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## Effect of an aqueous extract of *Ajuga iva* on glycaemia, reverse cholesterol transport and atherogenic ratios in rats with streptozotocin-induced diabetes

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Experimental evidence suggests that hyperglycaemia is commonly associated with hyperlipidaemia<sup>(1,2)</sup>. The present study was undertaken to investigate the effect of an aqueous extract of *Ajuga iva* L. Schreiber (Lamiaceae; *Ai*) on blood glucose, serum and lipoprotein lipid profiles and lecithin:cholesterol acyltransferase (LCAT) activity in rats with streptozotocin-induced diabetes.

Twelve rats with diabetes were divided into two groups that were fed a casein diet either with or without an *Ai* supplement (5 g/kg diet) for 4 weeks. Experimental diabetes was induced by intraperitoneal injection of streptozotocin as a single dose of 60 mg/kg body weight. HDL subfractions were separated by differential dextran sulphate–MgCl<sub>2</sub> precipitation and LCAT activity was determined by conversion of [<sup>3</sup>H]cholesterol (unesterified; UC) to [<sup>3</sup>H]cholesteryl esters (CE).

*Ai* treatment significantly decreased glycaemia (–41%) and liver total cholesterol (TC; –33%), TAG (–30%) and phospholipids (PL; –47%). In the *Ai*-treated rats compared with the untreated rats hypocholesterolaemia (–33%) and hypotriacylglycerolaemia (–72%) were observed with a concomitant reduction in LDL-HDL<sub>1</sub>-cholesterol (–50%), VLDL-cholesterol (–56%) and VLDL-TAG, whereas HDL-cholesterol remained unchanged for both groups. Moreover, plasma apoB concentration was 2-fold lower, while that of apoA was 2.4-fold higher.

Group.....	Untreated		<i>Ai</i> -treated	
	Mean	SE	Mean	SE
LCAT (nmol/ml per h)	10.0	1.93	14.86*	1.29
ApoA (g/l)	0.98	0.12	2.31***	0.24
HDL <sub>3</sub> -PL (mmol/l)	0.90	0.12	0.27**	0.08
HDL <sub>3</sub> -UC (mmol/l)	0.21	0.02	0.09**	0.03
HDL <sub>2</sub> -CE (mmol/l)	0.39	0.03	0.56	0.19

Mean values were significantly different from those for the untreated group: \* $P < 0.05$ , \*\* $P < 0.01$ , \*\*\* $P < 0.001$ .

LCAT activity was 1.5-fold higher in the *Ai*-treated rats than in the untreated rats. Moreover, HDL<sub>3</sub>-PL and HDL<sub>3</sub>-UC were decreased by 57% and 70% respectively, whereas HDL<sub>2</sub>-CE was similar for both groups. Also, the atherogenic ratios TC:HDL-cholesterol, VLDL-LDL-cholesterol:HDL-cholesterol and apoB:apoA were decreased by 31%, 46% and 79% respectively in *Ai* treated rats *v.* untreated rats.

These results suggest that *Ai* treatment is effective in decreasing the level of glycaemia and attenuating dyslipidaemia in rats with streptozotocin-induced diabetes by reducing plasma lipids and inversely increasing reverse cholesterol transport.

1. Annida B & Stanely Mainzen Prince P (2004) *J Med Food* 7, 153–156.
2. Tunali S & Yanardag R (2006) *Pharmacol Res* 53, 271–277.