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# Severe congenital heart defects: incidence, causes and time trends of preoperative mortality in Norway

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## ABSTRACT

**Background and aims** Severe congenital heart defects (CHDs) still represent one of the main causes of infant death. The risk factors associated with cardiac surgery and postoperative mortality are well known. We aimed to describe the rates, causes and time trends of mortality before surgery—including termination of pregnancies and palliative care—in fetuses and children below 2 years of age with severe CHDs.

**Methods and results** Data concerning all 789 345 pregnancies in Norway from 2004 to 2016 were retrieved from the Medical Birth Registry of Norway, the Oslo University Hospital's Clinical Registry for Congenital Heart Defects, the Norwegian Cause of Death Registry, the National Registry, Statistics Norway, autopsy reports and medical records. When including termination of pregnancy and stillbirths, the number of fetuses and children with severe CHDs that did not reach the age of 2 years was 31%. Mortality among the 2359 live-born infants with severe CHDs was 10%, of whom 58% died before surgery. Of the preoperative deaths, 81% died in a palliative care setting, and comorbidity and univentricular CHDs were common among these infants. Together, palliative care and termination of pregnancy accounted for 86% of deaths in cases of severe CHDs, and this proportion increased during the study period (annual percent changes 1.3, 95% CI 0.4 to 2.1,  $p < 0.001$ ), mainly due to an increased termination rate.

**Conclusions** Termination of pregnancy accounted for the majority of the deaths in fetuses and children with severe CHDs. Among live-born children, most preoperative deaths occurred in a palliative care setting and were strongly related to comorbidities and/or univentricular hearts.

## INTRODUCTION

Congenital heart defects (CHDs) are the most common birth defects, affecting approximately 1% of newborns.<sup>1–3</sup> The prevalence of live-born infants with severe CHDs has decreased, probably due to increased rates of prenatal detection and termination of pregnancy.<sup>1,4–7</sup> CHDs still represent one of the main causes of death in neonates and young children, despite major advancements in treatment during the last decades.<sup>8–10</sup> Several studies report high rates of comorbidity and mortality among infants with severe CHDs.<sup>2,3,11</sup> The mortality rate and risk factors (eg, CHD type, procedure, age, comorbidity and ethnicity) for early postoperative death and adverse outcomes in the year after discharge in children with severe CHDs have been extensively described.<sup>12–17</sup> However, population-based studies

## What is already known on this topic?

- ▶ The peroperative and postoperative mortality in children with congenital heart defects (CHDs) has decreased in recent decades.
- ▶ CHDs still represent one of the main causes of death in infants and young children.

## What this study adds?

- ▶ The majority of the preoperative deaths among live-born children with severe CHDs occurred in a palliative care setting.
- ▶ Unexpected deaths before surgery were rare.
- ▶ Terminated pregnancies and palliative care accounted for 86% of all deaths before 2 years of age.

investigating long-term outcomes of children born with severe CHDs are sparse.<sup>18</sup>

Moreover, there is limited knowledge on the rates and causes of preoperative mortality in infants with severe CHDs. Some of these infants die without surgical treatment in a palliative care setting. It is unknown whether the increased rate of terminated pregnancies has reduced the proportion of palliative care among infants with severe CHDs.

Norwegian national health registries provide an opportunity to conduct a nationwide population-based study of early mortality in children with severe CHDs.

The aims of the present study were to describe the rates, causes and time trends of mortality before surgery—including termination of pregnancies and palliative care—in fetuses and children below 2 years of age with severe CHDs during the period 2004–2016 in Norway.

## METHODS

### Classification of CHDs

CHDs are defined as structural abnormalities of the heart or intrathoracic vessels with functional or potentially functional significance.<sup>19</sup> Children with isolated cardiomyopathies, primary arrhythmias and primary pulmonary hypertension were not included in the study. Heterotaxia, transposition of the great arteries, tetralogy of Fallot, double-outlet right ventricle, truncus arteriosus, aortic arch hypoplasia, coarctation of the aorta, atrioventricular septal defects, anomalous pulmonary venous return,



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hypoplastic left heart syndrome, Shone's syndrome, aortic valve stenosis, pulmonary valve atresia, tricuspid valve atresia, Ebstein's anomaly, anomalies of the coronary arteries and other complex defects were classified as severe CHDs and included in the study.<sup>20 21</sup> Ventricular septal defects (VSDs) were classified as severe if treated by a surgical or catheter-based procedure or if the child with a non-restrictive VSD died in elective palliative care. Non-severe CHDs (atrial septal defects, restrictive VSDs, minor valve malformation, pulmonary valve stenosis and isolated patent ductus arteriosus) were not included. All individuals were assigned to only one CHD diagnosis according to the International Paediatric and Congenital Cardiac Code by two experienced paediatric cardiologists (HH and GW).<sup>22</sup>

### Data sources

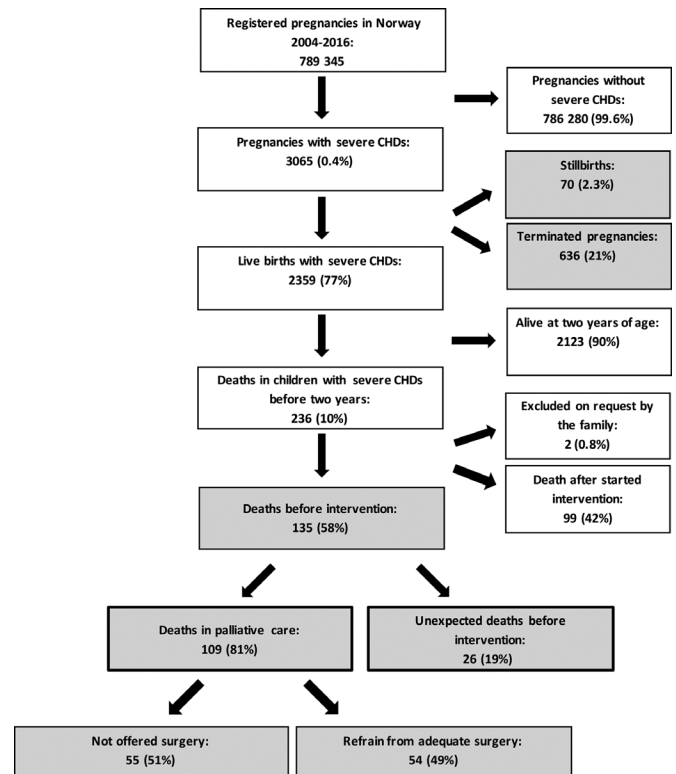
Information about all pregnancies from 1 January 2004 to 31 December 2016 was obtained from the Medical Birth Registry of Norway (MBRN).<sup>23</sup> Registration of pregnancy beyond 12 weeks is statutory in Norway. The MBRN contains information on all live births, stillbirths and terminations of pregnancies due to congenital anomalies after 12 weeks' gestation. Information on live-born children with severe CHDs was retrieved from the Oslo University Hospital's Clinical Registry for Congenital Heart Defects, which contains information on all children with CHDs who have been examined or treated at Oslo University Hospital. Oslo University Hospital is the national centre for invasive treatment of children with CHDs in Norway. This registry was linked to the Norwegian Cause of Death Registry and the National Registry, which contains death certificate information about the date and causes of death. Moreover, we reviewed the medical records and the autopsy reports for all live-born infants with severe CHDs who died during the study period. The overall number of deaths in children under the age of 2 years in Norway during the study period was retrieved from Statistics Norway.<sup>24</sup>

### Definitions

*Palliative care* was defined as supportive care with refrainment from cardiac surgery and other active life-prolonging treatment.<sup>25 26</sup> We differentiated between children who were *not offered surgery* and *refrainment from cardiac surgery*. *Not offered surgery* was defined as children in whom there was no feasible surgical option (such as univentricular hearts with severe pulmonary vein stenosis or pulmonary atresia with VSD and lacking ordinary pulmonary arteries). *Refrainment from cardiac surgery* was defined as palliative care chosen after dialogue with the parents and caretakers based on a total evaluation of the baby's condition and expected quality of life (eg, severe CHDs with comorbidities such as multiple malformations). *Unexpected death* was defined as preoperative death occurring in children with planned active treatment. *Comorbidity* was defined as a condition with anticipated effect on the child's health and/or development and included Trisomy 13, 18 and 21, other genetic syndromes, multiple malformations, intraventricular haemorrhage grade 3–4 or ischaemic brain injury,<sup>27</sup> organ failure (eg, severe kidney failure, liver failure and necrotising enterocolitis) and extreme prematurity (<28 weeks' gestation). *The number of fetuses with severe CHDs* was defined as the sum of terminated pregnancies, stillbirths and live births with severe CHDs

### Norwegian legislation and the process of counselling

In Norway, women have the right to terminate pregnancy on demand within the first 12 weeks of gestation. Thereafter, women can apply to a medical assessment board for termination



**Figure 1** Flow chart of the study population. CHDs, congenital heart defects.

until week 22. Terminations for medical reasons between weeks 12 and 22 will usually be approved if the child has poor prognosis and/or comorbidities with expected major impact on quality of life.

The routine ultrasound examination in Norway is offered to all pregnant women between weeks 18 and 20. Women carrying a fetus with a suspected CHD are referred to specialised fetal medicine centres at the university hospitals. Parental counselling, by a fetal medicine specialist and a paediatric cardiologist, is individualised, based on the diagnosed CHD and presence of comorbidities, and non-directive.

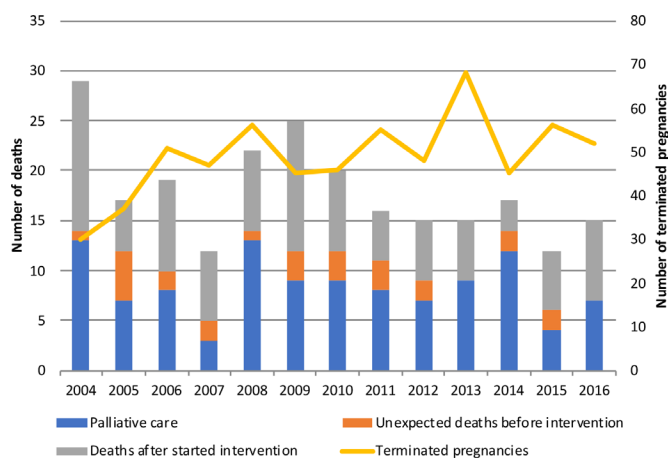
### Statistical analysis

Continuous variables are presented as median (range) and categorical variables as number and proportion. Time trends were analysed (log-linear model) using the Joinpoint Regression Program (version 4.0; SEER software, National Cancer Institute, USA) and are presented as the expected annual percent changes (APC) with a 95% CI. Other statistical analyses were performed using STATA/SE V.15.0.

### RESULTS

A total of 789 345 pregnancies were recorded in Norway from 2004 to 2016.<sup>23</sup> Severe CHDs were detected in 3065 (0.39%) cases, either prenatally or postnatally. Of these, 2359 (77.0%) children were live born (figure 1). The proportion of live-born children with severe CHDs decreased significantly (APC  $-1.9$ , 95% CI  $-2.6$  to  $1.1$ ,  $p < 0.001$ ). A total of 942 (31%) fetuses and children with severe CHDs were either stillborn ( $n = 70$ ), terminated ( $n = 636$ ) or died before 2 years of age ( $n = 236$ ). The rate of fetuses and children with severe CHDs that did not reach the age of 2 years increased significantly (APC  $1.7$ , 95% CI  $0.1$  to  $3.3$ ,  $p < 0.001$ ) during the study period. The annual number

## Original research



**Figure 2** Terminated pregnancies, palliative care and unexpected deaths before 2 years of age in children with severe congenital heart defects in Norway 2004–2016.

of deaths are presented in figure 2. Two live-born infants with severe CHDs were excluded from detailed analysis on request by their parents.

### Termination of pregnancy

Termination of pregnancy was chosen in 636 (21%) of the 3065 cases with severe CHDs. The proportion of termination of pregnancy increased from less than 15% in 2004 to approximately 25% in 2016 (APC 4, 95% CI 1.7 to 6.4,  $p < 0.001$ ). This accounted for 68% (636 of 942) of all deaths before 2 years of age in cases with severe CHDs.

### Mortality in live-born children

A total of 236 (10.0%) of 2359 live-born children with severe CHDs died before 2 years of age. In comparison, the mortality rate for all live-born children below 2 years of age in Norway was 0.3% ( $n = 2298$ ).<sup>24</sup> Infants with severe CHDs accounted for 10.3% of these deaths, the proportion was stable throughout the period (APC  $-0.5$ , 95% CI  $-4.1$  to  $3.1$ ,  $p = 0.7$ ), and most of the deaths in live-born infants with severe CHDs occurred during the first year of life ( $n = 219$ , 93%). Of the 234 children who died before 2 years of age, 135 (58%) died before surgery. The proportion of preoperative deaths was stable throughout the study period (APC  $2.0$ , 95% CI  $-1.4$  to  $5.4$ ,  $p = 0.2$ ).

### Unexpected preoperative deaths

Unexpected deaths before surgery occurred in 26 (1.1%) of 2359 live-born infants with severe CHDs, corresponding to 11% of the 234 live-born children with severe CHDs who died before 2 years of age and were included in the study. These infants comprised a heterogeneous group including 17 different CHDs. They have not been further analysed in this study.

### Palliative care

Of the 234 live-born children with severe CHDs who died before 2 years of age and were included in the study, 109 (47%) died in a palliative care setting. This accounts for 81% (109 of 135) of the preoperative deaths. Among these, 55 (50%) infants were *not offered surgery* (eg, hypoplastic left heart syndrome and additional pulmonary vein stenosis or severe valve-dysplasia ( $n = 9$ ), pulmonary atresia with coronary abnormalities or lacking native pulmonary arteries ( $n = 5$ ) and unbalanced atrioventricular septal defects with severe valve dysplasia or peripheral pulmonary

stenoses ( $n = 5$ )). *Refrainment from cardiac surgery* was chosen in 54 (50%) infants. The proportion of palliative care offered to live-born infants with severe CHDs was stable throughout the study period (APC  $-0.6$ , 95% CI  $-6.2$  to  $5.4$ ,  $p = 0.8$ ). The cardiac diagnoses in infants offered palliative care are listed in figure 3, and clinical characteristics are presented in table 1.

The median survival time in infants offered palliative care was 9 days with a range from 0 to 618 days. After the decision to provide solely palliative care, 20 (18%) of 109 infants were discharged and died at home.

### Comorbidity

A total of 73 (67%) infants offered palliative care had additional morbidities. Of these, five (7%) had trisomy 21, 20 (27%) had trisomy 13 or 18 and 10 (14%) had other genetic syndromes. Additional malformations were diagnosed in 29 (40%) children. Four (5%) infants had intraventricular haemorrhage grade 3–4 or ischaemic brain injury, whereas three (4%) infants had other organ failure as the main non-cardiac diagnosis. Five (7%) infants were born extremely preterm, of whom two had Down syndrome and one DiGeorge syndrome. Among the 14 infants with *refrainment from cardiac surgery* and no comorbidity, the vast majority ( $n = 12$ , 86%) had hypoplastic left heart syndrome.

### Terminated pregnancies and palliative care

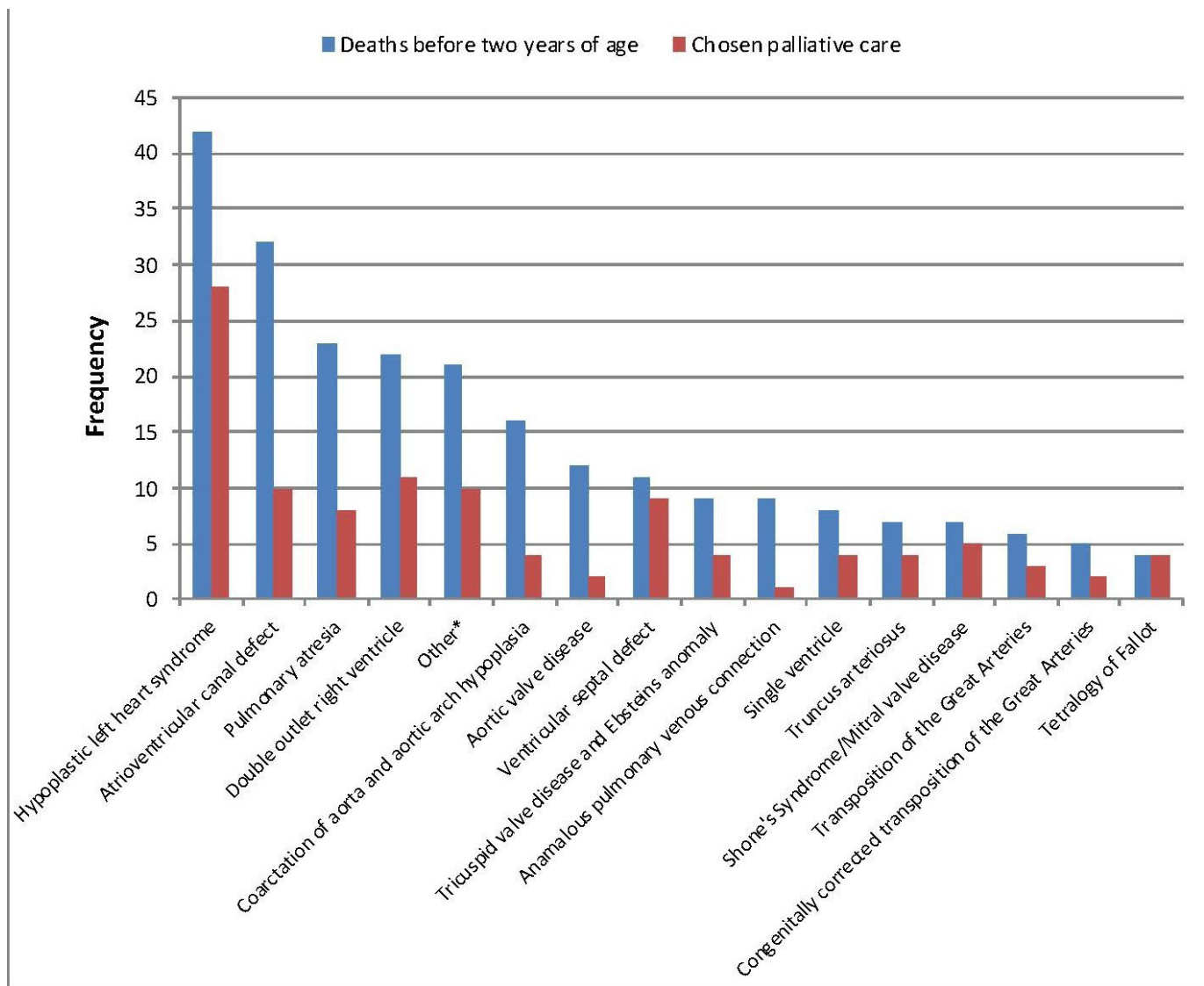
Together, terminated pregnancies ( $n = 636$ ) and infants offered palliative care ( $n = 109$ ) accounted for 86% ( $n = 745$ ) of all deaths in cases of severe CHDs ( $n = 870$ ), and this proportion increased significantly during the study period (APC  $1.3$ , 95% CI  $0.4$  to  $2.1$ ,  $p < 0.001$ ).

## DISCUSSION

In this nationwide study, the number of fetuses and children with severe CHDs that did not reach the age of 2 years was 31%. Termination of pregnancy accounted for about two-thirds of these cases, and the proportion of terminated pregnancies increased markedly from 2004 to 2016. Mortality among the live-born children with severe CHDs was 10%, of whom 58% died before surgery. More than 80% of the preoperative deaths occurred in a palliative care setting, and the proportion of palliative care was stable throughout the period. Comorbidities and univentricular CHDs were common among these infants. Approximately 10% of deceased live-born children with severe CHDs died unexpectedly before surgery.

Termination of about half of pregnancies with prenatally diagnosed severe CHDs has been recently reported.<sup>28</sup> We previously reported that the prenatal detection rate in Norway in 2016 was 58%.<sup>29</sup> In the present study, we were not able to include the prenatal detection rate for all fetuses with severe CHDs, but we registered an increased rate of terminated pregnancies during the study period. In fact, at the end of the study period, approximately one quarter of all pregnancies with severe CHDs were terminated. This finding might indicate that improved prenatal detection of CHDs may result in increasing termination of fetuses with severe CHDs.

The impact of prenatal detection of CHDs on neonatal outcome is unclear. Some studies reported increased survival in live-born infants with severe CHDs, partly as a consequence of the more frequent use of termination of pregnancy.<sup>4 30</sup> In contrast, a recent study reported no difference in mortality in infants with and without prenatal diagnosis of severe CHDs and questioned the impact of prenatal diagnosis on infant survival. However, the two groups (with and without prenatal diagnosis) differed in the types



**Figure 3** Diagnoses in deaths before 2 years of age in live-born children with severe congenital heart defects. Norway 2004–2016. \*Heterotaxia, pulmonary venous stenosis, coronary artery anomaly and other complex defects.

of heart diagnoses.<sup>28</sup> In the present study, the 2-year mortality rate among live-born infants was 10% and almost 50% died in palliative care. This proportion was stable despite increased termination rates. This result may illustrate how challenging it is to predict the severity of CHD and associated comorbidities during the second trimester and before the legal limit for termination of pregnancy. Moreover, the decision for the termination of pregnancy is ethically challenging. It is influenced by personal beliefs and probably involves more than just considerations around severity and survival. Genetic syndromes and coexisting malformations were major determinants when *refrainment from cardiac surgery* was chosen. Other comorbidities, such as severe intracerebral haemorrhage and extreme prematurity, were also important. The presence of additional cardiac defects was a determining factor when *not offering surgery*, particularly in children with univentricular heart defects. Several studies present a trend towards more cardiac surgery in infants with complex CHDs and severe comorbidity, such as trisomy 18 and 13.<sup>31 32</sup> In the study by Vincenti *et al*<sup>28</sup>, all infants received active treatment regardless of the severity of CHDs or comorbidities, but the mortality rate was similar to the mortality in our study. This may indicate that a palliative approach

mainly changes the manner of death without increasing the rate of mortality. Palliative care of the newborn is a challenging and developing arena,<sup>25 33 34</sup> and this study confirms that it is an important part of neonatal cardiology practice. An active approach involving heart surgery and intensive care is associated with painful procedures, stress and less parental involvement,<sup>35</sup> and a comprehensive palliative approach may be preferable in selected cases.

The main strength of this study is the complete national cohort of severe CHDs, including both live-born children and terminated pregnancies. The use of national registries and access to medical records ensure the inclusion of nearly all cases of severe CHDs with detailed medical information.

However, some limitations have to be acknowledged. The number of deaths was limited, and the time trends should be interpreted with caution. We cannot exclude the possibility that some patients may have died before surgery without being recorded in the registries. Furthermore, we used arbitrary definitions of comorbidity, and the classification was based on our assessment. The same applies for the causes of *refrainment from cardiac surgery*. The distinction between '*not offered surgery*' and '*refrainment from cardiac surgery*' was not clear in all cases. We

**Table 1** Characteristics of children with severe congenital heart defects and death in a setting of palliative care, Norway 2004–2016

IPCCC diagnoses	n	Prenatal detection, n (%)	Refrainment from adequate cardiac surgery, n (%)	Not offered cardiac surgery, n (%)	Comorbidity, n (%)	Prematurity,* n (%)	Median time to death (days (range))
Hypoplastic left heart syndrome	28	12 (43)	14 (50)	14 (50)	6 (21)	6 (21)	8 (1–154)
Double outlet right ventricle	11	7 (64)	6 (55)	5 (45)	9 (82)	7 (64)	5 (0–110)
Atrioventricular septal defect	10	5 (50)	5 (50)	5 (50)	8 (80)	5 (50)	5 (0–618)
Ventricular septal defect	9	2 (22)	9 (100)	0 (0)	9 (100)	2 (22)	13 (2–69)
Pulmonary atresia	8	4 (50)	1 (13)	7 (88)	6 (75)	1 (13)	10 (4–59)
Coarctation of aorta and aortic arch hypoplasia	4	1 (25)	4 (100)	0 (0)	4 (100)	2 (50)	7 (5–51)
Tricuspid valve disease and Ebstein's anomaly	4	4 (100)	0 (0)	4 (100)	4 (100)	2 (50)	5 (2–21)
Single ventricle	4	2 (50)	3 (75)	1 (25)	3 (75)	1 (25)	290 (11–447)
Truncus arteriosus	4	2 (50)	3 (75)	1 (25)	4 (100)	2 (50)	15 (12–36)
Tetralogy of Fallot	4	0 (0)	3 (75)	1 (25)	4 (100)	1 (25)	144 (12–282)
Transposition of the great arteries	3	1 (33)	1 (33)	2 (67)	2 (67)	1 (33)	8 (8–29)
Mitral valve disease	3	1 (33)	0 (0)	3 (100)	2 (67)	0 (0)	3 (0–3)
Other†	17	11 (65)	5 (29)	12 (71)	12 (71)	3 (18)	16 (1–416)
Total	109	52 (48)	54 (50)	55 (50)	73 (67)	33 (30)	9 (0–618)

\*Gestational age <37 weeks.

†Heterotaxia, aortic valve disease, anomalous pulmonary venous connection, congenitally corrected transposition of the great arteries, Shone's syndrome, pulmonary venous stenosis, coronary artery anomaly and other complex defects.  
IPCCC, International Paediatric and Congenital Cardiac Code.

have no information about comorbidity and specific diagnosis of the heart defect in the terminated pregnancies with severe CHDs. Finally, we emphasise that clinical practice varies between countries depending on factors like the healthcare system, socio-economic conditions, culture and religion. This study reflects the situation in Norway, and the findings are not automatically transferrable to other settings.

In conclusion, we found lower mortality than previously reported among live-born infants with severe CHDs. Moreover, the data revealed an increasing number of fetuses and infants not offered active life prolonging treatment.

We emphasise the importance of optimised antenatal service and decision-making processes, including the indications for surgery versus palliative care in live-born children. In our opinion, providing palliative care in the most complex cases may in fact improve quality of life for the affected infants and their families, without necessarily increasing the mortality rate.

**Contributors** GW contributed to the study concept and design, data collection and interpretation, literature review and revised and approved the manuscript. JJ contributed to the study concept and design, data interpretation, literature review and revised and approved the manuscript. VS contributed to the study concept, literature review and revised and approved the manuscript. GD and AER contributed to the study concept and revised and approved the manuscript. HH contributed to the study concept and design, data interpretation, literature review and revised and approved the manuscript.

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**Competing interests** None declared.

**Patient consent for publication** Not required.

**Ethics approval** The study was approved by the Regional Committee for Medical and Health Research Ethics South East Norway (REK nr. 2010/392).

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**Data availability statement** Data are stored in 'eReg Versjon 2.4.0 OUS-HF' and are available upon reasonable request. They are not to be reused without approval from the Regional Committee for Medical and Health Research Ethics South East Norway.

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