

Isolation of Brucella melitensis Biovar 3 from a Chamois (Rupicapra rupicapra) in the Southern French Alps

Authors: Garin-Bastuji, Bruno, Oudar, Jean, Richard, Yves, and

Gastellu, Jacques

Source: Journal of Wildlife Diseases, 26(1): 116-118

Published By: Wildlife Disease Association

URL: https://doi.org/10.7589/0090-3558-26.1.116

BioOne Complete (complete.BioOne.org) is a full-text database of 200 subscribed and open-access titles in the biological, ecological, and environmental sciences published by nonprofit societies, associations, museums, institutions, and presses.

Your use of this PDF, the BioOne Complete website, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at www.bioone.org/terms-of-use.

Usage of BioOne Complete content is strictly limited to personal, educational, and non - commercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

BioOne sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.

Isolation of *Brucella melitensis* Biovar 3 from a Chamois (*Rupicapra rupicapra*) in the Southern French Alps

Bruno Garin-Bastuji, 'Jean Oudar, 'Yves Richard, 'and Jacques Gastellu,' 'Centre National d'Etudes Vétérinaires et Alimentaires, L.C.R.V., B.P. 67, 94703 Maisons-Alfort, France; 'Ecole Nationale Vétérinaire de Lyon, B.P. 83, 69280 Marcy l'Etoile, France

ABSTRACT: Systemic brucellosis caused by Brucella melitensis biovar 3 was identified in a chamois (Rupicapra rupicapra) near the Parc National des Ecrins in the southern French Alps (France). Clinical signs included orchiepididymitis, polyarthritis, blindness and various neurological signs; necropsy findings included numerous calcified foci in testis, epididymis, kidney, subcutaneous connective tissue and brain. Brucella sp. were identified in brain by indirect immunofluorescence and B. melitensis biovar 3 was isolated from testis, kidney, eye, lung and joints. This report describes the first case of brucellosis and Brucella sp. isolation in chamois in France and the first case of B. melitensis isolation in this host species.

Key words: Brucellosis, Brucella melitensis biovar 3, chamois, Rupicapra rupicapra, isolation, case report.

Brucellosis is a highly contagious infection of many animal species including man and is caused by bacteria of the genus Brucella. Well known throughout the world in domestic ruminants, brucellosis has been reported in numerous wild species (Witter, 1981; Comité Mixtre FAO/OMS, 1986). It has been reported from roe deer (Capreolus capreolus), chamois (Rupicapra rupicapra), ibex (Capra ibex), and several species of deer (Cervus spp.) in Europe (Pastoret et al., 1988). In North America, brucellosis has been reported in caribou (Rangifer tarandus), elk (Cervus elaphus nelsoni), moose (Alces alces) and bison (Bison bison) (Moore and Schnurrenberger, 1981; Tessaro, 1986). Saïga (Saiga tatarica) has been found infected in the Union of Soviet Socialist Republics (Renoux, 1957). In most cases, diagnosis was by serology and confirmed only a few times by isolation of Brucella abortus, B. melitensis or B. suis. Infected wildlife usually cohabited with domestic ruminants with high prevalence of the disease.

In chamois, the disease was first reported in the 1950's in Switzerland in a few specimens and *B. abortus* was the only species isolated (Burgisser, 1952; Bouvier et al., 1954; Bouvier, 1961). Recently, several serological surveys were conducted in the mountains of France. A few isolated cases were identified by serology in chamois in the southern Alps (Baradel et al., 1988) and no case was described in isard (*Rupicapra rupicapra pyrenaica*) in the Pyrenees mountains (Candoussau-Luquet, 1987).

In North America and Europe, bovine, caprine and ovine brucellosis are usually characterized by abortion and orchitis (Comité Mixte FAO/OMS, 1986). The reported signs in chamois and roe deer, however, included orchitis, polyarthritis, and/or ocular (uveitis, blindness) and neurologic (ataxia) signs (Bouvier et al., 1954). Necropsy findings included widespread abscesses with thick pus evolving to mineralization (Burgisser, 1952; Bouvier et al., 1954).

This report is the first description of brucellosis and *Brucella* sp. isolation in chamois in France and the first case of *B. melitensis* infection recognized in this species.

In December 1988, a blind and ataxic 5-yr-old male chamois was caught on a road near Col du Lautaret in the southern French Alps (45°2'37"N, 6°25'31"E). The animal was transferred to the Veterinary School of Lyon (Marcy-l'Etoile, France) where it was examined and slaughtered. Clinical signs included bilateral keratoconjunctivitis and uveitis with blindness and nystagmus, polyarthritis including hocks, shoulders, carpus, sternal joints, etc., and orchiepididymitis. Necropsy findings included widespread calcified nodules in

subcutaneous connective tissue; orchiepididymitis with calcified or necrotic foci; calcified nodules and infarctions in the renal cortex; a small focus of chronic pleuritis and atelectasis of the right apical lung lobe; and serofibrinous and hemorrhagic polyarthritis. Histopathologic observations included: testicular cessation of spermatogenesis and necrosis of seminal epithelia around necrotic and calcified zones of parenchyma: necrotic and calcified zones associated with fibrosed infarctions of renal cortex; severe widespread corneal ulcerations, vascularization of limbus, fibrinopurulent exudate in the anterior chamber, and lymphoplasmacytic infiltration of iris and ciliary bodies; and degeneration, necrosis and calcification with perivascular lymphoplasmacytic and granulomatous inflammation in brain and pia mater. Brucella sp. were revealed in brain formalin-fixed sections by indirect immunofluorescence using fluorescein isothiocyanate-conjugated sheep anti-rabbit IgG (H + L) purified antibodies (Byosis, Paris, France). Smooth Brucella sp. antiserum was prepared as described by Corbel et al. (1983) except that a mixture of B. abortus strain 544 and B. melitensis strain 16 M (1 × 105 CFU of each strain per rabbit) was used as antigen.

Strongly positive results were obtained with standard serological tests including rose bengale plate test, complement fixation test and seroagglutination test (Alton et al., 1988). Bacteriological examinations were performed according to Alton et al. (1988) and staining methods (modified Ziehl-Nielsen and Köster's methods) revealed Brucella sp.-like organisms on fresh tissue imprints of orchitis lesions. Brucella sp. were isolated from testis, kidney, lung, eye and various joint fluids and were all biotyped as B. melitensis biovar 3. This biovar has been the most prevalent in domestic cattle, sheep and goats for over 10 yr in this region of France (Verger et al., 1989) where chamois populations usually mingle with domestic ruminant herds during summer. A preliminary epidemiological survey was conducted among the regional populations (1,200 specimens) of chamois but to date no clinical case has been found. Moreover, the breeding success stayed stable during recent years.

We thank J. M. Verger, C. Fleury, E. Richard, C. Sarrazin, A. M. Mahé and R. Keck for their help in epidemiological, microbiological and histopathologic observations.

LITERATURE CITED

- ALTON, G. G., L. M. JONES, R. D. ANGUS, AND J. M. VERGER. 1988. Techniques for the brucellosis laboratory. Institut National de la Recherche Agronomique, Paris, France, 190 pp.
- BARADEL, J. M., J. BARRAT, J. BLANCOU, J. M. BOUTIN, C. CHASTEL, G. DANNACHER, D. DELORME, Y. GERARD, J. M. GOURREAU, U. KHIM, B. LARENAUDIE, C. LE GOFF, P. P. PASTORET, P. PERREAU, A. SCHWERS, E. THIRY, D. TRAP, G. UILENBERG, AND P. VANNIER. 1988. Bilan d'une enquête sérologique effectuée sur différents mammifères sauvages de France. In Maladies de la faune sauvage. Revue Scientifique et Technique de l'Office International des Epizooties 7: 861–872.
- BOUVIER, G. 1961. Les possibilités de transmission de la brucellose du gibier. Schweizerische Medizinische Wochenschrift 91: 827-828.
- ———, H. BURGISSER, AND P. A. SCHNEIDER. 1954. Lésions oculaires d'un chamois dues à *Brucella abortus*. Schweizer Archiv für Tierheilkunde 96: 85–89.
- BURGISSER, H. 1952. Constatations sur la brucellose génitale du chamois. Schweizer Archiv für Tierheilkunde 94: 554–556.
- CANDOUSSAU-LUQUET, P. 1987. Pathologie de l'isard. Thèse de Doctorat Vétérinaire, Toulouse, France. 128 pp.
- COMITE MIXTE FAO/OMS D'EXPERTS DE LA BRU-CELLOSE. 1986. VIéme rapport, Séries de Rapports Techniques 740, Organisation Mondiale de la Santé, Genève, Switzerland, pp. 56–57.
- CORBEL, M. J., K. P. W. GILL, AND E. L. THOMAS. 1983. Methods for the identification of *Brucella*. Ministry of Agriculture, Fisheries and Food publications, Booklet No. 2085, Alnwick, United Kingdom, 18 pp.
- MOORE, C. G., AND P. R. SCHNURRENBERGER. 1981. A review of naturally occurring *Brucella abortus* infections in wild mammals. Journal of the American Veterinary Medical Association 179: 1105–1112.
- PASTORET, P.-P., E. THIRY, B. BROCHIER, A. SCHWERS, I. THOMAS, AND J. DUBUISSON. 1988. Maladies de la faune sauvage transmissibles aux animaux

- domestiques. In Maladies de la faune sauvage. Revue Scientifique et Technique de l'Office International des Epizooties 7: 661–704.
- Renoux, G. 1957. La brucellose des animaux sauvages et des insectes. Archives de l'Institut Pasteur de Tunis 34: 391-404.
- Tessaro, S. V. 1986. The existing and potential importance of brucellosis and tuberculosis in Canadian wildlife: A review. The Canadian Veterinary Journal 27: 119–124.
- VERGER, J. M., B. GARIN-BASTUJI, M. GRAYON, AND A. M. MAHE. 1989. La brucellose bovine à *Bru*cella melitensis en France. Annales de Recherches Vétérinaires 20: 93–102.
- WITTER, J. F. 1981. Brucellosis. In Infectious diseases of wild mammals. J. W. Davis, L. H. Karstad and D. O. Trainer (eds.). Iowa State University Press, Ames, Iowa, pp. 280-287.

Received for publication 26 May 1989.