Appropriateness of reference genes for normalizing messenger RNA in mouse 2,4-dinitrobenzene sulfonic acid (DNBS)-induced colitis using quantitative real time PCR

Nour Eissa^{1,2*}, Laëtitia Kermarrec^{1*}, Hayam Hussein³, Charles N. Bernstein^{4,5}, Jean-Eric Ghia^{1,2,4,5}

¹Immunology, University of Manitoba, Winnipeg, MB, Canada

²Children's Hospital Research Institute of Manitoba, University of Manitoba, Winnipeg, MB, Canada

³Large Animal Medicine, William R. Pritchard Veterinary Medical Teaching Hospital, School of Veterinary Medicine, University of California Davis, CA, USA

⁴Internal Medicine section of Gastroenterology, University of Manitoba, Winnipeg, MB, Canada

⁵IBD Clinical and Research Centre, University of Manitoba, Winnipeg, MB, Canada

* First co-authorship, authors contributed equally

Corresponding address:

Dr. Jean-Eric Ghia, Department of Immunology, College of Medicine, University of Manitoba, 431 Apotex Centre, 750 McDermot Avenue, Winnipeg, MB R3E 0T5 Canada Tel: +1 (204) 789-3802, Fax: +1 (204) 789-3921 Email: Jean-Eric.Ghia@umanitoba.ca or jeghia@yahoo.fr

Supplementary figures legend

Figure S1. Amplification efficiency of primers reference gene candidates. The X-axis denotes the log10 cDNA dilution series, and the Y-axis denotes the cycle threshold (Ct). The primer efficiency (E) is computed by $[10(1/-S)-1] \times 100\%$, where S represents the slope of the linear regression line. ROCH Light Cycler Software was used for analysis.

