

Infectious agents identified by real-time PCR, serology and bacteriology in blood and peritoneal exudate samples of cows affected by parietal fibrinous peritonitis after caesarean section

Djebala S.¹, Evrard J.², Gregoire F.², Thiry D.³, Bayrou C.¹, Moula N.⁴, Sartelet A.¹, Bossaert P.¹

¹. Clinical Department of Ruminants, University of Liège, Quartier Vallée 2, Avenue de Cureghem 7A-7D, Liège 4000, Belgium

². Gestion et Prévention de Santé, Regional Association of Health and Animal Identification, Allée des Artisans 2, Ciney 5590, Belgium

³. Bacteriology, Department of Infectious and Parasitic Diseases, University of Liège, quartier vallée 2, avenue Cureghem 6, B-4000 Liège, Belgium

⁴. Department of animal production, University of liege, Quartier Vallée 2, Avenue de Cureghem 6, Liège 4000, Belgium

Corresponding author: sdjebala@uliege.be

Abstract: The aim of the current study was to identify the pathogens that are potentially involved in parietal fibrinous peritonitis (PFP). PFP is a relatively common complication of cattle celiotomy, characterized by an accumulation of exudate in a fibrinous capsule within the abdominal or pelvic cavity; its aetiology is poorly understood.

We have studied 72 cases of PFP, confirmed by a standard diagnostic protocol. Blood were collected to evaluate the presence of antibodies for *Mycoplasma bovis* (*M. bovis*), *Coxiella burnetii* (*C. burnetii*) and *Bovine Herpesvirus 4* (*BoHV4*) by enzyme-linked immunosorbent assays. Peritoneal exudate samples were obtained from the PFP cavity to perform bacteriological culture, and to identify the DNA of *M. bovis*, *C. burnetii* and *BoHV4* using real time polymerase chain reaction (qPCR).

Bacteriological culture was positive in most of peritoneal samples (59/72); *Trueperella pyogenes* (*T. pyogenes*) (51/72) and *Escherichia coli* (*E. coli*) (20/72) were the most frequently identified. For *BoHV4*, the majority of cows showed a positive serology and qPCR result (56/72 and 49/72, respectively), in contrast to *M. bovis* (17/72 and 6/72, respectively) and *C. burnetii* (15/72 and 6/72, respectively), who were less frequently detected ($p < 0.0001$).

Our study proves that PFP can no longer be qualified as a sterile inflammation, since most PFP samples yielded a positive bacteriology and qPCR. Moreover, we herein describe the first identification of *BoHV4* and *C. burnetii* in cows affected by PFP. The exact role of these germs in the pathogenesis of PFP is not yet elucidated and requires further studies