

University of Groningen

Calprotectin instability may lead to undertreatment in children with IBD

Haisma, Sjoukje-Marije; van Rheenen, Patrick Ferry; Wagenmakers, Lucie; Muller Kobold, Anneke

Published in:
Archives of Disease in Childhood

DOI:
[10.1136/archdischild-2018-316584](https://doi.org/10.1136/archdischild-2018-316584)

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
2020

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):

Haisma, S-M., van Rheenen, P. F., Wagenmakers, L., & Muller Kobold, A. (2020). Calprotectin instability may lead to undertreatment in children with IBD. *Archives of Disease in Childhood*, 105(10), 996-998. <https://doi.org/10.1136/archdischild-2018-316584>

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.



OPEN ACCESS

Calprotectin instability may lead to undertreatment in children with IBD

Sjoukje-Marije Haisma,¹ Patrick Ferry van Rheenen,¹ Lucie Wagenmakers,² Anneke Muller Kobold²

¹Department of Pediatric Gastroenterology, Hepatology and Nutrition, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

²Department of Laboratory Medicine, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

Correspondence to

Dr Patrick Ferry van Rheenen, Department of Pediatric Gastroenterology, Hepatology and Nutrition, University of Groningen, University Medical Center Groningen, Groningen 9700 RB, The Netherlands; p.f.van.rheenen@umcg.nl

Received 21 November 2018

Revised 10 December 2018

Accepted 19 December 2018

ABSTRACT

Background Treatment decisions in children with inflammatory bowel disease (IBD) are increasingly based on longitudinal tracking of faecal calprotectin concentrations, but there is little known about the stability of this protein in stool.

Methods We stored aliquots of homogenised stool at room temperature and at 4°C, and measured the calprotectin concentration for 6 consecutive days with three different assays. In addition, we assessed calprotectin stability in assay-specific extraction buffers kept at room temperature.

Results After 6 days of storage at room temperature, mean percentage change from baseline calprotectin concentrations in stool and extraction buffer was 35% and 46%, respectively. The stability of calprotectin was significantly better preserved in samples stored at 4°C ($p=0.0066$ and 0.0011 , respectively).

Conclusions Calprotectin is not stable at room temperature. Children with IBD and their caretakers may be falsely reassured by low calprotectin values. The best advisable standard for preanalytical calprotectin handling is refrigeration of the stool sample until delivery at the hospital laboratory.

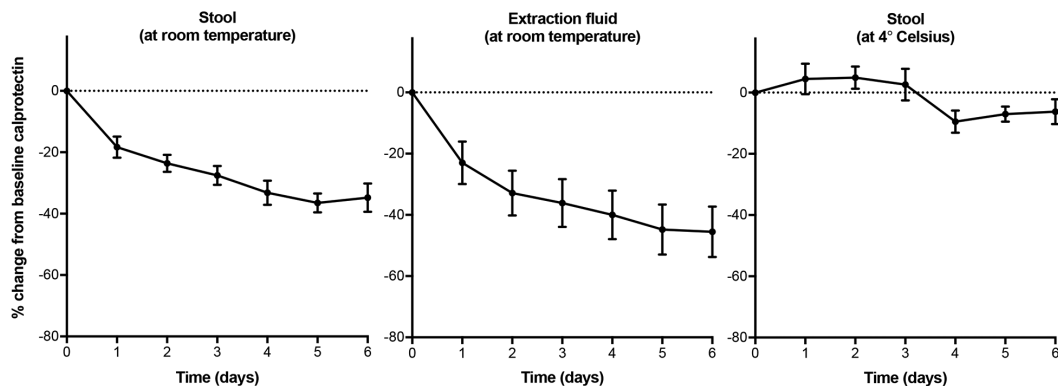
the use of faecal calprotectin as an objective and non-invasive measure of disease activity has gained a lot of interest in both paediatric practice and trial settings.^{1 2} Calprotectin is widely believed to be a stable marker, hence many physicians and researchers allow patients to send a stool sample to the laboratory by surface mail at ambient temperatures. However, evidence for stability at room temperature is scarce and all quotes can be traced back to one of the earliest scientific publications on stool calprotectin.³ We assessed calprotectin stability in stool samples stored at room temperature and at 4°C. In addition, we assessed calprotectin stability in extraction buffers.

METHODS

We collected 19 stool samples from patients with IBD and ensured that the whole range of potential calprotectin values was covered. Within 2 hours of collection spot samples were homogenised and split into 12 aliquots, six of which were kept at room temperature and six at 4°C for 1, 2, 3, 4, 5 and 6 days, respectively. Each stool sample was also extracted and split into six aliquots of extraction fluid and likewise kept at room temperature. We used ELISAs of BÜHLMANN Laboratories (fCAL), Preventis (IDK Calprotectin) and Calpro (CalproLab ALP), and pooled the calprotectin results per preanalytical condition. To minimise any batch-to-batch variation, all aliquots were analysed in the same analytical batch. A repeated measures analysis of variance was performed and results were plotted over time and expressed as percentage change from baseline calprotectin with 95% CI.

INTRODUCTION

Inflammatory bowel diseases (IBD), including Crohn's disease and ulcerative colitis, are lifelong conditions that often begin in childhood. Disease monitoring is usually based on clinical symptoms with an emphasis on altered bowel habits and rectal bleeding. As relying on subjective symptoms alone could lead to management errors,



Error bars represent 95% confidence intervals.

Figure 1 Calprotectin stability over time.



© Author(s) (or their employer(s)) 2019. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

To cite: Haisma S-M, van Rheenen PF, Wagenmakers L, et al. *Arch Dis Child Epub ahead of print: [please include Day Month Year].* doi:10.1136/archdischild-2018-316584

RESULTS

Baseline calprotectin concentration in the stool samples varied from 14 to 3270 $\mu\text{g/g}$. In aliquots kept at room temperature the mean change from baseline values on days 1, 3 and 6 was 18% (95% CI 11 to 25), 28% (95% CI 22 to 34) and 35% (95% CI 26 to 44), respectively. This decline was irrespective of the calprotectin concentration at baseline. In extraction fluids kept at room temperature the mean change from baseline values was 23% (95% CI 9 to 37), 36% (95% CI 21 to 51) and 46% (95% CI 30 to 62). In aliquots kept at 4°C calprotectin values remained stable during the observation period (see figure 1). Calprotectin stability over time was significantly diminished in stool and extraction fluids kept at room temperature compared with storage at 4°C ($p=0.0066$ and 0.0011 , respectively).

DISCUSSION

Our results indicate a steady decline in calprotectin concentration in the first 6 days after stool collection in samples kept at room temperature. A similar slope was seen in stool extracts kept at room temperature, while the stability of calprotectin was preserved in samples stored in a refrigerator at 4°C. We did not test the superiority of one ELISA kit over another. For that, we would have needed to perform a multitude of tests. The decline in calprotectin concentration over time was observed regardless of the ELISA kit used, which makes inadvertently smoothing of the curve unlikely.

Literature on calprotectin stability under various preservation conditions is scarce. A Swedish group observed a decline in calprotectin when stool samples were stored at room temperature for 7 days but claimed that the concentration remained unchanged in the first 3 days after collection.⁴ A Spanish group claimed that calprotectin remained stable in stool extracts stored at room temperature for 4 days,⁵ as opposed to our finding that extraction does not prevent a decline.

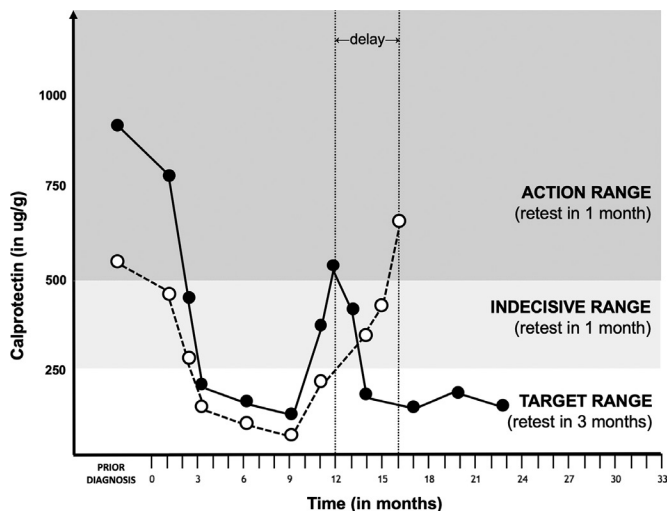


Figure 2 Stool calprotectin values of an imaginary patient with Crohn's disease plotted over time. Black dots represent true calprotectin values, white dots represent underestimated calprotectin values caused by protein degradation in the preanalytical phase. In month 12, the patient's true calprotectin measurement would have resulted in treatment intensification, while calprotectin degradation during transport may cause false reassurance and significant treatment delay.

Calprotectin instability in stool samples sent by surface mail at ambient temperatures may lead to management errors in treat-to-target strategies, in which physicians aim for calprotectin concentrations below 250 $\mu\text{g/g}$.⁶ Although there is no consensus on the optimal frequency of calprotectin retesting and cut-offs for treatment intensification, the authors of this article use the following monitoring strategy: patients with IBD with calprotectin values below 250 $\mu\text{g/g}$ are reassured and advised to retest in 3 months; patients with IBD with calprotectin values above 500 $\mu\text{g/g}$ are considered to have active disease, require treatment intensification and retesting in 1 month; and patients with IBD with calprotectin in the intermediate range are advised to retest in 1 month before progressing to a decision. In such a strategy, patients whose calprotectin values show a drift from the target range into the indecisive range may be falsely reassured, leading to a delay in treatment intensification (figure 2). The intermediate range is exactly where precision is of utmost importance and small deviations lead to misclassification of disease activity and wrong treatment decisions. Test results at the high and extreme low end of the calprotectin spectrum will not lead to management errors.

Currently, the best advisable standard for preanalytical calprotectin handling is refrigeration of the filled stool container until delivery at the hospital laboratory. Several manufacturers recently introduced a lateral flow-based test and a software application that turns an ordinary smartphone camera into a reader for quantitative calprotectin measurements.⁷ Although this innovation has great opportunities to take transport of stool samples out of the equation, home tests are not yet commonplace in monitoring disease activity in IBD. In the meantime, for those who continue shipping unrefrigerated stool samples to the laboratory, we emphasise the importance of documenting the time to arrival at the laboratory. Unrefrigerated stool samples that arrive with a delay exceeding 24 hours with a calprotectin concentration between 250 and 500 $\mu\text{g/g}$ need confirmation. We currently advise our patients to store the confirmation sample in the freezing compartment of the fridge until the face-to-face encounter with the physician and then deliver the confirmation sample to the laboratory by hand.

Contributors All authors contributed to the study concept and design. LW collected the data and analysed the data together with SMH, AMK and PFvR. SMH and PFvR drafted the first version of the research letter.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests PFvR received funding from BÜHLMANN Laboratories for other projects.

Patient consent Not required.

Provenance and peer review Not commissioned; externally peer reviewed.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

REFERENCES

- 1 Peyrin-Biroulet L, Sandborn W, Sands BE, *et al*. Selecting therapeutic targets in inflammatory bowel disease (stride): determining therapeutic goals for treat-to-target. *Am J Gastroenterol* 2015;110:1324–38.
- 2 Sandborn WJ, Panés J, Zhang H, *et al*. Correlation between concentrations of fecal calprotectin and outcomes of patients with ulcerative colitis in a phase 2 trial. *Gastroenterology* 2016;150:96–102.
- 3 Røseth AG, Fagerhol MK, Aadland E, *et al*. Assessment of the neutrophil dominating protein calprotectin in feces. A methodologic study. *Scand J Gastroenterol* 1992;27:793–8.

- 4 Lasson A, Stotzer PO, Öhman L, *et al.* The intra-individual variability of faecal calprotectin: a prospective study in patients with active ulcerative colitis. *J Crohns Colitis* 2015;9:26–32.
- 5 Acevedo D, Salvador MP, Girbes J, *et al.* Fecal calprotectin: a comparison of two commercial enzymeimmunoassays and study of fecal extract stability at room temperature. *J Clin Med Res* 2018;10:396–404.
- 6 Colombel JF, Panaccione R, Bossuyt P, *et al.* Effect of tight control management on Crohn's disease (CALM): a multicentre, randomised, controlled phase 3 trial. *Lancet* 2018;390:2779–89.
- 7 Heida A, Knol M, Kobold AM, *et al.* Agreement between home-based measurement of stool calprotectin and ELISA results for monitoring inflammatory bowel disease activity. *Clin Gastroenterol Hepatol* 2017;15:1742–9.