

Myotubular and Centronuclear Myopathy (MTM and CNM) Patient Registry

Principal Investigator/Data Controller: Dr Chiara Marini Bettolo, Institute of Genetic Medicine,
Newcastle University

Informed Consent for Female Carriers of XLMTM

	PLEASE NOW READ THE CONSENT STATEMENTS BELOW, AND GIVE YOUR CONSENT BY TICKING THE BOX NEXT TO EACH STATEMENT.	Tick box
1.	I confirm that I have read, and that I understand the above patient information (dated March 2018, version 1.1) for the MTM and CNM Patient Registry. I have had the opportunity to consider the information and to ask questions.	
2.	I give consent for my data to be stored in the MTM and CNM Registry and for it to be extracted (in anonymous form only identifiable by a code) so it may be used for research and for the planning of clinical trials.	
3.	I understand that allowing my data to be stored on the registry does not mean I will automatically be entered into future clinical trials.	
4.	I agree to be contacted periodically to provide an update about any changes in my medical condition, or about the management of my online account.	
5.	I agree to be contacted with information about relevant clinical trials, studies or surveys, to support research into my disease.	
6.	I give consent for the MTM and CNM Registry team to obtain access to relevant parts of my medical records (if needed) to confirm my diagnosis, including genetic test results, muscle biopsy results, and related correspondence.	
7.	I understand that the results from future research may not have any direct implications for myself or my family.	
8.	I understand that my participation is voluntary and that I am free to withdraw my data from the registry at any time, without giving any reason, and without my medical care or legal rights being affected.	
9.	The nature of the registry has been fully explained to me. I have understood the patient information and I agree to participate in this registry.	