

# Chorioamnionitis and neonatal morbidity: current perspectives

Gloria Mercedes Galán  
Henríquez  
Fermín García-Muñoz  
Rodrigo

Service of Neonatology, Complejo  
Hospital Universitario Insular  
Materno Infantil de Canarias, Las  
Palmas de Gran Canaria, Spain

**Abstract:** The term chorioamnionitis (CA) is commonly used to refer to different clinical or pathological conditions characterized by an infectious and/or inflammatory process that affects primarily the chorioamniotic membranes, but also the amniotic fluid, vessels of the chorionic plate, and, eventually, the umbilical cord (funisitis) and the fetus. Its incidence is higher at lower gestational ages, and the main mechanism is believed to be the ascending bacterial infection from the maternal genital tract. It can be diagnosed by clinical criteria, amniotic fluid examination for inflammatory mediators, and/or isolation of microorganisms, or by histopathological examination of the placenta. CA is an important cause of stillbirth and is related to an increased incidence of premature rupture of membranes, preterm delivery, and adverse maternal and neonatal outcomes such as early-onset neonatal sepsis and necrotizing enterocolitis. An independent causal association to other neonatal morbidities is more controversial. The heterogeneity in the diagnostic criteria and in the operative definitions for morbidity makes comparison of studies difficult, and results are inconsistent. In addition, the intensity and duration of the process are usually not considered. For all these reasons, evidence-based recommendations for the management of mother and infant under these circumstances are difficult to establish, and clinical practice varies widely.

**Keywords:** clinical chorioamnionitis, histological chorioamnionitis, stillbirth, prematurity, morbidity, mortality

## Introduction

During the past several decades, intrauterine infection has been pointed out as one of the principal causes of premature rupture of membranes (PROM) and preterm birth,<sup>1,2</sup> increasing the risk of fetal lesions, stillbirth, and morbidity and mortality after birth, including neonatal sepsis and pneumonia, respiratory distress syndrome (RDS), intraventricular hemorrhage, periventricular leucomalacia (PVL), and necrotizing enterocolitis (NEC), among others. These morbidities, in turn, have been linked to the development of chronic conditions such as bronchopulmonary dysplasia (BPD) and cerebral palsy (CP) and developmental delay.<sup>3,4</sup> However, a direct causal relationship between CA and these prematurity-related morbidities is still controversial, with some studies showing no effect or even opposite results. The reason for such discrepancies could be the different definitions of CA used in the studies, and the diversity of criteria to include patients. In addition, many of these studies were retrospective, and they reported crude data without considering potential confounders.

## Definitions

CA is an inflammatory condition of the intrauterine environment, frequently of infectious origin, that affects the chorion, amnion, or both.<sup>5,6</sup> It is present in 0.1%–2% of

Correspondence: Fermín García-Muñoz  
Rodrigo  
Service of Neonatology, Complejo  
Hospital Universitario Insular Materno  
Infantil de Canarias, Avenida Marítima  
del Sur S/N, 35016 Las Palmas de Gran  
Canaria, Las Palmas, Spain  
Tel/fax +34 928 444 694  
Email fgarciamu@gmail.com

all pregnancies,<sup>7,8</sup> but in 25%–40% of preterm births,<sup>9</sup> or even 40%–70% in cases of PROM or spontaneous labor.<sup>10</sup> It is more frequent at the lowest gestational age (GA)<sup>10,11</sup> (Table 1), but it also complicates 1%–13% of term births.<sup>6</sup> The diagnosis of CA can be made on clinical, histopathological, biochemical, or microbiological grounds, but disagreement in the criteria for diagnosis is common. A recent survey of US obstetricians found that 26% diagnose CA based only on the presence of maternal fever without taking into account any other clinical signs or symptoms.<sup>12</sup> Frequently the inflammation is subclinical, although histological, biochemical, or microbiological findings can be present. Edwards,<sup>13</sup> in a case series of preterm deliveries, reported a 5%–10% prevalence of clinical CA (CCA), while the prevalence of histological CA (HCA) was >50%. Likewise, in a personal observation (data not published) we also showed a high discrepancy between clinical and histological diagnosis of CA (Table 2).

Clinical diagnosis of CA is most frequently based on criteria adapted from Gibbs,<sup>14,15</sup> consisting of maternal fever >38°C in at least two occasions separated by 1 hour, plus two or more of the following: uterine tenderness defined as pain referred by the mother on abdominal examination in the absence of uterine contractions, leukocytosis (>15,000 cells/mm<sup>3</sup>), maternal tachycardia (>100 bpm), fetal tachycardia (>160 bpm), or foul smelling vaginal discharge. However, the term CCA is mainly used to express clinical suspicion before there is any laboratory or histological confirmation of the inflammation/infection, and there is growing concern about the excessive use of this term. To improve decision-making

in caring for mothers and newborns, and to unify definition criteria, in 2016 an expert panel of the American College of Obstetricians and Gynecologists (ACOG) proposed the use of the term “intrauterine inflammation or infection or both” (Triple I) with the following criteria, maternal fever with one or more of the following: 1) fetal tachycardia (>160 bpm for 10 minutes or longer); 2) maternal white blood cells (WBC) count >15,000 in absence of corticosteroids; 3) purulent fluid from the cervical os (cloudy or yellowish thick discharge confirmed visually on speculum exam to be coming from the cervical canal); and 4) biochemical or microbiological amniotic fluid (AF) results consistent with microbial invasion of the amniotic cavity (MIAC).<sup>5</sup> When there is an objective finding of AF infection or histopathological evidence of infection/inflammation of the placenta, membranes, or the umbilical cord, Triple I is confirmed. Otherwise, it should be categorized as “suspected.” More recently, an ACOG Committee Opinion stated that “the diagnosis of suspected intraamniotic infection is made when the maternal temperature is greater than or equal to 39.0°C or when the maternal temperature is 38.0–38.9°C and one additional clinical risk factor is present,” and “isolated maternal fever is defined as any maternal temperature between 38.0°C and 38.9°C with no additional risk factors present, and with or without persistent temperature elevation.”<sup>16</sup>

The microbiological diagnosis is based on the isolation of the causative microorganism from culture of AF obtained by amniocentesis, but this procedure is seldom performed during labor, when symptomatology usually begins, and the results can be delayed for some days. This makes the

**Table 1** Distribution of clinical and histological chorioamnionitis according to gestational age at delivery

Gestational age (weeks)	23–26	27–30	31–32 <sup>a</sup> or 31–34 <sup>b</sup>	Total	p-value
Clinical chorioamnionitis <sup>a</sup> n/N (%)	653/1,911 (34.2)	713/4,498 (15.9)	114/1,921 (5.9)	1,480/8,330 (17.8)	<0.001
Histological chorioamnionitis <sup>b</sup> n/N (%)	11/15 (73.3)	22/52 (42.3)	4/19 (21.1)	37/86 (43.0)	0.009

**Notes:** <sup>a</sup>Data from García-Muñoz Rodrigo<sup>11</sup>; N=8,330 (in this study, only patients ≤32 weeks gestational age were included); <sup>b</sup>personal observation (data not published), N=86.

**Table 2** Correlation between clinical and histological chorioamnionitis in a series of 84 anatomopathological studies of placentas from gestations ≤34 weeks (personal observation)

		Histological chorioamnionitis <sup>a</sup>		Total
		Present	Absent	
Clinical chorioamnionitis <sup>b</sup>	Positive	7	2	9
	Negative	30	45	75
	Total	37	47	84

**Notes:** <sup>a</sup>According to Redline criteria;<sup>19</sup> <sup>b</sup>adapted from Gibbs criteria;<sup>14</sup> sensitivity = 7/37 = 18.9%; specificity = 45/47 = 95.7%; positive predictive value (PPV) = 7/9 = 77.8%; negative predictive value (NPV) = 45/75 = 60%.

microbiological diagnosis less useful for decision-making during labor. It is estimated that the frequency of positive AF cultures for bacteria in women with labor and intact membranes is 12.8%, and 37.5% of them develop CCA. In contrast, in cases of preterm PROM a positive AF is found in 32.4%, and CCA will develop in 29.7% of them.<sup>17</sup> Another cause of underestimation of the rate of AF infections is that amniocentesis is less frequently carried out in women with oligohydramnios and in those admitted with labor, who usually have higher rates of infection in comparison to women admitted without labor.<sup>18</sup>

From a histological point of view, acute CA is characterized by diffuse infiltration of neutrophils into the chorioamniotic membranes<sup>19</sup> as a result of intra-amniotic inflammation, although not always of bacterial origin. The process begins in the subchorionic fibrin and at the membranous choriodecidual interface, extends to the fibrous chorion and amnion, and finally necrosis of the amniotic epithelium occurs.<sup>20</sup> Then, acute funisitis develops with involvement, in first instance, of the umbilical vein (phlebitis), which exhibits a higher expression of interleukin (IL)-8 mRNA and a pattern of gene expression suggestive of a greater predisposition to a proinflammatory response in comparison to arteries. Thereafter, polymorphonuclear (PMN) cells infiltrate arteries (arteritis), and the concentration of umbilical cord IL-6 increases, which is a marker of severity and is associated with a poorer infant prognosis. Finally, PMN cells infiltrate the Wharton's jelly (funisitis), initially as multifocal rings that subsequently coalesce as the process advances.<sup>21</sup> The origin of the WBC present in the AF of patients with CCA is mostly fetal.<sup>22</sup> An in-depth discussion of the histological aspects and the gradation of the inflammatory process that occurs during chorioamnionitis are beyond the scope of the present work. We refer the reader to the comprehensive chorioamnionitis grading and staging system as proposed by Redline and colleagues.<sup>19</sup> Interestingly, a recent study showed the correlation between MMP-8 concentration in the AF and the severity of HCA.<sup>23</sup>

## Pathophysiology

The principal mechanism of CA is ascending bacterial infection after PROM, but it may also occur with intact membranes. Four stages have been described.<sup>18</sup> In the first one, the normal vaginal and cervical microbial flora changes and/or pathological bacteria appear. Then (second stage) bacteria move toward the lower pole of the uterus, between the membranes and the chorion. Bacteria from the lower genital tract and their products are sensed by pattern recognition receptors, such as toll-like receptors (TLRs). Cervical dilatation and

shortening (cervical incompetence) has also been associated with a higher rate of AF infections, but it is unclear whether such incompetence is the cause or consequence of the infection.<sup>24</sup> In the third stage, bacteria invade the fetal vessels (choriovasculitis) or cross the amnion (amnionitis) into the amniotic cavity, stimulating the production of inflammatory mediators (prostaglandins and reactive oxygen radicals) and different proteases, like MMP. Finally, microorganisms reach the fetus through the respiratory or gastrointestinal tract, or through the mucous membranes (tympanic or conjunctiva). Other possible mechanisms, although less frequent, are hematogenous spread, invasion from the peritoneal cavity via the fallopian tubes, or accidental inoculation of bacteria during invasive procedures.

It is important to note that perinatal infection/inflammation, even if bacteria are not isolated, may lead to the fetal inflammatory response syndrome (FIRS), which is an endocrinological stress response with increased levels of fetal cortisol<sup>25</sup> and fetal cellular and humoral immune activation in response to the exposure to microorganisms and/or their endotoxins, with release of cytokines and chemokines (IL-1, IL-6, IL-8, tumor necrosis factor [TNF]- $\alpha$ , MMP) and upregulation of leucocyte adhesion molecules (ICAM1, E-Selectin, VCAM-1).<sup>26,27</sup> FIRS has been defined as a rise in fetal plasma concentrations of IL-6 >11 pg/mL.<sup>28</sup> Patients with FIRS exhibit funisitis and chorionic vasculitis,<sup>29</sup> and have an increased risk of neonatal sepsis,<sup>30</sup> BPD,<sup>31</sup> and CP.<sup>32</sup> FIRS also affects the hematopoietic system inducing the activation of the mononuclear phagocytic system with neutrophilia and neutrophil and monocyte activation. The elevation of IL-6 stimulates the increase in the number of nucleated red cells.<sup>33</sup> An involution of the fetal/neonatal thymus has also been observed,<sup>34</sup> even in cases of subclinical CA.<sup>35</sup> The grade of thymic involution may be proportional to the duration of the inflammatory process,<sup>36</sup> and it has been associated with lesions of the brain white matter.<sup>37</sup> In addition, the bacterial products and the inflammatory mediators released during FIRS might induce myocardial depression in the fetus.<sup>38</sup> Given its poor ability to maintain cardiac output, this leads to hypotension and cerebral hypoperfusion, which further contribute to brain damage.<sup>39</sup> In the skin, FIRS leads to dermatitis with increased expression of TLR-2 in epidermal keratinocytes of fetuses whose mothers had CA.<sup>40</sup>

## Microbiology

The AF is sterile under normal circumstances. When microbes are detected by culture or, more recently, by molecular techniques,<sup>41</sup> the condition has been termed MIAC.<sup>17</sup> The

prevalence of MIAC varies widely according to the clinical condition (term or preterm birth, intact membranes or not, with or without labor, presence of intra-uterine device, etc.).<sup>42</sup> In a study of women with CCA at term, the detection of microorganisms by conventional cultures and/or molecular techniques was 61%.<sup>43</sup> In this study, intra-amniotic inflammation was defined as a concentration of IL-6 in AF  $\geq 2.6$  ng/mL. Although 54% of the AF cases studied showed both microorganisms detection and inflammation, interestingly, 24% had inflammation without bacterial isolation, 6.5% only bacterial detection without elevation of IL-6, and, finally, 15% neither bacteria nor inflammation. Of note, HCA was significantly more frequent in patients with microbial-associated intra-amniotic inflammation than in those without intra-amniotic inflammation (70.8% [17/24] vs 28.6% [2/7];  $P=0.04$ ).

Frequently, MIAC is polymicrobial, with two or more bacteria isolated in the AF. Genital mycoplasmas (*Ureaplasma urealyticum* and *Mycoplasma hominis*) are the most frequent agents recovered. Other microorganisms include anaerobes (*Gardnerella vaginalis* and *Bacteroides* sp.), *Fusobacteria* sp., aerobes (Group B *Streptococcus*), and Gram-negative rods (*Escherichia coli*). In the rare occasions where pregnancy takes place in women with intrauterine contraceptive devices, these have been pointed out as a risk for *Candida albicans* infection.<sup>44</sup>

## Morbidity

### Preterm birth

Infection is probably the only pathologic process clearly related to prematurity, and the molecular mechanisms involved have been widely studied.<sup>45</sup> It is estimated that ~25%–40% of preterm births are related to intrauterine infections,<sup>2</sup> and the earlier the gestational age at delivery, the higher the frequency of infection, with fetuses of 23 weeks GA colonized in up to 79% of cases in some studies.<sup>46</sup> Furthermore, these figures may well underestimate the actual rate of intra-amniotic infections, since many of the pathogens involved, as mentioned before, do not grow in the usual cultures. The use of molecular microbiology techniques allows the isolation of DNA from bacteria in the AF of women with preterm labor in whom the conventional culture is negative.<sup>47</sup>

Preterm birth can also be triggered by changes in the normal vaginal flora (bacterial vaginosis), but although antimicrobial therapy can eradicate these bacteria, it has not proven to be useful to stop preterm birth,<sup>48</sup> which supports its multifactorial origin.<sup>49</sup> A mechanism of resistance of microorganisms (bacteria and fungi) to antibiotics was pos-

tulated by Donlan and Costerton<sup>50</sup> consisting in the creation of biofilms in which they are encapsulated and protected from those. This type of biofilm has been described, among others, in the urinary epithelium and in the amniotic fluid,<sup>51</sup> but its contribution to the onset of preterm birth is unknown. On the other hand, it is also important to point out that the presence of bacteria and/or inflammatory cells in the fetal membranes does not always cause premature labor.<sup>52</sup>

The inflammatory response associated with CA leads to the rupture of membranes or preterm labor by different mechanisms, including cytokine and MMP activation causing direct rupture of membranes, triggering apoptosis of the fetal membranes cells, activation of the extrinsic pathway in response to toxins and/or cytokines (TNF- $\alpha$ , IL-1 $\beta$ , etc.), or activating cell mediators, such as prostaglandins, that directly induce uterine contractions. A comprehensive review of the role of inflammation and infection in preterm birth can be found in the paper by Romero et al.<sup>53</sup>

## Infection-related complications

Most studies in this field have revealed a positive association between both CCA and HCA and the risk of an early-onset neonatal sepsis (EONS),<sup>11,54–60</sup> and this association persists after adjusting for GA, birth weight (BW), and other confounders. Fetal infection is the most advanced stage of an ascending intrauterine infection, and bacteremia have been detected in 33% of fetuses with positive AF cultures, in contrast to only 4% of fetuses with AF negative cultures.<sup>61</sup> It is estimated that about 1.5% of very low birth weight (VLBW) infants will be affected by an EONS, while 21% of them will develop at least an episode of late-onset neonatal sepsis (LONS).<sup>62</sup> Maternal infections transmitted before or during delivery seem to be the origin of EONS, while LONS are more frequently associated to pathogens acquired from nosocomial environment.<sup>63–65</sup> In cases of PROM, a recent systematic review has shown that maternal administration of broad-spectrum antibiotics prolongs pregnancy, and decreases the risk of chorioamnionitis and neonatal infection and, potentially, its related adverse fetal and neonatal outcomes.<sup>66</sup> Unfortunately, even with the administration of prenatal antibiotics, several studies showed that CA is still related to a 2- to 10-fold increase in the risk of EONS,<sup>11,59,64,65,67</sup> and the risk seems to be higher when funisitis is present.<sup>30</sup> In addition, the 7-year follow-up of children included in the ORACLE II trial found that the prescription of erythromycin for women in spontaneous preterm labor with intact membranes was associated with an increase in functional impairment among their children, and the risk of

cerebral palsy was increased by both erythromycin and co-amoxiclav, although the overall risk was low.<sup>68</sup>

Interestingly and in the opposite direction, two recent studies on CCA and HCA, respectively, have shown a reduction in the risk of LONS.<sup>11,60</sup> The authors speculated that early stimulation of the immune system from intrauterine exposure to less virulent pathogens, rather than to the more aggressive bacteria frequently associated with EONS, or the transfer of maternal cell mediators might be the possible explanation of this phenomenon. However, this hypothesis is now being challenged by the recent finding of an association between CA with aberrant neonatal gut colonization and a significant increase in the risk of the combined outcome of LONS and/or death.<sup>69</sup>

Regarding the future impact of perinatal infection in these patients, it is noteworthy that numerous epidemiological studies have demonstrated an association between systemic inflammation in the fetus/newborn of intrauterine or postnatal origin and brain damage,<sup>70–74</sup> which will be discussed in more detail in the following section.

## Necrotizing enterocolitis

NEC is less common than other complications in the preterm newborn, but remains a relevant problem for the VLBW infant due to its potential severity and its adverse consequences in both the short and long term. Although believed to be multifactorial, the exact mechanism of its development is not fully understood, and its incidence has not changed substantially over time.<sup>75</sup> It affects around 7% of neonates with BW 500–1,500 g, and 11.5% in the subgroup of  $\leq 750$  g.<sup>76</sup> Mortality varies between 20% and 30%, and it is higher in patients requiring surgery.<sup>77,78</sup> Evidence relating CA to NEC is not consistent, but several studies, summarized in the meta-analysis of Been et al,<sup>79</sup> support that CCA is significantly associated with NEC (odds ratio [OR] 1.24, 95% CI 1.01–1.52;  $P=0.04$ ), as well as HCA with fetal involvement (OR, 3.29; 95% CI, 1.87–5.78;  $P\leq 0.001$ ). In addition, a recent study showed that, after adjusting for GA, a high grade of fetal inflammation was significantly associated with chronic lung disease and necrotizing enterocolitis.<sup>80</sup> Of note, in the context of PROM and subclinical infection, it is well known that maternal antibiotic therapy might lessen infectious morbidity and delay labor, but co-amoxiclav should be avoided because of the increased risk of NEC (risk ratio [RR] 4.60, 95% CI 1.98–10.72).<sup>81</sup>

NEC basically consists of ischemic necrosis of the intestinal mucosa with important inflammation and invasion by gas-forming microorganisms, which can dissect the muscular layer

of the mucosa and reach the portal venous system. However, the clinical spectrum varies from this “classic” form to focal intestinal perforations with less inflammatory component that probably have a different etiopathogenesis.<sup>82</sup> The lack of consistent diagnostic criteria makes it difficult to compare results among studies. However, the gradation system of Bell et al in 1978,<sup>83</sup> with some subsequent adaptations<sup>84</sup> despite some possible limitations, has been very useful for decision-making with patients and to compare results between different centers. NEC is the first cause of short bowel syndrome in infancy. Apart from the potentially devastating local effects, the intense inflammatory processes that accompany its course have systemic repercussion, with damage to distant organs such as the brain. This adds to an increased risk of neurodevelopment impairment in this already vulnerable group of patients.<sup>85,86</sup> Martin et al recently reported an increased risk of neurodevelopmental dysfunction and microcephaly in patients with surgical NEC, especially when associated with late bacteremia.<sup>87</sup> Finally, a systematic review aimed at characterizing the neurodevelopment of extremely-low-birth-weight infants ( $<1,000$  g) with NEC and the possible relationship between NEC stage and the neurosensory outcomes found 10 studies comparing patients of the same GA with and without NEC and an average follow-up of 20 months.<sup>88</sup> Globally, 45% of infants with NEC had some form of neurodevelopment impairment. Table 3 summarizes the main results of this study.

## Respiratory morbidity

Again, the relationship between CA and respiratory morbidity in the preterm infants, although extensively studied, remains controversial. Fetuses make breathing movements in utero and may aspirate AF that reaches the distal airway and the alveoli. In Watterberg et al’s study,<sup>89</sup> infants prenatally exposed to HCA had greater concentrations of IL-1 $\beta$  in tracheal aspirates from the first day of intubation and exhibited

**Table 3** Risk of neurodevelopment and neurosensorial impairment in VLBW infants with NEC in comparison to patients of the same GA without NEC

Result	OR	95% CI	p-value
Altered neurodevelopment	1.6	(1.3–2.0)	<0.001
	2.3 <sup>a</sup>	(1.5–3.6) <sup>a</sup>	<0.001 <sup>a</sup>
Cerebral palsy	1.5	(1.2–2.0)	0.001
Visual impairment	2.3	(1.0–5.1)	0.04
Cognitive impairment	1.7	(1.4–2.2)	<0.001
Psychomotor impairment	1.7	(1.3–2.2)	<0.001

**Note:** Data from Rees et al;<sup>88</sup> <sup>a</sup>in patients with Bell’s stage 3.

**Abbreviations:** VLBW, very low birth weight; NEC, necrotizing enterocolitis; GA, gestational age; OR, odds ratio.

less respiratory distress syndrome (RDS), but thereafter they developed more frequently BPD. In animal experimentation, a maturative effect of intra-amniotic administration of IL-1 $\alpha$  has also been observed,<sup>90</sup> with increased expression of surfactant proteins A and B, and lipids. Likewise, Ghezzi et al studied the exposure of fetuses to subclinical intrauterine inflammation and found elevated concentrations of IL-8 in AF of mothers of 24–28 weeks who subsequently developed BPD.<sup>91</sup> All these studies support the hypothesis that HCA could stimulate lung maturation, reducing the incidence of acute RDS but, at the same time, increasing the lung susceptibility to postnatal damage, which finally leads to BPD. The maturational effect of prenatal inflammation may be accompanied not only by structural changes in the lungs,<sup>92</sup> but also by an alteration in the response to exogenous surfactant and the need of prolonged mechanical ventilation.<sup>93</sup> Nevertheless, it is important to remember that prematurity itself is characterized by lungs with a smaller number of alveoli, a slow microvascular development, and thickening of the arteriolar walls. Other mechanisms, such as a neonatal leukemoid reaction induced by CA, have also been suggested as important risk factor for BDP.<sup>94,95</sup> In contrast to HCA, Alexander et al reported an increased incidence of RDS among preterm babies exposed to CCA. Of note, many of these studies were conducted prior to the widespread use of prenatal corticosteroids for fetal lung maturation, exogenous surfactant administration, and/or noninvasive ventilation, which are currently considered standard practices with proven benefit for patients and improved respiratory outcomes.<sup>96</sup> In summary, although unadjusted and adjusted analyses showed that CA was significantly associated with BPD, a recent systematic review and meta-analysis by Hartling et al concluded that CA cannot be definitively considered a risk factor for BPD due to the strong evidence of publication bias with potential overestimation of the association between them<sup>97</sup> (Table 4).

## Patent ductus arteriosus

The relationship between the exposure to CCA or HCA and patent ductus arteriosus (PDA) is again conflicting. Most studies seem to show a positive relationship when crude data are analyzed.<sup>98</sup> In contrast, in our large multicenter study, after adjusting for confounders, we found that the risk of PDA was significantly lower among infants exposed to CCA (adjusted OR 0.831, 95% CI 0.711–0.971;  $P=0.02$ ).<sup>11</sup> However, a recent meta-regression analysis by Behbodi et al<sup>99</sup> concluded that the differences in GA or BW between the CA-exposed and nonexposed groups were significantly correlated with the

effect size of the association between CA and PDA and, consequently, CA appears not to be a risk factor for PDA in preterm infants (Table 4).

## Brain injury

In 1955, Eastman Eastman and Deleon first observed that intrapartum maternal fever increased seven times the risk of CP.<sup>100</sup> Subsequently, this same phenomenon was observed in VLBW infants born to mothers with CA.<sup>101</sup> Since then, numerous studies have tried to delve into the mechanisms by which the infection/inflammation produces brain damage. However, neither clinical trials nor large meta-analyses have yielded homogeneous results.<sup>102</sup> Although germinal matrix HIV is the most frequently acquired brain lesion in VLBW infants diagnosed by cerebral ultrasound (CUS), evaluation at discharge with magnetic resonance imaging (MRI) has shown that brain white matter damage (WMD) is much more common, affecting at least 50% of very preterm infants.<sup>103,104</sup> This injury is probably the leading cause of deficits observed in ~90% of survivors, including CP (5%–10%) and cognitive, attention, and behavioral disorders (about 50%).<sup>105,106</sup> Although other lesions, such as periventricular hemorrhagic infarction, post-hemorrhagic dilatation, or cerebellar lesion, may coexist, WMD seems to be the predominant one.<sup>107</sup> Two major factors are involved in the etiology of WMD, ischemia and infection/inflammation, along with the marked vulnerability of oligodendroglia to oxidative stress. The cerebral vasculature of the preterm infant is characterized by “borderline” zones,<sup>108</sup> where cerebral blood flow (CBF) is more vulnerable to decreases in blood pressure. In addition, sick preterm infants have a lower capacity for CBF autoregulation, making the brain more susceptible to diffuse ischemic damage.<sup>109,110</sup> Hypoxia-ischemia can lead to cytokine elevation even in the absence of infection.<sup>111</sup> Infection causes damage or death of oligodendroglial precursors,<sup>107</sup> mediated by proinflammatory cytokines, especially interferon- $\gamma$  and TNF- $\alpha$ ,<sup>112</sup> and by the activation of the microglia in the immature brain white matter, with the subsequent production of free radicals.<sup>113</sup>

Some clinical and epidemiological studies have shown an association between maternal-fetal infection and PVL (detected by CUS), CP,<sup>114,115</sup> and/or sensorineural involvement (visual impairment, hearing loss, and/or speech delay) during follow-up.<sup>116,117</sup> Among extremely low GA neonates (<27 weeks), Pappas et al showed that infants exposed to CA had a lower GA, and that HCA alone not, but HCA plus CCA increased the risk of cognitive impairment at 18–22 months’ corrected age (adjusted OR 2.00, 95% CI 1.10–3.64).<sup>118</sup>

**Table 4** Characteristics of studies evaluating the association between chorioamnionitis and neonatal morbidity

Author	Year	Study design	Population	N	Outcome	Results: odds ratios or relative risk (95% confidence intervals)	Notes
Been et al <sup>79</sup>	2013	Systematic review and meta-analysis	<37 weeks, CCA	28,490	NEC	OR 1.24 (1.01 – 1.52), <i>P</i> =.49	HCA with fetal involvement: OR 3.29 (95% CI 1.87 – 5.78), <i>P</i> =.12
Hartling et al <sup>97</sup>	2012	Systematic review and meta-analysis	<37 weeks, HCA	15,295	BDP	OR 1.39 (0.95 – 2.04), <i>P</i> =.10	Adjusted for publication bias
Park et al <sup>98</sup>	2015	Systematic review and meta-analysis	VLBW infants, CCA	13,027	PDA	OR 1.89 (1.56 – 2.3), <i>P</i> =.66.2	OR 1.07 (95% CI 0.74 – 1.54)
Behbodi et al <sup>99</sup>	2016	Systematic review and meta-analysis	<37 weeks, CCA	27,186	PDA	OR 1.28 (1.00 – 1.79), <i>P</i> =.64	HCA and CCA
Shatrov et al <sup>116</sup>	2010	Meta-analysis	<37 weeks, HCA	6,484	CP	OR 1.54 (1.10 – 2.15), <i>P</i> =.67	OR 1.75 (95% CI 1.07 – 2.86), <i>P</i> =.71
Shi et al <sup>122</sup>	2017	Meta-analysis	CCA	6,119,978	CP	OR 0.84 (0.70 – 0.91), <i>P</i> =0.0	Meta-analysis of adjusted data appearing in 11/45 studies (72% of infants)
Wu and Colford <sup>71</sup>	2000	Meta-analysis	CCA	1,883	CP	OR 1.21 (0.78 – 1.69), <i>P</i> =.73.7	All GA, most <34 weeks
Mitra et al <sup>131</sup>	2014	Systematic review and meta-analysis	CCA, HCA	10,590	ROP	OR 2.42 (1.52 – 3.84), <i>P</i> =.70.5	All GA, backward approach (CCA in CP, HCA in CP): RR 4.13 (95% CI 2.13 – 7.99) <sup>a</sup> , <i>P</i> =.34
García-Muñoz	2014	Multicenter prospective cohort	VLBW <32 weeks, CCA	8,330	PDA	OR 1.83 (1.17 – 2.89), <i>P</i> =.28.8	RR 1.12 (95% CI 0.98 – 1.28) <sup>a</sup> , <i>P</i> =.67
Rodrigo et al <sup>11</sup>	2009	Multicenter prospective cohort	<33 weeks, CCA	3,094	EONS	RR 1.9 (1.4 – 2.5), <i>P</i> =0.23	All GA, mostly VLBW infants or <32 weeks.
Soraisham et al <sup>127</sup>	2009	Multicenter prospective cohort	<33 weeks, CCA	3,094	EONS	RR 3.0 (2.2 – 4.0), <i>P</i> =0.48	Included also infants born to mothers with fever or infection
					Severe IVH	RR 1.6 (0.9 – 2.7), <i>P</i> =0.20	
					Severe IVH	RR 2.1 (1.5 – 2.9), <i>P</i> =0.87	
					Severe IVH	RR 1.33 (1.14 – 1.55)	
					Severe IVH	OR 0.83 (0.71 – 0.97)	
					Severe IVH	OR 3.10 (2.30 – 4.17)	
					Severe IVH	OR 0.84 (0.72 – 0.98)	
					Severe IVH	OR 1.30 (1.02 – 1.65)	
					Severe IVH	OR 5.54 (2.87 – 10.69)	
					Severe IVH	OR 1.62 (1.17 – 2.24)	

**Notes:** <sup>a</sup>Data available only for CA cases.

**Abbreviations:** CCA, clinical chorioamnionitis; HCA, histological chorioamnionitis; NEC, necrotizing enterocolitis; BPD, bronchopulmonary dysplasia; PDA, patent ductus arteriosus; CP, cerebral palsy; ROP, retinopathy of prematurity; EONS, early-onset neonatal sepsis; LONS, late-onset neonatal sepsis; GA, gestational age; CPR, cardiopulmonary resuscitation; RDS, respiratory distress syndrome; IVH, intraventricular hemorrhage; PVL, periventricular leukomalacia; VLBW, very low birth weight; *P* and *P*, heterogeneity; RR; risk ratio; OR, odds ratio; CRIB I, Clinical Risk Index for Babies I.

However, as stated before, CA is strongly associated with preterm birth and, after adjusting for GA, the association between CA and PVL decreases or disappears. For this reason, a causal relationship between them has been questioned by some.<sup>119</sup> Yliloki et al recently reported the lack of association between CCA and neurodevelopmental problems in very preterm infants.<sup>120</sup> In contrast, HCA was associated with a slightly less optimal performance at 5 years of age, although, in their opinion, further studies are needed to verify the clinical significance of these findings. Previously, in a systematic review by the same group that included 84 original papers, they concluded that the majority of the studies did not support the hypothesis that CA poses a direct risk on the central nervous system of preterm infants.<sup>121</sup> Similar conclusions seem to be derived from most recent reviews.<sup>122,123</sup>

An additional problem when evaluating the relationship between CA and WMD is that most of the initial studies, especially epidemiological ones, referred to cystic PVL, which is currently estimated to contribute to less than 10% of WMD cases. Modern technologies could provide new insight into the subject. In a recent study, magnetic resonance spectroscopy in healthy term newborns exposed to maternal CA showed evidence of occult neuroinflammation, and the author suggested that this finding might be associated to neurodevelopmental outcomes at 12 months.<sup>124</sup> Another study with MRI in patients exposed to CA revealed long-term alterations in brain morphology in areas related to cognitive and motor functions, leading to psychiatric and neurodevelopmental disorders, although this study was limited by the small sample size.<sup>125</sup> In summary, to date, the relationship between CA and neurodevelopmental outcomes is not completely understood, and the studies' results might be influenced by variables that covariate with or mediate the effect of the other.

## Mortality

Perinatal mortality related to maternal CA has been addressed in several studies, but results are still conflicting. Elimian et al found that HCA was independently associated with neonatal death, although in exposed infants the administration of antenatal steroids significantly reduced morbidity and mortality.<sup>126</sup> The risk of neonatal mortality seems to be higher when there is evidence of fetal inflammation (funisitis and/or fetal vessel angiitis).<sup>127</sup> In a population-based study of full-term infants born in the United States during 2008, the prevalence of CCA was 9.7 per 1,000 live births, and the neonatal mortality rate was higher among exposed infants (OR 1.72, 95% CI 1.20–2.45).<sup>7</sup> In contrast, several studies on CCA or HCA in VLBW infants, after correcting for confounders, showed no

effect on mortality<sup>11,60</sup> or, on the contrary, they showed a protective effect.<sup>128,129</sup> In the study by Lahra et al, both HCA and a histological fetal response to CA were observed to be more common in preterm survivors of the neonatal period.<sup>130</sup> In the same direction, Pappas et al found that HCA was associated with a reduced risk of death or neurodevelopmental impairment (adjusted OR 0.72, 95% CI 0.56–0.93).<sup>118</sup>

## Conclusion

Although some experimental evidence in animal models and the results of human research have demonstrated or suggest adverse effects of intra-amniotic infection/inflammation in the fetus/newborn, controversy still persists regarding a direct relationship between maternal CA and most of the neonatal outcomes studied. Table 4 summarizes the most relevant systematic reviews and meta-analyses. To date, the association between CA and preterm birth and EONS and NEC seems to be clear. Prematurity itself is a major risk factor for many different morbidities known to be of multifactorial origin. Prenatal infection/inflammation or both could have a modulating effect over them in both directions, sometimes increasing the final risk while sometimes reducing it, as it seems to be the case for RDS, for instance. The heterogeneity of the studies and the variety of clinical situations and management of patients in a changing field of knowledge (prenatal steroids, maternal antibiotics, postnatal surfactant administration, noninvasive ventilator support, etc.) precludes establishing a definitive relationship between CA and a particular morbidity. In the opinion of the authors, as new diagnostic tools develop and as new strategies of patient management are implemented, new associations could be detected or, on the contrary, some current beliefs will vanish.

## Disclosure

The authors report no conflicts of interest in this work.

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