Results from the ITP Natural History Study Registry
661 Respondants (Demographics) 460 (MedicalDx) May 2020

Top Six Countries Represented: (23 Countries in total included in registry)

<table>
<thead>
<tr>
<th>Country</th>
<th>Count</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>USA</td>
<td>562</td>
<td>85%</td>
</tr>
<tr>
<td>Canada</td>
<td>34</td>
<td>5%</td>
</tr>
<tr>
<td>Australia</td>
<td>12</td>
<td>2%</td>
</tr>
<tr>
<td>UK</td>
<td>12</td>
<td>2%</td>
</tr>
<tr>
<td>Brazil</td>
<td>6</td>
<td>1%</td>
</tr>
<tr>
<td>India</td>
<td>6</td>
<td>1%</td>
</tr>
</tbody>
</table>

**Gender**
- Female (76%)
- Male (23.5%)
- Other (0.5%)

**Age**
- Seniors (22.4%)
- Adults (64.7%)
- Teens (7%)
- Children (6%)

**Health Insured**
- Yes (93%)
- No (4.4%)
- Unknown (2.6%)

**Platelet Disorder**
- ITP (98%)
- Other (2%)

- 92% do not have a family history of their rare disease
- 2% reported multiple family members with the same rare disease.

**Family Members with Participants**

- Child <1%
- Niece/Nephew <1%
- Sister (<1%)
- Brother (<1%)
- Cousin (1.4%)
- Parent 1%
- Aunt/Uncle (<1%)
- Grandparent (<1%)