**Illness severity assessment for children with RTI: do parents and GPs agree?**

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**Abstract (N=250)**

**Background**

Children’s respiratory tract infection (RTI) severity assessments are known to differ between parents and clinicians, but determinants are unknown.

**Aim**

To investigate the agreement between, and compare the determinants of, parent and clinician severity scores.

**Design and Setting**

Secondary analysis of data from a prospective cohort study of 8394 children aged 3 months to 16 years presenting to primary care with acute (≤28 days) cough and RTI.

**Method**

Data on sociodemographic factors, parent reported symptoms, clinician reported physical examination findings and global illness severity assessments was used . Kappa-statistics used to investigate agreement and multivariable logistic regression to identify the factors associated with illness severity.

**Results**

Parents reported higher (mean 5.2 (SD= 1.8), median 5 (IQR 4-7)) illness severity than clinicians (mean 3.1 (SD= 1.7), median 3 (IQR 2-4), p<0.0001). There was low positive correlation between these scores (+0.43) and poor inter-rater-agreement between parents and clinicians (kappa 0.049). Number of clinical signs was highly correlated with clinician scores (+0.71). Parent-reported symptoms (in the past 24 hours) independently associated with higher illness severity scores (in descending order-of-importance) were severe fever, severe cough, breathing quickly, severe reduced eating, severe reduced fluid, severe disturbed sleep and change in cry. Three of these symptoms (severe fever, breathing quickly and change in cry) along with inter/subcostal recession, crackles/crepitations, nasal flaring, wheeze and drowsiness/irritability were associated with higher clinician scores.

**Conclusions**

Clinicians and parents use different factors and make different judgements about children’s RTI severity. Improved understanding of the factors concerning parents could improve parent-clinician communication and consultation outcomes.

**Keywords**

Primary health care, illness severity assessment, respiratory tract infection, fever

**How this fits in**

Parents’ medical advice seeking decisions are complex. An improved mutual understanding of the factors associated with illness severity assessment could facilitate better parent-clinician communication and improved parental understanding regarding the symptoms necessitating consultation.

**Introduction**

Respiratory tract infections (RTIs) are the most common reason parents take children to primary care worldwide,(1) and many are unnecessarily prescribed antibiotics. The decision to seek medical advice is complex(2, 3),as parents rely on the child’s physical expression of illness and their own interpretation of the symptoms.(4) Clinicians also report consultations for RTIs as complex as they manage clinical uncertainly regarding diagnosis and prognosis, which often results in ‘just in case’ antibiotic prescribing.(5)

Patient-centred care has been shown to improve the quality of doctor-patient communication, consultations and illness outcomes.(6) Reaching a shared understanding of the nature of the problem and its severity is central to patient-centred care.(7, 8) ‘Illness severity’ can best be referred to as the magnitude of the patient-perceived, clinically significant manifestations of disease processes that are associated with decrements in health related quality of life (HRQOL) or health status.(9) In children’s consultations, there are three interacting perceptions (child-parent-clinician) affecting illness severity assessment, with assessments likely to differ due to different factors taken into account.

Parent perceived severe illness in children with a RTI is one of the reasons parents choose to consult to primary care (10), and clinician illness severity assessment might be the guiding factor for antibiotic treatment choice. Misinterpretation of the child’s illness severity might not only lead to over-consultation but also to over-treatment. Therefore, in-depth knowledge of the differences in the factors that determine illness severity assessment and particularly (dis)agreement between parents and clinician’s assessments may be important in improving parent-clinician communication and management of children with RTI in primary care.

Our study aimed to: (1) investigate (dis) agreement between clinicians’ and parents’ illness severity scores; and (2) identify and compare the determinants of high parent and clinician severity scores in children presenting to primary care with cough and RTI.

**Methods**

**Design and study population**

Data were used from the ‘TARGET’ study (11), a multicentre prospective cohort study of children with acute (≤28 days) cough and RTI recruited between July 2011 and May 2013. The design of this study has been described in detail and the main results have been published.(11, 12)

Briefly, family physicians and prescribing nurse practitioners (hereafter referred to as ‘clinicians’) working in primary care centres (GP practice, Walk-in centres, GP Out of Hours centres or polyclinics) were recruited and trained by four UK hubs (Bristol, London, Oxford, and Southampton). They recruited children to the study if they were eligible, defined as aged between 3 months and 16 years and presented with the main symptom of acute (≤28 days) cough with other RTI symptoms (such as fever and coryza). Children with an infected exacerbation of asthma and those who were severely unwell (e.g. requiring same day hospital assessment or admission) were included.

Children were excluded if they presented with a non-infective exacerbation of asthma, were at high risk of serious infection (immunocompromised, for example with cystic fibrosis), required a throat swab for clinical management (which were taken for research purposes in a subgroup of children), had been previously recruited to the study, recently participated in other research, or had temporarily registered at the practice.

**Measurements**

After obtaining informed consent, clinicians completed a structured online (or paper) case report form (CRF) (appendix 1). The form recorded eight sociodemographic and four illness history/trajectory items, 33 parent-reported symptoms (including whether mild, moderate, or severe in the previous 24 hours), 14 physical examination signs (including vital signs), and the prescription of antibiotics (immediate or delayed) or referral to secondary care for acute assessment. Children’s diagnosis of current asthma was checked via a medical notes review, which was deemed present if asthma was noted in the medical notes and asthma medication was issued in the previous 12 months.

The Indices of Multiple Deprivation (IMD) (Source: National Statistics / Ordnance Survey Extracts) ranges from around zero (least deprived) to >90 (most deprived). Postcodes were used for the IMD.

Illness severity was measured independently by parents and clinician using a zero to ten visual-analogue scale(13), with zero being ‘well’ and ten being ‘very unwell’.

**Treatment of variables**

*Illness severity scores*

In the main analysis children were coded as having ‘high parent reported illness severity’ if their parent severity score was in the upper quartile (a score of ≥7). Similarly, ‘high clinician reported illness severity’ was coded if clinician score was in the upper quartile (a score of ≥4). For sub-analysis three Illness severity groups were used to determine agreement between parent and clinician scores: (1) parents score < clinician score (‘less ill’); (2) parents score = clinician score; and (3) parent score > clinician score (‘more ill’).

*Symptom severity*

For symptom severity (mild, moderate, severe) in the 24 hours prior to consultation the dichotomy for each variable was split depending on the prevalence, to either ‘severe’ if at least 5% of the whole cohort fell into this category or ‘moderate and severe’ if the proportion was smaller than this.

*Clinical cut-offs*

In case of dichotomising, commonly used clinical cut-offs were used for continuous data where possible (e.g. temperature >37.8°C) and age-related heart rate, respiratory rate and oxygen saturations were coded as raised or normal/low according to the Paediatric Life Support Manual, 2005.(14)

**Data analysis**

We compared parent and clinician illness severity scores using a non-parametric approach (as Figure 1 sows the clinician score data was heavily skewed' so a non-parametric approach that makes no assumption about distribution was chosen) for unpaired data, using medians and inter-quartile ranges [IQR] and the Mann-Whitney test to investigate differences between groups (lower/equal/higher).

To determine the association between the number of symptoms and signs reported on the global scores, correlation was determined between the parental illness severity score and the number of parent-reported symptoms, and between the clinician illness severity score and the number of symptoms and signs (both separately and added together). Correlation and agreement were calculated using Spearman’s rho and Kappa statistics respectively. To control for anchoring, the rounded mean difference between the parental score and the clinician score was added to the clinician score and the Kappa recalculated.

We used univariable and multivariable logistic regression to identify the sociodemographic and clinical factors independently associated with parent and clinician reported illness severity. Multivariable models were derived through several iterations using backward stepwise logistic regression including all variables significant in the univariable analyses (p<0.01) where missing data was <1%. We controlled for age, gender and ethnicity, as well as the presence of other symptoms or signs already in the model.

Two separate multivariable models were determined for high clinician score, one including demographic and symptoms (for comparison with the parental model) and one including demographics, symptoms and signs. The second clinician model included a combined variable for parent-reported severe fever OR clinician measured temperature ≥37.8°C, and combined variable for parent-reported breathing quickly OR clinician measured raised respiratory rate.

The final models were built in the following order: (1) sociodemographic, (2) clinical history, (3) parents reports symptoms and (for clinician model two) (4) clinical examination.

To reduce the problem of multiple comparisons in the analysis, the listed symptoms (CRF/Appendix 1) were only tested in terms of whether the symptom was present during the illness and whether the symptom was moderate or severe in the last 24 hours.

**Results**

*Descriptive statistics*

We recruited 8394 children to the study across the four centres. There was no difference in parental illness severity score between the 164 children declining participation (median 3, IQR 2-4) and the final recruited sample (median 3, IQR 2-4).

Children’s median age was 3 years [IQR: 1-6 years] with 1392 (16.6%) under one year old (Table 1). 52% were boys, 78% were white, mother’s median age at the child’s birth was 30 and 18% of recruited children’s mothers were current smokers. Families’ median deprivation score was 16.7 (IQR 8.8-29.5). Ethnicity, deprivation score and prevalence of maternal smoking were similar to national figures.(15, 16) The median illness duration prior to consultation was five days [IQR 3-10 days, range 0-28 days] and the median number of parent-reported symptoms was 7 (range 1-16, IQR 5-9) prior to consultation and 6 (range 0-16, IQR 4-8) within 24 hours of the consultation.

*Illness severity*

8360 (99.6%) children had complete parent and clinician illness severity score data. Parent severity scores (mean 5.2 (SD= 1.8), median=5, range 0-10, IQR 4-7) were higher (p<0.0001) than clinician scores (mean 3.1 (SD= 1.7), median 3, range 0-9, IQR 2-4, Figure 1, Table 2). Parents only scored illness severity lower than clinicians in 6% of the children, the same in 15% and higher in 79% (Table 2).

We found evidence of a low positive correlation (Spearman’s r: 0.43, p=<0.001) and poor inter-rater agreement between the parental- and clinician illness severity scores (Kappa 0.049, Figure 2), the parental scores and the number of parent-reported symptoms (Spearman’s r: 0.37), and the clinician scores and the number of parent-reported symptoms (Spearman’s r: 0.34). The clinician scores and the number of clinical signs are strongly positive correlated (Spearman’s r: 0.71) and moderate positive correlation was found between clinician scores and the number of parent-reported symptoms plus the number of clinical signs (Spearman’s r: 0.60). Anchor controlling for the inter-rater agreement did not improve the Kappa (0.064, Figure 2).

*Univariable analyses*

All but one variable (oxygen saturation) had <1% data missing. There was no evidence of differences between children with high parent illness scores compared to those with low scores regarding age, gender, ethnicity, number of consultations in the previous 12 months and current asthma. There was no evidence of differences between children with high clinician illness scores compared to those with low scores regarding ethnicity, number of consultations in the previous 12 months and parent-reported moderate-to-severe diarrhoea. Table 3 summarises the sociodemographic and clinical factors that were associated with higher and lower parent and clinician scores.

*Multivariable analyses*

Parent severity scores

The model included 8208/8394 (97.8 %) of the children in the cohort. Eight predictors were strongly associated with parent severity scores at p<0.001: one demographic variable (two or more children at home (OR:1.28, 95%CI:1.15-1.44)), and seven parent-reported symptoms ((n order of importance: severe fever (OR:2.58, 95%CI:2.12-3.13), severe dry cough (OR:1.93, 95%CI:1.60-12.34), breathing quickly (OR:1.88, 95%CI:1.69-2.10), severe reduction in eating (OR:1.26, 95%CI:1.26-1.98), moderate to severe reduced fluid intake (OR:1.55, 95%CI:1.34-1.80), severe disturbed sleep (OR:1.32, 95%CI:1.14-1.52) and change in cry (OR:1.30, 95%CI:1.13-1.49)). Together these gave an area under the ROC curve (AUROC) of 0.68 (95% C.I. 0.66-0.69, Table 4).

Clinician severity scores

Two multivariable regression models were derived, one model to compare to the parental model including demographics and parent reported symptoms only (model one), and a second model including demographics, parent reported symptoms and clinical signs (model two). Both models included 8198/8394 (97.7%) of the children in the cohort. Model one identified four predictors with p<0.001: illness deterioration recently before consultation (OR:2.11, 95%CI:1.89-2.34) and three parent reported symptoms (in order of importance: severe fever (OR:3.04, 95%CI:2.52-3.67), breathing quickly (OR:1.78 95%CI:1.61-1.98) and change in cry (OR:1.29 95%CI:1.14-1.46), ), with a combined AUROC of 0.67 (95% C.I. 0.66-0.68, Table 4). Adding clinical examination did not change model one factors included and identified six additional signs (in order of importance: inter/subcostal recession (OR:4.91 95%CI:3.54-6.82), crackles/crepitation (OR:4.79 95%CI:4.18-5.50), nasal flaring (OR:3.08 95%CI:1.76-5.41), wheeze (OR:2.31 95%CI:1.98-2.68), irritability/drowsiness (OR:1.91 95%CI:1.22-2.99), and pallor (OR:0.58 95%CI:0.49-0.70)). Together these gave an AUROC of 0.79 (95% C.I.0.78-0.80, Table 4). As clinicians were only able to take the oxygen saturation level for just over half the children recruited (n= 4194, 51%) a multivariable model was derived using this smaller number of children. This sensitivity analysis did not substantially change the final models.

**Discussion**

*Main findings*

Parent and clinician global severity illness assessment differed, with parents considering their child more severely unwell than clinicians. Factors associated with illness severity also differed between parents and clinicians, with parents relying on symptoms and clinician’s physical examination findings. That said, the symptoms ‘severe fever’ and ‘breathing quickly’ were both important for both parents and clinicians.

**Strengths and limitations**

We used data from a well characterised, large, representative cohort of children presenting to primary care with the most common problem managed by health services. Participating children had similar levels of overall illness severity than those who were invited but declined study participation. Baseline characteristics were pragmatic, measured according to routine clinical practice and with high level of completeness. The main outcome is clinically relevant, and we used a stringent model retention criterion (p<0.01) because of the many candidate predictors. The study question described in this paper was not the focus of the cohort study but a secondary hypothesis.

The main limitation is use of the visual analogue scale (VAS) to measure illness severity. VAS provide a simple technique for measuring subjective experience and has been established as valid and reliable in a range of clinical and research applications.(13) Although ease of use is frequently cited as a major advantage over other scales, others point out that it requires the ability to transform a complex assessment into a vision-spatial display, involving perceptual judgement and accuracy.(17) We used a VAS anchored by zero (well) and ten (very unwell). During decision making, anchoring occurs when individuals use an initial piece of information to make subsequent judgments. Once an anchor is set, other judgments are made by adjusting away from that anchor. Anchor points of parents and clinicians might differ as clinicians are probably used to seeing many more ill children (NB: none of the clinicians gave a 10/10 (very unwell)), which might lead to a lower score of illness severity in general. Therefore, the poor agreement we found between parents and clinicians might be partly due to anchoring. However, agreement remained poor when we adjusted for changes in anchoring, and we found different parent and clinician factors associated with high severity scores.

Information on any prior assessment (e.g. GP triage, NHS 111 triage, pharmacist recommendation) which might have influenced parent’s decision to consult a GP has not been included in our analyses, this, however, could have consequently influenced the parent’s illness severity assessment.

Another limitation is that we were not able comparing like with like in that the parents did not clinically examine the child and might not bet aware of signs such as ‘inter/subcostal recession’, although parents are taught to look out for signs of respiratory distress by GPs and 111 in general . Therefore, modelling the drivers of severity scores was always going to give different results. However, this reflects the actual situation parents and clinicians must navigate in.

**omparison with existing literature**

The poor agreement in parental and clinician illness of severity assessment found in the present study is consistent with the results of a previous systematic review of qualitative evidence on the interaction of primary care consultations for children with acute minor illnesses.(18) This review concluded that common misunderstandings occur when parents and clinicians speak about the ‘seriousness’ of the illness, with parents and clinicians talking at cross purposes: parents are seeking to justify their decision to consult, while family physicians seek to justify non-antibiotic treatment strategies.

Although our analysis showed that clinicians do consider parent reported symptoms, they seem to rely much on the clinical examination. Our exploratory analysis on the correlation between the number of symptoms and signs reported and the illness of severity scores showed that the number of parent-reported symptoms does not influence the parental score, but that the number of clinical signs does influence the clinician score. We therefore think that the type of parent-reported symptom is probably more important than the number, but that for clinicians it could be ‘the more clinical signs the more they worry’ (meaning a higher score). This is consistent with qualitative evidence (5) showing that GPs not only rely on the initial assessment but feel the need for a more deductive assessment, including physical examination to refine their diagnosis and to rule out serious illness. A study of Blacklock et al. on which symptoms and clinical features correctly identify serious respiratory infection in children attending a paediatric assessment unit shows that parent-reported symptoms were unreliable discriminators of serious acute respiratory infection in children. Nurse trial assessment of respiratory distress and some vital signs are found to be important predictors, which is in line with our findings.(19)

The main analyses from this cohort study found parent reported fever, and clinician measured inter- or sub-costal recession, as well as wheeze on auscultation are prognostically significant – these factors have predictive utility in identifying children at risk of future hospitalisation resulting from their RTI.(12) These factors are also included as intermediate/red flags in the NICE Traffic light system for identifying risk of serious illness (Fever (moderate and high), inter/sub-costal recession (high) and wheeze (moderate)) (20)

It is possible that parents have intuitively identified these factors as important, and therefore also use them to assess illness severity.

No explanation was found in literature that could explain the counterintuitive findings of pallor and one child in the home being associated with lower illness severity scores. Contrary to our finding, pallor is one of the red flag symptoms in NICE Traffic light system for identifying risk of serious illness in children with feverish illness.

**Implications for policy, teaching and clinical practice**

Interventions to promote self-care (one of the corner stones of NHS sustainability strategy) should take account of the parent concerns likely to be the main drivers of parent help seeking. For example, some symptoms of importance to parents (‘severe dry cough’, ‘reduced eating’ and ‘disturbed sleep’) have not shown to be predictive of poor outcome (12) and may be important targets for reassurance- e.g.odds of a parent perceiving illness to be severe is 14% to 52% higher for parents where the child's sleep has been disturbed but disturbed sleep was not a predictor of disease illness for clinicians.

Moreover, how to identify and address these concerns should be included in primary care clinician (medical and nursing) training programmes, as recommended by the recently published Health Education England ‘Tackling Antimicrobial Resistance’ strategy document (21).

Our study showed that fever is associated with both high parent and clinician severity scores. Education on fever management for parents on the one side and clinicians on the other side is key to make better informed decisions as to when (or if) to consult and to improve parent-clinician communication. In some cases, parent concerns regarding fever are the result of lack of experience and knowledge about fever.(22, 23) Empowering parents and teaching them about alarm symptoms could ameliorate illness anxiety and possibly improve use of primary care services.(24) Advice on self-management for parents and carers could be provided to help patients to self-manage the fever (25). However, as severe fever in a primary care setting is also associated with serious infection(26), parents might be correct to be concerned. Therefore, fever and fever management should be an important item in (NICE) guidelines and telephone triage protocols (NHS 111) on acute respiratory tract infections.

*Conclusions*

Clinicians need to recognise that parents reach different conclusions, using different factors, when making judgements about illness severity. Improved understanding of the factors concerning parents could improve parent-clinician communication, improve the quality of consultations and thereby child health outcomes. Understanding parents’ concerns and educating them about clinician’s concerns need to be part of day to day practice. Balancing the two is essential to successful management of patients with minor infection. Parent education should continue to be an important element of clinical care.

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**Ethical approval: body giving ethics approval with reference number where appropriate**

The study was approved by the South West Central Bristol Research Ethics Committee, UK (reference number 10/H0102/54)

**Competing interests**

All other authors declare no competing interests.

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**Table 1. Characteristics of children and parents (Total N=8394)**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **N** | **Median** | **IQR** |
| Age child (years) | 8394 | 3 | 1-6 |
| Number of children in home | 8355 | 2 | 1-2 |
| Family deprivation score\* | 8201 | 16.7 | 8.8-29.5 |
| Illness duration at recruitment (days) | 8390 | 5 | 3-10 |
| Number of parent reported symptoms prior to consultation\*\* | 8320 | 7 | 5-9 |
| Number of parent reported symptoms within 24 hours of consultation\*\* | 8229 | 6 | 4-8 |
|  | **N** | **Percentage** | |
| Child’s gender (male) | 8934 | 51.6 | |
| Child’s ethnicity (white) | 8349 | 78.4 | |
| Mother current smoker | 8285 | 18.0 | |
| Breastfeeding at 3 months | 7784 | 44.2 | |
| Illness got worse recently | 8383 | 66.0 | |

*\*The indices for multiple deprivation (IMD) score ranges from around zero (least deprived) to >90 (most deprived).*

*Ethnicity, deprivation score and prevalence of maternal smoking were similar to national figures.12 13*

*\*\* Median of positively reported parent-reported symptoms from the CRF out of the possible 20.*

**Table 2. Parent and clinician illness severity scores**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Severity of illness score** | **Mean\* (SD)** | **Median** | **Range** | **IQR** |
| Parent *(8360/8394, 99.6%)* | 5.2 (1.8) | 5 | 0-10 | 4-7 |
| Clinician *(8368/8394, 99.7%)* | 3.1 (1.7) | 3 | 0-9 | 2-4 |
|  | **N**  (total N= 8337\*\*) | **%** |  | |
| Parental score **<** clinician score | 478 | 5.7 |
| Parental score **=** clinician score | 1236 | 14.8 |
| Parental score **>** clinician score | 6623 | 79.4 |

*\*P<0.0001 (Wilcoxon test score); \*\* Number of children with parent and clinician score*

**Table 3. Univariable associations (p<0.01) with parental and clinician illness severity score**

|  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Significant Variables** | **Parents high severity score (≥ 7)\*** | | **Parents normal & low severity score (< 7)** | | **p-value** | **Clinician high severity score (≥ 4)\*** | | **Clinician normal & low severity score (< 4)** | | **p-value** |
| **N** | **%** | **N** | **%** |  | **N** | **%** | **N** | **%** |  |
| **Sociodemographics and past medical history** | | | | | |  |  |  |  |  |
| *Male* | † |  |  |  |  | 1578/2938 | 53.7 | 1735/5422 | 50.4 | <0.01 |
| *Two or more children at home* | 1,493/2134 | 70.0 | 4004/6196 | 64.6 | <0.001 | 2035/2925 | 69.6 | 3457/5397 | 64.1 | <0.001 |
| *Illness deteriorated recently before consultation* | 1742/2143 | 81.3 | 3778/6216 | 61.8 | <0.001 | 2300/2936 | 78.3 | 3214/5414 | 59.4 | <0.001 |
| *Current asthma\*\** |  |  |  |  |  | 308/2938 | 10.5 | 437/5422 | 8.1 | <0.001 |
| **Parent/carer-reported general symptoms** | | | | | |  |  |  |  |  |
| *Change in cry* | 498/2132 | 23.4 | 880/6207 | 14.2 | <0.001 | *634/2930* | *21.6* | *742/5403* | *13.7* | *<0.001* |
| *Breathing quickly* | 1085/2143 | 50.6 | 1885/6223 | 30.3 | <0.001 | *1419/2938* | *48.3* | *1549/5419* | *28.6* | *<0.001* |
| *Wheezing/ whistling in the chest* | 991/2143 | 46.3 | 2287/6220 | 36.8 | <0.001 | *1395/2937* | *47.5* | *1879/5418* | *34.7* | *<0.001* |
| *Vomiting including after a cough* | 738/2143 | 34.4 | 1603/6233 | 25.8 | <0.001 | *933/2938* | *31.8* | *1407/5419* | *26.0* | *<0.001* |
| **Parent/carer-reported symptoms (in the previous 24 hours)** | | | | | |  |  |  |  |  |
| *Severe dry cough* | 256/2137 | 12.0 | 302/6202 | 4.9 | <0.001 | 242/2931 | 8.3 | 319/5401 | 5.9 | <0.001 |
| *Severe fever* | 296/2134 | 13.9 | 243/6204 | 3.9 | <0.001 | 347/2922 | 11.9 | 193/5406 | 3.6 | <0.001 |
| *Severe disturbed sleep* | 539/2133 | 25.3 | 806/6193 | 13.0 | <0.001 | 625/2923 | 21.4 | 722/5394 | 13.4 | <0.001 |
| *Severe reduction in eating* | 217/2131 | 10.2 | 210/6201 | 3.4 | <0.001 | 252/2918 | 8.6 | 177/5405 | 3.3 | <0.001 |
| *Breathing quickly* | 712/2137 | 33.3 | 916/6216 | 14.7 | <0.001 | 852/2934 | 29.0 | 777/5410 | 14.4 | <0.001 |
| *Moderate-to-severe wheezing in chest* | 581/2141 | 27.1 | 1033/6211 | 16.6 | <0.001 | 784/2931 | 26.8 | 827/5412 | 15.3 | <0.001 |
| *Moderate-to-severe diarrhoea* | 126/2143 | 5.9 | 216/6216 | 3.5 | <0.001 |  |  |  |  |  |
| *Moderate-to-severe vomiting (incl. after a cough)* | 316/2138 | 14.8 | 521/6219 | 8.4 | <0.001 | 340/2934 | 11.6 | 496/5414 | 9.2 | <0.001 |
| *Moderate-to-severe reduced fluid intake* | 467/2133 | 21.9 | 688/6212 | 11.1 | <0.001 | 551/2924 | 18.8 | 601/5412 | 11.1 | <0.001 |
| *Moderate-to-severe reduction in urine passed* | 202/2132 | 9.5 | 260/6206 | 4.2 | <0.001 | 261/2931 | 8.9 | 203/5401 | 3.8 | <0.001 |
| **General clinical examination** | | | | | |  | | | | |
| *Pallor* | ‡ |  | ‡ |  |  | 331/2934 | 11.3 | 489/5416 | 9.0 | <0.001 |
| *Grunting* |  |  |  |  |  | 59/2933 | 2.0 | 16/5416 | 0.3 | <0.001 |
| *Nasal flaring* |  |  |  |  |  | 79/2934 | 2.7 | 22/5417 | 0.4 | <0.001 |
| *Temperature ≥ 37.8°C* |  |  |  |  |  | 690/2929 | 23.6 | 351/5409 | 6.5 | <0.001 |
| *Raised respiratory rate (age-related cut-offs)* |  |  |  |  |  | 681/2932 | 23.3 | 560/5395 | 10.4 | <0.001 |
| *Low oxygen saturation (≤95%)* |  |  |  |  |  | 234/1378 | 16.9 | 167/2804 | 6.0 | <0.001 |
| *irritable or drowsy* |  |  |  |  |  | 88/2936 | 3.0 | 29/5413 | 0.7 | <0.001 |
| *Clinician gut feeling that something was wrong* |  |  |  |  |  | 1213/2931 | 41.4 | 483/5413 | 8.9 | <0.001 |
| *Stridor* |  |  |  |  |  | 7/933 | 0.92 | 15/54145 | 0.8 | <0.001 |
| *Inflamed pharynx/tonsils* |  |  |  |  |  | 1096/928 | 37.4 | 187/5403 | 3.8 | <0.001 |
| **Chest examination** | | | | | |  | | | | |
| *Wheeze* | ‡ |  | ‡ |  |  | 776/2931 | 26.5 | 454/5416 | 8.4 | <0.001 |
| *Crackles/crepitations* |  |  |  |  |  | 1140/2932 | 38.9 | 452/5415 | 8.4 | <0.001 |
| *Raised respiratory rate (age-related cut-offs)* |  |  |  |  |  | 681/2932 | 23.3 | 560/5395 | 10.4 | <0.001 |
| *Inter/subcostal recession* |  |  |  |  |  | 350/2934 | 11.9 | 53/5416 | 1.0 | <0.001 |

*\* based on upper quartile of total dataset (N=8394); \*\*defined as present if asthma in medical notes and asthma drug issued in the previous 12 months;* † *not significantly associated;* ‡ p*arents were not expected to clinically examine their child.*

**Table 4. Final multivariable predictors of high parental illness of severity score and high clinician illness of severity score (all p<0.01)**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | | **Illness of severity score** | | | | | | |
|  | | **High parent score (≥ 7)**  (‡AUROC= 0.68,  95% CI= 0.66-0.69 ) | |  | **High clinician score (≥ 4)**  (‡AUROC= 0.67,  95% CI= 0.66-0.68) | | **High clinician score (≥ 4)**  (‡AUROC= 0.79,  95% CI=0.79-0.80 ) | |
|  | | **Model 1**  (Demographics & symptoms) | |  | **Model 1**  (Demographics & symptoms) | | **Model 2**  **(**Demographics, symptoms & clinical signs) | |
|  | **Source** | **Odds ratio** | **95% CI** |  | **Odds ratio** | **95% CI** | **Odds ratio** | **95% CI** |
| **Sociodemographic** | | | | | | | | |
| *2 or more children at home* | Parent | 1.28 | 1.15-1.44 |  | † |  | † |  |
| **Clinical history** | | | | | | | | |
| *Illness deteriorated recently before consultation* | Parent | † |  |  | 2.11 | 1.89-2.34 | 1.90 | 1.69-2.13 |
| **Symptoms** | | | | | | | | |
| *Change in cry* | Parent | 1.30 | 1.13-1.49 |  | 1.29 | 1.14-1.46 | 1.39 | 1.21-1.59 |
| *Breathing quickly/ raised respiratory rate\** | Parent/clinician | 1.88 | 1.69-2.10 |  | 1.78 | 1.61-1.98 | 1.43 | 1.28-1.59 |
| *Severe fever / temperature ≥ 37.8°C* | Parent/clinician | 2.58 | 2.12-3.13 |  | 3.04 | 2.52-3.67 | 3.58 | 3.12-4.10 |
| *Severe dry cough* | Parent | 1.93 | 1.60-2.34 |  | † |  | † |  |
| *Severe disturbed sleep* | Parent | 1.32 | 1.14-1.52 |  | † |  | † |  |
| *Severe reduced eating* | Parent | 1.58 | 1.26-1.98 |  | † |  | † |  |
| *Moderate-to-severe reduced fluid intake* | Parent | 1.55 | 1.34-1.80 |  | † |  | † |  |
| **Clinical examination** | | | | | | | | |
| *Pallor* | Clinician |  |  |  |  | | 0.58 | 0.49-0.70 |
| *Irritable or drowsy* | Clinician |  |  |  |  | | 1.91 | 1.22-2.99 |
| *Nasal flaring* | Clinician |  |  |  |  | | 3.08 | 1.76-5.41 |
| *Inter/subcostal recession* | Clinician |  |  |  |  | | 4.91 | 3.54-6.82 |
| *Wheeze* | Clinician |  |  |  |  | | 2.31 | 1.98-2.68 |
| *Crackles/crepitations* | Clinician |  |  |  |  | | 4.79 | 4.18-5.50 |

*\*age-related cut-offs:* † *p≥0.01;* ‡ *AUROC: Area Under the Receiver Operating Characteristic curve.*

**Figure 1. Distribution of illness severity scores**

***Figure 2. Comparing illness severity score of child between clinician and parent***

|  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| Parent-reported illness severity | Clinician-reported illness severity | | | | | | | | | | |  |
| 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | Total |
| 0 | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% |
| 1 | 0% | 1% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 2% |
| 2 | 0% | 3% | 3% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 6% |
| 3 | 0% | 3% | 4% | 3% | 1% | 0% | 0% | 0% | 0% | 0% | 0% | 12% |
| 4 | 0% | 3% | 5% | 4% | 2% | 1% | 0% | 0% | 0% | 0% | 0% | 15% |
| 5 | 0% | 3% | 5% | 5% | 4% | 3% | 1% | 0% | 0% | 0% | 0% | 22% |
| 6 | 0% | 1% | 4% | 4% | 4% | 3% | 2% | 0% | 0% | 0% | 0% | 18% |
| 7 | 0% | 1% | 3% | 3% | 3% | 2% | 2% | 1% | 0% | 0% | 0% | 15% |
| 8 | 0% | 1% | 1% | 2% | 1% | 1% | 1% | 1% | 0% | 0% | 0% | 8% |
| 9 | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 2% |
| 10 | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% | 0% |
| Total | 2% | 17% | 25% | 22% | 14% | 10% | 7% | 3% | 1% | 0% | 0% | 100% |
| *Percentages rounded to the nearest whole number/ ‘white’ representing 0% to the ‘darkest blue’ representing 5% of children got a similar score by parent as clinician.* | | | | | | | | | | | | |
| **Inter rater agreement:**  Parental score – Clinician score : 14.8%, Kappa= 0.049  Parental score – Clinician score + 2\* : 20.6%, Kappa= 0.064  *\*to control for anchoring, the rounded mean difference between the parental score and clinician score was added to the clinician score* | | | | | | | | | | | | |