TORCH screening in isolated IUGR: is it really necessary?

Christine Wade¹, Naomi Atkinson², Lisa Hui^{2,3}, Natasha Holmes^{1,3}

Background

The TORCH pathogens (*Toxoplasma gondii*, syphilis, rubella, cytomegalovirus (CMV), and herpes simplex virus (HSV)) can cause congenital infection.¹ However, isolated intrauterine growth restriction (IUGR) is rarely the sole manifestation of intrauterine TORCH infection.

There is limited evidence to support TORCH screening in the setting of isolated IUGR, yet routine screening for TORCH infections in isolated IUGR is common practice.^{2,3}. We aimed to review maternal TORCH screening and determine common indications for screening and evaluate its use in predicting congenital infections.

Methods

Data collection:

A retrospective chart review was conducted at two hospitals in Melbourne, Australia:

Hospital 1

- Specialist maternity centre
- Jan 2014 Dec 2018
- Hospital 2
 - Routine antenatal care
 - Jan 2014 Dec 2017

Eligible pregnant women were identified by extracting TORCH serology requests in pathology databases.

Definitions:

- Positive *screen*: any positive or indeterminate IgM.
- *Confirmed* maternal infection = clinically consistent illness + positive IgM + low avidity IgG
- *Possible* maternal infection = positive IgM + moderate or high IgG, without a consistent clinical illness

Results

- 718 pregnancies (760 fetuses) were reviewed. There were 676 were singleton births and 42 twin births.
- The mean maternal age was 30.2 +/- 5.2y.
- The average gestational age at the time of TORCH investigation was 36 weeks (+/- 4 weeks).
- The total cost of TORCH screens alone was AUD\$65,006.60.
- Zero cases of neonatal TORCH infection were identified in the setting of isolated IUGR (figure 1)

Table 1. Indications for TORCH testing

Indication	Hospital 1 n=425	Hospital 2 n=293	p value
Isolated IUGR	269 (63.3%)	242 (82.6%)	<0.001
Neurological abnormality	44 (10.4%)	5 (1.7%)	<0.001
Echogenic bowel	15 (3.5%)	10 (3.4%)	1.0
Abnormal fluid collection	13 (3.1%)	9 (3.1%)	1.0
Isolated polyhydramnios	29 (6.8%)	10 (3.4%)	0.07
Other US finding	28 (6.6%)	10 (3.4%)	0.089
Maternal illness	17 (4%)	5 (1.7%)	0.126
Other	10 (2.4%)	2 (0.7%)	0.156

Maternal serology results (table 2)

- 49 maternal IgM+ (outcomes in figure 1)
- 34 CMV, 15 *Toxoplasma;* 3 women positive for both
- No rubella IgM+ was detected
- Rubella IgG ordered on 506 women (84.3%) who had known previous immunity to rubella
- At both sites HSV testing was limited to IgG with no IgM performed

for testing

Indication

Isolated IUGR

Neurologic abnormality

Echogenic bowel

Abnormal fluid collection

Isolated polyhydramnios

Other ultrasound findings

Maternal illness Other Total

Department of Infectious Diseases, Austin Health, Melbourne, Australia Department of Obstetrics and Gynaecology, Northern Health, Melbourne, Australia Department of Perinatal Medicine, Mercy Hospital for Women, Melbourne, Australia

Investigations ordered (table 1)

Serology at hospital 2 was available as a single "TORCH screen" request.

Hospital 1 required all tests to be ordered individually. A 'full' panel* was ordered in 63%.of cases. *i.e. all 4 serologies

Table 2. Positive maternal serology by indication

No. IgM+ (% + screens for indication)	95% CI
38 (7.4%)	5.5-10.0%
2 (4.1%)	1.1-13.7%
1 (4.0%)	0.7-19.5%
1 (4.5%)	0.8-21.8%
4 (10.3%)	4.1-23.6%
1 (2.6%)	0.47-13.5%
2 (9.1%)	2.5-27.8%
0 (0%)	0-24.4%
49	

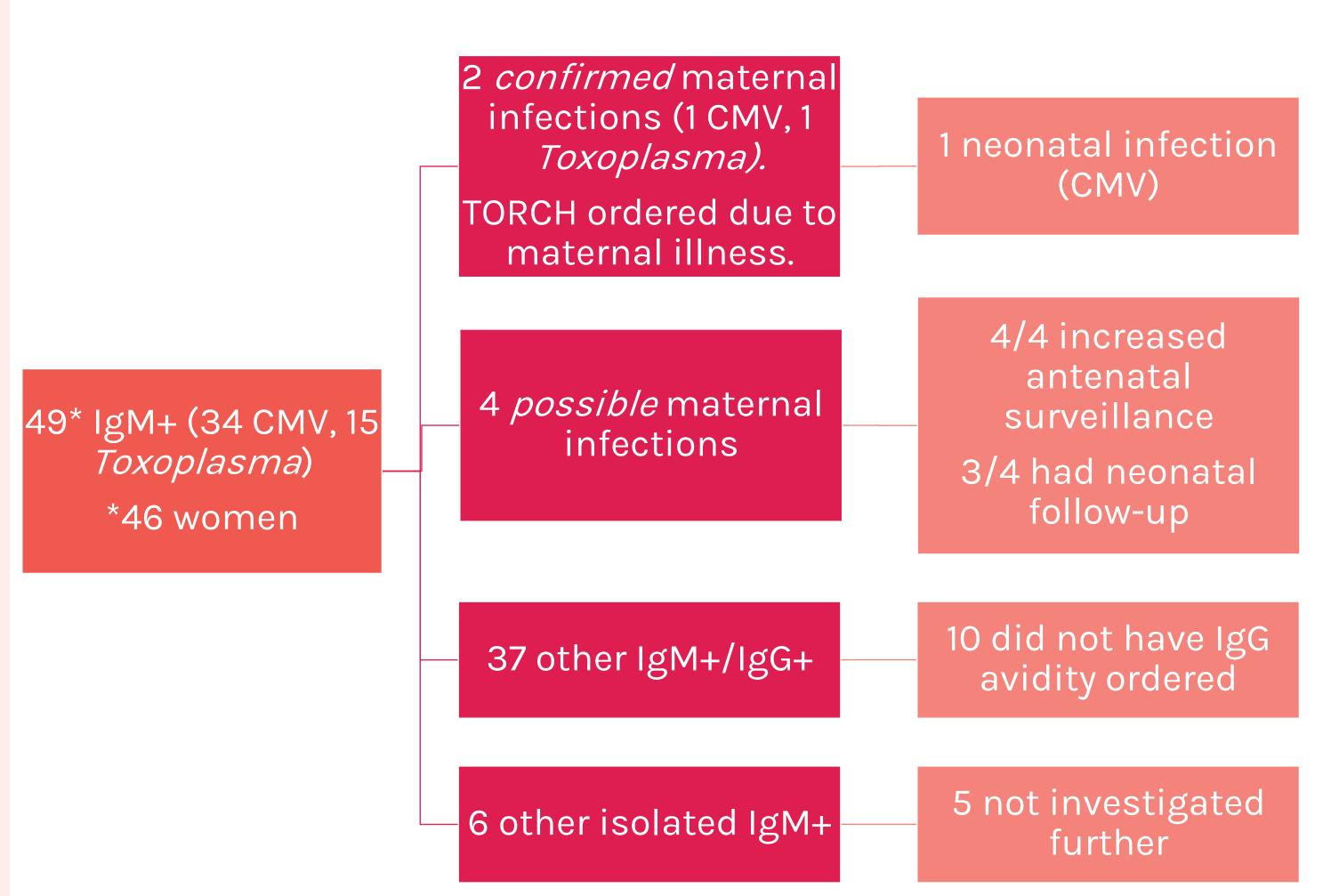


Figure 1. Outcomes of positive maternal IgMs. All six women with confirmed or possible maternal infection had increased antenatal surveillance. One of these women was not referred for neonatal follow-up. There was inconsistent interpretation and investigation of other positive IgM results.

Conclusion

Significant costs were expended to diagnose zer cases of TORCH infection isolated IUGR.

The only cases of materna infection were in women who were symptomatic of these illnesses. The only neonatal infection was in baby born to one of these women.

Investigations were order haphazardly, and positive results were interpreted inconsistently by clinicia with variable referral to a ID physician, maternalfoetal medicine specialis or paediatrician.



ero nin	TORCH screening in isolated IUGR is of no clinical utility and should be abandoned in routine workup of this diagnosis. Clinician education and directed guidelines are needed.
DT	
n a e	References 1. Nahmias AJ. The ToRCH complex-perinatal infections associated with toxoplasma and rubella, cytomegol- and herpes simplex virus. Pediatric Research. 1971;5:405-6.
red e ans, an st,	 2. Sharp A, Duong C, Agarwal U, Alfirevic. Screening and management of the small for gestational age fetus in the UK: A survey of practice. European journal of obstetrics, gynecology, and reproductive biology. 2018;231. 3. Halawa S, McDermott L, Donati M, Denbow. TORCH screening in pregnancy. Where are we now? An audit of use in a tertiary level centre. Journal of obstetrics and gynaecology : the journal of the Institute of Obstetrics and Gynaecology. 2014;34(4).