# BJ 6

Prof. Dr. Christine Van Broeckhoven, PhD DSc DNA Diagnostics Unit • Laboratory of Neurogenetics • Institute Born-Bunge University of Antwerp - Campus Drie Eiken • Building V - room 1.33 Universiteitsplein 1 • BE-2610 Antwerpen • Belgium Tel. +32 3 265 10 51 • Fax +32 3 265 10 12 • E-mail DNAdiagnostics@ua.ac.be

## Clinical questionnaire for Charcot-Marie-Tooth disease

PATIENT DATA:	CLINICIAN:
Name + first name:	Name + first name:
Date of birth + gender:	Hospital:

#### DIAGNOSIS

	classical CMT phenotype
	Dejerine-Sottas neuropathy
age at clinical investigation: yrs	congenital hypomyelination

#### SEVERITY

age at onset of symptoms:		mobility:			upper limbs:	
	<1 yrs		normal mobility		normal	
	1-5 yrs		abnormal mobility - no aids required (except		distal weakness	
	6-10 yrs		orthopedic shoes)			
	11-20 yrs		abnormal mobility - walking aids used		(hand-wrist)	
	>20 yrs		wheelchair	<b>u</b>	proximal weakness	

#### ADDITIONAL FEATURES

foot deformities	bulbar dysfunction	other (specify):
scoliosis	diaphragmatic dysfunction	
tremor	vocal cord paralysis	
sensory ataxia	ulcerations	
deafness	nystagmus	
cerebellar dysfunction	pupillary abnormalities	

#### NERVE CONDUCTION STUDIES (age at examination: \_\_\_\_\_\_ years) □ report included □ not done

motor nerves				
upper limb	nerve	NCV	distal latency	amplitude
	nerve	NCV	distal latency	amplitude
lower limb		NCV	distal latency	amplitude
	nerve		distal latency	amplitude
sensory nerves				
	nerve	NCV	distal latency	amplitude
		NCV	distal latency	amplitude
	nerve	NCV	distal latency	amplitude

NERVE BIOPSY PATHOLOGY (age at examination: \_\_\_\_\_\_ years) □ report included

Int done				
demyelination	other (specify):			
axonal damage				

### COMMENTS
