal's gold sublimate method for astrocytes.

Neurohistopathologic changes were characterized by widespread microcavitation or spongiform transformation of the neuropil, predominantly of gray matter (Fig. 2), single or multiple intracytoplasmic vacuoles in neuronal perikaryons (Fig. 3) and by neuronal degeneration. Spongiform change of varying severity was found in the gray matter of spinal cord, medulla oblongata, pons, mesencephalon, thalamus, hypothalamus and cerebellar cortex. Focal lesions occurred in cerebral cortex, particularly in olfactory regions of the frontal lobes. Specific nuclei of thalamus, mesencephalon and medulla oblongata were consistently and severely affected. In such areas, astrocytic hypertrophy and fibrillary hyperplasia were demonstrable in gold sublimate stained sections (Fig. 4). These were matched for comparison with identical regions of the brains of unaffected mule deer of similar age in which these changes were absent.

<table>
<thead>
<tr>
<th>TABLE 4. Cerebrospinal fluid (CSF) values for mule deer with histopathologically confirmed CWD&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Deernumber</th>
<th>Color</th>
<th>Appearance</th>
<th>Specific gravity</th>
<th>pH</th>
<th>Total protein (mg/dl)</th>
<th>Cells</th>
<th>8 lympha</th>
<th>10 cells</th>
<th>100 cells</th>
<th>3 monocyt</th>
<th>No cells</th>
<th>3 eosininflammatory cells</th>
</tr>
</thead>
<tbody>
<tr>
<td>77W/577</td>
<td>Colorless</td>
<td>Clear</td>
<td>1.005</td>
<td>6.5</td>
<td></td>
<td>22</td>
<td></td>
<td>0.06</td>
<td>ND</td>
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<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>77W/599</td>
<td>Slightly bloody</td>
<td>Clear</td>
<td>1.005</td>
<td>8.0</td>
<td></td>
<td>ND</td>
<td></td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
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</tr>
<tr>
<td>79W/259</td>
<td>Colorless</td>
<td>Clear</td>
<td>1.006</td>
<td>8.0</td>
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<td>ND</td>
<td></td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
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<td>79W/258</td>
<td>Slightly bloody</td>
<td>Clear</td>
<td>1.006</td>
<td>8.0</td>
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<td>ND</td>
<td></td>
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</tr>
</tbody>
</table>

<sup>a</sup>CSF sample taken immediately following euthanasia by electrocution
<sup>b</sup>ND = not determined
DISCUSSION

Morbidity and mortality from CWD have not been accurately recorded among mule deer in these facilities during the period 1967-74. From 1974-79 when accurate records are available, 54 (80%) of a total population of 67 mule deer and black-tailed deer held or presently in captivity for more than 2 years have shown typical clinical features of CWD and died or were euthanatized as a consequence of the disease.

The only alteration in clinical pathologic values for deer affected with CWD which we consider significant is low urine specific gravity. Degenerative encephalopathic changes are consistently identified in the hypothalamus, which is important in the regulation of antidiuretic hormone, and may be responsible for the diabetes insipidus-like clinical syndrome.

The insidious onset, protracted clinical course, fatal outcome and characteristic encephalopathic features of CWD of mule deer are qualitatively comparable to other established spongiform encephalopathies of animals and man. Spontaneously occurring diseases of this group include scrapie of sheep and goats, transmissible encephalopathy of mink, and kuru and Creutzfeld-Jakob disease of man. These represent widely acknowledged and demonstrably transmissible examples of “slow virus” infections which result in classical...
FIGURE 3. Intracytoplasmic vacuolation of neuronal perikaryons in the pons (×100, 7 μm section, HE).

FIGURE 4. Astrocytic hypertrophy and fibrillary hyperplasia (×160, 15 μm frozen section, Cajal's gold chloride impregnation).
spongiform encephalopathy. Recently a neurotropic retrovirus has been identified in wild mice which is associated with similar noninflammatory spongiform polioencephalopathy.2,4,5 Currently, studies are underway to characterize fully the clinical and pathologic features of CWD of mule deer, and to determine its epizootiology and transmissibility.

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LITERATURE CITED

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