UC Irvine

UC Irvine Previously Published Works

Title

Corrigendum: Averting the legacy of kidney disease - focus on childhood

Permalink

https://escholarship.org/uc/item/7kf68706

Journal

African Journal of Primary Health Care & Family Medicine, 9(1)

ISSN

2071-2928

Authors

Ingelfinger, Julie R Kalantar-Zadeh, Kamyar Schaefer, Franz

Publication Date

2017

DOI

10.4102/phcfm.v9i1.1706

Copyright Information

This work is made available under the terms of a Creative Commons Attribution License, available at https://creativecommons.org/licenses/by/4.0/

Peer reviewed





Corrigendum: Averting the legacy of kidney disease – focus on childhood



Authors:

Julie R. Ingelfinger¹ Kamyar Kalantar-Zadeh¹ Franz Schaefer¹

Affiliations:

¹World Kidney Day, International Society of Nephrology, in collaboration with the International Federation of Kidney Foundations, Brussels, Belgium

Corresponding author and email:

Julie Ingelfinger, myriam@worldkidneyday.org

Date:

Published: 13 Dec. 2017

How to cite this article:

Ingelfinger JR, Kalantar-Zadeh K, Schaefer F.
Corrigendum: Averting the legacy of kidney disease – focus on childhood. Afr J Prm Health Care Fam Med. 2017;9(1), a1706. https://doi.org/10.4102/phcfm. v9i1.1706

Copyright:

© 2017. The Authors. Licensee: AOSIS. This work is licensed under the Creative Commons Attribution License. In the version of this article initially published, the abbreviation HN located in the footnote for Table 2 was identified incorrectly as Hypertension. The correct definition for HN is Hereditary Nephropathy.

TABLE 2: Aetiology of chronic kidney disease in children.†

CKD aetiology	Percentage (range)	ESRD aetiology	Percentage (range)
CAKUT	48-59	CAKUT	34–43
GN	5–14	GN	15-29
HN	10–19	HN	12–22
HUS	2–6	HUS	2–6
Cystic	5–9	Cystic	6–12
Ischaemic	2–4	Ischaemic	2

Rare causes include congenital NS, metabolic diseases, cystinosis. Miscellaneous causes depend on how such entities are classified.

CAKUT, congenital anomalies of the kidney and urinary tract; GN, glomerulonephritis; HN, Hereditary Nephropathy; HUS, haemolytic uraemic syndrome.

†, from Harambat et al.² CKD data are from NAPRTCS, the Italian Registry and the Belgian Registry. ESRD data are from ANZDATA, ESPN/ERA-EDTA, UK Renal Registry and the Japanese Registry.

This correction does not alter the study's findings of significance or overall interpretation of the study results. The errors have been corrected in the PDF version of the article. The authors apologise for any inconvenience caused.

Read online:



Scan this QR code with your smart phone or mobile device to read online.

Note: DOI of original article: http://dx.doi.org/10.4102/phcfm.v8i1.1093

