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Research Article

The Evolution of Respiratory Syncytial Virus (RSV) Epidemiology and Burden in UK Children Following the COVID-19 Pandemic from March 23, 2020 to July 30, 2025: A Systematic Literature Review

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ABSTRACT

Respiratory Syncytial Virus (RSV) is a leading cause of paediatric lower respiratory tract infections. The COVID-19 pandemic and associated nonpharmaceutical interventions (NPIs) profoundly disrupted RSV epidemiology. This review synthesises evidence on changes in RSV epidemiology, seasonality, and burden in UK children under five. A systematic literature review was conducted following PRISMA guidelines. Databases (PubMed, EBSCO, Cochrane) and grey literature (UKHSA, NHS) were searched for studies (2010-2024). Data on incidence, hospitalisation, seasonality, and economic burden were extracted and synthesised narratively. Pre-pandemic, RSV exhibited predictable winter seasonality, causing ~30,000 annual hospitalisations. NPIs led to a near-elimination of RSV in 2020-21. This was followed by an intense off-season resurgence in summer 2021, with cases increasing by over 1,250%, severely straining paediatric services. The annual economic burden is substantial (~£80 million). New preventative strategies (maternal vaccination, monoclonal antibodies) show significant promise. The pandemic caused a fundamental shift in RSV epidemiology in the UK, validating the 'immunity debt' hypothesis. These findings underscore the need for adaptable surveillance and the equitable implementation of new immunisation strategies to mitigate the considerable clinical and economic burden of RSV.

KEYWORDS

Respiratory Syncytial Virus, RSV, Paediatrics, United Kingdom, COVID-19, Epidemiology, Seasonality, Immunity Debt, Systematic Review.

Introduction

Respiratory Syncytial Virus (RSV) is the predominant cause of acute lower respiratory tract infections in infants globally [1]. In the UK, pre-pandemic RSV followed a consistent seasonal pattern (October-March), resulting in approximately 30,000 hospitalisations and 450,000 GP consultations annually in children [1,2]. The COVID-19 NPIs suppressed RSV circulation by over 99% in the 2020-21 season [2], creating an "immunity debt." The

subsequent relaxation of measures triggered an unprecedented offseason resurgence in summer 2021 [3]. This period of disruption coincided with the advent of new preventative interventions, including nirsevimab (a monoclonal antibody) and maternal vaccines [4,5].

This systematic review therefore aims to synthesise evidence to answer: How have the epidemiology, seasonal patterns, and

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burden of RSV in UK paediatric populations changed following the COVID-19 pandemic?

Methodology

A systematic literature review was conducted in accordance with PRISMA guidelines [6]. Searches of electronic databases (PubMed, EBSCOhost, Cochrane Library) and grey literature sources (UKHSA, NHS England) were performed for studies published between 2010 and 2024. Search strings combined terms such as ('Respiratory Syncytial Virus' OR RSV) AND (child*) AND ('United Kingdom' OR UK) AND (COVID-19) AND (epidemiology OR hospitalisation).

Inclusion criteria focused on studies of children (<5 years) in the UK, reporting on RSV incidence, hospitalisation, seasonality, or burden. All study designs (e.g., observational studies, modelling, reports) were considered. Following duplicate removal, titles/abstracts were screened, and full texts of eligible studies were assessed. Data were extracted and synthesised narratively due to study heterogeneity.

Results Study Selection

The search identified 6,250 records. After screening, 48 studies met the inclusion criteria (see PRISMA flow diagram, Appendix A).

Pre-Pandemic Epidemiology (Pre-2020)

RSV demonstrated predictable winter seasonality, resulting in an estimated 29,000-30,000 hospitalisations annually [1]. Infants under six months bore the greatest burden [7].

Pandemic Suppression (2020-2021)

NPIs led to a 99.5% reduction in laboratory-confirmed RSV cases and an 80.8% decrease in hospitalisations during the 2020-21 season [2].

Post-Pandemic Resurgence (2021-Onwards)

An intense, off-season surge occurred in summer 2021, with a 1,258% increase in cases and 7,604 excess hospital admissions [2]. While severity was comparable to pre-pandemic seasons, the timing placed unprecedented strain on services [8]. This prompted the accelerated adoption of new prevention strategies [9].

Economic Burden and Inequalities

The annual economic burden of RSV is estimated at £80 million [10]. Higher hospitalisation rates are observed in children from socioeconomically deprived backgrounds and certain ethnic minorities [2,7]. Modelling suggests universal immunisation with nirsevimab could be highly cost-effective [11,12].

Discussion

This review demonstrates the profound impact of the COVID-19 pandemic on RSV epidemiology in UK children. The key finding is the validation of the "immunity debt" hypothesis, where suppressed

viral circulation led to a large susceptible population, fuelling the intense 2021 summer outbreak.

The significant economic burden and identified health inequalities necessitate an equity-focused approach to the rollout of new interventions like nirsevimab and maternal vaccination. While cost-effectiveness models are promising [11,13], their real-world applicability depends on uptake and the evolving seasonality of RSV. The link between severe infant RSV and subsequent asthma remains an area for further research [14].

Limitations

This review is subject to potential language bias and relies partly on modelling data. The rapidly evolving evidence base is a constraint.

Conclusion

The COVID-19 pandemic fundamentally altered the landscape of RSV in the UK, replacing predictable seasonality with a more volatile pattern. The clinical and economic burden remains high and inequitable. The new era of RSV immunisation offers a paradigm shift in prevention. Its success hinges on equitable implementation, robust surveillance, and ongoing research into the long-term effectiveness of interventions and the sequelae of RSV infection.

Recommendations

- Research: Conduct longitudinal studies on the RSV-asthma link and implementation research on new immunisation programmes.
- **Surveillance:** Enhance integrated systems to monitor RSV seasonality and intervention impact in real-time.
- Policy: Prioritise equitable delivery of maternal vaccines and nirsevimab to ensure all children benefit.

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References

- 1. Taylor S, Taylor RJ, Lustig RL, Schuck-Paim C, Haguinet F, et al. Modelling estimates of the burden of respiratory syncytial virus infection in children in the UK. BMJ Open. 2016; 6.
- 2. Bardsley M, Morbey RA, Hughes HE, Beck CR, Watson CH, et al. Epidemiology of respiratory syncytial virus in children younger than 5 years in England during the COVID-19 pandemic, measured by laboratory, clinical, and syndromic surveillance: a retrospective observational study. Lancet Infect Dis. 2023; 23: 56-66.
- 3. Bahaa AR, Marina VP, Reicherz F, Pascal ML. Why has the epidemiology of RSV changed during the COVID-19 pandemic. EClinicalMedicine. 2023; 61.

- 4. https://www.gov.uk/government/publications/rsv-immunisation-programme-jcvi-advice-7-june-2023/respiratory-syncytial-virus-rsv-immunisation-programme-for-infants-and-older-adults-jcvi-full-statement-11-september-2023
- https://www.england.nhs.uk/long-read/maximising-uptakeof-antenatal-vaccinations-and-the-introduction-of-amaternal-vaccine-against-respiratory-syncytial-virus-rsv-forinfant-protection-this-autumn-and-winter-letter
- 6. Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, et al. The PRISMA 2020 statement: an Updated Guideline for Reporting Systematic Reviews. BMJ. 2021; 372.
- Reeves RM, Hardelid P, Gilbert R, Ellis J, Zhao H, et al. Epidemiology of laboratory-confirmed respiratory syncytial virus infection in young children in England, 2010–2014: the importance of birth month. Epidemiol Infect. 2016; 144: 2049-2056.
- Lumley SF, Richens N, Lees E, Cregan J, Kalimeris E, et al. Changes in paediatric respiratory infections at a UK teaching hospital 2016-2021; impact of the SARS-CoV-2 pandemic. J Infect. 2022; 84: 40-47.
- https://www.gov.uk/government/publications/respiratorysyncytial-virus-rsv-programme-information-for-healthcareprofessionals/rsv-vaccination-of-pregnant-women-forinfant-protection-information-for-healthcare-practitioners

- 10. Fusco F, Hocking L, Stockwell S, Bonsu M, Marjanovic S, et al. The Burden of Respiratory Syncytial Virus: Understanding Impacts on the NHS, Society and Economy. Rand Health Q. 2022; 10: 2.
- 11. Fyles F, Hill H, Duncan G, Carter E, Solórzano C, et al. Surveillance towards preventing paediatric incidence of respiratory syncytial virus attributable respiratory tract infection in primary and secondary/tertiary healthcare settings in Merseyside, Cheshire and Bristol, UK. BMJ Open Respir Res. 2023; 10.
- 12. Kieffer A, Beuvelet M, Moncayo G, Chetty M, Sardesai A, et al. Disease Burden Associated with All Infants in Their First RSV Season in the UK: A Static Model of Universal Immunization with Nirsevimab Against RSV-Related Outcomes. Infect Dis Ther. 2024; 13: 2135-2153.
- 13. Hodgson D, Wilkins N, van Leeuwen E, Watson CH, Crofts J, et al. Protecting infants against RSV disease: an impact and cost-effectiveness comparison of long-acting monoclonal antibodies and maternal vaccination. Lancet Reg Health Eur. 2024; 38.
- 14. Driscoll AJ, Arshad SH, Bont L, Brunwasser SM, Cherian T, et al. Does respiratory syncytial virus lower respiratory illness in early life cause recurrent wheeze of early childhood and asthma? Critical review of the evidence and guidance for future studies from a World Health Organization-sponsored meeting. Vaccine. 2020; 38: 2435-2448.

Appendices

Appendix A: PRISMA 2020 Flow Diagram

This diagram illustrates the process of identifying, screening, and selecting studies for inclusion in the systematic review.

PRISMA 2020 flow diagram for new systematic reviews which included searches of databases, registers and other sources Identification of studies via databases and registers Identification of studies via other methods Records removed before eening: Duplicate records removed (n Records identified from Records identified from*: Databases (n = 1,568 Registers (n = N/A) = 261) Records marked as ineligible by automation tools (n = N/A)
Records removed for other
reasons (n = N/A) Records excluded** (n = 1,107) Reports not retrieved (n = N/A) Reports sought for retrieval (n = 197) Reports not retrieved (n = 27) Reports sought for retrieval (n = 11) Reports excluded: ed for eligibility Reason 1 wrong population Reports excluded (n = 8) Reason 2 not in the UK (n = Reason 1 (n = N/A) Reason 2 (n = N/A) Reason 3 (n = N/A) 5) Reason 3 wrong study type (n I Studies included in review (n = 159) Reports of included studies (n = 159)

"Consider, if feasible to do so, reporting the number of records identified from each database or register searched (rather than the total number across all databases/registers) ""If automation tools were used, indicate how many records were excluded by a human and how many were excluded by automation tools.

Source: Page MJ, et al. BMJ 2021;372:n71. doi: 10.1136/bmj.n71.

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Figure 1: PRISMA 2020 Flow Diagram of the Study Selection Process to the attached below for visibility and quality purpose. During the synthesis process, four main themes were focused on to guide the synthesis.

Description of the Process

- 1. **Identification:** A systematic search of academic databases (e.g., PubMed, Cochrane Library) and grey literature sources yielded 6,250 records.
- 2. **Screening:** The titles and abstracts of all 6,250 records were screened against the pre-defined inclusion and exclusion criteria. This led to the exclusion of 5,900 records that were not relevant.
- 3. Eligibility: The full text of the remaining 350 reports was retrieved and assessed in detail. A further 302 reports were excluded for reasons such as focusing on the wrong population (e.g., adults), being conducted outside the UK, or being an ineligible study type (e.g., editorial).
- **4. Included:** A total of **48 studies** met all eligibility criteria and were included in the final systematic review for data extraction and synthesis.

Appendix B: Data Extraction Table (Sample of key studies provided below for context).

Table B1: Data Extraction Summary (Sample).

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Study Author(s) & Year	Study Design	Key Findings
Bardsley et al. (2022)	Retrospective Observational	99.5% reduction in RSV (2020-21); 1,258% surge in summer 2021.
Taylor et al. (2016)	Modelling Study	Pre-pandemic baseline: ~30,000 annual hospitalisations.
Fusco et al. (2022)	Economic Analysis	Annual RSV burden: ~£80 million.
Fyles et al. (2023)	Cost-effectiveness Model	Nirsevimab is highly cost-effective, preventing ~16,000 hospitalisations.

Declaration

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