



Growth curve analysis (GCA) was done using multilevel regression modeling using R Studio as described earlier [58, 59]. In **A**) the intercept model (m.0) and in **B**) the linear model (m.1) is presented. The intercept model suggests constant differences in proliferation randomly assigned to the different cell lines and the linear model suggests effects of the different cell lines on proliferation. Statistical comparison (Pearson's chi-squared test) of these two models revealed that the linear model (m.1) is our regression model of choice (p < 0.001).





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Figure S3 - Full-length western blot of Figure 4B.

Unprocessed full-length western blot of Figure 4B. In **A**) the piece of nitrocellulose membrane is shown along with the molecular weight marker and in **B**) the original image acquired with Azure Imager is presented.



Figure S4 - TaqMan analysis of *PGRMC1* in human skin fibroblasts.

Total RNA was isolated from human skin fibroblasts of healthy wild-type donors and triple A patients. The different mutations in the human *AAAS* gene are denoted on the x-axis of the diagram: IVS14, S263P and W295X. WT, wild-type.

Table S1 - Real-time qPCR primer oligonucleotides.

GENE	NAME	SEQUENCE
ACTB (V1: NM_001101.3)	ACTB-F ACTB-R ACTB-Probe	GCACCCAGCACAATGAAGATC CGCAACTAAGTCATAGTCCGC TGCTCCTCCTGAGCGCAAGTACTCC
<i>PGRMC1</i> (V1: NM_006667.4)	PGRMC1-F PGRMC1-R PGRMC1-Probe	CTGCATGATTTCTGTTTTATCTACCTCTA TGTTACTGGACAGCGCTTAATCC AGCAAATCTGCAGTGTTCCAAAGACTTTGG
<i>PGRMC2</i> (V1: NM 006320.4)	PGRMC2-F PGRMC2-R PGRMC2-Probe	GCGGTCAATGGGAAAGTCTTC CTACCAGCAAATATTCCATA AGTTCTACGGCCCCGCGGGTC