

## **Omphalomesentric Duct Anomalies - our experience**

<sup>1</sup>Dr. Manisha Albal, Associate Professor, Department of Paediatric Surgery, N.K.P Salve Institute of Medical Sciences and Research Centre & Lata Mangeshkar Hospital, Hingna, Nagpur.

<sup>2</sup>Dr. Shriniket Sawarkar, Assistant Professor, Department of General Surgery, N.K.P Salve Institute of Medical Sciences and Research Centre & Lata Mangeshkar Hospital, Hingna, Nagpur.

**Corresponding Author:** Dr. Shriniket Sawarkar, Assistant Professor, Department of General Surgery, N.K.P Salve Institute of Medical Sciences and Research Centre & Lata Mangeshkar Hospital, Hingna, Nagpur.

**Citation this Article:** Dr. Manisha Albal, Dr. Shriniket Sawarkar, “Omphalomesentric Duct Anomalies - our experience”, IJMSIR- January - 2023, Vol – 8, Issue - 1, P. No. 116 – 120.

**Type of Publication:** Original Research Article

**Conflicts of Interest:** Nil

### **Abstract**

Om phalo mesentric ducts is a major embryonic anomaly encountered in paediatric surgery we intended to share our experience with other, aim of this study was to evaluate the clinical presentation and surgical management of om phalo mesenteric duct (OMD) remnants in paediatric population.

**Materials and method:** This an Observational study of Om phalo mesentric Duct remnants treated at the Department of Paediatric Surgery in our tertiary care Centre carried out under the period of three years. These anomalies occur in approximately 2% of the population and may remain silent throughout life, or may present incidentally with an intra-abdominal complication.

**Results:** A total of 26 (mean=6.5, median=6) patients were encounter in the time period amongst which 2 patients were below 1 month of age, 12 below 1 year, 5 below two years and 6 below 3 years of age. Depending upon the on clinical and radio logical examination patients provisional diagnosis were made and confirmed during surgical intervention.

**Discussion:** OMD remnants present as a wide spectrum of anomalies depending on the stage of arrest of normal process of involution, affecting the male population predominantly. Similar sex distribution was observed in the current series. The incidence of urachal anomalies is about 1:5000 in general population. The malignant epithelial neoplasm can arise from urachal remnants and form the urachus carcinoma, accounting for less than 1% of all bladder cancers.

However, most urachal carcinomas are always diagnosed in advanced stages and associated with a poor prognosis.

**Keywords:** Omphalomesentric, Embryonic, Malignancy, Benign, urachus, neoplasm.

### **Introduction**

The omphalomesentric duct (OMD) is an embryonic structure, which connects the yolk sac to the midgut and failure of its resorption results in various anomalies including Meckel's diverticulum, patent vitelline duct, fibrous band, sinus tract, umbilical polyp and cyst, enteric fistula with ileal intussusceptions prolapsing over the umbilicus or hemorrhagic umbilical mass. These anomalies occur in approximately 2% of the population

and may remain silent throughout life, or may present incidentally with an intra-abdominal complication (1). In newborns and infants these anomalies manifest as a mass, prolapsing ileal loop or discharge over the umbilicus and needs urgent surgical intervention (2). The major burden of these malformations is encountered in the paediatric population when they present with complication caused by an underlying remnant. The intra-abdominal components of OMD remnants may remain asymptomatic or are incidentally discovered during a laparotomy for other reasons. However, those who become symptomatic can have a wide range of presentation based on the underlying anomaly. This includes complications such as intestinal obstruction secondary to the band, intussusception, internal herniation, volvulus, or acute abdominal pain due to Meckel's diverticulitis (3).

Various surgical options have been described in the literature, ranging from wedge or segmental resection in case of Meckel's diverticulum to umbilical exploration in case of umbilical discharge. Most reports on symptomatic OMD focus on Meckel's diverticulum, whereas other related anomalies are given little attention. The basic aim of this study was to gather our institutional experience on the various OMD remnants in children with an emphasis on the age and clinical presentation, intra-operative findings, surgical intervention performed, and the histopathological outcome (4).

### Material and methods

This is an Observational study of paediatric patients of both sexes with some form of Omphalo mesenteric Duct remnants treated at the Department of Paediatric Surgery, N.K.P Salve institute of medical sciences and research Centre, Nagpur carried out under the period of three years from January 2019 to December 2021. Information about patient's sex, age at presentation, type of anomaly identified, and surgical intervention performed was

gathered for conducting the study. Apart from baseline Hema to logical workup, patients with acute abdomen had X-rays/ultrasound of the abdomen, and CT scan was also added wherever needed. Radio logical investigations were not required for umbilical anomalies except those in doubt of internal communication with the gut, where a contrast study was conducted. Child with intestinal obstruction or any acute abdominal episode underwent laparotomy after initial resuscitation. Surgical procedures were decided according to condition of the paediatric patient and peri-operative findings were noted. Those with a narrow base Meckel's diverticulum underwent wedge resection, while those with a wide base underwent resection and anastomosis. However, in case of gross peritoneal cavity contamination, friable, and edematous gut, ileostomy was made irrespective of the anatomy of Meckel's diverticulum. In patients with incidentally discovered Meckel's diverticulum, no intervention was performed. However, in case where it was required due to primary surgical condition requiring diversion, site of Meckel's diverticulum was chosen for ileostomy after its segmental resection. In cases of umbilical anomalies, polyps were excised under anesthesia, whereas patients with the patent OMD underwent laparotomy. Histopathological examination of specimen was done and recorded.

### Results

A total of 26 (mean=6.5, median=6) patients were encountered in the -----time period amongst which 2 patients were below 1 month of age, 12 below 1 year, 5 below two years and 6 below 3 years of age.

Table 1: age distribution.

Age	No. of Patients
0 to 1 month	2
1 to 12 months	12
12 to 24 months	7

24 to 36 months	6
Total	27

Amongst the sample size acquired 16 patients were male and remaining 10 were female.

Table 2: gender distribution.

Male	17
Female	10
Total	27

With a clinical variability in presentations of omphalo mesenteric duct remnants around 9 patients presented with umbilical anomalies, 6 with signs of intestinal obstruction, 5 with abdominal pain one with rectal bleeding and 4 were found incidentally.

Table 3: clinical presentation

Clinical Presentation	No. of Patients
Intestinal obstruction	6
Abdominal pain	5
Umbilical anomalies	10
Rectal bleeding	1
Incidental	4

Depending upon the on clinical and radiological examination patients' provisional diagnosis were made and confirmed during surgical intervention in which 9 patients were diagnosed with Meckel's diverticulum amongst which 4 were found incidentally. Three patients were diagnosed with patent vitello-intestinal duct, two each were diagnosed with Urachal cyst, Urachal sinus, Exomphalos minor, urachal abscess and bladder diverticulum respectively. One each with Patent urachus, Umbilical polyp, Umbilical Granuloma, Umbilical sinus and exomphalos major respectively.

Table 4: diagnosis.

Diagnosis	No. of patients
Patent Urachus	1
Urachal Cyst	2

Urachal Sinus	2
Urachal Abscess	2
Bladder diverticulum	2
Umbilical Polyp	1
Umbilical granuloma	1
Umbilical Sinus	1
Patent Vitello intestinal duct	3
Meckel's Diverticulum	9
Exomphalos Major	1
Exomphalos Minor	2
Total	27

### Surgical intervention

Amongst the sample size of 27, 10 patients underwent segmental resection and anastomosis, 6 patients underwent segmental resection and ileostomy. 11 patients underwent exploration of umbilicus.

Table 5: Surgical intervention

Surgery	No. of Patients
Segmental resection and Anastomosis	10
Segmental resection and ileostomy	6
Exploration	11
Total	27

### Discussion

OMD remnants present as a wide spectrum of anomalies depending on the stage of arrest of normal process of involution, affecting the male population predominantly (5). Similar sex distribution was observed in the current series. Intestinal obstruction is the most common presentation of OMD remnants with underlying mechanism such as bands, intussusception, or volvulus (6). In our study, more than half of the patients presented with intestinal obstruction. Meckel's diverticulum forming bands with surrounding small bowel or the

mesentery was the most predominant cause seen in 70.45% of cases. Meckel's diverticulum can act as a lead point in secondary intussusceptions (7).

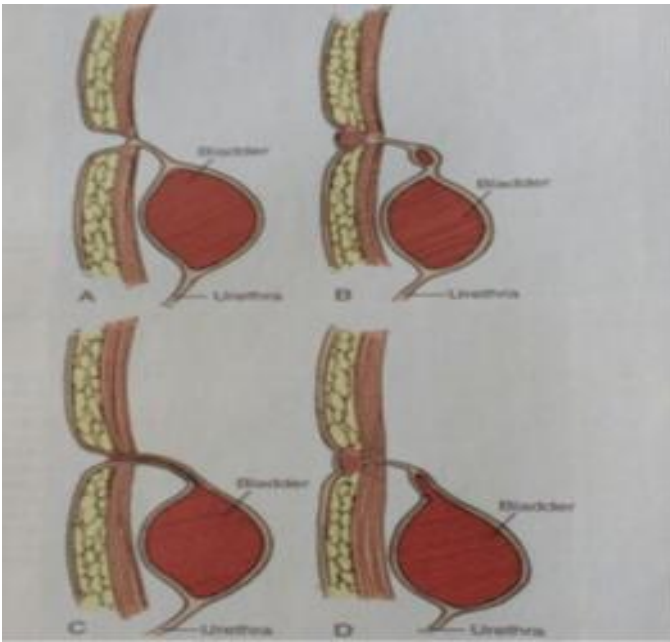


Fig 1: Classification of urachal anomalies.



Fig 2: Clinical photo.

Classification: Four distinct types of urachal anomalies are defined. In order of frequency, they are a patent urachus (50%), an urachal cyst (30%), an umbilical urachal sinus (15%), and vesicourachal diverticulum (3%-5%).

The incidence of urachal anomalies is about 1:5000 in general population (8). Based on the location of abnormal residual patency along the urachal tract, four types of congenital urachal anomalies are present, including patent urachus, urachal sinus, urachal cyst and vesicourachal diverticulum (9). Most of the urachal anomalies are detected incidentally and usually remain asymptomatic, while some may be misdiagnosed as other abdominal and pelvic diseases in the emergency room (10). The malignant epithelial neoplasm can arise from urachal remnants and form the urachus carcinoma, accounting for less than 1% of all bladder cancers (11). However, most urachal carcinomas are always diagnosed in advanced stages and associated with a poor prognosis (12)(13).



Fig 3: Patent urachus.

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