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The image depicts (left) Paul Wermer, who in 1954 was the first to suggest that what became the MEN1 phenotype was a genetic disorder, and (right) a hyperplastic pancreatic islet of a MENX mutant (p27 deficient) rat: α-insulin (red), α-glucagon (green).

Credit: (left) courtesy M Herrera; (right) Pellegata N, Mohr H & Wiedemann T (Helmholtz Zentrum München, Neuherberg, Germany)