Thrombosis in renal artery in a newborn. Presentation a review of a case

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SUMMARY

This report describes a newborn who presents high pressure values and absence of pulses in inferior extremities in his 4th day of life. Posterior to placement of umbilical arterial catheter, confirming diagnostic of thrombosis in aorta and left renal artery. A gamagraphy with MAG 3 was performed that gave evidence of left renal exclusion and aorta and left renal angioresonance. A management with no fractional heparin and streptokinase during 6 hours was initiated. The compromise in lower extremities was totally recovered, improving presure values in a few days. Following studies of renal function and growth give evidence of left renal artery flow and some function. This report describes an improvement with anticoagulant and thrombolitic treatment of a newborn kidney depite of a prolonged ischemia.

Keywords: Aorta thrombosis; Renal thrombosis; Newborn hypertension; Renal isquemia; Anticoagulan; Thrombolitic.

Trombosis de la arteria renal en un recién nacido. Presentación de un caso y revisión

RESUMEN

Se describe un neonato que presenta cifras tensionales elevadas y ausencia de pulsos en miembros inferiores en su cuarto día de vida después de la colocación de un catéter umbilical arterial, evidenciando trombosis en arteria aorta y renal izquierda. Se realizó gammagrafía con MAG 3 que mostró exclusión renal izquierda y angioresonancia de aorta y renal izquierda. Se inició manejo con heparina no fraccionada y estreptoquinasa durante 6 horas. Se recuperó totalmente el compromiso en miembros inferiores, mejorando cifras tensionales a los pocos días. Estudios de seguimiento de función renal y crecimiento evidencian flujo de arteria renal izquierda y algo de función. Se describe mejoría con tratamiento anticoagulante y trombolítico de un riñón neonatal a pesar de sufrir isquemia prolongada.

Palabras claves: Trombosis aorta; Trombosis renal; Hipertensión neonatal; Isquemia renal; Anticoagulante; Trombolítico.

This pathology has been recognized over the last decade due to the use of umbilical arterial and venous catheterization. Vascular spasms occur as a result of arterial constriction and in early stages may be difficult to distinguish them from thrombolytic states, which generally are followed by pallor and cyanosis of the limbs after the insertion of catheters^{1,2}. This complication may result in damages to different organs, migration of thrombus, embolization to other areas and infections. At artery level, compromise may extend to the aorta and its

ramifications to lower limbs, renal and cerebral. Systemic therapy with fibrinolytic agents continues to be controversial. However, treatment with these agents is considered to be positive in spite of their use several days after the diagnostic took place.

REPORT OF THE CASE

The case was present at the Neonatal Unit (CIRENA) of the Hospital Universitario del Valle. Corresponds to

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Figure 1. Abdominal echocardiography performed on the ninth day. Evidencing the thromb in the abdominal aorta partially obstructing it

a masculine neonatal of 34 weeks, weighing 2,035 g, hospitalized for presenting signs of breathing difficulties, for which the umbilical venous and arterial catheterization was performed. On his fourth day of life, the patient presents high blood pressure and on the seventh day acrocyanosis is evidenced, absence of pulsations and arterial tension in the lower limbs is not present. on the eighth day the umbilical catheters are removed, the patient continues to be hyper tense and changes in his lower limbs, previously described are present. Echocardiography is performed observing a thromb at the aorta level. The diagnostic was confirmed on the ninth day by means of a doppler echocardiography (Figures 1, 2) which evidenced a thromb partially compromising the aorta and completely compromising the left renal artery by not detecting blood flow to the left kidney.

The anticoagulants protocol was initiated with non fractioned heparin (initial Dose 75 U/k/hour and continued with 20 U/k/hour) and endovenous thrombolytic therapy

y with endovenous estreptokynase (4,000 U/k/initial dose and continued with 2000 U/k/hour for six hours). High blood tension persisted until the sixteenth day in spite of medical antihypertensive treatment and that the perfusion and flows to the lower limbs was completely recovered eight days after having initiated the thrombolytic handling of the patient. The patient was evaluated by a medical board which requested a renal gammagraphy with MAG3 (Figure 3) and an abdominal and renal angioresonance (Figures 4 a,b,c), which evidenced the absence of the left renal function and absence of blood flow to the left lower limb and left kidney respectively.

Antihypertensive treatment was continued (propanolol) and with oral anticoagulants (warfarin). At the age of three months a doppler renal echography control was performed which evidenced growth of the left kidney and with a DMSA Renal Gamagraphy which evidenced function of the left kidney. Arterial tension,



Figure 2. Doppler renal echography performed on the ninth day. Evidence of the absence of flow to the left renal artery is observed



Figure 3. Renal gammagraphy MAG 3. No evidence of left renal perfusion and secondary functional exclusion. Low tubular function compromise appears on the right kidney

creatinine, growth and development of the left limb were normal during the following six months. A doppler renal echography control was performed evidencing asymmetry in renal dimensions (right kidney length: 4.9 cm; left kidney 3.5 cm) evidencing flow in the left renal parenchyma (Figures 5 a, b).

DISCUSSION

The frequency of vascular complications associated with the use of arterial umbilical catheters in neonates is relatively high. Approximately 30%, depending of the diagnostic method and the symptomatology of the patient '1-5.



Figure 4. Abdominal aorta and renal angioresonance. Performed on the twenty eighth day of life. Shows evidence of blood flow to the lower left limb, persistence of the thromb at the level of the aorta, absence of blood flow to the left kidney and reduction in size of the kidney



Figure 5. Renal doppler echography at the age of six months. Asymmetric renal size

Clinical signs such as the absence of pulses, changes in color of the limbs, hypertension, hematuria and oliguria confirm the diagnostic. Verification may be made through angiographies but in some instances may prove to be of difficult access. Doppler Echography is also available but sensitivity and specificity may be uncertain^{1,2}. A detailed clinical history of ancestor and family members presenting thrombolyc and thrombolitic processes must be recorded.

The first step for the management of the process is to provide support care that must include paraclinic studies such as electrolytes, correction of anemia, sepsis, thrombocytopenia and the removal of catheters. The therapy with anticoagulants consist mainly of the use of standard or not fractioned heparin (HNF), low molecular weight heparins (HBPM) and thrombolytic therapy, with or without surgical intervention. Surgical intervention will be assessed accordingly with the degree of severity of the thrombus¹⁻³.

Arterial hypertension due to reno vascular lesion is

common in survivors of neonates presenting aortic lesions. A study, with a four year follow up performed by Seibert et al.⁶ documented renal growth impairment in 78% of the survivors of aortic thrombolysis. Many authors conclude that the absence of renal blood flow may early predict an irreversible renal lesion having high mortality rates. We initiated treatment in spite of the diagnostic made several days after the occurrence of the event, with previous valuations and strict follow up with fibrinogen, dimer D, coagulation times and hemograms. This case suggests a recovery of the perfusion in the left inferior limb and in the renal function, in spite of the appearance of a ischemia interval of nine days; accepting the theory that an immature kidney may tolerate better a hypoxic ischemia without a time limit for defining the treatment.

The management of this pathology in the neonatal stage is controverted and is performed based on cases reported and studies extrapolating the reports of descriptive studies on children. The therapy for neonates must be considered in an individual manner and the riskbenefit of each treatment must be assessed together with the possible compromise of vital organs¹⁻⁴.

Finally a strict follow up must take place during the infancy period evaluating the presence of hypertension, renal function anomalies and growth and development of lower limbs[.]

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