

51 cases reported in PID patients to 29/04.

M:F = 2.16:1.

Age groups:	pts.
0-2 yrs	3
3-12 yrs	7
13-18 yrs	3
19-24 yrs	2
25-34 yrs	9
35-44 yrs	10
45-54 yrs	3
55-64 yrs	8
65-74 yrs	3
75 yrs and older	3

PID diagnosis:

3 phagocyte defects: 1 CGD under immunosuppressive therapy

6 combined immunodeficiency: 1 Di George sdr also with chronic lung disease, 2 Down sdr, 1 WAS (5 months after gene therapy), 2 unknown

33 antibody deficiencies: 4 XLA, 2 AR agammaglobulinemia, 22 CVID (of which 2 NFKB1 def. and 1 NFKB2 def.), 5 generic hypogammaglobulinemia

2 Aicardi-Goutieres syndromes treated with Ig and JAK inhibitors

1 FMF treated with mAb

3 immune dysregulation defects: 1 LRBA deficiency with type 1 diabetes treated with Ig, insulin, abatacept; 1 ALPS-like disease (suspected CTLA4) under immunosuppressive treatment; 1 hyper IgE sdr. (PEPD mut.) under immunosuppressive treatment

1 autoinflammatory disease with lymphopenia and AIHA treated with steroids

1 with CMC and recurrent sepsis treated with Ig

1 not given.

Typical presentation symptoms: upper respiratory symptoms, cough, shortness of breath, fever. Seven pts had diarrhea, vomiting; seven pts had myalgia.

32 necessitating admission, 15 evolving into respiratory insufficiency, 5 succumbed. 5 pts requiring non-invasive ventilation, 10 invasive ventilation and 1 ECMO.

Complications: 4 HLH, 5 sepsis (1 Candida), 4 bacterial pneumonia, 4 hypotension/shock with kidney failure, 1 severe AIHA, 1 anemia, 1 neutropenia.

Therapy:

Antibiotics	32
Systemic steroids	11
Ig	6
(Hydroxy)chloroquine	23
mAb (tocilizumab)	6 (4)
Remdesivir	6
Lopinavir	13
Ritonavir	12
Enoxaparin	10