



INFORMAZIONI PERSONALI

GIULIETTA SCUVERA

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Sesso Femmina Data di Nascita 22/01/1982 Nazione ITALIA

POSIZIONE RICOPERTA DIRIGENTE MEDICO

ESPERIENZA PROFESSIONALE

Da Dicembre 2020 DIRIGENTE MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

U.O.S.D. Genetica Medica. Consulenza genetica in ambito preconcezionale e prenatale per la definizione del rischio riproduttivo di coppia. Pianificazione di eventuale diagnosi prenatale.

Attività o Settore SANITA'

Da Marzo 2020 a Novembre 2020 MEDICO

FONDAZIONE TELETHON

Consulente in regime di P.I. presso il Servizio Info_Rare della Fondazione Telethon

Attività o Settore Attivita' professionali, scientifiche e tecniche

Da Gennaio 2020 a Dicembre 2020 MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

Incarico professionale in regime di P.I. presso la UOSD Pediatria ad Alta Intensità di Cura come medico genetista: inquadramento diagnostico e follow up dei pazienti con neurofibromatosi 1 e sindromi genetiche.

Attività o Settore SANITA'

Da Gennaio 2019 a Dicembre 2019 MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

Contratto di collaborazione coordinata e continuativa (CO. CO. CO.) presso la UOSD Pediatria ad Alta Intensità di Cura come medico genetista: inquadramento diagnostico e follow up dei pazienti con neurofibromatosi 1 e sindromi genetiche.

Attività o Settore SANITA'

Da Gennaio 2018 a Dicembre 2018 MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

Borsista presso la UOSD Pediatria ad Alta Intensità di Cura come medico genetista: inquadramento diagnostico e follow up dei pazienti con neurofibromatosi 1 e sindromi genetiche.

Attività o Settore SANITA'

Da Novembre 2017 a Dicembre 2017 MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

Contratto di collaborazione occasionale presso la UOSD Pediatria ad Alta Intensità di Cura come medico genetista: inquadramento diagnostico e follow up dei pazienti con neurofibromatosi 1 e sindromi genetiche.

Attività o Settore SANITA'

Da Gennaio 2016 a Dicembre 2016 MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

Contratto di collaborazione coordinata e continuativa (CO. CO. CO.) presso la UOSD Pediatria ad Alta Intensità di Cura come medico genetista: inquadramento diagnostico e follow up dei pazienti con neurofibromatosi 1 e sindromi genetiche.

Attività o Settore SANITA'



Da Giugno 2015 a Dicembre 2015 MEDICO

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico Milano

Contratto di collaborazione a progetto (CO.CO.PRO) presso la UOSD Pediatria ad Alta Intensità di Cura come medico genetista: inquadramento diagnostico e follow up dei pazienti con neurofibromatosi 1 e sindromi genetiche.

Attività o Settore SANITA'

ISTRUZIONE E FORMAZIONE

Da Luglio 2009 a Gennaio 2015 Diploma di Specializzazione in Genetica Medica

Università degli Studi di Milano

Da Ottobre 2000 a Luglio 2008 Laurea in Medicina e Chirurgia

Università degli Studi di Pavia

COMPETENZE PERSONALI

Lingua madre Italiano

Altre lingue

	COMPRESIONE		PARLATO		PRODUZIONE SCRITTA
	Ascolto	Lettura	Interazione	Produzione orale	
Inglese	A2	B1	A2	B1	A2
Francese	B1	B2	B1	B1	A1

Livelli: A 1/2 Livello Base - B 1/2 Livello Intermedio - C 1/2 Livello Avanzato

Quadro Comune Europeo di Riferimento delle Lingue

Patente di guida B

ALLEGATI

Scuvera_publicazioni.pdf

Dati personali Autorizzo il trattamento dei miei dati personali ai sensi del Decreto Legislativo 30 giugno 2003, n. 196 'Codice in materia di protezione dei dati personali'.

Coautrice volume "Le Malattie Rare in età giovane-adulta: dal sospetto diagnostico alla gestione clinica". (casa editrice Hippocrates)

Family burden of children suffering from Epidermolysis Bullosa. De Stefano S, Grassi FS, Lalatta F, Scuvera G, Brena M, Grillo P, Peves Rios WE, Guez S. *G Ital Dermatol Venereol*. 2020 Oct 9. doi: 10.23736/S0392-0488.20.06613-4.

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Giulia Melloni, Marica Eoli, Claudia Cesaretti, Donatella Bianchessi, Maria Cristina Ibba, Silvia Esposito, Giulietta Scuvera, Guido Morcaldi, Roberto Micheli, Elena Piozzi, Sabrina Avignone, Luisa, Chiapparini, Chiara Pantaleoni, Federica Natacci, Gaetano Finocchiaro, Veronica Saletti. Risk of Optic Pathway Glioma in Neurofibromatosis Type 1: No Evidence of Genotype-Phenotype Correlations in A Large Independent Cohort. *Cancers (Basel)*, 2019 Nov 21. 11 (12)

Benedetta Beltrami, Elisabetta Prada, Gianluca Tolva, Giulietta Scuvera, Rosamaria Silipigni, Daniela Graziani, Gaetano Bulfamante, Cristina Gervasini, Paola Marchisio, Donatella Milani. Unexpected Phenotype in a Frameshift Mutation of *PTCH1*. *Mol Genet Genomic Med*. 2019 Oct 2, e987

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AustralianOvarianCancerStudy Group, Haiman CA, Schumacher F, Henderson BE, Le Marchand L, Lindblom A, Margolin S, Hooning MJ, Hollestelle A, Kriege M, Koppert LB, Hopper JL, Southey MC, Tsimiklis H, Apicella C, Slettedahl S, Toland AE, Vachon C, Yannoukakos D, Giles GG, Milne RL, McLean C, Fasching PA, Ruebner M, Ekici AB, Beckmann MW, Brenner H, Dieffenbach AK, Arndt V, Stegmaier C, Ashworth A, Orr N, Schoemaker MJ, Swerdlow A, García-Closas M, Figueroa J, Chanock SJ, Lissowska J, Goldberg MS, Labrèche F, Dumont M,

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