

International Journal of Pediatric Research

RESEARCH ARTICLE

Factors Associated with Neonatal Mortality at Kyeshero Hospital, Goma, North Kivu, DRC

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Abstract

Background: The neonatal mortality rate is still a significant public health problem in Sub-Saharan African countries and an indicator of the quality of prenatal and perinatal care. This study aimed to determine the neonatal intra-hospital neonatal mortality rate and the maternal and neonatal risk factors associated with neonatal mortality in Kyeshero Hospital.

Methods: This research was a cross-sectional analytical study, which took place over a period of 12 months, from January 1 to December 31, 2021, at Kyeshero Hospital in Goma, Eastern Democratic Republic of Congo.

Results: The Intra-hospital neonatal mortality rate was 17.0%. The multivariate analysis showed various risk factors associated with newborns that low birth weight < 2500 gr ORa = 4.11 (2.17-7.77), Presence of cyanosis ORa = 3.88 (1.97-7.65), absence of reactivity ORa = 20.07 (8.07-49.55), and neonatal infection ORa = 4.78 (2.47-9.22) are real predictor factors of neonatal mortality.

Conclusion: This study noted that neonatal mortality was influenced by neonatal factors. Proper care of the newborn should improve neonatal vital prognosis. The study of preventable causes of death and factors that lead to early neonatal death is fundamental to reducing the infant mortality rate in the world.

Keywords

Neonatal mortality, Risk factor, Goma

Introduction

The first 28 days of life is an extremely likely time for a newborn to make many physiological adjustments necessary for life outside the womb, making this the most dangerous time of life [1]. Newborns in need of serious medical attention are admitted to the neonatal intensive care unit, which incorporates innovative technology and trained staff to effectively provide specialized care to newborns [2]. Neonatal mortality is an indicator of a country's demographic, biological, and socioeconomic conditions as well as the health system, public health, and population growth rate [3]. Although the neonatal mortality rate has decreased worldwide, it is slower than the under-5 mortality rate and remains unacceptably high at 37 per 1000 live births [4]. Globally, approximately 7,000 babies die every day; Most of these deaths occur within the first week and nearly 2.6 million babies die within the first month of life [5].



Citation: Endanda ZE, Tshivwadi TA, Tongota NJ, Mulangu MA, Imani MP, et al. (2023) Factors Associated with Neonatal Mortality at Kyeshero Hospital, Goma, North Kivu, DRC. Int J Pediatr Res 9:115. doi.org/10.23937/2469-5769/1510115

Accepted: June 07, 2023: Published: June 09, 2023

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Approximately 78% of these neonatal deaths occur in developing countries, particularly in South Asia and sub-Saharan Africa [6], which account for 39% of all neonatal deaths worldwide and are among the top 10 countries with the highest neonatal mortality rate [7]. In Sub-Saharan Africa alone 1.2 million newborns died every year equivalent to 13,000 deaths per day or almost nine deaths every minute, 50% of neonatal deaths occur in just five countries: Nigeria, Democratic, Republic of Congo, Ethiopia, Tanzania, and Uganda [8]. The majority of neonatal deaths in this part of Africa occur at home, in rural communities, among the poor and poorest, less educated, and in war regions [9-11]. Also causes and risk factors of neonatal death vary by country with the availability in relation to the quality of health care [12]. The democratic Republic of Congo ranks second African Africa countries with higher neonatal rate more than 28 per 1000 live births [13].

This situation shows that neonatal mortality remains a public health problem in many countries. To accelerate the achievement of Millennium Development Goals (MDGs) 4 and 5, several countries, including the DRC, have implemented strategies to substantially reduce neonatal mortality. To target actions, an understanding of the causal factors of neonatal mortality is required in the context of the deaths occurring. This study aims to determine the neonatal intra-hospital neonatal mortality rate and the maternal and neonatal risk factors associated with neonatal mortality in Kyeshero Hospital. The hospital is located in Goma, the North Kivu province of the Eastern Democratic Republic of Congo.

Methods

Study design, setting, and period

We conducted a cross-sectional analytical study, which took place over a period of 12 months, from January 1 to December 31, 2021, at Kyeshero Hospital in Goma, Eastern Democratic Republic of Congo, which is one of the reference health facilities child health care in this region.

Study population, sample size, and inclusion criteria

All neonates who were admitted to the neonatal intensive care unit and registered as dead and alive during the first 28 days of the last one-year period (2021). The sample size was calculated based on a formula for cross-sectional study [$n \ge Z1-\alpha 2 p(1-p)/d2$], in which Z was the confidence interval at 95%, d was the margin of error of 5%, and p was the proportion of neonatal death at 50.0%. Therefore, the minimum sample size for this study was 484 participants. The simple random probability selection method was used as the sampling method. All newborns aged 0 to 28 days, hospitalized and deceased in the neonatology unit of Kyeshero Hospital, during the study period, and having

a complete medical file were included. Were excluded, all newborns who died on arrival in the neonatology department, all malformed newborns, and all newborns whose records were deemed unusable.

Data collection

The source of data for this study was the neonatology registers at Kyeshero Hospital which consisted of newborn information recorded on the characteristics of the newborn: the gestational age at birth, the age of the newborn at admission, the sex, the birth weight, the notion of crying, the APGAR score at the 5th and the 10th minute, the notion of resuscitation, the reason for hospitalization, the clinical signs on admission, the diagnosis, the treatment received and mode of feeding, the duration of hospitalization, the mode of discharge, the age and probable cause of death. But also, factors related to the mother: age, parity, origin, number of prenatal consultations, pathological pregnancy history, the notion of multipregnancies, and premature rupture of membranes, maternal fever 48 hours before delivery, antenatal treatment received, place and mode of delivery.

Statistical analyses

Variables were described as absolute frequencies and percentages. The determinants of the intention to vaccinate the child or adolescent were evaluated by univariate and multivariate analyses. The associations between independent variables and the primary outcomes (neonatal mortality) were tested using a t-test or Chi-square test as appropriate. The results of the multivariate analyzes were presented as odds ratio (OR) with standard error (SE) and 95% confidence interval (95% Cl). A step-by-step, bottom-up Wald analysis was performed to define the variables to be included in the final logistic regression model, according to the results of the univariate models and was complemented by analysis of the model's predictive power using the Receiver Operating Characteristic curve (ROC).

Ethical issues

Authorization to carry out the study, and ethical clearance were obtained from the hospital authorities and the hospital ethics committee. The data were kept confidential.

Results

Neonatal mortality rate

During our study period, among 501 newborns admitted in the neonatology unit; 85 newborns died. The neonatal intra-hospital neonatal mortality rate was 17.0%.

The study of the epidemiological characteristics of the newborns revealed that 92% of the newborns were aged between 0-6 days, the male sex was more
 Table 1: Descriptions of socio-demographic characteristics of newborns.

Characteristics of the newborn	Effective	Percentage	
Post-natal age in days			
0-6 days	461	92.0	
7-28 days	40	8.0	
Sex			
Feminine	205	40.9	
Male	296	59.1	
Birth weight			
2500-4000 g	300	59.9	
lower 2500 g	165	32.9	
Greater than 4000 g	36	7.2	
Type of pregnancy			
twin	33	6.6	
Monofetal	467	93.2	
Multiple greater than 2	1	0.2	
APGAR scored in the fifth minute			
Less than 7	104	20.8	
Greater than or equal to 7	397	79.2	
Presentation of the fetus			
Cephalic	437	87.2	
Headquarters	55	11.0	
Transverse	9	1.8	

 Table 2: Description of the maternal characteristics of newborns.

Characteristics of the mother	Effective	Percentage	
Under 18	21	4.2	
greater than or equal to 18-years-old	480	95.8	
Parity			
Grand Multipara	79	15.8	
Multipara	122	24.4	
Pauci parries	154	30.7	
Primiparous	146	29.1	
Gestational age			
32-34 weeks + 6 days	59	11.8	
35-36 weeks + 6 days	56	11.2	
37-40 weeks	309	61.7	
Less than 32 SA	36	7.2	
Greater than 40 SA	41	8.2	
Have followed the ANC			
No	17	3.4	
Yes	484	96.6	
The number of ANC followed			
Less than or equal to 3	330	68.2	
Greater than 3	154	31.8	
Mode of delivery			
Cesarean section	209	41.7	
Low way	292	58.3	
Place of birth			
Outside the health facility	5	1.0	
Sanitary structure	496	99.0	

represented with 59.1% vs. 40.9 for the female sex and in this about weight, and the 2500 to 4000 g bracket was more represented with 59.9% (Table 1).

The maternal age group greater than or equal to 18 years was more represented with 95.5%. The gestational age range from 37-40 SA was more represented with 61.7%, the number of CPN follow-ups less than or equal to 3 was more represented with 68.2% vs. 31.8 for the number greater than 3 The vaginal mode of delivery was the most practiced for these parturients, i.e., 58.3% vs. 41.7% vaginally. These patients were 30.7% parous pauci. For the place of delivery, 99% of these parturients were from Kyeshero hospital (Table 2).

Statistical analyzes of newborn characteristics associated with mortality with a statistically significant association were found in low birth weight newborns ($p = \le 0.0001$) with an Apgar score of less than 7 ($p = \le 0.0001$), with the presence of respiratory distress ($p \le 0.0001$) with the presence of cyanosis ($p = \le 0.0001$), neonatal infection ($p = \le 0.0001$), absence of sucking - swallowing ($p = \le 0.0001$) and absence responsiveness in neonates ($p = \le 0.0001$) (Table 3).

It appears from this Table 4 that the non-followed prenatal consultation, the premature rupture of the membranes, and the delivery at home were associated with the risk of neonatal mortality without this association being statistically significant.

Table 5 shows that low birth weight < 2500 gr aOR = 4.11 (2.17-7.77), presence of cyanosis aOR = 3.88 (1.97-7.65), absence of reactivity aOR = 20.07 (8.07-49.55), and neonatal infection aOR = 4.78 (2.47-9.22) were significantly associated with neonatal mortality.

The age of the parent being < 40 years (aOR: 2.40, CI: [1.50-3.83]), fearing that the parent or a member of his/her family will contract COVID-19 (aOR: 2.35, CI: [1.38-4.02]), thinking that the parent and the children are vulnerable to COVID-19 (aOR: 1.70, CI: [1.005-2.2881]), believing that the family or the parent could contract COVID-19 (aOR: 3.07, CI: [1.80-5.23]), and having been vaccinated against COVID-19 (aOR: 16.47, CI: [8.39-32.33]) were significantly associated with the intention to have the children or adolescents vaccinated.

The neonatal mortality prediction model is as follows (Figure 1):

P= (Y= Neonatal mortality/X=xi) =
$$\frac{e^{-3.56+1.41.x1+1.36.x2+3.00.x3+1.56.x4}}{1+e^{-3.56+1.41.x1+1.36.x2+3.00.x3+1.56.x4}}$$

Table 3: Description of the characteristics of newcomers associated with neonatal mortality.

Characteristics of the newborn		Mortality	GOLD	р	
	Yes No		[95% CI]		
Age in days					
0-6	80 (17.4)	381 (82.6)	1.47 [0.56-3.87]	0.43	
7-28	5 (12.5)	35 (87.5)			
Sex					
Feminine	34 (16.6)	171 (83.4)	0.96 [0.59-1.54]	0.85	
Male	51 (17.2)	245 (82.8)			
Low birth weight					
Yes	58 (35.2)	107 (64.8)	6.20 [3.74-10.30]	≤ 0.0001	
No	27 (8.0)	309 (92.0)			
Macrosomia					
Yes	4 (11.1)	32 (88.9)	0.59 [0.20-1.72]	0.33	
No	81 (17.4)	384 (82.6)			
Intake method					
Referred or brought by parents	30 (25.9)	86 (74.1)	2.09 [1.26-3.47]	0.004	
Self-hospitalization	55 (14.3)	330 (85.7)			
APGAR score in the fifth minute					
< 7	40 (38.5)	64 (61.5)	4.89 [2.96-8.08]	≤ 0.0001	
≥7	45 (11.3)	352 (88.7)			
Presence of respiratory distress	- (-)				
Yes	70 (27.1)	188 (72.9)	5.66 [3.14-10.21]	≤ 0.0001	
No	15 (6.2)	228 (93.8)			
Presence of pallor					
Yes	2 (18.2)	9 (81.8)	1.09 [0.23-5.14]	0.91	
No	83 (16.9)	407 (83.1)			
Presentation of the fetus					
Seat or transverse	18 (28.1)	46 (71.9)	2.16 [1.18-3.95]	0.011	
Cephalic	67 (15.3)	370 (84.7)	[]		
Presence of jaundice					
Yes	3 (16.7)	15 (83.3)	0.98 [0.28-3.46]	0.97	
No	82 (17.0)	401 (83.0)			
Presence of cvanosis	- (-)				
Yes	50 (47.6)	55 (52.4)	9.38 [5.59-15.72]	≤ 0.0001	
No	35 (8.8)	361 (91.2)			
Presence of sepsis					
Yes	0 (0.0)	10 (100.0)		0.15	
No	85 (17.3)	406 (82.7)			
Presence of neonatal infection					
Yes	41 (27.2)	306 (87.4)		≤ 0.0001	
No	44 (12.6)	110 (72.8)			
Sucking-swallowing					
Absent	24 (68.6)	11 (31.4)	14,49 [6,76-31,06]	≤ 0.0001	
Present	61 (13.1)	405 (86.9)			
Reactivity					
Absent	41 (83.7)	8 (16.3)	47.52 [20.95-107 78]	≤ 0.0001	
Present	44 (9.7)	408 (90.3)			
Monofetal	6 (17 6)	28 (82 4)	1 05 [0 42-2 62]	0.91	
Multifetal	79 (16 0)	388 (83 1)	1.00 [0.72-2.02]	0.01	
mannotai	10(10.0)	000 (00.1)			

Table 4: Maternal characteristics associated with neonatal mortality.

Characteristics of the mother		Mortality	GOLD	р	
	Yes	No	[95% CI]		
Age in years					
< 18	2 (9.5)	19 (90.5)	0.50 [0.12-2.20]	0.35	
≥ 18	83 (17.3)	397 (82.7)			
Parity					
Primiparous	19 (13.0)	127 (87.0)	0.66 [0.38-1.14]	0.13	
Multipara	66 (18.6)	289 (81.4)			
Prenatal consultation followed					
No	4 (23.5)	13 (76.5)	1.53 [0.49-4.81]	0.46	
Yes	81 (16.7)	403 (83.3)			
Premature rupture of membranes					
Yes	25 (19.1)	106 (80.9)	1.22 [0.73-2.04]	0.45	
No	60 (16.2)	310 (83.8)			
Delivery mode					
Cesarean section	32 (15.3)	177 (84.7)	0.82 [0.51-1.32]	0.40	
Low way	53 (18.2)	239 (81.8)			
Place of birth					
Residence	1 (20.0)	4 (80.0)	1.23 [0.14-11.11]	0.86	
Hospital	84 (16.9)	412 (83.1)			
Perpartal fever					
Yes	6 (10.7)	50 (89.3)	0.56 [0.23-1.34]	0.19	
No	79 (17.8)	366 (82.2)			
The appearance of amniotic fluid					
Pathologies	31 (15.0)	175 (85.0)	0.79 [0.49-1.28]	0.34	
Clear	54 (18.3)	241 (81.7)			

Table 5: Logistic regression.

	В	Fe	Mold	Ь	Exp (B)	CI for Exp (B) 95%	
Factors explaining neonatal mortality	D	ES	vvald	P		Inferior	Superior
Low birth weight (yes vs. no)	1.41	0.33	18.86	0.000	4.11	2.17	7.77
Presence of cyanosis (yes vs. no)	1.36	0.35	15.37	0.000	3.88	1.97	7.65
Reactivity (absent vs. present)	3.00	0.47	41.37	0.000	20.07	8.07	49.55
Neonatal infection (yes vs. no)	1.56	0.34	21.69	0.000	4.78	2.47	9.22
Constant	-3.56	0.54	44.24	0.000	0.028		

Discussion

The child mortality rate is considered the best proxy indicator of general population health and the level of socioeconomic development. The child mortality rate is also a useful marker of overall development and a Millennium Development Goal (MDG) indicator [14].

During our study period, the in-hospital neonatal mortality rate was 17.0%. However, this contrasts with the higher rates noted by Mashako K, et al. who conducted a study at the neonatal intensive care unit of the North Kivu Provincial Hospital in Goma, eastern Democratic Republic of Congo (19.7%). The decrease of neonatal mortality rate in Goma may be linked to differences in the quality of health care governance including early intervention which includes the availability, accessibility and affordability of health care systems for children. Quality of care plays an important role in improving neonatal outcomes, and the high levels of neonatal mortality in many resource-limited countries can, in part, be attributed to poor quality of care [15,16]. Effective implementation of quality improvement is possible in low income settings [17], with appropriate strategies including the identification of quality gaps, followed by the development of action plans (and their implementation) to overcome barriers [15].

However, the result was lower than studies in Ghana at 20.2% [18]; Mizan Tepi Teaching Hospital, South



West Ethiopia, at 22.8% [19]; Hiwot Fana University Specialty Hospital, Eastern Ethiopia, at 14.3% [20]; and Gondar Referral Hospital in northwestern Ethiopia at 23.1% [21]; University Clinics of Lubumbashi at 36.9% [22] and general referral hospital in Kamina at 25% [23]. Our prevalence of neonatal mortality was higher than in studies conducted in Somali region, Ethiopia, at 5.7% [24]; in Kolwezi, DRC, at 18.8% [25]; and in Mekelle, Ethiopia, at 6.6% [26]. The discrepancy could be due to the presence of socio-cultural and socioeconomic differences between Ethiopian regions. This variation could also be due to variation in sample sizes. Another possible justification may be the difference in the use of health services, including delivery in health facilities by skilled care providers and seeking care for sick newborns, variation in health facility configuration (equipment available and trained people) and economic disparities among study participants [27].

This study found that newborns with a birth weight were 4.11 times higher risk of death than those with a normal or higher birth weight (ORa = 4.11; Cl 95% [2.17-7.77]). The possible justification is that low birth weight increases children's susceptibility to infection and lowers their immune system and other body defense mechanisms, which control newborns' exposure to disease. As a result, neonatal survival is reduced [28]. Other possible explanations could also be due to low birth weight that newborns are expected to suffer from hypothermia, infection and poor immunological function, which increases the risk of neonatal death [29,30].

In the final logistic regression model of this study, we found neonatal infection was statistically associated with neonatal mortality. Thus, the risks of neonatal mortality were almost five times higher in newborns who had an early neonatal infection than in those who did not (ORa = 4.78; CI 95% [2.47-9.22]). This result is comparable with previous hospital studies conducted in the health zone of Kenge in the DRC [31] and in Eastern Ethiopia [32]. The possible explanation could be justified by newborns that had an infection in the neonatal period who risk dying during the first month of life because their immunity can be significantly affected by the course of the disease.

However, in a multivariate model, the association between cyanosis and neonatal mortality was also noted (ORa = 3.88; CI 95% [1.97-7.65]). For Mona A Simões, et al. the presence of cyanosis, gestational age, cause of death, first- and/or fifth-minute Apgar < 6, and pH value were associated with death in the first week of life [33]. Neonatal cyanosis is always a sign of serious pathological processes and may involve diverse organs and impose a significant diagnostic and therapeutic challenge.

Limitation of study

The limitation of this study was it might not indicate a

cause-effect relationship because the study design was cross-sectional. Even though this study investigates the most important determinants of neonatal mortality, our study encountered limitations like missing information both on mother and neonates. For instance, economic status and contextual were not assessed.

Conclusion

The intra-hospital frequency of neonatal deaths is 17.0% was justified by four main causes including: Low birth weight, neonatal infection, presence of cyanosis and absent reactivity. Neonatal mortality was influenced by neonatal factors. Proper care of the newborn should improve neonatal vital prognosis. The study of preventable causes of death and factors that lead to early neonatal death is fundamental to reduce the infant mortality rate in the world.

The results of the current study could be used to determine priorities, plan, evaluate services, and improve newborn health care.

References

- Raju SG, Rao SS (2019) A study of neonatal morbidity and mortality in government general hospital, Srikakulam Andhra Pradesh, India. International Journal of Contemporary Pediatrics 6: 1485.
- Chow S, Chow R, Popovic M, Lam M, Popovic M, et al. (2015) A selected review of the mortality rates of neonatal intensive care units. Front Public Health 3: 225.
- 3. Yaqub A, Ghani Z (2018) Frequency and outcome of neonatal diseases in neonatal intensive care unit at tertiary care hospital Islamabad. Isra Med J 10: 272-275.
- Islam A, Butt ZA, Sathi NJ (2022) Prevalence of neonatal mortality and its associated factors: A meta-analysis of demographic and health survey data from 21 developing countries. Dr. Sulaiman Al Habib Medical Journal 4: 145-152.
- Sharrow D, Hug L, You D, Alkema L, Black R, et al. (2022) Global, regional, and national trends in under-5 mortality between 1990 and 2019 with scenario-based projections until 2030: A systematic analysis by the UN Inter-agency Group for Child Mortality Estimation. Lancet Glob Health 10: e195-e206.
- Liu L, Oza S, Hogan D, Chu Y, Perin J, et al. (2016) Global, regional, and national causes of under-5 mortality in 2000-15: An updated systematic analysis with implications for the Sustainable Development Goals. Lancet 388: 3027-3035.
- Hug L, Alexander M, You D, Alkema L (2019) National, regional, and global levels and trends in neonatal mortality between 1990 and 2017, with scenario-based projections to 2030: A systematic analysis. Lancet Glob Health 7: e710-e720.
- Mashako RM, Ngbonda D, Alworong'a Oj, Bitwe MR, Mashako KY, et al. (2019) Predictive factors of neonatal mortality in intensive neonatal care unit at Goma Eastern Democratic Republic of Congo. J Pediatr Neonatal Care 9: 58-61.
- Kananura RM, Tetui M, Mutebi A, Bua JN, Waiswa P, et al. (2016) The neonatal mortality and its determinants in rural communities of Eastern Uganda. Reproductive Health 13: 1-9.

- Lindskog EE (2016) The effect of war on infant mortality in the Democratic Republic of Congo. BMC Public Health 16: 1059.
- 11. Ndayisenga T (2016) Maternal and newborn risk factors associated with neonatal mortality in gitwe district hospital in ruhango district, Rwanda. International Journal of Medicine and Public Health 6: 98-102.
- 12. Dare S, Oduro AR, Owusu-Agyei S, Mackay DF, Gruer L, et al. (2021) Neonatal mortality rates, characteristics, and risk factors for neonatal deaths in Ghana: Analyses of data from two health and demographic surveillance systems. Glob Health Action 14: 1938871.
- 13. Ndeba Prudence M, Mbeva Jean Bosco K, Kila Serge N, Kalemo Oscar K, Kasambula Polycarpe K, et al. (2018) Determinants of neonatal mortality in referral facilities of the health districts in North Kivu, Democratic Republic of Congo. International Journal of Sciences: Basic and Applied Research (IJSBAR) 38: 25-38.
- 14. Kandala NB, Mandungu TP, Mbela K, Nzita KPD, Kalambayi BB, et al. (2014) Child mortality in the Democratic Republic of Congo: Cross-sectional evidence of the effect of geographic location and prolonged conflict from a national household survey. BMC Public Health 14: 266.
- Ouedraogo P, Villani PE, Tubaldi L, Bua J, Uxa F, et al. (2022) Impact of a quality improvement intervention on neonatal mortality in a regional hospital in Burkina Faso. J Matern Fetal Neonatal Med 35: 4818-4823.
- van den Broek NR, Graham WJ (2009) Quality of care for maternal and newborn health: The neglected agenda. BJOG 116: 18-21.
- 17. Zaka N, Alexander EC, Manikam L, Norman ICF, Akhbari M, et al. (2018) Quality improvement initiatives for hospitalised small and sick newborns in low- and middle-income countries: A systematic review. Implement Sci 13: 20.
- Owusu BA, Lim A, Makaje N, Wobil P, Sameae A (2018) Neonatal mortality at the neonatal unit: The situation at a teaching hospital in Ghana. Afr Health Sci 18: 369-377.
- Mekonnen T, Tenu T, Aklilu T, Abera T (2018) Assessment of neonatal death and causes among admitted neonates in neonatal intensive care unit of Mizan Tepi University Teaching Hospital, Bench Maji Zone, South-West Ethiopia, 2018. Clinics Mother Child Health 15: 4.
- 20. Eyeberu A, Shore H, Getachew T, Atnafe G, Dheresa M (2021) Neonatal mortality among neonates admitted to NICU of Hiwot Fana specialized university hospital, eastern Ethiopia, 2020: A cross-sectional study design. BMC Pediatr 21: 125.
- 21. Kokeb M, Desta T (2016) Institution based prospective cross-sectional study on patterns of neonatal morbidity at Gondar University Hospital Neonatal Unit, North-West Ethiopia. Ethiop J Health Sci 26: 73-79.
- 22. Wakamb KA, Adonis NM, Toni LK, Oscar LN (2012) Mortalité á L'unité de Néonatologie des Cliniques Universitaires de Lubumbashi; Congo. Rev Méd Gd Lacs 1: 232-244.
- Kalonji DC, Mbayo PM, Kembo LN, Ngombe MI, Ngimbi SL, et al. (2018) Fréquence et causes de la mortalité néonatale précoce à Kamina, République Démocratique du Congo. Revue de l'Infirmier Congolais 2: 90-94.
- 24. Farah AE, Abbas AH, Ahmed AT (2018) Trends of admission and predictors of neonatal mortality: A hospital based retrospective cohort study in Somali region of Ethiopia. PLoS One 13: e0203314.

- 25. Michel KN, Bertin MK, Deddy KT, Elie KNU, Jack KB, et al. (2016) Risk factors for mortality of newborn at Kolwezi Hospital. Open Access Library Journal 3: 1-8.
- 26. Roro EM, Tumtu MI, Gebre DS (2019) Predictors, causes, and trends of neonatal mortality at Nekemte Referral Hospital, east Wollega Zone, western Ethiopia (2010-2014). Retrospective cohort study. PLoS One 14: e0221513.
- 27. Tadesse AW, Negussie YM, Aychiluhm SB (2021) Neonatal mortality and its associated factors among neonates admitted at public hospitals, pastoral region, Ethiopia: A health facility based study. PloS One 16: e0242481.
- 28. Shah R, Sharma B, Khanal V, Pandey UK, Vishwokarma A, et al. (2015) Factors associated with neonatal deaths in Chitwan district of Nepal. BMC Res Notes 8: 818.
- 29. Reyes JCL, Ramírez ROP, Ramos LL, Ruiz LMG, Vázquez EAB, et al. (2018) Neonatal mortality and associated factors in newborn infants admitted to a Neonatal Care Unit. Arch Argent Pediatr 116: 42-48.

- 30. Seid SS, Ibro SA, Ahmed AA, Akuma AO, Reta EY, et al. (2019) Causes and factors associated with neonatal mortality in Neonatal Intensive Care Unit (NICU) of Jimma University Medical Center, Jimma, South West Ethiopia. Pediatric Health Med Ther 10: 39-48.
- 31. Ngana WN, Piay FP, Mukinda Z, Kitonguna PN (2019) Determinants of neonatal mortality in the Health Zone Kenge, DR Congo (2013-2016). Biomedical Journal of Scientific & Technical Research 18: 13598-13608.
- 32. Roble AK, Ayehubizu LM, Olad HM (2022) Neonatal sepsis and associated factors among neonates admitted to neonatal intensive care unit in general hospitals, Eastern Ethiopia 2020. Clin Med Insights Pediatr 16: 11795565221098346.
- Simões MA, Pabis FC, Freitas AKE, Watanabe PK, Kayano RM, et al. (2016) Preventable causes of death and factors associated with newborn survival at a university hospital in Curitiba, Paraná, Brazil. J Bras Patol Med Lab 52: 338-344.

