8 VERÖFFENTLICHUNGEN

8.1 Wissenschaftliche Veröffentlichungen

Zeitz C., Scherthan H., Freier S., Feil S., Suckow V., Schweiger S. und Berger W. (2003) *NYX* (nyctalopin on chromosome X), the gene mutated in congenital stationary night blindness, encodes a cell surface protein. ***Invest. Ophthalmol. Vis. Sci.*** **44**(10): 4184-4191.

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Kirschner R., Erturk D., Zeitz C., Sahin S., Ramser J., Cremers F., Ropers H.-H. und Berger W. (2001) DNA sequence comparison of human and mouse retinitis pigmentosa GTPase regulator (*RPGR*) identifies tissue specific exons and putative regulatory elements***. Hum. Genet.*** **109**: 271-278.

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8.2 Kongressbeiträge

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Kirschner R., Erturk D., Zeitz C., Sahin S., Ramser J., Rosenberg T., Bergen A., Cremers F., Ropers H.-H. und Berger W. (2001) Comparative genomic sequencing of the human and mouse RPGR gene identifies tissue-specific coding sequences that are mutated in patients with X-linked retinitis pigemntosa. ***Europ. J. Hum. Genet.* 9 (Supplement)**; 430; 10th *International Congress of Human Genetics*, 15.-19. Mai. 2001, Wien, Österreich, Poster Präsentation.

Meindl A., Pusch C., Brandau O., Zeitz C., Wen G. P., Demirci Y., Achatz H., Wissinger B., Berger W., Hardcastle A., Platzer M., Gorin M und Ramser J. (2000) Towards a complete transcription map of the medically relevant human Xp11.4 region and isolation of the *CSNB1* gene. *German Human Genome Meeting*, 30.11.-01.12.2000, German Cancer Research Center (dkfz), Heidelberg, Deutschland, Poster Präsentation.

Kirschner R., Zeitz C., Knoebeloch K.-P., Rüther K., Rohde E., Hoffmann K., Horak I., Ropeman R., Rodenberg T., Cremers F., Ropers H.-H. und Berger W. (1999) X-linked retinitis pigmentosa 3 (RP3): identification of a retina-specific RPGR transcript and gene targeting of the mouse orthologue. ***Am. J. Hum. Genet. 65 (Supplement)*:** A305; 49nd *Annual Meeting of the American Society of Human Genetics*, 19.-23. Oktober 1999, San Francisco, California, USA, Poster Präsentation.