

Henry Ford Hospital Medical Journal

Volume 37 | Number 3

Article 3

9-1989

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Recommended Citation

Gagel, Robert F.; Jackson, Charles E.; Ponder, Bruce A. J.; Raue, Friedhelm; Simpson, Nancy E.; and Ziegler, Reinhard (1989) "Multiple Endocrine Neoplasia Type 2 Syndromes: Nomenclature Recommendations from the Workshop Organizing Committee," *Henry Ford Hospital Medical Journal* : Vol. 37 : No. 3 , 99. Available at: <https://scholarlycommons.henryford.com/hfhmedjournal/vol37/iss3/3>

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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.

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