MASTER PEDIATRIC ENDOCRINOLOGY AND DIABETES

Curriculum Vitae

Personal Information

Name	Stefano Stagi
Address	Via XXI Luglio 9 52022 Cavriglia (Arezzo)
Telephone	+39 0555662585
E-mail	stefano.stagi@meyer.it; stefano.stagi@unifi.it
Nationality	Italian
Date of birth	14/02/1969

Position

Assistant Professor in Pediatrics

Education and training

Date	17/12/2010 to date
Title of qualification awarded	Assistant Professor
Name and type of organisation providing education	Department of Health Sciences of the University of Florence, Auxoendocrinology Unit, Anna Meyer Children's University Hospital, Florence .
Date	01/11/2012-31/10/2013
Title of qualification awarded	II Level Master Degree in Metabolic Bone Diseases. From Gene to the Care.
Name and type of organisation providing education	University of Florence, Largo Brambilla 3, 50134 Firenze
Date	01/01/2010-30/6/2010
Title of qualification awarded	Postgraduate Master Degree in rare Paediatric Diseases
Name and type of organisation providing education	Department of Paediatrics, University of Florence, Anna Meyer Children's University Hospital, viale Pieraccini 24, Florence, Italy.
Date	01/11/2007-31/05/2009
Title of qualification awarded	Fixed term Research Fellow Name Project: "Epiphysis and hypophysis. Possible relation between the secretion of melatonin, growth and time of maturation".

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Name and type of organisation providing education	Department of Paediatrics, University of Florence, Anna Meyer Children's University Hospital, via Luca Giordano 13, Florence, Italy.
Date	01/11/2004-31/10/2007
Title of qualification awarded	Fixed term Research Fellow Name Project: "Endocrine and non-endocrine factors that influence the period of biological maturation".
Name and type of organisation providing education	Department of Paediatrics, University of Florence, Anna Meyer Children's University Hospital, via Luca Giordano 13, Florence, Italy.
Date	01/11/200431/10/20005
Title of qualification awarded	II Level Master in Clinical and Laboratory Genetics
Name and type of organisation providing education	Department of Paediatrics, University of Florence, Anna Meyer Children's University Hospital, via Luca Giordano 13, Florence, Italy.
Date	01/11/2003-31/10/2004
Title of qualification awarded	Research Fellowship Name project: "Variations of urinary melatonin during overexposure to television and its possible pathogenetic role in idiopathic precocious puberty".
Name and type of organisation providing education	Department of Paediatrics, University of Florence, Anna Meyer Children's University Hospital, via Luca Giordano 13, Florence, Italy.
Date	01/10/1998-20/10/2003
Title of qualification awarded	Graduated in Paediatrics Thesis title: "Deficit assoluto di IgA e livelli di immunoglobuline in pazienti con ipotiroidismo congenito"
Name and type of organisation providing education	Department of Paediatrics, University of Florence, Anna Meyer Children's University Hospital, via Luca Giordano 13, Florence, Italy.
Final degree	Final degree 70/70 cum laude
Date	10/1998
Title of qualification awarded	Medical Board Medical school
Name and type of organisation providing education	University of Florence, Largo Brambilla 3, 50134 Firenze
Final degree	90/90
Date	April 1998
Title of qualification awarded	Degree in Medicine and Surgery (106/110) from the University of Florence, School of Medicine and Surgery, defending my dissertation entitled "Screening and follow-up of congenital hypothyroidism. The experience of the Centre in Florence" Supervisor: Prof. Roberto Salti.
Name and type of organisation providing education	Medical School, School of Medicine and Surgery, University of Florence, Largo Brambilla 3, 50134 Firenze
Final degree	106/110
Date	1983-1988
Title of qualification awarded	High school leaving qualification in classical studies, Liceo Classico "F. Petrarca" in Arezzo.

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Name and type of organisation providing education	Liceo Classico “F. Petrarca”, Arezzo
Date	04/2014
Title of qualification awarded	Grant of the European Society for Pediatric Endocrinology (ESPE): ESPE diabetes/obesity school in Tel Aviv, 24-26 April 2014.
Date	09/2010
Title of qualification awarded	Member of the European Society for Paediatric Endocrinology (ESPE).
Date	09/2013
Title of qualification awarded	Member of the Italian Society of Pediatric Endocrinology and Diabetology (SIEDP).

Mother-tongue: Italian

Other

Self-evaluation	Understanding				Speaking				Writing	
European level (*)	Lissening		Reeding		Spoken interaction		Spoken production			
English	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate
Spanish	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate
French	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate	B2	Upper intermediate

(*) Quadro comune europeo di riferimento per le lingue

Computer skills: WORD, EXCEL, POWER POINT

Educational activities:

Time contract (from 2004 to date) assistant professor of Paediatric Endocrinology and Auxology at the Medical Degree Course of the University of Florence.

Time contract (from 2006 to 2007) assistant professor of Paediatric Endocrinology and Auxology at the Course of Sports Sciences and Techniques in the Field of Educational Theory and Techniques of the Stages of Development (E.T.T. Preventative and Compensatory Motor and Sports Activities Individual Sports). .

Time contract (from 2004 to 2009) assistant professor of Paediatric Endocrinology and Auxology at the Postgraduate School in Paediatrics of the University of Florence.

Assistant professor (from 2015 to date) of Auxology and Endocrinology at the Postgraduate School in Clinical Genetics, University of Florence.
Assistant professor (from 2015 to date) of Auxology and Endocrinology at the Postgraduate School in Paediatrics of the University of Florence.

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Pubblications

1. **Stagi S**, Lepri G, Rigante D, Matucci Cerinic M, Falcini F. Cross-Sectional Evaluation of Plasma Vitamin D Levels in a Large Cohort of Italian Patients with Pediatric Autoimmune Neuropsychiatric Disorders Associated with Streptococcal Infections. J Child Adolesc Psychopharmacol. 2017 Nov 7. doi: 10.1089/cap.2016.0159.
2. Serranti D, Indolfi G, Nebbia G, Cananzi M, D'Antiga L, Ricci S, **Stagi S**, Azzari C, Resti M; Italian Study Group for Treatment of Chronic Hepatitis C in Children. Transient Hypothyroidism and Autoimmune Thyroiditis in Children with Chronic Hepatitis C Treated with Pegylated-Interferon-α-2b and Ribavirin. Pediatr Infect Dis J. 2017 Sep 22. doi: 10.1097/INF.0000000000001791.
3. Pelosi P, Lapi E, Cavalli L, Verrotti A, Pantaleo M, de Martino M, **Stagi S**. Bone Status in a Patient with Insulin-Like Growth Factor-1 Receptor Deletion Syndrome: Bone Quality and Structure Evaluation Using Dual-Energy X-Ray Absorptiometry, Peripheral Quantitative Computed Tomography, and Quantitative Ultrasonography. Front Endocrinol (Lausanne). 2017 Sep 5;8:227. doi: 10.3389/fendo.2017.00227.
4. **Stagi S**, Scalini P, Farello G, Verrotti A. Possible effects of an early diagnosis and treatment in patients with growth hormone deficiency: the state of art. Ital J Pediatr. 2017 Sep 16;43(1):81. doi: 10.1186/s13052-017-0402-8.
5. Mori F, Sarti L, Barni S, Pucci N, Belli F, **Stagi S**, Novembre E. Donkey´s Milk Is Well Accepted and Tolerated by Infants With Cow´s Milk Food Protein-Induced Enterocolitis Syndrome: A Preliminary Study. J Investig Allergol Clin Immunol. 2017;27(4):269-271. doi: 10.18176/jiaci.0167.
6. **Stagi S**, Ricci F, Bianconi M, Sammarco MA, Municchi G, Toni S, Lenzi L, Verrotti A, de Martino M. Retrospective Evaluation of Metformin and/or Metformin Plus a New Polysaccharide Complex in Treating Severe Hyperinsulinism and Insulin Resistance in Obese Children and Adolescents with Metabolic Syndrome. Nutrients. 2017 May 20;9(5). pii: E524. doi: 10.3390/nu9050524.
7. Maggioli C, **Stagi S**. Bone modeling, remodeling, and skeletal health in children and adolescents: mineral accrual, assessment and treatment. Ann Pediatr Endocrinol Metab. 2017 Mar;22(1):1-5. doi: 10.6065/apem.2017.22.1.1.
8. Palazzo V, Provenzano A, Becherucci F, Sansavini G, Mazzinghi B, Orlandini V, Giunti L, Roperto RM, Pantaleo M, Artuso R, Andreucci E, Bargiacchi S, Traficante G, **Stagi S**, Murer L, Benetti E, Emma F, Giordano M, Rivieri F, Colussi G, Penco S, Manfredini E, Caruso MR, Garavelli L, Andrulli S, Vergine G, Miglietti N, Mancini E, Malaventura C, Percesepe A, Grosso E, Materassi M, Romagnani P, Giglio S. The genetic and clinical spectrum of a large cohort of patients with distal renal tubular acidosis. Kidney Int. 2017 May;91(5):1243-1255. doi: 10.1016/j.kint.2016.12.017.
9. **Stagi S**, Di Tommaso M, Scalini P, Sandini E, Masoni F, Chiarelli F, Verrotti A, Giglio S, Romano S, de Martino M. Cross-sectional study shows that impaired bone mineral status and metabolism are found in nonmosaic triple X syndrome. Acta Paediatr. 2017 Apr;106(4):619-626. doi: 10.1111/apa.13744.
10. Iughetti L, Tornese G, Street ME, Napoli F, Giavoli C, Antoniazzi F, **Stagi S**, Luongo C, Azzolini S, Ragusa L, Bona G, Zecchino C, Aversa T, Persani L, Guazzarotti L, Zecchi E, Pietropoli A, Zucchini S. Long-term safety and efficacy of Omnitrope®, a somatropin biosimilar, in children requiring growth hormone treatment: Italian interim analysis of the PATRO Children study. Ital J Pediatr. 2016 Nov 3;42(1):93.
11. **Stagi S**, Cavalli L, Cavalli T, de Martino M, Brandi ML. Peripheral quantitative computed tomography (pQCT) for the assessment of bone strength in most of bone affecting conditions in developmental age: a review. Ital J Pediatr. 2016 Sep 26;42(1):88.
12. **Stagi S**, Di Tommaso M, Manoni C, Scalini P, Chiarelli F, Verrotti A, Lapi E, Giglio S, Dosa L, de Martino M. Bone Mineral Status in Children and Adolescents with Klinefelter Syndrome. Int J Endocrinol. 2016;2016:3032759. doi: 10.1155/2016/3032759.
13. **Stagi S**, Manoni C, Scalini P, Chiarelli F, Verrotti A, Cecchi C, Lapi E, Giglio S, Romano S, de Martino M. Bone mineral status and metabolism in patients with Williams-Beuren syndrome. Hormones (Athens). 2016 Jul;15(3):404-412. doi: 10.14310/horm.2002.1683.
14. **Stagi S**, Lasorella S, Piccorossi A, Iapadre G, Verrotti A. Cessation of epilepsy therapy in children. Expert Rev Neurother. 2016 May;16(5):549-59. doi:10.1586/14737175.2016.1168296.
15. **Stagi S**, di Tommaso M, Scalini P, Lapi E, Losi S, Bencini E, Masoni F, Dosa L, Becciani S, de Martino M. Triple X syndrome and puberty: focus on the hypothalamus-hypophysis-gonad axis. Fertil Steril. 2016 Jun;105(6):1547-53. doi: 10.1016/j.fertnstert.2016.02.019.

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16. Verrotti A, Prezioso G, **Stagi S**, Paolino MC, Parisi P. Pharmacological considerations in the use of stiripentol for the treatment of epilepsy. Expert Opin Drug Metab Toxicol. 2016;12(3):345-52. doi: 10.1517/17425255.2016.1145657.
17. Bartolini E, **Stagi S**, Scalini P, Bianchi A, Ciccarone A, Mascalchi M. Central precocious puberty due to hypothalamic hamartoma in neurofibromatosis type 1.Hormones (Athens). 2016 Jan;15(1):144-6.
18. Giordano F, Spacca B, Danti A, Taverna M, Losi S, **Stagi S**, Genitori L. Amenorrhea after Endoscopic Third Ventriculostomy for a Failed Shunt in Spina Bifida: Case Report and Review of the Literature. Pediatr Neurosurg. 2016;51(1):35-41. doi: 10.1159/000441254.
19. **Stagi S**, Traficante G, Lapi E, Pantaleo M, Becciani S, Mortilla M, Seminara S, de Martino M. Agenesis of internal carotid artery associated with isolated growth hormone deficiency: a case report and literature review. BMC Endocr Disord. 2015 Oct 19;15:58. doi: 10.1186/s12902-015-0037-y.
20. **Stagi S**, Gulino AV, Lapi E, Rigante D. Epigenetic control of the immune system: a lesson from Kabuki syndrome. Immunol Res. 2016 Apr;64(2):345-59. doi: 10.1007/s12026-015-8707-4.
21. **Stagi S**, Lapi E, Pantaleo M, Carella M, Petracca A, De Crescenzo A, Zelante L, Riccio A, de Martino M. A new case of de novo 6q24.2-q25.2 deletion on paternal chromosome 6 with growth hormone deficiency: a twelve-year follow-up and literature review. BMC Med Genet. 2015 Aug 23;16:69. doi:10.1186/s12881-015-0212-z.
22. **Stagi S**, Lapi E, Martini E, de Martino M. Giant multiple bladder diverticula in Williams-Beuren syndrome. Kidney Int. 2015 Aug;88(2):416. doi:10.1038/ki.2014.312.
23. **Stagi S**, Cavalli L, Ricci S, Mola M, Marchi C, Seminara S, Brandi ML, decMartino M. Parathyroid Hormone Levels in Healthy Children and Adolescents. Horm Res Paediatr. 2015;84(2):124-9. doi: 10.1159/000432399.
24. **Stagi S**, Rigante D, Lepri G, Matucci Cerinic M, Falcini F. Severe vitamin D deficiency in patients with Kawasaki disease: a potential role in the risk to develop heart vascular abnormalities? Clin Rheumatol. 2016 Jul;35(7):1865-72. doi: 10.1007/s10067-015-2970-6.
25. **Stagi S**, Iurato C, Lapi E, Cavalli L, Brandi ML, de Martino M. Bone status in genetic syndromes: a review. Hormones (Athens). 2015 Jan-Mar;14(1):19-31.
26. **Stagi S**, Lapi E, Seminara S, Pelosi P, Del Greco P, Capirchio L, Strano M, Giglio S, Chiarelli F, de Martino M. Policaptil Gel Retard significantly reduces body mass index and hyperinsulinism and may decrease the risk of type 2 diabetes mellitus (T2DM) in obese children and adolescents with family history of obesity and T2DM. Ital J Pediatr. 2015 Feb 15;41:10. doi: 10.1186/s13052-015-0109-7.
27. **Stagi S**, Lapi E, Romano S, Bargiacchi S, Brambilla A, Giglio S, Seminara S, de Martino M. Determinants of vitamin d levels in children and adolescents with down syndrome. Int J Endocrinol. 2015;2015:896758. doi: 10.1155/2015/896758.
28. **Stagi S**, Cavalli L, Congiu L, Scusa MF, Ferlini A, Bigoni S, Benincasa A, Rossi B, Pini G. Thyroid function in Rett syndrome. Horm Res Paediatr. 2015;83(2):118-25. doi: 10.1159/000370066.
29. **Stagi S**, Gragnani SG, Michelotti F, de Martino M. A particular case of pubic pain. J Rheumatol. 2014 Dec;41(12):2487-9. doi: 10.3899/jrheum.140090.
30. **Stagi S**, Pelosi P, Strano M, Poggi G, Manoni C, de Martino M, Seminara S. Determinants of Vitamin D Levels in Italian Children and Adolescents: A Longitudinal Evaluation of Cholecalciferol Supplementation versus the Improvement of Factors Influencing 25(OH)D Status. Int J Endocrinol. 2014;2014:583039. doi: 10.1155/2014/583039.
31. **Stagi S**, Manoni C, Cirello V, Covelli D, Giglio S, Chiarelli F, Seminara S, de Martino M. Diabetes mellitus in a girl with thyroid hormone resistance syndrome: a little recognized interaction between the two diseases. Hormones (Athens). 2014 Oct-Dec;13(4):561-7. doi: 10.14310/horm.2002.1502.
32. **Stagi S**, Lapi E, Pantaleo M, Traficante G, Giglio S, Seminara S, de Martino M. A SOX3 (Xq26.3-27.3) duplication in a boy with growth hormone deficiency, ocular dyspraxia, and intellectual disability: a long-term follow-up and literature review. Hormones (Athens). 2014 Oct-Dec;13(4):552-60. doi: 10.14310/horm.2002.1523.
33. **Stagi S**, Del Greco P, Ricci F, Iurato C, Poggi G, Seminara S, de Martino M. Hydrocortisone malabsorption due to polyethylene glycols (Macrogol 3350) in a girl with congenital adrenal insufficiency. Ital J Pediatr. 2014 Sep 26;40:78. doi: 10.1186/s13052-014-0078-2.
34. **Stagi S**, Rigante D, Lepri G, Bertini F, Matucci-Cerinic M, Falcini F. Evaluation of autoimmune phenomena in patients with pediatric autoimmune neuropsychiatric disorders associated with streptococcal infections (PANDAS). Autoimmun Rev. 2014 Dec;13(12):1236-40. doi: 10.1016/j.autrev.2014.08.009.

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35. **Stagi S**, Pucci N, Di Grande L, de Libero C, Caputo R, Pantano S, Mattei I, Mori F, de Martino M, Novembre E. Increased incidence of thyroid dysfunction and autoimmunity in patients with vernal keratoconjunctivitis. Int J Endocrinol. 2014;2014:804870. doi: 10.1155/2014/804870.

36. **Stagi S**, Bertini F, Cavalli L, Matucci-Cerinic M, Brandi ML, Falcini F. Determinants of vitamin D levels in children, adolescents, and young adults with juvenile idiopathic arthritis. J Rheumatol. 2014 Sep;41(9):1884-92. doi:10.3899/jrheum.131421.

37. **Stagi S**, Pucci N, di Grande L, de Libero C, Caputo R, Pantano S, Seminara S, de Martino M, Novembre E. Increased prevalence of growth hormone deficiency in patients with vernal keratoconjuntivitis; an interesting new association. Hormones (Athens). 2014 Jul-Sep;13(3):382-8. doi: 10.14310/horm.2002.1499.

38. **Stagi S**, Cavalli L, Bertini F, Signorini C, Matucci Cerinic M, de Martino M, Brandi ML, Falcini F. Comparison of bone mass and quality determinants in adolescents and young adults with juvenile systemic lupus erythematosus (JSLE) and juvenile idiopathic arthritis (JIA). Lupus. 2014 Nov;23(13):1392-406. doi:10.1177/0961203314543916.

39. von Scheven E, Corbin KJ, **Stagi S**, Cimaz R. Glucocorticoid-associated osteoporosis in chronic inflammatory diseases: epidemiology, mechanisms, diagnosis, and treatment. Curr Osteoporos Rep. 2014 Sep;12(3):289-99. doi:10.1007/s11914-014-0228-x.

40. **Stagi S**, Lapi E, Cecchi C, Chiarelli F, D'Avanzo MG, Seminara S, de Martino M. Williams-beuren syndrome is a genetic disorder associated with impaired glucose tolerance and diabetes in childhood and adolescence: new insights from a longitudinal study. Horm Res Paediatr. 2014;82(1):38-43. doi: 10.1159/000360476.

41. **Stagi S**, Cavalli L, Seminara S, de Martino M, Brandi ML. The ever-expanding conundrum of primary osteoporosis: aetiopathogenesis, diagnosis, and treatment. Ital J Pediatr. 2014 Jun 7;40:55. doi: 10.1186/1824-7288-40-55.

42. **Stagi S**, Lapi E, D'Avanzo MG, Perferi G, Romano S, Giglio S, Ricci S, Azzari C, Chiarelli F, Seminara S, de Martino M. Coeliac disease and risk for other autoimmune diseases in patients with Williams-Beuren syndrome. BMC Med Genet. 2014 May 23;15:61. doi: 10.1186/1471-2350-15-61.

43. **Stagi S**, Bertini F, Rigante D, Falcini F. Vitamin D levels and effects of vitamin D replacement in children with periodic fever, aphthous stomatitis, pharyngitis, and cervical adenitis (PFAPA) syndrome. Int J Pediatr Otorhinolaryngol. 2014 Jun;78(6):964-8. doi: 10.1016/j.ijporl.2014.03.026.

44. **Stagi S**, Cavalli L, Bertini F, de Martino M, Cerinic MM, Brandi ML, Falcini F. Vitamin D levels in children, adolescents, and young adults with juvenile-onset systemic lupus erythematosus: a cross-sectional study. Lupus. 2014 Sep;23(10):1059-65. doi: 10.1177/0961203314532564.

45. **Stagi S**, Cavalli L, Signorini C, Bertini F, Cerinic MM, Brandi ML, Falcini F. Bone mass and quality in patients with juvenile idiopathic arthritis: longitudinal evaluation of bone-mass determinants by using dual-energy x-ray absorptiometry, peripheral quantitative computed tomography, and quantitative ultrasonography. Arthritis Res Ther. 2014 Mar 31;16(2):R83. doi: 10.1186/ar4525.

46. **Stagi S**, Losi S, Chiarelli F, de Martino M, Falcini F. Kawasaki disease in a girl with Turner syndrome: a remarkable association. Ital J Pediatr. 2014 Feb 28;40(1):24. doi: 10.1186/1824-7288-40-24.

47. **Stagi S**, Cavalli L, Iurato C, Seminara S, Brandi ML, de Martino M. Bone metabolism in children and adolescents: main characteristics of the determinants of peak bone mass. Clin Cases Miner Bone Metab. 2013 Sep;10(3):172-9.

48. **Stagi S**, Cavalli L, Iurato C, Seminara S, Brandi ML, de Martino M. Bone health in children and adolescents: the available imaging techniques. Clin Cases Miner Bone Metab. 2013 Sep;10(3):166-71.

49. **Stagi S**, Lapi E, Pantaleo M, Chiarelli F, Seminara S, de Martino M. Type II diabetes and impaired glucose tolerance due to severe hyperinsulinism in patients with 1p36 deletion syndrome and a Prader-Willi-like phenotype. BMC Med Genet. 2014 Jan 30;15:16. doi: 10.1186/1471-2350-15-16.

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50. **Stagi S**, Lapi E, Seminara S, Guarducci S, Pantaleo M, Giglio S, Chiarelli F, de Martino M. Long-term auxological and endocrinological evaluation of patients with 9p trisomy: a focus on the growth hormone-insulin-like growth factor-I axis. BMC Endocr Disord. 2014 Jan 8;14:3. doi: 10.1186/1472-6823-14-3.

51. **Stagi S**, Cavalli L, Bertini F, Matucci Cerinic M, Luisa Brandi M, Falcini F. Cross-sectional and longitudinal evaluation of bone mass and quality in children and young adults with juvenile onset systemic lupus erythematosus (JSLE): role of bone mass determinants analyzed by DXA, PQCT and QUS. Lupus. 2014;23(1):57-68. doi: 10.1177/0961203313511679.

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54. Gasperini S, **Stagi S**, Gasperini U, Guerrini R, la Marca G, Donati MA. Orange-colored diapers as first sign of Lesch-Nyhan disease in an asymptomatic infant. Pediatr Nephrol. 2010 Nov;25(11):2373-4. doi: 10.1007/s00467-010-1557-3.

55. **Stagi S**, Galli L, Cecchi C, Chiappini E, Losi S, Gattinara CG, Gabiano C, Tovo PA, Bernardi S, Chiarelli F, de Martino M. Final height in patients perinatally infected with the human immunodeficiency virus. Horm Res Paediatr. 2010;74(3):165-71. doi: 10.1159/000281018.

56. **Stagi S**, Lapi E, Gambineri E, Manoni C, Genuardi M, Colarusso G, Conti C, Chiarelli F, de Martino M, Azzari C. Bone density and metabolism in subjects with microdeletion of chromosome 22q11 (del22q11). Eur J Endocrinol. 2010 Aug;163(2):329-37. doi: 10.1530/EJE-10-0167.

57. **Stagi S**, Masi L, Capannini S, Cimaz R, Tonini G, Matucci-Cerinic M, de Martino M, Falcini F. Cross-sectional and longitudinal evaluation of bone mass in children and young adults with juvenile idiopathic arthritis: the role of bone mass determinants in a large cohort of patients. J Rheumatol. 2010 Sep;37(9):1935-43. doi: 10.3899/jrheum.091241.

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60. **Stagi S**, Lapi E, Sani I, Bindi G, Quaranta MR, Salti R, Chiarelli F. Growth hormone neurosecretory dysfunction in a boy with hypohidrotic/anhidrotic ectodermal dysplasia: definition of short stature, molecular characterization and long-term hGH replacement treatment to final height. J Pediatr Endocrinol Metab. 2009 Sep;22(9):853-8.

61. **Stagi S**, Lapi E, Gambineri E, Salti R, Genuardi M, Colarusso G, Conti C, Jenuso R, Chiarelli F, Azzari C, de Martino M. Thyroid function and morphology in subjects with microdeletion of chromosome 22q11 (del(22)(q11)). Clin Endocrinol (Oxf). 2010 Jun;72(6):839-44. doi: 10.1111/j.1365-2265.2009.03736.x.

62. Galluzzi F, Quaranta MR, Salti R, **Stagi S**, Nanni L, Seminara S. Diagnosis of growth hormone deficiency by using the arginine provocative test: is it possible to shorten testing time without altering validity? Horm Res. 2009;72(3):142-5. doi: 10.1159/000232487.

63. **Stagi S**, Bindi G, Lapi E, Giovannucci-Uzielli ML, Salti R, Chiarelli F. Congenital hypothyroidism in Young-Simpson syndrome. J Pediatr Endocrinol Metab. 2008 Nov;21(11):1089-92.

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65. **Stagi S**, Bindi G, Galluzzi F, Lapi E, Salti R, Chiarelli F. Partial growth hormone deficiency and changed bone quality and mass in type I trichorhinophalangeal syndrome. Am J Med Genet A. 2008 Jun 15;146A(12):1598-604. doi: 10.1002/ajmg.a.32348.

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66. **Stagi S**, Galluzzi F, Bindi G, Lapi E, Cecchi C, Salti R, Chiarelli F. Precocious puberty in a girl with floating-harbor syndrome. J Pediatr Endocrinol Metab. 2007 Dec;20(12):1333-7.

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Abstract

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