Acute kidney injury in ICU patients, relation with intraabdominal hypertension

Ehesis

Submitted for Partial Fulfillment of Master Degree in **Internal Medicine**

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List of Abbreviations

ACEIs Angiotensin-Converting Enzyme Inhibitors

ACS Abdominal Compartment Syndrome

ADH Antidiuretic Hormone

ADQI Acute Dialysis Quality Initiative

AKI Acute Kidney Injury

AKIN Acute Kidney Injury Network

ALT Alanine Transferase

ANCA Anti-Neutrophil Cytoplasmic Antibody

APP Abdominal Perfusion Pressure

ARBs Angiotensin II Receptor Blocker.

ARF Acute Renal Failure

ATIN Acute Tubulointerstitial nephritis

ATN Acute Tubular Necrosis

BUN Blood Urea Nitrogen

CBC Complete Blood Count

CGN Crescentic Glomerulonephritis

CNS Central nervous system

CrCl Creatinine Clearance

CRRT Continuous Renal Replacement Therapy

CT Computed Tomography

DIC Disseminated Intravascular Coagulopathy

FENa fractional excretion of sodium

FG Filtration Gradient

GBM Glomerular Basement Membrane

GFR Glomerular Filtration Rate

GN Glomerulonephritis

Gro-α Human Growth-Related Oncogene-α

HES Hydroxyl-Ethyl Starch

HSP Henoch-Schonlein Purpura

HUS Hemolytic Uremic Syndrome

IAH Intra-abdominal Hypertension

IAP Intra-abdominal Pressure

ICNARC Intensive Care National Audit and Research

Centre

ICU Intensive Care Unit

IL-18 Interleukin 18

IVP Intra-vesical pressure

KC Keratinocyte-derived chemokine

KDIGO Kidney Diseases Improving Global Outcomes

KIM-1 Kidney Injury Molecule-1

KUB Kidneys, ureter bladder

LDH Lactate Dehydrogenase

MAP Mean arterial blood pressure

MCP-1 Monocyte chemo attractant protein-1

MM Multiple Myeloma

MRA Magnetic Resonance Angiography

MPGN Membrano-Proliferative Glomerulonephritis

NGAL Neutrophil Gelatinin-Associated Lipocalin

NHE3 The sodium/hydrogen exchanged isoform 3

NKCC2 Sodium/potassium chloride cotransporter

NSAIDs Non-Steroidal Anti-Inflammatory Drugs

NV Necrotizing Vasculitis

PG Prostaglandin

PIGN Postinfectious Glomerulonephritis

PSGN Post-Streptococcal Glomerulonephritis

RPGN Rapid Progressive Glomerulonephritis

RRT Renal Replacement Therapy

SCr Serum Creatinine

SIRS Systemic Inflammatory Response Syndrome

SLE Systemic Lupus Erythematosus

SPSS Statistical Package for Special Science

TMA Thrombotic Microangiopathy

TTP Thrombotic Thrombocytopenic Purpura

UO Urine Output

WHO World Health Organization

WSACS World Society of the Abdominal Compartment

Syndrome

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Introduction

considered mostly a post-surgical condition, intra-abdominal hypertension (IAH) and the abdominal compartment syndrome (ACS) are now thought to increase morbidity and mortality in many patients receiving medical or surgical intensive care. Animal data and human observational studies indicate that oliquria and acute kidney injury are early and frequent consequences of IAH/ACS and can be present at relatively low levels of IAP. Among medical patients at particular risk are those septic shock and severe pancreatitis, but the adverse effects of IAH may also be seen in cardiorenal and hepatorenal syndromes. Factors predisposing to IAH/ACS include sepsis, large volume fluid resuscitation, polytransfusion, mechanical ventilation with high intrathoracic pressure, and acidosis, among others.(4)

Physical examination, with sensitivity of 40 to 60.9%, is unreliable for diagnosing IAH/ACS.(8)

Although a variety of direct and indirect techniques for measuring IAP have been developed and validated over the years,(9) transduction of urinary bladder pressure through an indwelling urinary catheter remains

the gold standard for measuring and monitoring IAP. The bladder pressure method has been shown when performed appropriately to strongly correlate with IAP measured directly, (1) while remaining cost-effective and safe, without any increased risk of catheter-associated urinary tract infection.(2)

In many institutions, screening patients at risk of developing IAH and serial monitoring of IAP every 4 to 6 hours is common practice.(3)

Several non-surgical methods can help reduce IAP. The role of renal replacement therapy for volume management is not well defined but may be beneficial in some cases. IAH/ACS is an important possible cause of acute renal failure in critically ill patients and screening may benefit those at increased risk.(4)

In recent years both IAH and ACS —the severe, advanced stage of IAH characterized by organ system failure— have increasingly been associated with morbidity and mortality in critically ill patients. (6)

Although multiorgan failure is also well recognized in ACS, what is much less appreciated and what some recent data suggest is that kidneys may be particularly at risk with much lower levels of IAP than would be seen in fully established ACS. These findings indicate that acute renal failure (ARF) resulting, at least in part, from lesser degrees of IAH may be present in a much larger population of critically ill patients than believed previously. (7)