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Improving the Management of Children with Bronchiolitis The Updated American Academy of Pediatrics Clinical Practice Guideline

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**Improving the management of children with bronchiolitis:
the updated American College of Pediatrics clinical practice
guideline**

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9 **Editorial**

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11 **Improving the management of children with bronchiolitis: the updated American**
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13 **Academy of Pediatrics clinical practice guideline**
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49 **Conflict of interest statements:** AC and SS have no financial conflicts. AC has undertaken
50 studies relating to the management of bronchiolitis and is a panel member for several
51 guidelines unrelated to bronchiolitis.
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3 Bronchiolitis is the most common cause of hospitalisation of very young children.¹ Although
4 the mortality associated with bronchiolitis is low in affluent countries, the related cost and
5 morbidity are high² especially in some minority groups such as Indigenous children.³ Despite
6 the burden of illness, there is a relative paucity of evidence-based guidelines on the
7 management of bronchiolitis in- and out- of hospitals. Prior to the updated bronchiolitis
8 guideline from the American Academy of Pediatrics (AAP),¹ the latest most comprehensive
9 collation of evidence for the management of bronchiolitis was that from the Scottish
10 Intercollegiate Guidelines Network (SIGN) guideline from the United Kingdom,² where the
11 latest search date was 2005. Thus, the updated AAP bronchiolitis guideline,¹ which has
12 incorporated relevant data from the last 10 years, is a document that should be welcomed by
13 clinicians worldwide.

24 25 26 27 28 29 What are the changes?

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31 There are many new and clinically important changes in the updated guideline;¹ only four (#1,
32 7, 8, 11) of the 14 recommendations have minimal or no changes from the 2006 guideline.⁴
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34 The rest of the recommendations are new or amended, reflecting the evidence obtained from
35 newer studies. Statements referring to the use of ribavirin and complementary/alternative
36 medicines, present in the previous guideline,⁴ have been omitted in the updated guideline.¹ All
37 but four (#6, 9, 12b and 14) recommendations have 'B' rated evidence (studies with minor
38 limitations; consistent findings from multiple observational studies) or higher.¹ The most
39 important cost-saving change is arguably the recommendation (#10a) against the use of
40 palivizumab prophylaxis for all healthy children born prematurely (≥ 29 weeks gestation).^{1,5}
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42 Not surprisingly, this was followed by 'big pharma' remonstrations
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44 (<http://bigstory.ap.org/article/virus-drugmaker-fights-pediatricians-new-advice>).

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3 Other important changes in the updated guideline¹ include: when to consider the use of
4 hypertonic saline (#4); the definitive statement not to use short acting beta₂ agonists (#2) or
5 epinephrine (#3) and; the alternate use of nasogastric fluids to intravenous fluids (#9). The
6 technical report is currently in preparation¹ will contain further information not reflected in
7 the recommendations, such as data relating to nasal suctioning and the use of high flow nasal
8 and home therapy oxygen.
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18 Like the 2006 AAP bronchiolitis guideline,⁴ important dissimilarities between the updated
19 2014 guideline¹ and SIGN guideline² are: when supplementary oxygen should be considered
20 (AAP: SpO₂ <90%;^{1,4} SIGN: SpO₂ ≤92%²), the use of continuous SpO₂ monitoring (AAP: not
21 routinely recommended;^{1,4} SIGN: recommended for 8-12 hours post cessation of O₂²); and
22 virology testing (AAP: recommends “laboratory testing should not be obtained”;^{1,4} SIGN:
23 rapid virological testing recommended for cohorting children²).
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34 Why use guidelines?

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36 Why should clinicians take note of, or use, clinical practice guidelines? The previous AAP
37 bronchiolitis guideline⁴ has been credited with a significant reduction in the use of diagnostic
38 and therapeutic resources⁶ in the USA, although variation in bronchiolitis management
39 persists even after almost 8 years of implementation.⁷ High quality and well-implemented
40 guidelines can reduce variance in clinical care, reduce cost⁶ and most importantly, improve
41 clinical outcomes.⁸ However the quality of guidelines are highly variable,⁸ including those
42 endorsed by academic societies,⁹ and perhaps unsurprisingly so, guidelines are not universally
43 popular. Furthermore, poorly developed guidelines, such as those overseen by panel members
44 with close relationships with ‘big pharma’, propagate discontent about guidelines⁹ and create
45 management dilemmas for doctors and possibly be harmful to patients.^{8,9} Good clinical
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3 guidelines are transparent, derived from a rigorous process, externally reviewed and
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5 disseminate “the most scientifically sound healthcare practice”⁸ undertaken by a
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7 multidisciplinary panel whose members are free of financial conflicts.^{8,10} The updated AAP
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9 guideline¹ has a multidisciplinary panel (including a methodologist, parent/consumer, family
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11 physician, neonatologist, general pediatricians, pediatric pulmonologists, pediatric
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13 hospitalists, emergency physicians, intensivist and infectious disease physicians) that has no
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15 financial ties to pharmaceutical companies and underwent an extensive peer review process
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17 (including review by American Academy of Family Physicians, the American College of
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19 Chest Physicians, the American Thoracic Society, and the American College of Emergency
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21 Physicians). The panel of the updated AAP guideline¹ thus responds to the eight items of
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23 guideline panel review outlined by Lenzer and colleagues.⁸
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29 The future

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31 The updated guideline¹ lists ideas for future research, highlighting the incompleteness of the
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33 guideline that relates to insufficient existing data. Other important clinical questions include:
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35 issues reflecting the differences in the AAP^{1,4} and the SIGN guidelines² described above; the
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37 diagnosis and management of possible co-existence of pneumonia in settings with a high
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39 incidence of pneumonia related deaths; post-discharge monitoring particularly for at-risk
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41 children and; post-bronchiolitis syndrome and its management.¹¹
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47 Guidelines should never represent ‘cookbook medicine’ and are not a substitute for
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49 individualised high quality clinical care as individuals, families and settings are heterogenous,
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51 necessitating individualised nuances and deviations in selected circumstances. However,
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53 when implemented well, the contribution of untainted high quality guidelines to improved
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55 clinical outcomes is undisputed.^{8,10} The field of guideline development and implementation
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3 has undergone substantial changes¹⁰ since defined in the 1990s as “systematically developed
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5 statements to assist practitioner and patient decisions about appropriate healthcare for specific
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7 clinical circumstances”.¹² The high standard of the 2014 AAP guideline is important in our
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9 current era and advances the field of guidelines and the management of bronchiolitis as
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11 described above. It is thus highly likely that the updated guideline¹ along with its
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13 accompanying technical report and that on the guidance for the use of palivizumab⁵ will be
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15 welcomed by clinicians worldwide. The value of the guideline in improving clinical
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17 outcomes and use of resources is now dependent on its implementation.
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