Agreement for the Transfer of Materials Implementing the ACR-EULAR Myositis Response Criteria

In response to the RECIPIENT's request for the 2016 ACR-EULAR Myositis Response Criteria, including supporting computer programs, user guide, data dictionary and validation data set, (MATERIAL) for work directed by the RECIPIENT SCIENTIST: (select all that apply)

- □ 2016 ACR/EULAR Criteria for Minimal, Moderate, and Major Clinical Response in Adult Dermatomyositis and Polymyositis and other candidate criteria
- □ 2016 ACR/EULAR Criteria for Minimal, Moderate, and Major Clinical Response in Juvenile Dermatomyositis and other candidate criteria

which is to be used for the purpose of

the PROVIDER asks that the RECIPIENT and the RECIPIENT SCIENTIST agree to the following before the RECIPIENT receives the MATERIAL:

- 1. The above MATERIAL is the property of the national Institute of Environmental Health Sciences (NIEHS), International Myositis Assessment and Clinical Studies Group (IMACS) and Paediatric Rheumatology International Trials Organisation (PRINTO) represented by the NIEHS (PROVIDER) and is made available as a service to the research community.
- 2. The MATERIAL will be used for teaching and non-commercial research purposes only. For an avoidance of doubt, the use of the MATERIAL to assist in the development of therapeutic products for the treatment of myositis does not constitute commercial use.
- 3. RECIPIENT represents that RECIPIENT SCIENTIST has adequate expertise and appropriate licensure to safely use the MATERIAL for the purpose described above.
- 4. RECIPIENT agrees to use the MATERIAL only for the purpose described above.
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- 6. The MATERIAL will not be further distributed to others without the PROVIDER's written consent. The RECIPIENT shall refer any request for the MATERIAL to the PROVIDER.
- 7. The RECIPIENT agrees to acknowledge Lisa G. Rider, MD, Frederick W. Miller, MD, PhD: NIEHS, NIH, Bethesda, Maryland; Rohit Aggarwal, MD, MSc, Howard Rockette, PhD, University of Pittsburgh, Pittsburgh, Pennsylvania; Angela Pistorio, MD, PhD, Istituto Giannina Gaslini, Servizio di Epidemiologia e Biostatistica, Genoa, Italy; Jiri Vencovsky, Vencovsky, MD, PhD: Charles University, Prague, Czech Republic. IMACS; PRINTO; NIEHS; ACR; and EULAR in any publications reporting use of it.
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- 9. The RECIPIENT agrees to use the MATERIAL in compliance with all applicable statutes and regulations.
- 10. The MATERIAL is provided at no cost.

RECIPIENT and **RECIPIENT** SCIENTIST must sign this agreement and email an electronic copy to PROVIDER Scientific and Administrative Contacts at the addresses below.

RECIPIENT INFORMATION and AUTHORIZED SIGNATURE

Recipient Scientist:
Recipient Organization:
Address:
Name of Authorized Official:
Title of Authorized Official:
Signature of Authorized Official:
Date:

Certification of Recipient Scientist: I have read and understood the conditions outlined in this Agreement and I agree to abide by them in the receipt and use of the MATERIAL.

Recipient Scientist

Date

PROVIDER INFORMATION and AUTHORIZED SIGNATURE

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Signature of Authorized Official:

ACR-EULAR Myositis Response Criteria Agreement