

BRONHO-PULMONARY CANCER: MORFO-PATHOLOGICAL FORMES

LIVIA MIRELA POPA¹

PhD candidate, "Lucian Blaga" University of Sibiu

Keywords:

histopathological classification, benign tumors, malignant tumors

Abstract: The primitive bronchopulmonary cancer includes the cases of cancer that develops primitively from the bronchic glandular epithelium, next invading the pulmonary parenchyma. It is the most important and frequent form of pulmonary tumor representing more than 90% of the primitive malignant and benign tumors. The histological classification of the bronchopulmonary cancer is complex and extremely important from the point of view of the neoplasia evolution, the choice of the possibilities of treatment and of the prognosis.

Cuvinte cheie:

clasificarea histopatologică, tumori benigne, tumori maligne

Rezumat: Cancerul bronhopulmonar primitiv include cazurile de cancer care se dezvoltă, în mod primitiv, de regulă din epiteliul glandular bronșic, invadând ulterior și parenchimul pulmonar. Este cea mai importantă și frecventă formă de tumoră pulmonară, reprezentând peste 90% din tumorile primitive maligne și benigne. Clasificarea histologică a cancerului bronho-pulmonar este complexă și este extrem de importantă din punctul de vedere al evoluției neoplaziei, a alegerii modalităților terapeutice și a prognosticului.

SCIENTIFIC ARTICLE OF BIBLIOGRAPHIC SYNTHESIS

The primitive bronchopulmonary cancer includes the cases of cancer that develops primitively from the bronchic glandular epithelium next invading the pulmonary parenchyma. It is the most important and frequent form of pulmonary tumor representing more than 90% from the malignant and benign tumors. (1)

Some authors (especially the bronchologists) consider that the correct name would be bronchogenoic or bronchogenetic cancer (Lemoine). Most use the name of pulmonary cancer or bronchopulmonary cancer that doesn't exclude the idea of the predominant bronchogenetic origin reflecting even better the coaffection bronchic and parenchymal and is not incorrectly from the anatomical point of view because the bronchi are constitutive part of the lung. Like wise correct is the name of carcinoma. (1)

Rarely met and mistaken for the consumption in the antiquity and with other consumptive lung diseases, was the lung cancer observed for the first time without being individualized in the antiquity, in the XVI-th century, by Paracelsus and by Agricola as *male metallorum*, at the miners from Schneeberg (1531) and from St. Joachimstal (1556). The diagnosis was established retrospectively later by Hesse and Härting (1879). Meantime Bayle (1810) described it under the name of cancerous consumption, considering it the sixth form of consumption. The worth of being individualized as a nosologic entity under the name of "the lung's encephaloid" is attributed few years later to Laennec. Stokes (1837) establishes the diagnosis procedure. Walsche (1843) gives it the first the name of lung cancer. Wolf (1895) points out the association with the tuberculosis. Waldayer mentions, the first, the epithelial origin of the cancerous tumor. (1)

Until the end of the XIX century are studied minutely, its clinical aspects (Jaccoud, Darolles, Marchiafan)

and morph- pathological ones (Virchow, Ménetrier). In the first decades of the XX century are specified the radiologic aspects of the disease in its manifest stadium, correlated with the morphologic ones (Letulle, Huguenin, Delarue). Later, by introducing new methods of bronchoscopic investigation, citodiagnosis, biopsic, funcțional respiratory ones becomes possible the diagnosis of the disease in a precocious phase, still operable (Adler, Lemoine).

Exeresis surgery developed and perfected on a large scale after 1946 (Overholt, Björk, Derra, Cărpinișan) and offers, for the first time, the possibility of a terapeutical solution of the cases, with the condition of realising an early sistematic diagnosis.

Morphopathological aspects of the bronchopulmonary cancer:

The morphopathological aspects of the bronchiopulmonary cancer are extremely varied raported to: the size of the tumor; the localization on a central or peripheral bronchia; degree of intratoracic and extratoracic extension and especially with; the histological type.

The statement of those elements is of a maximal importance for choosing the exploratory methods, establishing the optimal modalities of treatment and of the prognosis (3). From a morphopathological point of view two elements are essential: the localization of the tumor central or peripheral from the beginning; histological type.

The histological classification of the bronchopulmonary cancer is complex and extremely important from the point of view of the evolution of the neoplasia, of choosing the therapeutic possibilities and of the prognosis. Lung tumors have been classified by OMS, classification that was redacted in 1981 and 1999 (13, 15). Pulmonary tumors are composed from more different histological types with a varied degree of malignity, from complete benign to an extremely aggressively.

The next classification is redacted by OMS (From Travis

¹Corresponding Author: Livia Mirela Popa, 10 B-dul Corneliu Coposu street, ap.28, Sibiu, România; e-mail: liviamirelapopa@yahoo.com; tel +40-0723609030

Article received on 17.02.2010 and accepted for publication on 25.02.2010
ACTA MEDICA TRANSILVANICA September 2010; 2(3)210-211

CLINICAL ASPECTS

WD, Colby TV, Corrin B, et al. *WHO histological typing of lung and pleural tumors*, 3rd ed. Geneva: World Health Organization, 1999):

Epithelial tumors:

I. Benign

1. Papilloma: a. with squamous cells (exofitic, inverted); b. glandular, c. glandular and mixt squamous;
2. Adenoma: alveolar; papillar; of the salivary gland type (mucose, pleomorph); Mucinous Cyst adenoma.

II. *Pre -invasive lesions*: squamous dysplasia, carcinoma in situ, atypical hyperplasia adenomatous, diffuse idiopathic hyperplasia with neuro endocrine cells;

III. Malign:

1. Carcinoma with squamous: a. papillar; b. with clear cells; c. with small cells; d. Basaloid;
2. Carcinoma with small cells: a. carcinoma with mixt cells ;
3. Adenocarcinoma; a. Acinar; b. papillar; c. bronchioloalveolar: nonmucinous, mucinous, mixt: nonmucinous and mucinous or type of intermediate cells; d. solid with mucine; e.with mixt subtypes, variants: adenocarcinoma with well differentiated fetal cells, adenocarcinoma with mucinous cells(colloidal), cystadenocarcinom mucinous;
4. Carcinoma with large cells: a. carcinoma with neuro endocrine large cells; b. carcinoma combined with large neuro endocrine cells; c. basaloid carcinoma ; d. carcinoma limfo epithelial-like; e. carcinoma with clear cells; f. carcinoma with large cells with rhabdoid phenotype;
5. Adenosquamous Carcinoma;
6. Carcinoma with pleomorph elements, sarcomatoid or sarcomatous;
7. Carcinoma with giant cells or fusiform: a. pleomorph: with fusiform cells or large cells; b.carcinosarcoma; c.pulmonary blastoma ;
8. Carcinoid tumors: typical and atypical; I
9. Carcinomas of the salivary glands type: mucoepidermoid, adenoid cystic;
10. Unclassified

Tumors of soft tissue: Tumors localized fibrous, epitheloid hemangioendothelioma, pleuropulmonary blastoma, chondroma, pleural fibrous calcified pseudotumor, congenital peribronchial myofibroblastic tumor, diffuse pulmonary lymphangiomatosis, desmoplastic round cells tumor;

Mesothelial tumors: *Benign* – adenomatoid tumor, *Malignant* mesotheliom, Sarcomatoid mesotheliom (desmoplastic, biphasic);

Divers tumors:

1. Hamartoma,
2. Sclerosing hemangioma,
3. Tumor with clear cells,
4. Tumor with germinative cells (mature teratoma, immature teratoma, tumor with other cells),
5. Thymoma,
6. Melanoma malignant

Limfoproliferations: 1.Interstitial lymphoid pneumonia, 2.Nodular lymphoid hyperplasia, 3. Lymphoma with B cells type marginal area with a low degree of associated lymphoid tissue (MALT), 4.Lymphoid granulomatosis

Secondary tumors

Unclassified tumors

Tumoral type lesions: 1. Tumor, 2. meningothelioma multiple nodules, 3. Histiocytosis with Langerhans cells, 4.Inflammatory pseudotumors (mioinflamator), 5.Organized localized pneumonia, 6.Amyloid tumor (nodular amyloid), 7.Hyalinizing granuloma, 8.Lymphangioliomyomatosis, 9.Micronodular pneumocystic hyperplasia, 10.Endometriosis, 11. Inflammatory bronchial polyp

BIBLIOGRAPHY

1. Anastasatu C, Eskenasy A. Cancerul bronhopulmonar. În Colectia Enciclopedia Oncologică, Cluj-Napoca, 1986.
2. England DM, Hochholzer L. Truly benign „bronchial adenoma”; report of 10 cases of mucous gland adenoma with imunohistochemical and ultrastructural findings. Am J Surg Pathol.1995; 19: 887-889.
3. Gherasim L. Tumorile bronho-pulmonare. În *Medicină Internă*, Vol. I, Ed. Medicală, 1996; 341-375.
4. Hammond ME, Sause WT. Large cell neuroendocrine tumors of the lung: clinical significance and histopathologic definition. Cancer.1985; 56:1624-1629.
5. Kodama T, Shimosato Y, Kameya T,. Histology and ultrastructure of bronchogenic and bronchial gland adenocarcinomas (including adenoid cystic and mucoepidermoid carcinomas) in relation to histogenesis. In: Shimosato Y, Melamed MR, Nettesheim P, eds. Morphogenesis of lung cancer. Vol. I.Boca Raton, FL: CRC Press, 1982:65-69.
6. Miller RR, Nelems B, Evans KG, et al. Glandular neoplasia of the lung: a proposed analogy to colonic tumors. Cancer. 1988;61: 1009-1014.
7. Nakayama H, Noguchi M, Tsuchiya R, et al. Clonal growth of atypical adenomatous hyperplasia of the lung: cytofluorometric analysis of nuclear DNA content. Mod Pathol.1990; 3: 314-320.
8. Nomori H, Shimosato Y, Kodama T, et al. Subtypes of small cell carcinoma of the lung: morphometric, ultrastructural and imunohistochemical analyses. Hum Pathol.1986; 17:604-613.
9. Shimosato Y, Noguchi M. Pulmonary Neoplasms. In Sternberg's: Diagnostic Surgical Pathology; 4th ed., vol.I. ed. Lippincott, Williams&Wilkins. 2004; 1173-1217.
10. Shimosato Y, Noguchi M, Matsuno Y. Adenocarcinoma of the lung: its development and malignant progression. Lung Cancer. 1993;9: 99-108.
11. Shimosato Y, Miller RR. Biopsy interpretation of the lung. New York: Raven Press, 1994.
12. Travis Wd, Lubin J, Ries L, Devesa S. United States lung carcinoma incidence trends: declining for most histologic types among males, increasing among females. Cancer. 1996; 77; 2464-2470.
13. Travis WD, Colby TV, Corrin B, et al. WHO histological typing of lung and pleural tumors, 3rd ed. Geneva: World Health Organization, 1999.
14. Tsuchihashi T, Yamaguchi K, Miyake Y, et al. Parathyroid hormone-related protein in tumor tissues obtained from patients with humoral hypercalcemia of malignancy. J Natl Cancer Inst. 1990; 82: 40-44.
15. World Health Organization. Histological typing of lung tumors, 2nd ed. Geneva: World Health Organization, 1981.
16. Yokozaki M, Kodama T, Yokose T, et al. Differentiation of atypical adenomatous hyperplasia and adenocarcinoma of the lung by use of DNA ploidy and morphometric analysis. Mod Pathol.1996;9 : 1156-1164.