

Potential for functional replacement of methanogenic bacteria by acetogenic bacteria in the rumen environment

P Boccazzi, Ja Patterson

► To cite this version:

P Boccazzi, Ja Patterson. Potential for functional replacement of methanogenic bacteria by acetogenic bacteria in the rumen environment. Annales de zootechnie, 1996, 45 (Suppl1), pp.321-321. hal-00889643

HAL Id: hal-00889643 https://hal.science/hal-00889643

Submitted on 11 May 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Potential for functional replacement of methanogenic bacteria by acetogenic bacteria in the rumen environment

P Boccazzi, JA Patterson

Department of Animal Sciences, 3-101 Lilly Hall, Purdue University, West Lafayette, Indiana 47907, USA

Ruminal fermentation is influenced by interspecies H₂ transfer in which methanogens enhance energy production by fermentative microorganisms by converting H_2 to CH_4 . However, the energy contained in CH₄ represents a 5 to 15% loss of apparent digestible energy loss to the animal and contributes to global methane emissions. The objective of our research is to determine if alternative H₂-utilizing bacteria, acetogens are capable to replacing the interspecies H₂ transfer function of methanogens when methanogens are inhibited. We have isolated several acetogenic isolates, A10 and ³H, with low hydrogen thresholds (Boccazzi et al, 1993, Proc Conf Rumen Funct, 22-28). The purpose of this research was to determine if they could utilize H₂ in a mixed culture system.

Serum bottles (120 ml) containing 0.35 g ground alfalfa hay (1 mm) were inoculated with 10 ml of a solution consisting of 60% anaerobic dilution solution:40% fresh rumen contents from a cow consuming a 60% concentrate:40% corn silage diet. Bromoethane sulfonic acid (BES, 5 mM) was added to all serum bottles except the controls. Treatments were : control, control + BES, $_{3}$ H + BES and A10 + BES,

where the acetogenic cultures were added to give a final concentration of 4 x 10^8 acetogen cells/ml under a CO₂ atmosphere. Triplicate serum bottles for each treatment and time period were incubated at 37°C and shaken at 200 rpm. Hydrogen and CH₄ concentration in the headspace was measured by gas chromatography using thermal conductivity.

No CH₄ accumulated in serum bottles where BES had been added which indicates that 5 mM BES completely inhibited methanogenesis for up to 74 h. Hydrogen concentrations at 12 h were lower for cultures containing added acetogens. High levels of H₂ in the control of 12 h may have been due to low numbers of methanogens in the inoculum. Concentrations of H₂ were lower for control bottles and bottles containing acetogens at 24 and 74 h. Cultures containing acetogenic isolate ³H consistently had lower H₂ concentrations than cultures containing acetogenic isolate A10. These data show that acetogens can functionally replace the role of methanogens in interspecies H₂ transfer and that different levels of H₂ control may be obtained by using different acetogenic cultures.

Treatment		Η ₂ (μΜ)		CH ₄ (μM)	
	0h	12h	24h	74h	74h ¹
Control	10	166	16	0.5	258
Control + BES	8	171	112	23	0.8
3H + BES	8	48	12	2	1
A10 + BES	9	108	44	21	3

¹ Initial CH₄ concentrations were less than 2 µM.