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European Brown Hare Syndrome in Wild European Brown Hares from **Greece**

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ABSTRACT: From 1999 to mid-2003, 97 European brown hares (Lepus europaeus) found dead throughout Greece were examined by necropsy, histopathology, and reverse-transcription polymerase chain reaction (RT-PCR) for the presence of European brown hare syndrome (EBHS) and EBHS virus (EBHSV), respectively. Hare losses were sporadic, starting in the cold season and lasting for many months (December to May). The most prominent gross lesions were observed in the liver and included swelling and discoloration; congestion and hemorrhages were present mainly in lungs and tracheal mucosa. Necropsy findings were suggestive of EBHS, which was confirmed by histopathology and RT-PCR. This study documents, for the first time, EBHS in Greece.

Key words: European brown hare syndrome, Greece, histopathological findings, PCB

European brown hare syndrome (EBHS) affects wild and farmed hares of the species Lepus europaeus and Lepus timidus. The syndrome, a severe necrotic hepatitis, has affected up to 100% of a hare population in which animals usually died within 48–72 hr (Chasey et al., 1992). The disease was first reported during the 1980s and occurred simultaneously in many European countries (Morisse, 1988; Eskens and Volmer, 1989; Henriksen et al., 1989; Marcato et al., 1989; Okerman et al., 1989; Chasey and Duff, 1990; Sostaric et al., 1991; Gavier-Widen and Morner, 1993; Nauwynck et al., 1993; Salmela et al., 1993; Steineck and Nowotny, 1993; Gortazar and de Luco, 1995; Frolich et al., 1996, 2001, 2003; Slamecka et al., 1997). The viral etiology was first demonstrated in 1988 by electron microscopy (Lavazza et al., 1988). Sequencing the viral genome has demonstrated EBHS virus (EBHSV)

shares a similar genomic organization to rabbit hemorrhagic disease virus (RHDV), the cause of rabbit hemorrhagic disease (RHD) but that the two viruses are distinct caliciviruses (Wirblich et al., 1994; Le Gall et al., 1996).

European brown hare syndrome has significant similarities to RHD in its epidemiology, clinical signs, and pathology. Findings in the two diseases are characterized by rapid progression, mild nervous symptoms, severe necrotic hepatitis, and circulatory dysfunction in various organs (Capucci et al., 1991; Duff et al., 1994). Our objective was to determine whether EBHS occurs in Greece where an increased mortality in European brown hare populations has been observed in recent years.

Ninety-seven European brown hares found dead or shot throughout Greece were submitted by hunting federations from 1999 to mid-2003. All hares were adults and in fair or poor body condition. In five hares, postmortem findings were unrecognizable because of advanced putrefaction. Gross lesions were found in 85 hares. Edema, congestion, extensive hemorrhages in the lungs, and severe congestion of the tracheal mucosa were observed. The spleen was enlarged and dark redblack, and the kidneys were congested. In 83, the liver was swollen, discolored, and friable. Tissue samples from liver, lungs, spleen, and kidneys from all except the five decomposed hares were fixed in 10% buffered formalin and routinely processed for histopathologic examination. In addition, samples of liver and spleen from all hares

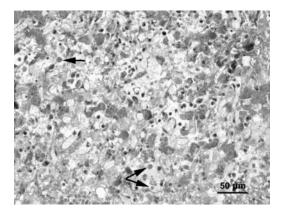
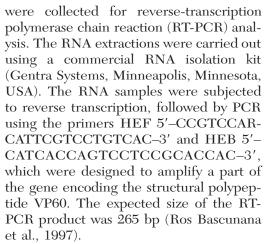


FIGURE 1. Liver. Periportal and midzonal necrosis with a mild inflammatory infiltration of mononuclear cells. Notice the acidophilic bodies (arrows). H&E stain.



Histopathologically, in the 85 hares with gross lesions, interstitial and alveolar hemorrhages and edema were observed in the lungs; and in the spleen, the red-pulp was congested and edematous. In 83, acute hepatic lesions, characterized by variably sized areas of periportal and midzonal hepatocellular necrosis, were the most prominent and consistent histopathologic findings. Acidophilic bodies were occasionally seen around or in the necrotic areas in connection with hepatocyte degeneration and calcium deposits (Fig. 1). Three of the seven hares with no gross lesions had a mild periportal inflammatory reaction, prominent periportal fibroplasia, and bile duct proliferation accompanied by peri-

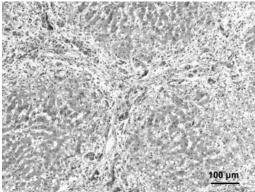


FIGURE 2. Liver. Fibrous tissue infiltrated with lymphocytes has formed in and around the portal tracts and bridged neighbouring portal tracts. Necrosis and degeneration of hepatocytes are seen. H&E stain

portal degeneration and necrosis of hepatocytes corresponding to chronic hepatitis (Fig. 2).

European brown hare syndrome virus RNA was detected in liver and spleen specimens from the 83 hares with acute hepatic lesions, the three hares with chronic hepatitis, and in spleen from the two hares with changes limited to the lung and spleen. In our study, EBHSV-positive hares were found most frequently between December and May; this observation also has been described by Poli et al. (1991).

Gross and histopathologic findings in these hares were consistent with EBHS, which was confirmed by RT-PCR. On the basis of the morphology of the liver changes, we recognized the two forms of the disease as described by Gavier-Widen (1994), an acute and a subacute form. Furthermore, we found alterations in the liver of three hares corresponding to the chronic form of the disease (Marcato et al., 1991). Detection of viral RNA in this form of the disease may imply that these animals could play a role in the epidemiology of EBHS by spreading and maintenance of the virus.

In conclusion, we have demonstrated that EBHS exists in Greece and may be one of the causes for increased mortality in the hare population in Greece over the past 6 yr. However, the ecological impact of EBHS needs to be further evaluated in conjunction with other factors, such as changes in habitat, agricultural techniques, and environmental pollution.

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