## Should wheat, barley, rye, and/or gluten be avoided in a 6-food elimination diet?



Eosinophilic esophagitis (EoE), a food antigen-mediated disease, is effectively treated with the dietary elimination of 6 foods commonly associated with food allergies (milk, wheat, egg, soy, tree nuts/peanuts, and fish/shellfish). Because wheat shares homologous proteins (including gluten) with barley and rye and can also be processed with these grains, some clinicians have suggested that barley and rye might also trigger EoE as a result of cross-reaction and/or cross-contamination with wheat. In this article, we discuss the theoretical risks of cross-reactivity and cross-contamination among wheat, barley, and rye proteins (including gluten); assess common practices at EoE treatment centers; and provide recommendations for dietary treatment and future studies of EoE. (J Allergy Clin Immunol 2016;137:1011-4.)

CrossMark

*Key words: Eosinophilic esophagitis, 6-food elimination diet, wheat, cross-reactivity, gluten* 

Eosinophilic esophagitis (EoE) is an immune-mediated clinicopathologic disease of the esophagus that manifests as vomiting,

Novartis, Receptos, Regeneron, and Roche and has received research funding from Miraca, Meritage, Receptos, and Regeneron. G. W. Falk has received a grant and travel support from CEGIR, has consultant arrangements with Aptalis, and has received grants from Meritage, Receptos, and Regeneron. G. T. Furuta is a medical advisor to Campaign Urging Research for Eosinophilic Disease (CURED), has consultant arrangements with Genentech and UpToDate, has received grants from the NIH, has a patent for Esophageal String Test, and is cofounder of EnteroTrack. N. Gonsalves has received lecture fees from Nutricia and receives royalties from UpToDate. S. K. Gupta has consultant arrangements with Receptos and Abbott Nutrition. I. Hirano has consultant arrangements with Receptos, Regeneron, and Meritage; has received grants from the NIH and the American Society of Gastrointestinal Endoscopy; and has received royalties from UpToDate. A. Kagalwalla has received payment for lectures from Nutricia. V. A. Mukkada has received grants from the Rare Disease Clinical Research Network (U54 AI117804) and the Patient Centered Outcomes Research Institute. J. M. Spergel has received grants from the NIH, DBV Technology, Food Allergy Research and Education, Aimmune Therapeutics, and the Stanford Food Allergy Research Center; is on the Scientific Advisory Board for DBV Technology; has consultant arrangements with Danone; has received payment for lectures from MEI; and has stock/stock options in DBV Technology. M. E. Rothenberg has received a grant from the NIH; has received money paid to his institution from the CURED Foundation, the Buckeye Foundation, Food Allergy Research and Education, and the APFED Foundation; is a board member for the International Eosinophil Society; is on the medical advisory panel for the American Partnership for Eosinophilic Disorders; has consultant arrangements with Immune Pharmaceuticals, Receptos, Celsus Therapeutics, Genentech/Roche, and Novartis; has patents submitted and owned by CCHMC for which he is an inventor; has received royalties from Teva Pharmaceuticals; and has stock/stock options in Immune Pharmaceuticals, Receptos, Celsus Therapeutics, and NKT Therapeutics, J. Leung declares that he has no relevant conflicts of interest.

Received for publication July 31, 2015; revised September 18, 2015; accepted for publication October 5, 2015.

Available online December 24, 2015.

- Corresponding author: Marc E. Rothenberg, MD, PhD, Division of Allergy and Immunology, Cincinnati Children's Hospital Medical Center, MLC 7028, 3333 Burnet Ave, Cincinnati, OH 45229. E-mail: rothenberg@cchmc.org.
- The CrossMark symbol notifies online readers when updates have been made to the article such as errata or minor corrections

0091-6749/\$36.00

© 2015 American Academy of Allergy, Asthma & Immunology http://dx.doi.org/10.1016/j.jaci.2015.10.040

From a the Division of Allergy and Immunology, Department of Pediatrics, and <sup>j</sup>the Division of Gastroenterology, Hepatology and Nutrition, Department of Pediatrics, Cincinnati Children's Hospital Medical Center; <sup>b</sup>the Division of Allergy and Immunology, Department of Pediatrics and Medicine, University of California, San Diego and Rady Children's Hospital, San Diego; <sup>c</sup>the Division of Gastroenterology, Tufts Medical Center, Boston; <sup>d</sup>the Division of Gastroenterology and Hepatology, University of North Carolina School of Medicine, Chapel Hill; ethe Division of Gastroenterology, Hospital of the University of Pennsylvania, University of Pennsylvania Perelman School of Medicine, Philadelphia; <sup>f</sup>the Section of Gastroenterology, Hepatology, and Nutrition, Department of Pediatrics, University of Colorado School of Medicine, Aurora; <sup>g</sup>the Division of Gastroenterology and Hepatology, Northwestern University Feinberg School of Medicine, Chicago; hthe Section of Pediatric Gastroenterology, Hepatology and Nutrition, Indiana University School of Medicine, Riley Hospital for Children, Indianapolis; <sup>i</sup>the Division of Gastroenterology, Hepatology and Nutrition, Ann and Robert H. Lurie Children's Hospital of Chicago; kthe Division of Allergy and Immunology, Children's Hospital of Philadelphia, University of Pennsylvania Perelman School of Medicine, Philadelphia.

<sup>\*</sup>Investigator/collaborator members of the Consortium of Eosinophilic Gastrointestinal Disease Researchers (CEGIR).

Supported in part by U54 AI117804, which is part of the Rare Disease Clinical Research Network (RDCRN), an initiative of the Office of Rare Disease Research (ORDR), National Center for Advancing Translational Sciences (NCATS), and funded through collaboration between the NCATS, National Institute of Allergy and Infectious Diseases, and National Institute of Diabetes and Digestive and Kidney Diseases, which have collectively resulted in the Consortium of Eosinophilic Gastrointestinal Disease Researchers (CEGIR).

Disclosure of potential conflict of interest: K. L. Kliewer has received a grant and travel support from the Consortium of Eosinophilic Gastrointestinal Disease Researchers (CEGIR; National Institutes of Health [NIH]). C. Venter has consultant arrangements with Danone, has received payment for lectures from Nestlé, and has received payment for development of education presentations from Mead Johnson. A. M. Cassin has consultant arrangements with and has received payment for lectures and development of educational presentations from Nutricia North America. J. P. Abonia has received a grant from the NIH (U54 Grant [CEGIR], PCORI grant). S. S. Aceves has received a grant from the NIH (U54 Consortium grant CEGIR, R01) and Raptor Pharmaceutical, is on the American Partnership for Eosinophilic Disorders Medical Advisory Board, has stock/stock options in Meritage Pharma, has received travel support from the NIH/ National Institute of Diabetes and Digestive and Kidney Diseases. P. A. Bonis is Chief Medical Officer at UpToDate. E. S. Dellon has consultant arrangements with Aptalis,

Abbreviations used CEGIR: Consortium of Eosinophilic Gastrointestinal Disease Researchers EoE: Eosinophilic esophagitis

6FED: Six-food elimination diet

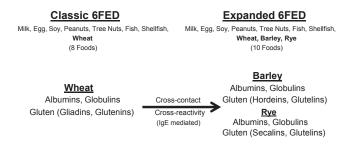
feeding difficulties, and food impaction, which vary as a function of a patient's age. Histologically, EoE is marked by esophageal eosinophilia that is unresponsive to proton pump inhibitor therapy.<sup>1</sup> A series of studies suggest allergic sensitization to food or aeroallergens underlies EoE.<sup>2</sup> Food elimination diets have been shown to be effective in achieving both clinical and histologic remission in patients with EoE,<sup>3-10</sup> providing evidence that EoE is, at least in part, food antigen mediated.<sup>11</sup>

In a retrospective study of children with EoE, Kagalwalla et al<sup>4</sup> found that the empiric elimination of 6 foods commonly associated with food allergies (cow's milk, wheat, soy, egg, nuts, and fish) significantly reduced esophageal eosinophilia in 74% of the patients. Dietary elimination of the same foods in subsequent prospective and retrospective studies also resulted in clinical and histologic remission in both adult<sup>5,9</sup> and pediatric<sup>6,7</sup> patients with EoE.

Kagalwalla's "classic" 6-food elimination diet (6FED)<sup>4</sup> is understood to technically eliminate 8 foods/food families: milk, wheat, soy, egg, tree nuts, peanuts, fish, and shellfish. Of the foods in the classic 6FED, wheat was identified as the most common trigger of EoE in adults<sup>5</sup> and the second most common trigger in children<sup>12</sup> in 2 US studies using food reintroduction to identify food antigens associated with EoE. Of foods in a "6FED-like" diet, wheat was also the second most common antigen associated with EoE in adults in a Spanish cohort.<sup>8</sup> Overall, wheat reintroduction reactivated EoE in 26% to 60% of patients in remission from dietary therapy.<sup>5,8,12</sup> Thus eliminating dietary wheat is necessary for remission in a significant number of patients with EoE. However, the extent to which wheat (and perhaps wheat-related grains) should be avoided for clinical and histologic remissions in patients with EoE remain unclear.

Wheat is a cereal grain composed of 4 fractions of proteins (ie, albumins, globulins, and "gluten" [gliadins and glutenins]),<sup>13</sup> any of which might elicit an IgE-mediated allergic response.<sup>14</sup> Wheat can be grown, harvested, stored, and/or processed with other grains, thereby contaminating these grains with wheat protein fractions.<sup>15,16</sup> In most countries, food allergen labeling regulations do not mandate that food manufacturers disclose cross-contamination risks on food labels.<sup>17</sup> Thus, patients advised to eliminate wheat on the classic 6FED might unintentionally consume trace contaminants of wheat when consuming other grains, especially grains at high risk of cross-contact with wheat, like barley, rye, and oats.<sup>16</sup>

In the absence of studies quantifying the clinical relevance of trace ingestions of wheat in patients with EoE, some clinicians have advocated a risk-averse approach. Prompted by concerns of wheat cross-contamination of barley, rye, and oats,<sup>16</sup> Doerfler et al<sup>18</sup> recently suggested that elimination diets for EoE be expanded from wheat free to exclude wheat, barley, rye, and conventional oats in practice to mitigate "unforeseen" risks of wheat contaminants to patients. Because wheat, barley, rye, and their crossbreeds are the only foods that inherently contain gluten,



**FIG 1.** Classic 6FED for dietary management of EoE modified to exclude all gluten-containing grains. Uncertainty about the risks posed by cross-contamination and cross-reaction of barley and rye with wheat have led some to expand the classic wheat-free 6FED to exclude wheat, barley, and rye.

this recommendation effectively suggests eliminating all gluten-containing grains in the 6FED.

In addition to concerns of wheat cross-contamination, concerns of possible cross-reactivity among related grains (barley, rye, and wheat) have also recently led other clinicians to exclude all gluten-containing foods in empiric elimination diets.<sup>10</sup> Barley and rye share homologous proteins with wheat, including the "gluten" proteins hordein (barley) and secalin (rye).<sup>19</sup> Several studies indicate wheat, barley, and rye also share cross-reacting proteins,<sup>20-22</sup> which might be of relevance in IgE-mediated disease. However, in an early study of cross-reactivity of cereal antigens, only 4 of 25 patients with wheat allergy clinically reacted to barley or rye.<sup>21</sup> In contrast, Pourpak et al<sup>23</sup> found 55% of pediatric patients with IgE-mediated hypersensitivity to ingested wheat clinically reacted to barley. A strong correlation between wheat and barley serum-specific IgEs was also observed, suggesting antigen cross-reactions.<sup>23</sup> Studies of cross-reactivity of food antigens in patients with EoE are lacking. However, the frequency of sensitization to cereal allergens with identifiable cross-reacting aeroallergens was found to be high (63%) in a study of adults with EoE,<sup>24</sup> suggesting the potential for cross-reactivity among ingested grains.

To date, there are no studies to indicate whether clinical or histologic outcomes in patients with EoE would improve if the classic wheat-free 6FED was broadened to exclude all glutencontaining grains (Fig 1). To assess active ongoing practices, we queried a set of leading US clinical centers treating EoE selected by their participation in the Consortium of Eosinophilic Gastrointestinal Disease Researchers (CEGIR), which is part of the National Institutes of Health-sponsored Rare Disease Clinical Research Consortium (http://rdcrn.org/cegir). CEGIR investigators, as well as a subset of other US-based EoE clinical practices, primarily excluded only wheat in the 6FED (Fig 2). However, a similar polling of EoE-treating international sites revealed that exclusion of all gluten-containing grains occurred more often (Fig 2). Concern over cross-reactivity of barley and rye with wheat was the most often cited rationale for eliminating all gluten-containing grains in the 6FED. It is interesting to speculate that in addition to a heightened concern about grain cross-reactivity, the reason for the difference in practice between the United States and other countries could also simply be a practical matter. In most countries outside the United States, food allergen labeling laws mandate disclosure of all gluten-containing grain ingredients (wheat, barley, and rye) on food labels. In the United States only wheat must be identified by name.<sup>17</sup> Thus in the United States, eliminating barley and

Download English Version:

## https://daneshyari.com/en/article/6062583

Download Persian Version:

https://daneshyari.com/article/6062583

Daneshyari.com