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RESEARCH

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# Increased creatinine clearance in polytrauma patients with normal serum creatinine: a retrospective observational study

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## Abstract

**Introduction:** The aim of this study, performed in an intensive care unit (ICU) population with a normal serum creatinine, was to estimate urinary creatinine clearance ( $CL_{CR}$ ) in a population of polytrauma patients (PT) through a comparison with a population of non trauma patients (NPT).

**Methods:** This was a retrospective, observational study in a medical and surgical ICU in a university hospital. A total of 284 patients were consecutively included. Two different groups were studied: PT ( $n = 144$ ) and NPT ( $n = 140$ ). Within the second week after admission to the ICU, renal function was assessed using serum creatinine, 24 h urinary  $CL_{CR}$ .

**Results:** Among the 106 patients with a  $CL_{CR}$  above  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ , 79 were PT and 27 NPT ( $P < 0.0001$ ). Only 63 patients had a  $CL_{CR}$  below  $60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$  with 15 PT and 48 NPT ( $P < 0.0001$ ). Patients with  $CL_{CR}$  greater than  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$  were younger, had a lower SAPS II score and a higher male ratio as compared to those having  $CL_{CR}$  lower than  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ . Through a logistic regression analysis, age and trauma were the only factors independently correlated to  $CL_{CR}$ .

**Conclusions:** In ICU patients with normal serum creatinine,  $CL_{CR}$  is higher in PT than in NPT. The measure of  $CL_{CR}$  should be proposed as routine for PT patients in order to adjust dose regimen, especially for drugs with renal elimination.

## Introduction

Early detection of renal dysfunction in intensive care unit (ICU) patients is important. Indeed, an increase of the glomerular filtration rate (GFR) was demonstrated in some ICU patients populations by using urinary creatinine clearance ( $CL_{CR}$ ) as a surrogate marker, and many of the drugs used in ICU patients need dose adjustment as a function of GFR [1]. Despite a normal serum creatinine measurement, a substantial number of burn patients demonstrated an increase in GFR with the need for increasing doses of renal elimination drugs to maintain therapeutic concentrations [1,2]. A study of our group showed that 42% of burn patients

had a creatinine clearance greater than  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^2$  [1]. Also, several studies performed in a general population of ICU patients suggested a poor correlation between serum creatinine concentration and GFR in polytrauma patients (PT) [3-5]. To the best of our knowledge, no study has specifically explored this population of PT patients.

The aim of this study, performed in a population of ICU patients with normal serum creatinine, was to estimate GFR, evaluated by measured  $CL_{CR}$ , in a population of PT patients through a comparison with a population of non-trauma patients (NPT).

## Materials and methods

### Patients

This observational study was conducted in the ICU of Toulouse University hospital during a five-year period

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(November 2002 to December 2007). The study was performed according to the Declaration of Helsinki. No change in our current clinical practice (measured creatinine clearance monitoring, at least once a week, is a part of the routine medical care of the patients) and no randomization was performed. As it was an observational retrospective study, in accordance with French law, neither approval of the ethics committee nor informed consent was required.

Ten days, on average, after admission in ICU, consecutive critically ill patients meeting the inclusion criteria were included. Inclusion criteria were: patients older than 18 years, with an arterial catheter, a urinary bladder catheter, a diuresis over 500 mL d<sup>-1</sup>. All patients had a tracheal tube and were mechanically ventilated. Patients were hemodynamically stable presenting with a stable serum creatinine in a normal range (40 to 125 μmol L<sup>-1</sup>).

Patients were excluded from the study: if they were hemodynamically unstable or if they needed a high dose of catecholamine (norepinephrine > 1 mg h<sup>-1</sup>); if they were recovering from acute kidney injury (AKI) or developing AKI; if they received histamine-2-receptor antagonist due to its interference with tubular creatinine secretion [6]; and if they had a medical history of diabetes, of chronic hepatic disease, cirrhosis or ongoing liver dysfunction with hepatitis [7,8]. Patients treated with diuretics were also excluded.

Baseline characteristics of patients were recorded at enrolment in the study and the SAPS II was obtained at ICU admission. PT patients had an ISS (Injury Severity Score) > 16. SOFA (Sequential Organ Failure Assessment) score was obtained on the day the urine 24-hour measure was sampled [9-11].

Urine was sampled over 24 hours to measure urinary creatinine concentration. Serum creatinine was also measured during the urine collection period.

The normal limits of CL<sub>CR</sub> were estimated between 60 and 120 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup> [9,12].

#### Serum creatinine measurement and calibration

Creatinine measurements were performed in the same laboratory of the University Hospital of Toulouse. Blood samples were obtained simultaneously with the CL<sub>CR</sub> measurement. A modified kinetic Jaffe colorimetric method was used with a COBAS MIRA (ABX Diagnostics, Montpellier, France) analyzer. A two-point calibration was applied in each assay.

Before measurement, ultrafiltration of plasma through a 20 kD cutoff membrane (MPS-1; Amicon, Beverly, MA, USA) was performed to discard chromogens that were linked to albumin-like bilirubinemia and other heavy proteins. In the absence of an international standard for creatinine assay, the linearity of the measurements was verified by using plasma samples from

normal subjects in which increasing amounts of desiccated creatinine hydrochloride (MW 149.6; Sigma Chemicals, Perth, Australia) had been added. Linear regression analysis showed that the relationship between measured and expected creatinine concentrations was  $1.0008 \pm 0.006$  (95% confidence interval, 0.997 to 1.020) and that the Y-intercept was  $0.014 \pm 0.013$  (95% confidence interval, -0.013 to 0.041). Squared Spearman rank coefficient of correlation was 0.998. Internal quality controls showed a coefficient of variation of 2.3% during the period.

#### Assessment of glomerular filtration rate

Creatinine clearance was measured according to the

formula  $CL_{CR} = \frac{UC_R \times V}{SC_R}$  where urine creatinine

(UCR) and serum creatinine (SCR) were expressed in μmol L<sup>-1</sup> and V corresponded to the urinary rate (diuresis) in mL minute<sup>-1</sup>.

At the same time, the GFR was estimated using the

Cockcroft Gault formula [13]  $CL_{CR} = \frac{(140 - \text{age}) \times \text{Weight}}{0.8 \times SC_R}$

for men, with age in years and weight in Kg. A correcting factor of 0.85 was used for women. The derivate formula proposed by Robert *et al.* [14] uses the ideal body weight and serum creatinine concentration corrected to 85 μmol L<sup>-1</sup> when the actual value is lower than 85 μmol L<sup>-1</sup>. Ideal body weight was determined as 50 kg for men and 45.5 kg for women, plus 2.3 kg for each inch over five feet. The simplified formula of the Modification of Diet in Renal Disease index (sMDRD) [15] was also calculated according to  $sMDRD = 186.3 \times SCR^{-1.154} \times \text{age}^{-0.203} \times (1.212 \text{ if black}) \times (0.742 \text{ if female})$  where serum creatinine was expressed in mg dL<sup>-1</sup>.

#### Statistical analysis

Statistical analyses were performed using StatView<sup>®</sup> software version 5.0 (SAS Institute Inc., Cary, NC, USA). Data are presented as mean ± standard deviation (SD) or ratio. Normal distribution of data was tested via Kolmogorov-Smirnov test. Chi-square test or Student's *t*-test was performed when appropriate. A logistic regression was performed to discriminate if trauma, age, SAPS II, ideal body weight and sex are independently correlated to the measured CL<sub>CR</sub>. A *P*-value < 0.05 was considered as statistically significant.

#### Results

Demographic and renal data are shown in Table 1. Two hundred, eighty-four patients were consecutively included in this observational study. The process of screening and inclusion in the study is shown in Figure 1 (flow chart).

**Table 1 Demographic data**

	NPT <sup>a</sup> (n = 140)	PT <sup>b</sup> (n = 144)	P*
Age (yr)	58 ± 17	42 ± 18	< 0.0001
Weight (kg)	72 ± 18	75 ± 14	NS
Height (cm)	170 ± 8	174 ± 9	NS
Sex (F/M)	52/88	36/108	0.03
Ideal body weight (Kg)	68 ± 11	72 ± 11	0.0006
SAPS 2	52 ± 14	42 ± 15	< 0.0001
Total SOFA score	3.7 ± 1.5	3.6 ± 1.4	NS
Respiratory system	0.3 ± 0.2	0.5 ± 0.6	
Coagulation	0.0 ± 0.0	0.0 ± 0.0	
Liver	0.0 ± 0.0	0.0 ± 0.0	
Cardiovascular system	1.3 ± 0.9	1.4 ± 0.9	
Neurological system	2.5 ± 1.5	2.4 ± 1.5	
Renal system	0.0 ± 0.0	0.0 ± 0.0	
Mean arterial blood pressure (mmHg)	83 +/- 10	82 +/- 11	NS
Systolic arterial blood pressure (mmHg)	126 +/- 16	125 +/- 17	NS
Diastolic arterial blood pressure (mmHg)	63 +/- 10	63 +/- 11	NS
Heart rate (bpm)	91 +/- 17	92 +/- 16	NS
Serum creatinine (μmol L <sup>-1</sup> )	74 ± 26	72 ± 19	NS
Measured creatinine clearance (mL minute <sup>-1</sup> 1.73 m <sup>-2</sup> )	85 ± 5	131 ± 5	< 0.0001
Diuresis/24H	2,700 ± 1,200	2,500 ± 1,200	NS
Serum Urea	8.7 ± 5	7 ± 3	NS
Measured creatinine clearance < 60 (mL minute <sup>-1</sup> 1.73 m <sup>-2</sup> )	48	15	0.0003
Measured creatinine clearance > 120 (mL minute <sup>-1</sup> 1.73 m <sup>-2</sup> )	27	79	< 0.0001

All data are expressed as ratio or mean ± SD.

\*The P-values correspond to comparison between NPT and PT groups.

<sup>a</sup>NPT, non-polytrauma patients.

<sup>b</sup>PT, polytrauma patients.

The group of 144 PT patients was compared with the group of 140 NPT patients. No difference was found concerning hemodynamic data (Table 1). No difference was found concerning ventilation pressure. All the patients were ventilated with a tidal volume of 6 to 8 ml/kg, the PEP value was set at  $5.8 \pm 3$  in PT vs  $5.5 \pm 3$  in NPT patients (NS). FiO<sub>2</sub> was  $45 \pm 16\%$  in PT vs  $45 \pm 15\%$  in NPT (NS), with Ph =  $7.38 \pm 0.8$  vs  $7.39 \pm 0.8$  (NS), PaO<sub>2</sub> =  $107 \pm 16$  in PT vs  $108 \pm 15$  in NPT (NS), PaCO<sub>2</sub> =  $39 \pm 8$  in PT vs  $40 \pm 9$ , SaO<sub>2</sub> =  $97 \pm 3$  vs  $97 \pm 3$  (NS). Glycemia was not different between groups ( $6.2 \pm 1.7$  vs  $6.1 \pm 1.8$ ; NS). Twenty-three percent of PT vs 24% of NPT received norepinephrine (NS).

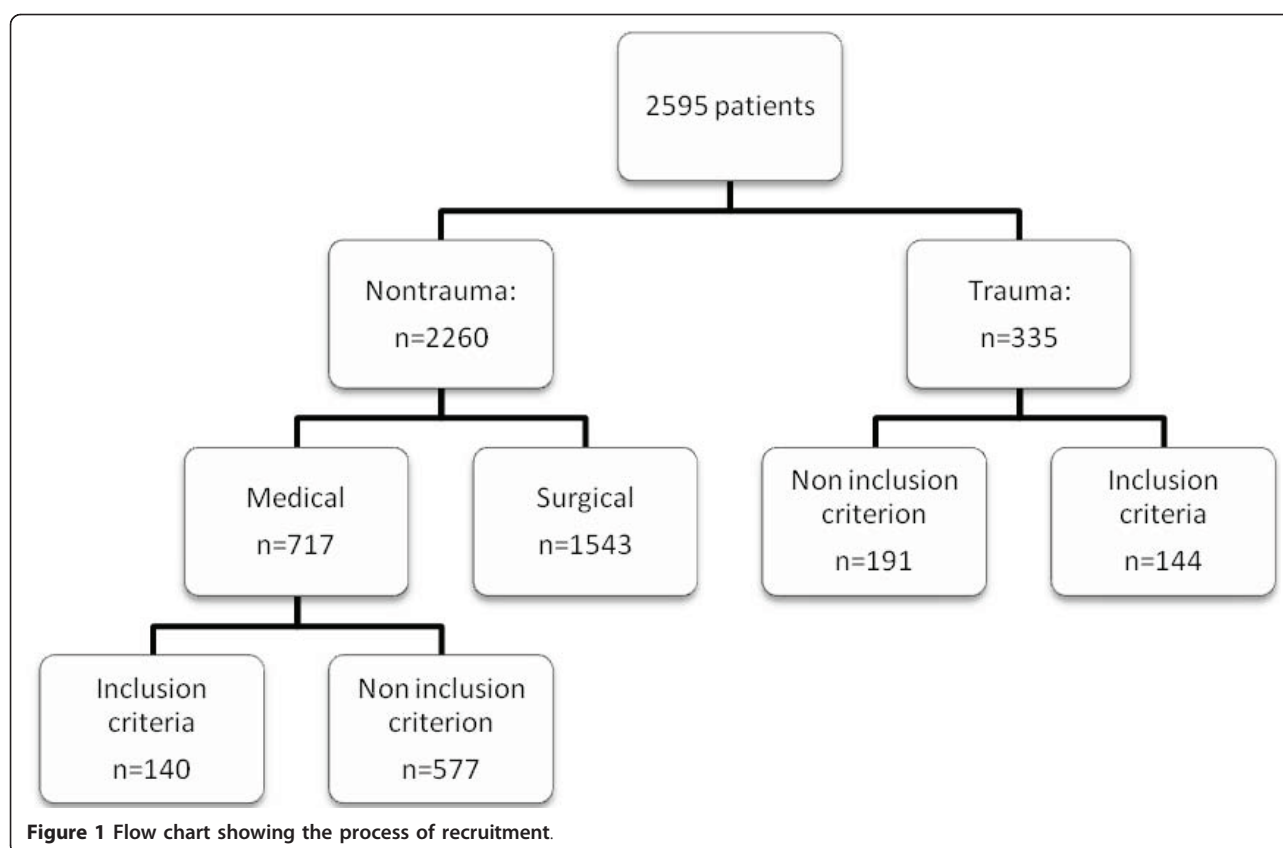
The overall results show that serum creatinine was normal ( $73 \pm 22 \mu\text{mol L}^{-1}$ ) and serum urea ( $8 \pm 4 \text{ mmol L}^{-1}$ ) was slightly higher than the normal limits, but with no difference between groups. One hundred, six patients had a CL<sub>CR</sub> above  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ , including 79 PT and 27 NPT ( $P < 0.0001$ ). Only 63 patients had a CL<sub>CR</sub> below  $60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$  with 15 PT and 48 NPT ( $P < 0.0001$ ), whereas nine patients had a CL<sub>CR</sub> below  $30 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ , including two PT.

The overall urinary creatinine excretion was  $929 \pm 678 \text{ mg } 24 \text{ h}^{-1} 1.73 \text{ m}^{-2}$  for women and  $1,369 \pm 685 \text{ mg } 24 \text{ h}^{-1} 1.73 \text{ m}^{-2}$  for men. There was a significant difference between the urinary creatinine excretion of PT and NPT patients ( $1,489 \pm 639$  vs  $969 \pm 688 \text{ mg } 24 \text{ h}^{-1} 1.73 \text{ m}^{-2}$  respectively,  $P < 0.001$ ). In the PT group, males had significantly higher urinary creatinine excretion than females ( $1,630 \pm 644$  vs  $1,067 \pm 392 \text{ mg } 24 \text{ h}^{-1} 1.73 \text{ m}^{-2}$ ,  $P < 0.001$ ).

The overall measured CL<sub>CR</sub> was  $108 \pm 57 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ . The CL<sub>CR</sub> was higher in PT patients than in NPT patients when using measured CL<sub>CR</sub> ( $131 \pm 56$  vs  $85 \pm 48 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$  respectively,  $P < 0.001$ ).

Most patients with increased CL<sub>CR</sub> (above  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ ) were PT patients as shown in Table 2. On the opposite, most patients with decreased CL<sub>CR</sub> (below  $60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ ) were NPT patients.

Patients with CL<sub>CR</sub> greater than  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$  were younger ( $40 \pm 16$  years vs  $56 \pm 18$  years), had a lower SAPS II score ( $43 \pm 14$  vs  $50 \pm 15$ ) and a higher male ratio as compared with patients presenting a CL<sub>CR</sub> lower than  $120 \text{ mL minute}^{-1}$ .



All factors presenting a statistical difference between hyperfiltration and hypofiltration subgroups (Table 2) were analyzed. Through a logistic regression analysis, including goodness of fit of the model, age and trauma were the only factors independently correlated to  $CL_{CR}$  (Table 3).

## Discussion

The present results comparing a population of hemodynamic stable PT patients to a population of hemodynamic stable NPT patients with steady state serum creatinine concentration with a normal creatinine serum value demonstrate that (i) PT patients exhibit dramatic variations of their  $CrCl$ ; (ii)  $CrCl$  is higher in PT patients than in NPT patients; (iii) Age and trauma are independently correlated factors to  $CL_{CR}$  in our study and in these study conditions.

Considering serum creatinine values, no significant difference was found between PT and NPT groups despite the variations of  $CrCl$ . These data demonstrated that a wide range of measured  $CL_{CR}$  variations exists and, therefore, confirm Hoste data obtained in critically ill patients with serum creatinine within normal range [16]. These authors demonstrated that “serum creatinine has a low sensitivity for detection of renal dysfunction”. Our results also revealed some opposite trends between  $CrCl$  and creatinine measurements, as some patients had

significantly lower values of serum creatinine for  $CL_{CR} > 60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$  than for  $CL_{CR} < 60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ . These data underline the inaccuracy of serum creatinine values in estimating the renal function.

Fifty-five percent of PT patients, and only 19% of NPT patients presented a measured  $CL_{CR}$  above  $120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ . In addition, only 10% of PT patients vs 34% of NPT patients presented a measured  $CL_{CR}$  below  $60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ . In clinical practice, the diagnosis of increased  $CL_{CR}$  as a surrogate marker of GFR is important and has largely been demonstrated in burn patients in the setting of antibiotics monitoring: ceftazidime, cefepime, vancomycin and amikacin [4,5,17,18]. In critically ill patients, high  $CL_{CR}$  required high doses of drugs, which are eliminated by the kidneys to obtain therapeutic concentration. Recently we confirmed the need for  $CL_{CR}$  monitoring in order to accurately monitor renal function and, therefore, to optimize the doses of antibiotics [4,19].

Our results demonstrate that age, gender, ideal body weight, severity index, trauma patients, and serum creatinine are factors for a  $CL_{CR}$  above normal ( $> 120 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ ), and for a moderate renal impairment ( $CL_{CR} < 60 \text{ mL minute}^{-1} 1.73 \text{ m}^{-2}$ ). Through a logistic regression analysis, only two factors (age and trauma patients) remained significantly correlated with a  $CL_{CR}$  above normal and for a moderate renal impairment. In the

**Table 2 Comparison of patients with different measured creatinine clearance (CL<sub>CR</sub>)**

	CL <sub>CR</sub> < 120 (n = 178)	CL <sub>CR</sub> > 120 (n = 106)	P*	CL <sub>CR</sub> < 60 (n = 63)	CL <sub>CR</sub> > 60 (n = 221)	P <sup>#</sup>	P <sup>§</sup>
Age (yr)	56 ± 18	40 ± 16	<0.0001	63 ± 15	46 ± 18	<0.0001	<0.0001
Weight (Kg)	74 ± 17	74 ± 14	NS	73 ± 19	74 ± 15	NS	NS
Sex (F/M)	64/114	24/82	0.03	35/28	161/60	NS	0.0032
Ideal body weight (Kg)	69 ± 12	72 ± 10	0.01	65 ± 11	71 ± 11	0.0002	<0.0001
SAPS 2	50 ± 15	43 ± 14	0.0002	54 ± 14	45 ± 15	<0.0001	<0.0001
Serum creatinine (μmol L <sup>-1</sup> )	76 ± 24	67 ± 19	0.0001	85 ± 25	69 ± 20	<0.0001	<0.0001
Diuresis/24H	2,400 ± 100	2,800 ± 1200	<0.0001	584 ± 224	1,418 ± 694		<0.0001
Measured creatinine clearance (mL min <sup>-1</sup> 1.73 m <sup>-2</sup> )	74 ± 30	166 ± 50	<0.0001	43 ± 11	127 ± 50	<0.0001	<0.0001
Simplified MDRD (mL minute <sup>-1</sup> 1.73 m <sup>-2</sup> )	96 ± 39	119 ± 45	<0.0001	81 ± 34	112 ± 43	<0.0001	<0.0001
Cockcroft-Gault (mL minute <sup>-1</sup> 1.73 m <sup>-2</sup> )	98 ± 39	130 ± 34	0.0005	80 ± 39	118 ± 36	<0.0001	<0.0001
Robert (mL minute <sup>-1</sup> 1.73 m <sup>-2</sup> )	72 ± 23	94 ± 23	<0.0001	59 ± 2	85 ± 2	<0.0001	<0.0001
PT <sup>b</sup> /NPT <sup>a</sup>	65/113	79/27	<0.0001	15/48	129/92	<0.0001	<0.0001

All data are expressed as ratio or mean ± SD.

\*The P-values correspond to comparison between subgroup with CL<sub>CR</sub> greater than 120 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup> versus subgroup with CL<sub>CR</sub> lower than 120 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup>.

<sup>#</sup>The P-values correspond to comparison between subgroup with CL<sub>CR</sub> greater than 60 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup> versus subgroup with CL<sub>CR</sub> lower than 60 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup>.

<sup>§</sup>The P-values correspond to comparison between less than 60 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup> more than 120 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup>.

<sup>a</sup>NPT = non-polytrauma patients.

<sup>b</sup>PT = polytrauma patients.

current results, 12% of elderly patients (over 65 years) have a CL<sub>CR</sub> greater than 120 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup>. The impact of age on CL<sub>CR</sub> is well known and this parameter was, therefore, introduced in the formulas estimating CL<sub>CR</sub> (Cockcroft-Gault, Robert and simplified MDRD) [13-15]. The decrease in glomerular filtration, the involution of nephronic units and the reduction of the renal blood flow explain the high frequency of renal impairment in elderly patients. However, it should be kept in mind that glomerular ageing is correlated to age in only

two-thirds of the patients, and this phenomenon accounts for the inaccuracy of the CL<sub>CR</sub> estimated by calculated formulae [20].

Current evidence suggests that PT (mainly, young patients without significant comorbidities) present with a CL<sub>CR</sub> increase. However, this phenomenon as received little attention in the literature, and dose modification are therefore rarely considered. The present results clearly demonstrate for the first time that trauma is a major factor for CL<sub>CR</sub> increase. Several factors may explain this increase in CL<sub>CR</sub> in PT patients. First, urinary creatinine excretion may be involved in such a phenomenon. A higher creatinine urinary excretion was observed in PT compared with NPT patients whereas serum creatinine was similar in both groups. However, the higher creatinine urinary excretion observed in PT patients was within a normal range. Serum protein variations may impact our results. However, all the patients had a serum protein value between 50 and 55 gL<sup>-1</sup>. It is, therefore, very unlikely that serum protein variations interfere with the present results. Also, regarding hemodynamics, CL<sub>CR</sub> were studied and measured at a steady state in both groups (that is, distant from the admittance). It should be noted that our patients were hemodynamically stable at the time of data collection with no sign of dehydration. Although interference due to some cephalosporin has been described when the creatininemia was measured by using the Jaffe method [21]; no changes in this parameter were observed during the overall period of the study. Sepsis can also reduce creatinine production as described in mice [22]. Moreover, in critically-ill patients, a positive

**Table 3 Logistic regression for different measured creatinine clearance**

	P*	Odd ratio (CI 95%)
CL <sub>CR</sub> > 120 mL minute <sup>-1</sup> 1.73 m <sup>-2</sup>		
Age	< 0.0001	0.95 (0.93 to 0.97)
SAPS 2	0.56	1.00 (0.98 to 1.04)
Ideal body weight	0.29	0.97 (0.93 to 1.02)
Gender	0.564	0.41 (0.12 to 1.4)
PT	0.0001	3.33 (1.8 to 6)
CL <sub>CR</sub> < 60 mL minute <sup>-1</sup> 1.73 m <sup>-2</sup>		
Age	< 0.0001	0.95 (0.93 to 0.97)
SAPS 2	0.36	0.98 (0.95 to 1.02)
Ideal body weight	0.3	1.03 (0.97 to 1.09)
Gender	0.8	0.84 (0.97 to 1.09)
NPT	0.02	2.39 (1.15 to 4.97)

PT = polytrauma patients, NPT non polytrauma patients.

\* The P-values correspond to logistic regression with CL<sub>CR</sub> greater than 120 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup> (C index = 0.56).

<sup>#</sup> The P-values correspond to logistic regression with CL<sub>CR</sub> lower than 60 mL minute<sup>-1</sup> 1.73 m<sup>-2</sup> (C index = 0.58).



fluid balance may lead to underestimation of the severity of AKI and delay the recognition of a 50% relative increase in sCr [23]. Finally, it should be hypothesized that humoral and inflammatory mechanisms encountered after severe trauma [24] or burn [1] are involved in the observed  $CL_{CR}$  increase.

The present study encountered some limitations. Increased  $CL_{CR}$  is related with enhanced renal elimination of circulating drugs. However, describing this phenomenon in terms of current available measures of the GFR at the bedside is still debated. In our study, the GFR was estimated by measured creatinine clearance on a 24-hour urine collection. However, the gold standard for GFR assessment is the measure of inulin clearance [25] but the cost and complexity of this tool limits its application in routine. Another limitation is the lack of consensus regarding the upper limit of normal GFR. However, increasing data support the concept of increased GFR in PT patients, and several reports demonstrated subtherapeutic concentrations of drugs in PT patients [5,26]. Also, a cross-sectional single 24-hr measure of  $CL_{CR}$  at 10 days in relatively stable patients was performed but fast modifications of kidney function may occur and there is a need for a continuous re-evaluation. Finally, some other factors may influence our results. In particular, fluid status, cardiac output may be significantly altered from baseline. However, whatever the causes of these alterations, the ATLS (Advanced Trauma Life support) principles are applied in our institution regarding resuscitation of PT patients. We, therefore, believe that the current results are broadly representative of the population of PT patients. It could be argued that the external validity of this single-center study may be limited. However, our findings may be relevant to the vast majority of level I trauma centers, provided that ATLS principles are applied in these institutions.

## Conclusions

In hemodynamic ICU stable patients with steady state serum creatinine concentration,  $CL_{CR}$ , which is a surrogate marker of GFR, is higher in polytrauma patients than in other critically ill patients. In ICU patients, the drug monitoring must take into account the glomerular filtration rate. The measure of  $CL_{CR}$  should be routinely proposed for PT patients in order to adjust dose regimen, especially for drugs with renal elimination (beta-lactams, ceftazidime, cefepime, piperacillin, vancomycin, aminoglycosides, and so on).

## Key messages

- In ICU patients with normal serum creatinine,  $CL_{CR}$ , is higher in trauma than in non-trauma patients.

- The measure of  $CL_{CR}$  should be proposed in routine for ICU patients in order to adjust dose regimen, especially for drugs with renal elimination.
- Age and trauma were the only factors independently correlated to  $CL_{CR}$ .
- Glomerular filtration rate should be measured in ICU patient to detect renal filtration abnormalities.
- Serum creatinine is not a good marker for renal function estimation.

## Abbreviations

AKI: Acute Kidney Injury;  $CL_{CR}$ : creatinine clearance; GFR: glomerular filtration rate; ICU: intensive care unit; NPT: non polytrauma patients; PT: polytrauma patients; sMDRD, Modification of Diet in Renal Disease index; SOFA: score, Sequential Organ Failure Assessment score.

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## Authors' contributions

AJ and IT carried out the serum creatinine measurement and calibration. SR, AB and TS carried out the patients' inclusions. KA helped to draft the manuscript and reviewed the intellectual content. BG, SS, OF and KS participated in the design of the study and helped to draft the manuscript. JMC and VM conceived of the study, and participated in its design and coordination and helped to draft the manuscript and performed the statistical analysis. All authors read and approved the final manuscript.

## Competing interests

The authors declare that they have no competing interests.

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## References

1. Conil JM, Georges B, Fourcade O, Seguin T, Lavit M, Samii K, Houin G, Tack I, Saivin S: **Assessment of renal function in clinical practice at the bedside of burn patients.** *Br J Clin Pharmacol* 2007, **63**:583-594.
2. Loirat P, Rohan J, Baillet A, Beaufils F, David R, Chapman A: **Increased glomerular filtration rate in patients with major burns and its effect on the pharmacokinetics of tobramycin.** *N Engl J Med* 1978, **299**:915-919.
3. Conil JM, Georges B, Breden A, Segonds C, Lavit M, Seguin T, Coley N, Samii K, Chabanon G, Houin G, Saivin S: **Increased amikacin dosage requirements in burn patients receiving a once-daily regimen.** *Int J Antimicrob Agents* 2006, **28**:226-230.
4. Conil JM, Georges B, Lavit M, Laguerre J, Samii K, Houin G, Saivin S: **A population pharmacokinetic approach to ceftazidime use in burn patients: influence of glomerular filtration, gender and mechanical ventilation.** *Br J Clin Pharmacol* 2007, **64**:27-35.
5. Conil JM, Georges B, Mimoz O, Dieye E, Ruiz S, Cougot P, Samii K, Houin G, Saivin S: **Influence of renal function on trough serum concentrations of piperacillin in intensive care unit patients.** *Intensive Care Med* 2006, **32**:2063-2066.
6. Larsson R, Bodemar G, Kagedal B, Walan A: **The effects of cimetidine (Tagamet) on renal function in patients with renal failure.** *Acta Med Scand* 1980, **208**:27-31.
7. Hull JH, Hak LJ, Koch GG, Wargin WA, Chi SL, Mattocks AM: **Influence of range of renal function and liver disease on predictability of creatinine clearance.** *Clin Pharmacol Ther* 1981, **29**:516-521.

8. Papadakis MA, Arief AL: **Unpredictability of clinical evaluation of renal function in cirrhosis. Prospective study.** *Am J Med* 1987, **82**:945-952.
9. Ferreira FL, Bota DP, Bross A, Melot C, Vincent JL: **Serial evaluation of the SOFA score to predict outcome in critically ill patients.** *JAMA* 2001, **286**:1754-1758.
10. Antonelli M, Moreno R, Vincent JL, Sprung CL, Mendoca A, Passariello M, Riccioni L, Osborn J: **Application of SOFA score to trauma patients. Sequential Organ Failure Assessment.** *Intensive Care Med* 1999, **25**:389-394.
11. Vincent JL, de Mendonca A, Cantraine F, Moreno R, Takala J, Suter PM, Sprung CL, Colardyn F, Blecher S: **Use of the SOFA score to assess the incidence of organ dysfunction/failure in intensive care units: results of a multicenter, prospective study. Working group on "sepsis-related problems" of the European Society of Intensive Care Medicine.** *Crit Care Med* 1998, **26**:1793-1800.
12. Froissart M, Rossert J, Jacquot C, Paillard M, Houillier P: **Predictive performance of the modification of diet in renal disease and Cockcroft-Gault equations for estimating renal function.** *J Am Soc Nephrol* 2005, **16**:763-773.
13. Cockcroft DW, Gault MH: **Prediction of creatinine clearance from serum creatinine.** *Nephron* 1976, **16**:31-41.
14. Robert S, Zarowitz BJ, Peterson EL, Dumler F: **Predictability of creatinine clearance estimates in critically ill patients.** *Crit Care Med* 1993, **21**:1487-1495.
15. Levey AS, Bosch JP, Lewis JB, Greene T, Rogers N, Roth D: **A more accurate method to estimate glomerular filtration rate from serum creatinine: a new prediction equation. Modification of Diet in Renal Disease Study Group.** *Ann Intern Med* 1999, **130**:461-470.
16. Hoste EA, Damen J, Vanholder RC, Lameire NH, Delanghe JR, Van den Hauwe K, Colardyn FA: **Assessment of renal function in recently admitted critically ill patients with normal serum creatinine.** *Nephrol Dial Transplant* 2005, **20**:747-753.
17. Conil JM, Georges B, Lavit M, Seguin T, Tack I, Samii K, Chabanon G, Houin G, Saivin S: **Pharmacokinetics of ceftazidime and cefepime in burn patients: the importance of age and creatinine clearance.** *Int J Clin Pharmacol Ther* 2007, **45**:529-538.
18. Conil JM, Favarel H, Laguerre J, Bouchet A, Chabanon G, Cazal L, Bodnar M, Rouge D, Virenque C, Costagliola M: **[Continuous administration of vancomycin in patients with severe burns].** *Presse Med* 1994, **23**:1554-1558.
19. Conil JM, Georges B, de Lussy A, Khachman D, Seguin T, Ruiz S, Cougot P, Fourcade O, Houin G, Saivin S: **Ciprofloxacin use in critically ill patients: pharmacokinetic and pharmacodynamic approaches.** *Int J Antimicrob Agents* 2008, **32**:505-510.
20. O'Connell MB, Dwinell AM, Bannick-Mohrland SD: **Predictive performance of equations to estimate creatinine clearance in hospitalized elderly patients.** *Ann Pharmacother* 1992, **26**:627-635.
21. Grottsch H, Hajdu P: **Interference by the new antibiotic ceftazidime and other cephalosporins in clinical laboratory tests, with special regard to the "Jaffe" reaction.** *J Clin Chem Clin Biochem* 1987, **25**:49-52.
22. Doi K, Yuen PS, Eisner C, Hu X, Leelahavanichkul A, Schnermann J, Star RA: **Reduced production of creatinine limits its use as marker of kidney injury in sepsis.** *J Am Soc Nephrol* 2009, **20**:1217-1221.
23. Macedo E, Bouchard J, Soroko SH, Chertow GM, Himmelfarb J, Ikizler TA, Paganini EP, Mehta RL: **Fluid accumulation, recognition and staging of acute kidney injury in critically-ill patients.** *Crit Care* 2010, **14**:R82.
24. Hoen S, Asehnoune K, Brailly-Tabard S, Mazoit JX, Benhamou D, Moine P, Edouard AR: **Cortisol response to corticotropin stimulation in trauma patients: influence of hemorrhagic shock.** *Anesthesiology* 2002, **97**:807-813.
25. Stevens LA, Coresh J, Greene T, Levey AS: **Assessing kidney function-measured and estimated glomerular filtration rate.** *N Engl J Med* 2006, **354**:2473-2483.
26. Georges B, Conil JM, Seguin T, Ruiz S, Minville V, Cougot P, Decun JF, Gonzalez H, Houin G, Fourcade O, Saivin S: **Population pharmacokinetics of ceftazidime in intensive care unit patients: influence of glomerular filtration rate, mechanical ventilation, and reason for admission.** *Antimicrob Agents Chemother* 2009, **53**:4483-4489.

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