Molecular Epidemiology of Human Rhinovirus Infections in Kilifi, Coastal Kenya

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This study reports pediatric surveillance over 3 years for human rhinovirus (HRV) at the District Hospital of Kilifi, coastal Kenya. Nasopharyngeal samples were collected from children presenting at outpatient clinic with no signs of acute respiratory infection, or with signs of upper respiratory tract infection, and from children admitted to the hospital with lower respiratory tract infection. Samples were screened by real-time reverse transcriptase polymerase chain reaction (real-time RT-PCR) and classified further to species by nucleotide sequencing of the VP4/VP2 junction. Of 441 HRV positives by real-time RT-PCR, 332 were classified to species, with 47% (155) being HRV-A, 5% (18) HRV-B, and 48% (159) HRV-C. There was no clear seasonal pattern of occurrence for any species. The species were present in similar proportions in the inpatient and outpatient sample sets, and no significant association between species distribution and the severity of lower respiratory tract infection in the inpatients could be determined. HRV sequence analysis revealed multiple but separate clusters in circulation particularly for HRV-A and HRV-C. Most HRV-C clusters were distinct from reference sequences downloaded from GenBank. In contrast, most HRV-A and HRV-B sequences clustered with either known serotypes or strains from elsewhere within Africa and other regions of the world. This first molecular epidemiological study of HRV in the region defines species distribution in accord with reports from elsewhere in the world, shows considerable strain diversity and does not identify an association between any species and disease severity. J. Med. Virol. 84:823–831, 2012. © 2012 Wiley Periodicals, Inc.

KEY WORDS: acute respiratory illness; rhinovirus; RT-PCR; VP4/VP2 protein; genetic sequencing

INTRODUCTION

Human rhinoviruses (HRVs), members of the Picornaviridae family [Whitton et al., 2005], are recognized as the most frequent viral agents in humans presenting with symptoms of the common cold [Denny, 1995]. The profile of HRVs has been raised as a result of widening use of sensitive molecular methods of detection compared with in vitro cultivation [Kammerer et al., 1994; Arruda et al., 1997; Pitkaranta et al., 1997; Andeweg et al., 1999; Vesa et al., 2001; Renwick et al., 2007], with an increase in the observations of association with lower respiratory tract infection [Lamson et al., 2006; Lau et al., 2007, 2009; Khetsuriani et al., 2008; McErlean et al., 2008; Linsuwanon et al., 2009; Piralla et al., 2009]. A number of other studies have suggested that respiratory illness, presenting with wheezing, rales and respiratory distress may be associated with HRVs [Camara et al., 2004; Cheuk et al., 2007; Singh et al., 2007; Miller et al., 2007b, 2009].

Sensitive RT-PCR and sequencing has been used to type rhinoviruses as HRV-A, HRV-B, and HRV-C, with HRV-C being discovered as recently as 2006 [Lau et al., 2007; McErlean et al., 2007]. These PCR methods have also enabled studies on HRV disease burden in hospitalized infants and children under 5 years old [Midulla et al., 2010; Miller et al., 2009, 2007b], including sub-Saharan Africa [Niang et al., 2010; O’Callaghan-Gordo et al., 2011; Venter et al., Abstract]
2011]. Molecular studies suggest almost equal prevalence of HRV-A and HRV-C with similar global distribution patterns [reviewed in Simmonds et al., 2010]. Although most of the epidemiological studies on HRV-C have focused on hospitalized or asthmatic children, the clinical outcomes of HRV-C infection, which was initially controversial, shows association with severe disease including asthma and acute wheezing [McErlean et al., 2007; Renwick et al., 2007; Miller et al., 2009; Peltola et al., 2009; Piralla et al., 2009; Smuts et al., 2011].

HRVs are non-enveloped viruses with an icosahedral capsid enclosing a single-stranded, positive sense RNA genome that is translated in the cytoplasm [Rueckert, 1996]. The viral polyprotein is divided into P1, P2, and P3 regions, with the P1 region encoding the capsid proteins VP4, VP2, VP3, and VP1, while the P2 and P3 regions encode non-structural proteins 2APro, 2B, 2C, 3A, 3B (VPg), 2CPro and 3DPol [reviewed by Kirchberger et al., 2007]. Sequencing of the VP1 led to the classification of HRVs into three species (HRV-A-C) [Ledford et al., 2004] and recently, the VP1 led to the classification of HRVs into three species (HRV-A-C) [Wisdom et al., 2009] were used at a concentration of 1 μM each per reaction in a total volume of 25 μL. PCR products were purified using Qiagen PCR purification Kit and sequenced with both the outer and inner primers in a BigDye chain terminator reaction and analyzed in an ABI 3130xl instrument.

**Materials and Methods**

**Patients and Samples**

The study was undertaken with samples from children presenting to the Kilifi District Hospital, which is located on the coast of Kenya approximately 60 km from Mombasa city. The District comprises a largely rural population of subsistence farmers and experiences an equatorial climate with rains predominantly falling in the months of April to July and November to December. Further details of the study area and respiratory virus disease surveillance through Kilifi District Hospital can be found in previous reports [Nokes et al., 2009; Berkley et al., 2010]. Nasopharyngeal wash or aspirate residues collected from 1912 children aged 1 day—12 years between January 2007 and December 2008, and stored in viral transport medium at −80°C, were tested for the presence of HRV. The children were outpatients (254) presenting either for routine immunization and categorized as without respiratory infection or with signs and symptoms of upper respiratory tract infection [Berkley et al., 2010], or inpatients (1,759) with signs and symptoms consistent with lower respiratory tract infection and categorized as either mild, severe or very severe following WHO syndromic criteria [Nokes et al., 2008]. Written informed consent was obtained from the parent/guardian. This study was approved by the Kenya National Ethical Review Committee and the University of Warwick Biomedical Research Ethics Sub-Committee.

**RNA Extraction and PCR**

RNA was extracted from HRV positive samples using the Qiagen Viral RNA miniprep kit (Qiagen, West Sussex, UK) following the manufacturer's instructions. Diagnostic PCR to confirm the presence of HRV genetic material was performed using a real-time RT-PCR with primers and probe targeting the 5’ untranslated region [Hammitt et al., 2011]. For VP4/VP2 amplification, one tube RT-PCR was performed using Qiagen OneStep RT-PCR kit. Primers amplifying VP4/VP2 [Wisdom et al., 2009] were used at a concentration of 1 μM each per reaction in a total volume of 25 μL. PCR products were purified using Qiagen PCR purification Kit and sequenced with both the outer and inner primers in a BigDye chain terminator reaction and analyzed in an ABI 3130xl instrument.

**Sequence Analysis and Phylogenetics**

The raw sequences were edited using BioEdit software, translated in Lasergene (DNASTar) and phylogenetic analysis performed with MEGA 5.01 software [Tamura et al., 2011]. The evolutionary history was inferred using Neighbor-Joining method [Saitou and Nei, 1987] using 1,000 bootstrap replicates [Felsenstein, 1985]. Evolutionary distances were computed using the Kimura 2-parameter [Kimura, 1980]. HRV typing of the samples was determined by comparison with reference sequences from GenBank encoding VP4/VP2 protein. Intra-species patristic distances (p-distances) were tested by Neighbor-Joining method using 2,000 bootstrap replicates in MEGA 5.01 software, together with sequences downloaded from the global knowledge base of these viruses.
from http://www.picornaviridae.com/news.htm (Tables SI and SII). Strains as used in this study refer to those reference sequences identified genetically without supporting serotypic information. Samples were considered to form a monophyletic grouping if they had a bootstrap value ≥ 90%. Statistical analyses were done in Stata version 11.0 (StataCorp LP, College Station, TX, www.stata.com) and graphs drawn in GraphPad Prism version 5.01 for Windows (GraphPad Software, San Diego, CA, www.graphpad.com”). Fisher’s exact test was used to test for independence of two categorical variables, while Spearman’s correlation was used to test the associations between HRV and climatic conditions.

RESULTS

Diagnostic real-time PCR identified 23% (441/1,912) of samples as HRV positive. Of these 86% (380/441) were from inpatient children with lower respiratory tract infection, while 14% (61/441) were from outpatient children. The median age (IQR) for non-acute respiratory infection, upper respiratory tract infection, mild lower respiratory tract infection, severe lower respiratory tract infection, and very severe lower respiratory tract infection children was 3.4 months (m) (3.2–11.8 m), 10.4 m (2.0–16.6 m), 18.8 m (9.5–26.9 m), 9.1 m (3.2–19.0 m), and 7.1 m (1.7–23.3 m) respectively. In general, the prevalence of HRV in inpatients was 22% (380/1,759) and 24% (61/254) in outpatients.

Distribution of HRV Species

The VP4/VP2 region of the genome was amplified from the 441 HRV positive samples with 75% (332/441) success. While 25% (109) samples in total were unclassified to species, the proportions differed significantly between outpatients and inpatients (44% (27/61) vs. 22% (82/380)), as shown in Table I (P < 0.001). In the un-classified category, 90% (98/109) failed to amplify and 10% (11) had poor sequence readout. Of the 98 samples that failed to amplify, 65% (69) had late cycle threshold (Ct) values (above 30 cycles), within the real-time diagnostic assay. The distribution of HRV in 298 classified inpatient samples was 47% (141) HRV-A, 4.4% (13) HRV-B, and 48% (144) HRV-C; this distribution did not differ significantly by lower respiratory tract infection severity (P = 0.175). In the outpatients, of 34 classified viruses, the proportion of HRV-A was 41%, HRV-B 15%, HRV-C 44% and the distribution did not differ significantly between children with and without upper respiratory tract infection (P = 0.221).

Seasonal Distribution of the HRV Species

Cases of HRV were identified in Kilifi District Hospital throughout the year over the 3-year study period with higher proportions being seen in the months February 2007, June–July in 2007, August–September 2008, February 2009, April–June 2009, and November 2009, while lower proportions were observed between October–November 2007 and November–December 2008 (Fig. 1).

The distribution of the HRV species and the proportion of samples positive, in the inpatients, over the 3-year period is shown in Figure 1. HRV-A and C co-circulated in most months of surveillance, and while either was rarely absent (e.g., in the months of November 2008–January 2009), the numbers fluctuated considerably on an irregular basis. In contrast, HRV-B occurred only rarely and sporadically throughout the 3 years. Peak admissions of HRV-A and HRV-C cases did not obviously segregate or alternate and were not regularly seasonal. Numbers of HRV-C cases were higher in July 2007, and again in September 2008; while HRV-A cases were higher in July 2007, February and June 2008. There was no clear association between the monthly number of HRV cases and mean relative humidity (r² = 0.1507, P = 0.380) or daily mean rainfall (r² = −0.1399, P = 0.416) over the period of this study (Fig. 2).

Relatedness of Kilifi Sequences to Other HRV Sequences

The Kilifi sequences for each of the species formed numerous unique clusters with mean nucleotide identity of 76.4% (range 52.2%–97.6%) for HRV-A, 74.2% (range 44.5–99.7%) for HRV-B, and 68% (range 51.0–98.9%) for HRV-C, suggesting considerable diversity in strains/serotypes present in the population. In the cases of HRV-A and HRV-B most sequences clustered with known serotypes or strains from around the world (Figs. 3 and 4). In contrast, many of the sequences of HRV-C identified in circulation in Kilifi (Fig. 5) did not cluster with any published strains that also

| Category | HRV-A | HRV-B | HRV-C | Un-typed | Total |
|----------|-------|-------|-------|----------|-------|
| Non-ARI  | 8 (34.78) | 1 (4.35) | 4 (17.39) | 10 (43.48) | 23 |
| URTI     | 6 (15.79) | 4 (10.53) | 11 (28.95) | 17 (44.74) | 38 |
| Mild     | 18 (43.9) | 2 (4.88) | 15 (36.59) | 6 (14.63) | 41 |
| Severe   | 96 (34.91) | 7 (2.55) | 111 (40.36) | 61 (22.18) | 275 |
| V. Severe| 27 (42.19) | 4 (6.25) | 18 (28.12) | 15 (23.44) | 64 |
| Total    | 155 (35.15) | 18 (4.08) | 159 (36.05) | 109 (24.72) | 441 |

URTI, upper respiratory tract infection; ARI, acute respiratory illness; un-typed, unclassified.

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are based on sequencing of the VP4/VP2 junction. The Kilifi sequences (most easily seen for HRV-A and HRV-B) showed reasonable genetic deviation from nearest known strains/serotypes, generally with mean nucleotide differences of 25% (range 15.1–48.0%) for all the three HRV species.

One sequence (KI-5607), that seemed an offshoot in HRV-B (Fig. 4) was found to be human enterovirus 68 when compared with other sequences from the GenBank. The KI-5607 had 95% nucleotide identity to strain JPOC10-396 isolated in Osaka Japan with GenBank accession number AB601884.

DISCUSSION

The distribution of HRV species in sub-Saharan Africa is not well described although there are some reports on the overall prevalence of HRV. Recently, epidemiological studies in sub-Saharan Africa showed the prevalence of HRV in children <5 years presenting to a primary care center with acute respiratory infection as 17% in rural Senegal [Niang et al., 2010], 41% in rural Mozambique [O’Callaghan-Gordo et al., 2011], and 33% in Pretoria South Africa [Venter et al., 2011]. The equivalent prevalence in inpatients...
The current study reports the species typing of 441 Kilifi HRV positive samples by the genetic analysis of the VP4/VP2 junction. Failure to type 25% of HRV positive samples was largely attributed to PCR failure (65%) due to low viral load (inferred from Ct value) than the severity category. However, it cannot be excluded that certain genetic variants were missed in the typing assay due to their variability in the primer-annealing sites, leading to PCR failure.

All the three species of HRV were found to be circulating in children under 12 years of age in Kilifi, with occurrence of 47% HRV-A, 5% HRV-B, and 48% HRV-C. Such patterns of species distribution have been reported elsewhere in the USA [Miller et al., 2011] and Cape Town, South Africa [Smuts et al., 2011].

The current study also identified a human enterovirus 68, which was diagnosed initially as HRV by real-time PCR assay. This discrepancy in diagnosis was resolved after BLAST search correctly identified the virus. This was also supported by its unique topology on the HRV-B phylogenetic tree (Fig. 4). The identification of non-HRV is not unique as the PCR primers used the Kilifi HRV diagnostic platform are

Fig. 3. A phylogenetic analysis of HRV-A sequences from Kilifi and reference sequences from Genbank. Sequences with closed circles denote known serotypes, while those with open circles denote previously described strains. Bootstrap values are (2,000 replicates) shown on the branches, with values <90% omitted from the tree. The tree was drawn to scale, with branch lengths in the same units (p-distance) used to infer the phylogenetic tree. Kilifi sequences are designated K-number (KI for inpatient sequences and KO for outpatient sequences).
capable of amplifying other non-HRV human enteroviruses [Wisdom et al., 2009].

There was neither a clear seasonal circulation pattern nor an identifiable influence of weather (rainfall and humidity) on HRV occurrence in Kilifi. This is in contrast to a study from Germany, which showed an association between relative humidity and rhinovirus infection in a hospitalized cohort [du Prel et al., 2009].

This study also noted lack of an association between inpatient severity and any of the HRV species. Other studies elsewhere have shown evidence of the involvement of HRV-C in infants and toddlers with wheeze, and asthma in older children [Lau et al., 2007; Khetsuriani et al., 2008; Miller et al., 2009] and a shorter duration of asthma symptoms and less cough in HRV-A infected children [Arden and Mackay, 2010]. The discrepancy in associating HRV-C with disease severity in the current study may be as a result of either the broad case definition used hence masking the broad definitions of non-acute respiratory

Fig. 4. A phylogenetic analysis of HRV-B sequences from Kilifi alongside reference serotypes and strains from Genbank. Sequences with closed circles denote known serotypes, while those with open circles denote previously described strains. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (2,000 replicates) is shown next to the branches. Bootstrap values <90% have been omitted from the tree. The tree is drawn to scale, with branch lengths in the same units (p-distance) used to infer the phylogenetic tree. Kilifi sequences are designated K-number (KI for inpatient sequences and KO for outpatient sequences).
infection and lower respiratory tract infection or as a result of the emergence of HRV-C strains with altered disease causing potential.

This study did not attempt to compare inpatient and outpatient cases to identify HRV disease severity by species due to the bias in design. It is therefore important that studies in African settings focus on case–control studies as well as collecting adequate clinical history to help decipher the clinical relevance of HRVs.

There was high diversity of sequences in circulation in all the Kilifi HRV species, although sequences of HRV-A and HRV-B clustered with known serotypes and strains. However there were a large number of clusters for HRV-C that had no closely associated strain from the literature. The formation of few clusters between the known HRV-C and Kilifi sequences compared with HRV-A and HRV-B was mostly likely due to the availability of few HRV-C VP4/VP2 sequences in the GenBank.

**ACKNOWLEDGMENTS**

Our sincere thanks to the clinical and laboratory staff for their hard work in collection and processing of the specimens. Special thanks to Dr. Jay Berkley Fig. 5. A phylogenetic analysis of HRV-C sequences from Kilifi alongside reference serotypes and strains from Genbank. Sequences with closed circles denote known strains. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (2,000 replicates) is shown next to the branches. Bootstrap values <90% have been omitted from the tree. The tree is drawn to scale, with branch lengths in the same units (p-distance) used to infer the phylogenetic tree. Kilifi sequences are designated K-number (KI for inpatient sequences and KO for outpatient sequences).
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REFERENCES

Andeweg AC, Bestebroer TM, Huybrechs M, Kimman TG, de Jong JC. 1999. Improved detection of rhinoviruses in clinical samples by using a successfully developed reverse transcription-PCR assay. J Clin Microbiol 37:594–530.

Arends KE, Mackay IM. 2010. Newly identified respiratory viruses in children with non-hospitalised asthma exacerbation. J Med Virol 82:1458–1461.

Arruda E, Pitkaranta A, Witek TJ, Jr., Doyle CA, Hayden FG. 1997. Frequency and natural history of rhinovirus infections in adults during summer. J Clin Microbiol 35:2864–2868.

Berkley JA, Munywoki P,Maga, Kanzugu S, Abwao J, Bett A, Lasauzierre N, Kresfelder T, Cane PA, Venter M, Scott JA, Nokes DJ. 2010. Viral etiology of severe pneumonia among Kenyan infants and children. JAMA 303:2021–2027.

Camara AA, Silva JM, Ferriani VP, Tobias KR, Macedo IS, Padovani MA, Harsi CM, Cardoso MR, Chapman MD, Arruda E, Platts-Mills TA, Arruda LK. 2004. Risk factors for wheezing in a tropical environment: Role of respiratory viruses and allergen sensitization. J Allergy Clin Immunol 113:551–557.

Cheuk DK, Tang IW, Chan KH, Woo PC, Peiris MJ, Chiu SS. 2007. Rhinovirus infection in hospitalized children in Hong Kong: A prospective study. Pediatr Infect Dis J 26:995–1000.

Denny FW, Jr. 1985. The clinical impact of human respiratory virus infections. Annu Respir Crit Care Med 152:524–512.

du Preb JF, Puppe W, Grondahl B, Knuf M, Weigl JA, Schaaff F, Erdman DD. 2008. Novel human rhinoviruses and exacerbation of respiratory wheezing in children. Emerg Infect Dis 10:1095–1101.

Felsenstein J. 1985. Confidence limits on phylogenies: An approach using the bootstrap. Evolution 39:783–791.

Gwaltney JM, Jr., Hendley JO, Simon G, Jordan WS, Jr. 1966. Rhinovirus infections in an industrial population. I. The occurrence of illness. N Engl J Med 275:1261–1268.

Hantunen JL, Welch S, Betts A, Orton C, Gunson R, Scott J, Nokes D. 2011. Added value of an oropharyngeal swab in the detection and rapid identification of human picornaviruses. J Clin Microbiol 49:861–867.

Mills TA, Arruda LK, Scagnolari C, Bonci E, Pierangeli A, Antonelli G, Arruda LB, Scagnolari C, Bonci E, Pierangeli A, Antonelli G, Midulla F, Cane PA, Medley GF. 2008. Distinguishing respiratory viruses and clinical characteristics of a putative new rhinovirus species, human rhinovirus C (HRV C), PLoS ONE 3:e1847.

Miller EK, Williams JV, Shehabi AA, Faouri S, Al Jundi I, Chen Q, Heil L, Mohamed Y, Morin LL, Ali A, Halasa NB. 2009. Human rhinovirus C associated with wheezing in hospitalized children in the Middle East. J Clin Virol 46:85–89.

Miller EK, Krusi-Bulos N, Williams JV, Shehabi AA, Faouri S, Al Jundi I, Chen Q, Heil L, Mohamed Y, Morin LL, Ali A, Halasa NB. 2009. Human rhinovirus C associated with wheezing in hospitalized children in the Middle East. J Clin Virol 46:85–89.

Mills TA, Arruda LK, Scagnolari C, Bonci E, Pierangeli A, Antonelli G, Midulla F, Cane PA, Medley GF. 2008. Distinguishing respiratory viruses and clinical characteristics of a putative new rhinovirus species, human rhinovirus C (HRV C), PLoS ONE 3:e1847.

Nang MN, Diop OM, Sarr FD, Goudiaby D, Malou-Sompy H, Niayi K, Vahret A, Baril L. 2010. Viral etiology of respiratory infections in children under 5 years old living in tropical rural areas of Senegal: The EVIRA project. J Med Virol 82:866–872.

Nokes DJ, Okiro EA, Ngama M, Ochola R, White LJ, Scott PD, English M, Cane PA, Medley GF. 2008. Respiratory syncytial virus infection and disease in infants and young children observed from birth in Kilifi District, Kenya. Clin Infect Dis 46:50–57.

Nokes DJ, Ngama M, Bett A, Abwao J, Munywoki P, English M, Scott JA, Cane PA, Medley GF. 2009. Incidence and severity of respiratory syncytial virus pneumonia in rural Kenyan children identified through hospital surveillance. Clin Infect Dis 49:1341–1349.

O’Callaghan-Gordo C, Bassat Q, Morais L, Díez-Padrisa N, Machado S, Nhampossa T, Nhalungo D, Sanz S, Quinto L, Alonso PL, Roca A. 2011. Etiology and epidemiology of viral pneumonia among hospitalized children in rural Mozambique: A malaria endemic area with high prevalence of human immunodeficiency virus. Pediatr Infect Dis J 30:39–44.

Peltola V, Ruuskanen O. 2008. Editorial commentary: Respiratory viral infections in developing countries: Common, severe, and unrecognized. Clin Infect Dis 46:85–89.

Peltola V, Jartti T, Putto-Laurila A, Mertesdal J, Vainionpaav J, Waris M, Hyytiä T, Ruuskanen O. 2009. Rhinovirus infections in children: A retrospective and prospective hospital-based study. J Med Virol 81:1831–1838.

Lamson D, Renwick N, Kapoor V, Liu Z, Palacios G, Ju J, Dean A, St George K, Briese T, Lipkin WI. 2006. MassTag polymerase-chain-reaction detection of respiratory pathogens, including a new rhinovirus genotype, that caused influenza-like illness in New York State during 2004–2005. J Infect Dis 194:1398–1402.

Lau SK, Yip CS, Tsoi HW, Lee RA, So LY, Lau YL, Chan KH, Woo PC, Yuen KY. 2007. Clinical features and complete genome characterization of a distinct human rhinovirus (HRV) genetic clade, probably representing a previously undetected HRV species, HRV-C, associated with acute respiratory illness in children. J Clin Microbiol 45:3655–3664.

Lau SK, Yip CS, Lin AW, Lee RA, So LY, Lau YL, Chan KH, Woo PC, Yuen KY. 2009. Clinical and molecular epidemiology of human rhinovirus C in children and adults in Hong Kong reveals a possible distinct human rhinovirus C subgroup. J Infect Dis 200:1096–1103.

Ledford RM, Patel NR, Demencuzuk TM, Watanyar A, Herbert T, Gofft MS, Peverar DC. 2004. VP1 sequencing of all human rhinovirus serotypes: Insights into genus phylogeny and susceptibility to antiviral capsid-binding compounds. J Virol 78:3665–3674.

Onyango et al.
Piralla A, Rovida F, Campanini G, Rognoni V, Marchi A, Locatelli F, Gerna G. 2009. Clinical severity and molecular typing of human rhinovirus C strains during a fall outbreak affecting hospitalized patients. J Clin Virol 45:311–317.

Piralla A, Baldanti F, Gerna G. 2011. Phylogenetic patterns of human respiratory picornavirus species, including the newly identified group C rhinoviruses, during a 1-year surveillance of a hospitalized patient population in Italy. J Clin Microbiol 49:373–376.

Pitkaranta A, Arruda E, Malmberg H, Hayden FG. 1997. Detection of rhinovirus in sinus brushings of patients with acute community-acquired sinusitis by reverse transcription-PCR. J Clin Microbiol 35:1791–1793.

Renwick N, Schweiger B, Kapoor V, Liu Z, Villari J, Bullmann R, Miething R, Briese T, Lipkin WI. 2007. A recently identified rhinovirus genotype is associated with severe respiratory-tract infection in children in Germany. J Infect Dis 196:1754–1760.

Rueckert R. 1996. Picornaviridae: The virus and their replication. In: Fields BN, Knipe DM, Howley PM, editors. Fundamental virology, 3rd edition. Philadelphia: Lippincott-Raven. pp 477–522.

Saitou N, Nei M. 1987. The neighbor-joining method: A new method for reconstructing phylogenetic trees. Mol Biol Evol 4:406–425.

Simmonds P, McIntyre C, Savolainen-Kopra C, Tapparel C, Mackay IM, Hovi T. 2010. Proposals for the classification of human rhinovirus species C into genotypically assigned types. J Gen Virol 91:2409–2419.

Singh AM, Moore PE, Gern JE, Lemanske RF, Jr., Hartert TV. 2007. Bronchiolitis to asthma: A review and call for studies of gene-virus interactions in asthma causation. Am J Respir Crit Care Med 175:108–119.

Smuts HE, Workman L, Zar HJ. 2011. Human rhinovirus infection in young African children with acute wheezing. BMC Infect Dis 11:65.

Tamura K, Peterson D, Peterson N, Stecher G, Nei M, Kumar S. 2011. MEGA5: Molecular evolutionary genetics analysis using maximum likelihood, evolutionary distance, and maximum parsimony methods. Mol Biol Evol 28:2731–2739.

Venter M, Lassauniere R, Kresfelder TL, Westerberg Y, Visser A. 2011. Contribution of common and recently described respiratory viruses to annual hospitalizations in children in South Africa. J Med Virol 83:1458–1468.

Vesa S, Kleemola M, Blomqvist S, Takala A, Kilpi T, Hovi T. 2001. Epidemiology of documented viral respiratory infections and acute otitis media in a cohort of children followed from two to twenty-four months of age. Pediatr Infect Dis J 20:574–581.

Whitton JL, Cornell CT, Fuhr R. 2005. Host and virus determinants of picornavirus pathogenesis and tropism. Nat Rev Microbiol 3:765–776.

Wisdom A, Leitch EC, Gaunt E, Harvala H, Simmonds P. 2009. Screening respiratory samples for detection of human rhinoviruses (HRVs) and enteroviruses: Comprehensive VP4-VP2 typing reveals high incidence and genetic diversity of HRV species C. J Clin Microbiol 47:3958–3967.