

**Supplemental Table 1: Patient baseline characteristics and current status**

	<b>All subjects (n = 10)</b>
<i>Age (years), mean (SD)</i>	
<i>At diagnosis</i>	52.5 (19.6)
<i>Current age or at time of death</i>	56.5 (18.9)
<i>Female, n (%)</i>	8 (80.0%)
<i>Years of disease, mean (SD)</i>	3.9 (3.21)
<i>Expert classification, n (%)</i>	
<i>Classic DM</i>	5 (50.0%)
<i>Amyopathic DM</i>	3 (30.0%)
<i>Anti-synthetase syndrome</i>	2 (20.0%)
<i>Autoantibodies identified, n</i>	
<i>Anti-Jo1</i>	1
<i>Anti-PL12</i>	1
<i>Anti-SAE1</i>	1
<i>Anti-MDA5</i>	4
<i>Anti-TIF-1</i>	1
<i>Anti-Ro52</i>	6
<i>Negative MSA and MAA</i>	1
<i>Myositis antibody panel not done</i>	1
<i>Systems involved, n (%)</i>	
<i>Muscular</i>	6 (60.0%)
<i>Pulmonary</i>	5 (50.0%)
<i>Articular</i>	5 (50.0%)
<i>Cutaneous</i>	7 (70.0%)
<i>Previous therapies, mean (SD)</i>	3.90 (1.91)
<i>Rituximab, n (%)</i>	1 (10.0%)
<i>IVIg, n (%)</i>	4 (40.0%)
<i>Mycophenolate, n (%)</i>	7 (70.0%)
<i>Azathioprine, n (%)</i>	5 (50.0%)
<i>Methotrexate, n (%)</i>	8 (80.0%)
<i>Leflunomide, n (%)</i>	1 (10.0%)
<i>Tacrolimus, n (%)</i>	3 (30.0%)
<i>Hydroxychloroquine or Quinacrine, n (%)</i>	5 (50.0%)
<i>Tofacitinib, n (%)</i>	3 (30.0%)
<i>Cyclophosphamide, n (%)</i>	1 (10.0%)
<i>Currently on upadacitinib, n</i>	9
<i>As monotherapy, n (%)</i>	3 (33.3%)
<i>With concurrent treatment, n (%)</i>	6 (67.7%)
<i>Prednisone, n (%)</i>	3 (33.3%)
<i>IVIg, n (%)</i>	1 (11.1%)

<i>Mycophenolate, n (%)</i>	<i>2 (22.2%)</i>
<i>Methotrexate, n (%)</i>	<i>3 (44.4%)</i>
<i>Adverse events, total, n (%)</i>	<i>1 (10.0%)</i>
<i>Acneiform rash, n (%)</i>	<i>1 (10.0%)</i>