Sociedad Latinoamericana de Immunodeficiencias

Estatísticas - Registro de IDPs

Intravenous Immunoglobulin

Sumário

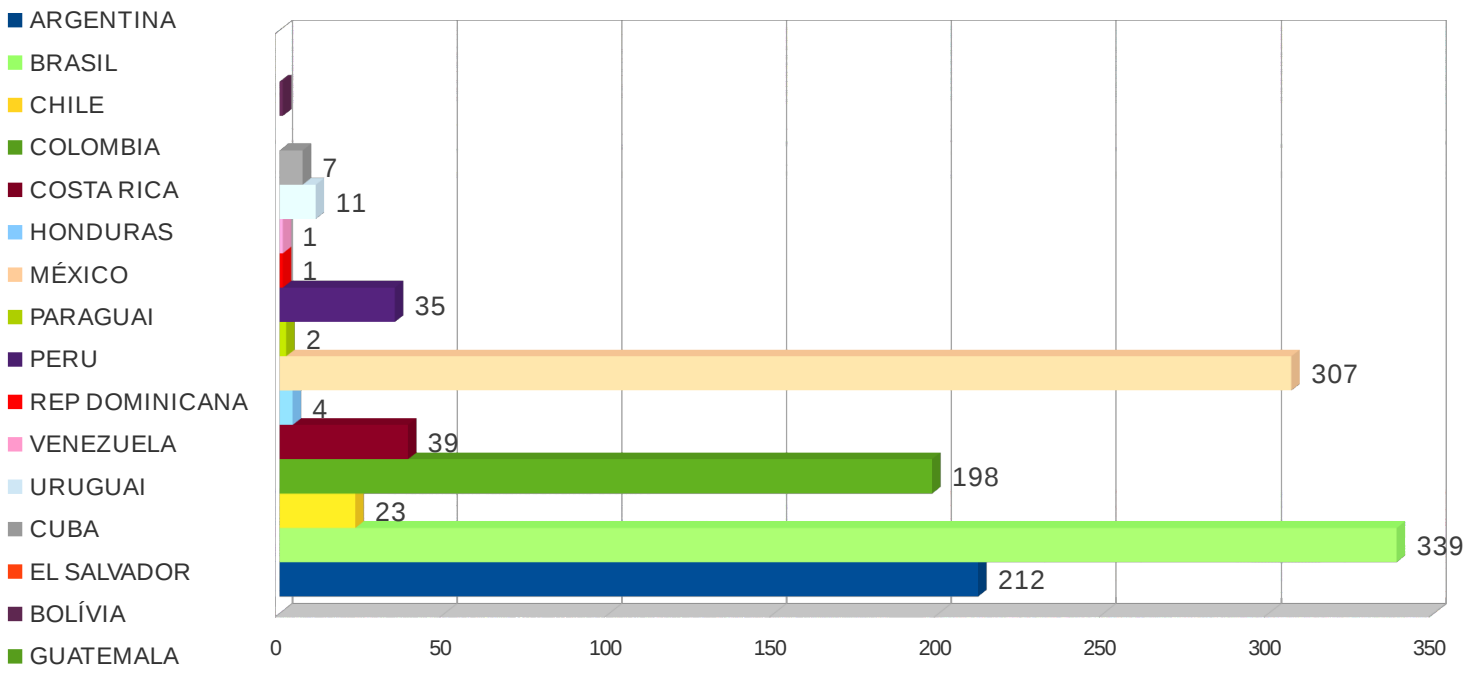
Objetivo

Este relatório apresenta dados estatísticos extraídos do Registro de Imuno Deficiências Primárias do LASID, referente ao mês de Junho de 2017.

São apresentados os casos registrados por IDP que fazem uso de '**intravenous immunoglobulin**' em sua terapia. Na ocasião da extração desses dados haviam **7.095** pacientes registrados no total.

1. Intravenous immunoglobulin por país

PAÍS	ABR / 2017	MAIO / 2017	JUNHO / 2017
ARGENTINA	212	212	212
BRASIL	328	333	339
CHILE	23	23	23
COLOMBIA	193	195	198
COSTA RICA	39	39	39
HONDURAS	4	4	4
MÉXICO	293	302	307
PARAGUAI	2	2	2
PERU	32	32	35
REP DOMINICANA	1	1	1
VENEZUELA	1	1	1
URUGUAI	11	11	11
CUBA	7	7	7
EL SALVADOR	0	0	0
BOLÍVIA	0	0	1
GUATEMALA	0	0	0
	1146	**1162**	**1180**



2. Intravenous immunoglobulin por PID

Uso intravenous immunoglobulin por PID	JUNHO / 2017
pdt; ada_ada - ADA deficiency	9
pad; ag_unknown - Agammaglobulinemias with unknown genetic cause	73
pad; ag_xla - XLA (Btk.)	168
aid; alps_fas - Fas (CD95)	1
aid; alps_unknown - ALPS with unknown genetic cause	6
pdt; cd3_cd3g - CD3 gamma deficiency	2
pdt; cd4_cd4 - Selective CD4 cell deficiency	1
pd; cgd_unknown - CGD with unknown genetic cause	2
pd; cgd_xlinkedcybb - CGD X-linked (CYBB)	1
owd; chh_pmrpchh - PMRP/CHH	1
pd; chs_chs1 - Chediak-Higashi syndrome (CHS1/Lyst)	4
pd; chs_unknown - CHS with unknown genetic cause	2
pdt; cmc_apeced - APECED (AIRE)	1
pdt; cmc_unknown - Other CMC	2
pad; csr_aid - AID deficiency (AICDA)	2
pad; csr_cd40 - CD40 deficiency	1
pad; csr_cd40l - CD40L deficiency (CD154)	18
pad; csr_uhigm - CSR defects and HIGM syndromes with unknown genetic cause	22
pdt; dgs_chrom22 - DiGeorge Syndrome	11
owd; dnab_atm - Ataxia telangiectasia (ATM)	76
owd; dnab_other - Other DNA-breakage disorder	1
pd; dwstmi_other - Susceptibility to mycobacterial infection and unknown genetic defect	1
pd; dwstmi_stat1 - STAT1 deficiency	2
pd; griscs_unknown - Griscelli syndrome with unknown genetic cause	5
pad; hg_cd19 - CD19 deficiency	6
pad; hg_cvid - CVID	333
pad; hg_cviddi - Secondary hypogammaglobulinemia	9
pad; hg_goods - Good-syndrome (associated with thymoma)	2
pad; hg_iggspecdef - Deficiency of specific IgG	61
pad; hg_iggsub - IgG subclass deficiency	29
pad; hg_siga - Selective IgA deficiency	10
pad; hg_sigm - Selective IgM deficiency	1
pad; hg_taci - TACI deficiency	1
pad; hg_transhypinf - Transient hypogammaglobulinemia of infancy	92
pad; hg_unknown - Other Hypogammaglobulinemias	13

owd; hige_hies - HIES	26
pdt; hla_ciita - Bare lymphocyte syndrome (CII TA)	6
idouc; idouc_idouc - Unclassified immunodeficiencies	5
pd; lad_lad1 - LAD1 = CD11/CD18 (CD18/ITGB2)	4
pd; nedcn_ela2 - ELA2 defective neutropenia	2
owd; ostp_clcn7 - Osteopetrosis (CLCN7)	1
pd; other_other - Other phagocytic disorders	8
pdt; outd_outd - Other unclassified T-cell disorders	30
pd; perfd_prf1 - PRF1 deficiency	1
pdt; pnp_pnp - PNP deficiency	4
pd; scn_scn - Severe congenital neutropenia	1
pdt; tbscid1_artemis - Artemis deficiency	1
pdt; tbscid1_rag1 - RAG 1 deficiency	3
pdt; tbscid1_unknown - T-B- SCID with unknown genetic cause	23
pdt; tbscid2_il7r - IL7R deficiency (IL-7R-alpha)	1
pdt; tbscid2_unknown - T-B+ SCID with unknown genetic cause	25
pdt; tbscid2_xlinkedgammac - SCID X-linked (gamma-c) (CD132)	12
owd; tlr_nemo - NEMO/IKK-gamma defiency (XED)	1
owd; was_unknown - WAS with unknown genetic cause	26
owd; was_wasp - WAS with mutations in WASP	22
owd; xlp_sh2d1a - XLP (SH2D1A)	5
owd; xlp_unknown - XLP with unknown genetic cause	3
pdt; zap_zap70 - ZAP 70	1
	1180