Sociedad Latinoamericana de Immunodeficiencias

Estatísticas - Registro de IDPs

Intravenous Immunoglobulin

Sumário

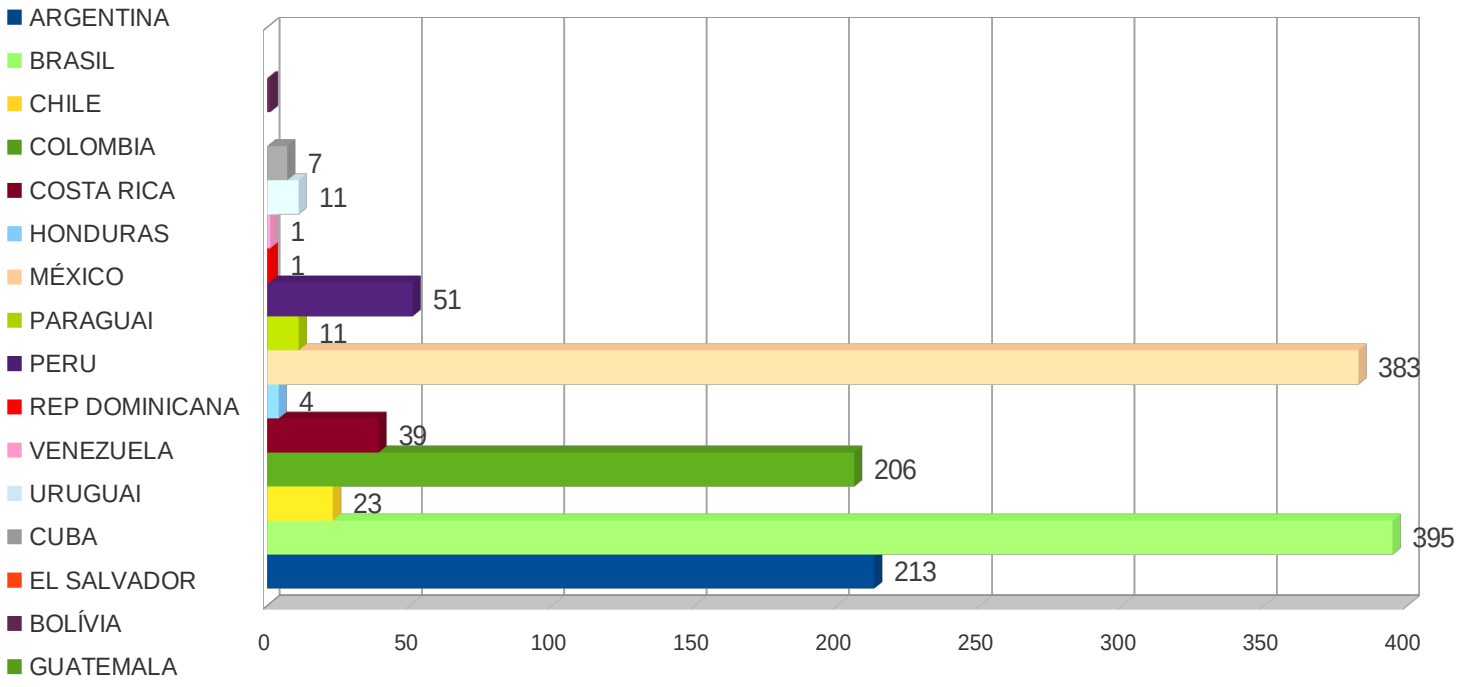
Objetivo

Este relatório apresenta dados estatísticos extraídos do Registro de Imuno Deficiências Primárias do LASID, referente ao mês de Novembro de 2018.

São apresentados os casos registrados por IDP que fazem uso de '**intravenous immunoglobulin**' em sua terapia. Na ocasião da extração desses dados haviam **8.064** pacientes registrados no total.

1. Intravenous immunoglobulin por país

PAÍS	SETEMBRO / 2018	OUTUBRO / 2018	NOVEMBRO / 2018
ARGENTINA	212	212	213
BRASIL	393	393	395
CHILE	23	23	23
COLOMBIA	206	206	206
COSTA RICA	39	39	39
HONDURAS	4	4	4
MÉXICO	369	377	383
PARAGUAI	11	11	11
PERU	51	51	51
REP DOMINICANA	1	1	1
VENEZUELA	1	1	1
URUGUAI	11	11	11
CUBA	7	7	7
EL SALVADOR	0	0	0
BOLÍVIA	1	1	1
GUATEMALA	0	0	0
	1329	**1337**	**1346**



2. Intravenous immunoglobulin por PID

Uso intravenous immunoglobulin por PID	NOVEMBRO / 2018
pdt; ada_ada - ADA deficiency	9
pad; ag_unknown - Agammaglobulinemias with unknown genetic cause	94
pad; ag_xla - XLA (Btk.)	188
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	2
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	4
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	26
pdt; dgs_chrom22 - DiGeorge Syndrome	14
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	2
pd; dwstmi_stat1 - STAT1 deficiency	2
pd; griscs_unknown - Griscelli syndrome with unknown genetic cause	6
pad; hg_baffr - BAFFR deficiency	1
pad; hg_cd19 - CD19 deficiency	8
pad; hg_cvid - CVID	380
pad; hg_cviddi - Secondary hypogammaglobulinemia	11
pad; hg_goods - Good-syndrome (associated with thymoma)	3
pad; hg_iggspecdef - Deficiency of specific IgG	69
pad; hg_iggsub - IgG subclass deficiency	32
pad; hg_siga - Selective IgA deficiency	16
pad; hg_sigm - Selective IgM deficiency	1
pad; hg_taci - TACI deficiency	1
pad; hg_transhypinf - Transient hypogammaglobulinemia of infancy	95

pad; hg_unknown - Other Hypogammaglobulinemias	24
owd; hige_hies - HIES	36
pdt; hla_ciita - Bare lymphocyte syndrome (CII TA)	6
idouc; idouc_idouc - Unclassified immunodeficiencies	6
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	35
pd; perfd_prf1 - PRF1 deficiency	1
pdt; pnp_pnp - PNP deficiency	4
pd; scn_scn - Severe congenital neutropenia	3
pdt; tbscid1_artemis - Artemis deficiency	1
pdt; tbscid1_rag1 - RAG 1 deficiency	3
pdt; tbscid1_unknown - T-B- SCID with unknown genetic cause	29
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)	13
owd; tlr_nemo - NEMO/IKK-gamma defiency (XED)	1
owd; was_unknown - WAS with unknown genetic cause	27
owd; was_wasp - WAS with mutations in WASP	25
owd; xlp_sh2d1a - XLP (SH2D1A)	5
owd; xlp_unknown - XLP with unknown genetic cause	3
pdt; zap_zap70 - ZAP 70	1
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