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Maryam Rafati, Seyedeh Zahra Tara, Fatemeh Hoseininasab, Saeed Reza Ghaffari

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Identification of a Novel *ALMS1* Pathogenic Variant in a Family with Cone-Rod Dystrophy using Whole Exome Sequencing, Followed by Prenatal Diagnosis

Authors:

Maryam Rafati^{1,2}, rafati@hopegen.org Seyedeh Zahra Tara², sztara88@yahoo.com Fatemeh Hoseininasab¹, hoseininasab.f@gmail.com Saeed Reza Ghaffari^{2,3}*, saeed@ghaffari.org

Affiliations:

¹Reproductive Biotechnology Research Center, Avicenna Research Institute, ACECR, Tehran, Iran; ²Fetal Health Research Center, Hope Generation Foundation, Tehran, Iran;

³Gene Clinic, Tehran, Iran

*Correspondence:

Saeed Reza Ghaffari

Postal address: Gene Building, No. 344, Keshavarz Blvd, Tehran, Iran

Postal code: 1418894161 Tel: +98 21 66935655 Fax: +98 21 66933336 Email: saeed@ghaffari.org

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