

IMER- Indagine sulle Malformazioni congenite in Emilia Romagna

Pubblicazioni

- 1) LORENTE C, CORDIER S, BERGERET A, DE WALLE HE, GOJUARD J, AYME S, KNILL-JONES R, CALZOLARI E, BIANCHI F. (2000): Maternal occupational risk factors for oral clefts. Occupational Exposure and Congenital Malformation Working Group. *Scand J Work Environ Health.* Apr;26(2):137-45.
- 2) VINCETI M, CANN CI, CALZOLARI E, VIVOLI R, GARAVELLI L, BERGOMI M. (2000): Reproductive outcomes in a population exposed long-term to inorganic selenium via drinking water. *Sci Total Environ.* 2000 Apr 24;250(1-3):1-7.
- 3) LORENTE C, CORDIER S, GOJUARD J, AYME S, BIANCHI F, CALZOLARI E, DE WALLE HE, KNILL-JONES R. (2000): Tobacco and alcohol use during pregnancy and risk of oral clefts. Occupational Exposure and Congenital Malformation Working Group. *Am J Public Health.* 2000 Mar;90(3):415-9.
- 4) GARAVELLI L, DONADIO A, BANCHINI G, MAGNANI C, CALZOLARI E, AND FRYNS JP, (2000): Marden Walker Syndrome : case report, nosologic discussion and aspects of counselling. *Genetic Counselling;* 11(2): 111-118
- 5) SOLLAZZO V, BERTOLANI G, CALZOLARI E, ATTI G, SCAPOLI C, (2000): A two-locus model for non-syndromic congenital dysplasia of the hip (CDH). *Ann Hum Genet,* 64(Pt 1): 51-59
- 6) CLAUSER L, GALIE M, HASSANIPOUR A, CALABRESE O, (2000): Saethre-Chotzen syndrome: review of the literature and report of a case. *J Craniofac Surg.* Sep; 11(5): 480-486
- 7) BIANCHI F, CALZOLARI E, CIULLI L, CORDIER S, GUALANDI F, PIERINI A, MOSSEY P, (2000): Environment and genetics in the etiology of cleft lip and cleft palate with reference to the role of folic acid. *Epidemiol Prev.* Jan-Feb; 24(1): 21-27. Review. Italian.
- 8) VINCETI M, ROVESTI S, CALZOLARI E, VIVOLI R, ASTOLFI G, BERGOMI M. (2000): Teratogenic effects of environmental lead exposure: a population case-control study.
- 9) GARAVELLI L, CALZOLARI E, (2000): Congenital cardiopathy in a data-based population. *Acta Biomed. Ateneo Parmense.* 71 (1): 483-486. Italian.
- 10) MAGNANI C, BUSSOLATI G, GAMBINI L, GARANI GP, BOSI G, COCCHI G, GARAVELLI L, CALZOLARI E, (2000): Congenital cardiopathy in a data-based population. *Acta Biomed. Ateneo Parmense.* 71 (1): 483-486. Italian.
- 11) MAGNANI C, BUSSOLATI G, GAMBINI L, GARANI GP, BOSI G, COCCHI G, GARAVELLI L, CALZOLARI E.(2000): Congenital cardiopathy in a data-based population. *Acta Biomed Ateneo Parmense.*;71 Suppl 1:483-6. Italian.
- 12) GARAVELLI L, DONADIO A, SIGORINI M, GRASSI L, BANCHINI G. (2000): Genetics of type 1 neurofibromatosis. *Acta Biomed Ateneo Parmense.*;71(3-4):89-95. Italian.
- 13) DONADIO A, GARAVELLI L, LORENZETTI ME, VILLANI AR, BANCHINI G, VIRDIS R. (2000): Care recommendations for type 1 neurofibromatosis. *Acta Biomed Ateneo Parmense.* 71(3-4):83-7. Italian.
- 14) VIRDIS R, SIGORINI M, LAIOLO A, LORENZETTI E, STREET ME, VILLANI AR, DONADIO A, PISANI F, TERZI C, GARAVELLI L. (2000): Neurofibromatosis type 1 and precocious puberty. *J Pediatr Endocrinol Metab.* Jul;13 Suppl 1:841-4.
- 15) GARAVELLI L, DONADIO A, BANCHINI G, MAGNANI C, CALZOLARI E, FRYNS JP. (2000): Marden-Walker syndrome: case report, nosologic discussion and aspects of counseling. *Genet Couns.* 2000;11(2):111-8.
- 16) MIRABELLI D, MAGNANI C. (2003): Interaction between occupational and environmental exposure to asbestos. Epidemiologic survey. *G Ital Med Lav Ergon.* Jul-Sep;25(3):402-4. Italian.
- 17) VINCETI M, CANN CI, CALZOLARI E, VIVOLI R, GARAVELLI L, BERGOMI M. (2000): Reproductive outcomes in a population exposed long-term to inorganic selenium via drinking water. *Sci Total Environ.* Apr 24;250(1-3):1-7.
- 18) DONADIO A, GARAVELLI L, BANCHINI G, NERI G. (2000): Kabuki syndrome and diaphragmatic defects: a frequent association in non-Asian patients? *Am J Med Genet.* Mar 13;91(2):164-5..
- 19) STOLL C, ROSANO A, BOTTO LD, ERICKSON D, KHOURY MJ, OLNEY RS, CASTILLA EE, COCCHI G, CORNEL MC, GOJUARD J, BERMEJO E, MERLOB P, MUTCHINICK O, RITVANEN A, ZAMPINO G, MASTROIACOVO P, (2001): On the symmetry of limb deficiencies among children with multiple congenital anomalies. *Ann Genet.* Jan-Mar;44(1):19-24.
- 20) ARPINO C, BRESCIANINI S, ROBERT E, CASTILLA EE, COCCHI G, CORNEL MC, DE VIGAN C, LANCASTER PA, MERLOB P, SUMIYOSHI Y, ZAMPINO G, RENZI C, ROSANO A, MASTROIACOVO P. (2000): Teratogenic effects of antiepileptic drugs: use of an International Database on Malformations and Drug Exposure (MADRE). *Epilepsia.* Nov;41(11):1436-43.
- 21) ROSANO A, BOTTO LD, OLNEY RS, KHOURY MJ, RITVANEN A, GOJUARD J, STOLL C, COCCHI G, MERLOB P, MUTCHINICK O, CORNEL MC, CASTILLA EE, MARTINEZ-FRIAS ML, ZAMPINO G, ERICKSON JD, MASTROIACOVO P. (2000): Limb defects associated with major congenital anomalies: clinical and epidemiological study from the International Clearinghouse for Birth Defects Monitoring Systems. *Am J Med Genet.* Jul 17;93(2):110-6.
- 22) VINCETI M, ROVESTI S, BERGOMI M, CALZOLARI E, CANDELA S, CAMPAGNA A, MILAN M, VIVOLI G, (2001): Risk of birth defects in a population exposed to environmental lead pollution. *Sci Total Environ.*, Oct 20; 278(1-3): 23-30
- 23) GUIDETTI D, SABADINI R, FERLINI A, TORRENTE I, (2001): Epidemiological survey of X-linked bulbar and spinal muscular atrophy, or Kennedy disease, in the provincia of Reggio Emilia, Italy. *Eur J Epidemiol.* 17(6):587-591

- 24) VINCETI M, ROVESTI S, CALZOLARI E, VIVOLI R, ASTOLFI G, BERGOMI M, (2001) : Teratogenic effects of environmental lead exposure: a population case-control study. Ann Ig. 20 Mar-Apr; 13(2): 129-138
- 25) MASCALCHI M, COSOTTINI M, LOLLI F, SALVI F, TESSA C, MACUCCI M, TOSETTI M, PLASMATI R, FERLINI A, TASSINARI CA, VILLARI N, (2002) : Proton MR spectroscopy of the cerebellum and pons in patients with degenerative ataxia. Radiology, 223(2) : 371-3
- 26) BIANCHI F, LINZALONE N, CATALANO S, BIANCA S, CALZOLARI E, MAMMI I, SCARANO G, (2002): Parental agricultural work and risk of congenital anomalies: a registry based case-control study in Italy. La Medicina del Lavoro 93(5):399-400
- 27) STORTI S, VITTORINI S, LASCONE MR, SACCHELLI M, COLLAVALI A, RIPOLI A, COCCHI G, BIAGINI A, CLERICI A. (2003): Association between 5,10-methylenetetrahydrofolate reductase C677T and A1298C polymorphisms and conotruncal heart defects. Clin Chem Lab Med. Mar;41(3):276-80.
- 28) BARONCINI A, GARAVELLI L, GUALANDI F, SCORRANO M, BOSI G; IMER WORKING GROUP. (2003) Congenital heart defects: 15 years of experience of the Emilia-Romagna Registry (Italy). Eur J Epidemiol.;18(8):773-80.
- 29) BOSI G, GARANI G, SCORRANO M, CALZOLARI E; IMER WORKING PARTY. (2003): Temporal variability in birth prevalence of congenital heart defects as recorded by a general birth defects registry. J Pediatr. 2003 Jun;142(6):690-8.
- 30) CALZOLARI E, GARANI G, COCCHI G, MAGNANI C, RIVIERI F, NEVILLE A, ASTOLFI G, BARONCINI A, GARAVELLI L, GUALANDI F, SCORRANO M, BOSI G, ; imer Working group (2003): Congenital heart defects: 15 years of experience of the Emilia-Romagna Registry (Italy). Eur J Epidemiol.; 18(8): 773-780
- 31) GUALANDI F, MARTINI A, CALZOLARI E, (2003): Progress in understanding GJB2-linked deafness: Community Genet. ; 6(3): 125-132
- 32) RIVIERI F, MIOLO G, ASTOLFI G, BIANCA S, INGEGNOSI C, PALAZZI P, CALZOLARI E, (2003): Esophageal atresia and tracheoesophageal fistula in two Italian Registers of Congenital Malformations. Abstr 7th European Symposium on the Prevention of Congenital Anomalies - Heidelberg
- 33) VOLANTE E, BRAIBANTI S, MUSETTI M, MAGNANI C, PISANI F, NERI F. (2000): Retinopathy of prematurity: incidence and risk factors. Acta Biomed Ateneo Parmense.;71 Suppl 1:615-20. Italian.
- 34) CALZOLARI E, GARANI G, COCCHI G, MAGNANI C, RIVIERI F, NEVILLE A, ASTOLFI G, BARONCINI A, GARAVELLI L, GUALANDI F, SCORRANO M, BOSI G, IMER Working Group. (2003): Congenital heart defects: 15 years of experience of the Emilia-Romagna Registry (Italy). Eur J Epidemiol.;18(8):773-80.
- 35) NEVILLE A, COCCHI G AND CALZOLARI E, (2003): EUROCAT Special Report: Prevention of Neural Tube Defects by Periconceptional Folic Acid Supplementation in Europe. EUROCAT Central Registry, University of Ulster. Chapter Italy
- 36) CALZOLARI E, NEVILLE A, COCCHI G. (2003): Dietary folates are not enough. British Medical Journal; 326: 1054-c. Rapid response.
- 37) TESSA A, SALVI S, CASALI C, GARAVELLI L, DIGILIO MC, DOTTI MT, DI GIANDOMENICO S, VALOPPI M, GRIECO GS, COMANDUCCI G, BIANCHINI G, FORTINI D, FEDERICO A, GIANNOTTI A, SANTORELLI FM. (2003): Six novel mutations of the RUNX2 gene in Italian patients with cleidocranial dysplasia. Hum Mutat. Jul;22(1):104.
- 38) VIRDIS R, STREET ME, BANDELLO MA, TRIPODI C, DONADIO A, VILLANI AR, CAGOZZI L, GARAVELLI L, BERNASCONI S. (2003): Growth and pubertal disorders in neurofibromatosis type 1. J Pediatr Endocrinol Metab. Mar;16 Suppl 2:289-92. Review.
- 39) GARAVELLI L, DONADIO A, ZANACCA C, BANCHINI G, DELLA GIUSTINA E, BERTANI G, ALBERTINI G, DEL ROSSI C, ZWEIER C, RAUCH A, ZOLLINO M, NERI G. (2003): Hirschsprung disease, mental retardation, characteristic facial features, and mutation in the gene ZFHX1B (SIP1): confirmation of the Mowat-Wilson syndrome. Am J Med Genet A. Feb 1;116(4):385-8
- 40) TARUSCIO D, FALBO V, FLORIDIA G, SALVATORE M, PESCUCCI C, CANTAFORA A, MARONGIU C, BARONCINI A, CALZOLARI E, CAO A, CASTALDO G, BRICARELLI FD, GUANTI G, NITSCH L, PIGNATTI PF, ROSATELLI C, SALVATORE F, ZUFFARDI O, (2004): Quality assessment in cytogenetic and molecular genetic testing: the experience of the Italian Project on Standardisation and Quality Assurance: Clin Chem Lab med. ; 42(8): 915-921
- 41) KROES HY, OLNEY RS, ROSANO A, LIU Y, CASTILLA EE, COCCHI G, DE VIGAN C, MARTINEZ-FRIAS ML, MASTROIACOVO P, MERLOB P, MUTCHINICK O, RITVANEN A, STOLL C, VAN ESSEN AJ, COBBEN JM, CORNEL MC. (2004): Renal defects and limb deficiencies in 197 infants: is it possible to define the "acrorenal syndrome"? Am J Med Genet A. Aug 30;129(2):149-55.
- 42) DOLK H, VRIJHEIDM, SCOTT JE, ADDOR MC, BOTTING B, DE VIGAN C, DE WALLE H, GARNE E, LOANE M, PIERINI A, GARCIA-MINAUR S, PHYSCK N, TENCONI R, WIESEL A, CALZOLARI E, STONE D, (2004): Toward the effective surveillance of hypospadias. Environ health Perspect. Mar; 112(3): 398-402
- 43) COHEN N, RIMESSI P, GUALANDI F, FERLINI A, MUNTONI F, (2004): In vivo study of an aberrant dystrophin exon inclusion in X-linked dilated cardiomyopathy. Biochem Biophys Res Commun. May 14; 317(4): 1215-1220
- 44) GUALANDI F, RAVANI A, BERTO A, BURDO S, TREVISI P, FERLINI A, MARTINI A, CALZOLARI E, (2004): Occurrence of del (GIB6-D13S1830) mutation in Italian non-syndromic hearing loss patients carrying a single GJB2 mutated allele. Acta Otolaryngol Suppl. May: (552): 29-34
- 45) CALZOLARI E, BIANCHI F, RUBINI M, RITVANEN A, NEVILLE AJ, EUROCAT Working Group (2004): Epidemiology of cleft palate in Europe: implication for genetic research: Cleft palate Craniofac J. May: 41(3): 244-249
- 46) NEVILLE A, CALZOLARI E, BUSBY A, ABRAMSKY L, DOLK H, AND THE EUROCAT WORKING GROUP (2004): EUROCAT European Surveillance of Congenital Anomalies Working Group on Periconceptional Folic Acid
- 47) Supplementation. 1° Riunione Network Italiano . Promozione Acido Folico per la prevenzione Primaria dei difetti congeniti. Libro degli Abstracts. Istituto Superiore di Sanita'- Roma-Aprile

- 48) NEVILLE A, CALZOLARI E, ABRAMSKY L. (2004): Folic acid supplementation: long term safety is not an excuse for inaction. British Medical Journal; 328: 211- Rapid Response.
- 49) RIVIERI F, MIOLO G, NEVILLE A, ASTOLFI G, CALZOLARI E, BIANCHI F, (2004): Investigation of two clusters of Downs sindrome from live births and induced abortions in the year 2000, Emilia Romagna Italy. Anomalies. Reproductive Toxicology 18(1):142,
- 50) RIVIERI F, MIOLO G, NEVILLE A, ASTOLFI G, PALAZZI P, MAGNANI C, FERRARI P, GARAVELLI L, GARANI G, COCCHI G, CALZOLARI E E BIANCHI F. Cluster di Sindrome di Down nell' Agosto 2000 in Emilia Romagna. XIX Convegno IMER. 2 aprile 2004. Poster.
- 51) RIVIERI F, MIOLO G, NEVILLE A, ASTOLFI G, PALAZZI P, MAGNANI C, FERRARI P, GARAVELLI L, GARANI G, COCCHI G, CALZOLARI E Atresia esofagea: Registro Imer (1981-2000) XIX Convegno IMER. Approccio integrato al bambino con anomalie congenite. Bologna 2 Aprile 2004. Poster.
- 52) RIVIERI F, MIOLO G, NEVILLE A, ASTOLFI G, PALAZZI P, MAGNANI C, FERRARI P, GARAVELLI L, GARANI G, COCCHI G, CALZOLARI E Ipospadia: 20 anni di sorveglianza XIX Convegno IMER Bologna 2 Aprile 2004. Poster.
- 53) NEVILLE A RIVIERI F, PALAZZI P, ASTOLFI G, CALZOLARI E. Cambiamento demografico in Emilia Romagna: Prevalenza di malformazione congenite nella popolazioni stranieri. XIX Convegno IMER Bologna 2 Aprile 2004. Poster
- 54) ZOLLINO M, LECCE R, SELICORNI A, MURDOLO M, MANCUSO I, MARANGI G, ZAMPINO G, GARAVELLI L, FERRARINI A, ROCCHI M, OPITZ JM, NERI G. (2004): A double cryptic chromosome imbalance is an important factor to explain phenotypic variability in Wolf-Hirschhorn syndrome. Eur J Hum Genet. Oct;12(10):797-804.
- 55) GARAVELLI L, ZANACCA C, CASELLI G, BANCHINI G, DUBOURG C, DAVID V, ODENT S, GURRIERI F, NERI G. (2004): Solitary median maxillary central incisor syndrome: clinical case with a novel mutation of sonic hedgehog. Am J Med Genet A May 15;127(1):93-5.
- 56) GURRIERI F, SCARANO G, GARAVELLI L, DELLA MONICA M, LONARDO F, CUDA D, BANCHINI G, OPITZ JM, NERI G. (2004): Mental retardation, Robin sequence, and brachydactyly: further confirmation of a new syndrome. Am J Med Genet A. Apr 15;126(2):204-7.
- 57) ORRICO A, GALLI L, CAVALIERE ML, GARAVELLI L, FRYNS JP, CRUSHELL E, RINALDI MM, MEDEIRA A, SORRENTINO V. (2004): Phenotypic and molecular characterisation of the Aarskog-Scott syndrome: a survey of the clinical variability in light of FGD1 mutation analysis in 46 patients. Eur J Hum Genet. 2004 Jan;12(1):16-23.
- 58) GARAVELLI L, PEDORI S, ZANACCA C, CASELLI G, LOIODICE A, MANTOVANI G, AMMENTI A, VIRDIS R, BANCHINI G. (2005): Albright's hereditary osteodystrophy (pseudohypoparathyroidism type Ia): clinical case with a novel mutation of GNAS1. Acta Biomed Ateneo Parmense. Apr;76(1):45-8.
- 59) GARAVELLI L, LEASK K, ZANACCA C, PEDORI S, ALBERTINI G, DELLA GIUSTINA E, CROCI GF, MAGNANI C, BANCHINI G, CLAYTON-SMITH J, BOCIAN M, FIRTH H, GOLD JA, HURST J. (2005): MRI and neurological findings in macrocephaly-cutis marmorata telangiectatica congenita syndrome: report of ten cases and review of the literature. Genet Couns.;16(2):117-28.
- 60) ZWEIER C, THIEL CT, DUFKE A, CROW YJ, MEINECKE P, SURI M, ALA-MELLO S, BEEMER F, BERNASCONI S, BIANCHI P, BIER A, DEVRIENDT K, DIMITROV B, FIRTH H, GALLAGHER RC, GARAVELLI L, GILLESSEN-KAESBACH G, HUGDINS L, KAARIAINEN H, KARSTENS S, KRANTZ I, MANNHARDT A, MEDNE L, MUCKE J, KIBAEK M, KROGH LN, PEIPPO M, RITTINGER O, SCHULZ S, SCHELLEY SL, TEMPLE IK, DENNIS NR, VAN DER KNAAP MS, WHEELER P, YERUSHALMI B, ZENKER M, SEIDEL H, LACHMEIJER A, PRESCOTT T, KRAUS C, LOWRY RB, RAUCH A. (2005): Clinical and mutational spectrum of Mowat-Wilson syndrome. Eur J Med Genet. Apr-Jun;48(2):97-111. Epub 2005 Feb 25.
- 61) GARAVELLI L, CERRUTI-MAINARDI P, VIRDIS R, PEDORI S, PASTORE G, GODI M, PROVERA S, RAUCH A, ZWEIER C, ZOLLINO M, BANCHINI G, LONGO N, MOWAT D, NERI G, BERNASCONI S. (2005): Genitourinary anomalies in Mowat-Wilson syndrome with deletion/mutation in the zinc finger homeo box 1B gene (ZFXH1B). Report of three Italian cases with hypospadias and review. Horm Res.;63(4):187-92. Epub 2005 May 20.
- 62) CALZOLARI E, BARONCINI A, (2005): International cooperation and networking in genetic health care provision: issue arising from the genetic services plan for the Emilia-Romagna region, Italy. Community Genet. ; 8(2): 122-129
- 63) BUSBY A, ARMSTRONG B, DOLK H, ARMSTRONG N, HAEUSLER M, BERGHOLD A, GILLEROT Y, BAGUETTE A, GJERGJA R, BARISIC I, CHRISTIANSEN M, GOJUARDA J, STEINBICKER V, ROSCH C, MCDONNELL R, SCARANO G, CALZOLARI E, NEVILLE A, COCCHI G, BIANCA S, GATT M, WALLE HD, BRAZ P, LATOS-BIELENSKA A, GENNER B, PORTILLO I, ADDOR MC, ABRAMSKY L, RITVANEN A, ROBERT-GNANSIA E, DALTEVEIT AK, ANEREN G, OLARS B, EDWARDS G, (2005): Preventing neural tube defects in Europe: a missed opportunity. Reprod Toxicol. Sep-Oct; 20(3): 393-402
- 64) BARONCINI A, RIVIERI F, CAPUCCI A, CROCI G, FRANCJI F, SENSI A, BATTAGLIA P, AIELLO V, CALZOLARI E, (2005). FISH screening for subtelomeric rearrangements in 219 patients with idiopathic mental retardation and normal karyotype. Eur J Med Genet. Oct-Dec; 48(4): 388-396
- 65) STUPPIA, ANTONUCCI I, BINNI F, BRENDI A, GRIFONE N, COLOSIMO A, DE SANTO M, GATTA V, GELLI G, GUIDA V, MAJORE S, CALABRESE G, PALKA C, RAVANI A, RINALDI R, TIBONI GM, BALLONE E, VENTUROLI A, FERLINI A, TORRENTE I, GRAMMATICO P, CALZOLARI E, DALLAPICCOLA B, (2005). Screening of mutations in the CFTR gene in 1195 couples entering assisted reproduction technique programs. Eur J Hum genet. Aug; 13(8) 959-964
- 66) RUBINI M, BRUSATI R, GARATTINI G, MAGNANI C, LINIERO F, BIANCHI F, TARANTINO E, MASSEI A, POLLASTRI S, CARTURAN S, AMADORI A, BERTAGNIN E, CAVALLARO A, FABIANO A, FRANCHELLA A, CALZOLARI E, (2005):Cistationine beta-synthase c.844ins68 gene variant and non-syndromic cleft lip and palate. Am J Med Genet A. Aug 1; 136(4): 368-372

- 67) NEVILLE A, ASTOLFI G, RIVIERI F, PALAZZI P, COCCHI G, MAGNANI C, GARAVELLI L, GARANI GP, FERRARI P, CALZOLARI E, (2005): Maternal obesity and congenital anomalies: temporal change in BMI in Emilia Romagna, Italy, Archives of Perinatal Medicine, suppl.
- 68) CALZOLARI E, ASTOLFI G, BIANCHI F, NEVILLE A, PALAZZI P, PIERINI A, RIVIERI F, and EUROCAT Working Group (2005). Epidemiology of cleft lip and palate in EUROCAT registries. Archives of Perinatal Medicine, suppl.
- 69) NEVILLE A, RIVIERI F, ASTOLFI G, PALAZZI P, GARAVELLI L, GARANI GP, FERRARI P, CALZOLARI E, (2005): Understanding demographic change in a birth defects registry: prevalence, risk factors and monitoring of an immigrant population in Emilia Romagna. Archives of Perinatal Medicine, suppl
- 70) NEVILLE A, COCCHI G, AND CALZOLARI E, (2005): EUROCAT Special Report: Prevention of Neural Tube Defects by Periconceptional Folic Acid. Supplementation in Europe: Chapter Italy. EUROCAT Central Registry, University of Ulster
- 71) NEVILLE A, CALZOLARI E, (2005): EUROCAT Special Report: Review of environmental Risk : Chapter :Maternal obesity and risk for birth defects . EUROCAT Central Registry, University of Ulster.
- 72) ASTOLFI G , CALZOLARI E , COCCHI G, FERRARI P, GARANI G, GARAVELLI L, MAGNANI C, NEVILLE A ,PALAZZI P, RIVIERI F, (2005).Rapporto annuale sulle malformazioni congenite 2002. IMER
- 73) COCCHI G, CAPELLI M, GIURA F, VITALI F, GUALDI S, MAZZONI E,: 2005: Short distal limbs, polydactyly and nail hypoplasia. Italian Journal of Pediatrics December; Vol.31; 6:345-346
- 74) COCCHI G, VITALI F, CAPELLI M, GUALDI S, GIURA F, ASTOLFI G, RIVIERI F, (2005): Ritardo di crescita intrauterina (iugr) e malformazioni congenite. Minerva Pediatrica, 57 (Suppl. 1 al N. 3): 19-21
- 75) COCCHI G, CAPELLI M, GARAVELLI L, GUALDI S, MAZZONI E, (2003): Thumb and facial abnormalities, microcephaly and cardiac defects. Italian Journal of Pediatrics;29:254-256
- 76) PRANDSTRALLER D, COCCHI G, CAPELLI M, CANZI A, MAZZONI E, CAPUCCI A, GUALDI S, TRABANELLI C, BARONCINI A, CALZOLARI E, PICCHIO FM, SALVIOLI GP,(2002) : Phenotypic variability in patients with conotruncal diseases and del22q11.2. Atti deletion 22q11.2 - Third International Meeting - Rome, 7/8 June
- 77) TRABANELLI C, VITTORIANI S, SACCHELLI M, ROSATO S, COCCHI G, CAPELLI M, PICCHIO FM, PRANDSTRALLER D, BARBONCINI A, CALZOLARI E, (2002): 22q11 deletions in 135 patients with conotruncal heart defects. Atti deletion 22q11.2 - Third International Meeting - Rome, 7/8 June
- 78) CAPELLI M, DI CARLO R, GUALDI S, MASTROCOLA M, MAZZONI E, RIVIERI F, MIOLO GM, ASTOLFI G, CALZOLARI E, COCCHI G, (2003): Displasie scheletriche: dati del registro imer. Atti 9° congresso nazionale SIN - Napoli, 21-24 maggio
- 79) CALZOLARI E, PIERINI A, ASTOLFI G, BIANCHI F, NEVILLE A, RIVIERI F AND EUROCAT WORKING GROUP. Associated anomalies in multi-malformed infants with Cleft Lip and Palate: An epidemiological study based on nearly 6 million births in 23 EUROCAT Registries American Journal of Medical Genetics. (Submitted Feb 2006)
- 80) BERTINO E, GIULIANI F, TONETTO P, FABRIS C, PROFETI C, MAGNANI C, MORO GE, ARSLANOGLU S. (2006): Randomized, controlled trial of breastfeeding versus formula feeding in extremely low birth weight infants. Pediatrics. Mar;117(3):985-6; author reply 986-7.