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The prevalence of eosinophilic esophagitis in pediatric patients with IgE-mediated food allergy

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Abstract

BACKGROUND—Eosinophilic esophagitis (EoE) is an allergic inflammatory disease that is triggered by food allergens and characterized by progressive esophageal dysfunction. Recently, EoE has been identified in patients undergoing oral immunotherapy (OIT) for IgE-mediated food allergy, suggesting an association.

OBJECTIVE—We sought to ascertain whether significant associations exist between IgE-mediated food allergies and EoE.

METHODS—Utilizing analysis of EMR data and manual chart review, we examined our subspecialty care network of 35,528 children and adolescents to identify and characterize patients with IgE-mediated, and EoE food allergy. The most common food allergens were defined, and the prevalence of EoE in patients with IgE-mediated food allergy was determined. Logistic regression was used to measure the extent to which IgE-mediated food allergy to specific foods is associated with EoE.

RESULTS—The most common causes of EoE were milk, grains, meats, peanut, tree nuts, egg, and soy, an allergen pattern that is distinct from that of IgE-mediated food allergy. The prevalence of EoE in patients with IgE-mediated food allergy was higher than that reported in the general population (4.7% vs 0.04%). The distribution of IgE-mediated food allergens in patients with EoE was similar to that of the general population, and IgE-mediated allergy to egg (2.27; 1.91–2.64),

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milk (4.19; 3.52–4.97), or shellfish (1.55; 1.24–1.92) was significantly associated with EoE diagnosis.

CONCLUSIONS—Our findings support a clinical association between these conditions that has implications for the management of children with food allergy, and particular relevance to patients undergoing OIT.

Keywords

IgE; Food Allergy; Eosinophilic Esophagitis; Oral Immunotherapy; Prevalence

Introduction

IgE-mediated food allergies are common in the United States, and their prevalence and severity may be increasing.^(1–5) Childhood food allergy is associated with impaired quality of life, limited social interactions, comorbid allergic conditions, and significant economic cost to our healthcare system. Importantly, a severe allergic reaction resulting in anaphylaxis can be life threatening, and food allergens are the most common cause of anaphylaxis and anaphylaxis-related mortality in children and adolescents.⁽⁶⁾ Prior studies have identified an association between IgE-mediated food allergy and the development of comorbid atopic conditions including asthma and allergic rhinitis.^(7, 8) However, the extent to which IgE-mediated food allergy is associated with eosinophilic esophagitis (EoE) is unknown.

Eosinophilic esophagitis is a chronic allergic inflammatory disease of the esophagus that, if left untreated, can result in significant impairment in quality of life. EoE should be considered in older children and adults who present with symptoms of esophageal dysfunction and/or fibrosis such as dysphagia, odynophagia, or a history of food impaction and/or esophageal stricture formation. Diagnosis in younger children can be difficult as symptoms often mimic those of gastro-esophageal reflux disease, however, EoE-related symptoms do not respond to gastric acid suppression alone. On histology, esophageal tissue of patients with EoE is characterized by esophageal epithelial barrier defects and eosinophil infiltrates. Diagnosis requires the presence of 15 or more eosinophils per high power field in one or more of at least four esophageal biopsy specimens, while the patient is on optimal gastric acid suppression.^(9–11) When an EoE diagnosis is made, it is important that measures be taken to eliminate or control esophageal inflammation, through allergen identification and avoidance or topical corticosteroid therapy, to relieve symptoms and prevent fibrotic complications of esophageal stricture formation and food impaction.⁽¹²⁾

The pathophysiology of EoE is complex and an active area of research.⁽¹³⁾ However, there are several lines of experimental and clinical evidence from both animal and human studies to suggest that while IgE-mediated food allergy and EoE both lie on the atopic spectrum, they have distinct pathophysiology.^(14–16) We have shown that eosinophilic inflammation is independent of both B cells and IgE in animal models.^(17, 18) These findings are consistent with the clinical observations that 1) IgE-mediated food allergy and EoE have distinct natural histories (patients with IgE-mediated allergy frequently outgrow their disease, while those with EoE to the same allergen typically do not),^(19–21) 2) testing for allergen-specific IgE molecules is not helpful when attempting to identify EoE-causative foods,⁽²²⁾ 3) patients

can develop EoE after resolution of IgE-mediated food allergy,⁽²³⁾ and 4) omalizumab (anti-IgE) therapy is not beneficial in EoE.⁽²⁴⁾ Despite these observations, the extent to which IgE-mediated food allergy is associated with the development of EoE is not known.

We perform a retrospective cohort study of electronic medical record (EMR) data from one of the largest pediatric EoE cohorts available. We define and compare children with IgE-mediated or EoE food allergy in terms of demographics, clinical characteristics, and most common food allergens. We determine the prevalence of EoE in patients with IgE-mediated food allergy, and perform multivariate logistic regression analysis to measure the extent to which IgE-mediated food allergy is associated with EoE diagnosis. Our results identify an important association between these conditions, have implications for the management of children with food allergy, and are of particular relevance to patients undergoing oral immunotherapy (OIT).

Methods

Cohort generation and data extraction

The Allergy Section at The Children's Hospital of Philadelphia (CHOP), which includes six care locations, is the largest provider of sub-specialty allergy care services to patients residing in the eastern Pennsylvania, New Jersey, and Delaware region. In addition, we are an international referral center for eosinophilic esophagitis and have developed one of the largest pediatric cohorts for this condition through our Center for Pediatric Eosinophilic Diseases (CPED). We extracted data from the electronic medical record (EMR; Epic Systems, Verona, WI) of 35,528 children who obtained specialty care in our division between 1/1/2010 and 12/31/2015 (Fig 1). Direct patient identifiers (e.g., medical record number, name, etc.) were removed to create a dataset with limited identifiers (e.g., date of birth, dates of healthcare encounters, diagnoses, etc.) and data analyses tasks were completed using SAS v9.4 (Cary, NC). Three EMR study cohorts were defined: a cohort of 11,513 children with IgE-mediated food allergy in the absence of EoE (Fig 1, a), 570 children with concurrent IgE-mediated food allergy and EoE (Fig 1, b), and 266 children with EoE in the absence of IgE-mediated food allergy (Fig 1, c). We also utilized a manual cohort with detailed information on IgE and EoE allergens for 1795 patients with EoE who were evaluated in the CPED between 1/1/2006 and 5/1/2016. The CHOP Institutional Review Board (IRB) reviewed the retrospective cohort portion of our study and determined that it was exempt from requiring ethics approval or subject consent as it did not meet the definition of "human subject" research. The CPED cohort was approved by the CHOP IRB and performed under protocol number 07-005420.

Definitions of conditions studied

Diagnosis of IgE-mediated food allergy and EoE was made in accordance with established practice parameters.^(10, 25) ICD-9 and ICD-10 codes were used to identify children with IgE-mediated food allergy (ICD-9 codes 477.1, 693.1, 995.60-995.69 and V15.05; ICD-10 codes J30.5, L27.2, T78.0%, T78.1%, and Z91.01%, where % includes all variations that follow the preceding characters), and EoE (ICD-9 code 530.13; ICD-10 code K20.0). The allergy module of our EMR was used to identify specific food allergens. The presence of both a

respective ICD diagnosis code and a food allergen in the allergy module were required for inclusion in the EMR study cohorts. Requirement for inclusion in the CPED EoE cohort was diagnosis of EoE and identification of trigger food by biopsy while on an appropriate PPI regimen, in accordance with consensus guidelines.⁽¹⁰⁾ For prevalence calculations, individual children with each diagnosis were enumerated. For determination of the most common food allergens, individual instances of food allergy were enumerated (even if multiple allergies occurred in the same patient).

Validation of ICD codes and food allergy data via chart review

To determine the extent to which ICD diagnosis codes and food allergy data correctly represented the presence of each of the conditions of interest, a manual review of 100 charts was performed. Additionally, to ensure accurate estimation of the association between IgE-mediated food allergy and EoE, we performed a manual chart review of 100 patients with IgE-mediated food allergy and EoE. Evidence of IgE-mediated allergy was defined as presence of consistent history and laboratory analysis, skin testing, and/or food challenge. The chart review portion of our study was approved by the CHOP IRB and performed under protocol number 07–005420.

Statistics

We used multivariate logistic regression analysis of our EMR cohort to assess the relationship between demographic variables or IgE-mediated food allergy and EoE status. Analysis of IgE-mediated food allergy was adjusted for gender, race, and ethnicity. The top five IgE-mediated food allergens for patients with EoE (peanut, egg, tree nut, milk, shellfish) were included in our analysis. Odds ratios, 95% confidence intervals, and p values were calculated and reported.

Results

Cohort demographics and ICD code validation

A graphical representation of our EMR study cohorts is shown (Fig 1), as are the demographic characteristics of patients with IgE-mediated, and EoE food allergy (Table 1). Patients with IgE-mediated food allergy were found to be approximately 61% male and 58% white, while patients with EoE were found to be approximately 76% male and 74% white. These findings are consistent with prior studies that have found that EoE is most prevalent in Caucasian males.⁽²⁶⁾ Both cohorts had a similar ethnic distribution, while patients with EoE were slightly older (21% born before the year 2000 compared to 9% in the IgE cohort) consistent with the natural history of these conditions.⁽¹⁹⁾ The demographic characteristics of the manual CPED cohort were similar to those of the EMR EoE cohort (data not shown). To determine the accuracy of our coded EHR data, a manual chart review was performed for 100 patients. The percent confirmed diagnosis for all conditions studied (true positive rate for ICD codes) was 92% overall (92% for EoE, and 92% for IgE-mediated food allergy).

Analysis of food allergens in patients with IgE-mediated food allergy or EoE

To determine the most common food allergens in patients with IgE-mediated food allergy (in the absence of EoE), we utilized our EMR cohort. Consistent with previous reports, the most

common causal foods of IgE-mediated allergy were peanut, tree nut, egg, milk, and shellfish (Fig 2 a, b). To determine the most common causal foods of EoE, we examined our CPED cohort of 1795 patients with EoE. The most common causal foods of EoE were milk, soy, egg, grains, and meats (Fig 2 c, d). To determine the most common causes of IgE-mediated food allergy in patients with concurrent EoE, we performed a manual evaluation of our CPED cohort. Of patients with EoE, the most common causes of IgE-mediated food allergy were peanut, egg, tree nut, milk, and shellfish (Fig 2e, f).

Prevalence of EoE in patients with IgE-mediated food allergy

We next sought to determine the rate at which IgE and EoE food allergy co-occur. It has previously been shown that patients with EoE have a high rate of IgE-mediated food allergy.⁽²⁷⁾ We measured a rate of IgE-mediated food allergy in patients with EoE of 68%, a prevalence that is higher than previously reported.^(27, 28) To ensure accurate measurement of IgE-mediated food allergy in patients with EoE, we performed a manual chart review of 100 patients. In this review, 94% of patients with both an IgE-mediated food allergy and EoE diagnosis were found to have evidence of IgE-mediated allergy to at least one food.

To measure the prevalence of EoE in patients with IgE-mediated food allergy, we determined the number of patients in our EMR IgE cohort who had a diagnosis of EoE (Fig 1b). Of the 12,083 patients with IgE-mediated food allergy, 570 patients had a diagnosis of EoE indicating a prevalence of 4.72% (~1 in 20). In comparison, the prevalence of EoE in the general population nationally is estimated to be 1 in 2500 when utilizing either a similar ICD-based methodology⁽²⁹⁾ or a national questionnaire of physicians.⁽³⁰⁾ Together, these findings indicate that the co-occurrence of IgE-mediated food allergy and EoE is high, and that the prevalence of EoE in patients with IgE-mediated food allergy is higher than that observed in the general population (Table 2).

Association between IgE-mediated allergy to specific foods and EoE

We next used multivariate logistic regression analysis of our EMR cohort of 12,083 patients with IgE-mediated food allergy to determine whether demographic variables, or the presence of IgE-mediated allergy to specific foods, were associated with a diagnosis of EoE. Consistent with previous reports, we found that EoE is more commonly diagnosed in males (OR 2.27, 95% CI 1.89–2.78) and Caucasians (OR 1.99, 95% CI 1.55–2.56) (Table 3).⁽³¹⁾

We next sought to measure the extent to which IgE-mediated food allergy to specific foods was associated with EoE. We took advantage of different causal allergen patterns for IgE-mediated food allergy and EoE to differentiate foods causing IgE-mediated food allergy from foods causing EoE in our EMR cohort. To ensure that we were accurately measuring specific IgE-mediated food allergens in patients with EoE, we performed a manual chart review of 100 patients in our EMR cohort with both IgE-mediated food allergy and EoE. EMR food allergen data for the top five causes of IgE-mediated food allergy in patients with EoE (peanut, egg, tree nut, milk, shellfish) had a true positive rate (positive predictive value) of 91% for foods causing IgE-mediated allergy. There were no false-negatives (i.e. cases where a child was allergic to a specific food that was not identified in the EMR food allergy data).

We next performed multivariate logistic regression analysis of our EMR cohort dataset to determine the extent to which IgE-mediated food allergy to specific foods is associated with EoE. We found that IgE-mediated allergy to egg (2.27; 1.91–2.64), milk (4.19; 3.52–4.97), or shellfish (1.55; 1.24–1.92) was significantly associated with a diagnosis of EoE (Table 3). Furthermore, we found that more than one IgE-mediated allergy diagnosis to a top 5 food (peanut, egg, tree nut, milk, shellfish) was associated with an increased risk of EoE (Table 3). Together, these findings indicate that individuals with IgE-mediated allergy to specific foods are at increased risk of developing EoE, and that individuals are at increased risk of EoE with each additional IgE-mediated food allergy.

Discussion

Eosinophilic esophagitis is an increasingly common cause of esophageal symptoms in pediatric and adult patient populations. The pathophysiologic relationship between IgE-mediated food allergy and EoE continues to be clarified. Here we provide one of the largest retrospective cohort studies of patients with EoE ever performed. We find the most common causal foods of EoE to be milk, soy, egg, grains, and meats. In comparison, the pattern of IgE-mediated food allergens in patients with EoE are distinct, and similar to those observed in the general population (peanut, tree nut, egg, milk, and shellfish). Together, these findings support different pathophysiology for these two conditions.

While IgE-mediated and EoE food allergy have been reported to occur together,^(14, 15, 23, 28) it is difficult to estimate simultaneous occurrence of these conditions to the same food as avoidance of the causal food in IgE-mediated food allergy ameliorates EoE symptoms. Additionally, patients with EoE and IgE-mediated food allergy may display distinct clinical symptoms as compared to those without IgE-mediated food allergy, making EoE diagnosis difficult.⁽²⁸⁾ As such, there are likely a proportion of patients with concurrent IgE-mediated food allergy and EoE in which EoE is not detected.⁽²³⁾ We observed a higher rate of IgE-mediated food allergy in patients with EoE as compared with prior studies,^(27, 28) a finding which was confirmed by manual chart review. While differences in methodology may account for the higher rates of IgE-mediated food allergy observed in our study, it is possible that IgE-mediated food allergy is more common in patients with EoE than previously appreciated. As such, further study of the rate of IgE-mediated food allergy in patients with EoE is warranted.

Recently, diagnoses of EoE in patients undergoing OIT for IgE-mediated food allergy has raised concern that OIT could trigger EoE, at rates ranging from 1 to 10% depending on methodology.⁽³²⁾ The association between OIT and EoE was the focus of a recent meta-analysis which reported a positive correlation between OIT and EoE, with new onset of EoE occurring in 2.7% of patients undergoing OIT.⁽³³⁾ However, this meta-analysis defined EoE by biopsy, excluding patients with symptoms of EoE (such as abdominal pain or vomiting) that did not undergo biopsy. In our review of studies that documented discontinuation of OIT due to EoE symptoms and/or consistent biopsy findings, we found that symptoms of EoE occur at a rate between 8 and 14% (Table 4). While it should be noted that abdominal pain or vomiting are non-specific symptoms that are not diagnostic for EoE, many patients who

discontinue OIT due to these symptoms do not obtain biopsies. As such, actual rates of EoE during OIT may be higher than previously appreciated.

Our findings indicate that the prevalence of EoE in patients with IgE-mediated food allergy is higher than the rate of EoE in the general population,⁽³¹⁾ and similar to rates of biopsy-confirmed EoE observed during trials of OIT (Table 4).^(33–35) Furthermore, we found that allergy to three of the most common IgE-mediated food allergens (milk, egg, and shellfish) was highly associated with diagnosis of EoE. Together, our findings suggest that IgE-mediated food allergy is a marker of risk for development of EoE, possibly because of shared genetic predisposition or environmental modulators. Furthermore, the concern for iatrogenic EoE in patients undergoing OIT is likely warranted, and patients undergoing OIT should be closely monitored for development of EoE symptoms.

Limitations

This study is a secondary analysis of health records of a sub-specialty clinic at a single institution collected as part of routine care. Utilization of a sub-specialty cohort can result in selection bias. Additionally, we relied on diagnosis codes to identify conditions of interest, and choice of diagnosis codes can be affected by billing or administrative constraints. However, in a chart review of a subset of our cohorts we found a high degree of diagnosis code accuracy when compared to commonly accepted practice parameters. In patients with history consistent with IgE-mediated food allergy, diagnosis of allergy to a food not yet consumed may be made based on positive testing. With this methodology, an over-estimation of IgE-mediated food allergy rates could occur. Finally, our patients primarily reside in an urban and suburban setting which should be considered when extrapolating to rural communities or comparing to national studies.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations

EoE	Eosinophilic Esophagitis
OIT	Oral Immunotherapy
CPED	Center for Pediatric Eosinophilic Diseases

EMR	Electronic Medical Record
IRB	Institutional Review Board
CHOP	Children's Hospital of Philadelphia

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What is already known about this topic?

IgE-mediated food allergy is associated with comorbid atopic conditions such as asthma and allergic rhinitis. However, it is not known whether patients with IgE-mediated food allergy are at risk for developing eosinophilic esophagitis (EoE).

What does this article add to our knowledge?

We find that the causal allergens for IgE-mediated and EoE food allergy are distinct, that the prevalence of EoE is higher in the IgE-mediated food allergy population, and that allergy to common IgE-mediated food allergens is associated with developing EoE.

How does this study impact current management guidelines?

The association between IgE-mediated food allergy and EoE suggests a shared predisposition, and has implications for the management and monitoring of children with food allergy. Our results identify a very real risk of EoE in patients with IgE-mediated food allergy, and have implications for patients undergoing oral immunotherapy.

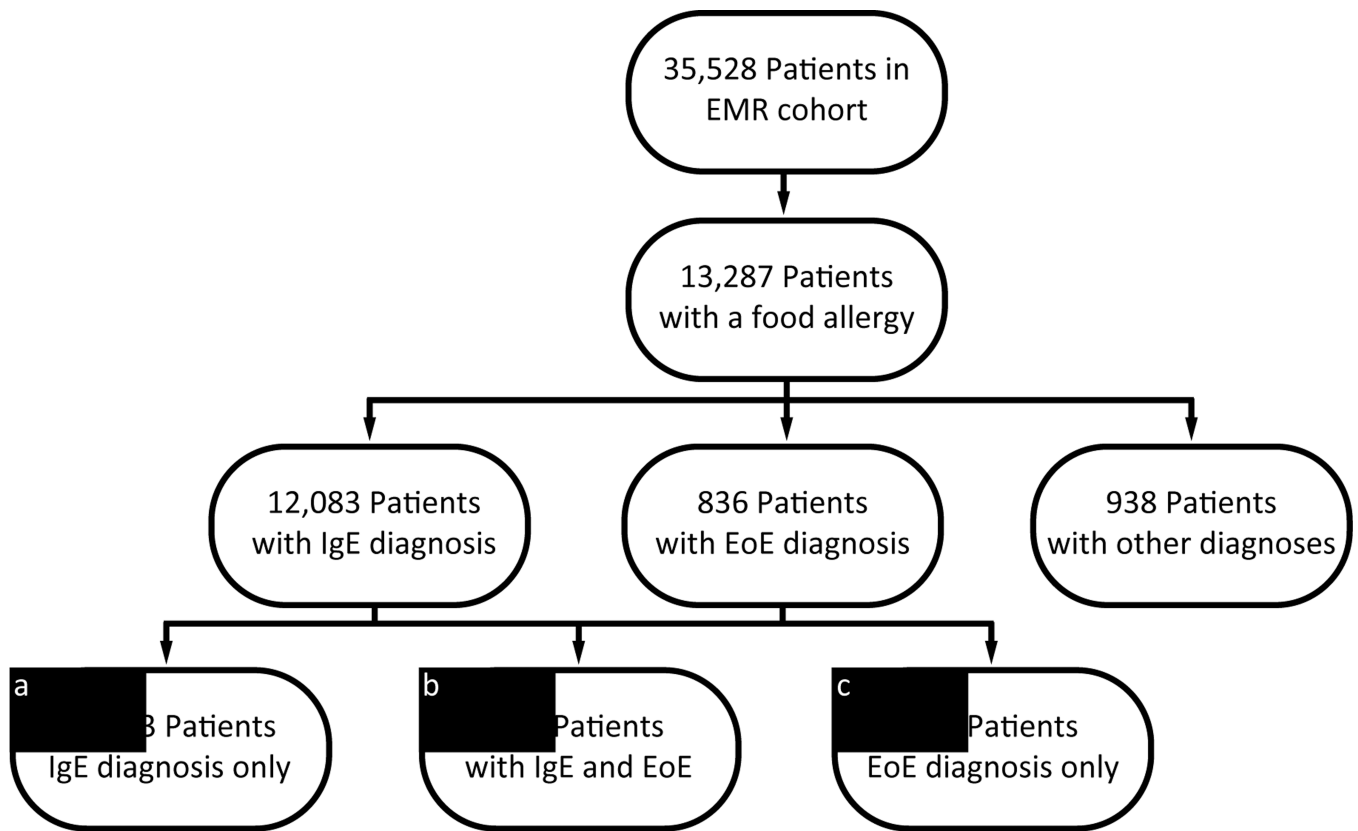


Figure 1.

EMR study cohorts. Three study cohorts were identified: **(a)** A cohort of 11,513 children with IgE-mediated food allergy in the absence of EoE, **(b)** a cohort of 570 children with concurrent IgE-mediated food allergy and EoE, and **(c)** a cohort of 266 children with EoE in the absence of IgE-mediated food allergy.

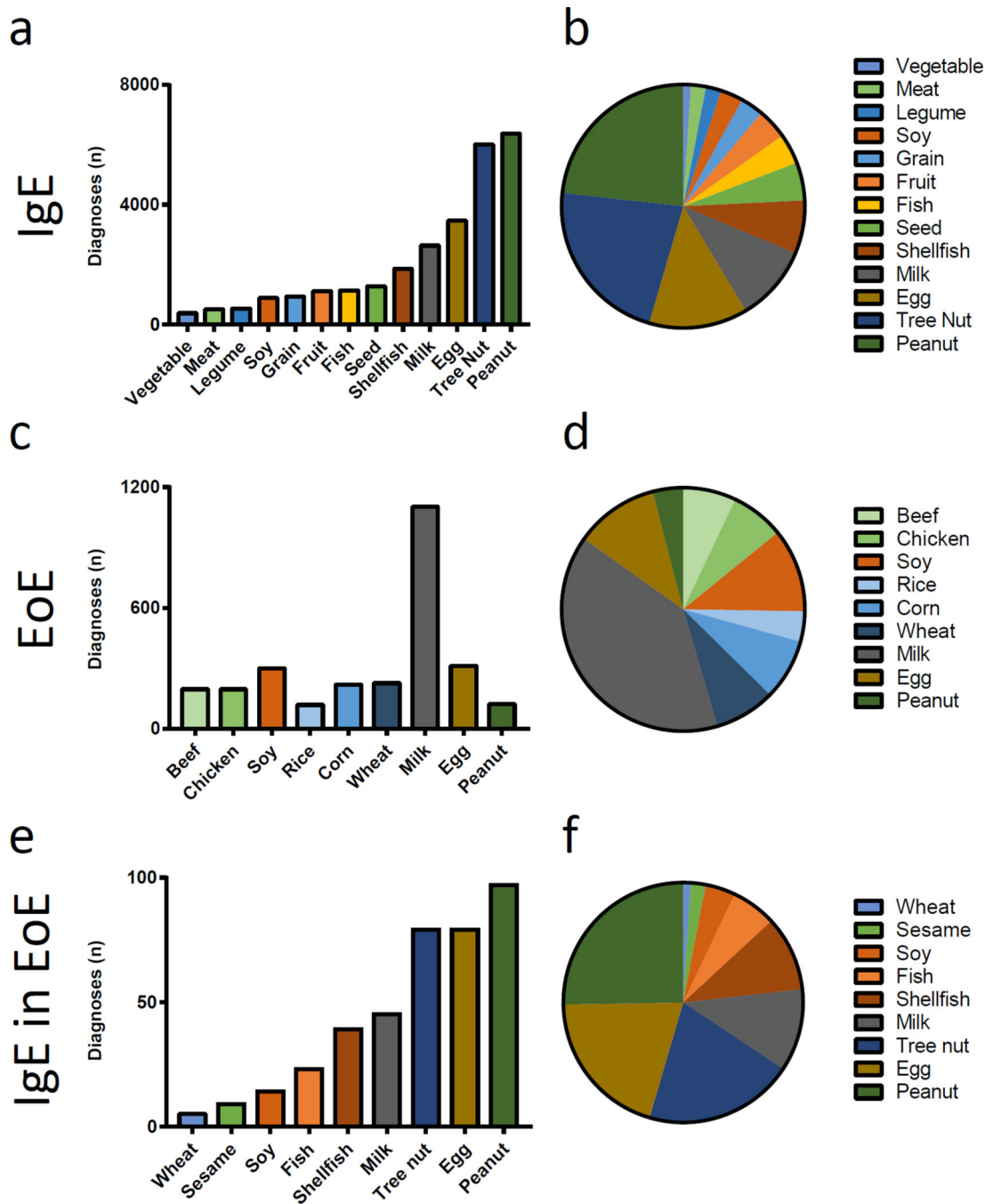


Figure 2. The most common food allergens in patients with IgE-mediated food allergy or EoE. (a) Number of diagnoses for the most common food allergens in patients with IgE-mediated food allergy. (b) Frequency of the most common food allergens in patients with IgE-mediated food allergy. (c) Number of diagnoses for the most common food allergens in patients with EoE food allergy. (d) Frequency of the most common food allergens in patients with EoE food allergy. (e) Number of diagnoses for the most common IgE-mediated food

allergens in patients with EoE. (f) Frequency of the most common IgE-mediated food allergens in patients with EoE.

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Table 1

Demographic characteristics of the EMR cohorts

Characteristic	Frequency, % (n)	
	IgE (12,083)	EoE (836)
<i>Gender</i>		
Male	61 (7,393)	76 (638)
Female	39 (4,690)	24 (198)
<i>Race</i>		
White	58 (6,994)	74 (621)
Black	20 (2,463)	11 (88)
American Indian or Alaska Native	<1 (15)	<1 (1)
Asian	6 (750)	2 (14)
Native Hawaiian or Other Pacific Islander	<1 (11)	0 (0)
Other	13 (1620)	12 (99)
Multiple	1 (168)	1 (12)
Unknown	<1 (11)	<1 (1)
Refused	<1 (51)	0 (0)
<i>Ethnicity</i>		
Non-Hispanic or Latino	88 (10,692)	85 (714)
Hispanic or Latino	5 (591)	5 (43)
Unknown	5 (599)	9 (76)
Refused	2 (201)	<1 (3)
<i>Birth year</i>		
Before 2000	9 (1044)	21 (178)
2000 to 2004	20 (2421)	31 (261)
2005 to 2009	36 (4319)	34 (283)
2010 to 2015	36 (4299)	14 (114)

Table 2

Rate of EoE

	Rate	Source	References^a
General Population	0.0004	ICD, Survey	1, 2
IgE Population	0.0472	ICD	

^aSee supplemental references

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Table 3

Association between patient characteristics and EoE in the EMR cohort

Characteristic	Odds Ratio	95% CI	P value
<i>Gender</i>			
Male vs Female	2.27	1.89 – 2.78	< 0.0001
<i>Race</i>			
White vs Black	1.99	1.55 – 2.56	< 0.0001
<i>Ethnicity</i>			
Hispanic vs Non-Hispanic	0.99	0.67 – 1.47	ns
<i>Specific Food Allergens</i>			
Peanut	1.07	0.90 – 1.27	ns
Egg	2.27	1.91 – 2.64	<0.0001
Tree nut	1.04	0.88 – 1.23	ns
Milk	4.19	3.52 – 4.97	<0.0001
Shellfish	1.55	1.24 – 1.92	<0.0001
<i>Number of Food Allergies</i>			
2 foods	2.93	2.16 – 4.05	<0.0001
3+ foods	5.29	3.82 – 7.32	<0.0001

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Table 4

Rate of abdominal pain and biopsy-confirmed EoE during OIT

Food	Abdominal pain	Biopsy	References^a
Milk	14.3%	4.7%	3–11
Egg	7.9%	2.5%	12–21
Peanut	13.4%	7.3%	22–33

^aSee supplemental references

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