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CURRICULUM VITAE

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RESEARCH GRANTS/AWARDS/CONTRACTS/PROJECTS:

Prior

Peer Review

Title:	MAB21L Family in Human Ocular Disease and Development
Source:	NIH/NEI
PI:	Semina, Elena V, PhD
Dates:	03/01/2017 - 02/28/2021
Direct Funds:	$1,394,474

Title:	WDR37: a novel factor in human congenital multisystem disease
Source:	NIH/NICHHD
PI:	Semina, Elena V, PhD
Dates:	07/20/2019 - 06/30/2021
Direct Funds:	$418,000

Title:	Retinopathy of Prematurity: Genetic Mechanisms
Source:	Children’s Wisconsin/Children’s Research Institute
PI:	Semina EV, PhD
Dates:	01/01/2020 - 12/31/2020
Direct Funds:	$75,000

Title:	Genetic Studies of Human Development
Source:	Children’s Wisconsin/Children’s Research Institute
PI:	Semina, Elena V, PhD
Dates:	01/01/2021 - 12/31/2021
Direct Funds:	$150,000

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Refereed Journal Publications/Original Papers

1. Semina EV, Reiter R, Leysens NJ, Alward WL, Small KW, Datson NA, Siegel-Bartelt J, Bierke-Nelson D, Bitoun P, Zabel BU, Carey JC, Murray JC. Cloning and characterization of a novel bicoid-related homeobox transcription factor gene, RIEG, involved in Rieger syndrome. Nat Genet. 1996 Dec;14(4):392-9.
2. Semina EV, Datson NA, Leysens NJ, Zabel BU, Carey JC, Bell GI, Bitoun P, Lindgren C, Stevenson T, Frants RR, van Ommen G, Murray JC. Exclusion of epidermal growth factor and high-resolution physical mapping across the Rieger syndrome locus. Am J Hum Genet. 1996 Dec;59(6):1288-96. PMCID: PMC1914874
3. Semina EV, Reiter RS, Murray JC. Isolation of a new homeobox gene belonging to the Pitx/Rieg family: expression during lens development and mapping to the aphakia region on mouse chromosome 19. Hum Mol Genet. 1997 Nov;6(12):2109-16.
4. Alward WL, Semina EV, Kalenak JW, Héon E, Sheth BP, Stone EM, Murray JC. Autosomal dominant iris hypoplasia is caused by a mutation in the Rieger syndrome (RIEG/PITX2) gene. Am J Ophthalmol. 1998 Jan;125(1):98-100.
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negative homeodomain mutation in Rieger syndrome. J Biol Chem. 2001 Jun 22;276(25):23034-41.
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