Insulinomas : about 9 cases

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Introduction:

Insulinomas are rare, their incidence is estimated at 4/million/an [1]. They are the most common tumors of the endocrine pancreas [2]. They are frequently small lesions, malignant in 10% of cases [3]. We report a series of 9 cases of insulinoma.

Objective:

The purpose of this work is to analyze the clinical, morphological biological and aspects Of insulinomas.

According to biology:

Prolonged diagnosis fast



 \rightarrow Diagnosis fast was positive in 85.71% of cases with 83.33% from the first day.

Patients and methods:

This is a retrospective study spanning 29 years 1984 until December 2013) concerning (from patients with insulinoma collected in our service.

By age:

Results:



Localisation:



 \rightarrow Insulinoma was essentially located in the head or body of the pancreas.

 \rightarrow It was visualized by ultrasound endoscopy in 100% of cases and by CT or MRI with less sensitivity.

Treatment and evolution:

→ Modal age: 18 and 34 years \rightarrow Average age: **35 years** (18 to 66 years)



- → Male predominance 77.77% of cases
- → 66,66% benign
- \rightarrow Sporadic in **100%** of cases studied.

Delay in diagnosis::



 \rightarrow The treatment was either medical, surgical or medical + surgical with equal frequency of **22.22%** of cases. \rightarrow Resolution of hypoglycemia was obtained in 33.33% of cases.

Discussion and conclusion:

In our series of insulinomas, we note the male predominance, it was often small, unique and sporadic malignant tumors were more frequent tumors, compared with literature data 33.33% vs 10% [3], finally ultrasound endoscopy has excellent sensitivity to diagnose insulinomas [4].



 \rightarrow Pathology revealed all was in cases by neuroglucopenic signs with a delay to diagnosis ranging from months to years.

References:

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