A Two Year Serological Surveillance of Coronavirus Infections in Hamburg

Introduction

Various surveillance programmes undertaken in recent decades have provided evidence that viral infection is the major identifiable cause of upper and lower respiratory tract disease in man.

Material and Methods

Population: A total of 3,016 blood samples was collected between October 1974 and September 1976 (minimum of 100 sera monthly) from "normal" healthy individuals divided into four age-groups: Group I (0 to 14 years old); Group II (15 to 24 years old); Group III (25 to 59 years old); Group IV (60 years old). We collected a second sample of blood two months later from 331 girls, 12 to 17 years old. After inactivation for 30 minutes at 56 °C, the sera were stored at -20 °C until used. All paired sera were tested simultaneously.

Antigen: Coronavirus strain OC-43 has been isolated by McIntosh et al. (3) in human embryo trachea organ culture from an adult with a cold. The strain was subsequently adapted to suckling mouse brain. Strain OC-43 antigen was prepared from infected suckling mouse brain according to the technique described by Kaye and Dowdle (4).

Serology: Hemagglutination inhibition (HI) tests were performed according to the Sever microtitre technique (5), using chicken red blood cells (4). Fifty positive sera (titre 1:16 – 1:64) were treated with receptor destroying enzyme (RDE) (6), kaolin (7) or ethacridin (8) for removal of nonspecific inhibitors.

Results

The incidence of seropositive reactions in the total population studied is presented in Table 1. From the 3,016 sera investigated, 1,755 (58.2%) showed HI antibody titres.

As some cases did not appear to be linked with infections by known viruses, it was hoped that the human coronaviruses discovered relatively recently might account for a proportion of these unexplained illnesses. That coronaviruses are potential respiratory pathogens has been demonstrated in experimental studies (1, 2). However, data on the extent and significance of infections by these viruses in human illnesses are still limited. Serological surveillance carried out at regular intervals on a large number of sera taken from representative groups reveals the total infection rates (apparent and non-apparent) and the level and distribution of the seroimmunity in the population.

The following report of seroepidemiological studies of coronavirus infections is based upon observations made in the course of a surveillance programme in Hamburg (carried out since October 1974) in various sections of the population in which all age groups were represented.
Table 1: HI-Coronavirus antibodies in the population of Hamburg.

| Age        | 0-14 | 15-24 | 25-59 | 60+  |
|------------|------|-------|-------|------|
| October 1974 | 403  | 368   | 571   | 282  |
| September 1975 | 329  | 350   | 369   | 344  |
| September 1976 | 1324 | 999   | 1624  | 1392 |

| HI-antibody titer | October 1974 | September 1975 | September 1976 |
|-------------------|--------------|----------------|----------------|
| 1:8               | 90           | 54             | 112            |
| 1:16              | 49           | 113            | 112            |
| 1:32              | 49           | 85             | 40             |
| 1:64              | 31           | 30             | 10             |
| 1:128             | 7            | 2              | 5              |

As far as the seasonal pattern is concerned, we observed frequent infections in November 1974; after that period the seroimmunity level of the population studied dropped continuously till June 1975. An increased infection rate, with a peak in October to November 1975, was evident. From December 1975 to February 1976, the infection rates dropped markedly, followed by a smooth increase till the end of our present study in September 1976.

Table 2: HI-Coronavirus antibodies in paired sera (Hamburg 1974 to 1976).

| Age   | Paired sera | Seropositive reactions | Constant HI-titer | Declining HI-titer | Rising HI-titer |
|-------|-------------|------------------------|-------------------|--------------------|-----------------|
|       | n | % | n | % | n | % | 2x | % | 4x | % |
| 0-14  | 131 | 95 | 72.5 | 53 | 40.4 | 9 | 6.9 | 25 | 19.1 | 8 | 6.1 |
| 12-17 | 200 | 185 | 92.5 | 133 | 66.5 | 14 | 7 | 24 | 12.1 | 14 | 7 |
| Total | 331 | 280 | 84.6 | 186 | 56.2 | 23 | 6.9 | 49 | 14.8 | 22 | 6.6 |

Discussion

Certain conclusions can be drawn from the data obtained during our two year surveillance period. Infections with human coronaviruses are common in all four groups studied, especially in Group II (15-24 years old). The higher incidence of seropositive reactions in 15 to 24 year olds (67.7% and 82.9%, respectively) confirms our previous data (9, 10, 11), those of Henigst (12), Monto and Lira (13), and Riski and Estola (14) who also found a higher percentage of seropositive reactions in the 15 to 19 year old group (12, 13) and in individuals of 21 to 50 years (14).

Considering the results obtained in Group I and II, our data are in agreement with those of Kaye and Dowdle (15). The higher level of coronavirus antibodies and the
The seasonal pattern suggests that coronaviruses are circulating widely in the urban population, with prevalence in the cold season, a decrease in the warm season and sporadic cases at other times of the year. Other authors (13, 17, 18, 19) also found coronavirus infections usual in the period January to April. In our study we observed peaks in November 1974, October to November 1975, and in September 1976, followed by low infection rates in the spring-summer season. Similar data were published by Bradburne and Somerset (20), and by McIntosh et al. (16).

We may assume that in the peak periods Hamburg witnessed a wave of infection caused by an OC-43 virus or a related virus. Neither the number of human respiratory types nor the extent of their serological cross-reactivity are well known as far as coronaviruses are concerned. It is known however that several members cross-react in one or more types of serological tests and that heterotypic antibody rises occur (21).

As we did not isolate coronaviruses in our patients and in view of the possibility of heterologous responses of the antibody titers that we have found, we cannot interpret them as being specifically due to the OC-43 virus alone. The antibody response found in this seroepidemiological survey may be an expression either of past or present infections or reinfections with an agent identical or closely related to the antigen used in the test. Reinfections with coronaviruses are not a rare event considering that over 50% of the titer rises encountered in the 331 paired sera occurred in individuals with prior antibodies. The high rates of seropositive reactions provide complete evidence that coronaviruses are circulating widely in the urban population of Hamburg.

Literature

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