The feasibility and acceptability of collecting oral fluid from healthy children for anti-HCV testing

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This pilot study investigated the feasibility of surveying, anonymously, HCV infection among healthy children using an oral fluid specimen. Seventy seven per cent of children provided their assent, or where appropriate, consent to participate; 2.8% were anti-HCV positive. Oral fluid collection is acceptable to children and more extensive studies are indicated.

METHODS

Study setting and population

A cross sectional survey involving the collection of non-identifying epidemiological data and a saliva sample for voluntary anonymous anti-HCV testing was performed in the General Anaesthetic (GA) Department of the Glasgow Dental Hospital and School. The hospital serves the people of Glasgow and its surrounding areas. The setting was chosen because it was considered to be a non-intrusive environment in which to request an oral fluid sample and because Glasgow is known to have the highest prevalence of HCV in the UK and Europe.

Sample size and eligibility

In the context of what was a pilot study, the investigators aimed to recruit an arbitrary figure of 100 children. All children attending the GA Department between 14 and 24 June 2002 were asked to participate; children attending without a guardian were excluded.

Consent and ethics

Following an explanation of the study by a medically qualified researcher, written consent was sought from guardians. Assent from the children aged less than 7 years, and written consent from children aged 7 years and older, of consenting guardians was then sought. An information sheet for school age children was provided, while cartoons and play were used to describe the procedure to younger children (fig 1). If children refused to participate, verbally or non-verbally, no attempt was made to change their decision.

Ethical approval for the study was obtained from the Area Dental Ethics Committee, NHS Greater Glasgow.

Data collection

A standardised questionnaire was used to collect limited demographic information, including age, gender, and postcode sector of residence of the child. Risk information was not collected, although questions were asked of the guardians to determine whether they would be prepared to answer questions about, for example, injecting drug use behaviour.

Oral fluid collection and anti-HCV testing

Oral fluid specimens for anonymous anti-HCV testing were collected using the OraSure device (Altrix Healthcare plc, Warrington, UK). The collection of an oral fluid specimen has been shown to be of minimal risk to a child, and no adverse reactions to the collection device have been described. Salivary anti-HCV was detected using the Murex anti-HCV 4.0 ELISA (Abbot Diagnostics, Maidenhead, UK), with the following modifications: the specimen volume was increased to 1 vol of saliva (100 μl) to 1 vol of diluent (100 μl), and the incubation period was increased to overnight (~20 h) at room temperature; this assay has a sensitivity of 96% (personal communication, S Cameron). A positive salivary antibody test correlates well with serum HCV RNA status and is considered a good marker for hepatitis C viraemia in epidemiological studies.

RESULTS

From a total of 91 guardians approached, 84 (92%) consented to their child participating. Of the 84 children approached, 70 (83.3%) assented and provided an oral fluid specimen. Ten children read the information sheet and signed the consent form. The overall participation rate was 77% (70/91; 95% CI 68–86%). All guardians felt it was laudable that the child was directly involved in the decision to participate.

Table 1 presents the demographic characteristics of the study population. The sample was considered representative...
of the child population attending the GA Department. The median age of the children who participated (5 years) was significantly greater than those who did not (95% CI of the difference: 1–2 years, p = 0.0027). Participation was not associated with gender or deprivation category. Although a number of children found the collection device distasteful, only one child terminated their participation by removing the device from their mouth. All guardians, except one, indicated that they would be comfortable answering questions about their own risk behaviours.

The prevalence of HCV infection among the sample was 2.9% (2/70; 95% CI 0.35–9.9%).

**DISCUSSION**

Since the early 1990s, several epidemiological surveys of HIV and hepatitis virus infection among intravenous drug users in
the UK have used anonymous testing of oral fluid. This pilot study is the first attempt in the UK to use such a methodology to investigate the prevalence of HCV infection among a group of apparently healthy children, and showed that the voluntary collection of oral fluid specimens for anonymous HCV testing from children as young as 4 years is possible and acceptable to both the children and their parents. No conclusions can be drawn on the prevalence of anti-HCV among children in Glasgow, as the sample size of the pilot study was small.

In undertaking the study, the child’s rights, Scottish Law, ethical principles, and paediatric guidelines for ethical conduct of research with children were considered. The researchers were satisfied that: (1) the parents were happy with the adequacy of the consent procedures; and (2) the children were capable of providing their assent and, where appropriate, consent to their involvement. The investigators conclude that larger studies among children are feasible, and as the resultant prevalence indicates, necessary to determine HCV prevalence among young populations.

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REFERENCES