Infection with cytomegalovirus in homosexual men

R A COUTINHO,* P WERTHEIM-vAN DILLEN,† P ALBRECHT-vAN LENT,*
N NAGELKERKE,‡ H KUIPERS,§ A VAN BENTUM-vAN HAAGEN,† T RIJSDEIJK,*
AND J VAN DER NOORDAA†

From the *Municipal Health Service, the †Laboratorium voor de Gezondheidsleer, the ‡Laboratory of
Medical Physics, and the §Bureau of Statistics, Amsterdam, The Netherlands

SUMMARY  The prevalence and incidence of cytomegalovirus (CMV) infections were studied in a
group of homosexual men. Of the 710 participants, 501 (70.6%) had complement fixing anti-
body to CMV on entry to the study. During the follow up (maximum 23 months) 69 CMV
infections were found: 50 primary infections among the 209 seronegative men (attack rate 27.3%),
and 19 recurrent infections among the 501 seropositive men (attack rate 6.2%).

The prevalence of antibody to CMV was correlated with four characteristics of the participants’
lifestyles: duration of homosexual activity, number of different sexual partners, history of
syphilis, and anal sexual contact. Among the seronegative men, the incidence of primary infection
with CMV correlated with a history of syphilis and anal sexual contact.

We conclude that infections with CMV are highly prevalent among homosexual men, and that
anal sexual contact plays an important part in the transmission of this virus.

Introduction

There is evidence that infections with cytomegalovirus (CMV) can be transmitted by sexual contact,
especially among homosexual men. A study of male homosexuals in San Francisco found a very high
(94%) prevalence of antibodies to CMV, and 14% of
the men under 30 years old had CMV viruria.1 In a
Danish study the prevalence of antibodies among
homosexual men was related to the duration of
homosexual activity.2 CMV has also been suggested
as having a role in the occurrence of the acquired
immune deficiency syndrome (AIDS) and in the
aetiology of Kaposi’s sarcoma.3

During a study of the efficacy of a hepatitis B
vaccine we recently followed a large group of homo-
osexual men over a period of nearly two years. This
gave us an opportunity to study the prevalence and
incidence of infection with CMV and its relation with
a number of risk factors in this group of men.

Patients, materials, and methods

STUDY POPULATION

The study population consisted of male homosexuals
who had participated in an efficacy trial with a heat
inactivated hepatitis B vaccine between November
1980 and December 1982.4 Men could enter this trial
if they were between 16 and 50 years old, were
negative for hepatitis B virus markers, had serum alanine transferase (serum ALT) activity <50 IU/l
(normal activity <21 IU/l), no serious illnesses, and
had had at least two different male sexual partners in
the preceding six months. Men who lived in and
around Amsterdam were selected for the study of
CMV.

CONDUCT OF THE STUDY

At the beginning of the study (month 0) the partici-
pants were questioned about their medical history
and lifestyle. Age was taken in years at the time of
the interview. The duration of homosexual activity
was defined as the number of years from the first
homosexual contact until the time of the interview. A
history of syphilis was noted. Each man was asked to
give an estimate of the number of different sexual
partners in the six months preceding the interview.
The question about anal sexual contact did not specify
whether intercourse was active or passive.

Blood samples were taken at monthly intervals
during the first five months and every three months
thereafter. The first and last blood samples were
tested for the presence of antibodies to cytomegalo-
virus (anti-CMV). If a seroconversion or a significant
(more than fourfold) rise in titre was found, all other
samples were tested for anti-CMV.
CMV infections were defined as: primary CMV infection (seroconversion for anti-CMV in at least two sequential blood samples, confirmed by the presence of anti-CMV IgM antibodies); or recurrent CMV infection (a more than fourfold rise in titre in a man with anti-CMV (complement fixation >8), confirmed in at least one subsequent blood sample). A man was considered to have had clinical signs of infection with CMV if he had had either viral hepatitis (serum ALT activity ≥50 IU/l at least once with negative serological markers for hepatitis B and A) or a mononucleosis-like illness of more than one week's duration at the time of the seroconversion or the rise in titre.

**Laboratory Tests**

Antibodies to CMV were measured by complement fixation, using AD 169 as the antigen (Flow laboratories, Virginia, USA). An enzyme linked immunosorbent assay was used to test for IgM antibodies specific to CMV. Serum samples from each man were titrated simultaneously. Serum ALT activity was measured by an automated kinetic method.

**Statistical Analyses**

To study the association between the prevalence of anti-CMV and potential risk factors stepwise logistic regression was used. The incidence of CMV during follow up was investigated with life table methods. Attack rates were estimated with the product limit method, and the relation to risk factors was evaluated with Cox's proportional hazards regression model. These statistical methods and the formulae have been described previously.

**Results**

**Characteristics of Participants**

A total of 710 homosexual men participated in this study. Table I shows that their mean (SD) age was 30·1 (7·0) years, the mean (SD) duration of their homosexual activity was 11·0 (7·0) years, 35% had more than 10 different sexual partners in the preceding six months, 18·6% had a history of syphilis, and 74·2% had had anal sexual contact.

**CMV Infections Among Participants**

Of the 710 participants, 501 (70·6%) were found to have complement fixing antibodies to CMV at entry to the study. A total of 69 infections with CMV were diagnosed during follow up: 50 primary infections among the 209 seronegative men and 19 recurrent infections among 501 seropositive men. Only 12 men had clinical signs of CMV infection: nine had a mononucleosis-like illness, and three had viral hepatitis. Of these 12 men, 10 had a primary and two a recurrent infection.

Figure 1 shows that at the end of the study (after 23 months) the attack rate for primary infections was 27·3% and for recurrent infections 6·2%. As the curves climbed steadily there was no evidence of clustering of the CMV infections.

**Association Between Prevalence of Antibody and Potential Risk Factors**

Table II shows that four characteristics of the participants’ lifestyles correlated (by stepwise logistic regression) with seropositivity for CMV. The duration of homosexual activity had the highest influence (p<0·002), the probability of being seropositive

**Table I** Characteristics of homosexual men participating in a study of cytomegalovirus

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>710</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number entering study</td>
<td>710</td>
</tr>
<tr>
<td>Mean (SD) age (years)</td>
<td>30·1 (7·0)</td>
</tr>
<tr>
<td>Mean (SD) duration of homosexual activity (years)</td>
<td>11·0 (7·0)</td>
</tr>
<tr>
<td>More than 10 different sexual partners in preceding six months (%)</td>
<td>35·0</td>
</tr>
<tr>
<td>History of jaundice (%)</td>
<td>11·3</td>
</tr>
<tr>
<td>History of syphilis (%)</td>
<td>18·6</td>
</tr>
<tr>
<td>Anal sexual contact (%)</td>
<td>74·2</td>
</tr>
<tr>
<td>Oral sexual contact (%)</td>
<td>97·7</td>
</tr>
<tr>
<td>Hepatitis A antibodies (%)</td>
<td>40·8</td>
</tr>
</tbody>
</table>

**Figure 1** Life table showing the attack rates for (a) primary and (b) recurrent infections with cytomegalovirus.
Infection with cytomegalovirus in homosexual men

**TABLE II** Correlation of characteristics of lifestyles with seropositivity for cytomegalovirus in homosexual men (evaluation by stepwise logistic regression)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Co-efficient (β)</th>
<th>Standard error (SE)</th>
<th>β/SE</th>
<th>p value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duration of homosexual activity</td>
<td>0.047</td>
<td>0.015</td>
<td>3.20</td>
<td>&lt;0.002</td>
</tr>
<tr>
<td>Number of different sexual partners in preceding six months</td>
<td>0.128</td>
<td>0.058</td>
<td>2.21</td>
<td>&lt;0.03</td>
</tr>
<tr>
<td>History of syphilis</td>
<td>0.474</td>
<td>0.142</td>
<td>3.34</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Anal sexual contact</td>
<td>0.204</td>
<td>0.100</td>
<td>2.04</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>Constant</td>
<td>0.166</td>
<td>0.269</td>
<td>0.62</td>
<td>NS</td>
</tr>
</tbody>
</table>

*two tailed

Increasing by 1-2% with each year of homosexual activity (fig 2). This effect was independent of age. The next important risk factor was the number of different sexual partners in the preceding six months (p<0.03), the risk increasing with an increasing number of partners (fig 3). A history of syphilis significantly increased the odds P(+)P(-) by 61%, and anal sexual contact by 23%. Other variables such as age, oral sexual contact, and history of jaundice had no appreciable influence in this study.

ASSOCIATION BETWEEN INCIDENCE OF CMV DURING FOLLOW UP AND POTENTIAL RISK FACTORS

Table III shows that two variables correlated with the risk of contracting a primary CMV infection among the 209 antibody negative homosexual men: a history of syphilis (relative risk 2.21), and anal sexual contact (relative risk 2.49). The other characteristics of lifestyle (including the number of different sexual partners in the preceding six months) had no significant influence on the rate of primary infection with CMV.

None of the characteristics of lifestyle of the seropositive participants was found to be appreciably associated with the risk of acquiring a recurrent CMV infection during follow up.

Discussion

The prevalence of antibodies to CMV among adults in Europe and North America generally varies between 30% and 60%, depending on the socioeconomic background of the population, with a much higher prevalence among male homosexuals in San Francisco (94%) and Copenhagen (87%). In our study of 710 homosexual men living in and around Amsterdam a...
somewhat lower prevalence (71%) was found. The reason for this relatively low prevalence is probably the selection of the participants who were all hepatitis B marker negative. Hepatitis B is sexually transmitted among homosexual men, and we therefore selected those men with a relatively short duration of homosexuality and a relatively low total number of different sexual partners.

Among the 209 seronegative men, 50 primary CMV infections were detected during follow up, the attack rate being 27·3% at the end of the study (after 23 months). This attack rate is very high when compared with incidences in other large prospective studies among healthy adults. Stern et al followed 254 seronegative English women during pregnancy and found eight (3%) primary CMV infections, and Griffith et al in a study of 1608 seronegative pregnant women found 14 (0·9%) primary CMV infections. Among 713 seronegative English students, 10 (1·4%) seroconverted during a follow up period of seven months.

Among the 501 seropositive men, 19 recurrent CMV infections were detected serologically during the follow up period of 23 months (attack rate 6·2%).

It is difficult to draw conclusions about the recurrent infections as we have only serological data, and viral cultures were not performed. The majority of recurrent CMV infections represent reactivation of latent virus, although reinfection with a new virus strain occurs occasionally. From a Danish study it appears that a high percentage of antibody positive homosexual men excrete CMV in their seminal fluid. There is therefore ample opportunity for promiscuous homosexual men to become reinfected with a new virus strain, but our serological data indicate that this may be an uncommon event.

We found that the prevalence of antibody to CMV correlated with four characteristics of the participants' lifestyle: duration of homosexual activity, number of different sexual partners in the preceding six months, history of syphilis, and anal sexual contact. Among the seronegative men, the primary incidence of infection with CMV correlated with two variables: a history of syphilis and anal sexual contact. These correlations indicate that CMV infections are sexually transmitted among homosexual men. It is not completely clear why no correlation was found in this study between the number of different sexual partners in the preceding six months and the incidence of CMV among those susceptible. This may indicate that the number of sexual partners is of less importance than anal sexual contact in the transmission of CMV. CMV is known to be present in semen and it seems likely from this study that anal sexual contact plays an important part in the transmission of CMV among this group of men. Another sexually transmitted disease among male homosexuals, hepatitis B, is also related to anal sexual contact. In a recent study of homosexual men hepatitis A was also found to be sexually transmitted but no relation was found with anal sexual contact alone, which indicates a different means of transmission (by orogenous contact). We conclude that CMV infections are highly prevalent among homosexual men and that anal sexual contact plays an important part in the transmission of the virus.

We thank the participants for their cooperation, and W Maruana, J van Marle, N Lelie, and H Ross for their help. This study was supported by the Dutch ministry of health and in part by the Netherlands foundation for preventive medicine (grant no 28-440).

References


Infection with cytomegalovirus in homosexual men.

R A Coutinho, P Wertheim-van Dillen, P Albrecht-van Lent, N Nagelkerke, H Kuipers, A van Bentum-van Haagen, T Rijsdijk and J van der Noordaa

Br J Vener Dis 1984 60: 249-252
doi: 10.1136/sti.60.4.249

Updated information and services can be found at: http://sti.bmj.com/content/60/4/249

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to: http://group.bmj.com/group/rights-licensing/permissions
To order reprints go to: http://journals.bmj.com/cgi/reprintform
To subscribe to BMJ go to: http://group.bmj.com/subscribe/