Clinical Neuroendocrinology and Neuroendocrine Tumors



Neuroendocrinology 2008;88:235–242 DOI: 10.1159/000149356 Received: January 18, 2008 Accepted after revision: May 9, 2008 Published online: July 29, 2008

Management of Neuroendocrine Tumors: A Meeting of Experts from Latin America

F. Costa^a E. Domenichini^c G. Garavito^e R. Medrano^g G. Mendez^d J. O'Connor^c W. Rojas^f S. Torres^h R.N. Younes^{a, b} G. Delle Faveⁱ K. Öberg^j

^aHospital Sírio Libanês, ^bUniversity of São Paulo, São Paulo, Brazil; ^cInstituto Alexander Fleming, ^dHospital de Gastroenterología Dr. Carlos Bonorino Udaondo, Buenos Aires, Argentina; ^eInstituto Nacional de Cancerología, ^fHospital San Jose, Bogotá, Colombia; ^gCentro Médico Naval, Secretaria de Marina Armada de México, ^hHospital de Oncologia Centro Medico Nacional Siglo XXI I.M.S.S., Mexico, Mexico; ⁱUniversita Degli Studi di Roma 'La Sapienza', Rome, Italy; ^jUppsala University, Uppsala, Sweden

Key Words

Interferon- α · Latin America · Neuroendocrine tumors · Octreotide · Radionuclide imaging · Somatostatin receptors

Abstract

A panel of experts from Latin America convened in Brazil, in May of 2007, for consensus recommendations regarding the management of neuroendocrine tumors (NETs) of the gastrointestinal tract and pancreas. The recently introduced World Health Organization classification of NETs represents a step forward, but the former classification of carcinoids into foregut, midgut and hindgut is still likely to be useful in the near future. Macroscopic description of the tumor should be followed by light microscopic examination and immunohistochemical staining, whereas other techniques might not be widely available in Latin America. Surgery remains the mainstay of treatment for patients with potentially curable tumors, and adequate selection is paramount in order to optimize treatment results. Regarding systemic therapy, patients with well-differentiated tumors or islet-cell carcinomas may be categorized as having indolent disease, while patients with poorly differentiated, anaplastic, and smallcell carcinomas, or with atypical carcinoids, may be approached initially as having aggressive disease. Somatostatin analogues play a cytostatic role in indolent tumors, and chemotherapy may play a role against other, more aggressive NETs. Obviously, there is an urgent need for novel therapies that are effective against NETs.

Copyright $\ @$ 2008 S. Karger AG, Basel

Introduction

A panel of experts from four Latin-American and two European countries convened in São Paulo, Brazil, in May of 2007, for a consensus meeting on the management of neuroendocrine tumors (NETs) of the gastrointestinal tract and pancreas. This paper presents the opinions of the panelists on subjects related to the classification, diagnosis, treatment and follow-up of patients with NETs in Latin America, a world region that currently comprises approximately 550 million inhabitants. Given the fact that prospective data from controlled trials for many of the questions discussed in the meeting are currently scarce [1], evidence-based medical data have been used by panel members whenever possible, but many of the rec-