



INSTITUTO NACIONAL DE PEDIATRIA

PUBLICACIONES REALIZADAS

PERIODO ENERO - DICIEMBRE 2010

No.	NOMBRE	GRUPO
1	de Beaucoudrey L, Samarina A, Bustamante J, Cobat A, Boisson-Dupuis S, Feinberg J, Al-Muhsen S, Jannièrè L, Rose Y, de Suremain M, Kong XF, Filipe-Santos O, Chappier A, Picard C, Fischer A, Dogu F, Ikinciogullari A, Tanir G, Al-Hajjar S, Al-Jumaah S, Frayha HH, AlSum Z, Al-Ajaji S, Alangari A, Al-Ghonaïum A, Adimi P, Mansouri D, Ben-Mustapha I, Yancoski J, Garty BZ, Rodriguez-Gallego C, Caragol I, Kutukculer N, Kumararatne DS, Patel S, Doffinger R, Exley A, Jeppsson O, Reichenbach J, Nadal D, Boyko Y, Pietrucha B, Anderson S, Levin M, Schandené L, Schepers K, Efir A, Mascart F, Matsuoka M, Sakai T, Siegrist CA, Freceirova K, Blüetters-Sawatzki R, Bernhöft J, Freiherst J, Baumann U, Richter D, Haerynck F, De Baets F, Novelli V, Lammás D, Vermeylen C, Tuerlinckx D, Nieuwhof C, Pac M, Haas WH, Müller-Fleckenstein I, Fleckenstein B, Levy J, Raj R, Cohen AC, Lewis DB, Holland SM, Yang KD, Wang X, Wang X, Jiang L, Yang X, Zhu C, Xie Y, Lee PP, Chan KW, Chen TX, Castro G, Natera I, Codoceo A, King A, Bezrodnik L, Di Giovanni D, Gaillard MI, de Moraes-Vasconcelos D, Grumach AS, da Silva Duarte AJ, Aldana R, Espinosa-Rosales FJ, Bejaoui M, Bousfiha AA, Baghdadi JE, Ozbek N, Aksu G, Keser M, Somer A, Hatipoglu N, Aydogmus C, Asilsoy S, Camcioglu Y, Gülle S, Ozgur TT, Ozen M, Oleastro M, Bernasconi A, Mamishi S, Parvaneh N, Rosenzweig S, Barbouche R, Pedraza S, Lau YL, Ehlayel MS, Fieschi C, Abel L, Sanal O, Casanova JL. Revisiting human IL-12Rβ1 deficiency: a survey of 141 patients from 30 countries. <i>Medicine (Baltimore)</i> . 2010 Nov;89(6):381-402.	5
2	García-Ortiz H, Velázquez-Cruz R, Espinosa-Rosales F, Jiménez-Morales S, Baca V, Orozco L. Association of TLR7 copy number variation with susceptibility to childhood-onset systemic lupus erythematosus in Mexican population. <i>Ann Rheum Dis</i> . 2010 Jun 4. [Epub ahead of print]	5
3	Jacinta Bustamante, Andres A Arias, Capucine Picard, Guillaume Vogt, Iizbeth blancas galicia, jean Laurent Casanova. Germiline but macrophage-tropic CYBB mutations in kindreds with x linked predisposition to tuberculous mycobacterial diseases. <i>Nat immunol</i> . Aceptado paraa publicaciones.	5
4	Engelhardt KR, McGhee S, Winkler S, Sassi A, Woellner C, Lopez-Herrera G, Chen A, Kim HS, Lloret MG, Schulze I, Ehl S, Thiel J, Pfeifer D, Veelken H, Niehues T, Siepermann K, Weinspach S, Reisli I, Keles S, Genel F, Kutukculer N, Camcioglu Y, Somer A, Karakoc-Aydiner E, Barlan I, Gennery A, Metin A, Degerliyurt A, Pietrogrande MC, Yeganeh M, Baz Z, Al-Tamemi S, Klein C, Puck JM, Holland SM, McCabe ER, Grimbacher B, Chatila TA. Large deletions and point mutations involving the dedicator of cytokinesis 8 (DOCK8) in the autosomal-recessive form of hyper-IgE syndrome. <i>J Allergy Clin Immunol</i> . 2009 Dec;124(6):1289-302.e4.	5
5	Zander JF, Münster-Wandowski A, Brunk I, Pahner I, Gómez-Lira G, Heinemann U, Gutiérrez R, Laube G, Ahnert-Hilger G. Synaptic and vesicular coexistence of VGLUT and VGAT in selected excitatory and inhibitory synapses. <i>J Neurosci</i> . 2010 Jun 2;30(22):7634-45.	5