**CURRICULUM VITAE**

Barbara BARDONI
Born in Montebello della Battaglia (Italy), June 20, 1963.
Married – Two children

PRESENT POSITION:

Research Director INSERM Institute of Cellular and Molecular Pharmacology - CNRS UMR7275, Valbonne (France) bardoni@ipmc.cnrs.fr

EDUCATION

**1986** Faculty of Science-University of Pavia (Italy) - Ph.D. in Biology (Cum Laude).
**1987-88** Specialization School of Genetics - Faculty of Science-University of Pavia (Italy)
**2004** Habilitation à Diriger la Recherche (HDR) Université Louis Pasteur (Strasbourg)

PROFESSIONAL ACTIVITY

**1988-90** Research fellow at the Dep. of Human and Hereditary Pathology, Faculty of Medicine, University of Pavia (Italy).
**1991-2002** Researcher at the Department of Human and Hereditary Pathology, Faculty of Medicine, University of Pavia (Italy)
**1997-2002** Visiting scientist at IGBMC - Illkirch (France), in Prof. J-L Mandel's laboratory
**2002-2004** Researcher (CR1) INSERM at IGBMC - Illkirch (France)
**2005 - December 2007** Group leader at the CNRS UMR6543- Faculty of Medicine, Nice, (France)
**January 2008 to present** Group leader at the CNRS UMR7275 - Valbonne (France)

HONORS AND GRANTS

**Award and Fellowship**

**1988:** Price "Rino Minoprio" for the Excellence in Human Genetics; **1988-90**: Fellowship "AnnaVilla Rusconi"; **1994**: EMBO Short-Term Fellowship; **1996-98**: Italian Telethon Fellowship; **1998-99**: FRM Fellowship; **2004:** ATIP**; 2007:** ATIP PLUS; **2014**: 6ème Prix du Conseil General06.

**GRANTS:**

**PI: NIMH/NICHD/FRAXA cofunding** (2001-04), **ATIP** (2004-07); **FRM New laboratory** (2005) **GIS Maladies Rares** (2005-08); **ANR-Neurosci** (2006-09; 2013-17); **ATIP Plus** (2007-09); **BQR** (2009); **MRT Thèse Flechée** (2009-12); **FRM TEAM** (20010-13; 2014-17); **FRAXA Foundation** (2000; 2006); **Jerôme Lejeune Foundation** (2004/10; 2014/16; 2018-20); **IngFRM** (2014); **FRC** (2019-20); **AFM** (2014-16); **Prematuration UCA-SATT** (2019)

**Co-Investigator: EEC (**1995-96)**; HFSP** (2002-2005)**; ULP** (2002)**; ANR Blanc** (2006-09)**; Marie Curie-RI** (2009-12)**; ANR E-RARE** (2010-13)**; LabEX "Signalife** (2012-19)**; Fondation Maladies Rares** (2014-16)**; ANR-AAP** (2015-19)**; FRAXA Foundation** (2016-18)**.**

**Exchanging grants: France-Canada cooperation grant**. PI: E.W. Khandjian (2004); **« Ambassade de Canada » Québec**, Prof. E. Khandjian **(**2008)**; « G. Galilei » (Italy)** Dr. M.V. Catania (2007-08); **FAST (Australia)** Prof. J. Gecz (2009-10); **Ulysse** (Ireland) Prof. D. Tropea (2019).

**Fellowshipw for members of the Team : 2007-09**: «Marie Curie –Post doc» L. Davidovic ; **2007:** «Telecom Autisme- Post doc» M. Bensaid; **2007**: « **Boehringer Ingelheim Fonds**» M. Melko; **2007:** «FRM –Fin thèse» E. Bechara; **2008**: «FRM – Fin thèse» M. Melko; **2009**: «La Ligue contre le cancer – Fin thèse» M. Melko; **2010-11**: «Ville de Nice - Post doc» O. Khalfallah; **2010**: «Embo Short Term»

M. Melko; **2012** « Boehringer Ingelheim Fonds» 2 months stage - D. Masini; «ARC – Fin thèse» S. Abekhoukh; **2013-15**: AFM post-doc: S. Zongaro.

INVITED SPEAKER IN INTERNATIONAL MEETINGS (A selection)

**1.** Banbury Meeting – Fragile X Syndrome: Basic mechanisms and Treatment Implications - New York (USA) April 9-12, 2006
**2.** FRAXA Research Foundation/Investigators Meeting - Durham NH (USA), September 21-24, 2008
**3.** FRAXA Research Foundation -Investigators Meeting, Durham NH (USA), Mai 2-5, 2010.
**4.** Jacques Monod Conference «Mental Retardation: from genes to synapses, functions and dysfunctions ». Roscoff (France), October 15-19, 2010.
**7.** Jacques Monod Conference " Mechanism of Intellectual disability: from genes to treatment", – Roscoff (France) October 4-7, 2012.
**6.** 16th International Workshop on Fragile X and Other Early-Onset Cognitive Disorders, Adelaide (Australia) September 17-20, 2013
**7**. Dynamics of synaptic signaling: from molecular to integrated levels - Societé Française de Neuroscience, Montpellier, May 19- 22, 2015.
**8**. ECNP Targeted Network Meeting (TNM) on "Common mechanisms and therapeutic avenues for Down syndrome and other genetic developmental disorders" September 2016, Vienna, (Austria)
**9.** ECNP Targeted Network Meeting (TNM) on "Common mechanisms and therapeutic avenues for Down syndrome and other genetic developmental disorders" September 2017, Paris (France)
**10**. XVI Workshop on Fragile X and other early-onset neurodevelopmental disorders, 12-16 October, 2017 – Montreal (Canada)
**11.** Spring Hippocampus Research Conference, 2-6 June, 2019 – Taormina (Italy).

EDITORIAL ACTIVITY :

**Board member:** *PLOS ONE*; *Frontiers in Molecular Neuroscience*; *Frontiers in Synaptic Neuroscience*; *Frontiers in Molecular Bioscience* section *RNA Proteins Network.*
**2013 Guest Editor** : *Neuroscience and Biobehavioural Reviews* : « Common Mechanisms in Intellectual Disabilities: A Challenge for Translational Outlooks »

OTHER PROFESSIONAL EXPERIENCES

Referee «ad hoc» for the following journals***:*** *Cell Rep ; E.J. Hum. Genet.; E.J. Neurosci.; HMG; J. Mol. Biol., JMG; J. Neurosci.; Mol. Psychiatry;Nature; MCB; Neuroscience; Neurobiol. Dis.; NAR; PLOS Genet; PLOS One; PNAS; Science, Trends Genet; PNAS.*
Referee «ad hoc» for *AFM (Association Française contre les Myopathies); ANR (Agence Nationale de la Recherche); Horizon Program (Netherland Genomics Initiative); )*; *ISF (Israel Science Foundation); Italian Telethon; Italian Ministry of Health;*NWO Division for Earth and Life Sciences *(The Netherlands); European Science Foundation; Health Research Council of NZ*
**2008** Member of the evaluation committee for Life Science Master of University of Nice
**2010** Member of a visiting committee of AERES
Member of an evaluation committee for a researcher position (MCU) at the University of Nice
**2012-17** Member of Laboratory Council of Institute of Molecular and Cellular Pharmacology
**2014-2016** Member of the Selection committee of LabEx Signalife PhD Program
**2015- to present**: Member of Scientific Committee of « Fondation J. Lejeune »
**2016-19** Member of Scientific Committee of LabEx « Signalife »
**2016** Member of Evaluation Committee ATIP –AVIESAN
**2016-21** Member of Evaluation Committee CSS2-INSERM
**2018** Member HCERES visiting committee of Imagine Institute – Paris
Teacher at the Big@UCA Summer School "*Social Cognition*", Nice (France), August 27/31.

CONFERENCE & WORKSHOP ORGANIZATION

**2010** Chair of the Jacques Monod Conference *«Mental Retardation: from genes to synapses, functions and dysfunctions »* Roscoff (France), October 7/11.

**2013** Chair of *« G-Quadruplex Day»*, Nice (France), March 22
Member of the Scientific Board of the Meeting *« XVI Workshop on Fragile X and other early-onset neurodevelopmental disorders »* Adelaide (Australia), 17/20 September.
Chair of the conference *«RNA metabolism:Cancer, development &disease »* Nice (France), December 4/6

**2016** Chair of the Workshop *« RNA Metabolism from transcription to Degradation»,* Valbonne (France), May 26

**2018** Chair of the conference *«The multiple facets of RNA in development and in disease»*, Nice (France), February 7/8
Member of Scientific Board of *SIFRARN Meeting*, Nancy (France), November 22

**2019** Member of Scientific Board of *« XIX Workshop on Fragile X and other early-onset neurodevelopmental disorders »* Sorrento (Italy), September 18/21

**List of Publication**

Profile URL : http://www.researcherid.com/rid/F-9918-2013
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**1.** Raimondi E, **Bardoni B**, Rinaldi E, Camerino G. (1987) - A TaqI RFLP detecting single copy fragment (G80) from chromosome 7 p13-p15 (D7S373). *Nucleic Acid Res.,* 15, 7653

**2. Bardoni B**, Sampietro M, Romano M, Crapanzano M, Mannucci PM, Camerino G. (1988) - Characterization of a partial deletion of the factor VIII gene in one hemophiliac with inhibitor. *Hum. Genet.,* 79, 86-88.

**3. Bardoni B**, Guioli S, Raimondi E, Heilig R, Mandel JL, Ottolenghi S, Camerino G. (1988) - Isolation and characterization of a family of sequences dispersed on the human X chromosome. *Genomics,* 3, 32-38.

**4. Bardoni B**, Guioli S, Maserati E, Maraschio P, Camerino G. (1988) - A highly conserved sequence on the short arm of chromosome 7 detects multiple polymorphisms. *Hum. Genet*., 81, 23-25

**5.** Guioli S, Arveiler B, **Bardoni B**, Notarangelo LD, Panina P, Duse M, Ugazio A, Oberlé I, de Saint Basile B, Mandel JL, Camerino G. (1989) - Close linkage of probe p212 (DXS178) to X-linked agammaglobulinemia. *Hum. Genet*., 84, 19-21.

**6.** Caiulo A, **Bardoni B**, Camerino G, Guioli S, Minelli A, Piantanida M, Crosato F, Dalla Fior T, Maraschio P. (1989) - Cytogenetic and molecular analysis of an unbalanced translocation (X;7)(q28;p15) in a dysmorphic girl. *Hum. Genet*., 84, 51-54.

**7.** Ballabio A, **Bardoni B**, Carrozzo R, Andria G., Persico G, Bick D, Campbell L, Ropers HH, Ferguson-Smith MA, Gimelli G, Fraccaro M, Maraschio P, Zuffardi O, Guioli S, Camerino G. (1989) - Contigous gene syndromes due to deletions in the distal short arm of the human X chromosome. *Proc. Nat. Acad. Sci. USA*, 86, 10001-10010.

**8.** Ballabio A, **Bardoni B**, Guioli S, Basler E, Camerino G. (1990) -Two families of low copy number repeats are interspersed on Xp22.3: implication for the high frequency of deletions in this region. *Genomics*, 8, 263-270.

**9.** Caiulo A, Nicolis S, Bianchi P, Zuffardi O, **Bardoni B**, Maraschio P, Ottolenghi S, Camerino G, Giglioni B. (1991) - Mapping the gene encoding the human erythroid transcriptional factor NFE1-GF1 to Xp11.23. *Hum. Genet*., 86, 388-390.

**10.** Mostacciuolo ML, Mueller E, Fardin P, Micaglio GF, **Bardoni B**, Guioli S, Camerino G, Danieli GA. (1991)- X-linked Charcot-Marie-Tooth disease. A linkage study in a large family by using 12 probes of the pericentromeric region. *Hum. Genet*., 87, 23-27.

**11**. **Bardoni B**, Zuffardi O, Guioli S, Ballabio A, Simi P, Cavalli P, Grimoldi MG, Fraccaro M, Camerino G. (1991) - A deletion map of the human Yq11 region: implications for the evolution of the Y chromosome and tentative mapping of a locus involved in spermatogenesis. *Genomics*, 11, 443-451.

**12**. Franco B, Guioli S, Pragliola A, Incerti B, **Bardoni B**, Tonlorenzi R, Carrozzo R, Maestrini E, Pieretti M, Taillon-Miller P, Brown C, Willard HF, Lawrence C, Persico G, Camerino G, Ballabio A. (1991) - A gene deleted in Kallmann's syndrome shares homology with neural cell adhesion and axonal pathfinding molecules. *Nature*, 353, 529-533.

**13**. Guioli S, Incerti B, Zanaria E, **Bardoni B**, Franco B, Taylor K, Ballabio A, Camerino G (1992) - Kallmann syndrome due to a translocation producing an X/Y fusion gene. *Nature Genet.*, 1, 337-340.

**14.** Incerti B, Guioli S, Pragliola A, Zanaria E, Borsani G, Tonlorenzi R, **Bardoni B**, Franco B, Ballabio A, Camerino G. (1992) - The Kallmann syndrome gene on the X and Y chromosomes: implications for the evolutionary divergence of human sex chromosomes. *Nature Genet*., 2, 311-314.

**15. Bardoni B**, Floridia G, Guioli S, Peverali G, Anichini C, Cisternino M, Casalone R, Danesino C, Fraccaro M, Zuffardi O, Camerino G. (1993) - Functional disomy of Xp22-pter in three males carrying a portion of Xp22 translocated to Yq11. *Hum. Genet*., 91, 333-338.

**16. Bardoni B**, Zanaria E, Guioli S, Floridia G, Worley K.C, Tonini G, Ferrante E, Chiumello G, McCabe E.R.B, Fraccaro M, Zuffardi O, Camerino G (1994) - A dosage sensitive locus at chromosome Xp21 is involved in male to female sex reversal. *Nature Genet*., 7, 497- 501

**17.** Zanaria E, Muscatelli F, **Bardoni B**, Strom T, Guioli S, Guo W, Lalli E, Moser C, Walker A.P, McCabe E.R.B, Meitinger T, Monaco A.P, Sassone-Corsi P, Camerino G. (1994) - An unusual member of the nuclear hormone receptor superfamily is resposible for X-linked adrenal hypoplasia congenita. *Nature*, 372, 635-641.

**18**. Muscatelli F, Strom T.M, Walker A. P, Zanaria E, Recan D, Meindl A, **Bardoni B**, Guioli S, Zehetner G, Rabl W, Schwarz H.P, Kaplan J, Camerino G, Meitinger T, Monaco A.P. (1994) - Mutations in the DAX-1 gene give rise to both X-linked adrenal hypoplasia congenita and hipogonadotropic hypogonadism. *Nature,* 372, 672-676.

**19.** Schmitt-Ney M, Thiele H, Kaltwasser P, **Bardoni B**, Cisternino M, Scherer G. (1995) - Two novel SRY missense mutations reducing DNA binding identified in XY femals and their mosaic fathers. *Am.J.Hum.Genet.,* 56, 862-869.

**20.** Genuardi M, **Bardoni B**, Floridia G, Chiurazzi P, Scarano G, Zollino M, Garcea M, Martini-Neri M.E, Neri G. (1995) - A dicentric chromosome Yp in a pseudohermaphrodite subjet with leydig cell agenesis. *Clin. Genet*., 47, 38-41.

**21.** Dabovic B, Zanaria E, **Bardoni B**, Lisa A, Bordignon C, Matessi C, Traversari C, Camerino G. (1995) - A family of rapidly evolving genes from the sex reversal critical region in Xp 21. *Mammalian Genome*, 6, 571-580.

**22.** Carrozzo R, Arrigo G, Rossi E, **Bardoni B**, Cammarata M, Gandullia P, Gatti R, Zuffardi O. (1997) - Multiple congenital anomalies, brain hipomyelination, and ocular albinism in a female with dup (X) ( pter-->q24: :q21.32-->qter ) and random X inactivation. *Am. J. Med. Genet.,* 72, 329-334.

**23.** Lalli E, **Bardoni B**, Zazopoulos E, Wurtz JM, Strom T, Moras D, Sassone-Corsi P. (1997) - A transcriptional silencing domain in DAX-1 whose mutation causes Adrenal Hypoplasia Congenita. *Mol. Endocrinol*., 11, 1950-1960.

**24. Bardoni B,** Sittler A, Shen Y, Mandel J.L. (1997) -Analysis of domains affecting intracellular localization of FMRP protein. *Neurobiology of Disease*, 4, 329-336.

**25.** Khandjian EW, **Bardoni B**, Corbin F, Sittler A, Giroux S, Heitz D, Tremblay S, Pinset C, Montarras D, Rousseau F, Mandel JL (1998) - Novel isoforms of the fragile X related protein FXR1P are expressed during myogenesis. *Hum. Mol. Genet*., 7, 2121-2128

**26. Bardoni B***, Schenck A, Mandel JL (1999) - A novel RNA binding nuclear protein that interacts with the fragile X mental retardation (FMR1) protein. *Hum. Mol. Genet*., 8, 2557-2566.

**27**. Parigi GB, **Bardoni B**, Avoltini V, Caputo MA, Bragheri R (1999) - Is bilateral congenital anorchia genetically determined? *Eur. J. Pediatr. Surg.,* 19, 312-315

**28.** Giglio S, Pirola B, Arrigo G, Dagrada P, **Bardoni B,** Bernardi F, Russo G, Argentiero L, Forabosco A, Carrozzo R, Zuffardi O. (2000) Opposite deletions/duplications of the X chromosome: two novel reciprocal rearragements. *Eur. J. Hum. Genet.,* 8, 63-70.

**29.** **Bardoni B**, Giglio S, Schenck A, Rocchi M, Mandel J.L. (2000). Assigment of the *NUFIP* (Nuclear FMRP Interacting Protein) gene to chromosome 13q13-14 and assigment of a pseudogene to chromosome 6q12-q13. *Cytogen. Cell Genet*., 89, 11-13.

**30.** Salat U, **Bardoni B**, Wörle D, Steinbach P. (2000) - Increase of FMRP expression, raised levels of *FMR1* mRNA, and clonal selection in proliferating cells with unmethylated fragile X repeat expansions: a clue to the sex bias in the transmission of full mutations? *J. Med. Genet*., 37, 842-850.

**31.** Schenck A, **Bardoni B,** Moro A, Bagni C, Mandel JL. (2001) - A highly conserved protein family interacting with the fragile X mental retardation protein (FMRP) and displaying selective interactions with FMRP-related proteins FXR1P and FXR2P. *Proc. Natl. Acad. Sci. U S A*, 98, 8844-8849.

**32.** Schaeffer C, **Bardoni B**, Mandel J.L., Ehresmann B, Ehresmann C, Moine H. (2001) - The fragile X mental retardation protein binds specifically to its mRNA via a purine quartet motif. *EMBO J*., 20, 4803-4813.

**33.** Larizza D, Maraschio P, **Bardoni B,** Calcaterra V, Manfredi P, Gemma A. (2002) - Two sisters with 45,X karyotype: influence of genomic imprinting of phenotype and cognitive profile. *Eur. J. Pediatr*., 161, 224-225.

**34.** Schenck A, Van de Bor V, **Bardoni B,** Giangrande A (2002) - Novel features of dFMR1, the *Drosophila* orthologue of the Fragile X Mental Retardation Protein (FMRP). *Neurobiology of Disease*, 11, 53-63.

**35.** Schenck A, **Bardoni B***, Langmann C, Harden N, Mandel J-L, Giangrande A. (2003) - CYFIP/Sra-1 controls neuronal connectivity in *Drosophila* and links the Rac1 GTPase pathway to the Fragile X protein. *Neuron*, 38, 887-898.

**36.** **Bardoni B***, Castets M, Huot M-T, Schenck A, Adinolfi A, Corbin F, Pastore A, Khandjian EW, Mandel J-L. (2003) - 82-FIP, a novel FMRP (Fragile X Mental retardation Protein) interacting protein, shows a cell-cycle dependent intracellular localization. *Hum. Mol. Genet*., 12, 1689-1698. COVER

**37.** **Bardoni B*,** Willemsen R, Weiler IJ, Schenck A, Severijnen L-A, Lalli E, Mandel J-L. (2003) - NUFIP1 (Nuclear FMRP Interacting Protein 1) is a nucleo-cytoplasmic shuttling protein associated with active synaptoneurosomes. *Exp. Cell Res*., 289, 95-107.

**38.** Adinolfi S, Ramos A,Martin S.R., Dal Piaz F., Pucci P, **Bardoni B,** Mandel JL, Pastore A. (2003) -The N-terminus of thefragile X mental retardation protein contains a novel domain involved in dimerisation and RNA-binding. *Biochemistry*, 42, 10437-10444.

**39**. Mayne M, Moffatt T, Kong H, McLaren PJ, Fowke KR, Becker KG, Namaka M, Schenck A, **Bardoni B**, Bernstein CN, Melanson M. (2004) - CYFIP2 is highly abundant in CD4+ cells from multiple sclerosis patients and is involved in T cell adhesion. *Eur J Immunol*., 34, 1217-1227.

**40.** Mientjes E.J., Willemsen R., Kirkpatrick L.L., Nieuwenhuizen I.M., Hoogeveen-Westerveld M., Verweij M., Reis S., **Bardoni B,** Hoogeveen A.T., Oostra B.A., Nelson D.L. (2004) - Fxr1

knockout mice show a striated muscle phenotype: implications for Fxr1p function *in vivo***.** *Hum. Mol. Genet*., 13, 1291-1302.

**41.** Khandjian EW, Huot M-E, Tremblay S, Davidovic L, Mazroui R, **Bardoni B**. (2004) - Biochemical evidence for the association of fragile X mental retardation protein with brain polyribosomal ribonucleoparticles. *Proc. Nat. Acad. Sci. U.S.A.,* 101, 13357-13362.

**42**. Schenck A, Qurashi AA, **Bardoni B**, Schejter ED, Mandel J-L, Giangrande A. (2004) - WAVE/SCAR, a multifunctional complex required in fly neuronal connectivity. *Dev. Biol.*, 274, 260-270.

**43.** Castets M, Schaeffer C, Bechara E, Schenck A, Khandjian EW, Luche S, Moine H, Rabilloud T, Mandel J-L and **Bardoni B**.*****(2005) - FMRP interferes with the Rac1 pathway and controls actin cytoskeleton dynamics in murine fibroblasts. *Hum. Mol. Genet.*, 14, 835-844.

**44.** Ramos A**,** Hollingworth D, Adinolfi S, Castets M, Kelly G, Frenkiel TA, **Bardoni B,** Pastore A. (2006) - The N-terminal domain of the Fragile X Mental retardation Protein forms a novel platform for protein-protein interaction. *Structure,* 14, 21-31

**45.** Davidovic L., Bechara E, Gravel M, Jaglin XH, Tremblay S, Sik A, **Bardoni B**, Khandjian EW (2006). - The nuclear MicroSpherule protein 58 is a novel RNA-binding protein that interacts with fragile X mental retardation protein in polyribosomal mRNPs from neurons. *Hum. Mol. Genet.,* 15, 1525-1538

**46.** Kim Y, Sung JY, Ceglia I, Lee K-W, Ah J-H, Halford JM, Kim AM, Kwak SP, Park JB, Ryu SH, Schenck A, **Bardoni B**, Scott JD, Nairn AC, Greengard, P. (2006) - Phosphorylation of WAVE1 regulates actin polymerization and dendritic spine morphology. *Nature,* 442, 814-817.

**47**. Bechara E., Davidovic L., Melko M., Bensaid M., Grosgeorge J., Tremblay S., Khandjian E.W., Lalli E., **Bardoni B.*** (2007) – Fragile X Related Protein 1 isoforms differentially modulate the affinity of Fragile X Mental Retardation Protein for G-quartet mRNA structure*. Nuc. Acid Res.*, 35, 299-306.

**48.** Davidovic L., Jaglin X.H., Lespagnol-Bestel A.M., Tremblay S., Simonneau M., **Bardoni B.**, Khandjian E.W. (2007) - The Fragile X Mental Retardation Protein is a molecular adaptor between the neurospecific KIF3C kinesin and dendritic RNA granules. *Hum. Mol. Genet*., 16, 3047-3058.COVER

**49.** Boulon S, Marmier-Gourrier N, Pradet-Balade B, Wurth L, Verheggen C, Jády B, Rothé J, Pescia C, Robert M-C, Kiss T, **Bardoni B**, Krol A, Branlant C, Allmang C, Bertrand E, and Charpentier B. (2008) - The HSP90 chaperone controls the biogenesis of L7Ae RNPs through aconserved machinery**.** *J.Cell Biol.,* 180*,* 579-595

**50.** Davidovic L, Sacconi S, Bechara E, Delplace S, Allegra M, Desnuelle C and **Bardoni B*** (2008) - Alteration of expression of muscle-specific isoforms of the Fragile X Related Protein 1 (FXR1P) in facio-scapulohumeral muscular distrophy patients. *J. Med. Genet.,* 45, 679-85

**51**. Bensaid M, Melko M, Bechara E, Davidovic L, Beretta A, Catania MV, Gecz J, Lalli E, **Bardoni B*** (2009) - FRAXE-associated mental retardation protein (FMR2) is an RNA-binding protein with high affinity for G-quartet RNA forming structure. *Nucleic Acids Res.,* 37,1269-79

**52**. Bechara E, Didiot, MC, Melko M, Davidovic L, Bensaid M, Martin P, Castets M, Pognonec P, Khandjian E, Moine H, **Bardoni B*** (2009) - A Novel Function of Fragile X Mental Retardation Protein in translational activation. *PLoS Biology*, Jan 20;7(1):e16

**53.** Khalfallah O, Rouleau M, Barbry P, **Bardoni B**, Lalli E. (2009) - Dax-1 knockdown in mouse embryonic stem cells induces loss of pluripotency and multilineage differentiation. *Stem Cells*, 27, 1529-37.

**54.** Melko M, Douguet D., Bensaid M, Zongaro S, Verheggen C, Gecz J, **Bardoni B*** (2011) - Functional characterization of the AFF (AF4/FMR2) family of RNA binding proteins: insights into the molecular pathology of FRAXE intellectual disability. *Hum. Mol. Genet*. 20,1873-85. COVER

**55**. Davidovic L, Navratil V, Bonaccorso C.M., post-transcriptional RNA-metabolism and RNA-binding proteins: 3 actors for a complex scenario. Catania M.V., **Bardoni B**, Dumas M. (2011) – A metabolomic and system biology perspective on the brain of the Fragile X syndrome mouse model. *Genome Res*. 21, 2190- 2202.

**56.** Zongaro S, Hukema R, D'Antoni S, Davidovic L, Barbry P, Catania M.V., Willemsen R, Mari B, **Bardoni B*** (2013) - The 3' UTR of FMR1 mRNA is a target of miR-101, miR-221 and miR-129-5p:implications for the molecular pathology of FXTAS at the synapse. *Hum Mol Genet*., 22,1971-1982.

**57**. Davidovic L, Durand N, Khalfallah O, Tabet R, Barbry P, Mari B, Sacconi s, Moine H, **Bardoni B*** (2013) - A novel role for the RNA-binding protein FXR1P in myoblasts cell-cycle progression by modulating *p21/Cdkn1a/Cip1/Waf1* mRNA stability. *PLoS Genet*., 9(3):e1003367.

**58**. Melko M, Nguyen LS, Shaw M, Jolly L, **Bardoni B***, Gecz J (2013) - Loss of FMR2 further emphasizes the link between deregulation of immediate early response genes FOS and JUN and intellectual disability. *Hum Mol Genet*., 22, 2984-91

**59**. Ugolotti R, Mesejo P, Zongaro S, **Bardoni B**, Berto G, Bianchi F, Molineris I, Giacobini M, Cagnoni S, Di Cunto F. (2013) - Visual search of neuropil-enriched RNAs from brain in situ hybridization data through the image analysis pipeline hippo-ATESC. *PLoS One*, 8(9):e74481.

**60.** Matik K, Eninger T, **Bardoni B**, Davidovic L, Macek B. (2014) - Quantitative phosphoproteomics of murine Fmr1-Ko cell lines provides new insights into FMRP-dependent signal transduction mechanisms. *J Proteome Res*, [Epub ahead of print]

**61.** Schertzer M, Jouravleva K, Perderiset M, Dingli F, Loew D, Le Guen T, **Bardoni B**, de Villartay JP, Revy P, Londoño-Vallejo A. (2015) - Human regulator of télomère élongation helicase 1 (RTEL1) is required for the nuclear and cytoplasmic trafic of pre-U2 RNA. *Nuc. Acids Res.,* 43, 1834-47.

**62.** Bonaccorso CM, Spatuzza M, Di Marco B, Gloria A, Barrancotto G, Cupo A, Musumeci SA, D'Antoni S, **Bardoni B**, Catania MV. (2015) Fragile X Mental Retardation Protein (FMRP)

interacting proteins exhibit different expression patterns during development. *Int J Dev Neurosci,* 42, 15-23.

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