B6.129P2–Il10 tm1Cgn/J

Additional Use Restrictions Apply

002251 | Il-10 KO

Congenic, Targeted Mutation

AVAILABLE NOW

B6.129P2–Il10 tm1Cgn/J mutant mice spontaneously develop a chronic inflammatory bowel disease (IBD). Il10-deficiency is associated with altered lymphocyte and myeloid profiles, elevated serum amyloid A levels, altered responses to inflammatory or autoimmune stimuli, increased prevalence of colorectal adenocarcinoma, and spontaneous development of chronic enterocolitis. Il10-deficient mice exhibit a significant increase in peripheral blood granulocyte populations upon lesion development.

Genetic overview

C57BL/6

N13F10
(05-AUG-14)
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