Pain Diary

for Autosomal Dominant Polycystic Kidney Disease (ADPKD)



Often, doctors ask patients to describe their pain using a scale from 1 to 10, where 0 is no pain and 10 is extreme pain. This scale also describes how pain is impacting your quality of life and/or ability to complete daily activities. You can use the infographic included in this pain diary to determine how to rate your pain.

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Pain Intensity	0 No pain		-2 Pain	1	3-5 Moderate Pai	'n	6 Sever	-7 e Pain		-9 vere Pain	10 Worst Pain Possible
-	Quality of Life										Non- Functioning
Life (QoL)	Work/ volunteer, normal activities each day Social life outside of work Take an active part in family life	 Work/ volunteer, be active 8 hours daily Outside social activities limited Take part in family life 		Work/volunteer, a few hours daily Active at least 5 hours a day Do simple activities on the weekend	Work/volunteer, limited hours Take part in limited social activities on weekends	Work/volunteer, minimal hours possible Do simple chores around house Minimal activities outside of home 2 days a week	 Not able to work/ volunteer Struggle but fulfill daily home responsibilities No outside activity 	 Get dressed in morning Minimal activities at home Contact with friends via phone, email 	Get out of bed but don't get dressed Stay at home all day	Stay in bed at least half the day Have no contact with outside world	Stay in bed all day Feel hopeless and helpless about life

Link to references

Pain Diary

for Autosomal Dominant Polycystic Kidney Disease (ADPKD) BCRena Provincial Health Services Autosomal Dominant Polycystic Kidney Disease (ADPKD)



Date & Time	Rate pain and impact on quality of life (0-10)	Describe pain (e.g. where is it? how does it feel? how long did it last?)	What triggered the pain?	How did you treat it medication strategie (e.g. rest, exercise, si	How did you feel after treating it? What worked? What didn't work?	
	*see image and descriptions on pg 1			Non-Medications	Medications (if required)	

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Date & Time	Comments, Notes or Questions

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