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Clinical questionnaire for hereditary spastic paraplegia

PATIENT DATA:		CLINICIAN:				
Name + first name: Name		Name + fi	ame + first name:			
Date of birth + gender: H		Hospital:				
DIAGNOSIS age at clinical investigation: years			AGE AT ONSET		years	
				□ 21-4	,	
			□ 1-5 yrs			
			□ 6-10 yrs □ >60 yrs □ 11-20 yrs		yı s	
			u 11-20 j	113		
SEVERITY						
mobility:			ower limbs:	upper limbs:		
normal mobility			normal			
□ abnormal mobility - no aids required			hyperrefle		reflexia	
abnormal mobility - walking aids usedwheelchair			weaknessatrophy	□ weakı □ atrop		
□ wheelchair □ atrophy □ atrophy						
ADDITIONAL FEATURES						
	skin anomalies deafness			□ bladder dysfunction		
	· · · · · · · · · · · · · · · · · · ·			bowel dysfunctionsensory abnormalities:		
□ scoliosis□ foot deformities				sensory abnormalities:		
☐ foot deformities ☐ cognitive impairment ☐ adducted thumbs ☐ epilepsy				o vibration o position o pain o touch o tem	acratura	
distal atrophy lower limbs	, , ,			other (specify):	berature	
□ abnormal saccades/nystagmus □ tremor				•		
□ optic atrophy						
□ cataract	extrapyramida	al signs				
NERVE CONDUCTION STUDIES (age at examination:years) □ report included □ not do					one	
motor nerves						
□ axonal neuropathy □				□ abnor	mal	
EVOKED POTENTIALS (age at examination:years)			report included	□ not d	one	
IEP 🔲 normal		SSEP	SEP normal			
□ abnormal			□ abnorn	nal		
MRI (age at examination:years)			report included	□ not d	one	
Brain normal		Spine	□ norma			
□ abnormal (specify):			□ abnorn	nal (specify):		
NERVE BIOPSY PATHOLOGY (age at examination:years)			report included	□ not d	one	
□ demyelination			other (specify):			
axonal damage						
MUSCLE BIOPSY PATHOLOGY (age at examination:years)			report included	□ not d	one	
□ atrophy			other (specify):			
□ mitochondrial abnormalities						
COMMENTS						