1	Preeclampsia Treatment in SARS-CoV-2
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We have all faced unprecedented challenges caring for pregnant women in the SARS-CoV-2 pandemic		
ith limited experience and rapidly evolving guidelines. We took great interest in AJOG's re	ecent Boelig	
al "Labor and Delivery Guidance for COVID-19" (1). They note a paucity of experience v	with	
agnesium for neuroprotection or seizure prevention in patients with SARS-CoV-2. Given p	ootential	
spiratory complications associated with magnesium sulfate, there is a theoretical concern the	hat	
eatment could exacerbate SARS-CoV-2 infection.		
Ve present the first reported case of management of severe pre-eclampsia with known mater	rnal SARS-	
oV-2 infection, including magnesium sulfate administration.		
26-year-old woman at 37 weeks gestation diagnosed with SARS-CoV-2 for symptoms of		
nd "allergies" was also diagnosed with pre-eclampsia based on sustained elevated blood pre-	essures	
140/90 and proteinuria.		
trapartum, she reported dyspnea and a sensation of "drowning", although she maintained of	oxvgen	
turation greater than 97% on room air and lung exam was clear to auscultation bilaterally v		
rackles or wheezes. She began to experience sustained severe range blood pressures of 175/		
66/101 with mild headache. Serum labs were notable for AST 131 U/L, ALT 133 U/L, crea		
g/dL, and platelets 199 k/uL. Thromboelastography (TEG) notable for increased platelet at		
brinogen activity. There was brief pause for consideration if intravenous labetalol could be		
atients with SARS-CoV-2, given the recommendation to avoid with reactive airway disease	_	
F bronchoconstriction (2,3). Similarly, a quick literature review was conducted regarding m		
alfate infusion in this at-risk patient population given its possibility to worsen respiratory st	atus (4).	
iven normal oxygenation and benign lung exam, the decision was made to manage severe-	range blood	
ressure with standard first-line agent of 20mg of intravenous labetalol. Next, a loading dose	e of 4g	
travenous magnesium sulfate was initiated for seizure prevention, followed by a maintenar	nce rate of	
g per hour infusion. Her blood pressure improved to 147/85 and remained on average 130s/	/80s	
llowing these interventions, and portable AP chest x-ray revealed no acute cardiopulmonal	ry	
ocess. The patient had no reported exacerbation of pulmonary symptoms during magnesium	ım sulfate	
lministration and was able to maintain oxygen saturation greater than 97% on room air dur	ing	
eatment.		

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70 She progressed to 10cm cervical dilation and pushed for 120 minutes with a Category 2 fetal heart 71 tracing due to recurrent variable decelerations with slow return to baseline, with subsequent 72 uncomplicated forceps-assisted vaginal delivery for fetal indication and maternal exhaustion. 73 74 She delivered a vigorous male infant weighing 3042g with Appar scores of 7 and 9 at 1 and 5 minutes. Delayed cord clamping was performed without placing infant skin-to-skin. The awaiting Pediatrics team 75 76 took infant to the NICU for assessment where SARS-CoV-2 testing resulted negative. The patient declined separation from her infant; therefore, the infant remained in her postpartum isolation room in a 77 bassinet six-feet away from bed. The patient initially hand-expressed then moved to breastfeeding after 78 washing her hands well and while wearing a mask. The infant was incidentally noted to have penile 79 80 torsion and was referred to outpatient Pediatric Urology. 81 82 Blood pressures remained mild-range following delivery, and intravenous magnesium sulfate therapy at 83 a maintenance rate of 2g per hour was continued for 24 hours following delivery. After evaluation by dedicated SARS-CoV-2 ICU team, the patient did not meet inclusion criteria for clinical trial or 84 85 compassionate use of Remdesivir given clinical stability. She was immediately ambulatory after delivery thus we elected against VTE pharmacoprophylaxis in favor of mechanical prophylaxis. The 86 87 patient was discharged home on postpartum day two with no symptoms suggestive of SARS-CoV-2 infection, and did not require oral medication for blood pressure control. 88 89 There is currently a lack of data regarding the safety of magnesium sulfate administration in patients 90 91 with SARS-CoV-2 infection. In this case, the patient had mild respiratory symptoms with normal oxygenation on room air and with normal clinical exam and chest x-ray. Given severely elevated blood 92 93 pressures with headache in the setting of pre-eclampsia, the decision was made to proceed with magnesium sulfate administration. We observed that this patient was able to tolerate a 4g magnesium 94 95 sulfate loading dose followed by 2g/hour maintenance rate without issue. Additionally, there was concern for administering intravenous labetalol for blood pressure control given the possibility of 96 97 respiratory compromise; this drug was fortunately administered without adverse consequence and successfully lowered blood pressure with one 20mg dose. Our limited clinical experience supports the 98 authors' expert opinion that "Magnesium sulfate may be used as indicated in patients with 99

mild/moderate symptoms" in SARS-CoV-2 infection.

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