

**Effects of almond consumption on metabolic function and liver fat in overweight and obese adults with elevated fasting blood glucose: A randomised controlled trial.**

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**Abstract:**

Background: Almonds are a rich source of bioactive components. This study examined the effects of daily almond consumption on glycaemic regulation, liver fat concentration and function, adiposity, systemic inflammation and cardiometabolic health. Methods: 76 adults with elevated risk of type 2 diabetes (T2D) or T2D (age:  $60.7 \pm 7.7$  years, body mass index:  $33.8 \pm 5.6$  kg/m<sup>2</sup>) were randomly assigned to daily consumption of either 2 servings of almonds (AS:56 g/day) or an isocaloric, higher carbohydrate biscuit snack (BS) for 8 weeks. Glycosylated haemoglobin (HbA1c), glycaemic variability (GV), liver fat, serum aminotransferases, body weight and composition, markers of cardio-metabolic risk and systemic inflammation were assessed at baseline and week 8. Results: No group differential effects were observed on HbA1c, GV, body weight and composition, liver fat and aminotransferases, cardio-metabolic health and inflammatory markers (all  $P > 0.05$ ). For serum TC/HDL-C ratio a significant gender x treatment x time interaction occurred ( $P < 0.01$ ), such that in women TC/HDL-C ratio was significantly reduced after AS compared to BS ( $-0.36$  [0.26] mmol/L [ $n = 14$ ] vs.  $-0.14$  [0.32] mmol/L [ $n = 17$ ];  $P = 0.05$ ), but not in men ( $P = 0.52$ ). Conclusions: Compared to BS, AS consumed between meals did not substantially alter glycaemic regulation, liver fat or function, adiposity, and metabolic health and inflammatory markers. Serum TC/HDL-C ratio improved in women, but not in men with AS; but as this sub-analysis was not defined a priori the results should be interpreted with caution. Further research should examine the longer-term health effects of regular almond consumption and differential gender responses.