Purpose of this report

The purpose of this report is to provide feedback to the neonatal-perinatal medicine community regarding content areas of strength and weakness, information which may be useful for identifying potential gaps in knowledge and guiding the development of educational materials. Using data from the American Board of Pediatrics' (ABP) Maintenance of Certification Assessment for Pediatrics (MOCA-Peds), this report summarizes diplomate performance on the questions within each of the 47 content areas assessed in 2020.

MOCA-Peds content areas

In 2020, MOCA-Peds—Neonatal-Perinatal Medicine consisted of questions from a total of 47 content areas, broken down as follows:

- 45 learning objectives¹ Each diplomate initially received one question from each of the 45 specific content areas drawn from the neonatal-perinatal medicine content outline.
- Two featured readings¹ Each diplomate also received two questions per featured reading (eg, clinical guidelines, journal articles) for a total of four featured reading questions.

A pool of questions was developed for each learning objective and for each featured reading. Questions were then drawn from the pool and administered to diplomates throughout 2020 according to the specifications described in the bulleted list above.

Understanding this report

This report provides a graphical summary of diplomate performance on each of the 47 content areas assessed in 2020. Within the graphic and in the example below, the point (•) reflects the average percent correct for all questions within that learning objective or featured reading. The bar (—) reflects the range of percent correct values for the questions within that learning objective or featured reading. More specifically, the bar's lower endpoint indicates the most difficult question (ie, answered correctly by the lowest percentage of diplomates) and the bar's upper endpoint indicates the easiest question (ie, answered correctly by the highest percentage of diplomates).



¹Each diplomate also received 15 "repeat" questions selected from their original subset of learning objective and featured reading questions. Performance on the repeat administrations is not included in this report.

A note of caution

Many factors (eg, specific content of the question, wording of the question, plausibility of the incorrect answers) can impact diplomate performance on any question. It is thus difficult to determine if poor performance on a single question, or small set of questions, within a given content area reflects a true gap in diplomate knowledge or if the question(s) associated with that content area were difficult for other reasons (or some combination of both). Collectively, the entire set of MOCA-Peds questions (across all content areas) constitutes a psychometrically valid assessment of the diplomate's overall level of knowledge. Performance within a given content area is based on fewer questions, however, and is therefore less useful for making inferences about diplomate knowledge in that specific content area.

It is important to note again that for security reasons, a pool of questions was developed for each content area so that each diplomate received a unique set of questions. In addition, the number of questions can vary from one content area to the next. In cases where a content area had a relatively large pool of questions, the number of diplomates who answered each question was reduced, which diminished the statistical precision of each question's percent correct value. In cases where a content area had a relatively small number of questions, each question was answered by a larger number of diplomates, but the overall breadth of the content being assessed within that content area was constrained, which limits the generalizability of the results.

In other words, MOCA-Peds was designed to assess individual diplomates with respect to their overall level of knowledge in neonatal-perinatal medicine. It was not designed to provide the pediatric community with diagnostic feedback pertaining to specific content areas within neonatal-perinatal medicine. The results within this report may be informative and useful for that secondary purpose, but they should be interpreted with a degree of caution.

Additional notes

- To protect the security of the content of the assessment, the questions themselves, along with information about the number of questions in the pool for any particular learning objective or featured reading, are not provided in this report.
- This report contains data aggregated across many diplomates participating in the MOCA-Peds program and cannot be used to make inferences or draw conclusions regarding any particular diplomate.

2020 Content Area Feedback Report Neonatal-Perinatal Medicine

	Learning Objective	0	Perc 25	ent Coi	rect	10
Learning Objective 1. Recognize and understand the strengths and limitations of a cohort study, case control study, and	<u> </u>		-		_	
ra	ndomized controlled clinical trial.				•	
va	evelop a management plan for a preterm or term infant exposed to a communicable disease such as ricella (using knowledge of placental immunoglobulin transfer).				-	-
ОХ	now how to interpret arterial blood gas measurements and noninvasive methods for estimating arterial rygenation.				-	_
	ecognize the association of cholestasis with parenteral nutrition and describe its management.				-	
Kr	ormulate a differential diagnosis for a cyanotic newborn infant. now how to set up and conduct a quality improvement (QI) project, and how to determine whether it is accessful or not.		:	:	•	
	andomized Trial of Platelet–Transfusion Thresholds in Neonates (Featured Reading)					_
Kr	now the risk factors for development, proposed mechanisms, diagnosis, and consequences of traventricular hemorrhage in the preterm infant.			-		Þ
	now the treatment and complications of group B streptococcal infections.					•
	eview the evaluation and management of common arrhythmias in the fetus and newborn infant.				-	•
l. Id	entify the clinical, laboratory, and imaging features of neuroblastoma in the newborn infant.				-	•
2. De	escribe best practices for management of a newborn with persistent pulmonary hypertension (PPHN).					•
B. Fo	ormulate an evaluation and management plan for a neonate with acute kidney injury.		:		:	•
I. Co	ompare the indications for and limitations of various neuroimaging studies; recognize normal and				•	_
ab	onormal findings that occur during development and after brain injury. Sociation Between Oxygen Saturation Targeting and Death or Disability in Extremely Preterm Infants in		:	:	- : _	•
	e Neonatal Oxygenation Prospective Meta-analysis Collaboration (Featured Reading)		- :			
	ecognize the significance of abnormalities in fetal heart rate patterns during labor.					4
	scuss current recommendations regarding suctioning meconium from the newborn infant airway.					-0
	nderstand the fetal/neonatal significance of oligohydramnios or polyhydramnios.					-
	evelop an evaluation and management plan for an infant whose newborn screen was positive for congenital					-
	rpothyroidism.		- :		- :	
m	ecognize the effects of environmental factors (for example: maternal bonding, socioeconomic conditions, edia exposure) on infant outcomes.				-	-
ar	cplain the clinical and echocardiographic features of a physiologically significant patent ductus teriosus in a preterm neonate.					1
	escribe the indications for and proper administration of supplemental oxygen immediately after birth.					_
he	ecognize the skin manifestations of common neonatal infections including cytomegalovirus, candidiasis, erpes, staphylococcus. ecognize the effects of fetal programming and nutrition on the prevalence and types of adult onset					4
	sorders.					4
	elate the protein requirements of preterm and term infants to parenteral and enteral nutrition uidelines.					4
	emonstrate an understanding of inheritance patterns and recurrence risks for autosomal dominant sorders.					•
	now the factors associated with increased bilirubin production and decreased serum bilirubin excretion the neonate.					•
	now the maternal factors, incidence, clinical manifestations, common complications, and prognosis of own syndrome.					4
	escribe an evaluation and management plan for an infant with suspected choanal stenosis/atresia.					1
). Re	ecognize the clinical features of a neonate with a left-sided cardiac obstructive lesion.					-(
se	fferentiate the clinical manifestations of disorders of amino acid metabolism from other causes of rious neonatal illness.					
us	escribe indications for use, clinical effects, side effects, and toxicity for classes of drugs commonly sed in the neonate (eg, antibiotics, analgesics, anticonvulsants).					-
CC	nderstand the clinical manifestations, diagnostic criteria, treatment, and complications of exactivities, echovirus, and enterovirus infections.					
be	now the evolution of neurodevelopmental impairments over time, and when and with what tools they are set assessed.					
	now the indications for phototherapy and exchange transfusion in the preterm and term neonate.				:	
ge	evelop a management plan for an infant born to a mother with active genital herpes or with a history of enital herpes.					
as	ecognize the characteristics and consequences of congenital defects with non–genetic etiologies (such amniotic band syndrome).					
	escribe the prognosis and long-term complications of bronchopulmonary dysplasia.					
	cplain the pathophysiologic effects of maternal diabetes on the newborn.				:	
	st the most effective therapeutic interventions for decreasing insensible water loss.					
. Kr	scuss best practices for management of a newborn infant with respiratory distress syndrome (RDS). now the clinical and diagnostic features, evaluation, management, and complications of necrotizing					
	terocolitis.					
	ecognize the neonatal systemic complications of perinatal asphyxia.		:		:	
	now the causes, diagnosis, management, and outcomes of hypoxic-ischemic encephalopathy. I compare the indications for and techniques, effects, and risks of extracorporeal membrane oxygenation					
(E	CMO). nalyze the potential causes and treatment options for neonatal anemia in preterm and term neonates.					
. Ar						