Changes in Estimated Glomerular Filtration Rate, Biochemical and Hematological Profile of Workers in Spray Painting Industry: A Cross-Sectional Study

Christopher E. Ekpenyong, PhD; Nseobong N. Nkereuwem, BS

Department of Physiology, University of Uyo, Uyo, Akwa Ibom, Nigeria

*Corresponding author
Christopher E. Ekpenyong, PhD
Faculty of Basic Medical Sciences, Department of Physiology, University of Uyo, Uyo, Akwa Ibom, Nigeria;
E-mail: chrisvon200@yahoo.com; chrisvon300@yahoo.com

ABSTRACT

Introduction
There is a global increase in occupational exposure to solvents, some of which are suspected to cause acute or chronic toxic nephropathies in humans. However, limited studies have been done to evaluate the systemic effects of exposure to some of the commonly used solvents such as paints.

Aim
The aim of the present study was to assess the effect of chronic exposure to paint fumes on renal and hepatic functions of industrial spray painters.

Methodology
In this cross-sectional study, 49 occupationally exposed male industrial spray painters who had served for greater than 5 years were evaluated for changes in renal, hepatic and hematological indices using standard instruments and results were compared with levels in the unexposed (sex and age-matched) participants.

Results
Significant changes in markers of renal, hepatic and hematological functions were observed in the exposed compared with unexposed participants including significant decrease in estimated glomerular filtration rate (eGFR) and serum levels of potassium (K⁺) and chloride (Cl⁻), and significant increases in serum levels of creatinine (Cr), sodium (Na⁺), urea (Ur) and uric acid (UA) in the exposed compared to levels in the unexposed group. Abnormal serum levels of hepatic enzymes (AST, ALT and ALP) and hematological indices (PCV, total-RBC, neutrophils, basophils, monocytes and lymphocytes) were also observed in the exposed compared to levels in the unexposed participants.

Conclusion
Prolonged exposure to paint fumes may be associated with a significant risk for hepato-renal dysfunction and hematotoxicity. Preventive measures should include limiting exposure and using antioxidant medications.

Keywords
Spray painting; Toxicity; Workers; Kidney; Liver; Blood cells.

Abbreviations
ALT: Alanine transaminase; ALP: Alkaline phosphatase; AST: Aspartate transaminase; eGFR: Estimated glomerular filtration rate; C-G: Cockcroft-Gault; MDRD: Modification of diet in renal disease; Na+/K+/ATPase: Sodium potassium adenosine triphosphatase; PH: Hydrogen Concentration; PCV: Packed cell volume; RBC: Red blood cells; ROS: Reactive oxygen species; UA: Uric acid; Ur: Urea.
INTRODUCTION

O ccupational and environmental factors have been recognized as important contributors to the rising incidence of kidney disease in the general population, especially occupations that exposed workers to nephrotoxic substances. Exposure to nephrotoxic substances accounts for about 25% of all cases of acute or chronic renal failure. Some nephrotoxic substances commonly encountered at workplace include organic solvents, copper, chromium, tin, lead, mercury, welding fumes and spray painting fumes.

Some of these chemicals may exist alone or in combination in pigments or in solvents. A typical example is paint which consist of a mixture of pigments and solvents. Some pigments found in paints include: white lead, titanium white, red lead, venetian red, chrome red, zinc chromate and chrome yellow; while solvents such as hydrocarbon solvents (aliphatic and aromatic hydrocarbons), halogenated solvents (chlorinated hydrocarbons) and oxygenated solvents (alcohols, glycol ethers, ketones and esters). Furthermore, some volatile organic compounds (VOCs) are added to paints to attain the desired quality such as to enhance dryness due to their volatile nature. VOCs and other constituents vaporized easily and emit gasses that cause widespread environmental pollution. The percentage composition of these chemicals in paints is dependent on the physiochemical needs of the paints.

Millions of people around the world are exposed to the toxic constituents of spray painting fumes that contaminate the air around them and the worst affected are workers who are in occupations with the highest burden of paint fumes exposure such as the spray painters. When inhaled, the chemicals are readily absorbed, metabolized and distributed to various organs in the body. The active metabolites undergo further toxico-kinetic modifications and leading to generation of reactive oxygen species (ROS), oxidative tissue damage and altered structure and function. It is now known that exposure to chemicals with very diverse chemical structures and compositions such as paints may produce a wide range of health effects due to the synergistic actions of the different components, including effect on the kidney, liver and blood cells.

In view of the drawbacks of previous studies such as isolated use of serum biochemical indices of renal function (creatinine, blood urea nitrogen, uric acid) with no adjustment for known confounders such as age, gender, body mass index and dietary factors, and the fact that the toxicokinetic properties of nephrotoxic chemicals may be modified by individual susceptibility factors (age, gender, genetic susceptibility, exposure concentration, duration of exposure, race and ethnicity).

The present study aimed to assess the effect of exposure to spray painting fumes on estimated glomerular filtration rate (eGFR), biochemical and hematological indices of spray painters in two cities in Akwa Ibom State.

MATERIALS AND METHODS

Study Design and Participants

This cross-sectional study was carried out between June 2017 and May 2018 among industrial spray painters whose workshops are located in Uyo and Ikot Ekepene Local Government Areas, Akwa Ibom State, Nigeria. A total of 97 (49 exposed and 48 unexposed) male participants of the 122 initially invited to participate were recruited for the study.

Others were excluded from the study for not meeting the inclusion criteria.

Exclusion criteria were as follows; age outside the study age (18 to 40 years), workers who had served for <5 years on the job, decline participation, inappropriate completion of the study questionnaire, presence of any metabolic syndrome clusters or on medication for any of the metabolic syndrome, concomitant exposure to other solvents besides paints, smoking, binge drinking of alcohol, dehydration, dieting and suffering from any other debilitating disease.

Study design and experimental protocols were approved by the Institutional Human Ethics Committee and participants signed written informed consent. Participants were briefed about the study and participation was voluntary. They were also allowed to withdraw from participation at any stage of the study.

The study was conducted according to the rules set forth in the Declaration of Helsinki, governing the conduct of human research.

Anthropometric Measures

Anthropometric indices measured in this study were weight (kg) and height (m) with light clothing and without shoes. Weight was measured using a portable weighing scale to the nearest 0.1 kg and height was measured using a height measuring tape to the nearest 0.1 cm. All measurements were performed twice and the average of the two measurements was used. Body mass index was calculated from the formula BMI= weight (kg)/height(m)^2.

Biochemical Measures

Venous blood samples were obtained from all participants for biochemical analysis including renal, hepatic and hematological evaluations.

Renal indices measured included electrolytes (potassium (K’), sodium (Na’), and chloride (Cl’)). Others were serum creatinine (Cr), urea (Ur) and uric acid (UA) levels. Serum Na’ and K’ were measured using flame photometer, while serum Cr levels were assessed according to Jaffe Kinetic method as described elsewhere. Serum liver enzymes (Alanine aminotransferase (ALT), Aspartate aminotransferase (AST), Alkaline phosphatase (ALP)
and bilirubin total and conjugated) were also measured.

Hematological indices measured included packed cell volume (PCV), hemoglobin concentration (Hb), Total red blood cell (RBC-total), Mean cell hemoglobin (MCH), Mean cell hemoglobin concentration (MCHC) white blood cell (WBC)-total and differentials and platelets. Measurements were carried out within 2 hrs of blood sample collection using the SYSMEX KX-21 Automated Haematology Analyzer (Kobe, Japan).

All measurements were done using standard methods and the quality assurance was strictly complied with. eGFR was calculated using Modification of Diet in Renal Disease (MDRD) and Cockroft and Gault Equations as follows:

**MDRD Equation:**

eGFR (mL/minute) = 186×(Plasma Creatinine\(^{-1.154}\))×Age\(^{-0.203}\)

multiply by 0.742 if female

and

multiply by 1.210 if African/American

**Cockroft-Gault Equation:**

eGFR (mL/minute) = \frac{140-\text{age (years)}×\text{weight (kg)}}{72×\text{serum creatinine}}

multiply by 0.85 if female

**STATISTICAL ANALYSIS**

The hepato-renal and hematological indices of the participants (exposed and unexposed) were computed, and results expressed as mean±standard deviation (SD). Significant differences in hepato-renal and hematological indices between groups were established using independent *t*-test with *p*<0.05 level of significance. Computations of results were done using Statistical Package for Social Sciences (SPSS) version 22.0, while graphical representations of results were done throughout using graph pad Prism.

**RESULTS**

The socio-demographic characteristics of the 97 participants showed that 49 (50.5%) were occupationally exposed to paint fumes while 48 (49.5%) were not exposed. Of the exposed participants, 36.7% were married, 55.1% were single while 8.2% were divorced. Most of the exposed participants (52.1%) were aged between 26 and 35 years, worked for 5-10 years, and had secondary level of education (77.6%). Socio-demographic variables of the exposed participants were not significantly different from those of the unexposed participants (Table 1).

Table 1. Socio-Demographic Characteristics of Study Participants

| Socio-Demographic Variables | Exposed (49) | Unexposed (48) | Total | p-value |
|-----------------------------|-------------|---------------|-------|---------|
| Marital Status              |             |               |       |         |
| Married                     | 18 (36.7)   | 21 (43.8)     | 39 (40.2) | 0.618   |
| Single                      | 27 (55.1)   | 25 (52.1)     | 52 (53.6) |         |
| Divorced                    | 04 (8.2)    | 02 (4.2)      | 06 (6.2)  |         |
| Age (years)                 |             |               |       |         |
| 18-25                       | 15 (30.6)   | 18 (37.5)     | 33 (34.0) | 0.720   |
| 26-35                       | 27 (55.1)   | 25 (52.1)     | 52 (33.6) |         |
| >35                         | 04 (8.2)    | 04 (8.3)      | 08 (8.2)  |         |
| Mean Age                    | 27.45 years | 27.02 years   | 54.47   | 0.454   |
| Duration on the Job         |             |               |       |         |
| 5-10 years                  | 38 (77.6)   | 33 (66.8)     | 71 (73.2) |         |
| >10 years                   | 11 (22.4)   | 14 (31.2)     | 26 (26.8) |         |
| Educational Level           |             |               |       |         |
| Primary                     | 02 (4.1)    | 01 (2.1)      | 03 (3.1)  | 0.152   |
| Secondary                   | 38 (77.6)   | 44 (91.7)     | 82 (84.5) |         |
| Tertiary                    | 09 (18.4)   | 03 (6.2)      | 12 (12.4) |         |

Table 2. Anthropometric Indices

| Adiposity Indices | Exposed | Unexposed | p-value |
|-------------------|---------|-----------|---------|
| Weight (Kg)       | 67.59±3.42 | 66.89±5.26 | 0.4381 |
| Height (m²)       | 1.65±0.96  | 1.66±0.88  | 0.9575 |
| BMI (Kg/m²)       | 24.83±6.43 | 24.27±8.01 | 0.7047 |

Table 3 shows significant (*p*<0.05 or *p*<0.01) differences in some serum biochemical indices of renal function between the exposed and unexposed participants including serum K⁺ which significantly (*p*<0.01) decreased in the exposed group compared to level in the unexposed group. Also, serum Na⁺ level significantly (*p*<0.05) increased in the exposed compared to level in the unexposed group.

Table 3. Serum Biochemical Indices of Renal Function

| Adiposity Indices | Exposed | Unexposed | p-value |
|-------------------|---------|-----------|---------|
| Na⁺ (mmol/L)      | 144.27±6.48 | 140.70±4.11 | 0.034* |
| K⁺ (mmol/L)       | 3.32±0.25   | 4.36±0.07   | 0.0001**|
| Cl⁻ (mmol/L)      | 99.90±3.95  | 100.55±4.30 | 0.378  |
| Glucose (mmol/L)  | 6.33±1.22   | 6.39±1.31   | 0.8159 |
| Uric Acid (mmol/L)| 7.89±0.95   | 6.92±1.02   | 0.0001**|
| pH                | 7.35±0.08   | 7.39±1.12   | 0.8583 |

Values are reported as Mean±Standard deviation (SD).

*Significant at 5% (*p*<0.05), **Significant at 1% (*p*<0.01)
Furthermore, chronic exposure to paint fumes was associated with a significant increase ($p<0.05$) in serum levels of hepatic enzymes (ALT, AST, ALP), total bilirubin and unconjugated bilirubin compared to levels in the unexposed group (Table 4).

Table 4. Liver Function Indices of Exposed and Unexposed Participants

| Variables | Exposed (µ/L) | Unexposed (µ/L) | $p$-value |
|-----------|--------------|----------------|-----------|
| Alanine Amino transferase (ALT) | 21.73±14.69 | 5.70±2.49 | <0.001*** |
| Aspartate Amino transferase (AST) | 29.82±14.64 | 6.80±1.20 | <0.001*** |
| Alkaline Phosphatas (ALP) | 110.37±27.02 | 56.15±15.29 | <0.0001** |
| Bilirubin Total (µmol/L) | 14.42±7.5 | 10.26±4.00 | 0.028* |
| Conjugated (µmol/L) | 5.96±3.14 | 6.42±8.54 | 0.787 |
| Unconjugated (µmol/L) | 8.53±5.43 | 7.73±3.11 | 0.042 |

Values are reported as Mean±Standard deviation (SD).
*Significant at 5% ($p<0.05$), **Significant at 1% ($p<0.01$)

Table 5 shows that there were significant ($p<0.05$) decreases in some hematological indices including packed cell volume (PCV) and total red blood cells (RBC) compared to levels in the unexposed group. However, MCH significantly ($p<0.05$) increased in the exposed compared to levels in the unexposed group.

Table 5. Hematological Indices of Exposed and Unexposed Participants

| Variables | Exposed | Unexposed | $p$-value |
|-----------|---------|-----------|-----------|
| PCV (1%) | 41.63±3.26 | 43.85±3.00 | 0.019* |
| WBC-T otal (x109/L) | 5.22±1.20 | 5.73±0.91 | 0.195 |
| RBC-T otal (x1012/L) | 4.69±0.53 | 5.02±0.25 | 0.014* |
| Platelet (x109/L) | 153.10±15.70 | 156.25±13.13 | 0.463 |
| MCHC (g/dL) | 33.48±1.62 | 31.75±1.37 | 0.350 |
| MCV (fl) | 90.10±12.64 | 87.35±3.75 | 0.350 |
| MCH (pg) | 30.00±3.92 | 27.65±1.50 | 0.014* |
| Basophil (%) | 0.18±0.39 | 0.51±0.49 | 0.009** |

Values are reported as Mean±Standard deviation (SD).
*Significant at 5% ($p<0.05$), **Significant at 1% ($p<0.01$)

Figure 1 shows that serum Cr was significantly ($p<0.01$) increased in the exposed than unexposed group, whereas eGFR using G-C and modification of diet in renal disease (MDRD) equations were significantly ($p<0.01$) lower in the exposed than unexposed group (Figure 2).

Serum UA and Ur were significantly ($p<0.01$) higher in the exposed compared to levels in the unexposed group (Figure 3).

Total basophil and neutrophil count significantly ($p<0.01$) decreased in the exposed than unexposed group (Figure 4). Total lymphocyte counts significantly ($p<0.01$) increased in the exposed than unexposed group whereas total monocyte counts significantly ($p<0.01$) decreased in exposed than unexposed (Group 5).
RESULTS

Results of the present study showed that prolonged exposure to paint fumes could be detrimental to hepato-renal endpoints and hematopoietic functions.

Evidences supporting this notion are three fold; first, the observed significant alterations in renal indices in the exposed compared to levels in the unexposed group, including the significant increase in mean serum Na+, Cr, UA and Ur levels and significant decrease in mean serum levels of K+ and eGFR in exposed compared to levels in the unexposed group; second, the significant increase in mean serum levels of hepatic enzymes (ALT, AST, ALP), total bilirubin and un-conjugated bilirubin; third, the significant derangements in several hematological indices in the exposed compared with the unexposed participants.

The above findings are consistent with previous studies that showed that prolonged exposures to environmental/occupational solvents are important contributors to the rising incidences of hepato-renal and hematologic disorders in the general population.3

Paint is a mixture of pigments (white lead, venetian red, zinc, chromate, etc.) and solvents (gasoline, turpentine and coal hydrocarbons). These substances have been shown to have demonstrable nephrotoxic, hepatotoxic and hematotoxic effects similar to the results of the present study. Accordingly, Al-Ghamdi,1 studied the kidney function of car painters who had prolonged exposure to paint fumes and found a significant higher mean serum value of Cr, Ur and UA than levels in the unexposed groups.

Ravnklov4 and Alsuwaida5 provided strong evidences for the association between prolonged exposure to solvents and nephrotoxicity. Their study findings were in good agreement with recent findings demonstrating solvent-induced decrease in eGFR, azotemia, electrolyte abnormalities and hematotoxicity.3 Likewise a recent study that showed significant increases in serum levels of liver enzymes, including AST, ALT, ALP, total bilirubin and marked histo-morphological changes following prolonged exposure to solvent.9 Also, evidence of increased lipid peroxidation (raised serum MDA) was reported.

Similar to findings of the present study, a plethora of research shows significant correlation between exposure to solvents and hematotoxicity.10,12 Although largely hypothetical, a consideration of existing research findings suggest a number of pathophysiological mechanisms underlying the adverse hepato-renal effects of solvents (e.g., paints and constituents), including the generation of reactive oxygen species (ROS) following their breakdown to reactive metabolites by Cytochrome P450 monooxygenase and beta-lyase mediated pathways and subsequent interaction of ROS with renal tissues to cause damage to renal glomerules, tubular epithelial cells, and macromolecules such as sodium/potassium/adenosine triphosphatase (Na+/K+/ATPase) and alter their activities with a resultant renal injury.13

Na+/K+/ATPase is an enzyme involves in electrolyte and water homeostasis. It regulates serum Na+, K+ and Cl by pumping 3 Na+ out of the cell for every K+ pumped into the cell, hence it’s regarded as a 3 ion co-transport system. Available evidence suggests that this enzyme is inhibited by metabolites from solvent constituents and leading to altered electrolytes, water and acid-base homeostasis including increased serum Na+ and decreased serum K+ as observed in the present study.

Reactive metabolites from paints and other solvents can also cause physiochemical damage to the glomerular membrane and tubules due to their high lipophilicity.7

Such damage has been shown to alter the functional integrity of the membrane, impairs Na+/K+/ATPase activity and several other enzymes involved in ionic and acid-base balance. Also, associated tubular damage can cause tubular dysfunction, impairs renal excretory function, macula densa cell signaling mechanisms, renal hemo-dynamics/autoregulation and poor handling of metabolic waste products with a resultant increase in serum Cr, Ur and UA as observed in the present study.

Likewise, glomerular damage could interfere with dynamics of glomerular filtration and ultimately leading to decrease GFR as observed in the present study.

Induction of OS by metabolites from paints has also been reported to cause disruption of the immune system and induced inflammation. There is associated auto-immunity, T-cell dysfunction, suppression of immune function, increased post-glomerular resistance and decreased GFR following exposure to solvents as reported previously.13 At this point time, it should be noted that solvent-induced kidney injury is not an isolated process, evidence has been garnered suggesting that solvent-induced kidney injury could be extended to cause the dysfunction of other remote organs such as the liver, blood cells, heart, lungs, intestines and brain via a pro-inflammatory mechanism mediated by inflammatory mediators such as neutrophil cell migration and cytokine expression and increased oxidative stress.13 This can
partly explain the associated hepatic and hematological disorders including cirrhosis, hepatitis, necrotic inflammatory hepatitis, cholestatic liver disorders, anemia, leukocytosis, leukopenia and changes in liver enzymes reported previously in solvent exposed workers (AST, ALT, ALP), and some of which are observed in the present study. These findings are consistent with data of Golab et al., White et al. and Lane et al. who independently demonstrated hepatic dysfunction as a consequence of acute kidney injury. The hematological profile of the exposed subjects showed that there were significant decreases in the erythropoiesis modulated hemocytic variables (PCV, total RBC, MCHC, MCH and MCV), total WBC, acute inflammatory cells (neutrophils, basophils and monocytes) and platelets, whereas a significant increase was found in the level of chronic inflammatory cells (Lymphocytes). These findings indicate that induction of chronic inflammation could serve as one of the pathophysiological mechanisms of paints-induced hematotoxicity. Other mechanisms of solvent-induced hepatic and hematologic disorders include induction of cellular degeneration and down regulation of gene expression.

**CONCLUSION**

Results of the present study suggest that prolonged exposure to paint fumes may be associated with a significant risk for hepato-renal dysfunction and blood cell disorders.

Preventive measures should include limiting exposure and using antioxidant medications. Pre-employment evaluation of renal and hepatic functions should be conducted and repeated periodically in potentially exposed and the already exposed workers since some paint constituents have been shown to initiate hepato-renal injury and hematotoxicity or worsen extant hepato-renal function impairment or both.

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**CONFLICTS OF INTEREST**

The authors declare that they have no conflicts of interest.

**REFERENCES**

1. Ekpenyong CE, Inyang MAE. A cross-sectional survey of estimated glomerula filtration rate, acid-base balance and electrolyte status among workers exposed to petroleum products. *J Nephrol Kid Dis.* 2017; 1 (2): 1008.

2. Voss JU, Roller M, Mangelsdorf I. Nephrotoxicity of organic solvents. A literature survey. Federal Institute for Occupational Safety and Health. https://www.baua.de/DE/Angebote/Publikationen/Berichte/Gd8.pdf?__blob=publicationFile. 2003. 1-25. Accessed May 30, 2019.

3. Nerbass FB, Pecoits-Filho R, Clark WF, Sontrup JM, McIntyre CW, Mois L. Occupational heat stress and kidney health from farms to factories. *Kidney Int Rep.* 2017; 2(6): 998-1008. doi: 10.1016/j.ekir.2017.08.012

4. Ravnkov U. Hydrocarbon exposure may cause glomerulonephritis and worsen renal failure: Evidence based on Hill’s criteria of causality. *QJM.* 2000; 93: 551-556. doi: 10.1093/qjmed/93.8.551

5. Perez CA, Bosia JD, Cantore MS, et al. Liver damage in workers exposed to hydrocarbons. *Gastroenteral Hepatol.* 2006; 29: 334-337.

6. Okoro AM, Ani Ej, Ibu JO, Akpogomeh BA. Effect of petroleum products inhalation on some haematological indices of fuel attendants in Calabar Metropolis, Nigeria. *Nig J Physiol Sci.* 2006; 21: 71-75. doi: 10.4314/njps.v21i1-2.53954

7. Al-Ghamdi SS. Occupational exposure to paints causes impairment of kidney function. *J Environ Protect.* 2011; 2: 533-536. doi: 10.4236/jep.2011.25061

8. Alsuwaida A, Jet fuel intoxication and acute renal failure. *Saudi J Kidney Dis Transpl.* 2010; 21(1): 150-152.

9. Ekpenyong CE, Bassey G. Investigation of the effect of cymbopogon citratus decoctions on gasoline vapour-induced lipid peroxidation and hepatotoxicity in rats. *J Inter Res Med Pharm Sci.* 2016; 10(4): 171-181.

10. Ajugwo AO, Adias TC, Aghatise K, Fadairo JK, Nyenke CU. Reduced haematological indices in auto-mechanics and fuel attendants in Elele Nigeria. *Am J Med Biol Res.* 2014; 2: 1-4. doi: 10.12691/ajmr-2-1-1

11. Sahb AAA. Hematological assessment of gasoline exposure among petrol filling workers in Baghdad. *J Fac Med Baghdad.* 2011; 53: 396-400.

12. Tunsaringkarn T, Soogarun S, Palasuwa A. Occupational exposure to benzene and changes in hematological parameters and urinary trans, trans-muconic acid. *Int J Occup Environ Med.* 2013; 4: 45-49.

13. Ekpenyong CE, Asuquo EA. Recent advances in occupational and environmental health hazards of workers exposed to gasoline compounds: Toxicology review. *Int J Ocs Med Environ Health.* 2017; 30(1): 1-26. doi: 10.13075/ijomeh.1896.00800

14. Makris K, Loukia S. Acute injury: Definition, pathophysiology and clinical phenotypes. *Clin Biochem Rev.* 2016; 37 (2): 85-98.

15. Golab F, Kadhodaee M, Zahmatkesh M, et al. Ischemic and non-ischemic acute kidney injury cause hepatic damage. *Kidney Int.* 2009; 75: 783-792. doi: 10.1038/ki.2008.683

16. White LE, Chaudhary R, Moore LJ, Moore FA, Hassoum HT.
Surgical sepsis and organ crosstalk: the role of the kidney. *J Surg Res.* 2011; 167: 306-315. doi: 10.1016/j.jsr.2010.11.923

17. Lane K, Dixon JJ, MacPhee IA, Philip BJ. Renohepatic crosstalk: Does acute kidney injury cause liver dysfunction? *Nephrol Dial Transplant.* 2013; 28: 1634-1647. doi: 10.1093/ndt/gft091

18. Ravnskov U. Experimental glomerulonephritis induced by hydrocarbon exposure: A systemic review. *BMC Nephrol.* 2005; 6: 15-18. doi: 10.1186/1471-2369-6-15