

Risk factors affecting postoperative pulmonary function in congenital diaphragmatic hernia

저자 (Authors)	Qianqian Wang, Quanhua Liu, Jingyu Zang, Jun Wang, Jie Chen
출처 (Source)	Annals of Surgical Treatment and Research 98(4), 2020.4, 206–213 (8 pages)
발행처 (Publisher)	대한외과학회 The Korean Surgical Society
URL	http://www.dbpia.co.kr/journal/articleDetail?nodeId=NODE09323192
APA Style	Qianqian Wang, Quanhua Liu, Jingyu Zang, Jun Wang, Jie Chen (2020). Risk factors affecting postoperative pulmonary function in congenital diaphragmatic hernia. <i>Annals of Surgical Treatment and Research</i> , 98(4), 206–213.
이용정보 (Accessed)	이화여자대학교 203.255.***.68 2020/05/18 03:54 (KST)

저작권 안내

DBpia에서 제공되는 모든 저작물의 저작권은 원저작자에게 있으며, 누리미디어는 각 저작물의 내용을 보증하거나 책임을 지지 않습니다. 그리고 DBpia에서 제공되는 저작물은 DBpia와 구독 계약을 체결한 기관소속 이용자 혹은 해당 저작물의 개별 구매자가 비영리적으로만 이용할 수 있습니다. 그러므로 이에 위반하여 DBpia에서 제공되는 저작물을 복제, 전송 등의 방법으로 무단 이용하는 경우 관련 법령에 따라 민, 형사상의 책임을 질 수 있습니다.

Copyright Information

Copyright of all literary works provided by DBpia belongs to the copyright holder(s) and Nurimedia does not guarantee contents of the literary work or assume responsibility for the same. In addition, the literary works provided by DBpia may only be used by the users affiliated to the institutions which executed a subscription agreement with DBpia or the individual purchasers of the literary work(s) for non-commercial purposes. Therefore, any person who illegally uses the literary works provided by DBpia by means of reproduction or transmission shall assume civil and criminal responsibility according to applicable laws and regulations.

Risk factors affecting postoperative pulmonary function in congenital diaphragmatic hernia

Qianqian Wang^{1,2,*}, Quanhua Liu^{3,*}, Jingyu Zang^{1,2}, Jun Wang^{1,2}, Jie Chen^{1,2}

¹Department of Pediatric Surgery, Xinhua Hospital Affiliated to Shanghai Jiao Tong University School of Medicine, Shanghai, China

²Department of Pediatric Surgery, Jiaying Maternity and Child Health Care Hospital, Jiaying, China

³Department of Pediatric Respiration, Xinhua Hospital Affiliated to Shanghai Jiao Tong University School of Medicine, Shanghai, China

Purpose: It is well known that congenital diaphragmatic hernia (CDH) in infants impacts pulmonary function rehabilitation after surgery. However, the risk factors of postoperative pulmonary function are still unclear. In this research, we analyzed the potential risk factors of postoperative pulmonary function in CDH patients in order to improve the clinical management of CDH patients.

Methods: Thirty-three cases CDH infants followed were enrolled from November 2016 to September 2018. Clinical data were reviewed. Tidal breathing pulmonary function testing was performed after surgery. Correlation between pulmonary function and clinical characteristics was evaluated using multivariate analysis of variance.

Results: Pulmonary dysfunction was detected in 87.9% patients (29 of 33). The defect size was found to be significantly larger in patients with obstructed and mixed ventilatory disorders ($P = 0.001$). Diagnosis of gestational age (GA) was also significantly earlier compared to restrictive ventilatory disorders ($P = 0.001$). Larger defect size, and earlier prenatal diagnosis of GA were detected in severe obstructive ventilatory disorders ($P = 0.007$, $P = 0.001$, retrospectively).

Conclusion: Most patients had various degrees of pulmonary dysfunction after surgery. Patients with larger defect size and earlier diagnosis time might be vulnerable to severe obstructive and mixed ventilatory disorders.

[Ann Surg Treat Res 2020;98(4):206-213]

Key Words: Airway obstruction, Infant, Tidal volume

INTRODUCTION

Congenital diaphragmatic hernia (CDH) is a group of conditions characterized by developmental defects of the diaphragm with an incidence of 1/2500–1/4000. The developmental defects are usually associated with pulmonary hypoplasia, abnormal pulmonary vascular development, and vasoreactivity. Both abnormal lung development in embryonic stage and improper treatment after birth could lead to morbidity after birth [1]. Most survivors showed reduced

forced vital capacity for several weeks after the repair surgery, and mild-to-moderate airway obstruction may still persist into adolescence and young adulthood [2,3].

Several studies have examined lung mechanics, functional residual capacity, and forced expiratory flows over the tidal range of breathing in CDH infants [2,4,5]. Advances in infant pulmonary function techniques have improved the assessment of flow rates in infants and toddlers by avoiding airway intubation. In our study, noninvasive tidal breathing pulmonary function testing was applied to evaluate respiratory

Received August 27, 2019, Revised October 21, 2019,
Accepted January 11, 2020

Corresponding Author: Jie Chen

Department of Pediatric Surgery, Xinhua Hospital Affiliated to Shanghai Jiao Tong University School of Medicine, No. 1665 of Kongjiang Road, Yangpu District, Shanghai 200092, China

Tel: +86-21-25078999, Fax: +86-21-25078999

E-mail: jiechen1974@163.com

ORCID: https://orcid.org/0000-0003-0266-0929

*Qianqian Wang and Quanhua Liu contributed equally to this study as co-first authors.

Copyright © 2020, the Korean Surgical Society

© Annals of Surgical Treatment and Research is an Open Access Journal. All articles are distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0/>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

function. The aim of this study was to investigate the risk factors affecting postoperative lung function and ameliorate perioperative therapy in CDH.

METHODS

Study design and ethics approval

A retrospective study for CDH was performed from November 2016 through September 2018. The registration number of this study is ChiCTR1800017627 on the Chinese clinical trial center (<http://www.chictr.org.cn>). The study was performed with strict accordance with Helsinki Declaration. The research was reviewed by the Ethics Committee of Xin Hua Hospital Affiliated to Shanghai Jiao Tong University School of Medicine, and the approval number is XHEC-D-2018-053.

Clinical data acquisition

The medical records of all candidates were reviewed to obtain clinical data, including sex, prenatal diagnosis of gestational age (GA), birth weight, surgical approach, sidedness, defect size, ventilation period, days of neonatal intensive care unit and combined anomalies, including liver herniation, atrial septal defect, patent ductus arteriosus, and pulmonary arterial hypertension. All cases of CDH infants were diagnosed prenatally and underwent repair operation postnatally. Lung function was assessed by tidal volume analysis 3 weeks after surgery when most patients were in stable situation.

Tidal breathing pulmonary function testing

Pulmonary function testing was performed in all enrolled CDH infants within 3 weeks after operation. The method of tidal breathing was used to determine pulmonary function and assist in ruling out severe expiratory airway obstruction. Measurements were performed according to European Respiratory Society/American Thoracic Society guidelines [6,7]. Informed consent was given by the children's legal guardians before testing. Flows and volumes were measured using a commercially available system (Master Screen Paed, JAEGGER, Friedberg, Germany) connected via a low-dead-space face mask secured tightly. In this test, we used tidal breathing flow-volume analysis to determine the pulmonary function, through mask-to-monitor, by applying the flow sensor during quiet breathing, thus analyzing the all respiratory function changes. Measured parameters include: tidal volume by per kilogram (V_T/kg), time to peak tidal expiratory flow as a proportion of expiratory time (TPTEF/TE), volume to peak expiratory flow as a proportion of exhaled volume (VPEF/VE), expiratory flow when 75%, 50%, and 25% of tidal volume remains in the lungs (TEF 75%, TEF 50%, TEF 25%). The normal value of V_T/kg was 6–10 mL/kg. The normal value of TPTEF/TE and VPEF/VE was 28–55 mL/sec.

All infants were insured under quiet breathing before testing. The state of body temperature and pressure saturated conditions of the testing environment were 37°C and 760 mmHg, respectively. Each lung function testing was repeated 3 times, and the average value was recorded as definitive result in the research. The possible correlations between clinical characteristics and postoperative lung function were analyzed to identify the risk factors correlated with postoperative lung function. All the infants with pulmonary dysfunction were treated with atomizing airway management to recover pulmonary function. The pulmonary function of the CDH patients was followed for 3 months after discharge from hospital.

Data analysis

All patients were classified into 4 groups according to pulmonary function: obstructive ventilatory disorder group (group 1), mixed ventilatory disorder group (group 2), restrictive ventilatory disorder group (group 3), and normal lung function group (group 4). We used statistical R programming to analyze the possible correlation between operative characteristics and lung function in all 4 groups. Demographic variables were analyzed to detect potential risk factors for pulmonary function and severity of disease. Patient characteristics and pulmonary function parameters were presented as mean \pm standard error of the mean. The data of all groups were analyzed by multivariate analysis of variance in IBM SPSS Statistics ver. 24.0 (IBM Co., Armonk, NY, USA). The group-wise changes in lung function and correlation with patient characteristics were determined using Pearson correlation. Statistical significance for all analysis was determined at the $P < 0.05$ level.

Table 1. Patient characteristics (n = 33)

Characteristic	Value
Sex, male:female	14:19
Operation age (day)	4.55 \pm 0.46
Birth age (wk)	38.62 \pm 0.43
Birth weight (g)	3,176 \pm 84.5
Sidedness, left:right	26:7
Diagnosis time (wk)	26.97 \pm 0.73
Defect size (cm ²)	16.48 \pm 1.11
Operation, open:thoracoscopic	16:17
Combined anomalies, yes:no ^{a)}	15:18
Patch use, yes:no	7:26
Ventilation period (day)	11.97 \pm 1.11
NICU time (day)	25.79 \pm 2.67

Values are presented as number or mean \pm standard error of the mean.

NICU, neonatal intensive care unit.

^{a)}Anomalies include liver herniation, and congenital heart disease, mainly atrial septal defect, patent ductus arteriosus and pulmonary arterial hypertension.

RESULTS

Patients and operative characteristics

Clinical statistics are shown in Table 1. Most infants received surgeries such as thoracoscopic surgeries and open surgeries within 2 weeks after birth. Amongst them, 14 were male and 19 were female; 26 were left-sided and 7 right-sided; 16 were open surgeries and 17 underwent thoracoscopic surgeries. The mean birth age was around 38 weeks.

Tidal breathing pulmonary function after operation

Pulmonary function tests were performed at 3 weeks after operation. The results are shown in Table 2. Lower numbers of V_T /kg were detected in 7 infants (21.2%). Decreased VPEF/

VE and TPTEF/TE levels were detected in 24 infants (72.7%). Among all the patients, 21 (64%) had obstructive ventilatory disorder (group 1), 3 (9%) had mixed ventilatory disorder (group 2), 4 (12%) had restrictive ventilatory disorder (group 3), and 5 (15%) showed normal lung function (group 4).

Prognostic factors affecting pulmonary function

Details are shown in Table 3. The results showed that GA at diagnosis was significantly increased in restrictive and normal groups ($P = 0.001$). The diagnosis age of children with obstructive ventilatory disorders and mixed ventilatory disorders was much younger compared with restrictive ventilatory disorders. Defect size in children with mixed ventilatory disorders and obstructive ventilatory disorders

Table 2. Pulmonary function classification analysis

Variable	Obstructive	Mixed	Restrictive	Normal	P-value
No. of patients	21 (64)	3 (9)	4 (12)	5 (15)	-
V_T /kg (mL/kg)	7.81 ± 0.26	4.27 ± 0.88	4.43 ± 0.69	6.78 ± 0.42	0.001*
TPTEF/TE (%)	20.10 ± 0.95	17.90 ± 0.92	40.38 ± 3.35	38.00 ± 5.37	0.001*
VPEF/VE (%)	22.56 ± 0.97	21.77 ± 0.38	37.78 ± 3.68	37.90 ± 5.36	0.001*
TEF 75% (mL/sec)	83.62 ± 6.24	51.33 ± 12.22	39.50 ± 5.55	67.80 ± 5.50	0.011*
TEF 50% (mL/sec)	70.81 ± 5.04	41.67 ± 11.46	40.00 ± 3.98	67.20 ± 5.27	0.020*
TEF 25% (mL/sec)	51.24 ± 4.08	28.67 ± 9.84	30.75 ± 2.87	57.20 ± 5.71	0.030*

Values are presented as number (%) or mean \pm standard error of the mean.

V_T /kg, tidal volume by per kilogram; TPTEF/TE, time to peak tidal expiratory flow as a proportion of expiratory time; VPEF/VE, volume to peak expiratory flow as a proportion of exhaled volume; TEF 75%, TEF 50%, TEF 25%, expiratory flow when 75%, 50% and 25% of tidal volume remains in the lungs.

Reference value: V_T /kg, 6–10 mL/kg; TPTEF/TE, 29%–35%; VPEF/VE, 29%–35%.

* $P < 0.05$, statistically significant differences.

Table 3. Multivariate analysis of variance of prognostic factors in pulmonary function

Variable	Different types of ventilation disorders				P-value
	Obstructive	Mixed	Restrictive	Normal	
Sex, male:female	9:12	2:1	2:2	1:4	0.605
Operation age (day)	4.6 ± 0.65	5.0 ± 1.15	4.3 ± 0.75	4.4 ± 1.21	0.986
Birth age (wk)	38.4 ± 0.65	39.1 ± 0.93	39.4 ± 0.10	38.8 ± 0.59	0.873
Birth weight (g)	$3,108 \pm 110$	$3,150 \pm 284$	$3,515 \pm 191$	$3,207 \pm 214$	0.516
Sidedness, left:right	17:4	2:1	2:2	4:1	0.302
Patch use, yes:no	5:16	1:2	0:4	1:4	0.642
Diagnosis of GA (wk)	25.00 ± 0.46	23.67 ± 1.45	31.75 ± 1.18	33.4 ± 1.17	0.001*
Defect size (cm ²)	16.81 ± 1.10	26.33 ± 4.49	15.50 ± 1.26	10.0 ± 1.27	0.001*
Combined anatomies ^{a)}	12	2	1	0	0.088
Surgery (O/T) ^{b)}	12:9	2:1	2:2	0:5	0.125
Ventilation period (day)	13.19 ± 1.62	14.0 ± 2.00	7.75 ± 0.63	9.00 ± 0.95	0.282
NICU time (day)	28.60 ± 3.98	24.7 ± 1.20	17.50 ± 2.40	21.2 ± 3.56	0.524

Values are presented as number or mean \pm standard error of the mean.

GA, gestational age; NICU, neonatal intensive care unit.

^{a)}Anomalies include liver herniation, and congenital heart disease, mainly atrial septal defect and pulmonary arterial hypertension.

^{b)}Open surgery (O); thoracoscopic surgery (T).

* $P < 0.05$, statistically significant differences.

was also significantly increased compared to that in restrictive ventilatory disorders ($P = 0.001$). The details of defect size and diagnosis time were shown in Fig. 1. However, no significant correlation was detected between days of intensive care unit (ICU), ventilation period, and lung dysfunction ($P = 0.282$). No significant differences were detected in gender ($P = 0.605$), surgical approach ($P = 0.125$), sidedness ($P = 0.302$), or combined anomalies ($P = 0.088$), either.

Interestingly, ventilation time and days of ICU appeared to be increased in both obstructive ventilatory disorder group and mixed ventilatory disorder group, while decreased in restrictive ventilatory disorder group. However, no significant differences were detected in the 4 groups.

Factors related to the severity of respiratory function disorders

Most CDH patients showed obstructive ventilatory disorders. The ventilatory disorders were classified into 3 groups based on the severity of obstruction (mild, moderate, severe). Normal pulmonary function and restrictive ventilatory disorders were amalgamated into mild ventilatory disorder group. Mixed and serious obstructive ventilatory disorders were included in severe ventilatory disorder group. The details were shown in Table 4. Mild or moderate ventilatory disorders were detected in 29 patients (87.9%). The ICU time was significantly longer in patients with severe obstructive ventilatory disorders compared with that in patients with mild-moderate ventilatory disorders ($P = 0.022$). The defect size was significantly different among the 3 groups ($P = 0.001$). Also, the diagnosis times of mixed ventilatory disorder and obstructive ventilatory disorder

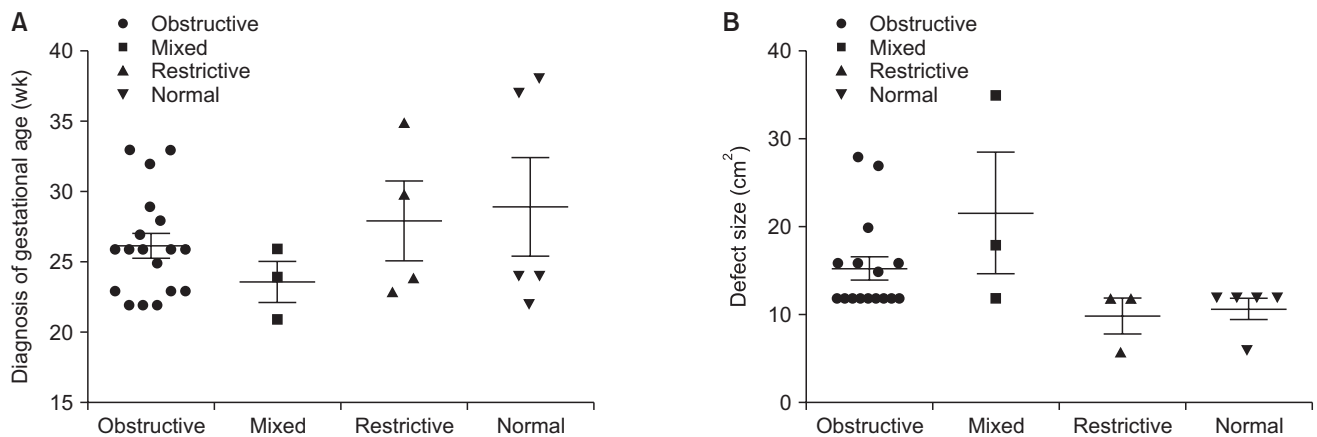


Fig. 1. Analysis of risk factors related to different ventilatory disorders. (A) Analysis of the diagnosis of gestational age related to different ventilatory disorders. (B) Analysis of defect size related to different ventilatory disorders.

Table 4. Factors related to the severity of airway obstruction

Variable	Severity of obstructive respiratory function disorders			P-value
	Mild (n = 16)	Moderate (n = 13)	Severe (n = 4)	
Ventilation time (day)	10.93 ± 1.31	11.92 ± 1.63	16.25 ± 5.71	0.342
NICU time (day)	21.88 ± 2.01	24.77 ± 3.85	44.75 ± 14.5	0.222
Defect size (cm ²)	13.56 ± 1.06	18.08 ± 1.78	23.00 ± 2.65	0.007*
Diagnosis of GA (wk)	29.88 ± 0.98	24.38 ± 0.64	23.75 ± 0.48	0.001*
Sex, male:female	7:9	5:8	2:2	0.910
Combined anomalies ^{a)}	7	6	2	0.549
Operation age (day)	4.31 ± 0.58	4.92 ± 0.90	4.25 ± 0.95	0.812
Birth age (wk)	39.13 ± 0.2	38.32 ± 1.03	37.55 ± 1.06	0.465
Birth weight (g)	3,332 ± 115	3,016 ± 128	3,070 ± 290	0.200
Sidedness, left:right	13:3	10:3	3:1	0.942
Surgery (O/T) ^{b)}	5:11	6:7	2:2	0.643

Values are presented as mean ± standard error of the mean or number.

^{a)}Anomalies include liver herniation, and congenital heart disease, mainly atrial septal defect and pulmonary arterial hypertension.

^{b)}Open surgery (O); thoracoscopic surgery (T).

* $P < 0.05$, statistically significant differences.

were found significantly earlier than in the other 2 groups ($P = 0.001$). Larger defect size was the influential factor in severe obstructive ventilatory disorders. Severe obstructive ventilatory disorders were combined with earlier diagnosis time in 4 patients (12.2%). Spearman correlation on Rank test showed that defect size was positively correlated with severity of obstruction ($\rho = 0.543$, $P < 0.05$); on the contrary, diagnosis time was negatively correlated with severity of obstruction ($\rho = -0.702$, $P < 0.05$), which was shown in Fig. 2.

Follow-up pulmonary function at 3 months after discharge from hospital

The follow-up pulmonary function results at 3 months after discharge from hospital were shown in Table 5. Amongst these patients, only 14 (42.1%) had obstructive ventilatory disorders, 4 (12.1%) had mixed ventilatory disorders, 5 (15.2%) had restrictive ventilatory disorders. 10 patients (30.3%) showed normal pulmonary function. Compared to postoperative

pulmonary function in Table 2, obstructive ventilatory disorder was relieved in 7 patients, and 6 more patients recovered from ventilatory disorders. Compared to postoperative pulmonary function, there was a trend of improvement in follow-up pulmonary function (Fig. 3). However, it was not statistically significant.

DISCUSSION

CDH is one of the most severe and life-threatening congenital diseases in the neonatal period characterized by unilateral or bilateral phrenic dysplasia, abdominal viscera herniation, and a major pathophysiological change. It usually combines a developmental defect of the diaphragm with pulmonary hypoplasia, abnormal pulmonary vascular development, and vasoreactivity. As a result, infants with CDH usually have variable degrees of pulmonary hypoplasia at birth. Functional and microstructural changes in lungs can persist into adulthood

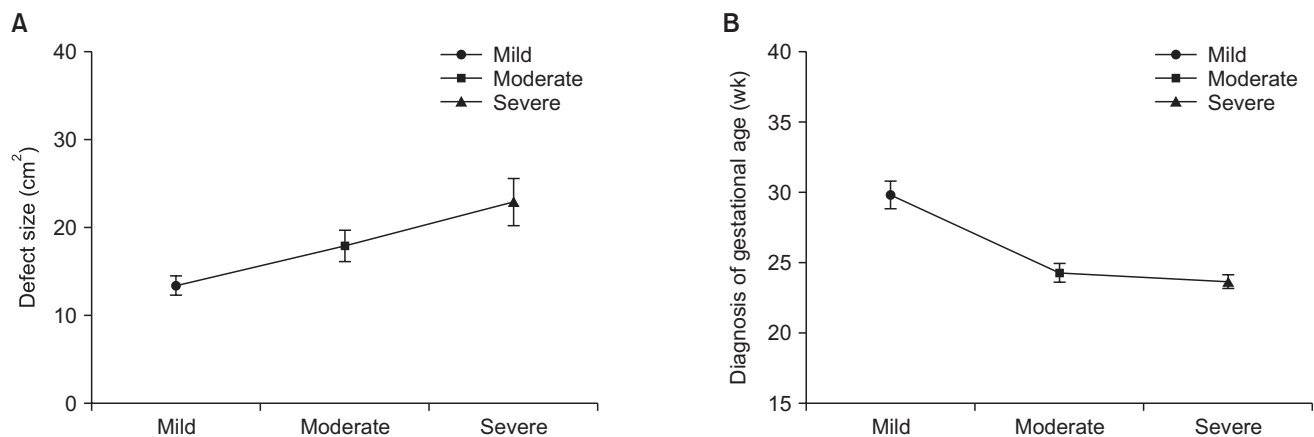


Fig. 2. Potential factors related to the severity of airway obstruction. (A) Defect size was positively related with the severity of airway obstruction ($\rho = 0.543$, $P < 0.05$). (B) Diagnosis of gestational age was negatively related with the severity of airway obstruction ($\rho = -0.702$, $P < 0.05$).

Table 5. Follow-up pulmonary function at 3 months after discharge

Variable	Obstructive	Mixed	Restrictive	Normal	P-value
No. of patients	14 (42.4)	4 (12.1)	5 (15.2)	10 (30.3)	-
V_T/kg (mL/kg)	7.79 ± 0.38	5.9 ± 1.13	5.16 ± 0.11	7.45 ± 0.34	0.002*
TPTEF/TE (%)	22.85 ± 1.34	22.25 ± 1.77	36.88 ± 2.58	38.02 ± 2.58	0.001*
VPEF/VE (%)	25.06 ± 0.88	22.53 ± 1.19	34.26 ± 2.34	37.05 ± 2.59	0.001*
TEF 75% (mL/sec)	83.92 ± 6.64	71.25 ± 5.11	40.8 ± 7.32	83.4 ± 8.06	0.006*
TEF 50% (mL/sec)	74.31 ± 5.24	62.75 ± 6.29	39.6 ± 5.90	86.10 ± 8.00	0.002*
TEF 25% (mL/sec)	54.86 ± 4.16	48.25 ± 5.65	31.00 ± 3.74	74.10 ± 7.79	0.001*

Values are presented as number (%) or mean \pm standard error of the mean.

V_T/kg , tidal volume by per kilogram; TPTEF/TE, time to peak tidal expiratory flow as a proportion of expiratory time; VPEF/VE, volume to peak expiratory flow as a proportion of exhaled volume; TEF 75%, TEF 50%, TEF 25%, expiratory flow when 75%, 50% and 25% of tidal volume remains in the lungs.

Reference value: V_T/kg , 6–10 mL/kg; TPTEF/TE, 29%–35%; VPEF/VE, 29%–35%.

* $P < 0.05$, statistically significant differences.

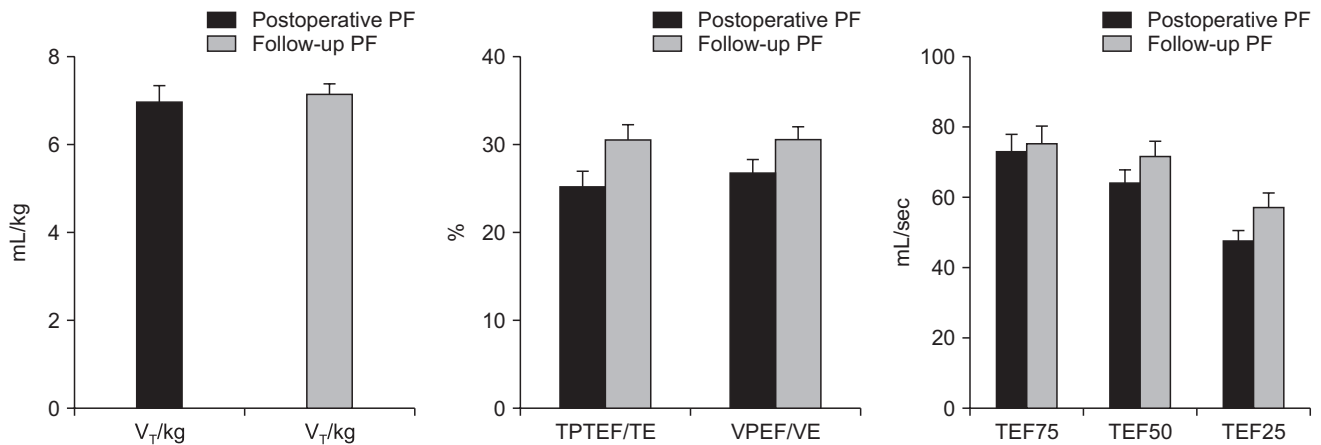


Fig. 3. Comparison between postoperative pulmonary function (PF) and follow-up PF at 3 months after discharge.

of most CDH patients [3]. Perioperative relative factors may affect pulmonary function in CDH infants. However, few studies of lung function over the first years of CDH infants have been reported. Herein, we analyzed the correlation between operative characteristics and pulmonary function parameters and found that defect size, diagnosis time, and ventilation time may be associated with postoperative pulmonary function.

The most significant result of our work is to verify risk factors of pulmonary dysfunction in CDH infants after surgery consistent with previous reports [2,3,5]. Generally, normal values of V_T/kg (tidal volume per kilogram) in children are 6–10 mL/kg, and decreased values indicate restrictive pulmonary disease. Obstructive disease, especially moderate or severe, tidal volume could decrease. TPTEF/TE is one of the main indicators reflecting airway obstruction. VPEF/VE is another parameter reflecting airway obstruction, and its changes are nearly consistent with the ratio of TPTEF/TE. The correlation between them reaches 90%. TPTEF/TE and VPEF/VE are the most sensitive indicators in neonatal respiratory function measurement. The heavier the obstruction is, the lower the ratio is. Normal values of TPTEF/TE and VPEF/VE in infants are 28%–55%. There is also a point of view that the expiratory flow velocity when exhaling different tidal volume, such as TEF 25%, TEF 50%, TEF 75%, is also a sensitive indicator for detection of small airway obstruction. Patients with obstructive ventilatory disorders generally showed lower TPTEF/TE and VPEF/VE values. Patients with restrictive ventilatory disorders showed lower V_T/kg values. In our research, larger defect size was found in patients with mixed ventilatory disorders and obstructive ventilatory disorders. Diagnosis time was also detected much earlier in patients with severe obstructive ventilatory disorders. Recently, a long-term study of CDH repair also demonstrated that larger defect size was associated with higher rates of early recurrence in CDH infants [8]. Other studies also demonstrated that prenatal diagnosis was correlated with poor outcome of

CDH infants [9,10].

The degree of pulmonary function impairment was also correlated both with markers of the initial degree of pulmonary hypoplasia and the duration of mechanical ventilation [2]. Meanwhile, children with CDH have primary disease with severe lung dysplasia, and such patients require ventilator-assisted respiration after birth, which may significantly aggravate lung injury of children and cause severe tissue damage. However, there is no significant difference between ventilation time and ventilatory disorders in our research.

The mechanisms of pulmonary hypoplasia in CDH infants have been controversial. It has been suggested that an early embryogenic alteration of the pulmonary vasculature and parenchymal (and diaphragmatic) development, as well as external compression from herniated intra-abdominal contents, stifle pulmonary vasculature/parenchymal development [11]; Guilbert et al. [12] showed that pulmonary hypoplasia and the intercept in lung development occur before the normal closure of the diaphragm and they are partially independent from the mechanical compression of the ascended abdominal organs. These procedures are amalgamated into the dual-hit hypothesis. The first hit, possibly genetic or environmental, probably occurs at an early phase before the formation of the diaphragm, which would arrest pulmonary development in both lungs. Lung compression by the abdominal organs at a later phase (second hit) would then worsen the ipsilateral hypoplasia, explaining why both lungs are usually hypoplastic, though the ipsilateral lung is further influenced than the contralateral [11]. Meanwhile, human postmortem studies confirmed that CDH patients had severe dysplasia in the ipsilateral lung, and only mild dysplasia of the contralateral lung [13]. Researches in newborns with CDH had shown that CDH was composed of fewer alveoli and reduced gas-exchange surface area, significantly thickened alveolar walls, arteries of a smaller diameter than normal, larger alveoli, and increased interstitial tissue in alveolar development

[12]. It indicated that pulmonary hypoplasia and dysfunction existed in CDH patients, and these patients might be vulnerable to poor pulmonary function outcome.

Our pulmonary function test results confirmed that different types of ventilation disorder existed in CDH patients after surgery. Children with mild diaphragmatic hernia may be prone to restrictive ventilatory disorder postoperatively; while moderate or severe diaphragmatic hernia may be prone to obstructive lung ventilation or mixed disorders. Meanwhile, long-term follow-up study has identified pulmonary function disorder existed in most patients with CDH after surgery [3,14,15]. Earlier diagnosis time was also associated with more severe ventilatory disorder. It is important to note that antenatal care is of great significance for observing some severe diseases. Therefore, postoperative pulmonary function rehabilitation is of great significance in improving pulmonary function abnormality after diaphragmatic hernia.

Our study preliminarily analyzed risk factors associated with postoperative outcomes of pulmonary function in infants with CDH. Our results showed that defect size, ventilation time, and prenatal diagnosis of GA were risk factors correlating with postoperative pulmonary function of CDH infants. To our knowledge, this is the first report of risk factors affecting postoperative pulmonary function in CDH infants. Long-term follow-up study should further explore the pulmonary function modification in CDH patients at several years after surgery, and whether it might enhance medicine intervention in pulmonary rehabilitation. One limitation of our work is insufficient number of cases; however, the small number of infants with CDH managed in our center further highlights the attention needed

for the rehabilitation of CDH patients' respiratory function after operation. Also, a collaborative multidisciplinary research ought to be established to facilitate pulmonary function rehabilitation and improve clinical outcomes of CDH patients.

ACKNOWLEDGEMENTS

Fund/Grant Support

This study was supported by grants from Science and Technology Planning Project of Jiaxing City, No. 2018AY32019 and the Foundation of Xinhua Hospital, No. 18YJ01-XH2406.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

ORCID iD

Qianqian Wang: <https://orcid.org/0000-0003-1962-289X>

Quanhua Liu: <https://orcid.org/0000-0002-9581-1265>

Jun Wang: <https://orcid.org/0000-0003-3455-0302>

Jie Chen: <https://orcid.org/0000-0003-0266-0929>

Author Contribution

Conceptualization: JC

Formal Analysis: QL

Investigation: QW

Methodology: QL, JZ

Project Administration: JW

Writing – Original Draft: QW

Writing – Review & Editing: JC

REFERENCES

1. Ameis D, Khoshgoo N, Keijzer R. Abnormal lung development in congenital diaphragmatic hernia. *Semin Pediatr Surg* 2017;26:123-8.
2. Panitch HB, Weiner DJ, Feng R, Perez MR, Healy F, McDonough JM, et al. Lung function over the first 3 years of life in children with congenital diaphragmatic hernia. *Pediatr Pulmonol* 2015;50:896-907.
3. Spoel M, van der Cammen-van Zijp MH, Hop WC, Tibboel D, de Jongste JC, Ijsselstijn H. Lung function in young adults with congenital diaphragmatic hernia; a longitudinal evaluation. *Pediatr Pulmonol* 2013;48:130-7.
4. American Thoracic Society; European Respiratory Society. ATS/ERS statement: raised volume forced expirations in infants: guidelines for current practice. *Am J Respir Crit Care Med* 2005;172:1463-71.
5. Spoel M, van den Hout L, Gischler SJ, Hop WC, Reiss I, Tibboel D, et al. Prospective longitudinal evaluation of lung function during the first year of life after repair of congenital diaphragmatic hernia. *Pediatr Crit Care Med* 2012;13:e133-9.
6. Bates JH, Schmalisch G, Filbrun D, Stocks J. Tidal breath analysis for infant pulmonary function testing. *ERS/ATS Task Force on Standards for Infant Respiratory Function Testing. European Respiratory Society/American Thoracic Society. Eur Respir J* 2000;16:1180-92.
7. Frey U, Stocks J, Coates A, Sly P, Bates J. Specifications for equipment used for infant pulmonary function testing. *ERS/ATS Task Force on Standards for Infant Respiratory Function Testing. European Respiratory Society/American Thoracic Society. Eur Respir J* 2000;16:731-40.
8. Putnam LR, Gupta V, Tsao K, Davis CF, Lally PA, Lally KP, et al. Factors associated with early recurrence after congenital

- diaphragmatic hernia repair. *J Pediatr Surg* 2017;52:928-32.
9. Brindle ME, Cook EF, Tibboel D, Lally PA, Lally KP; Congenital Diaphragmatic Hernia Study Group. A clinical prediction rule for the severity of congenital diaphragmatic hernias in newborns. *Pediatrics* 2014;134:e413-9.
10. Long AM, Bunch KJ, Knight M, Kurinczuk JJ, Losty PD; BAPS-CASS. Early population-based outcomes of infants born with congenital diaphragmatic hernia. *Arch Dis Child Fetal Neonatal Ed* 2018;103:F517-22.
11. Keijzer R, Liu J, Deimling J, Tibboel D, Post M. Dual-hit hypothesis explains pulmonary hypoplasia in the nitrofen model of congenital diaphragmatic hernia. *Am J Pathol* 2000;156:1299-306.
12. Guilbert TW, Gebb SA, Shannon JM. Lung hypoplasia in the nitrofen model of congenital diaphragmatic hernia occurs early in development. *Am J Physiol Lung Cell Mol Physiol* 2000;279:L1159-71.
13. Kitagawa M, Hislop A, Boyden EA, Reid L. Lung hypoplasia in congenital diaphragmatic hernia. A quantitative study of airway, artery, and alveolar development. *Br J Surg* 1971;58:342-6.
14. Stefanutti G, Filippone M, Tommasoni N, Midrio P, Zucchetta P, Moreolo GS, et al. Cardiopulmonary anatomy and function in long-term survivors of mild to moderate congenital diaphragmatic hernia. *J Pediatr Surg* 2004;39:526-31.
15. Choi SY, Lee HW, Hong J. Late lung function in the repaired congenital diaphragmatic hernia. *J Korean Surg Soc* 2010;79:143-8.