Diagnostic Imaging of Retinoblastomas: About 54 Cases
M. Badraoui*, D. Basraoui, H. Jalal

Department of Pediatric Radiology, Mother and Child Hospital, Marrakech, Morocco

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*Corresponding author: Marouane Badraoui

Abstract

Retinoblastoma is the frequent ocular malignant tumor of the genes in children. It mainly affects infants and young children with a very high cure rate of around 95%, provided that an early radiological diagnosis is made with an accurate extension report. Show the contribution of imaging in the diagnosis and management of retinoblastomas. Retrospective study between January 2012 and October 2019, 54 cases of retinoblastoma in children under 15 years.

RESULTS

The average age of our patients was 3 years (2 months - 15 years) with a male predominance (sex ratio 1.4). Leukocoria followed by exophthalmia were the most frequent telltale signs. It was unilateral in 84% of the cases.

Ultrasound showed a mass more echoic than vitreous, vascularized with color Doppler, with very hyperchoic areas with or without posterior shadow cone (micro or macro calcifications) in 90% of cases (Figure 1). Orbital CT showed extra-scleral orbital extension in 5% of cases (Figure 2), retro-laminar invasion particularly of the optic nerve in 5% of cases (Figure 2), calcifications in 93% of cases (Figure 3) and intense contrast enhancement with retinal detachment in 5% of cases.

INTRODUCTION

Retinoblastoma is the frequent ocular malignant tumor of the genes in children. It mainly affects infants and young children with a very high cure rate of around 95%, provided that an early radiological diagnosis is made with an accurate extension report.
DISCUSSION

Retinoblastoma is a neuroectodermal tumor developed from the retina [1], it is malignant of genetic origin by constitutional mutation of the gene Rb 1 (13q14) this mutation predisposes to other types of cancers including sarcomas [1], its incidence is in the order of 1/15000 to 1/20000 births [1], occurred before the age of 3 years in 98%. Its seat is unilateral in 60% and bilateral in the family forms [1]. The diagnosis of retinoblastoma is clinically suspected by leukocoria and / or exophthalmos [2], is confirmed by a fundus examination which objectifies a whitish tumor with angiomatous dilatation of the retinal vessels [2], there are three forms of retinoblastoma the form endophytic interesting the inner layers of the retina, an exophytic layer interesting the outer layers of the retina and a diffuse infiltrative form that is misleading is often affects older children.

When the fundus examination is not characteristic or impossible, imaging plays a role in the confirmation of diagnosis, evaluation of the size of the tumor process, search for calcifications, assessment of local extension and the search for metastases to the central nervous system. Ultrasonography is the first examination to be performed after the fundus examination ideally performed under general anesthesia and objective a solid heterogeneous echogenic mass vascularized with color Doppler and having calcifications [2]. Orbital CT is essential to study the extra-scleral orbital extension, the retro-laminar invasion in particular of the optic nerve, the presence of calcifications which are present in 93% of the cases grouped in clusters in the vitreous and the intense contrast enhancement with or without detachment of the retina [3]. Orbital CT can also be used to search for other intra-cranial abnormalities such as intracranial tumor extension, midline tumors and malformative abnormalities [3].

CONCLUSION

Imaging plays a key role in the positive diagnosis of retinoblastoma when the fundus is not characteristic or impossible (intra-vitreous haemorrhage), and in the pre-therapeutic extension report to adapt the therapeutic management and improve the prognosis in the long term.

REFERENCES