Etiologies of Pediatric Cervical Lymphadenopathy: A Systematic Review of 2687 Subjects

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Abstract

Background. Cervical lymphadenopathy in children is common and its etiologies diverse. No systematic review of the differential diagnosis of pediatric cervical lymphadenopathy has been conducted. Objective. To determine the prevalence rate of specific etiologies of pediatric cervical lymphadenopathy. Data Sources. EMBASE, PubMed, and SCOPUS were searched electronically. Bibliographies of select studies were reviewed as well. Study Selection. (1) Any clinical trial, observational study, or cross-sectional case series with 10 or more subjects that included delineation of etiologies and/or associated conditions with lymphadenopathy; (2) subjects aged 0 to 21 years with enlarged lymphoid tissue on body; (3) lymphadenopathy was confirmed by clinical evaluation; and (4) no specific diagnoses were excluded. Data Extraction. Year and location of publication, definition of lymphadenopathy, percentage of lymphadenopathy that was cervical, total number of subjects, gender distribution of subjects, age range of patients, and specific etiologies. Results. Of the 1790 studies, 7 studies that were combined resulted in 2687 subjects that were selected. Nonspecific benign etiology was the most common diagnosis occurring at a rate of 67.8%. Epstein-Barr virus was the next most prevalent (8.86%), followed by malignancy (4.69%) and granulomatous disease (4.06%). The most common malignancy etiology was non-Hodgkin’s lymphoma (46.0%), and the most common granulomatous disease was tuberculosis (73.4%). Conclusions. This systematic review and meta-analysis provides a rate-based differential diagnosis of pediatric cervical lymphadenopathy. Although the most common causes of pediatric cervical lymphadenopathy are nonspecific, the etiologies are diverse. Rates and credible intervals are provided to enable a probability-based diagnostic approach to palpable cervical lymphadenopathy in this age group.

Keywords
cervical lymphadenopathy, pediatrics, differential diagnosis

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Introduction

Cervical lymphadenopathy is a common complaint in the pediatric population. The prevalence rate of cervical lymphadenopathy, commonly defined as cervical nodal tissue measuring >1 cm in diameter, is estimated at 38% to 45% in otherwise healthy children.1 The differential diagnosis for cervical lymphadenopathy in children is broad, including both common benign etiologies and much more rare malignant causes.1,2 While myriad clinical trials, observational studies, and case series appear in the literature, a systematic review delineating the etiologies of pediatric cervical lymphadenopathy does not presently exist. The objectives of this study were to (1) identify the most common etiologies associated with pediatric cervical lymphadenopathy and (2) determine the relative prevalence of etiologies across a wide spectrum of patients.

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Methods

Protocol

This study followed the guidelines of the Preferred Reporting Items in Systematic Reviews and Meta- Analyses. \(^3\)

Eligibility

In order to be included in this review, the following criteria were developed a priori: (1) any clinical trial, observational study, or cross-sectional case series with 10 or more subjects that included delineation of etiologies and/or associated conditions with lymphadenopathy; (2) ages 0 to 21 years with enlarged lymphoid tissue and lymphadenopathy confirmed by clinical evaluation; and (3) no specific diagnoses were excluded. Case reports, review articles, and editorials were excluded. There were no restrictions regarding date of publication or language; however, abstracts were required to be in English.

Information Sources

EMBASE, PubMed, and SCOPUS were searched electronically. Bibliographies of selected studies were reviewed manually as well.

Search

The main search terms were “pediatric,” “lymphadenopathy,” and “children.” The following filters were used during the search: human, no language restriction, no publication status restriction, all dates, ages 0 to 24 years. The search was performed individually by authors AD and KD. The results of the searches were pooled and duplicates were eliminated.

Study Selection

Each article was evaluated by authors AD and KD. Any differences in judgment were resolved by consensus. All selected studies were reviewed by a third author MD.

Data Collection

For each selected article, the following information was recorded: year and location of study population, definition of lymphadenopathy, percentage of lymphadenopathy that was cervical, total number of subjects, gender distribution of subjects, age range of patients, and specific etiologies. Major categories for specific etiologies for pediatric cervical lymphadenopathy included malignancy, nonspecific changes, and granulomatous, infectious, and autoimmune diseases. Within each of these categories more specific etiologies were developed from the data. Data extraction was performed by authors AD and KD and reviewed by MD.

Synthesis of Results

A Bayesian methodology for point estimation was used as previously described. \(^4\) Posterior probabilities are reported as rates with the associated 95% credible intervals. All calculations were performed in the R environment.

Sources of Bias Across Studies

Potential sources of bias in this study include the following: inconsistency in individual studies regarding data in the text when compared with data presented in tables, inconsistencies in definition and terms across studies, and a percentage of patients with lymphadenopathy in areas other than cervical. To identify articles in which cervical lymphadenopathy was studied in conjunction with other regional lymphadenopathies, a broad search term of “lymphadenopathy” was used in the initial search terms. Once the source literature was pooled, eliminations were made to narrow the focus to cervical lymphadenopathy. Articles describing patients with peripheral lymphadenopathy in addition to cervical were accepted only if \(\geq 90\%\) of the subjects had cervical lymphadenopathy.

Ethical Approval and Informed Consent

This study did not involve any direct contact with patients. Public data sources were solely used thus not requiring the institutional review board approval or patient consent.

Results

Study Selection

The results of the literature search are summarized in Figure 1. Searches of EMBASE, PubMed, and SCOPUS resulted in 1790 articles. Reviews of the bibliographies of selected articles and a search of the OVID Cochrane Reviews failed to yield any additional source references. A total of 1750 studies were excluded after review of the title. An additional 21 were excluded after review of the abstract. The full texts of the remaining 19 articles were assessed in detail; 12 were excluded. Three of the discarded studies were review articles and 5 had exclusion and inclusion criteria that eliminated certain etiologies
of cervical lymphadenopathy from the study. Another study was excluded because it did not specify the age range of the patients. Attempts to contact the author were not successful. Three articles were excluded because the percentage of children with cervical lymphadenopathy was less than 90%. Attempts to contact these authors were also not successful. The remaining 7 articles were selected for review. Though the final articles selected for review were only 7 of the original 1790, this was after carefully eliminating articles that failed to meet inclusion and exclusion criteria, and had sufficient data to draw conclusions specifically about pediatric cervical lymphadenopathy. Though a prevalent presenting symptoms in pediatric patients globally, and articles were reviewed from countries across the world including the United States, only these 7 met our criteria.

Study Characteristics and Outcomes

The 7 studies selected for this review are summarized in Table 1. These studies ranged in size from 49 to 1700 subjects and had a combined total of 2687 children. The subjects represent a global pediatric population from Nigeria, Turkey, Italy, India, and Egypt.

Figure 1. Flow diagram of literature search.
One study consisted of patients referred to an oncology clinic. In another study, multiple lymph node aspirates were taken from subjects and the etiologies were reported by the number of aspirates (n = 238) not by the number of subjects (n = 217). Another series reported both cervical and peripheral lymphadenopathy; the diagnoses specific to cervical lymphadenopathy were identifiable, but age and gender data specific to these patients were not.

Table 1. Summary of Included Studies.

| Study                                    | No. of Patients | No. of Females/Males | Age Range   | Country   |
|------------------------------------------|-----------------|----------------------|-------------|-----------|
| Ejeckam and Nwabueze (1984)             | 100             | 32/68                | <20 years   | Nigeria   |
| Reddy et al (2002)                       | 100             | 21/79                | 1 month to 12 years | India    |
| Citak et al (2011)                       | 282             | 124/158              | 6 months to 16 years | Turkey   |
| De Corti et al (2014)                    | 238             | 91/126               | 3 months to 17 years | Italy    |
| Gwili et al (2015)                       | 49              | 35/85                | 1 month to 18 years | Egypt    |
| Bozlak et al (2016)                      | 218             | 75/143               | 13 months to 18 years | Turkey   |
| Sarsu and Sahin (2016)                   | 1700            | 697/1003             | <18 years   | Turkey    |

Discussion

Pediatric cervical lymphadenopathy is a common problem with a broad differential diagnosis. Synthesized data obtained from 7 studies representing 5 countries and including 2687 patients demonstrates that two thirds of cases have no identifiable etiology. Of the patients who received a definitive diagnosis, Epstein-Barr virus, malignancy, and granulomatous disease accounted for 8.86%, 4.69%, and 4.06% of cases, respectively. Non-Hodgkin’s lymphoma was the most common malignant etiology (46.0% of malignancies), while tuberculosis was the most common granulomatous disease (73.4% of cases of granulomatous origin).

While every article selected for this study did provide an age range of their subjects, age range of each specific etiology was not always indicated. In the articles with specific malignant etiology identified, subjects were under the age of 18 years, and the youngest subject was...
between 1 month and 13 months. Also common to all these articles was the presenting symptom of cervical lymphadenopathy. Then depending on other presenting symptoms and examinations, there was further workup of laboratory tests, diagnostic imaging such as ultrasound, fine-needle aspiration, and/or biopsy, which established the malignant etiology.

Common signs found in history included cough, fever, changes in weight, or sore throat. In Citak et al., 3 of the patients with Hodgkin’s or non-Hodgkin’s lymphoma had weight loss and night sweats; common to all of the patients with lymphoma was the physical examination finding of rubbery and fixed lymph nodes. Other physical examination findings

Table 2. Summary by Disease Category.

| Etiology                                | No. of Patients | Prevalence Rate (% of Total) | 95% Confidence Interval |
|-----------------------------------------|-----------------|------------------------------|-------------------------|
|                                         |                 |                              | Lower (%) | Upper (%) |
| Nonspecific diagnosis                   | 1822            | 67.8                         | 66         | 69.6      |
| Epstein-Barr virus                      | 238             | 8.86                         | 7.81       | 9.96      |
| Malignant                               | 126             | 4.69                         | 3.92       | 5.52      |
| Granulomatous                           | 109             | 4.06                         | 3.34       | 4.83      |
| Cytomegalovirus                         | 108             | 4.02                         | 3.31       | 4.79      |
| Group A/β Hemolytic Streptococcus       | 78              | 2.9                          | 2.3        | 3.57      |
| Rubella                                 | 46              | 1.71                         | 1.26       | 2.23      |
| Toxoplasmosis                           | 32              | 1.19                         | 0.816      | 1.63      |
| Abscess                                 | 28              | 1.04                         | 0.694      | 1.46      |
| Systemic lupus erythematosus            | 22              | 0.819                        | 0.514      | 1.19      |
| Cat scratch disease                     | 17              | 0.633                        | 0.369      | 0.966     |
| Kawasaki disease                        | 14              | 0.521                        | 0.285      | 0.826     |
| Sarcoïdosis                             | 10              | 0.372                        | 0.179      | 0.635     |
| Staphylococcal pharyngitis              | 8               | 0.298                        | 0.129      | 0.536     |
| Streptococcal tonsillitis               | 7               | 0.261                        | 0.105      | 0.486     |
| Phlegmons                               | 6               | 0.223                        | 0.082      | 0.434     |
| Mycoplasma pneumonia                    | 3               | 0.112                        | 0.023      | 0.269     |
| Parvovirus B19                          | 2               | 0.0744                       | 0.00902    | 0.207     |
| Familial Mediterranean fever            | 2               | 0.0744                       | 0.00902    | 0.207     |
| AIDS                                    | 2               | 0.0744                       | 0.00902    | 0.207     |
| Dermatomyositis                         | 2               | 0.0744                       | 0.00902    | 0.207     |
| Streptococcal pneumoniae                | 1               | 0.0372                       | 0.000943   | 0.137     |
| Rosai-Dorfman disease                   | 1               | 0.0372                       | 0.000943   | 0.137     |
| Kilkuchi-Fugimoto disease               | 1               | 0.0372                       | 0.000943   | 0.137     |
| PFAPA syndrome                          | 1               | 0.0372                       | 0.000943   | 0.137     |
| Lymphomatoid papillomatosis             | 1               | 0.0372                       | 0.000943   | 0.137     |

Table 3. Malignant Etiologies in 126 Patients With Cervical Lymphadenopathy.

| Etiology                                | No. of Patients | Prevalence Rate (% of Total) | 95% Confidence Interval |
|-----------------------------------------|-----------------|------------------------------|-------------------------|
|                                         |                 |                              | Lower (%) | Upper (%) |
| Unspecified malignant                   | 58              | 46                           | 37.4       | 54.7      |
| Non-Hodgkin’s lymphoma                  | 34              | 27                           | 19.6       | 35        |
| Hodgkin’s lymphoma                      | 27              | 21.4                         | 14.7       | 29        |
| Langerhans cell histiocytosis           | 3               | 2.38                         | 0.498      | 5.66      |
| Acute lymphocytic leukemia              | 1               | 0.794                        | 0.0203     | 2.91      |
| Hemophagocytic lymphohistiocytosis      | 1               | 0.794                        | 0.0203     | 2.91      |
| Leukemia                                | 1               | 0.794                        | 0.0203     | 2.91      |
| Neuroblastoma                           | 1               | 0.794                        | 0.0203     | 2.91      |
common to lymph nodes of malignant etiology were bilateral, multiple, non-tender nodes that did not present with any skin changes. These findings in the history and physical led to further workup, and some of these predicative factors for malignancy included enlargement of lymph node in follow-up, elevated serum C-reactive protein, lactate dehydrogenase, and uric acid, and leucopenia. Ultrasound findings showed no hilum and undetectable cortical and medullary contours.5,11

Regarding the specific etiologies of malignancy found across the articles, unspecified was the most common followed by non-Hodgkin’s lymphoma at 27% and Hodgkin’s lymphoma at 21%. This is not to say necessarily that non-Hodgkin’s is the more common malignant etiology when presenting with cervical lymphadenopathy. Had the other articles specified beyond malignant versus nonmalignant, another malignant etiology may have been made more prevalent. Common to all the articles is a careful history paired with the above physical examination findings and laboratory/diagnostics that then led to a procedure confirming the malignant etiology. These findings help practitioners differentiate between benign and malignant etiology of cervical lymphadenopathy, as the vast majority of the time the etiology will be nonmalignant in nature. Reassuring to both the clinical provider and the pediatric patient and their family is that in the vast majority of cases, the etiology of cervical lymphadenopathy is benign and nonspecific, rather than malignant.

Risk of Bias

The largest study in this review contributed 63.3% of the total number of subjects. This article was included after analysis of its data showed consistency with the smaller studies. There were many variances in nomenclature to define nonspecific, benign disease. The breakdown of these terms can be seen in Table 5. The large number of tuberculosis cases reflects the countries of origin included in this study; no large studies from the United States were included in this study.

Limitations

The main limitations of this study are variations in terminology and diagnostic evaluation across studies illustrated in Table 5. However, all articles included were required to demonstrate a thorough diagnostic evaluation.
Another limitation of this study was the need to extrapolate data specific to the cervical lymph node region from articles in which etiology and workup of peripheral sites of lymphadenopathy were included. In some articles it was possible to extract the diagnoses pertinent to cervical lymphadenopathy from the included tables. In others, it was not possible to differentiate between the cases of cervical and peripheral lymphadenopathy. In the last instance, only articles in which more than 90% of cases were identified as cervical lymphadenopathy were included.

Finally, given that many of the leading specific etiologies identified in this study are of infectious origin, the locations where the included studies were performed may have influenced the prevalence of certain infectious diseases, primarily tuberculosis. Of the 7 studies included in this meta-analysis, 2 were in tuberculosis endemic regions (Nigeria and India), while the remaining 5 studies took place in regions not endemic for tuberculosis. However, while the tuberculosis rates differed based on study location, the rates of other infections etiologies remained consistent among studies.

Conclusions
Cervical lymphadenopathy in the pediatric population is a common presenting complaint with myriad possible etiologies. The 3 most common etiologies identified include nonspecific diagnosis, Epstein-Barr virus, and malignancy. As with any systematic review, limitations and potential sources of bias exist. By combining these data with the patient’s clinical picture, the treating physician can estimate the probability of a specific diagnosis and more accurately evaluate the need for further diagnostic evaluation.

Author Contributions
A.D helped conceive the topic for study, performed the original search, aided in data collection and writing the manuscript;
K.D helped conceive the topic for study, performed original searches, aided in data collection and in writing the manuscript;
S.A aided in conception of the topic, performed the statistical analysis and aided in final manuscript preparation;
M.D.V conceived the topic of study, directed the search methods and aided in final manuscript preparation.

Declaration of Conflicting Interests
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