

Joint Theater Trauma System Clinical Practice Guideline

UROLOGIC TRAUMA MANAGEMENT

Original Release/Approval	18 Dec 2004	Note: This CPG requires an annual review.	
Reviewed:	Jun 2010	Approved:	30 Jun 2010
Supersedes:	Urologic Trauma Management, 7 Nov 2008		
<input type="checkbox"/> Minor Changes (or)	<input type="checkbox"/> <i>Changes are substantial and require a thorough reading of this CPG (or)</i>		
<input checked="" type="checkbox"/> Significant Changes	Exposure of the kidney in trauma is best performed lateral to medial vice obtaining vascular pedicle control prior to mobilization		

- 1. Goal.** To provide guidance for management of genitourinary (GU) trauma in combat casualties.
- 2. Background.** GU trauma accounts for approximately 5% of all combat casualties. As with all operative management, treatment of these injuries adheres to established surgical principals of hemostasis, debridement, and drainage. Whenever possible, proper radiographic evaluation of the GU system should be undertaken prior to operative intervention. For far forward surgical units, the preservation of as much tissue as possible – particularly when dealing with the external genitalia - is indicated, followed by rapid evacuation to a higher level of care where definitive urological management can be undertaken.
- 3. Recommendations.** See [Appendix A](#).
- 4. Responsibilities.** It is the trauma team leader's responsibility to ensure familiarity and appropriate compliance with this CPG.
- 5. Reference.**

¹ *Emergency War Surgery Handbook*

Approved by CENTCOM JTTS Director and Deputy
Director and CENTCOM SG

Opinions, interpretations, conclusions, and recommendations are those of the authors
and are not necessarily endorsed by the Services or DoD

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APPENDIX A

	GENERAL MANAGEMENT PRINCIPLES
LABS	CBC, Chem 10, PT, PTT, UA. Type and Screen or Type and Cross x 4 units.
HEMATURIA	<ul style="list-style-type: none"> • During trauma evaluation, place foley catheter unless contra-indicated. Perform RUG first if blood at the meatus, high riding prostate or other evidence of urethral injury. RUG - Obtain a KUB plain film first, then a 14-16 Fr foley, primed with contrast to rid air, is placed in the urethra past the balloon. Use 1-2 ml saline to fill the balloon snugly in the fossa navicularis. A pelvic film in a semi-lateral position is obtained after injecting approximately 30ml of straight contrast (con-ray) under steady, gentle pressure. Study is considered normal only if contrast enters the bladder without any extravastaion. • If anterior urethral injury, plan to repair in OR. If posterior urethral injury, attempt to gently place a foley catheter. If unable, then place supra-pubic tube in EMT or in OR. • If catheter passes, and gross hematuria noted, proceed with GU diagnostic evaluation for bladder injury or a renal/ureteral source. CT scan with delayed images + a CT cystogram is ideal imaging study (see technique description following).
RENAL INJURY Penetrating renal injury = Abdominal exploration	<ul style="list-style-type: none"> • Blunt Trauma: All patients with gross hematuria (regardless of initial SBP) <i>and</i> those patients with microscopic hematuria <i>whose initial SBP is less than 90 mmHg</i> should undergo contrast enhanced CT scan <i>if/when they become hemodynamically stable.</i> • <u>Renal Injury Grading</u> <u>Grade 1:</u> Sub-capsular hematoma <u>Grade 2:</u> Small parenchymal laceration <u>Grade 3:</u> Deeper parenchymal laceration without entry into collecting system <u>Grade 4:</u> Laceration into collecting system with extravasation; vascular injury with contained hemorrhage <u>Grade 5:</u> Shattered kidney or renal pedicle avulsion • Hemodynamically stable patients can usually be managed without operation. • Vascular repair is indicated for salvageable kidneys with renal artery or vein injury (<i>see vascular CPG for more details</i>). • Ureteral stent may need to be placed for persistent urinary extravasation.

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June 2010

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RENAL EXPLORATION DURING ABDOMINAL OPERATION	<ul style="list-style-type: none"> • Absolute indications: persistent bleeding or expanding/pulsatile hematoma • Relative indications: urinary extravasation, nonviable tissue (> 20%), and segmental arterial injury on pre-op study. • Urinary extravasation from a grade IV parenchymal laceration or forniceal rupture can be managed nonoperatively in most patients.
RENAL REPAIR AND PARTIAL NEPHRECTOMY PRINCIPLES	<ul style="list-style-type: none"> • Complete renal exposure, debridement of nonviable tissue, hemostasis by individual suture ligation of bleeding vessels, watertight closure (absorbable suture), drainage of the collecting system, and coverage/approximation of the parenchymal defect. Perform partial nephrectomy if reconstruction not possible: the collecting system must be closed and the parenchyma covered with omentum. • Place ureteral stent for persistent urinary extravasation
NEPHRECTOMY	<ul style="list-style-type: none"> • Total nephrectomy is immediately indicated in extensive renal injuries when the patient's life would be threatened by attempted renal repair. The preferred approach in these situations is mobilization of the kidney from lateral to medial. This approach has been shown to be faster and is associated with less blood loss compared to attempting vascular control of the renal pedicle prior to exploration. This approach is identical to the medial visceral rotation (Mattox maneuver). • Damage control by packing the wound to control bleeding and attempting to correct metabolic and coagulation abnormalities, with a plan to return for corrective surgery within 24 hours is an option.
URETERAL INJURIES	<ul style="list-style-type: none"> • Hematuria not universal: a high index of suspicion must be maintained. • Can be diagnosed with IV contrast and a delayed KUB or CT, but should only be done after the patient is fully resuscitated and hemodynamically stable. • Middle and upper 1/3 ureteral contusion is treated by excision and ureteroureterostomy: mobilize injured ureter, sparing adventitia widely to prevent devascularization; débride ureter liberally until edges bleed; repair ureter (absorbable suture) under magnification with spatulated, tension-free, stented, watertight anastomosis, and drain. Consider omental interposition to isolate repair. • Ureteral Pelvic Junction (UPJ) avulsion injuries should undergo re-anastomosis of the ureter to the renal pelvis. A stent and drains need to be placed. • Lower 1/3 ureteral injuries should be reimplanted into the bladder. Use a psoas hitch or Boari flap if required.

Guideline Only/Not a Substitute for Clinical Judgment

June 2010

Joint Theater Trauma System Clinical Practice Guideline

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BLADDER INJURIES	<ul style="list-style-type: none"> • Most patients will present with gross hematuria. If CT is planned for other injuries, a CT cystogram (<i>use DILUTED conray</i>) should be performed. If no CT, obtain a plain film cystogram (<i>need minimum 300ml to be adequate study.</i>) • Cystogram: Obtain scout film. Fill bladder via foley by gravity with at least 350ml contrast (7 ml/kg for pediatrics). Obtain AP image ± Oblique view. Drain bladder completely and obtain AP image. Many bladder injuries are detected only on the post-drainage film. • Extraperitoneal extravasation of contrast can be managed with foley catheterization alone, unless: bone fragment projecting into the bladder, open pelvic fracture, or rectal perforation. Open repair is indicated in these cases (see below). • Intraperitoneal ruptures require open repair, two-layer closure with absorbable suture and perivesical drain placement. Last, place a large-bore suprapubic catheter and a urethral catheter to maximize bladder drainage of blood and clots
POSTERIOR URETHRAL INJURIES	<ul style="list-style-type: none"> • Initial Management – Surgeon to attempt foley placement, consider urethroscopic assisted stenting of the injury with a urethral catheter. • If unable to pass a urethral foley catheter, operatively place an open suprapubic tube. At the time of open s-p tube placement, inspect the bladder to rule out injury.
ANTERIOR URETHRAL INJURIES	<ul style="list-style-type: none"> • Diagnosis – As with posterior urethral injury, a high index of suspicion must be maintained in all patients with blunt or penetrating trauma in the urogenital region, and a RUG should be performed in any case of suspected urethral injury. • Anterior urethral injuries may also be associated with large hematoma or swelling from extravasated urine. In severe trauma, Buck's fascia may be disrupted, resulting in blood and urinary extravasation into the scrotum. • Management- Initial suprapubic urinary diversion is recommended after high-velocity gunshot wounds to the urethra, followed by delayed reconstruction.
EXTERNAL GENITALIA INJURIES	<ul style="list-style-type: none"> • Penis - superficial wounds can be irrigated and closed primarily. Corporal injures are repaired by approximation of the tunical margins with absorbable sutures. Associated anterior urethral injuries should be closed primarily with a watertight, spatulated, catheter-stented technique and absorbable suture; posterior urethral injuries should be managed in staged fashion with suprapubic catheterization.

Guideline Only/Not a Substitute for Clinical Judgment

June 2010

Joint Theater Trauma System Clinical Practice Guideline

	GENERAL MANAGEMENT PRINCIPLES
EXTERNAL GENITALIA INJURIES (CONT)	<ul style="list-style-type: none">• Scrotum/Testicle - Diagnosis by physical exam and ultrasound. Equivocal cases should be explored. Explore testicles with overlying shrapnel on pelvic film or if there is a scrotal laceration and any abnormality on exam. Necrotic testicular tissue should be débrided and the capsule closed with running absorbable suture. In some cases, loss of capsule requires removal of intratesticular tissue to allow closure.

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June 2010

APPENDIX B

ADDITIONAL INFORMATION REGARDING OFF-LABEL USES IN CPGs

A. Purpose.

The purpose of this Appendix is to ensure an understanding of DoD policy and practice regarding inclusion in CPGs of “off-label” uses of U.S. Food and Drug Administration (FDA)–approved products. This applies to off-label uses with patients who are armed forces members.

B. Background.

Unapproved (i.e., “off-label”) uses of FDA-approved products are extremely common in American medicine and are usually not subject to any special regulations. However, under Federal law, in some circumstances, unapproved uses of approved drugs are subject to FDA regulations governing “investigational new drugs.” These circumstances include such uses as part of clinical trials, and in the military context, command required, unapproved uses. Some command requested unapproved uses may also be subject to special regulations.

C. Additional Information Regarding Off-Label Uses in CPGs.

The inclusion in CPGs of off-label uses is not a clinical trial, nor is it a command request or requirement. Further, it does not imply that the Military Health System requires that use by DoD health care practitioners or considers it to be the “standard of care.” Rather, the inclusion in CPGs of off-label uses is to inform the clinical judgment of the responsible health care practitioner by providing information regarding potential risks and benefits of treatment alternatives. The decision is for the clinical judgment of the responsible health care practitioner within the practitioner-patient relationship.

D. Additional Procedures.

1. Balanced Discussion. Consistent with this purpose, CPG discussions of off-label uses specifically state that they are uses not approved by the FDA. Further, such discussions are balanced in the presentation of appropriate clinical study data, including any such data that suggest caution in the use of the product and specifically including any FDA-issued warnings.

2. Quality Assurance Monitoring. With respect to such off-label uses, DoD procedure is to maintain a regular system of quality assurance monitoring of outcomes and known potential adverse events. For this reason, the importance of accurate clinical records is underscored.

3. Information to Patients. Good clinical practice includes the provision of appropriate information to patients. Each CPG discussing an unusual off-label use will address the issue of information to patients. When practicable, consideration will be given to including in an appendix an appropriate information sheet for distribution to patients, whether before or after use of the product. Information to patients should address in plain language: a) that the use is not approved by the FDA; b) the reasons why a DoD health care practitioner would decide to use the product for this purpose; and c) the potential risks associated with such use.