Primary Ciliary Dyskinesia
(Kartagener syndrome, ICS)

Call for collaboration

• This long-term project (since 1994) seeks to identify and characterize genes involved in PCD

• Sporadic as well as familial cases are sought for this multi-center collaborative study

• Needed for participation in the project:
  -- clinically well-characterized patients (ideally including electron microscopic analysis of cilia)
  -- blood samples from affected individuals; if possible, samples from parents (even for sporadic cases)
  -- your help, to keep patients informed of progress

For additional information, or to receive protocols:

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