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PO-01	Concetta Micalizzi, Johanna Svahn, Francesca Scuderi, Tiziana Lanza, Maura Faraci, Laura Banov, Michaela Calvillo, Oriana Burlando, Francesca Fioredda, Daniela Caprino, Carlo Dufour.	Chronic urticaria as first sign of MDS
PO-02	Noha M.El Hossainy(1)Mervat M. Mattar (2)	prohepcidin in myelodysplasia
PO-03	*A Piccin, ^U Rovigatti, *C Mc Mahon, #J Kelly, ##B Bourke, *A OíMarcaigh, *OP Smith	THIOPURINE S-METHYL TRANSFERASE POLYMORPHIC STATUS, 6-MERCAPTOPYRIMIDINE, AND TNF-ALPHA ANTAGONISM IN THE DEVELOPMENT OF MYELOYDYSPLASTIC SYNDROME / MONOSOMY 7 IN A CHILD WITH CROHN'S DISEASE
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PO-05	Rekha Chaubey ,Sudha Sazawal , Manoranjan Mahapatra ,Renu Saxena	RAS and FLT3-ITD gene mutations in patients with Myelodysplastic Syndromes in India: AIIMS experience
PO-06	Shaoyan Hu1* Weiyang Gu2 Hailong He1 Yihuan Chai1 Shaoyan Hu and Weiyang Gu	The features of 65 Chinese pediatric patients with de novo myelodysplastic syndromes
PO-07	Nao Yoshida 1), Hiroshi Yagasaki 2), Yoshiyuki Kosaka 3), Ryoji Kobayashi 4), Hiromasa Yabe 5), Takashi Kaneko 6), Masahiro Tsuchida 7), Akira Ohara 8), Tatsutoshi Nakahata 9), Seiji Kojima 2)	Predicting response to immunosuppressive therapy in childhood aplastic anemia
PO-08	Muñoz-Villa A , Diaz de Heredia , Diaz MA , Badell I , Martinez A , Gonzalez-Valentin M , Dasi MA . Gomez P ,Bureo E , Olive T , Perez-Hurtado JM , Vicent MG and Maldonado MS on behalf of GETMON	Very severe acquired aplastic anemia: historical outcome of patients treated by allogeneic bone marrow transplantation from matched sibling donors. A study by the Spanish Group for Blood and Marrow Transplantation in Children (GETMON)
PO-09	Masafumi Ito, Ryoichi Ichihashi, Masahiko Fujino, Seiji Kojima*	Absence of Hemoglobin F expression on bone marrow erythroblasts identifies a distinct subset of acquired aplastic anemia patients with good response to immunosuppressive therapy
PO-10	Ferrara M., Capozzi L., Russo R.	INCIDENCE OF MALIGNANCY IN A FAMILY OF A PATIENT WITH FANCONI ANAEMIA (FA)
PO-11	Perolla.A , Caja.T, Xhumari.P , Pulluqi.P , Pushi. A	PANCYTOPENIA- First expression of several blood diseases [a 10 years study]
PO-12	M.Sukova(1),E.Mejstrikova(2),S.Pekova(3),J.Cermak(4),D.Pospisilova(5),J.Stary(1)	Childhood paroxysmal nocturnal hemoglobinuria/aplastic anemia (PNH/AA) syndrome - analysis for PNH clonal evolution after immunosuppressive therapy
PO-13	Isabelle Meyts, Marleen Renard, Johan Maertens*, Ann Uyttebroeck	Salvage allogeneic stem cell transplantation in a child with severe aplastic anemia in the course of life-threatening potential invasive fungal lung infection with peri- and myocardial infiltration
PO-14	RAHIM F 1,2,PANAHI M 3,RASHIDI 14	Incidence of Aplastic Anaemia in Khuzestan Province, Iran: A Retrospective Single-Centre Study

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PO-15	Veronica M.E. Neefjes(1), Rienk Y.J. Tamminga(2)	ATG IN ACQUIRED APLASTIC ANAEMIA: IS LESS MORE?
PO-16	Pallota, Ronald 1; Gomes, Laís 1	Aplastic Anemia secondary to Dengue virus infection: A Case Report
PO-17	Vallero S, Quarello P, Lorenzati A, Renga D, Rivetti E, Laudati R, Farinasso L, Timeus F, Ramenghi U, Saracco P	Use of pegylated granulocyte-colony stimulating factor (PG-CSF) in childhood severe aplastic anemia.
PO-18	Dr. Maitreyee Bhattacharyya, Dr. Seema Tyagi, Dr. M Mahapatra, Dr. V.P. Chaudhury, Dr. Renu Saxena	Etiopathogenesis of childhood aplastic anemia -- a single center study of 94 cases.
PO-19	F Timeus, N Crescenzo, P Quarello, A Doria, L Foglia, S Pagliano, A Lorenzati, S Vallero, L Farinasso, P Saracco.	Paroxysmal nocturnal hemoglobinuria clones in children with acquired aplastic anemia: a single centre study.
PO-20	Saracco P, Quarello P, Iori AP, Zecca M, Longoni D, Svahn J, Varotto S, Del Vecchio GC, Dufour C, Ramenghi U, Bacigalupo A, Locasciulli A; Bone Marrow Failure Study Group of the AIEOP	Cyclosporin A response and dependence in children with acquired aplastic anaemia: an update of a multicentre retrospective study with long-term observation follow-up.
PO-21	1 Mahfuzah Mohamed, 1 Eni Juraida A. Rahman, 2 Bina S Menon, 1 Caroline Ho, 1 Raja Khuzaiyah, 1 Zulaiha Muda, 1 Ida Shahnaz Othman, 1 Lim Yin Sear, 1 Asohan Thevarajah, 1 Aisyah M. Rivai, 1 Hishamshah M. Ibrahim	Immunosuppression therapy in children with severe aplastic anaemia
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PO-26	Tim Ripperger (+), Doris Steinemann (+), Gudrun G^hring (+), Charlotte Niemeyer (*), Brigitte Strahm (*), and Brigitte Schlegelberger (+)	A further pedigree with germline RUNX1 mutation and propensity to myeloid malignancies
PO-27	Daisuke Hasegawa, Chitose Ogawa, Shinsuke Hirabayashi, Myoung-ja Park, Yasuhide Hayashi, Atsushi Manabe, Ryota Hosoya.	A Japanese pedigree with RUNX1 mutation resulting in familial platelet disorder with propensity to acute myelogenous leukemia (FPD/AML)
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PO-29	Ferrara M., Capozzi L., Russo R.	A case of Congenital Dyserythropoietic Anemia type II (CDA II)
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PO-34	Vallero S, Aschero S, Garbarini L, Mazzone R, Crescenzo N, Iavarone A, Ricotti E, Vassallo E, Nesi F, Cordero di Montezemolo L, Farinasso L	Del(5q) in a patient with JMML, neurofibromatosis 1 and Parvovirus B19 infection.
PO-35	A.C.H. de Vries (1), R.G.M. Bredius (2), A.C. Lankester (2), M. Bierings (3), M. Trebo (4), P. Sedlacek (5), C.M. Niemeyer (6), M. Zecca (7), F. Locatelli (7), M.M. van den Heuvel-Eibrink (1)	HLA-IDENTICAL UMBILICAL CORD BLOOD TRANSPLANTATION FROM A SIBLING DONOR IN JUVENILE MYELOMONOCYTIC LEUKEMIA
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PO-38	Arzu Akyay*, Lale Olcay*, Mehmet Ertem**, I nsu Kuzu**, Nazan Bozdo an*, Elif knal nce**, Talia leri**, Bet l Tavl*, Ay e Say l **, Z mr t Uysal**	A CASE OF MYELOYDYSPLASTIC SYNDROME WITH HYPOCELLULER FIBROSIS
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PO-43	Sudha Sazawal, Jyoti Bajaj, Bharti Sharma, Rahul Bhargava, Rekha Chaubey, Manoranjan Mahapatra, Renu Saxena	Frequency of JAK2 V617F in Indian patients with Chronic myeloproliferative disorders: AIIMS experience
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PO-51	D.Wojcik ^{1,2} ,J.Musia ¹ ,E.Gorczy ska ¹ ,M.Ussowicz ¹ ,A.Dyla ¹ ,J.Owoc-Lempach ¹ ,M. S ociak ¹ ,D.Sega-Pondel ¹ ,W.Pietras ¹ and K.Ka wak ¹	Haematopoietic Stem Cell Transplantation (HSCT) for pediatric MDS-Single center study-
PO-52	Igor Resnick, Polina Stepensky, Michael Weintraub, Dani Engelhard, Diana Averbukh, Jacqueline Feinberg, StÉphanie Boisson-Dupuis, Jacinta Bustamante, Jean-Laurent Casanova.	A NOVEL FORM OF MYELODYSPLASTIC SYNDROME (MDS) ASSOCIATED WITH MENDELIAN SUSCEPTIBILITY TO MYCOBACTERIAL DISEASE (MSMD).