



This is a repository copy of *1060: Renal safety of zoledronic acid in patients with borderline kidney function at a metabolic bone centre In the United Kingdom.*

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/192353/>

Version: Accepted Version

Proceedings Paper:

Schini, M. orcid.org/0000-0003-2204-2095, Peel, N., Toronjo-Urquiza, L. et al. (5 more authors) (2022) 1060: Renal safety of zoledronic acid in patients with borderline kidney function at a metabolic bone centre In the United Kingdom. In: Journal of Bone and Mineral Research. ASBMR Annual Meeting 2021, 01-04 Oct 2021, San Diego, California. Wiley , S271-S271.

<https://doi.org/10.1002/jbmr.4515>

This is the peer reviewed version of the following Abstract: [ASBMR Annual Meeting Abstract Supplement 2021. J Bone Miner Res, 37: S271, which has been published in final form at <https://doi.org/10.1002/jbmr.4515>. This article may be used for non-commercial purposes in accordance with Wiley Terms and Conditions for Use of Self-Archived Versions. This article may not be enhanced, enriched or otherwise transformed into a derivative work, without express permission from Wiley or by statutory rights under applicable legislation. Copyright notices must not be removed, obscured or modified. The article must be linked to Wiley's version of record on Wiley Online Library and any embedding, framing or otherwise making available the article or pages thereof by third parties from platforms, services and websites other than Wiley Online Library must be prohibited.

Reuse

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.



eprints@whiterose.ac.uk
<https://eprints.whiterose.ac.uk/>

Renal Safety Of Zoledronic Acid In Patients With Borderline Kidney Function At A Metabolic Bone Centre In The United Kingdom

Authors

Schini M1, Peel N², Toronjo-Urquiza L³, Thomas E4, Salam S5, Khwaja A5, Eastell R1, Walsh J1

Affiliations

1. Academic Unit of Bone Metabolism, The University of Sheffield
2. Metabolic Bone Centre, Sheffield Teaching Hospitals
3. Chemical Engineering Department, The University of Sheffield
4. Pharmacy, Sheffield Teaching Hospitals
5. Sheffield Kidney Institute, Sheffield Teaching Hospitals

Abstract

Background:

Zoledronic acid is a bisphosphonate widely used for the treatment of osteoporosis that leads to improvements in bone mineral density and reductions in fractures. A potential side effect is nephrotoxicity and acute kidney injury (AKI). Advice from the UK Medicines and Healthcare products Regulatory Agency (MHRA) in 2019 stated that creatinine clearance (CrCl) and not estimated glomerular rate (eGFR) should be used to guide and make decisions about treatment and that patients should not receive zoledronic acid if their CrCl is below 35 ml/min. The objective of this study was to review the safety of our previous practice using eGFR and the clinical impact of implementing the MHRA recommendations.

Methods:

The study was performed at the Metabolic Bone Centre (MBC) in Sheffield Teaching Hospitals, UK. Data on all the patients who had zoledronic acid infusions from the 1/09/2015 to the 1/10/2020 at the centre were retrieved and evaluated.

Results:

Data on 4405 patients were retrieved. Serum creatinine in the 14 days post- infusion was available for a total of 969 infusions and amongst them, 160 (16%) infusions were given with baseline CrCl < 35 ml/min. AKI was observed within 14 days following 45

infusions (4.5%). Only 9 infusions resulted in AKI with a pre-treatment of CrCl < 35 ml/min.

If the MHRA rules had been followed (calculating CrCl for patients aged ≥ 75 years and/or extreme BMI <18 or >40 kg/m²), 996 infusions with baseline CrCl <35 ml/min would not have been given.

Logistic regression showed that both CrCl and eGFR were significant factors in predicting AKI within 14 days, but that the current recommended cut-off of CrCl 35 ml/min had a poor sensitivity. The areas under the curve for each marker were 0.608 and 0.627 respectively, suggesting that neither are sensitive in predicting AKI.

Conclusions:

This study suggested that zoledronic acid could be contributing to the development of AKI in some patients. Estimated GFR is better validated than CrCl and so is preferable to use. Since low eGFR is at least as good a predictor of AKI as CrCl, it should be used in every day clinical practice.

Conflicts of Interest

MS receives consultancy from Kyowa Kirin International and grant funding from Roche Diagnostics

SS received grant funding from IDS.