



## Effects of colloidal bismuth tartrate on colitis induced by immune-complex in rabbits

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### Abstract

**AIM:** To observe the therapeutic effect of colloidal bismuth tartrate in an animal colitis model.

**METHODS:** Immune-complex colitis was induced in groups of rabbits by formalin, and two hours later 0.85 mL heat aggregated rabbit IgG was given intravenously through the ear cannula. Animals were intracolonicly treated with colloidal bismuth tartrate (BITNAL), and its effect was compared with sulfasalazine (SASP), indomethacin (IND) and bifidobiogen (BIFG). Animals were killed, the mucosal appearance was scored (0-4), and tissue saved for histological studies, the number of neutrophils present in inflamed colonic tissue was quantitated by the myeloperoxidase (MPO) activity assay, the

production of lipoxygenase and cyclo-oxygenase products was monitored and eicosanoid production were assayed by incubation colonic specimens and the media for prostaglandin E2 (PGE2), leukotriene (LTB4), thromboxane B2 (TXB2) were examined by radiomunoassay.

**RESULTS:** Immune complex colitis was induced by formalin and IgG, colonic damage persisted for at least 1 wk by macrography. Histologically, the inflammatory response included mucosal and submucosal infiltration by polymorphonuclear leukocytes, macrophages, lymphocytes and fibroblasts, the macroscopic, percent 2 wk after IgG, was correlated with greatly increased PGE2, LTB4 and TXB2 compared with levels in controls. Treatment with BITNAL (500 mg/kg) resulted in a lowered inflammation index, lowered MPO activity and inhibited the increased formation of PGE2, LTB4 and TXB2 by the inflamed colon, and IND (500 mg/kg) markedly inhibited prostanoid formation in both inflamed and control colon but did not reduce tissue damage, SASP (500 mg/kg) also inhibited the formation of PGE2, LTB4 and TXB2 but the effects were less marked. BIFG (400 mg/kg) did not significantly reduce the colonic injury and the media synthesized by the rabbit colon.

**CONCLUSION:** BITAL provides better therapeutic effects in experimental colitis than anti-inflammatory drug IND or SASP.

**Key words:** Colitis/therapy; Bitnal; Rabbits; Models; Formaldehyde; Immunoglobulins

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