

# THE LANCET

## Global Health

### Supplementary appendix

This appendix formed part of the original submission and has been peer reviewed.  
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Supplement to: The WHO Global Maternal Sepsis Study (GLOSS) Research Group.  
Frequency and management of maternal infection in health facilities in 52 countries  
(GLOSS): a 1-week inception cohort study. *Lancet Glob Health* 2020; **8**: e661–71.

## Supplementary materials

**Table S1. Reference list of infections\* associated with systemic repercussions during pregnancy, childbirth, post abortion and postpartum period (modified from ICD-MM, the WHO Application of ICD-10 to deaths during pregnancy, childbirth, and the puerperium)**

<p><b>Pregnancy-related infection (ICD-MM Group 4)</b></p> <ul style="list-style-type: none"><li>• Acute pyelonephritis</li><li>• Infection of amniotic sac and membranes (amnionitis, chorioamnionitis, membranitis, placentitis)</li><li>• Retained products of conception</li><li>• Endometritis, endomyometritis</li><li>• Pelvic abscess</li><li>• Uterine microabscess or necrotizing myometritis</li><li>• Necrotizing fasciitis</li><li>• Necrotizing vulvitis</li><li>• Infection of obstetric surgical wound (caesarean section, perineal repair)</li><li>• Episiotomy infection or dehiscence</li><li>• Other infection of genital tract following delivery (cervicitis, vaginitis following delivery, genital tract laceration)</li><li>• Pyrexia of unknown origin following delivery</li><li>• Infections of breast associated with childbirth (abscess of the nipple, abscess of the breast, subareolar abscess, mastitis, lymphangitis of breast)</li><li>• Tetanus</li></ul> <p><b>Maternal infectious and parasitic diseases classifiable elsewhere but complicating pregnancy, childbirth and the puerperium</b></p> <ul style="list-style-type: none"><li>• Pneumonia</li><li>• Other pulmonary infections (mycoplasma, legionella)</li><li>• Acute viral infections (influenza, H1N1, herpes with systemic repercussion, varicella, acute infectious hepatitis, Encephalitis, dengue, chikungunya, yellow fever, other haemorrhagic fever)</li><li>• Malaria</li><li>• Complicated tuberculosis</li><li>• Listeriosis</li><li>• Leptospirosis</li><li>• Rickettsioses (scrub typhus, murine typhus)</li></ul> <p><b>Exclusion criteria. Women presenting the following conditions were excluded, unless they present with systemic repercussion due to infection:</b></p> <ul style="list-style-type: none"><li>• Any non-severe, localized, uncomplicated infection<ul style="list-style-type: none"><li>◦ Vaginosis, candidiasis</li><li>◦ Lower tract urinary infection</li><li>◦ Fungal infections of the skin (athlete's foot, jock itch, ringworm, and yeast infections)</li><li>◦ Otitis</li><li>◦ Pharyngitis</li><li>◦ Herpes simplex, herpes zoster (shingles)</li></ul></li><li>• Any uncomplicated chronic infection<ul style="list-style-type: none"><li>◦ Sexually transmitted infections (gonorrhoea, syphilis, trichomonas, chlamydia, hepatitis, HIV)</li><li>◦ Tuberculosis</li></ul></li><li>• Any colonization (presence of microorganisms without clinical signs/symptoms)<ul style="list-style-type: none"><li>◦ Known GBS vaginal, urethral and/or rectal colonization</li><li>◦ Asymptomatic bacteriuria</li><li>◦ Known oropharyngeal colonization</li></ul></li><li>• Any iatrogenic hypothermia/hyperthermia (e.g. related to epidural, thyroid storm, prostaglandin administration) during hospital stay</li><li>• Use of any prescription of prophylactic antibiotics (e.g. for GBS colonization, after caesarean section, manual removal of the placenta, vaginal delivery)</li></ul>
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\*study eligibility was not limited to those

**Table S2. Characteristics of participating facilities**

Characteristics	n (%)
Public facility (yes)	417 (78.5%)
Location	
Urban	514 (80.7%)
Peri-urban	80 (12.6%)
Rural	43 (6.8%)
Level	
Tertiary	220 (34.5%)
Secondary	305 (47.9%)
Primary	220 (34.5%)
Country income level*	
Low income	82 (11.5%)
Lower-middle income	200 (28.1%)
Higher-middle income	126 (17.7%)
High income	305(42.8%)
Maternity exclusive hospital (yes)	95 (14.6%)
Median number of births/year in 2016 (Q1-Q3)	1993 (861-4211)
Number of births per year in 2016	
<1000	137 (26.7)
1000-2499	156 (30.4)
2500-4499	103 (20.1)
>4500	117 (22.8)
Median number of births during identification week (Q1-Q3)	40 (16.78)
Adult intensive care or high dependency unit (yes)	540 (84.8%)

\*using the 2018 World Bank classification (<https://datatopics.worldbank.org/worlddevelopment-indicators/theworld-by-income-and-region.html>)

**Table S3: Distribution of underlying causes of maternal deaths**

	Total maternal deaths		Maternal deaths that occurred during the week of identification	
	Primary cause of maternal death <sup>b</sup>	Infection-related maternal deaths <sup>a,b,c</sup>	Primary cause of maternal death <sup>b</sup>	Infection-related maternal deaths <sup>a,b,c</sup>
	(n=46)	(n=26)	(n=39)	(n=19)
Obstetric haemorrhage	9 (20%)	2 (8%)	9 (23%)	2 (11%)
Hypertensive disorder	8 (17%)	1 (4%)	7 (18%)	0 (0%)
Infection/Sepsis	7 (15%)	7 (27%)	6 (15%)	6 (32%)
Other direct	1 (2%)	1 (4%)	1 (3%)	1 (5%)
Abortion	5 (11%)	5 (19%)	3 (8%)	3 (16%)
Any indirect cause	11 (24%)	8 (31%)	9 (23%)	6 (32%)
Unknown	5 (11%)	2 (8%)	4 (10%)	1 (5%)

<sup>a</sup>Infection is underlying or contributing cause of maternal death; <sup>b</sup>Includes all maternal deaths reported in the study: 20 maternal deaths not related to infections; <sup>c</sup>Seven maternal deaths occurred after the identification week.

**Table S4: Distribution of organ dysfunction by system**

	<b>Severe maternal outcomes related to infections</b>	<b>Infection-related near-miss</b>	<b>Infection-related maternal death</b>
	<b>(n=381)</b>	<b>(n=355)</b>	<b>(n=26)</b>
Cardiovascular	160 (42.0%)	139 (39.1%)	21 (80.8%)
Respiratory	150 (39.4%)	130 (36.6%)	20 (76.9%)
Renal	66 (17.3%)	57 (16.1%)	9 (34.6%)
Coagulation	105 (27.6%)	97 (27.3%)	8 (30.8%)
Hepatic	46 (12.1%)	42 (11.8%)	4 (15.4%)
Neural	34 (8.9%)	25 (7.0%)	9 (34.6%)
Uterine	79 (20.7%)	75 (21.1%)	4 (15.4%)
None	1 (0.3%)	0 (0.0%)	1 (3.8%)
Multiple organ system dysfunction	143 (37.5%)	122 (34.4%)	21 (80.8%)