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PUBLICATIONS:

- Watts GDJ, Wymer J, **Kovach MJ**, Mehta SG, Mumm S, Darvish D, Pestronk A, Whyte MP, Kimonis VE (2004) Inclusion body myopathy associated with Paget disease of bone and frontotemporal dementia is caused by mutant valosin-containing protein. *Nat. Genet.* 23:377-381.
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- Kimonis VE, **Kovach MJ**, Waggoner B, Leal S, Salam A, Rimer L, Davis K, Khardori R, Gelber D (2000). Clinical and molecular studies in a unique family with autosomal

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Chen AS, **Kovach MJ**, Herman K, Frank W, Forrester S, Avakian A, Lin J-P, Kimonis VE (2000). Clinical heterogeneity in autosomal dominant optic atrophy in two 3q28-pter linked central Illinois families. *Genet. Med.* 2:283-289.

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ABSTRACTS:

Chen AS, **Kovach MJ**, Kimonis VE (1999) Neurological and dysmorphic features in a family with brachial plexus neuropathy. **49th Annual Meeting of the American Society of Human Genetics.**

Kovach MJ and Landy A (1997) Characterization of protein:DNA interactions of λ -integrase utilizing photocrosslinking and 4-thio-T modified DNA oligomers. **Gordon Research Conference on Plasmid and Chromosome dynamics.**

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