

A Journey Through the Hidden Depths of Onco-Nephrology

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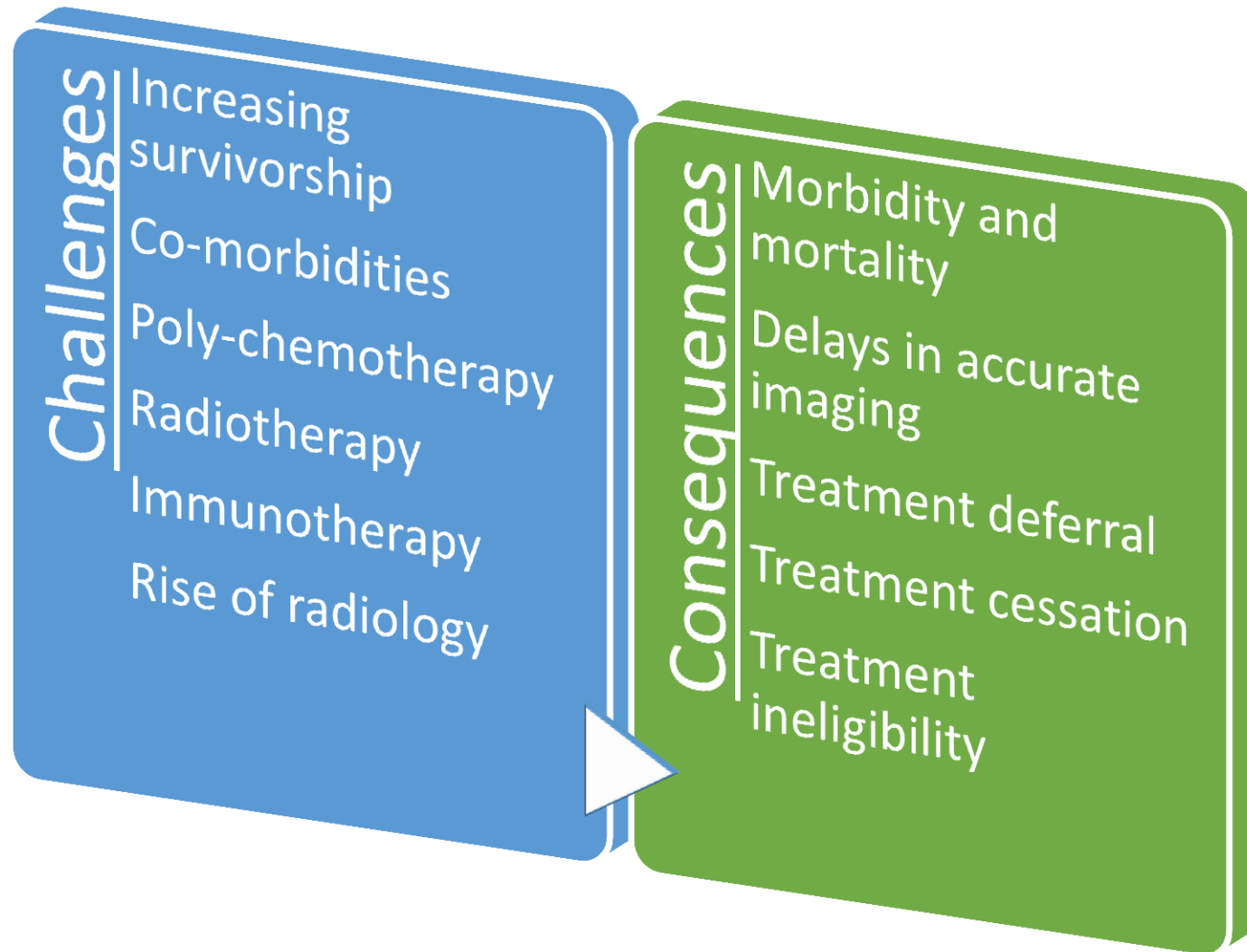
Consultant in Acute & Renal Medicine

The Christie NHS Foundation Trust

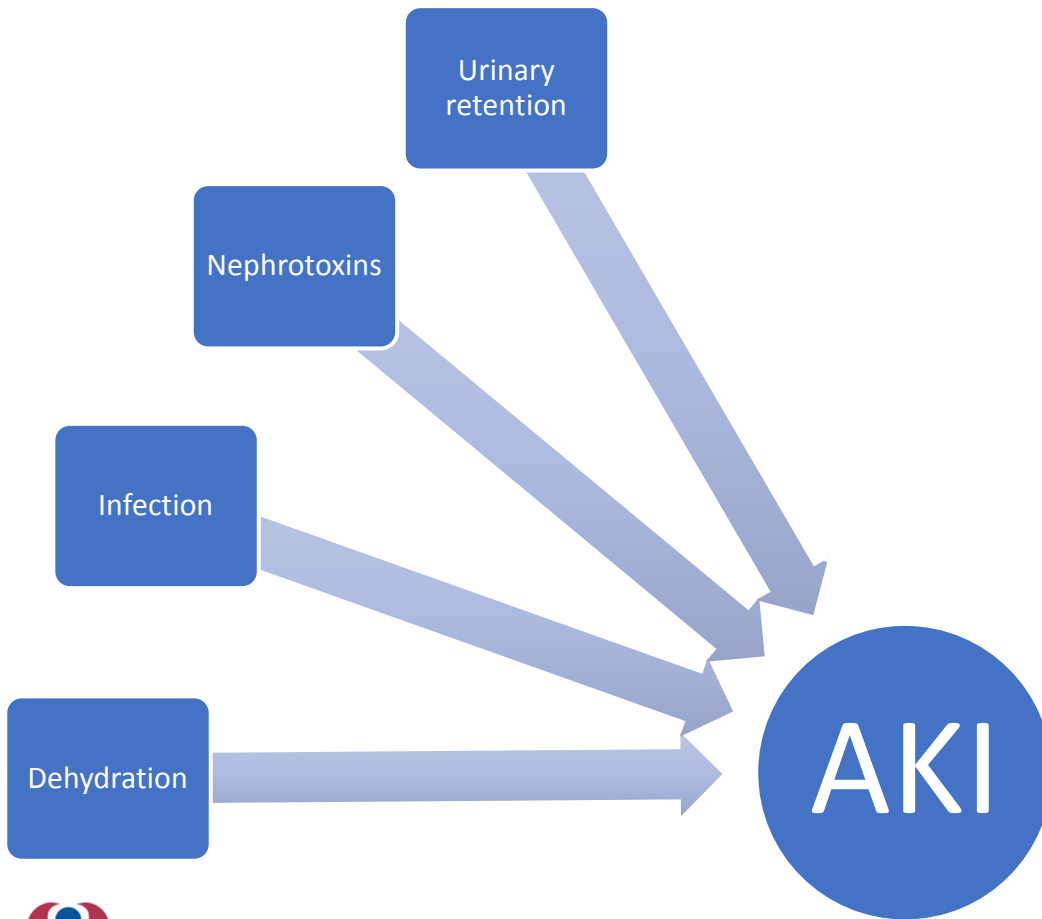
Wednesday 23rd November 2016



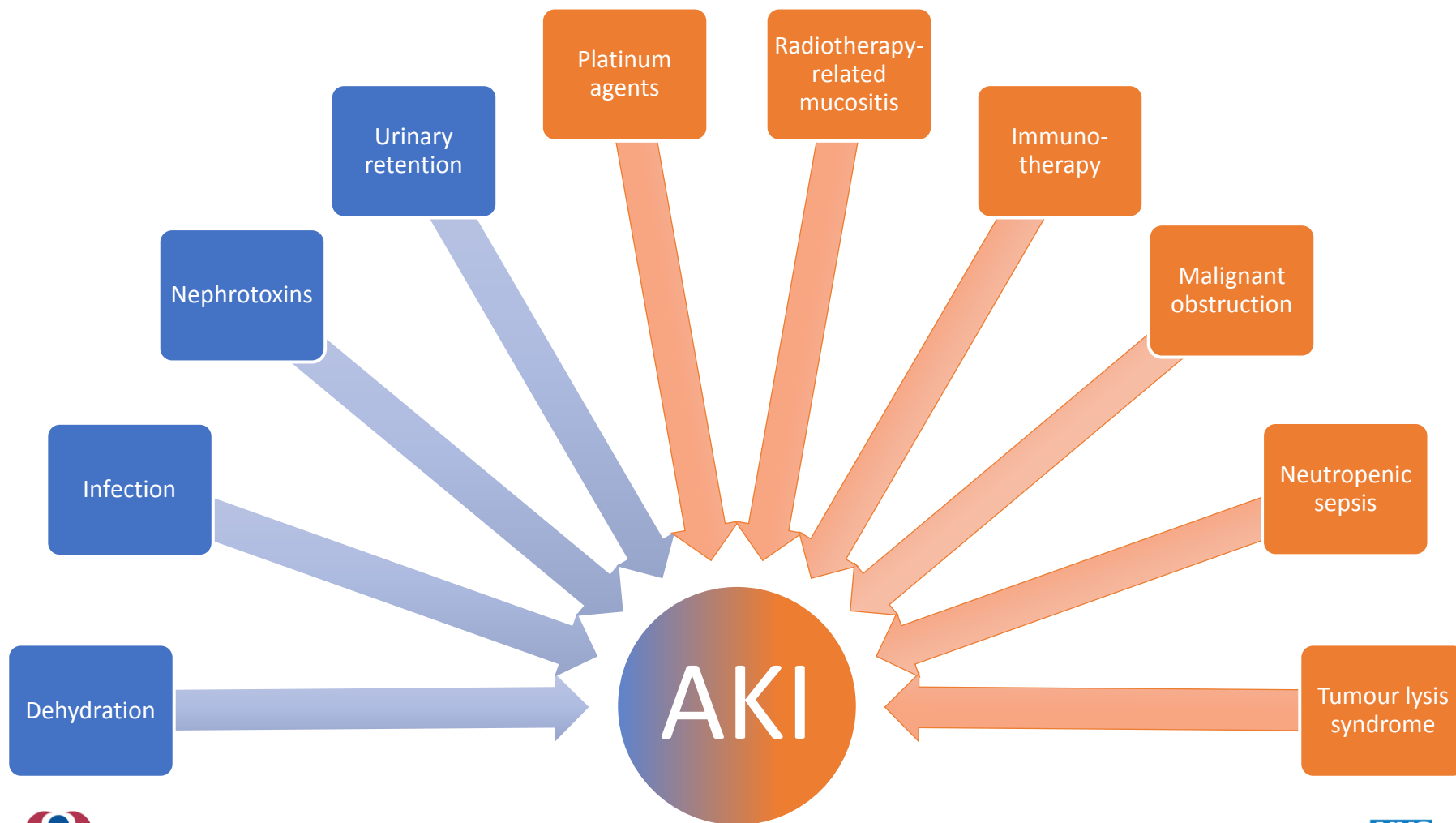
The Challenges of Modern Onco-Nephrology



Aetiology



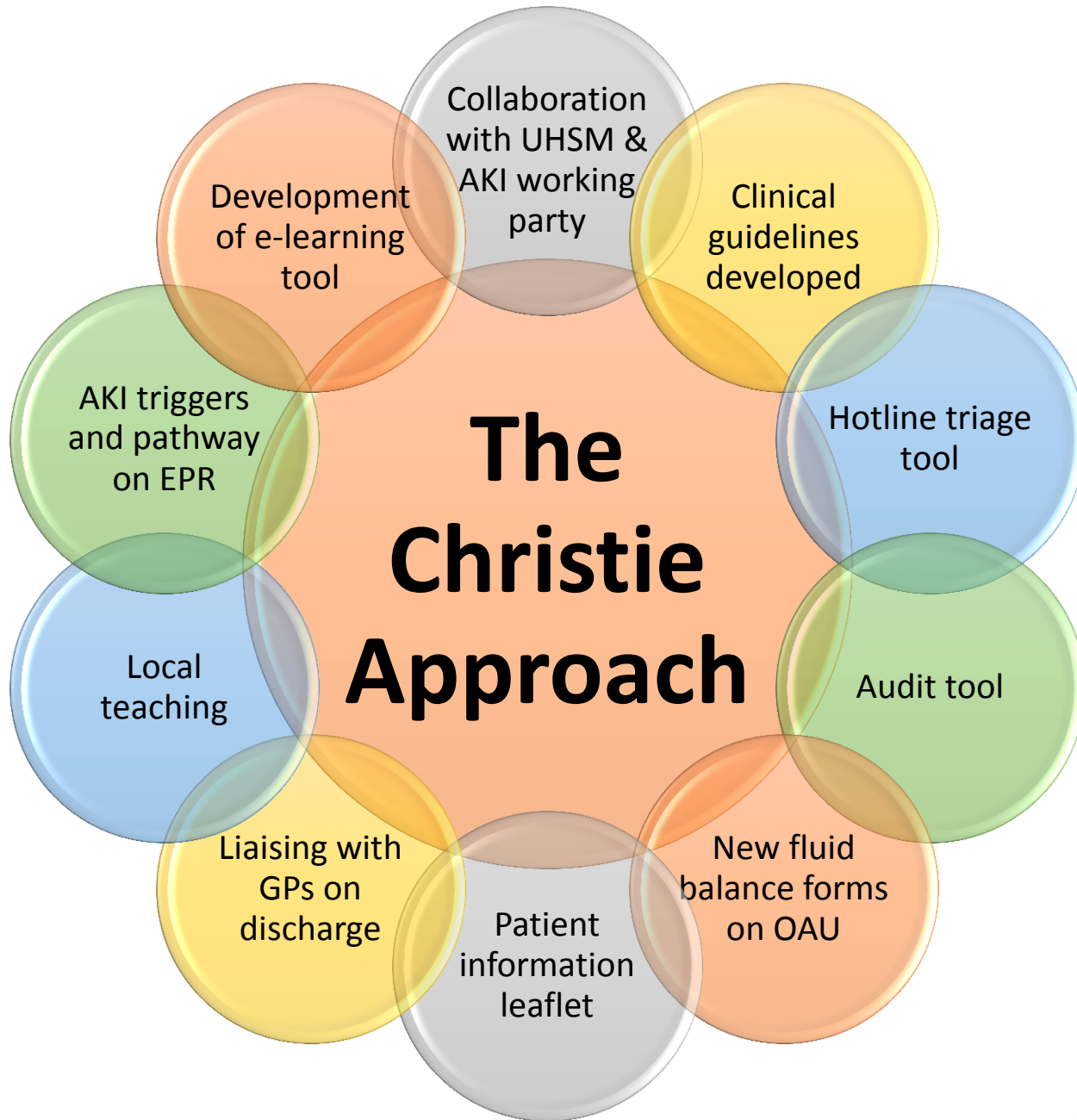
Aetiology

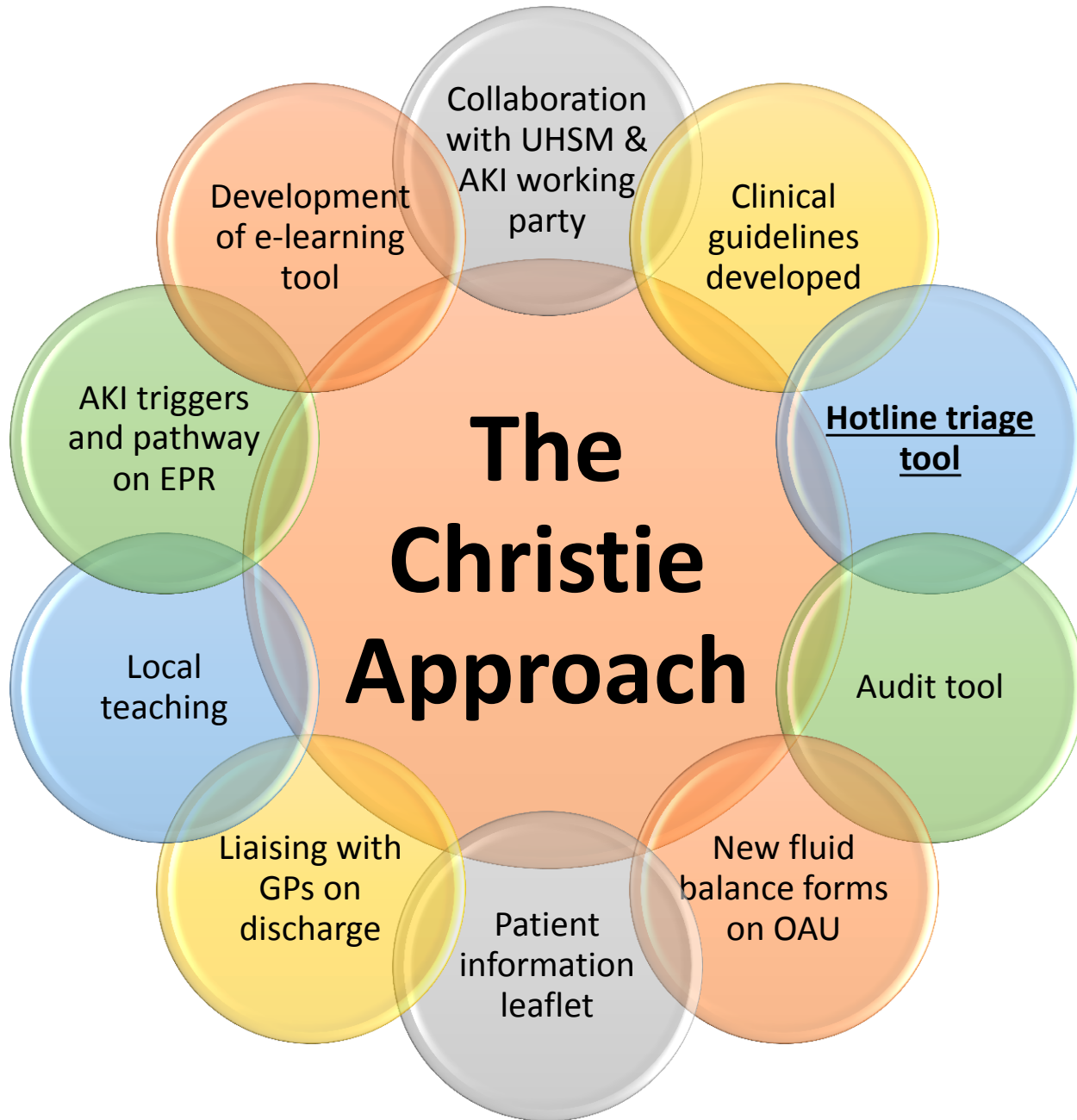


Acute Kidney Injury (AKI) at The Christie

- The Christie NHS FT
 - Largest single site cancer centre in Europe treating more than 44,000 patients per year
- 18 months ago, new Acute medicine role appointed
 - *“As a Consultant treating acutely unwell patients, I have worked with an expert team to develop guidelines for staff and patients in the treatment of acute kidney injury (AKI),” – Dr Al-Sayed*







DIARRHOEA

Initial Assessment

Always review last U&E results in patient electronic notes

Questions:

- What chemotherapy is the patient on and when was the last treatment/tablet?
- Are they receiving radiotherapy and when was their last treatment?
- Number of recent episodes?
- How often do the bowels usually move?
- How many stools a day is the patient passing or how much stoma output is there above normal amount?
- Are stools/stoma output formed, loose or watery? Any faecal incontinence or urgency? Nocturnal movements?
- Is there any abdominal pain e.g., cramping pains coming in waves?
- For how many days has the patient had diarrhoea? Is it interfering with activities of daily living?
- Are they able to eat and drink normally? Are they passing plenty of clear urine?
- Do they have any other chemotherapy related toxicities, e.g. mouth ulcers, N/V, red hands/feet, stomatitis, mucositis?
- Any recent antibiotics or recent hospital admissions?
- Have they taken any laxatives or anti-sickness medication **or** any anti-diarrhoeal medication in the last 24 hours? What?

Advice:

If taking Capecitabine chemotherapy follow the Capecitabine management protocol

Grade 1 (Green)	Grade 2 (Amber)	Grade 3 (Red)	Grade 4 (Red)
Increase to 2-3 bowel movements a day over pre-treatment baseline or mild increase in stoma output	Increase of 4-6 bowel movements a day over pre-treatment baseline, moderate increase in stoma output. Moderate cramping Nocturnal stools	Increase of 7-9 bowel movements a day over pre-treatment baseline or incontinence. Severe increase in stoma output. Severe cramping Nocturnal stools Interfering with ADL	Increase to > 10 bowel movements a day over pre-treatment baseline and/or grossly bloody diarrhoea and/or need for parenteral support

DRUGS - NB. Has the patient had a platinum based chemotherapy?

Is the patient taking:

- NSAIDs e.g. Diclofenac, Ibuprofen
- ACE inhibitors e.g. Ramipril, Lisinopril.

NB if patient taking any of the above drugs advise to **omit** until management plan agreed.



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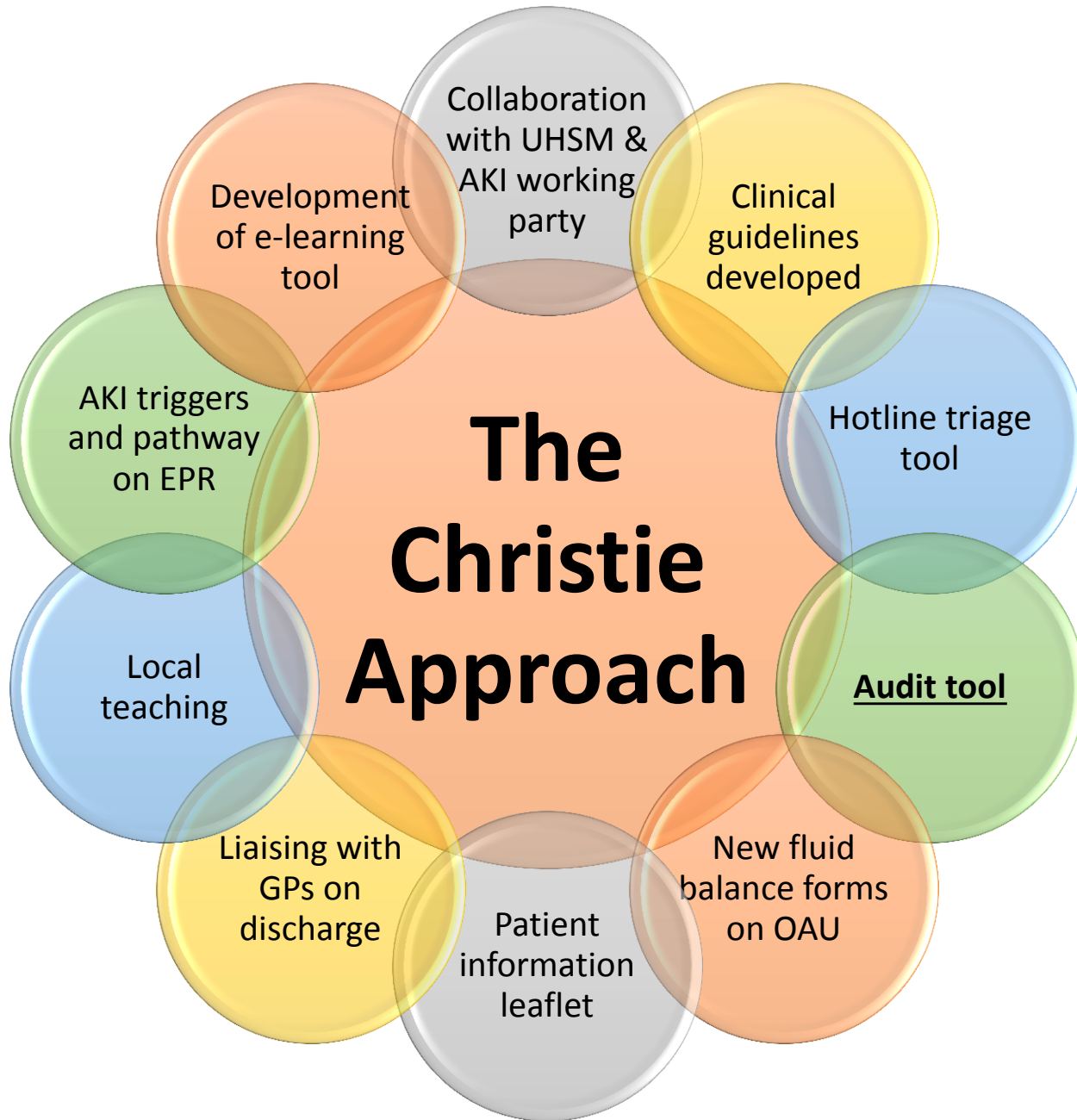
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The Christie Approach

Development of e-learning tool

Collaboration with UHSM & AKI working party

Clinical guidelines developed

Hotline triage tool

AKI triggers and pathway on EPR

Audit tool

Local teaching

New fluid balance forms on OAU

Patient information leaflet

Liaising with GPs on discharge



Affix patient sticker here
(form to be retained in notes)

Date of admission	
Time	
Dr(grade)/ANP	
Pt referred from	
Hospital number	

AKI risk recognition tool and preventative care bundle

ADULTS AT RISK (any of the following):	Tick		Tick
CKD (eGFR <60ml/min)		Nephrotoxic drugs including :	
Heart failure		Iodinated contrast agents in last 14 days	
Chronic Liver Disease (Cirrhosis)		Platin chemotherapy within 30 days	
Diabetes		Non-platin chemotherapy in last 30 days	
Previous AKI		ACE inhibitors	
Oliguria (urine output < 0.5ml/kg/hr)		NSAIDS	
Reduced/impaired conscious level (GCS)		Diuretics	
Hypovolaemia			
Symptoms/history of urological obstruction			
Sepsis			
Deteriorating MEWS			

If patient is at risk of developing AKI or has established AKI commence the following measures:

	Tick
- Fluid balance and daily weight charts initiated	
- Urinary catheter considered (or used)	
- Correct hypovolaemia (see NICE guidance for fluid management)	
- Regular biochemistry (Compare admission creatinine with baseline)	
- Urine dipstick & analysis	
- Stop nephrotoxic medications	
- Review acid/base balance	
- Seek senior input (Acute Physician/Critical Care)	
- Ultrasound KUB within 24 hours if kidney function not improving	
- Cause of AKI (if established):	
- Seek renal opinion if the cause of AKI is not apparent and/or the patient is deteriorating to the point they require escalation of care	
AKI diagnosis communicated to GP on discharge	
Patient information leaflet offered upon discharge	

Classification of AKI

AKI Stage	Serum Creatinine criteria	Tick	Urine output criteria	Tick
1	Rise of $\geq 26.5 \mu\text{mol/l}$ or 1.5-2 fold increase from baseline		<0.5 mls/kg/hr for 6 hours	
2	>2-3 fold increase from baseline		<0.5 mls/kg/hr for 12 hours	
3	>3.0-fold from baseline OR serum creatinine $\geq 354 \mu\text{mol/l}$ with an acute increase of at least $44 \mu\text{mol/l}$ OR need for RRT		<0.5 mls/kg/hr for 24 hours or anuria for 12 hours or need for RRT	





AKI identification and risk

[Patient Menu](#)[Patient Forms](#)[Clinical Notes](#)[Lab Results](#)

* Date and time of assessment 18-Nov-2016 14:17

Serum creatinine This is the most recent result reported. If blank, there are no results within 3 days

Date: [View Results](#)

Risk factors and present AKI status

* AKI present Is there evidence of AKI

 Yes No Not yet known

* AKI risk factors (chronic) Complete on admission, change as necessary.

- No chronic risk factors
- CKD (eGFR less than 60ml/min)
- Heart failure
- Liver disease
- Diabetes
- Previous AKI
- Chronic neurological or cognitive impairment
- History of urological obstruction

* AKI risk factors (acute) Present now

- No acute risk factors
- Oligouria
- Acute neurological or cognitive impairment
- Hypovolaemia
- Nephrotoxic drugs (include iodine containing contrast)
- Symptoms or new urological obstruction
- Sepsis
- Deteriorating EWS

Initial management

* Renal management plan Mark all completed or planned

- Fluid balance and daily weight charts requested
- Correction of hypovolaemia
- Regular haematology and Christie profile planned
- Urine dip requested
- Nephrotoxic medications reviewed
- Review of acid/ base balance

* Actions taken and plan

* Senior doctor aware Has a senior doctor on the team or on call team been made aware of the findings and actions?

- No senior review required
- No senior review yet
- Registrar
- Consultant

[Submit](#)

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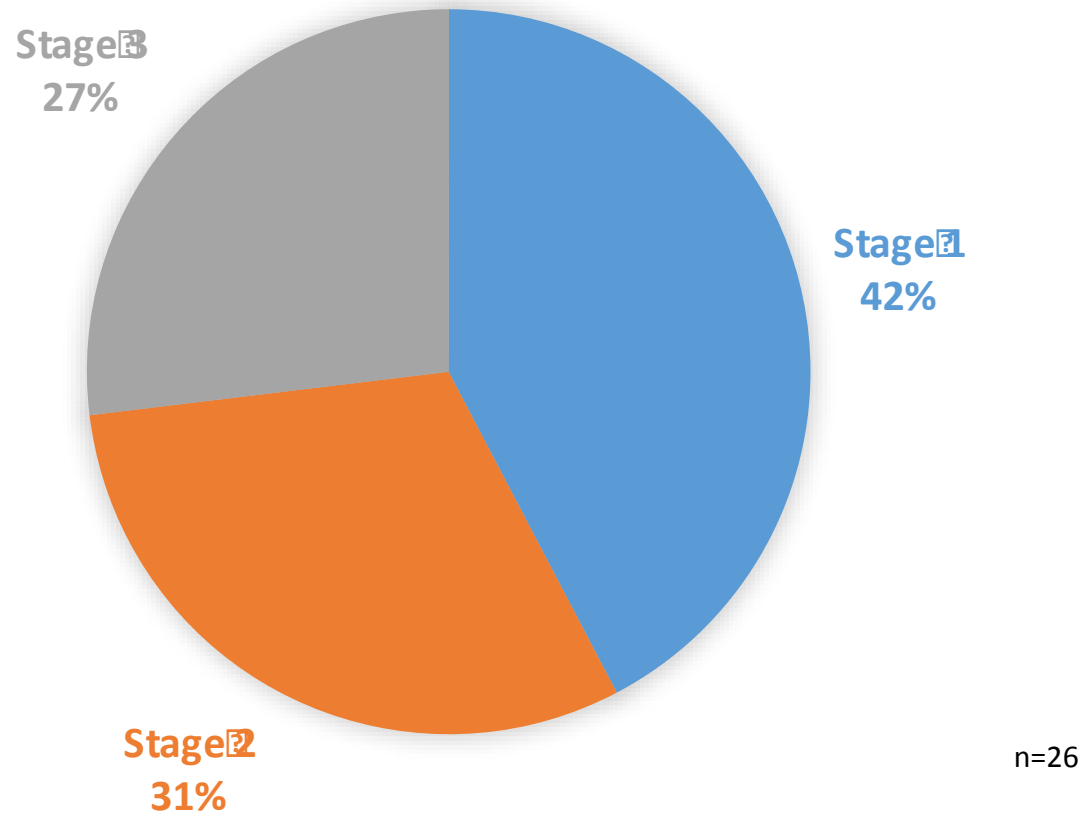
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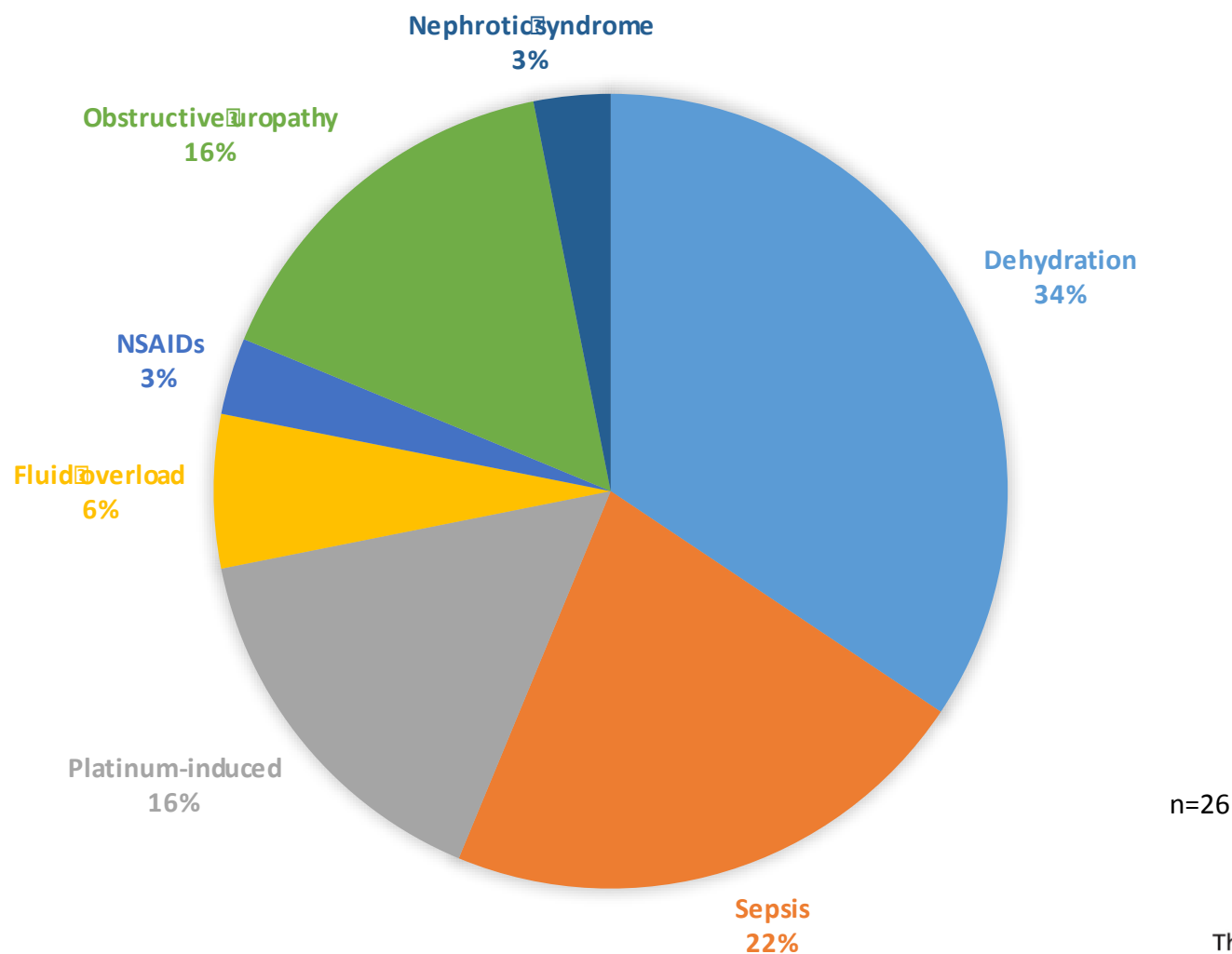
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[Submit](#)

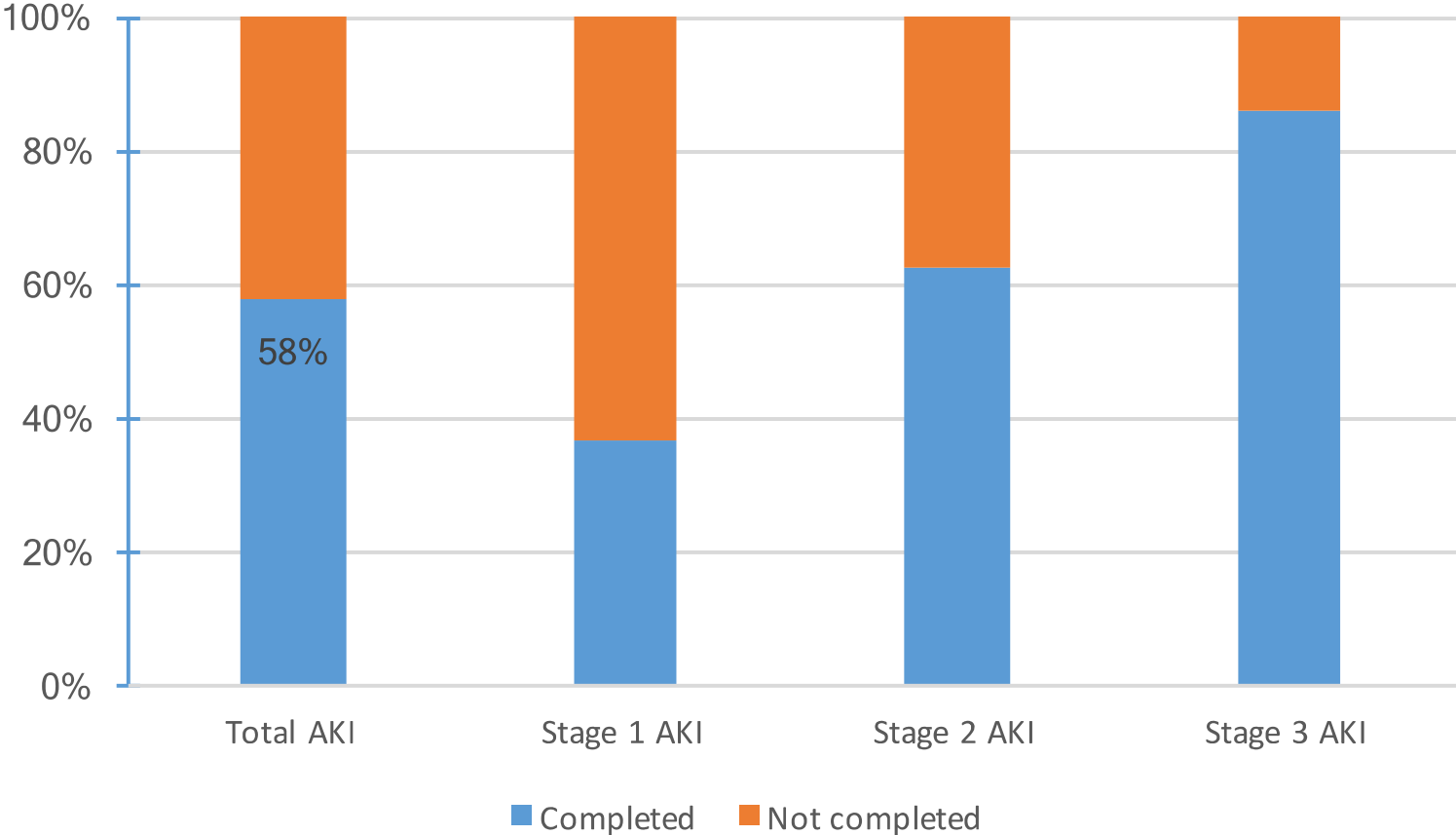
AKI Incidence by Stage (inpatients only)



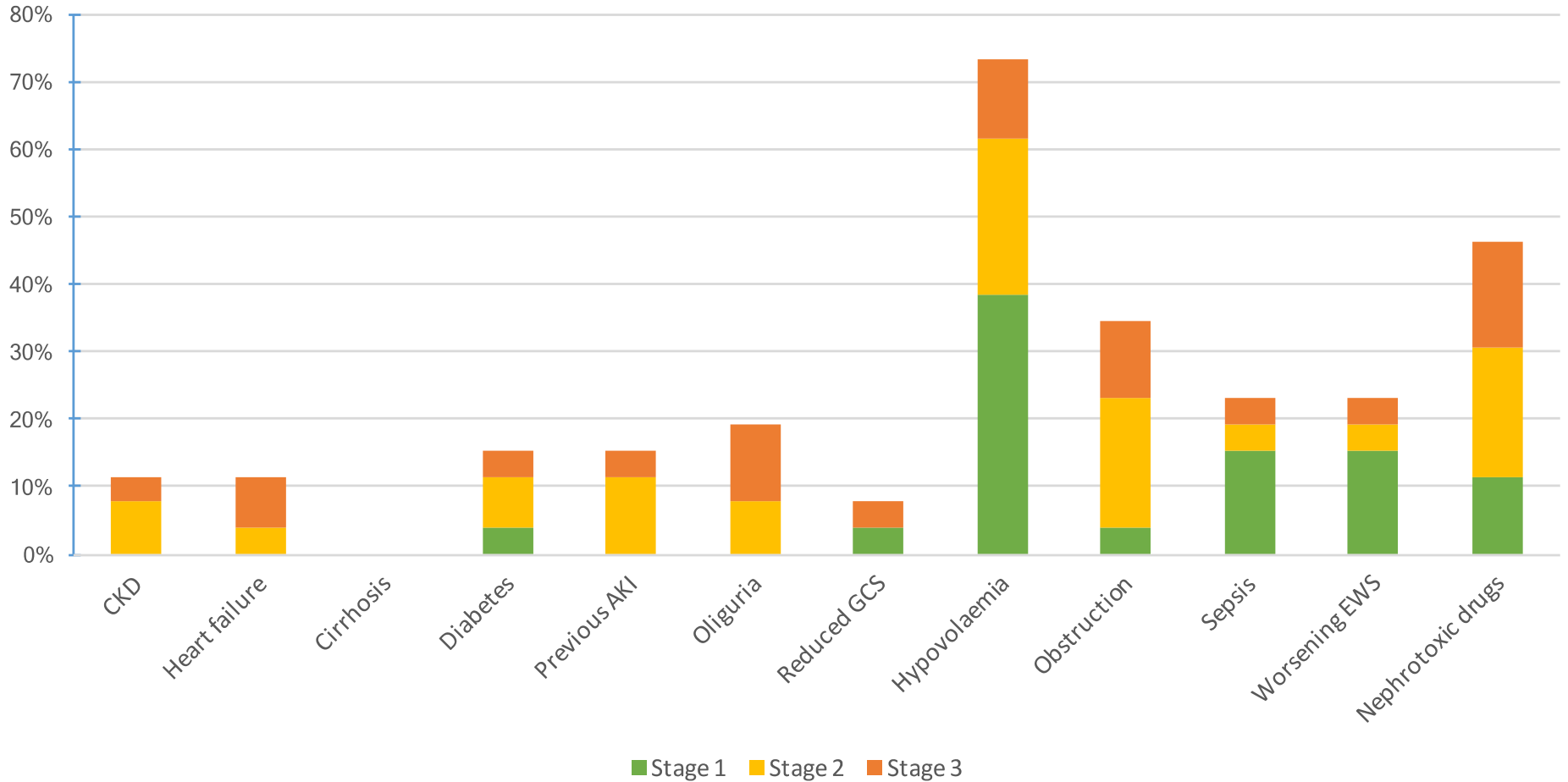
AKI Aetiology in Oncology Patients



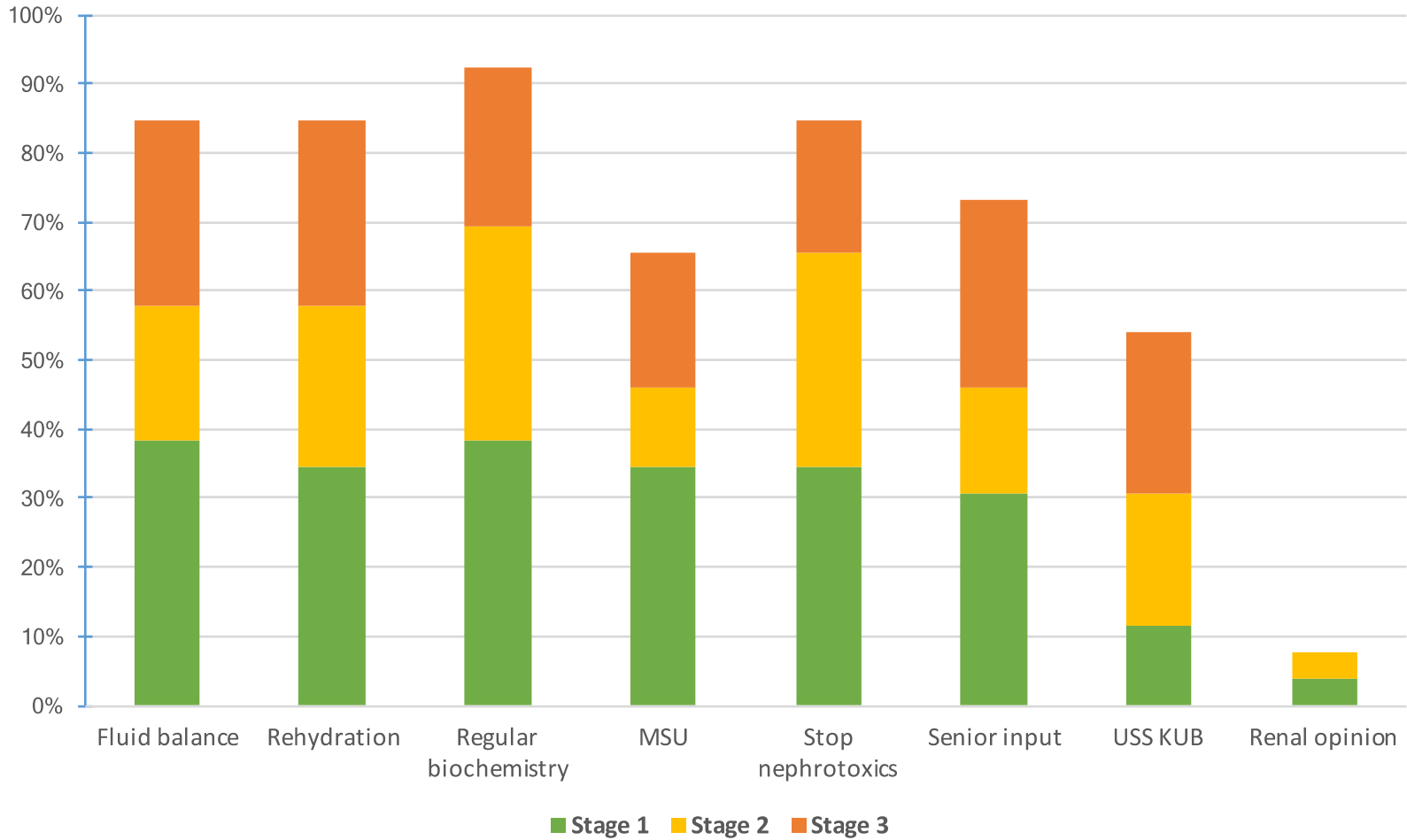
Compliance with AKI Care Bundle



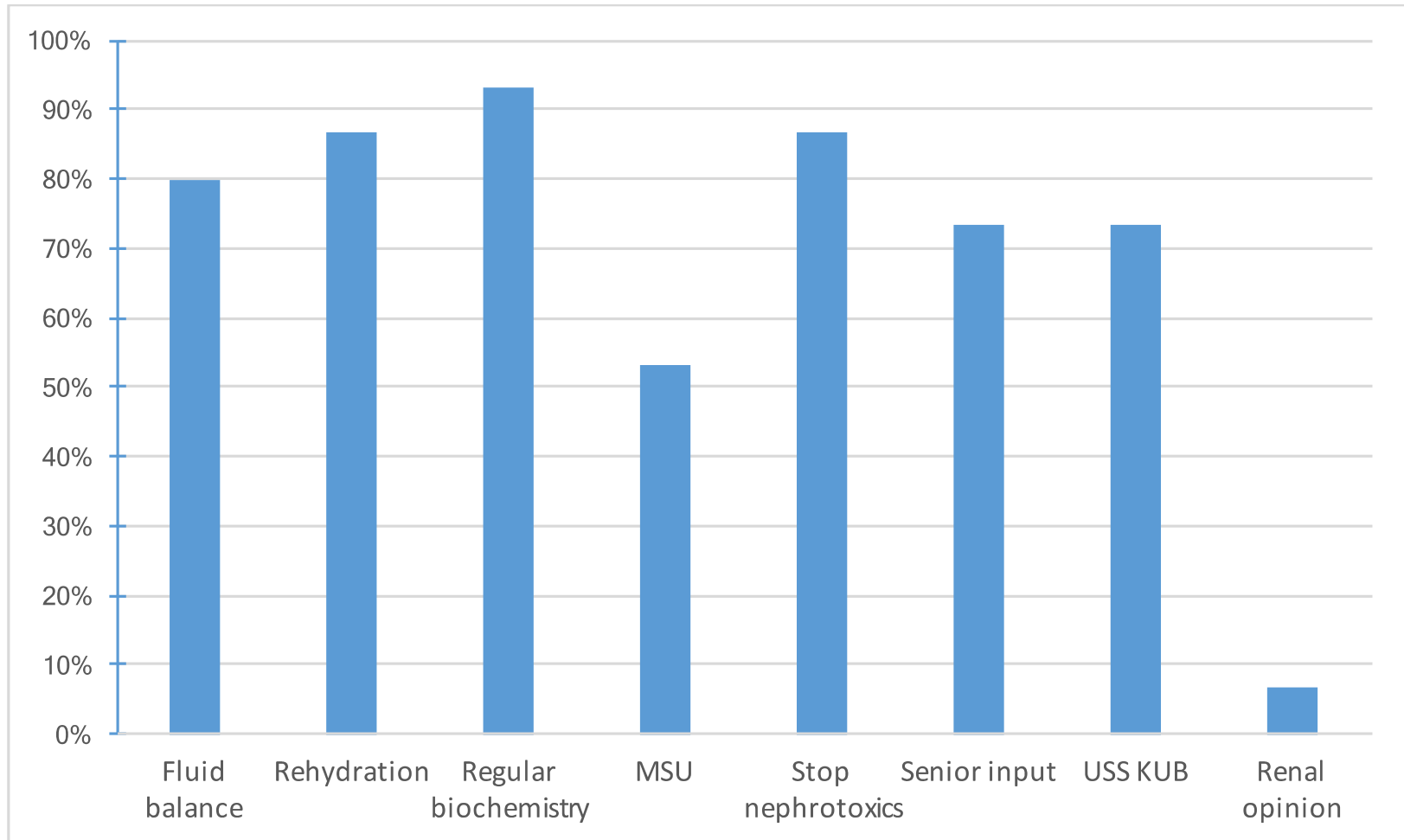
Risk Factors for AKI in Oncology Patients



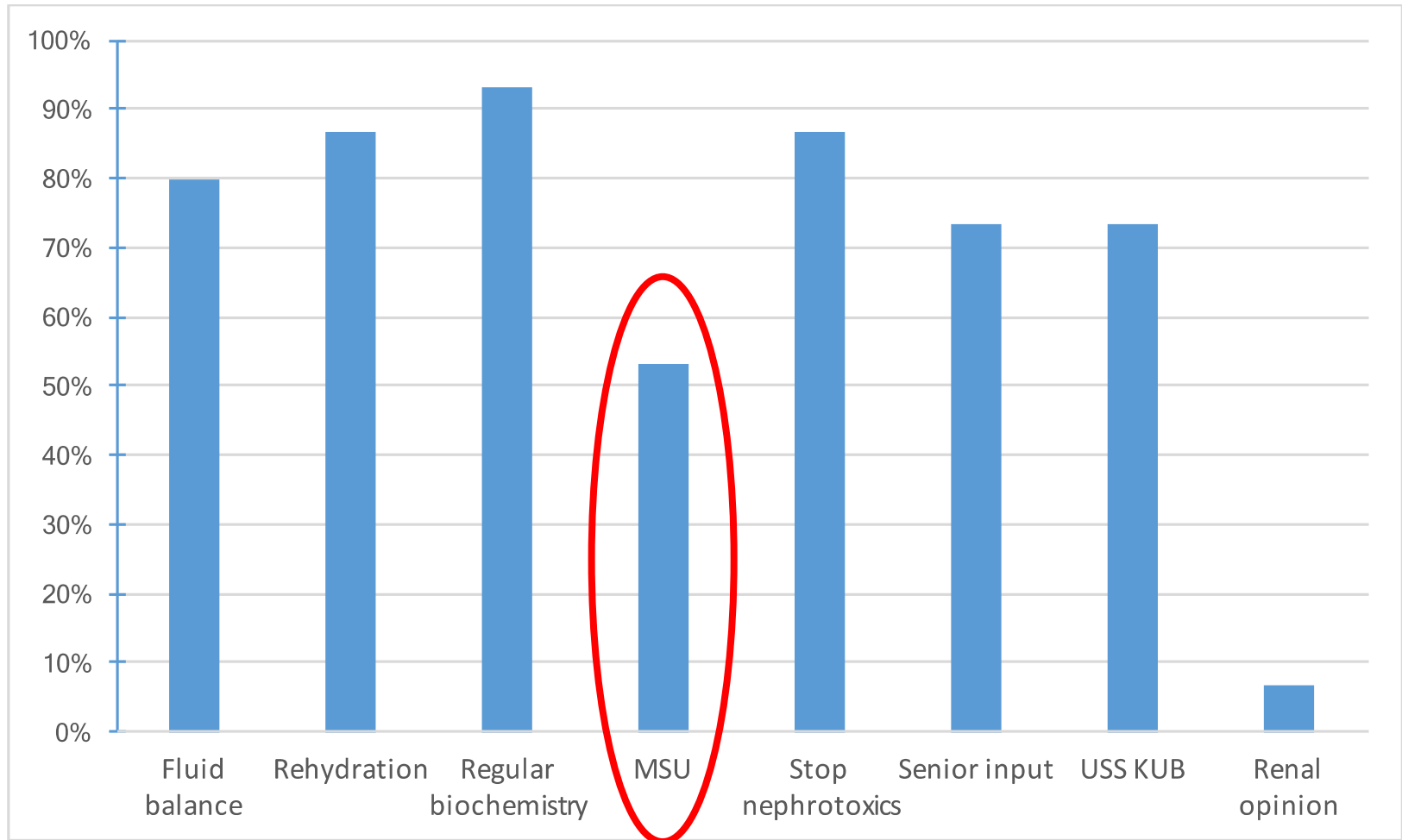
Management of AKI by stage



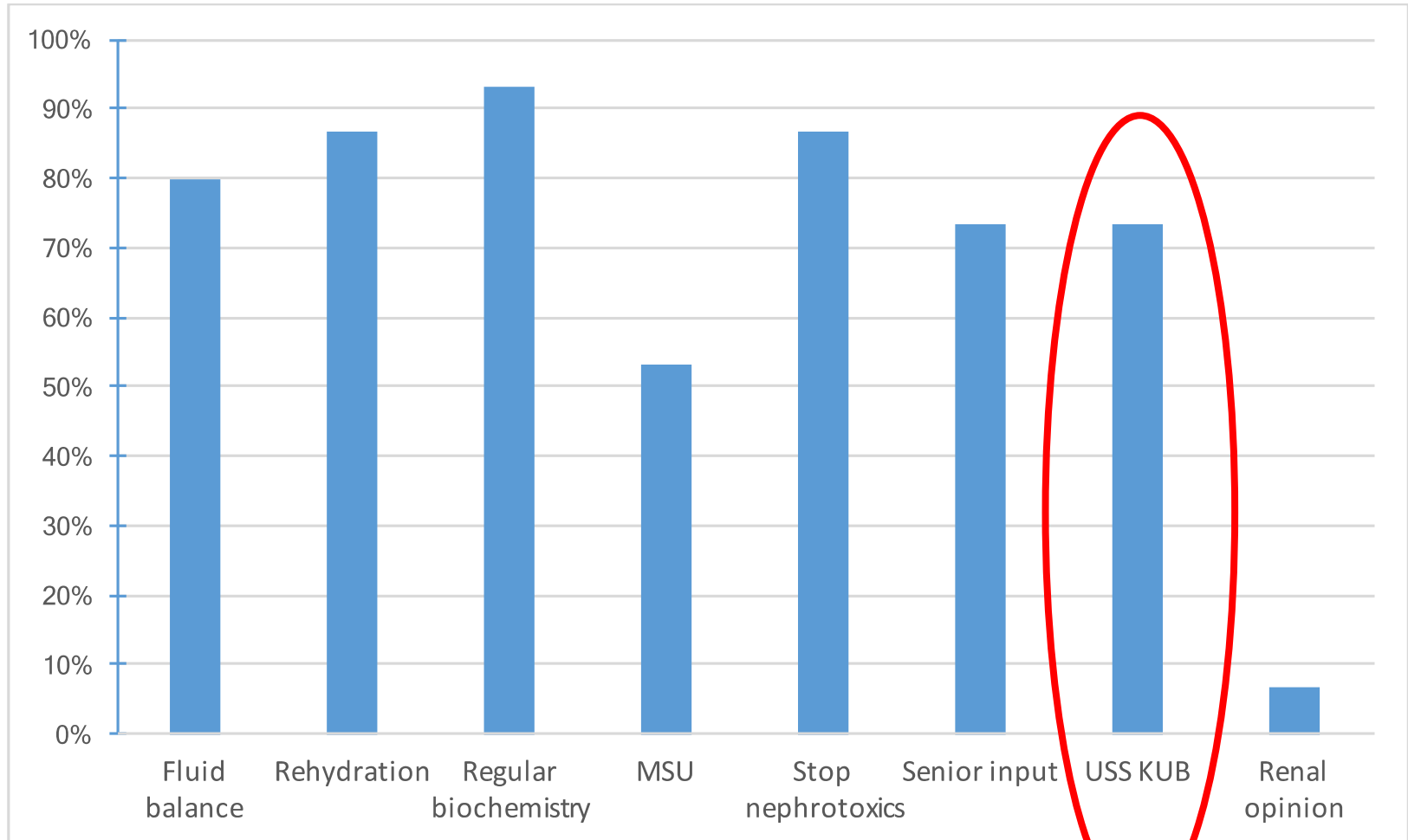
Management of Stage 2-3 AKI



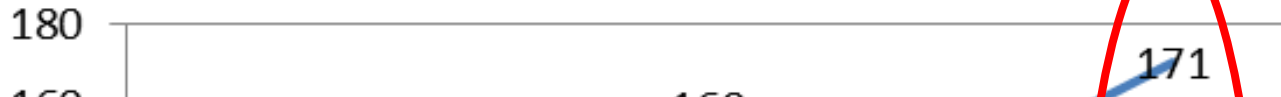
Management of Stage 2-3 AKI



Management of Stage 2-3 AKI



No. individual patients with AKI alerts by month



Patient Activity (OAU)	July 2016	August 2016	September 2016
Total activity	364	380	383
Elective admissions	4	1	1
Emergency admissions	297	317	305
Bed occupancy	82%	85%	81%
Direct discharges	151	150	121

KI level 1
 KI level 2
 KI level 3

Jun-16 Jul-16 Aug-16 Sep-16 Oct-16



Difficulties & Potential Solutions

- Data collection erratic & therefore not 100% reliable
 - EPR, e-prescribing & live dashboards are potential game changers
- Lack of human resource
 - Employing a full time AKI nurse
- AKI risk precedes SACT
 - AKI assessment needs to occur in the OPD before prescribing chemotherapy
- Multiple areas for AKI risk including surgery & haematology
 - Identify AKI champions for different clinical areas
- Lack of community integration
 - AQuA 2017

Learning Points

- AKI is common and has significant morbidity, including prolonged LOS and precluding further SACT
 - Often complex aetiologies especially with emerging new therapies
- Obstruction and nephro-toxic medications potentially confer a more severe degree of AKI, albeit reversible if intervened upon early
- Greater emphasis on care bundle completion & data collection
- Immediate medicines optimisation vital
- Integrating AKI tools with other protocols is key e.g. sepsis & IV fluids
- Executive support has been crucial
- Education – PowerPoint e-learning bytes



Thank you for listening...

Any questions?

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