Case report

Atypical Progress of Primary Undiagnosed Hepatoblastoma

Romaniuk A¹, Piddubnyi A², Sikora V³

Abstract:

Introduction: Hepatoblastoma is a disease that has a variety of clinical manifestations. This fact complicates the diagnosis of a large number of cases of this disease. Case presentation: A 64-year-old female Ukrainian patient was admitted to the anesthesiology and reanimation department in critical condition with a diagnosis: dissecting abdominal aortic aneurysm. Despite the complex of treatment and diagnostic measures after 10 hours was established a biological death. At autopsy of the patient were detected changes that characteristic for hepatoblastoma. Histological examination of the liver revealed the next diagnosis: poorly differentiated hepatocellular liver cancer. Conclusions: The patient had non-typical progress of the hepatoblastoma, which led to the erroneous diagnosis and incorrect chosen of the treatment strategy.

Keywords: hepatoblastoma; atypical progress; hepatocellular cancer; erroneous diagnosis

Introduction

Hepatocellular liver cancer (HLC) is a malignant tumor, that develops from hepatocytes. The most important etiological factors of this disease include infectious agents (viral hepatitis B and C), chronic alcoholism (consumption of more than 50 grams per day, in alcohol equivalent), metabolic disorders (hereditary hemochromatosis, tyrosinemia, Wilson’s disease, deficiency of alpha-one-antitrypsin), the use of drugs and toxins (aflatoxin B, androgens, progestogens, thorium dioxide), cirrhosis, etc [2]. It was also investigated the relationship of liver cancer with the usage of xenobiotics, such as organochlorine pesticides, polychlorinated biphenyls and vinyl chlorides [6]. HLC ranks the fifth place among the structure of malignant tumor and the third place among the causes of death. [6]. According to the WHO, this pathology is often found among people of South Africa, Asia, North and South America, North Europe, where the incidence can range from 21.1 [2] to 30 [3] per 100 000 population. Analysis of the reproductive structure of morbidity indicates the major lesion of men, than women (7.4% and 3.2% according to all the malignancies) [2]. In the majority of patients HLC is asymptomatic or accompanied by pain, weight loss, bleeding, fever, malaise, ascites, sub- and icteric of the skin and mucous membranes, the presence of distant and regional metastases [3, 6]. For intravital diagnosis one uses ultrasound examination, magnetic resonance tomography, computed tomography, scintigraphy, biopsy research, determination of the alpha-fetoprotein in blood. The high level of the alpha-fetoprotein (> 1000 ng / mL) is found in 2/3 patients with HLC [3]. At the autopsy HLC manifests in the form of single or multiple soft, yellow-green or mottled nodes of different forms and sizes (diameter from 0.5 to 15 cm) [3]. According to the type of growth one allocates multinodular, single-nodular and diffuse forms of the HLC [6].

Case presentation

A 64-year-old female Ukrainian patient was admitted to the anesthesiology and reanimation department with complaints of pain in all the parts of the abdomen, lumbar pain, general weakness, nausea.

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Undiagnosed hepatoblastoma

From anamnesis we know that the patient has felt discomfort in the right upper quadrant for about 4 days. On the eve patient was consulted by urologist and general surgeon because of overall health deteriorating. Urgent surgical and urological pathologies at the time of inspection were not found. The patient has had rheumatoid arthritis for 17 years and has regularly taken NSAIDs. She was operated on about fibromyoma 16 years ago, suffered from hypertension, received treatment repeatedly because of atherosclerosis, but she did not receive a systemic therapy.

When the patient was admitted to the hospital, she had a broken mind (stupor type), a slurred speech, a pale-pink skin, arterial pressure was 25/0 mm Hg, heart rate was 100/min, a frequency of breathing movements was 20 per minute, oxygen saturation was of 90%. The level of total bilirubin was to 89.31 mmol/l (22.3 mmol/l - direct and 67.01 mmol/l - indirect fractions), total cholesterol - 11.2 mmol/l (HDL - 1.42 mmol/l, LDL - 8.14 mmol/l, TG-1.64 mmol/l, IA - 6.89), total protein 68.4 g /l (albumins - 54.1%, α1-globulins - 4.6%, α2-globulins - 8.3 %, β-globulins - 19.9%, γ-globulins - 13.1%).

Based on the data from anamnesis, laboratory results and ultrasound examination of the abdomen (conclusion: in the projection of the abdominal aorta there was revealed a formation of 120 x 70 mm irregular shape with a wall thickness of 15 mm), the negative dynamics of clinical blood parameters (Tab. 1), hemodynamics and other important parameters (Fig. 1) it was diagnosed: Dissecting abdominal aortic aneurysm.

For the differential diagnosis with acute coronary syndrome there was made an electrocardiographic research (Fig. 2).

The patient received a medical treatment: infusion therapy of plasma substitutes (NaCl 0.9% 200 ml 1 time per day, Ringer’s solution 200 ml 1 time per day, 5% glucose solution 200 ml 1 time per day, 6% hydroethyl-amylum 400 ml1 time per day, 8% gelatin polysuccinate 200 ml 1 time per day), antibiotics (ceftazidime 1.0 2 time per day), steroid hormones (dexamethasone 0.016 per day), catecholamine hormones (4% dopamine 10 ml per day, 0.1% epinephrine 6 ml).

After 10 hours despite the medical therapy there was established a biological death. At the autopsy abdominal and pleural cavity was free, without adhesions. Lungs were with edema, pink foamy liquid ran on the cut (Fig. 3 - E). Bronchial mucosa had a pink color. Size of the heart was 12x10x5,5 cm, with signs of obesity, small and large focal areas of cardiosclerosis, concentric hypertrophy of myocardial, fibrotic changes of valvular, especially mitral(Fig. 3 - F). Coronary artery was winding, with signs of atherosclerosis, unevenly calcified. The aorta was with pronounced...
symptoms of atherosclerosis and atherocalcinosis, aneurysms were not found. Mucosa of the stomach had a single small erosion and was smoothed and pale. Intestinal mucosa was hyperemic, folding was saved. Examination of liver drew the attention because of its enlarged size (31x20x17x15 cm), tuberosity of the surface, multiple formations in the blood vessels and bile ducts of whitish-yellow color, that had a diameter of 2 cm, fuzzy contour, soft texture and a tendency to germinate. These nodal formations were evenly distributed throughout the liver parenchyma, were surrounded by small hemorrhages at liver tissue (Fig. 3 - A, 3 - B).

The left kidney was small (9x4x3,5 cm), with signs of fatty degeneration. The right kidney was increased (15x9x7cm), with a mottled surface (Fig. 3 - C) large mottled kidney), on the incision - with signs of obesity and blood supply disorders - infarcts (Fig. 3 - D). The mucous membrane of the renal pelvis had a grayish-red color and the ureters were cyanotic-pale (Fig. 3 - D). The spleen was enlarged (18x9x7cm) and had a soft consistency. Internal female genital organs were missing.

With the further research we didn’t find metastases in other

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Fig. 3. Autopsy of patient: Liver: A - organ surface; B - internal structure. Right kidney: C - appearance; D - surface cut. Lungs: E – surface cut. Heart: F - organ surface.
Undiagnosed hepatoblastoma organs.

Histological examination revealed that the formations in liver tissue were atypical hepatocytes with a high mitotic index. Architectonics of the organ was broken, tumor cells had round or irregular polygonal forms and exaggerated the size of typical hepatocytes (1.5-2 times), also, they were with a high nuclear-cytoplasmic index, had a basophilic cytoplasm with fatty and granular inclusions. The nuclei had a round or irregular shape and were hyperchromic, optically - uniform.

In the preparation there were optically empty vessels and bile ducts in the portal areas. Many cells were presented by atypical hepatocytes with several (2 to 5) nuclei of different size, abnormal mitosis. According to the results of the histological examination it was determined that the tumor belonged to poorly differentiated HLC (Fig. 4 - A, 4 - B).

Histological examination of the kidneys revealed that the right kidney was hypertrophied with the phenomena of pyramids blood seepage and cortex substance of the organ, perivascular edema, vessels hyperemia (Fig. 4 - C, 4 - D). In the wall of the aorta there was revealed a deposit of calcium salts, proliferation of connective tissue, degradation of elastic fibers (Fig. 4 - E). Also, the myocardial research showed the hypertrophy of cardiomyocytes (Fig. 4 - F).

**Discussion**

HLC is a polyetiological disease, which can clinically show a variety of symptoms - from asthenic-vegetative syndrome to the symptoms of hepatocellular insufficiency. There is a problem of early diagnosis and a treatment of this disease. Literature data about the similarity of clinical symptoms of HLC progress and dissecting aneurysm in abdominal part of aortic are absent.

A 64-year-old female Ukrainian patient was admitted to the anesthesiology and reanimation department, where a multinodular form of low differentiated HLC was found. A clinical picture of this pathology in this case resembled the symptoms of dissecting aneurysm in abdominal part of aortic. The possibility of aortic lesions testified –an advanced age of the patient, a female sex, atherosclerotic, hypertensive, rheumatic anamnesis and also the results of additional research methods, including ultrasound of the abdomen, the deterioration of the blood pressure dynamics and clinical blood indicators. Despite the complex diagnostic and therapeutic manipulation - treatment of the patient was without a positive result.

The lethal result was caused by the extremely difficult general condition of the patient, the late stage of the disease, insufficiency and inaccuracy of the diagnostics and, as a result, an incorrectly chosen treatment.

Fig. 4. Histological examination of the liver, kidneys, aorta and heart. Staining by hematoxylin – eosin. A. Liver zoom × 150; B. Liver zoom × 400; C. Left kidney zoom × 150; D. Right kidney zoom × 150; E. Aorta zoom × 400; F. Heart zoom × 150.
Conclusions
The clinical course of hepatoblastoma was accompanied by non-typical signs of liver disease. A non-typical course of the disease led to the erroneous diagnosis: dissecting abdominal aortic aneurysm. An ultrasound lifetime picture of morphological changes in the patient was assessed incorrectly, causing the discrepancy between the clinical and final posthumous diagnosis.

Consent
Written informed consent was obtained from the patient’s next of kin for publication of this case report and any accompanying images.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
AR was responsible for final approval of the article, analysis and interpretation of the data. AP and VS were involved in the writing of the manuscript. AR, AP and VS participated in writing and editing the manuscript. AP and VS performed the autopsy and anatomopathological evaluation. All authors read and approved the final manuscript.

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Conflict of interest: None

References: