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Informed Consent for Genetic Testing

Document for patient's file of the referring physician

Patient:		Referring Physician:					
Name Prena Date			Name: Address / Stamp:				
		I have received genetic counselli ses (LAGH) and adequate time w	•	o the Swiss federal law on human questions and reflection.			
I he	reby agre	e to have the indicated genetic te	st(s) performe	d:			
Molecular genetic testing for (name of disease):							
	My decision for the sample after the test is completed:						
	If possible,	possible, my samples(s) should be stored for future analysis in my interest, only on my request.					
	My samples(s) may be used anonymously (whithout knowing my name) for medical research purposes, the results of which are withheld and remain confidential.						
	Other:						
Place	e and date, sig	nature of the patient	Place and date, sig	gnature of the referring physician:			
			I have given an a	appropriate explanation of the test to this			

Genetic Testing – Legal Background

Since April 1, 2007 the federal law on human genetic analyses (LAGH; SR 810.12) and the regulation on human genetic analyses (RAGH; SR 810.122.1) regulate the preconditions to carry out genetic tests. In addition to the requirements for the laboratories to carry out the respective analyses, preconditions are the obligation of the referring physician to inform the patient, the guarantee of sufficient genetic counselling and the written informed consent of the patient. The above written consent allows us to comply with

these federal regulations. For further information, please contact Prof. A R. Miserez, M.D. **dia**grapa I aboratories in **Switzerland**

diagene gmbh, Routine Laboratories
Kaegenstrasse 17; CH-4153 Reinach (BL)
Tel.:+41 61 7118 580, +41 79 7086 617 (24hrs); Fax: +41 61 7118 581
E-mail: info@diagene.com, http://www.diagene.com, http://www.diagene.ch
Bank Account: Basellandschaftliche Kantonalbank, CH-4153 Reinach (BL)
Account no. 16 2.062.395.92, Clearing 769, EBAN: CH6100769016206239592

diagene gmbh, Routine Laboratories
Innocel Innovation Center, Marie Curie-Strasse 8, D-79539 Lörrach
Tel: +49 7621 5500 580, Fax: +49 7621 5500 581
E-mail: info@diagene.com, http://www.diagene.de

Bank Account: Deutsche Bank, D-79539 Lörrach, Deutschland