

Sero-Detection of Cytomegalovirus and Rubella Virus IgG Antibodies Among Sudanese Pregnant Women in Khartoum State-Sudan

T. Badri¹, M. Javed²

Lecturer of Internal Medicine and Clinical Hematology, Faculty of Medicine, Al-Azhar University, Egypt^{1,2}.



Abstract— Cytomegalovirus and Rubella infection are the most well-known reasons for innate contaminations, which increment dreariness and mortality during childbirth and one of the normal reasons for fetus removal in creating nations. Plasma tests acquired from 87 pregnant womens, the examples were gotten from Omdurman Friendship Hospital 53(61%), AL-Saudi Specialized Hospital 21(25%) and Ultra lab Diagnostic Center 13(14%).all examples were inspected for nearness of CMV and Rubella infection IgG antibodies by utilizing an ELISA test. The outcome demonstrated that out of 87 blood tests explored, 64(73.6%) were sure for CMV, while the reslt 23(26.4%) were negative and 85(97.7%) were sure for rubella, while the reslt 2(2.3%) were negative, where was 62(71.2%) examples had both CMV and Rubella infection IgG antibodies, 25(28.8%) had either CMV or Rubella infection IgG antibodies and there was no example negative for both CMV and Rubella infection IgG antibodies. The present examination watched the high commonness rate of CMV and rubella infection IgG antibodies among pregnant ladies in Khartoum State. The degree of diseases is higher in pregnant ladies without history of unsuccessful labor than those prematurely ended ladies.

Keywords—Cytomegalovirus; Rubella Virus; Pregnant Women; IgG Antibody; Sero-Detection.

1. Introduction

Inherent CMV contamination is one of the TORCH contaminations (toxoplasmosis, rubella, CMV, and HSV), which convey a danger of noteworthy symptomatic illness and formative deformities in babies [1]. Human cytomegalovirus (HCMV) has a place with the β -herpesvirus subfamily, and albeit most solid people stay asymptomatic ensuing to contamination, the pathogen is a noteworthy supporter of birth abandons and to hazardous ailment in immunocompromised patients [2-4]. Likewise with all infections, HCMV relies upon the host cell to give macromolecular structure squares to virion creation, and over the span of its development, HCMV has adjusted to control various basic cell forms, including RNA aggregation [5], interpretation [6], digestion [7-8], secretory pathways [9], and the cell cycle [10]. Cytomegalovirus contamination during pregnancy is a noteworthy reason for inborn disease worldwide with an occurrence of 0.2 – 2.2% of live births. Up to 15% of such kids have infants following intrauterine CMV contamination [11]. Disease in the infant can be gained through close contact (by means of polluted blood, pee, and discharges), vertically through Trans placental transmission, and postnatal through bosom milk [12]. Rubella infection (RuV) is a little wrapped single-stranded RNA infection and the sole individual from the Rubivirus variety. Rubivirus and alphaviruses together include the Togaviridae [8]. While alphaviruses are commonly transmitted by mosquito vectors, RuV spreads via airborne transmission between people [13]. The main known host [9]. RuV causes a mellow youth illness generally alluded to as German measles [8-10]. Rubella (at first known as German measles) is related with a 80% danger of generally various inherent variations from the norm whenever gained in the initial 12 weeks of pregnancy

[14], particularly the initial 8-10 weeks, and prompts fetal development issues or still birth [15]. The infection at first recreates in the nasopharyngeal mucosa and nearby lymph hubs, and in pregnancy taints the placenta and creating baby.

2. Materials and Methods

2.1 Study structure and length

An expressive Cross-sectional examination was led to identify Human Cytomegalovirus and Rubella infection IgG Antibodies among pregnant ladies going to Omdurman Friendship Hospital, Al-Saudi Maternity Hospital and Ultra Lab Diagnostic Center, Khartoum, Sudan. During the period from May to June 2016.

2.2 Collection and planning of tests

A fringe blood example was gathered from each pregnant lady into EDTA-containing vacutainer tubes, centrifuged at 3000 RPM for 5 minutes and the acquired plasma was put away at - 20°C until utilized.

2.3 Immunoassay for HCMV and Rubella infection IgG antibodies identification

Plasma tests were inspected for against HCMV and Rubella infection IgG antibodies by a circuitous Enzyme-connected Immunosorbent Assay (ELISA) unit (prescience, Acon research centers, Inc., 10125Mesa Rim Road, San Diego, CA92121, and USA).

2.4 Data Analysis

Factual investigation was finished by utilizing Statistical Package for Social Science program (SPSS-adaptation 16).

3. Results

A sum of 87 blood tests (n=87) were gathered from pregnant ladies in Khartoum State. All examples were inspected for the nearness of CMV and Rubella infection IgG antibodies utilizing an ELISA packs.

Out of 87 blood tests explored, 85(97.7%) and 64(73.6%) were responsive for hostile to CMV and against Rubella IgG antibodies, separately. Exceptionally compelling, 62(71.2%) examples were receptive for both enemy of CMV and hostile to Rubella IgG antibodies, while there was no example demonstrated non responsive outcome for neither enemy of CMV nor against Rubella infection IgG antibodies (Table 1).

Result	CMV IgG antibodies		Rubella IgG antibodies		CMV and Rubella IgG antibodies	
	No.	percentage	No.	percentage	No.	Percentage
Positive	64	73.6%	85	97.7%	62	71.2%
Negative	23	26.4%	2	2.3%	25	28.8%
Total	87	100%	87	100%	87	100%

Table 1: Prevalence of CMV and Rubella IgG antibodies among pregnant women

Out of 38 ladies with history of premature birth 31(81.6%) and 37(97.4%) were receptive for hostile to CMV and against Rubella infection IgG antibodies, individually, with P-esteem (0.882). Also out of 49

ladies without history of premature birth 32(65.3%) and 47(95.9%) were responsive for against CMV and hostile to Rubella infection IgG antibodies, separately, with P-esteem (0.106) (Table 2).

Result		CMV IgG antibodies		Rubella IgG antibodies	
		No.	percentage	No.	Percentage
Abortion (n=38)	Positive	31	81.6%	37	97.4%
	Negative	7	18.4%	1	2.6%
No abortion (n=49)	Positive	32	65.3%	47	95.9%
	Negative	17	34.7%	2	4.1%

Table 2: distribution of CMV and Rubella virus IgG antibodies according to history of abortion

4. Discussion

Human cytomegalovirus and Rubella infection are two of the vertically transmitted contaminations that lead to innate variations from the norm and pregnancy issues. Studies demonstrated that ladies who are presented to cytomegalovirus or potentially Rubella infection just because during pregnancy may have a higher danger of unsuccessful labor. These contaminations can prompt significant entanglements on pregnancy for maternal and fetal wellbeing [16,17]. The present examination went for recognition of against CMV and hostile to Rubella infection IgG antibodies among pregnant ladies in Khartoum State.

A sum of 87 blood tests examined, 67(73.6%) and 85(97.7%) were certain for hostile to CMV and against Rubella infection IgG antibodies, individually. Our CMV result (73.6%) was like that acquired in western Sudan [18] who detailed that 72.2% of pregnant ladies were hostile to CMV-IgG antibodies receptive, yet higher than result got in Mexico, (65.6%) by Luis et al. [19] and not exactly those acquired in Nigeria (91.1%) by Hamid et al. [20], in Palestine (96.6%) by Tahani et al. [21] and in China (98.7%) by Lingqing et al. [22]. These distinctions may be credited to endemic varieties of these nations with CMV diseases and diverse wellbeing approaches took a crack at these nations. Specifically, noteworthy, the most astounding frequencies of receptive enemy of CMV IgG antibodies were seen among the main trimester of pregnancy and those without history of premature delivery. Nonetheless, no huge distinction ($P > 0.05$) was seen among the three trimesters of pregnancy. While Rubella infection results (97.7%) were higher than that acquired in Nigeria and Sudan 70% and 65.3%, individually [18,23], and in accordance with the outcome got in Mozambique is practically 100% [24,25]. Various examinations uncover variable consequences of the seroprevalence of rubella over various landmasses; 54.1% in Nigerian [26], 76% in Sri Lanka [27], 77.5% in Russian [28] and 93% in Eritrea [29]. These distinctions might be because of endemicity varieties of these nations with rubella diseases and ongoing presentation of Rubella antibody alone or in mix as MMR immunizations in national inoculation plan.

In the present examination, the higher (54%) occurrence of seropositivity for rubella infection IgG immune response was seen in pregnant ladies without history of premature delivery than that of the ordinary pregnancy (43.8%) results gathering, proposed that rubella could be a reason for rehashed pregnancy wastage in those ladies. Comparable proof was found in Punjab, India that higher (73.2%) occurrence was found in the unfavorable pregnancy result bunch than the typical (69.5%) obstetric result bunch [30].

Our finding identified that the most elevated seropositivity of rubella infection was accounted for in those pregnant ladies inside the third trimester (55.2%) of incubation than others. Nonetheless, no huge distinction ($P > 0.05$) was seen among the three pregnancy trimesters.

5. Conclusion

As per our outcomes we presume that there is high predominance rate CMV and rubella infection IgG antibodies among pregnant ladies in Khartoum State. The degree of contaminations is higher in those pregnant ladies without history of unsuccessful labor than those prematurely ended ladies and factual examination demonstrates that there is no huge relationship among CMV and rubella infection diseases and fetus removal.

From this investigation, we saw that the high predominance rate of CMV and rubella infection diseases were found in those ladies in the third trimester of development and those inside the age bunch \geq multi year-old. The present examination additionally presumes that there is a high rate of blend contaminations by CMV and Rubella infection among pregnant ladies.

4. References

1. Ljungman P, Griffiths P, and Paya C. Definitions of cytomegalovirus infection and disease in transplant recipients. *Clin Infect Dis* 34 (2002): 1094-1097.
2. Dollard SC, Grosse SD, Ross DS. New estimates of the prevalence of neurological and sensory sequelae and mortality associated with congenital cytomegalovirus infection. *Rev Med Virol* 17 (2007): 355-363.
3. Ross SA, Boppana SB. Congenital cytomegalovirus infection: Outcome and diagnosis. *Semin Pediatr Infect Dis* 16 (2005): 44-49.
4. Gerna G, Baldanti F, Revello MG. Pathogenesis of human cytomegalovirus infection and cellular targets. *Hum Immunol* 65 (2004): 381-386.
5. Yurochko AD. Human cytomegalovirus modulation of signal transduction. *Curr Top Microbiol Immunol* 325 (2008): 205-220.
6. McKinney C. Global reprogramming of the cellular translational landscape facilitates cytomegalovirus replication. *Cell Reports* 6 (2014): 9-17.
7. Yu Y, Clippinger AJ, Alwine JC. Viral effects on metabolism: Changes in glucose and glutamine utilization during human cytomegalovirus infection. *Trends Microbiol* 19 (2011): 360-367.
8. Hobman TC (2013). Rubella virus. In Knipe DM, Howley PM, Cohen JI, Griffin DE, Lamb RA, Martin MA, Racaniello VR, Roizman B (ed), *Fields virology*, 6th ed, vol 1. p 687-711. Lippincott Williams & Wilkins, Philadelphia, PA.
9. Lee JY, Bowden DS. Rubella virus replication and links to teratogenicity. *Clin Microbiol Rev* 13 (2000): 571- 587.
10. Frey TK. Molecular biology of rubella virus. *Adv Virus Res* 44 (1994): 69-160.
11. Adler SP. Screening for Cytomegalovirus during Pregnancy. *Infectious Diseases in Obstetrics and Gynecology* 9 (2011): 100-115.
12. Bhide A. Managing primary CMV infection in pregnancy. *BJOG* 115 (2008): 805-807.

13. Alzeidan RA, Wahabi HA, Fayed AA, Esmaeil SA, Amer YS. Postpartum rubella vaccination for sero-negative women (Protocol). *Cochrane Database of Systematic Reviews* (2013).
14. Best JM. Rubella. *Seminars in Fetal & Neonatal Medicine* 12 (2007): 182-92.
15. World Health Organization. Rubella vaccines: WHO position *Weekly Epidemiological Record* No 29 86 (2011): 301-16. 33.
16. Best JM, Banatvala JE. Rubella. In: *Principles and Practice of Clinical Virology*. Eds. Zuckerman AJ, Banatvala JE, Pattison JR, Griffiths PD, Schoub BD), Fifth Edition, John Wiley and Sons, Ltd., West Sussex, England (2004): 427-457.
17. Griffiths PD. Cytomegalovirus (In: *Principles and Practice of Clinical Virology*. Eds. Zuckerman AJ, Banatvala JE, Pattison JR, Griffiths PD, Schoub BD), Fifth Edition, John Wiley and Sons, Ltd., West Sussex, England (2004): 85-122.
18. Hamdan Z, Ismail E, Nasser M and Ishag A. Seroprevalence of cytomegalovirus and rubella among pregnant women in western Sudan. *Virol J* 8 (2011): 217-223.
19. Luis FS, Cosme AE, Jesus HT, Sandra MC, Sergio EM, et al. Seroepidemiology of cytomegalovirus infection in pregnant women in Durango City, Mexico. *BMC Infect Dis* 14 (2014):1471-2334.
20. Hamid KM, Onoja AB, Tofa UA and Garba KN. Seroprevalence of cytomegalovirus among pregnant women attending Murtala Mohammed Specialist Hospital Kano, Nigeria. *Afr Health Sci* 14 (2014): 125-130.
21. Tahani N, Ayda Q, Niveen Sh, Areej AR, Eman AZ, et al. Seroprevalence of Cytomegalovirus among pregnant women and hospitalized children in Palestine. *BMC Infect Dis* 13 (2013): 528-534.
22. Lingqing H, Shu Z, Jie C, Biyun X, Hua Z, et al. Cytomegalovirus Seroprevalence in Pregnant Women and Association with Adverse Pregnancy/Neonatal Outcomes in Jiangsu Province, China. *PLoS One* 9 (2014): 100– 117.
23. Onyenekwe CC, Kehinde-Agbeyangi TA, Ofor US, Arinola OG. Prevalence of rubella-IgG antibody in women of childbearing age in Lagos, Nigeria. *West African Journal of Medicine* 19 (2000):23-26.
24. Barreto J, Sacramento I, Robertson SE et al. Antenatal rubella serosurvey in Maputo, Mozambique. *Trop Med Into Health* 11 (2006): 559-564.
25. Lawn JE, Reef S, Baffoe-Bonnie B, Adadevoh S, Caul EO, et al. Unseen blindness, unheard deafness, and unrecorded death and disability: congenital rubella in Kumasi, Ghana. *Am J Public Health* 10 (2000): 1555- 1561.
26. Bukbuk DN, El nafty AU, Obed JY. Prevalence of rubella-specific IgG antibody in non-immunized pregnant women in Maiduguri, North Eastern Nigeria. *Cent Eur J Public Health* 10 (2002): 21-23.
27. Palhawadan AP, Wickremasingha AR, Perera J. Seroprevalence of rubella antibodies among pregnant females in Sri Lanka. *Southeast. Asian J Trop Med Public Health* 34 (2003): 398-404.
28. Odland JO, Sergejeva IV, Ivaneev MD, Jensen IP, Stray-Pedersen B. Seropositivity of cytomegalovirus, parvovirus and rubella in pregnant women and recurrent aborters in Leningrad County, Russia. *Acta Obstet Gynecol Scand* 80 (2001): 1025-1029.
29. Tolfvenstamt, Enbomm, Ghebrenkh, Rudenu, Lindea, et al. Seroprevalence of viral childhood infections in Eritrea. *J Clin Virol* 16 (2000): 49-54.
30. Singla N, Jindal N, Aggarwal A. The seroepidemiology of rubella in Amritsar (Punjab). *Indian Journal of Medical Microbiology* 22 (2004): 61-63.



This work is licensed under a Creative Commons Attribution Non-Commercial 4.0 International License.