TITLE:
Development of Signs and Symptoms of Congenital Choledochal Dilatation: Its Relation to Pregnancy with Special Reference to Experimental Study in Guinea Pigs

AUTHOR(S):
AOKI, YOZO; SHIMADA, KOSUKE; KAWASHIMA, HIROAKI; KATSUMI, MASAHARU

CITATION:
AOKI, YOZO ...[et al]. Development of Signs and Symptoms of Congenital Choledochal Dilatation: Its Relation to Pregnancy with Special Reference to Experimental Study in Guinea Pigs. 日本外科宝函 1984, 53(2): 338-344

ISSUE DATE:
1984-03-01

URL:
http://hdl.handle.net/2433/208771

RIGHT:
Development of Signs and Symptoms of Congenital Choledochal Dilatation: Its Relation to Pregnancy with Special Reference to Experimental Study in Guinea Pigs

Yozo Aoki, Kosuke Shimada, Hiroaki Kawashima and Masaharu Katsumi

Department of Gastroenterological Surgery, Wakayama Medical College, Wakayama City,
(Director: Prof. Masaharu Katsumi)
Received for Publication, Oct. 25, 1983.

Introduction

Many theories have been proposed regarding the etiology of the congenital choledochal dilatation. Among them, extrinsic or acquired factors have been defended by some surgeons.

In this paper, the experimental study on the relationship of development of this disease to pregnancy and two cases of congenital choledochal dilatation developed on pregnancy are reported.

Part I. Experimental Study

The following experiments were made to elucidate the effect of pregnancy on the bile excretion dynamics of the gallbladder.

Materials and Methods

(1) Female Hartley strain guinea pigs weighing 400 to 500 g were used. The animals at the 30th and 50 to 60th day of pregnancy were employed as a pregnant model. Non-pregnant female guinea pigs were served as control. Each group consisted of 2 guinea pigs.

(2) After fasting overnight, the jugular vein was exposed, the blood was drawn for the measurement of estrogen and progesterone levels, and 2 mCi of technetium-99 m-N-(2,6-dimethylacetanilide)-iminodiacetic acid (99mTc-E-HIDA) was injected, and then the wound was closed under ether anesthesia.

(3) The animals were positioned under pinhole collimator of a LFOV scintillation camera (Searle) in the supine position under conscious state. Data from the gamma camera were stored in at 5 min intervals for 1 hr and processed by the analyser. Thereafter, 0.26 g/kg b.w. of yolk preparation was administered orally, and the scanning was continued for more 1 hr. These images were played back and the scans from each animal were inspected and the gallbladder area

Key words: congenital choledochal dilatation, pregnancy, hepatobiliary scintigraphy, estrogen, progesterone, 索引語: 先天性胆道拡張症, 妊娠, 肝・胆道シンチグラフィー, エストロゲン, プロゲステロン.
Present address: Yozo Aoki, M.D., Department of Gastroenterological Surgery, Wakayama Medical College, Wakayama City, Wakayama 640, Japan.
of interest was marked out on each. Then the radioactivity in the area was counted and the activity counts against time, i.e., the time-activity curve (T-A curve) was obtained.

Results

(1) Estrogen and progesterone levels in serum

The results obtained are shown in Table 1. Estrone (E1), esteradiol (E2) and progesterone in serum were extremely elevated in both gestational subgroups though there was no relation between estriol (E3) and pregnancy.

(2) Hepatobiliary scintigram and its T-A curve

Figure 1 shows the hepatobiliary scintigrams of pregnant and non-pregnant animals 50 min after the injection of $^{99m}$Tc-E-HIDA. In both subgroups on the 30th and 50 to 60th day of gestation, the gallbladder markedly enlarged compared to control.

T-A curve of the gallbladder area appeared to support this phenomenon as shown in Figure 2. In both gestational subgroups, the $^{99m}$Tc activity continued to rise at a rapid rate and reached its maximum within 60 min. The maximum activities were about twofold of that of the non-pregnant group. The administration of yolk preparation prompted the bile excretion from the gallbladder.

Part II. Clinical Cases

Case 1.

The patient, 27 year old multipara, was admitted under suspicion of congenital choledochal dilatation because she had experienced right hypochondrial pain at and after the 30th week gestation of her first and second pregnancies. At the same time, she had been pointed out by a physician a mass in the right upper quadrant. These signs and symptoms disappeared after each delivery. Various examinations yielded a diagnosis of congenital choledochal dilatation (Alonso-Lej I type).

She underwent a cholecystectomy and Roux-Y cystojejunostomy. The dilated common bile duct contained a calcium bilirubinate gallstone.

Follow-up of the patient 6 years later shows normal liver functions.

<table>
<thead>
<tr>
<th>Table 1. Estrogen and progesterone in serum of experimental animals.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Estrone (E1) (pg/ml)</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Estradiol (E2) (pg/ml)</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Estriol (E3) (pg/ml)</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Progesterone (ng/ml)</td>
</tr>
<tr>
<td></td>
</tr>
</tbody>
</table>
Case 2.

The patient, 39 year old multipara, was admitted with a chief complaint of intermittent colicky right hypochondrial pain and pyrexia. These symptoms developed at the 16th week of her every 3 gestations and continued till the end of each gestation. The diagnosis of congenital choledochal
CONGENITAL CHOLEDOCHAL DILATATION AND PREGNANCY

Dilatation was made and the patient underwent a resection of the dilated common bile duct and Roux-Y hepaticojejunostomy. The dilated common bile duct contained a calcium bilirubinate gallstone.

Three-year-follow-up has revealed she has been doing well.

Discussion

In the last 11 years from 1972 to 1982, fifteen cases of congenital choledochal dilatation complicated with pregnancy have been reported in Japan. As listed in Table 2, mean patient age was 26.2 years ranging from 23 to 33. Among these patients, six had formerly experienced similar signs and/or symptoms in their childhood or previous pregnant periods. The relationship between the developmental period of the clinical pictures and pregnant stages was very interesting. Some cases of them developed their signs and symptoms in the early

Table 2. A review of cases in Japanese literature.

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Age (yrs.)</th>
<th>Wks. of gestation on onset</th>
<th>Signs and symptoms*</th>
<th>Operation*</th>
<th>Prognosis of gestation</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>1⁴⁻</td>
<td>23</td>
<td>28</td>
<td>J, M</td>
<td>Cystojejunostomy</td>
<td>not described</td>
<td></td>
</tr>
<tr>
<td>2⁴⁻</td>
<td>23</td>
<td>28</td>
<td>J, M</td>
<td>Cystectomy, HJ</td>
<td>not described</td>
<td></td>
</tr>
<tr>
<td>3⁴⁻</td>
<td>26</td>
<td>24</td>
<td>J, M, P</td>
<td>Cystectomy, HJ</td>
<td>Artificial abortion</td>
<td></td>
</tr>
<tr>
<td>4⁵</td>
<td>not described</td>
<td>32</td>
<td>P, V</td>
<td>Cystojejunostomy</td>
<td>Cesarean section</td>
<td>Epigastric pain in former times</td>
</tr>
<tr>
<td>5⁵</td>
<td>27</td>
<td>not described</td>
<td>J, M, P</td>
<td>not described</td>
<td>Cesarean section</td>
<td></td>
</tr>
<tr>
<td>6⁴⁻</td>
<td>23</td>
<td>28</td>
<td>J, M, P</td>
<td>Cystojejunostomy</td>
<td>Premature delivery</td>
<td></td>
</tr>
<tr>
<td>7⁴⁻</td>
<td>26</td>
<td>32</td>
<td>J, P</td>
<td>Cystojejunostomy</td>
<td>Cesarean section</td>
<td></td>
</tr>
<tr>
<td>8⁷</td>
<td>33</td>
<td>24</td>
<td>J, M, P</td>
<td>Cystectomy, CD</td>
<td>Artificial abortion</td>
<td>Treated with a diagnosis of cholecystopathy at her age of 16 yrs.</td>
</tr>
<tr>
<td>9⁷</td>
<td>25</td>
<td>20</td>
<td>J, M, P</td>
<td>Cystectomy, HJ</td>
<td>Artificial abortion</td>
<td>Some time epigastric pain since her childhood</td>
</tr>
<tr>
<td>10⁴⁻</td>
<td>24</td>
<td>40</td>
<td>J, P, V</td>
<td>Cystoduodenostomy</td>
<td>Cesarean section</td>
<td>Primi para</td>
</tr>
<tr>
<td>11⁴⁻</td>
<td>24</td>
<td>36</td>
<td>J, P, V</td>
<td>Cystectomy, HJ</td>
<td>Cesarean section</td>
<td>Primipara</td>
</tr>
<tr>
<td>12⁴⁻</td>
<td>26</td>
<td>24</td>
<td>P, V</td>
<td>Cystectomy, HJ</td>
<td>Normal delivery</td>
<td>Primipara</td>
</tr>
<tr>
<td>13²⁻</td>
<td>27</td>
<td>28</td>
<td>J, P, V</td>
<td>Cystojejunostomy</td>
<td>Cesarean section</td>
<td></td>
</tr>
<tr>
<td>14**</td>
<td>27</td>
<td>32</td>
<td>P</td>
<td>Cystojejunostomy</td>
<td>Normal delivery</td>
<td>Multipara, similar symptoms in previous pregnancy</td>
</tr>
<tr>
<td>15**</td>
<td>33</td>
<td>12</td>
<td>P</td>
<td>Cystectomy, HJ</td>
<td>Normal delivery</td>
<td>Multipara, similar symptoms in previous pregnancies</td>
</tr>
</tbody>
</table>

* Abbreviations are: J, jaundice; M, abdominal mass; P, pain; V, vomiting; HJ, hepaticojejunostomy; CD, choledochoduodenostomy.
** Our cases
gestational stage in which their uteri were still small and they never pressed on the biliary system.

Many theories have been proposed regarding the etiology of the congenital choledochal dilatation\textsuperscript{1,2,13,22}. Alonso-Lej et al.\textsuperscript{1} stated that this disease is caused by two factors—weakness of the wall of the common bile duct in a specific segment and obstruction distal to it. They also described that one group defended extrinsic or acquired factors (pregnancy, abdominal trauma etc.) and the other congenital origin: obstructive factor localized at the junction of the choledochus with the duodenum, and a factor originating in the common bile duct proper. Many authors tend to support the second theory, however, some cases developed the signs and symptoms in an early gestational stage and this fact supports the possibility that compression to the dilated common bile duct by enlarged pregnant uterus is not always necessary in the development of the disease.

Imamoglu et al.\textsuperscript{8} reported that fifty young women were subjected to intravenous cholangiographic studies 4 to 10 hrs after delivery and that 56 per cent of the subjects showed delayed emptying of the gallbladder. Based on these clinical data, they succeeded to produce gallstones by prolonged administration of progesterone and estradiol in rabbits, and concluded that the increased incidence of gallstones in women following pregnancy perhaps was owing to changes in gallbladder functions secondary to elevated titers of the placental hormones.

Our experimental (Table 1, Figures 1 and 2) and clinical studies clearly demonstrate these facts. Case 1 in our series showed clinical symptoms at and after the 30th week gestation. These symptoms were suspected to be partly due to a compression to the biliary tract by pregnant uterus. Clinical signs and symptoms in Case 2 developed at the 16th week gestation when the uterus was still small. In this case, increased serum progesterone and estrogen might play one of the causative factors in the development of clinical symptoms\textsuperscript{4,5,11,14,16,18}.

Hence, it is possible that pregnancy especially in an early gestational stage, adding to the preexistence of congenital choledochal dilatation with anomalous choledochal junction, may lead to stagnation of bile in the extrahepatic biliary system and reflux of the pancreatic juice to the biliary system, thus predispose to cholangitis and provoke clinical signs and symptoms in selected cases.

**Summary**

From two experiences of congenital choledochal dilatation developed in pregnancy, experimental and clinical studies on the relationship of the sign and symptom of this pathological condition to pregnancy were investigated.

Pregnant guinea pigs revealed a dilated gallbladder and delayed emptying of the gallbladder on hepatobiliary scintigraphy regardless of their gestational stage. These phenomena were suspected to make the ground to lead to cholangitis under the pre-existence of congenital dilatation of the common bile duct.
References


22) Yotuyanagi S: Contributions to the etiology and pathology of idiopathic cystic dilatation of the common bile-duct with report of there cases; A new aetiological theory based on supposed unequal epithelial proliferation at the stage of the physiological epithelial occlusion of the primitive choledochus. Gann 30: 601–652, 1936.
先天性胆道拡張症における症状発現と妊娠との関係

和歌山県立医科大学消化器外科

青木 洋三，嶋田 浩介，川嶋 寛昭，勝見 正治

我々の教室で妊娠を契機に発症した先天性胆道拡張症を2例経験した。第1例は過去2回の分娩歴がある症例で，それぞれ妊娠時にも8か月目（30週）に右上腹部痛をもって発症した。第2例は過去3回の分娩歴がある症例で，それぞれの妊娠時にも3〜4か月目（16週）に右上腹部痛と発熱をもって発症し，以後分娩時までこれらの症状が間欠的に出現した。

妊娠モルモットを用いて肝・胆道シンチグラフィを行い，胆囊での胆汁排出動態を解析すると，胆囊の拡張と胆汁のうっ滞が妊娠初めに関係なく認められ，また同時に血中エストロゲン，プロゲステロン値を測定すると何れも高値を示した。

従来から本症における医療症状の発現機序に関しては，拡張した胆管の存在に加え，何らかの後天的因子が加わって発症するものと考えられてきた。その後の1つに妊娠子宮による胆道の機械的圧迫，あるいは妊娠に伴うホルモン変動の変化による胆汁の流出障害，胆汁の濃縮などが挙げられている。我々が今回経験した2症例はその発症時期に明らかに相違があり，妊娠期における本症の発症には妊娠子宮による胆道の物理的圧迫の他，妊娠に伴う母体の液性変化，すなわち増加するエストロゲン，プロゲステロンの関与が実験的にも推定された。