

Supplemental Table 3. Review of reports describing features of hemophagocytic lymphohistiocytosis (HLH) and culture proven invasive salmonellosis (typhoidal or non-typhoidal).

S. No.	Authors, year of publication [reference]	Country	Microbe, No. of patients	Features of HLH (as per HLH-2004 criteria). Comments	Therapy{Dose, duration}
1.	Kim BS et al., 1971 [1]	United States	<i>S. Typhi</i> *, 2	Bone marrow examination (BME) suggestive of HLH	Not available (NA)
2.	Macias EG, 1975 [2]	United States	<i>S. Typhi</i> , 2	BME at presentation: histiocytic proliferation with hemophagocytosis and granuloma. BME after 3 weeks of antimicrobial therapy: normal	Chloramphenicol {~3 weeks}
3.	Fernandes-Costa F et al., 1979 [3]	South Africa	<i>S. Typhi</i> *, 2	Fever, pancytopenia, splenomegaly, and BME showing histiocytic proliferation with hemophagocytosis. Repeat BME in 1 patient was normal	Antibiotics. Details: NA
4.	Benz-Lemoine E et al., 1983 [4]	France	<i>S. Typhimurium</i> septicemia, 1 pediatric patient	Child with chronic granulomatous disease (CGD). Article in French language. Details: NA	NA

			<18 years (P)		
5.	Fame TM et al., 1986 [5]	United States	<i>S. Typhi</i> , 1 (P)	Fever, hepatosplenomegaly, pancytopenia, and BME showing increased histiocytes with hemophagocytosis (overall hypocellular marrow). Readmitted again after 3 weeks for typhoid, no features of HLH	Intravenous (IV) ampicillin {100mg/kg/d, 10 days} followed by oral amoxicillin {1.5 g/d, 2 weeks}. IV and oral cotromoxazole {10/50 mg/kg/d, 2 weeks}
6.	Udden MM et al., 1985 [6]	United States	<i>S. Typhi</i> , 5 (P=1)	In all 5: fever, cytopenias, and BME showing histiocytosis and hemophagocytosis. Splenomegaly was seen in 2 and hypofibrinogenemia in 1.	Ampicillin or chloramphenicol. Details: NA
7.	Mallouh AA et al., 1987 [7]	Saudi Arabia	<i>S. Typhi</i> , 3 (P=3)	Fever, cytopenia(s), and BME showing histiocytosis and hemophagocytosis. Other details: NA	NA
8.	Chien YH et al., 1999 [8]	Taiwan	<i>S. Typhi</i> , 1 (P)	Fever, pancytopenia, elevated triglycerides, and	IV ceftriaxone. Details:

				BME showing hemophagocytosis. Other details: NA.	NA.
9.	Karthik R, 2007 [9]	India	<i>S. Typhi</i> , 1 (P)	Fever, cytopenia, elevated triglycerides, elevated ferritin, and BME showed hemophagocytosis.	IV ceftriaxone {2 weeks}
10.	Koul PA et al., 2010 [10]	India	<i>S. Typhi</i> , 3	Fever, cytopenia, splenomegaly, and BME showing hemophagocytosis (same features in all)	Antibiotics. Details: NA
11.	Khalaf D et al., 2011 [11]	Saudi Arabia	<i>S. Typhi</i> , 1	Fever, splenomegaly, cytopenia, hypofibrinogenemia, hyperferritinemia, and BME showing hemophagocytosis.	Floroquinolone, IV dexamethasone {8 days}, and IVIg {4 days}
12.	Shah PA et al., 2011 [12]	India	<i>S. Typhi</i> , 1	Fever, splenomegaly, pancytopenia, elevated ferritin, elevated triglycerides, and BME showing hemophagocytosis	IV ceftriaxone {2 g/day, 10 days}
13.	Nath UK et al., 2013 [13]	India	<i>S. Paratyphi</i> , 1	Fever, splenomegaly, pancytopenia, elevated ferritin, elevated triglycerides, decreased fibrinogen, and BME showed hemophagocytosis (overall normocellular marrow)	IV ceftriaxone {10 days}. Dexamethasone {10 mg/m ² /d, 3 weeks} was employed upfront

14.	Iqbal N et al., 2013 [14]	India	<i>S. Typhi</i> , 2	Both fulfilled HLH-2004 criteria. Details: NA	NA
15.	Non LR et al., 2015 [15]	India	<i>S. Typhi</i> , 1	Fever, splenomegaly, cytopenias, elevated ferritin, and elevated soluble interleukin-2 receptor (sIL-2R). Additionally, features of rhabdomyolysis	IV ceftriaxone followed by cotrimoxazole {total 2 weeks}
16.	Song M et al., 2017 [16]	China	<i>S. Choleraesuis</i>	NA	NA
17.	Uribe-Londono J et al., 2018 [17]	Colombia	<i>S. Typhi</i> , 1 (P)	Fever, splenomegaly, pancytopenia, elevated ferritin, and elevated triglycerides. BME did not show hemophagocytosis	Ciprofloxacin {~2 weeks}. Intravenous immunoglobulin (IVIg) {1 g/kg/d, 2 days} was given upfront.
18.	Abbas A et al., 2018 [18]	Pakistan	<i>S. Typhi</i> , 1 (P)	Fever, pancytopenia, elevated ferritin and triglycerides, decreased fibrinogen, and BME showed hemophagocytosis (overall suppressed hematopoiesis)	IV ceftriaxone {2 weeks}
19.	Sánchez-Moreno P et al.,	Spain	<i>S. Typhi</i> , 1 (P)	Fever, splenomegaly, pancytopenia, elevated	IV ceftriaxone {80

	2019 [19]			ferritin, and decreased fibrinogen.	mg/kg/d, 2 weeks} and methylprednisolone {initially 2 mg/kg, 2 weeks}
20.	Şahin Yaşar A et al., 2019 [20]	Turkey	<i>S. Typhi</i> , 1 (P)	Fever, pancytopenia, elevated ferritin and triglycerides, decreased fibrinogen, and BME showing histiocytes with hemophagocytosis. Additionally, findings of rhabdomyolysis	IV ceftriaxone {2 weeks}. Dexamethasone {10 mg/m ² /d} and IVIg {0.5 g/kg} given upfront
21.	Durmuş SY et al., 2020 [21]	Turkey	<i>S. Typhimurium</i> septicemia, 1 (P)	Fever, cytopenias, elevated ferritin and triglycerides, decreased fibrinogen, and splenomegaly (ultrasonography). BME did not show hemophagocytosis	IV ceftriaxone {100 mg/kg/d, 10 days}. IVIg {1 g/kg/d, 2 days} started on day 2
22.	Wei A et al., 2020 [22]	China	<i>S. Typhimurium</i> septicemia, 1 (P)	Fever, splenomegaly, pancytopenia, elevated ferritin, elevated sIL-2R α , decreased fibrinogen, decreased natural killer cell activity, and BME showed hemophagocytosis. Diagnosed as CGD	IV Meropenem {~2 weeks}. Methylprednisolone {initial dose 10 mg/kg/d}

					given upfront.
23.	Banday AZ et al., 2020 [our report]	India	<i>S. Enteritidis</i> , 1 (P)	Fever, splenomegaly, cytopenia, elevated ferritin, decreased fibrinogen, and BME showing histiocytic proliferation with marked hemophagocytosis	IV Meropenem {120 mg/kg/d, 2 weeks}. IV dexamethasone {initially 10 mg/m ² /d, 2 weeks} was given upfront

Literature search was performed using PubMed database and references of retrieved articles were checked for relevant articles.

Abbreviations used in the table elaborated at first use. * Culture positivity not explicitly mentioned in the report.

References for Supplemental Table 3:

1. Kim BS, Sanders DY, 1971. Bone-marrow studies in two cases of typhoid fever. *NC Med J* 32:339–343.
2. Macias EG, 1975. Letter: Typhoidal cells. *Lancet* 2(7941):927–928.
3. Fernandes-Costa F, Eintracht I, 1979. Histiocytic medullary reticulosis. *Lancet* 2(8135):204–205.
4. Benz-Lemoine E, Bordigoni P, Schaack JC, Briquel E, Chiclet AM, Olive D, 1983. Systemic reactive histiocytosis with hemophagocytosis and hemostasis disorders associated with septic granulomatosis. *Arch Fr Pediatr* 40(3):179–182. Article in French.
5. Fame TM, Engelhard D, Riley HD Jr., 1986. Hemophagocytosis accompanying typhoid fever. *Pediatr Infect Dis* 5(3):367–369.
6. Udden MM, Bañez E, Sears DA, 1986. Bone marrow histiocytic hyperplasia and hemophagocytosis with pancytopenia in typhoid fever. *Am J Med Sci* 291(6):396–400.
7. Mallouh AA, Sa'di AR, 1987. White blood cells and bone marrow in typhoid fever. *Pediatr Infect Dis J* 6(6):527–529.
8. Chien YH, Lee PI, Huang LM, Lee CY, Lin DT, Lin KH, 1999. Typhoid fever presenting as infection-associated hemophagocytic syndrome: report of one case. *Acta Paediatr Taiwan* 40(5):339–340.
9. Karthik R, 2007. Infectious causes of macrophage activation syndrome. *J Assoc Physicians India* 55:877–878.
10. Nath UK, Sinha N, De D, 2013. Hemophagocytic lymphohistiocytosis secondary to *Salmonella Paratyphi A* infection presenting with severe pancytopenia and multiorgan dysfunction: the first case report. *J Bone Marrow Res* 1:114.

11. Koul PA, Khan U, Shah S, Jan R, Wani A, Qadri A, Masoodi ZA, 2010. Adult hemophagocytic lymphohistiocytosis: a 25-year experience at a tertiary care hospital. *WebmedCentral Infectious Diseases* 1(9):WMC00674.
12. Khalaf D, Toema B, Al-sadadi S, Al-jehani F, Sammak M, 2011. Salmonella Typhi associated hemophagocytic lymphohistiocytosis in a previously healthy 23 years old woman. *WebmedCentral Infectious Diseases* 2(3):WMC001751.
13. Nath UK, Sinha N, De D, 2013. Hemophagocytic lymphohistiocytosis secondary to Salmonella Paratyphi A infection presenting with severe pancytopenia and multiorgan dysfunction: the first case report. *J Bone Marrow Res* 1:114.
14. Non LR, Patel R, Esmaeeli A, Despotovic V, 2015. Typhoid fever complicated by hemophagocytic lymphohistiocytosis and rhabdomyolysis. *Am J Trop Med Hyg* 93(5):1068–1069.
15. Iqbal N, et al., 2015. Clinicopathological profile of Salmonella Typhi and Paratyphi infections presenting as fever of unknown origin in a tropical country. *Mediterr J Hematol Infect Dis* 7(1):e2015021.
16. Song M, Qiu H, 2017. Clinical analysis of Gram-negative bacillisepticemia-associated hemophagocytic lymphohistiocytosis. *Blood* 130 (Supplement 1):2292.
17. Uribe-Londono J, Castano-Jaramillo LM, Penagos-Tascon L, Restrepo-Gouzy A, Escobar-Gonzalez AF, 2018. Hemophagocytic lymphohistiocytosis associated with Salmonella Typhi infection in a child: a case report with review of literature. *Case Rep Pediatr* 2018:6236270.

18. Abbas A, Raza M, Majid A, Khalid Y, Bin Waqar SH, 2018. Infection-associated hemophagocytic lymphohistiocytosis: an unusual clinical masquerader. *Cureus* 10(4):e2472.
19. Sánchez-Moreno P, Olbrich P, Falcón-Neyra L, Lucena JM, Aznar J, Neth O, 2019. Typhoid fever causing haemophagocytic lymphohistiocytosis in a non-endemic country - first case report and review of the current literature. *Enferm Infect Microbiol Clin* 37(2):112–116.
20. Şahin Yaşar A, Karaman K, Geylan H, Çetin M, Güven B, Öner AF, 2019. Typhoid fever accompanied with hematopoietic lymphohistiocytosis and rhabdomyolysis in a refugee child. *J Pediatr Hematol Oncol* 41(4):e233–e234.
21. Durmuş SY, Tanır G, Öz FN, Teke TA, Kaman A, 2020. Salmonella ser. Typhimurium bacteremia related hemophagocytic lymphohistiocytosis: a case report. *J Pediatr Inf /Cocuk Enfeksiyon Dergisi* 14(1):e35–e37.
22. Wei A, et al., 2020. Hemophagocytic lymphohistiocytosis resulting from a cytokine storm triggered by septicemia in a child with chronic granuloma disease: a case report and literature review. *BMC Pediatr* 20(1):100.