

Letters to the Editor – Brief papers about basic research or clinical experiences

Capturing the diagnosis of chronic kidney disease in the electronic medical record and its influence on therapeutic management[☆]

Registro del diagnóstico de insuficiencia renal crónica en la historia clínica electrónica y su influencia en el manejo terapéutico

Dear Editor,

Chronic kidney disease (CKD) affects 9% of the adult population in Spain.^{1,2} The majority of patients with CKD are managed by primary care physicians,³⁻⁶ and early detection is essential for adopting the measures required to slow its progression.^{7,8} At the primary care clinics, the physician includes in the electronic medical record the diagnosis of chronic kidney disease (CKD) manually when the estimated glomerular filtration rate (GFR) is lower than 60 ml/min/1.73 m². The objective of the study is to determine, in patients with a GFR < 60 ml/min/1.73 m², whether documenting a diagnosis of CKD in the record encompasses different attitudes with respect to the prevention and treatment of CKD.

The study was performed in the 16 primary care teams from the Muntanya Primary Care Service (SAP) in Barcelona, belonging to the Catalan Health Institute (ICS), where 212,321 adult patients were covered for health care in 2012. Patients with a minimum of 2 estimated glomerular filtration rates (eGFRs) using the CKD-EPI formula lower than 60 ml/min/1.73 m² in 2012 were enrolled. We identified two groups of patient depending of whether to diagnosis of CKD had been recorded in the medical history. A between-groups comparison was performed using the chi-squared test for qualitative variables and Student's t test for quantitative variables. A level of statistical significance of 5% ($p \leq 0.05$) was accepted.

In 2331 patients, at least 2 eGFR measurements < 60 ml/min/1.73 m² were recorded. The diagnosis of CKD

had been recorded in 1344 patients (57%), but not in 987 patients (43%). [Table 1](#) summarises the patients' characteristics. The patients with a recorded diagnosis of CKD had a lower eGFR and greater albuminuria. The proportion of diabetics was similar in the two groups. Body weight, LDL cholesterol and glycated haemoglobin in patients with diabetes were lower in the group of patients with a diagnosis of CKD. [Table 2](#) summarises the treatment in both groups. The use of renin-angiotensin system inhibitors was similar in the both groups. By contrast, the use of non-steroidal anti-inflammatory drugs was lower in patients with the diagnosis of CKD. In diabetic patients the use of metformin and sulphonylureas was also lower in the group with a diagnosis of CKD.

The main observation of the present study is that a diagnosis of CKD is made late and that the diagnosis is not shown in the medical history of approximately 40% of patients with an eGFR < 60 ml/min/1.73 m². Despite the greater severity of kidney disease, patients with a diagnosis of CKD recorded had less overweight and better control of LDL cholesterol and glycated haemoglobin (in diabetic patients). The use of non-steroidal anti-inflammatory drugs was lower in the group of patients with a diagnosis of CKD. Metformin and sulphonylureas were used less often in diabetic patients with a diagnosis of CKD. Taken together, these results suggest that a diagnosis of CKD is associated with a more proactive attitude aimed at decreasing risk factors for cardiovascular problems and for progression of kidney disease and also with greater prevention of acute renal failure. These findings raise the question of whether automatically including a diagnosis of CKD in the medical history of patients with an eGFR < 60 ml/min/1.73 m² might foster a more

[☆] Please cite this article as: Espinel E, Benavides F, Feijóo MV, Fernández-Liz E, Cossio Y, Serón D, et al. Registro del diagnóstico de insuficiencia renal crónica en la historia clínica electrónica y su influencia en el manejo terapéutico. Nefrología. 2016;36:315-317.

Table 1 – Sociodemographic data and mean baseline values.

Group	Without a diagnosis	With a diagnosis	p
N	987	1344	
Age (years) (SD)	78.09 (8.74)	79.27 (8.74)	<0.001
Women (n, %)	647 (65.6)	770 (57.3)	<0.001
SBP (mmHg) (SD)	132.58 (19.94)	133.62 (19.47)	NS
DBP (mmHg) (SD)	71.58 (12.16)	71.33 (10.94)	NS
Malb (mg/g) (SD)	24.68 (58.59)	41.02 (10.88)	<0.001
Creatinine (mg/dl) (SD)	1.21 (0.32)	1.62 (3.50)	<0.001
GFR, ml/min/1.73 m ² (SD)	48.90 (7.99)	41.02 (10.88)	<0.001
Haemoglobin (g/dl) (SD)	12.48 (1.73)	12.44 (3.23)	NS
Diabetes (n, %)	496 (50.3)	641 (47.7)	NS
HbA1c (%) (in DM) (SD)	7.6 (3.11)	7.3 (1.28)	0.035
BMI (kg/m ²)	30.16 (5.76)	29.46 (4.91)	0.024
LDL-C (mg/dl) (SD)	110 (34.88)	105 (34.64)	0.004
No. of visits ^a (2012) (SD)	18.07 (17.75)	18.72 (17.42)	NS
Referrals ^b (n, %)	26 (2.6)	123 (9.2)	<0.001

DM, diabetes mellitus; GFR, glomerular filtration rate; HbA1c, glycated haemoglobin; BMI, body mass index; LDL-C, LDL cholesterol; Malb, microalbuminuria; NS, $p > 0.05$; DBP, diastolic blood pressure; SBP, systolic blood pressure.

^a Any visit to the site for any reason has been counted.

^b Referrals to the Nephrology Department of the referral hospital.

Table 2 – Treatment differences according to group.

Group	Without a diagnosis (n, %)	With a diagnosis (n, %)	p
Total number of cases	987	1344	
ACE inhibitors	393 (39.9)	505 (37.6)	NS
AIIRAs	373 (37.8)	550 (40.9)	NS
ACE inhibitors + AIIRAs	5 (0.5)	4 (0.3)	NS
Aliskiren	4 (0.4)	3 (0.2)	NS
NSAIDs	312 (31.6)	345 (25.7)	0.002
Total diabetic patients	497	642	
Diab:metformin	186 (37.4)	167 (26)	<0.001
Diab:sulphonylureas	77 (15.5)	74 (11.5)	0.05

NSAIDs, non-steroidal anti-inflammatory drugs; AIIRAs, angiotensin II receptor antagonists; Diab, diabetic; ACE inhibitors, angiotensin converting enzyme inhibitors; NS, $p > 0.05$.

Bold values represent the number of patients.

proactive attitude in the treatment of these patients.^{9,10} However, these results should be interpreted with caution since this is a retrospective study and therefore does not allow causal relationships to be established. Conforming or refuting this hypothesis would require a prospective randomised study.

Conflicts of interest

The authors declare that they have no conflicts of interest.

Appendix. Grupo clínico de Nefrología del Servei d'Atenció Primària SAP Muntanya y Hospital Universitari Vall d'Hebrón. Institut Català de la Salut (ICS) Barcelona.

Carme Nebot. Servei d'atenció primària Muntanya Dreta.
 Pilar Algueró. Centre d'atenció primària Turó.
 Maribel Colas. Centre d'atenció primària Turó.
 Joan Atmetlla Andreu. Centre d'atenció primària Xafarinas.
 Dolors Llauradó. Centre d'atenció primària Turó.
 Araceli Martín. Centre d'atenció primària Ciutat Meridiana.

Andrea Sánchez Callejas. BASIQ. Unitat d'Avaluació, Sistemes d'Informació i Qualitat.

Francisca Benavides. Centre d'atenció primària Rio de Janeiro.

Eladio Fernández-Liz. Area del Medicament Àmbit Barcelona Ciutat.

Daniel Serón. Servei de Nefrologia. Hospital Universitari Vall d'Hebrón. Barcelona.

Eugenia Espinel. Servei de Nefrologia. Hospital Universitari Vall d'Hebrón. Barcelona.

REFERENCES

- Eckardt K, Coresh J, Devuyst O, Johnson RJ, Köttgen A, Levey AS, et al. Evolving importance of kidney disease: from subspecialty to global health burden. *Lancet*. 2013;382:158–69.
- Otero A, de Francisco AL, Gayoso P, García F, on behalf of the EPIRCE Study. Prevalence of chronic renal disease in Spain. Results of the EPIRCE study. *Nefrologia*. 2010;30:78–86.
- Martínez-Castelao A, Górriz JL, Bover J, Segura-de la Morena J, Cebollada J, Escalada J. Consensus document for the detection and management of chronic kidney disease. *Nefrologia*. 2014;34:243–62.

4. Gorostiti M, Santamaria R, Alcazar R, Fernández-Fresnedo G, Galcerán JM, Goicoechea M, et al. Spanish Society of Nephrology document on KDIGO guidelines for the assessment and treatment of chronic kidney disease. *Nefrología*. 2014;34:302-16.
5. Kidney disease: Improving Global Outcomes (KDIGO) blood pressure work group. KDIGO Clinical practice guideline for the management of blood pressure in chronic kidney disease. *Kidney Int Suppl*. 2012;2:337-414.
6. Kidney Disease Outcomes Quality initiative (K/DOQI). Clinical practice guidelines on hypertension and antihypertensive agents in chronic kidney disease. *Am J Kidney Dis*. 2004;43:S1-290.
7. García de Vinuesa S. Factores de progresión de la enfermedad renal crónica. Prevención secundaria. *Nefrología*. 2008;17-21.
8. Weir MR. Effects of renin-angiotensin system inhibition on end-organ protection: can we do better? *Clin Ther*. 2007;29:1803-24.
9. Chase HS, Radhakrishnan J, Shirazian S, Rao MK, Vawdrey DK. Under-documentation of chronic kidney disease in the electronic health record in outpatients. *J Am Med Inform Assoc*. 2010;17:558-94.
10. Navaneethan SD, Jolly SE, Sharp J, Jain A, Schold JD, Schreiber MJ, et al. Electronic health records: a new tool to combat chronic kidney disease. *Clin Nephrol*. 2013;79:175-83.

Eugenia Espinel^{a,f,*}, Francisca Benavides^b,
M. Victoria Feijóo^c, Eladio Fernández-Liz^d, Yolima Cossio^e,
Daniel Serón^{a,f}, on behalf of the Grupo Clínico de Nefrología
del Servicio de Atención Primaria SAP Muntanya y Hospital
Universitari Vall d'Hebrón de Barcelona[◇]

^a Servei de Nefrologia, Hospital Universitari Vall d'Hebrón,
Barcelona, Spain

^b Centre de Atenció Primària Rio de Janeiro, Spain

^c BASIQ Unitat d'Avaluació, Sistemes d'Informació i Qualitat,
Àmbit Barcelona Ciutat, Institut Català de la Salut, Barcelona,
Spain

^d Area del Medicament, Àmbit Barcelona Ciutat, Institut Català de
la Salut, Spain

^e Servei de Medicina Preventiva i Epidemiologia, Hospital
Universitari Vall d'Hebrón, Barcelona, Spain

^f Instituto Carlos III, Red de Investigación en Enfermedades
Renales, REDinREN, RD12/0021/0013, Spain

* Corresponding author.

E-mail addresses: eespinel@vhebron.net, 13047eeg@comb.cat
(E. Espinel).

◇ The names of participants in the study are available on
[Appendix](#).

2013-2514/© 2016 Sociedad Española de Nefrología. Published
by Elsevier España, S.L.U. This is an open access article under
the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).