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This is an ongoing study of the clinical, epidemiologic and genetic features of patients with autosomal dominant polycystic liver disease (ADPLD). The investigation involves screening probands of apparently unrelated families for ADPLD. All probands are evaluated clinically to evaluate the probability of ADPLD. If ADPLD is diagnosed, an index case is then screened and ADPLD diagnosed in a second generation family member.

Patients with ADPLD undergo a clinical evaluation and have their liver and kidney evaluated by ultrasound, hepatic venous pressure gradient measurement and liver biopsy. All patients have an ultrasound of the abdomen done at the time of their evaluation for ADPLD. Further testing and evaluation are done as needed including a chest x-ray, bone densitometry and liver biopsy. Genealogical information is collected and tested for a relationship with ADPLD. A registry of patients

with ADPLD is maintained and a patient follow-up program initiated. Current studies are: to identify the spectrum of manifestations of ADPLD; to identify and assess the cause of mortality in patients with ADPLD; to delineate the natural history of ADPLD. It's August. It's the middle of the month, I think, but who can remember. It's the time of year when most book reviewers are updating their schedules to squeeze a couple more months into the season before admitting

defeat and going on holiday. My guess is that, by this time of year, they are all exhausted. It's been a long and tiring summer, and the end of it is approaching. With the autumn and winter around the corner, now is the time to pause for a rest, and possibly a summer holiday. We all have plans to attend or plan to write or plan to do something else. It's a good time to rest, reflect and take stock of what we

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