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Robert F. Gagel

Charles E. Jackson

Bruce A. J. Ponder

Friedhelm Raue

Nancy E. Simpson

See next page for additional authors

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Authors

Robert F. Gagel, Charles E. Jackson, Bruce A. J. Ponder, Friedhelm Raue, Nancy E. Simpson, and Reinhard Ziegler

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Robert F. Gagel,* Charles E. Jackson,† Bruce A.J. Ponder,‡ Friedhelm Raue,§ Nancy E. Simpson,|| and Reinhard Ziegler§

For this issue of the *Journal* and as a recommendation to other authors and journals for uniformity of preferred nomenclature, the following suggestions were made:

- MEN 2 The multiple endocrine neoplasia type 2 syndromes (*not* MEA, MEN-II, MEN-2, or M.E.N. 2).
- MEN 2A The MEN 2 syndrome without the mucosal neuroma phenotype (*not* MEN 2a).
- MEN 2B The MEN 2 syndrome with the mucosal neuroma phenotype (*not* MEN 2b). This has also been referred to as MEN 3.
- MEN2A The designation of the mutated gene causing MEN 2A. As recommended by the 1989 Human Gene Mapping conference (1), gene designations are no longer italicized or underlined.
- MEN2B The designation of the mutated gene causing MEN 2B.

MTC Medullary thyroid cancer or carcinoma, to be used for consistency rather than MCT. The two types of MTC are hereditary and nonhereditary.

Reference

1. McAlpine PJ, Shows TB, Boucheix C, et al. Report of the nomenclature committee and the 1989 catalog of mapped genes. *Human gene mapping 10* (1989): Tenth International Workshop on Human Gene Mapping. *Cytogenet Cell Genet* 1989;51:13-66.

*Laboratory of Molecular and Cellular Endocrinology, Baylor College of Medicine and Veterans Administration Medical Center, Houston, TX.

†Department of Internal Medicine, Division of Clinical & Molecular Genetics, Henry Ford Hospital.

‡Department of Pathology, CRC Human Cancer Genetics Research Group, Cambridge, England.

§Department of Internal Medicine I, Endocrinology & Metabolism, University of Heidelberg, West Germany.

||Departments of Biology and Paediatrics, Queens University, Kingston, Ontario, Canada.

Address correspondence to Dr. Gagel, Laboratory of Molecular and Cellular Endocrinology (111E), Veterans Affairs Medical Center, 2002 Holcombe Blvd, Houston, TX 77030.