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TORCHLIGHT DOWNLOAD TORCHLIGHT UNLOCKED[Mutation analysis of the fibrinogen gamma-chain gene in a Chinese family with ligneous conjunctivitis]. To identify the mutation in the fibrinogen gamma-chain gene in a Chinese family with ligneous conjunctivitis (LC). Blood samples were obtained from the proband and his family members. DNA was extracted from the proband's leukocytes and those of his family members. The coding sequences of the exons and flanking regions of the fibrinogen gamma-chain gene were amplified by PCR and analyzed by direct sequencing. A novel mutation was identified in the fibrinogen gamma-chain gene in the proband. This mutation, including substitution of guanine for adenine (G2198A), would lead to the amino acid substitution of glycine for arginine (R697K). This mutation was found not only in the proband but also in his father and brother, who were clinically asymptomatic. This study has identified a novel mutation in the fibrinogen gamma-chain gene, which may be associated with LC.

Metabolism of the antimalarial agent, artemisinin, in humans. 1. The pharmacokinetics of artemisinin (sodium artesunate) were investigated in 12 healthy adults. 2. Following oral administration of 1 mg kg⁻¹ of artemisinin, its two major metabolites, dihydroartemisinin and desferriartemzine, were monitored by high performance liquid chromatography and by an immunoassay procedure based on a monoclonal antibody. 3. Plasma concentration profiles of artemisinin and the two metabolites were fitted to a four-compartment open model. Mean (SD) parameters estimated for artemisinin and dihydroartemisinin were as follows: volume of distribution at steady state (V_{ss}), 11.9 (1.6) litre kg⁻¹; clearance (Cl), 20.8 (3.6) ml min⁻¹ kg⁻¹; total body clearance (Cl_{tot}), 23.1 (3.7) ml min⁻¹ kg⁻¹; half-life (t_{1/2}), 0.6 (0.1) h; and area under the plasma concentration time curve (AUC), 36.8 (6.3) mg h litre⁻¹

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