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European molecular epidemiology and strain diversity of feline calicivirus


Feline calicivirus (FCV) causes a variable syndrome of upper respiratory tract disease, mouth ulcers and lameness. A convenience-based prospective sample of oropharyngeal swabs (n=426) was obtained from five countries (France, Germany, Greece, Portugal and the UK). The prevalence of FCV by virus isolation was 22.2 per cent. Multivariable analysis found that animals presenting with lymphoplasmacytic gingivitis stomatitis complex were more likely to test positive for FCV infection. Furthermore, vaccinated cats up to 48 months of age were significantly less likely to be infected with FCV than unvaccinated animals of similar ages. Phylogenetic analysis based on consensus sequences for the immunodominant region of the capsid gene from 72 FCV isolates identified 46 strains. Thirteen of the 14 strains with more than one sequence were restricted to individual regions or sites in individual countries; the exception was a strain present in two sites close to each other in France. Four strains were present in more than one household. Five colonies, four of which were rescue shelters, had multiple strains within them. Polymerase sequence suggested possible rare recombination events. These locally, nationally and internationally diverse FCV populations maintain a continuous challenge to the control of FCV infection and disease.
level, a highly complex genetic landscape can exist with many 10 s of strains co-circulating, with only rare cases of transmission between practices at the national (UK) level (Coyne and others 2012).

The primary aim of this study was to see whether the population structure of viruses previously seen in the UK is mirrored when data from multiple countries is included.

Materials and methods
Sample collection
Samples were collected from five countries: France (five sites in the Paris area), UK (two sites: Bangor (Northern Ireland) and Northampton), Portugal (two sites: Mem Martins and Lisbon area), Greece (one site: Thessaloniki) and Germany (three sites: Lippstadt, Berlin and Achim). At each site, a private veterinary practitioner was recruited based on convenience by the project sponsor, and asked to sample two types of cats; healthy cats not vaccinated against FCV for at least 1 month (to reduce the probability of isolating vaccine strains), and those cats with clinical signs typical of FCV infection (for example corza or LGSC).

For each cat whose owner gave consent, a plain cotton-tipped swab was used to collect an oropharyngeal saliva sample and coughed up in 2 ml viral transport medium (40 per cent L15/Mac Coy’s (v/v) cell culture media, 60 per cent freezing excipient). These were stored locally at −18°C then shipped as a batch to the laboratory on dry ice. With each sample a short questionnaire was completed by the attending veterinary surgeon and owner, capturing demographic data, disease and vaccination history.

The non-invasive nature of this project and anonymised data collection meant that formal ethical committee approval was not necessary. However, informed owner consent was a stipulation before samples were collected.

Virus isolation
Diagnosis of FCV and feline herpesvirus type 1 (FeHV-1) was by demonstrating typical cytopathic effect on confluent monolayers of feline embryo A cells. Samples were only reported negative following a second negative passage. Positive isolates were stored at −80°C (Knowles and others 1990).

Statistical analysis
A parametrical bootstrap approach was used to test country and sites as random effects based on a hypothesis of the form H0: σ2=0. The estimate P value was 0.99 for country and 0.09 for sites. Thus, the level of between-country and between-site variability was not considered sufficient to incorporate random effects in a multivariable model.

A multivariable logistic regression analysis was conducted to model the probability of being infected with FCV. The model was fitted in R (V3.01) using the ‘glm’ function from ‘lme4’ package. Risk factors considered within the model included the country and site within the country where the samples were collected, sex, whether a cat lived outdoors, neutering status, FCV vaccination status, age, and presence of the following clinical signs: ocular discharge, sneezing, gingivitis, nasal discharge, coughing, oral ulcers, conjunctivitis, stomatitis and LGSC. Visual inspection of the data indicated a curvilinear relationship before samples were collected.

The probability of being infected with FCV. The model was fitted in R (V3.01) using the ‘glm’ function from ‘lme4’ package. Risk factors considered within the model included the country and site within the country where the samples were collected, sex, whether a cat lived outdoors, neutering status, FCV vaccination status, age, and presence of the following clinical signs: ocular discharge, sneezing, gingivitis, nasal discharge, coughing, oral ulcers, conjunctivitis, stomatitis and LGSC. Visual inspection of the data indicated a curvilinear relationship before samples were collected.

The multivariable model was only available for 299 samples. The number of cats living in the same household (NCH) was only available for 66 per cent of the observations in the multivariable analysis. Therefore, to avoid reducing the numbers included in the multivariable model, the impact of NCH on FCV infection was only assessed by univariable logistic regression.

RNA extraction, reverse transcription-PCR
RNAs were extracted from positive samples (second passage or less) (QIamp viral RNA Mini Kit; Qiagen). For every three samples, one negative control was included from uninfected cell cultures. Reverse transcription was performed using 200 ng random hexamers (Superscript III, Invitrogen), according to manufacturer’s guidelines.

Capsid amplification: A 529-nucleotide region of the capsid gene, equivalent to residues 6406-6934 of FCV strain F9 (Carter and others 1992) and incorporating immunodominant regions C and E (Seal and others 1993, Radford and others 1999b), was amplified as previously described (Coyne and others 2007b, 2012). Briefly, each 50 μl reaction contained 2 μl cDNA, 45 μl 1.1×Reddy mix (Thermo Scientific), 1 μl nuclease-free water and 3.2 ng each of forward and reverse primers (Table 1). In early experiments, primers M13cap2F/T7cap2R were most cross-reactive and so all PCR products were initially conducted with these: only samples testing negative were subsequently attempted with the additional primers.

Thermal cycling consisted of DNA denaturation (95°C, 2 min), followed by 40 cycles of denaturation (95°C, 30 s), primer annealing (45–55°C, 60 s) and primer extension (72°C, 90 s). A final extension was performed at 72°C (5 min).

Polymerase amplification: A 486-nucleotide region corresponding to the 5’ end of the FCV POL region was amplified from the same cDNA template using Reddy-Mix (A/βgene) according to the manufacturers’ instructions, in 50 μl reactions containing 100 ng each of the primers M13-53D and T7-33D (Coyne and others 2006b) (Table 1). Thermal cycling consisted of denaturation (95°C, 4 minutes), followed by 40 cycles of denaturation (95°C, 60 s), primer annealing (55°C, 60 s) and primer extension (72°C, 3 minutes). A final extension was performed at 72°C (5 minutes).

Nucleotide sequence and phylogenetic analysis
Amplicons were purified (QIAquick PCR purification, Qiagen), quantified (Genequant) and sequenced bidirectionally (Source Bioscience). The authors have previously found this method to be >99 per cent reproducible (Coyne and others 2007b). Forward and reverse sequences were aligned and manually corrected (Chromas Pro, Technelysium). Pairwise p-distances between sequences, and Kimura 2-parameter Neighbour-joining trees. A 486-nucleotide region corresponding to the 5’ end of the FCV POL region was amplified from the same cDNA template using Reddy-Mix (A/βgene) according to the manufacturers’ instructions, in 50 μl reactions containing 100 ng each of the primers M13-53D and T7-33D (Coyne and others 2006b) (Table 1). Thermal cycling consisted of denaturation (95°C, 4 minutes), followed by 40 cycles of denaturation (95°C, 60 s), primer annealing (55°C, 60 s) and primer extension (72°C, 3 minutes). A final extension was performed at 72°C (5 minutes).

Results
Study sample
A total of 426 samples were collected from 13 sites in five countries. For 17 samples, the viral status could not be assessed due to bacterial overgrowth. For the remaining 409, FCV and FeHV-1 were isolated from 91 (22.2 per cent) and 18 (4.4 per cent), respectively (Table 2). For FCV, 16.2 per cent and 34.2 per cent of healthy and sick (at least one clinical sign) cats tested positive for FCV, respectively. For FeHV-1, the figures were 2.6 per cent and 8.0 per cent.

Risk factors for FCV infection
Complete information about all the variables considered within the multivariable model was only available for 299 samples. The
final model included the continuous variables age and a quadratic term for age (age²), as well as the categorical variables vaccination status against FCV (FCV-V) and whether or not the animal was presenting with LGSC, and the two-way interactions FCV-V by age and FCV-V by age² (Table 3). Other individual clinical signs were not found to be significant in this population.

Cats presenting with LGSC were 9.33 (95% CI 3.18 to 29.45) times more likely to present with FCV infection than cats without LGSC. The relationship between the probability of FCV infection and age and vaccination status was more complex due to an interaction between these risk factors. In short, in unvaccinated animals (n=119) there was a reduction in risk each month for the first 8 years, followed by a plateau for the next 2.5 years, an increase every month for the next 1.5 years and a decrease every month for the next 10 years (Fig 2). In vaccinated animals (n=218), there was an increase in risk each month for the first 8.5 years, followed by a plateau for the next 1.5 years and a decrease every month for the next 12.5 years (Fig 2). The 95% CIs of the probability for FCV infection in vaccinated and unvaccinated animals of similar ages. From 48 months of age, the probability of being FCV infected was lower than in the small sample size, especially for older animals, reflected by wide CIs in the predicted probability of FCV infection for those animals (Fig 3).

Visual inspection of the data indicated a linear relationship between the percentage of cats testing positive for FCV and NCH (Fig 1b). The number of cats living in a household was found to be significantly associated with the probability of being FCV infected on the univariable analysis. Thus, for one-unit increase in NCH, the odds of being FCV infected increase 1.5% (OR=1.015, 95% CI 1.004 to 1.034, p=0.043).

Sequence analysis
In total, 72 consensus sequences of the major immunodominant region of the FCV capsid were obtained from the 91 FCV isolates (Table 2). This included 71 clear consensus sequences containing a small number (<5 per cent) of ambiguous nucleotide positions typical for a calicivirus (Fig 3a). In contrast, one sample (FR1_11), contained many ambiguities strongly suggesting a mixed infection (Fig 3b). One of the two sequences in this sample was present in a clear majority allowing FR1_11 sequence to be manually ‘split’ into its predicted major and minor sequences, as well as its consensus (average). Whilst the accuracy of these three individual sequences is likely to be lower, the authors included them to gain further insight into the diversity of viruses circulating in this population.

Two additional samples (FR1_37 and FR1_40) gave very clear sequence, and were also identified as mixed infections. However, unlike FR1_11, the two sequences were present in approximately equal proportions making it impossible to manually resolve them (Fig 3c). These sequences were not available for further analyses and would require a cloning approach to separate them.

The final capsid phylogeny (Fig 4a) included 74 usable sequences (71 consensuses, three for FR1_11). In total, 46 strains (pairwise genetic distance >20 per cent) were identified; the number of strains in each country and site ranged from 5 to 12 and 1 to 9, respectively (Table 2). There was no evidence of wide-scale clustering at the geographical level; rather strains from each country were dispersed throughout the tree.

Of the 46 strains identified, 32 were represented by single sequences, the remaining 14 each being represented by more than 1 isolate and supported by bootstrap values of >75 per cent (A–N on Fig 4a). The most numerous of these was strain A (9 samples and 11 sequences, including 3 for FR1_11). Of these 14 strains, 13 were geographically restricted to single sites, the exception being strain M, found in two sites in France (FR1_43 and FR4_02). Eight of these strains had low levels of diversity (<5 per cent) suggestive of relatively acute or recent infection. The remaining six strains (A, C, D, G, M and N) showed greater diversity (5–20 per cent); these high levels of strain diversity have been found previously in endemically infected households

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**TABLE 1: Primers used for partial capsid and polymerase gene amplification**

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence (5’→3’)</th>
<th>Binding site (nt)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Capsid primers</td>
<td></td>
<td></td>
</tr>
<tr>
<td>M13cap2F</td>
<td>CAGGAAACAGCTATGACCCCTTTGTCTTCCARGCHAAYCG</td>
<td>6406-6428</td>
</tr>
<tr>
<td>T7cap2R</td>
<td>TAATGACCTCAATAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6934-6913</td>
</tr>
<tr>
<td>A1</td>
<td>CCGTCTGCTTGCAGGCAAGCGG</td>
<td>6406-6428</td>
</tr>
<tr>
<td>forward</td>
<td></td>
<td></td>
</tr>
<tr>
<td>A2</td>
<td>CCTGCCAATCCGATGACG</td>
<td>6934-6913</td>
</tr>
<tr>
<td>reverse</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P1</td>
<td>CGTTTGTGTTTCAAGAAACCG</td>
<td>6406-6428</td>
</tr>
<tr>
<td>forward</td>
<td></td>
<td></td>
</tr>
<tr>
<td>g2</td>
<td>CCTGCTACAATACCGATGAAACCG</td>
<td>6934-6913</td>
</tr>
<tr>
<td>reverse</td>
<td></td>
<td></td>
</tr>
<tr>
<td>M13cap1F</td>
<td>CAGGAAACAGCTACGCTGTTGNNITTCCYTCGCHAAYCG</td>
<td>6406-6428</td>
</tr>
<tr>
<td>T7cap1F</td>
<td>TAATGACCTCAATAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6934-6913</td>
</tr>
<tr>
<td>Polymerase primers</td>
<td></td>
<td></td>
</tr>
<tr>
<td>M13_53D</td>
<td>T7_33D</td>
<td></td>
</tr>
<tr>
<td>T7_33D</td>
<td>TAAAGACCTGCTTAAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6428</td>
</tr>
<tr>
<td>M13_53D</td>
<td>T7_33D</td>
<td></td>
</tr>
<tr>
<td>M13_53D</td>
<td>TAAAGACCTGCTTAAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6428</td>
</tr>
<tr>
<td>M13_53D</td>
<td>TAAAGACCTGCTTAAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6428</td>
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<tr>
<td>Polymerase primers</td>
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<td>M13_53D</td>
<td>T7_33D</td>
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<td>TAAAGACCTGCTTAAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6428</td>
</tr>
<tr>
<td>M13_53D</td>
<td>TAAAGACCTGCTTAAGGCGCCCTCACAAACCCCGTTCNCCCC</td>
<td>6428</td>
</tr>
</tbody>
</table>

Underlined regions illustrate the M13 forward and T7 reverse primer binding site sequences used for sequencing amplicons produced with these otherwise degenerate primers. Nucleotide sequences are shown using the single-letter IUB codes for degeneracy: R=A/G purine; Y=T/C pyrimidine; K=T/G. Nucleotide numbers of the binding sites relate to feline calicivirus (FCV) strain F9 (GenBank accession no. M86379) (Carter and others 1992).

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**FIG 1: Percentage of cats testing positive for feline calicivirus (FCV) by age—years (a) and by number of cats living in the same household (b)**

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TABLE 2: Summary of samples, isolates, amplification and strains identified in each country and site

<table>
<thead>
<tr>
<th>Country Site</th>
<th>France FR1</th>
<th>FR2</th>
<th>FR3</th>
<th>FR4</th>
<th>FR5</th>
<th>Total</th>
<th>UK UK1</th>
<th>UK2</th>
<th>Total</th>
<th>Portugal PT1</th>
<th>PT2</th>
<th>Total</th>
<th>Greece GE1</th>
<th>GE2</th>
<th>GE3</th>
<th>Total</th>
<th>Germany Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Samples (N)</td>
<td>41</td>
<td>7</td>
<td>33</td>
<td>7</td>
<td>14</td>
<td>102</td>
<td>49</td>
<td>34</td>
<td>83</td>
<td>48</td>
<td>32</td>
<td>80</td>
<td>42</td>
<td>28</td>
<td>34</td>
<td>102</td>
<td>409</td>
</tr>
<tr>
<td>FCV Positive, N (%)</td>
<td>14 (14)</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td>6</td>
<td>27</td>
<td>9</td>
<td>4</td>
<td>13</td>
<td>14</td>
<td>4</td>
<td>18</td>
<td>10</td>
<td>10</td>
<td>8</td>
<td>5</td>
<td>91</td>
</tr>
<tr>
<td>FCV Capsid PCR+</td>
<td>16</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td>2</td>
<td>5</td>
<td>25</td>
<td>6</td>
<td>2</td>
<td>14</td>
<td>3</td>
<td>17</td>
<td>9</td>
<td>8</td>
<td>4</td>
<td>3</td>
<td>74</td>
</tr>
<tr>
<td>FCV Capsid sequence Strains</td>
<td>5</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td>4</td>
<td>12</td>
<td>3</td>
<td>2</td>
<td>5</td>
<td>9</td>
<td>3</td>
<td>12</td>
<td>8</td>
<td>4</td>
<td>3</td>
<td>2</td>
<td>46</td>
</tr>
</tbody>
</table>

FCV, feline calicivirus

(Coyne and others 2007b). Four of the strains (D, J, L and M) were present in more than one household/colony (data not presented). Five of the households had more than one strain of virus (range two to six; Fig 4a).

None of the sequences obtained clustered with live vaccine strain FCV-F9 (as represented by a published sequence M86579) or with the inactivated vaccine strain FCV_255 (data not presented). The median uncorrected nucleotide distances between F9 and sequences obtained in this study was 39.2 per cent (SD 6.9). This was similar to that obtained when the sequences previously reported, possibly reflecting different sampling methodologies. In two previous UK studies, in which cats were not selected based on clinical signs, isolation rates of 10 per cent (Porter and others 2008) and 9–11 per cent (Coyne and others 2012) for FCV and <1 per cent for FeHV-1 were achieved. In contrast, in the study reported here, some of the 426 cats were recruited specifically based on disease, notably the presence of LGSC (N=26; 6 per cent). This inclusion rate is much higher than would be expected in the general population where the prevalence of LGSC is estimated to be 0.7 per cent (Healey and others 2007). Since previous studies have shown this condition to be strongly associated with FCV (Belgard and others 2010), with 80 per cent or more of cats testing positive (Knowles and others 1989), over-representation of such cases in the study population might be expected to increase the FCV prevalence over that found when cats were more randomly recruited (Radford and others 2007).

Discussion

Previously the authors have been reconstructing the molecular epidemiology of FCV at a local and national level in the UK. These studies have pointed to both high FCV prevalence and strain diversity, produced by a combination of rapid evolution and occasional geographical dispersal. Here the authors report for the first time how FCVs are dispersed internationally, showing that, subject to limitations associated with the authors’ sampling methodology, FCV prevalence remains high across Europe, with little evidence for wide geographical dispersal of individual strains.

The isolation rates reported here are somewhat higher than previously reported, possibly reflecting different sampling methodologies. In two previous UK studies, in which cats were not selected based on clinical signs, isolation rates of 10 per cent (Porter and others 2008) and 9–11 per cent (Coyne and others 2012) for FCV and <1 per cent for FeHV-1 were achieved. In contrast, in the study reported here, some of the 426 cats were recruited specifically based on disease, notably the presence of LGSC (N=26; 6 per cent). This inclusion rate is much higher than would be expected in the general population where the prevalence of LGSC is estimated to be 0.7 per cent (Healey and others 2007). Since previous studies have shown this condition to be strongly associated with FCV (Belgard and others 2010), with 80 per cent or more of cats testing positive (Knowles and others 1989), over-representation of such cases in the study population might be expected to increase the FCV prevalence over that found when cats were more randomly recruited (Radford and others 2007).

TABLE 3: Final multivariable logistic regression model of factors associated with FCV infection

<table>
<thead>
<tr>
<th>Variable</th>
<th>Beta</th>
<th>OR</th>
<th>95% CI</th>
<th>P value</th>
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<tr>
<td>LGSC</td>
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<td></td>
<td>Reference</td>
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</tr>
<tr>
<td>No</td>
<td>2.23</td>
<td>9.33</td>
<td>3.18 29.45</td>
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</tr>
<tr>
<td>Yes</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Vaccinated</td>
<td></td>
<td></td>
<td>Reference</td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>0.35</td>
<td>1.42</td>
<td>0.54 4.07</td>
<td>0.5</td>
</tr>
<tr>
<td>Yes</td>
<td>−0.014</td>
<td>0.99</td>
<td>0.98 1.00</td>
<td>0.003</td>
</tr>
<tr>
<td>Age centered (months)</td>
<td>0.0002</td>
<td>1.0002</td>
<td>1.0001 1.0003</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Age centered (months)</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
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<tr>
<td>Vaccinated, age centered</td>
<td>0.03</td>
<td>1.03</td>
<td>1.016 1.05</td>
<td>&lt;0.001</td>
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<tr>
<td>Not vaccinated, age centered</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Vaccinated, age centered</td>
<td>−0.0004</td>
<td>0.9996</td>
<td>0.9993 0.9998</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

*Age centered: mean age was 71.7 months

FCV, feline calicivirus; LGSC, lymphoplasmacytic gingivitis stomatitis complex

FIG 2: Predicted probability of a cat being infected with feline calicivirus by age and vaccination status. Black and grey lines depict the predicted probability in a vaccinated and unvaccinated cat, respectively. Shaded boundaries represent 95% CIs.
Other risk factors associated with FCV shedding included age and vaccination status, confirming results of previous studies (Porter and others 2008, Coyne and others 2012). In this population, vaccinated cats up to 48 months of age were less likely to shed FCV than unvaccinated animals of similar ages, suggesting in this age group, vaccination may have reduced infection rates. Experimental studies have shown both vaccinated cats (Redersen and Hawkins 1995, Poulet and others 2005) and cats recovered from acute FCV infection (Povey and Ingersoll 1975) remain susceptible to subsequent infection. However, such previous exposure is often associated with a reduction in the shedding of subsequent heterologous challenge (Povey and Ingersoll 1975, Poulet and others 2005), consistent with the reduced risk of testing FCV positive identified here. The loss of a demonstrable protective effect of vaccination on FCV infection over 48 weeks of age could represent the acquisition of immunity in the unvaccinated cats following field infection, a consequence of the high prevalence of FCV in the population. However, since the authors did not model here the time since last vaccine, the apparent loss of protective effect may also reflect increasing time since the last vaccination in older animals.

Sequence analyses identified 46 strains among the 72 isolates sequenced, confirming the high levels of strain diversity seen previously (Coyne and others 2012, Prikhodko and others 2014). Although the authors have not assessed the antigenicity of these viruses, it seems likely that because of the known localisation of neutralising epitopes in the regions sequenced, such genetic diversity would be associated with similar antigenic diversity. These levels of diversity pose a continual challenge to FCV companies evaluate in relation to the antigens they choose.

An interesting finding of the present study in several countries best exemplified by a Portuguese cluster of five isolates including those in strain D (Fig 4a), suggesting occasionally more widespread and longer-term virus transmission as suggested previously in Japan (Sato and others 2002). However, the overall picture remains one of lack of international promiscuity, very distinct from human caliciviruses where some strains or variants such as human norovirus genotype ii.4 are widely distributed internationally (Ramani and others 2014). The authors’ current hypothesis is that the rate of FCV evolution is so high that evolutionary signals become saturated rapidly, leading to a loss of phylogenetic resolution over relatively short time periods (Coyne and others 2012). This will only be resolved when longer sequences are obtained.

Some of the strains the authors identified were associated with high levels of within-strain diversity, notably strains A, C, D and G. These levels of diversity are reminiscent of those seen in endemically infected, multicat households, where the authors hypothesised that high levels of population immunity lead to rapid evolution by positive selection (Coyne and others 2007b).

In five cases, individual households/communities of cats were infected with more than one strain (Fig 4a). The most extreme example was a Greek rescue shelter where all six isolates sequenced were distinct strains. This observation suggests lots of viruses coming together into a single population but with little transmission, a pattern either attributable to very good internal biosecurity (Radford and others 2001b, Coyne and others 2007a), or suggesting that the cats were sampled on arrival in the shelter before transmission had a chance to occur. A more detailed understanding of the population demographics and husbandry of these populations may shed more light on the behaviours that underlie these phylogenetic patterns.

In the present study, the authors found no FCV F9-like sequences, a common strain used in live vaccines. This is in contrast to previous studies which have reported albeit rare occurrences of F9-like viruses in the general cat population (Radford and others 2001a, Coyne and others 2007b, 2012). The origins of these F9-like viruses are unknown but because of their reported close sequence similarity to the original F9 sequence, it seems reasonable they would have originated from regular use of live vaccines containing this strain. The authors’ failure to find them in this study may reflect the fact that cats within 1 month of receiving a live vaccination were excluded from the study population. Together, this suggests that on the rare occasions...
FIG 4: Unrooted Kimura 2-parameter Neighbour joining tree of (a) 74 partial capsid sequences (including three sequences for FR1_11) and (b) 73 partial polymerase sequences obtained in this study. Each sequence has a unique ID made up of country code (FRance, GErmany GReece, PorTugal and UK) and site number (1–5) followed by sample number. Those strains represented by more than a single sequence (<20 per cent capsid divergence) are boxed, additionally labelled A–N, and the intrastrain capsid diversity indicated in the box. Where multiple sequences come from a single household they are indicated by an additional symbol (□, ○, Δ, *, †). Two new strain clusters O and P with high bootstrap support are also indicated. Where clustering varies between capsid and polymerase phylogenies, the sequences are linked in each phylogeny by a thin grey line. The feline calicivirus (FCV) vaccine strain F9 is also included (GenBank accession No. M86379). The percentage of replicate trees in which the associated taxa clustered together in bootstrap tests (1000 replicates) is shown next to the branches; only bootstrap values >75 per cent are shown. Distances are drawn to scale and relate to the distance bar.
cats do shed vaccine-derived virus following vaccination, the duration is short, and the potential for onward transmission limited, consistent with reversion to virulence assays and dissemination experiments conducted for registration of live vaccines. This allows these vaccine-derived viruses to sporadically appear at a low level in the population in a state of evolutionary stasis.

The close co-circulation of distinct strains, and lack of sterility in individual cats, provides an ideal opportunity for mixed infections (Coyne and others 2006b). Sequence data suggested such mixed infections both at the quasispecies (within-strain) level and cats infected with more than one strain (FR1 11, 57 and 40). These diverse populations within an individual host provide an ideal opportunity for recombination (Lai 1992), and the identification of several incongruences between capsid and polymerase phylogenies presented here provide some evidence of recombination in these populations. Recombination is a feature of the evolution of many RNA viruses (Lai 1992), and has been described previously for caliciviruses in general and FCV in particular (Coyne and others 2006b); there is some evidence that a hot spot for the necessary template switch exists at the junction between ORF1 and ORF2, in between the polymerase capsid sequence generated in this study (Oliver and others 2004, Bull and others 2005), driven by predicted RNA secondary structure (Jiang and others 1993, Porter 2004, Coyne and others 2006b, Prichodko and others 2014). Formal conformation of recombination requires sequencing across the putative recombination site, but was beyond the scope of this project.

For 19 (21 per cent) of the 91 samples that showed FCV-like cytopathic effect in cell culture the authors were unable to generate PCR products, despite using several primer combinations for RNA viruses, especially when trying to amplify the most variable regions of the virus. In a previous study this has been described previously for caliciviruses in general and FCV (FR1_11, 37 and 40). These diverse populations within an individual host provide an ideal opportunity for recombination (Lai 1992), and evidence of recombination in these populations. Recombination requires sequencing across the putative recombination site (Jiang and others 1993, Porter 2004, Coyne and others 2006b, Prichodko and others 2014). Formal conformation of recombination requires sequencing across the putative recombination site, but was beyond the scope of this project.

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Study limitations: This study is the largest of its kind to provide molecular epidemiology of a prospective sample of FCV isolates from a European country, and the lack of dominant strains within the sample of European countries, con...
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