KARYA ILMIAH:

SEMINAR NASIONAL

Congenital Colonic Atresia

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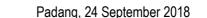


"DEALING WITH GLOBAL ISSUES IN PEDIATRIC SURGERY" PADANG, 18 – 20 OKTOBER 2018

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Hal : Pengumuman Poster PIT PERBANI

: 50/PIT PERBANI/IX/2018

Kepada Yth.

Nomor

dr. Azka Darajat

di

Tempat

Dengan hormat,

Dengan surat ini kami menginformasikan kepada Sejawat bahwasanya abstrak dengan judul :

"Congenital Colonic Atresia"

Telah diterima oleh panitia Pertemuan Ilmiah Tahunan (PIT) ke XXVI PERBANI Padang untuk presentasi **Poster** pada :

Hari/Tanggal : Jum'at-Sabtu / 19-20 Oktober 2018

Kode Poster : PO19

Dengan aturan presentasi sebagai berikut :

- 1. Presentasi poster menggunakan elektronik poster (e-poster) yang dikemas dalam format jpg. Dengan layout portrait dan resolusi 1920x1080 dpi.
- 2. Bahasa tertulis dalam presentasi menggunakan bahasa Inggris
- 3. Presentasi ditayangkan dalam bentuk slide show selama acara ilmiah berlangsung
- 4. E-poster paling lambat diterima panitia pada hari Jum'at / 12 Oktober 2018

Demikianlah kami sampaikan, atas perhatiannya di ucapkan terimakasih.

Hormat Kami,

idi Pratama Arnofyan, SpB.SpBA

Ketua Panitia

CONGENITAL COLONIC ATRESIA: A RARE CASE REPORT

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ABSTRACT

Background: Colonic Atresia is a rare form of congenital abnormanilies in

children. The incidence is approximately 1:66.000 live birth. Method: This report

is a retrospective review of one patient with colonic atresia presented to our

hospital. Results: A 2-day-old female baby transferred to our hospital from

peripheral hospital. The baby diagnosis from the peripheral hospital was low type

obstruction with differential diagnosis congenital megacolon. At the time of

admission to our hospital, the child presented with history of abdominal distention

and didn't pass meconium after 24 hours. Abdominal X-ray was taken and showed

distended bowel and air fluid level. Laparotomy exploration surgery revealed

colonic atresia type II in descending colon and colostomy was performed.

Conclusions: Colonic atresia is a rare case that may cause a miss diagnosed. A

good management will determine the outcome

KEY WORDS: Colonic atresia, surgery

INTRODUCTION

Colonic Atresia is a rare form of congenital abnormanilies in children. The incidence is approximately 1:66.000 live birth. Colonic atresia can occur in ascending colon, hepatic flexure, transverse colon, splenic flexure, descending and sigmoid colon. The authors were encouraged to report this case because it was rare.

METHOD

This report is a retrospective review of one patients with colonic atresia presented to our hospital.

RESULT

Case Presentation

A 2-day-old female baby transferred to our hospital from peripheral hospital. The baby diagnosis from the peripheral hospital was low type obstruction with differential diagnosis congenital megacolon. The baby was born to a 33- year old multigravida at 39 weeks gestation by spontaneous vaginal delivery with a birth weight of 3.3 kgs. At the time of admission to our hospital, the child presented with history of abdominal distention and didn't pass meconium after 24 hours. Oxygen and oral gastric tube were administered.

Abdominal X-ray was taken and showed distended bowel and air fluid level. A diagnosis of colonic atresia was made. Emergency surgery was planned. Fluid, electrolytes, parenteral nutrition and intravenous antibiotics were administered before the surgery. Laparotomy exploration surgery was performed through

tranverse incision. Surgery revealed colonic atresia type II in descending colon and colostomy was performed.



Image 1. Abnominal X-Ray show distended bowel and air fluid level



Image 2. Surgery revealed colonic atresia type II in descending colon

DISCUSSION

Colonic atresia is a rare case. Utero vascular insufficiency after organogenesis was considered as the etiology. There are some possible causes that can disrupt the vascular such as volvulus, intussusception, embolic or thrombotic events, and incarceration or strangulation secondary to hernias or abdominal wall defects.² Colonic atresia can be classified base on Grosfeld classification.

Because of its rarity, it is usually not thought of in the differential diagnosis of neonatal intestinal obstruction.³ The diagnosis is still likely to be easily missed and the surgery could be delayed.⁴ Sign of distal bowel obstruction present in the colonic atresia. We can be suspicious of the diagnosis of colonic atresia if the baby is failed to pass meconium and radiological investigations can play role in early detection.^{4,5} The degree of distention proximal to the obstruction is more marked in the colon than in small intestine.

Laparotomy is usually performed. The obstructing lesion and dilated intestine are resected. Minimizing size mismatch for the anastomosis is important to facilitate the intestinal fuction. Because of the association with Hirschsprung disease, it is important to do rectal biopsy to evaluate ganglion cells and prevent complication.^{2,5}

CONCLUSION

Colonic atresia is a rare case that may be challenging to be diagnosed. Because of the rarity, colonic atresia can be miss diagnosed. A good management will determine the outcome of treatment.

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Congenital Colonic Atresia ARare Case Report



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1) Background

- Colonic atresia is a rare form of congenital abnormalities.
- The incidence is approximately 1:66.000¹.

2) Method

This report is a retrospective review of one patients with colonic atresia presented to our hospital.

3) Result

- A 2-day-old female baby presented with history of abdominal distention and didn't pass meconium after 24 hours.
- Abdominal X-ray was taken and showed distended bowel and air fluid level.
- Laparotomy exploration surgery revealed colonic atresia type II and colostomy was performed.







(4) Discussion

- Colonic atresia is still likely to be easily missed and the surgery could be delayed^{3,4}.
- Radiological investigations can play role in early detection^{2,4}.

5 Conclusion

Colonic atresia is a rare case that may be challenging to be diagnosed. A good management will determine the outcome of treatment.

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Congenital Colonic Atresia ARare Case Report



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References: