Monitoring intra-abdominal pressure : should it be a routine for critically ill?

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Abdomen was deemed a neglected compartment¹ but not anymore! The consensus definitions², clinical practice guidelines³, and recommendations for research⁴ reflect a decade of sustained efforts of World Society of the Abdominal Compartment Syndrome (WSACS) to bring intra-abdominal hypertension (IAH) and the abdominal compartment syndrome (ACS) in the spotlight. Clinical care and outcomes of ACS have improved over the last decade along with an exponential growth in research⁵. It is now known that the normal or mean intra-abdominal pressure (IAP) within the non-diseased abdominal cavity is between 2 and 5 mmHg but can run as high as 12 mmHg in the obese adult without causing any organ dysfunction. Critically ill patients are almost always fluid overloaded and edematous, therefore an IAP of 5 to 7 mmHg is considered normal. Intraabdominal hypertension (IAH), is defined as a sustained or repeated pathologic elevation of the IAP \geq 12 mmHg. When the sustained IAP >20 mmHg is associated with new onset of organ dysfunction or failure it is labeled as Abdominal Compartment Syndrome (ACS). An abdominal perfusion pressure (APP) of at least 60 mmHg is desirable to maintain adequate perfusion to the viscera and prevent end-organ dysfunction (APP = MAP - IAP). (MAP=Mean Arterial Prsssure)

IAH occurs in 20% to 40% of intensive care patients⁶ depending upon the sample population but there is a possibility that it is under reported because physicians ascribe the organ dysfunction to progression of the primary disease rather than labeling it as IAH or ACS. Measuring IAP is the key to understanding the evolution from IAH to ACS and avoids or reduces the harmful effects of raised IAP by timely interventions7. The Sensitivity of physical examination to detect high IAP is only 40% to 60%, therefore it is not considered reliable ^{8,9}. Measuring vesical pressure transmitted through a Foley catheter following the WSACS guidelines is the Gold standard method of determining IAP10. However measuring IAP has not become a standard of care in ICUs6. One survey showed that 25% physicians had never measured bladder pressure, and bladder pressure was measured routinely by only 31%^{11,12}. Most physicians measure IAP only when one or more risk factors for ACS are present⁶. Identifying the risk factors for developing ACS is challenging because there is a long list of possible clinical conditions and predisposing factors associated with IAH² and it does not help the physicians in decision-making as to which patients will benefit from early IAP measurement. The most substantiated independent risk factors¹³ for development of ACS are massive fluid resuscitation (> 3.5-5 L/24 hours), multiple transfusions (>10 U packed RBC 24 hours), hypothermia (core temperature $< 33^{\circ}$ C), base deficit/ acidosis (pH <7.2), and BMI >30. Mechanically ventilated patients with sepsis, severe burns, severe trauma, severe acute pancreatitis, major abdominal surgery, ruptured aortic aneurysm repair or liver failure also represent a high risk population⁶. *Routine IAP measurement in patients with these well-defined risk factors is strongly recommended in order to recognize IAH early, institute timely interventions and halt progression to ACS*.

As measuring IAP requires training and additional expense even if the specially designed disposable equipment is not used, routine measurement in all patients is not feasible. Risk of developing IAH is minimal in mechanically ventilated patients with a positive end-expiratory pressure (PEEP) < 10 cm H₂O, PaO₂/FiO₂ > 300, BMI < 30 kg m-2 and without pancreatitis, hepatic failure/cirrhosis, gastrointestinal bleeding or laparotomy and the use of vasopressors/inotropes on admission¹⁴. In these patients, IAP measurement is not mandatory. ICU patients in whom IAP measurements are not done on admission should be closely monitored clinically for signs of IAH and there should be a very low threshold for starting IAP monitoring. Developing local checklists might be the first step in decision making process for IAP monitoring.

Routine measurement of IAP in all critically ill patients cannot be currently recommended and stronger evidence is required to develop clear guidelines to identify patients in whom IAP measurement is mandatory. However routine clinical monitoring for signs IAH in all critically ill patients and a low threshold to frequently measure bladder IAP in patients with known risk factors is strongly recommended to diagnose IAH before it progresses to ACS.

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