TIME LINE OF DISCOVERY OF MAJOR TYPES OF PI AND MAJOR ADVANCES IN TREATMENT AND SCIENTIFIC UNDERSTANDING

1922	Neutropenia
1926	Ataxia-Telangiectasia
1929	Chronic mucocutaneous candidiasis
1937	Wiskott-(Aldrich) syndrome
1944	Purification of γ-globulin
1950	Lymphocytophthisis (SCID)
1952	Agammaglobulinemia (XLA) and treatment with γ-globulin
1953	Alymphocytosis (SCID)
1954	Acquired agammaglobulinemia in an adult woman (CVID)
1957	Chronic granulomatous disease
1957	Swiss-type agammaglobulinemia and lymphopenia (SCID)
1958	"Combined humoral and cellular" deficiency (SCID)
1960	Complement deficiency
1961	Hyper IgM Syndrome
1963	Thymic alymphoplasia (X-linked SCID)
1964	Selective IgA Deficiency
1965	DiGeorge Syndrome
1968	"Two component" concept for development of immune system (T and B cells)
1968	Bone marrow transplantation for SCID
1969	XLP (1975 "Duncan syndrome")
1970	Classification of Primary Immune Deficiencies by WHO
1972	ADA deficiency as a cause of SCID
1974	IgG Subclass deficiency
1974	Hyper IgE syndrome
1980	Leukocyte Adherence Protein deficiency (LAD)
1982	Recognition of AIDS
1982	IVIG in the US
1991	Gene therapy trials for ADA deficiency
1992	"10 Warning Signs of PI" published
1993	Identification of Btk as site of mutation in XLA
1997	SCID as "pediatric emergency"
2003-2004	Only 10 genes account for >93% of SCID

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Compiled February 2006 by Melvin Berger M.D., Ph.D.