Supplemental Table 1: Evolution of SCNIR Patients to Lymphoproliferative Disorders

| Patient # | Diagnosis at Enrollment | FACS*, Pre G-CSF | FACS, Most current, on G-CSF | Patient Evolved To: |
|--------------|------------------------------|---|--|---|
| 1 | Idiopathic / Autoimmune | Monoclonal T cell population detected, T cell predominance | Not Tested | T-cell Lymphoma Disorder |
| 2 | Idiopathic / Autoimmune | No clonal population detected | Clonal T cell receptor gene rearrangement detected | T-cell Lymphoproliferative Disorder |
| 3 | Idiopathic / Autoimmune | Not Tested | Predominantly T-cells, CD57+ | T-cell Lymphoproliferative Disorder |
| 4 | Idiopathic / Autoimmune | Not Tested | Not Tested | Non Hodgkins Lymphoma |
| 5 | Idiopathic / Autoimmune | Not Tested | Normal CD8+ cells | Non Hodgkins Lymphoma |
| 6 | Idiopathic / Autoimmune | Not Tested | Increase CD8+ cells | T-cell Lymphoproliferative Disorder |
| 7 | Idiopathic / Autoimmune | T-cell gene rearrangements negative | T-cell gene rearrangements negative | T-cell Lymphoproliferative Disorder |
| 8 | Idiopathic / Autoimmune | Flow negative | No aberrant T cells, NK cells 25% | Multiple Myeloma |
| 9 | Idiopathic / Autoimmune | 36% lymphocytes | Increase CD8+ cells | T-cell Lymphoproliferative Disorder |
| 10 | Idiopathic / Autoimmune | Not Tested | Increase CD8+ cells | T-cell Lymphoproliferative Disorder |
| 11 | Idiopathic / Autoimmune | No immunophenotypic evidence of clonal cell population | No immunophenotypic evidence of lymphoproliferative disorder | T-cell Lymphoproliferative Disorder |
| 12 | Idiopathic / Autoimmune | Lymphoid aggregates, possible B cell neoplasm | Not Tested | Non Hodgkins Lymphoma |
| 13 | Lymphoproliferative / LGL† | Not Tested | Increase CD8+ cells | Primary diagnosis: Lymphoproliferative Disorder |
| 14 | Lymphoproliferative / LGL | T cells normal expression | Large # Polyclonal cells, mixed CD4+ and CD8+ | Primary diagnosis: Lymphoproliferative Disorder |
| 15 | Lymphoproliferative / LGL | Not Tested | Increase CD8+ cells | Primary diagnosis: Lymphoproliferative Disorder |
| 16 | Lymphoproliferative / LGL | T-cell clonality detected, clonal rearrangement detected | Abnormal T cell population | Primary diagnosis: Lymphoproliferative Disorder |

| 17 | Lymphoproliferative / LGL | Aberrant CD7+ expression by monocytes | Not Tested | Primary diagnosis: Lymphoproliferative Disorder |
|----|------------------------------|---|--|---|
| 18 | Lymphoproliferative / LGL | Not Tested | T-NK cells, CD8+ cells | Primary diagnosis: Lymphoproliferative Disorder |
| 19 | Lymphoproliferative / LGL | Not Tested | Increase CD8+ cells | Primary diagnosis: Lymphoproliferative Disorder |
| 20 | Lymphoproliferative / LGL | T-cell gene rearrangements positive | T-cell clonality detected, Increase CD8+ cells | Primary diagnosis: Lymphoproliferative Disorder |
| 21 | Lymphoproliferative / LGL | Flow consistent with LGL / T suppressor cell lymphoproliferative disorder | Not Tested | Primary diagnosis: Lymphoproliferative Disorder |
| 22 | Lymphoproliferative / LGL | Not Tested | Not Tested | Primary diagnosis: Lymphoproliferative Disorder |
| 23 | Lymphoproliferative / LGL | Not Tested | Not Tested | Non Hodgkins Lymphoma |
| 24 | Lymphoproliferative / LGL | Not Tested | 30% T8 cells | Primary diagnosis: Lymphoproliferative Disorder |

^{*}FACS – Fluorescence-activated cell sorter †LGL – Large granular lymphocyte syndrome