Syphilis and HIV infection in patients with hepatitis A: a preliminary study from one centre in Poland

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In 2017, there was a significant increase in hepatitis A incidence in Poland and several dozen times more cases were reported in comparison to previous years [1]. For instance, in the reference centre for infectious diseases for the Greater Poland Voivodeship in the period from the beginning of February to the end of August 2017, more patients with hepatitis A were hospitalized than in the previous 10 years in total.

As hepatitis A virus (HAV) infections are transmitted through the faecal-oral route, hepatitis A can spread during intimate contacts and some specific sexual practices [2, 3]. For this reason, we hypothesized that it could be an indicator of risky sexual behaviours and diagnosis of hepatitis A may be an opportunity to test patients for other sexually transmitted infections (STI).

Persons hospitalized at the Department of Infectious Diseases in Poznan (Jozef Strus Multidisciplinary City Hospital) for symptomatic HAV infection from February to mid-August 2017 were included in the present analysis.

The objectives of this study were a simple demographic description of patients with hepatitis A and their screening for some STI that may be asymptomatic (syphilis, HIV infection, hepatitis B, hepatitis C).

The study was approved by the local Bioethics Committee. All patients gave their informed consent.

One hundred patients aged 20–64 (median: 29.5 years) with symptomatic hepatitis A have been screened for the above mentioned STI.

Testing for HAV (HAVAb-IgM), HBV (HBsAg Qualitative), HCV (anti-HCV) and HIV (HIV Ag/Ab Combo) infections was performed using the ARCHITECT system (Abbott Laboratories). Each positive result of the screening test for HIV infection was confirmed with the western blot technique.

RPR Carbon (Hydrex Diagnostics Sp. z o.o., Warszawa, Poland) and Immuntrep TPHA (Omega Diagnostics Ltd, Alva, Scotland) assays were used for screening and confirmation of syphilis, respectively. Syphilis was diagnosed in the following circumstances: the presence of symptoms or signs consistent with an early stage of disease (primary or secondary syphilis) and reactive RPR and TPHA tests, or a 4-fold increase from baseline in RPR titres with a reactive TPHA in patients who had syphilis in the past, or positive RPR (at least 1 : 8 titre in quantitative assessment) and TPHA in patients who have never been tested for syphilis before.

The majority of patients were men (93%). A significant proportion of male study participants (76.3%; 71/93) described themselves as men having sex with men (MSM).

Twenty patients (20%; 19 men including 17 MSM) travelled abroad in the period of 2 months preceding the first symptoms of hepatitis A. All but one visited only European countries.

The peak values of hepatic biochemistry in patients from our centre are shown in the Table 1.

All but 1 patient (woman) were tested for HIV infection, and it was only found in men. In most cases (79.3%; 23 out of 29 HIV-positive men with hepatitis A), it was diagnosed and treated with antiretroviral therapy before hospitalization for symptomatic HAV infection. The new HIV infection diagnosis concerned 6/70 men (8.6%) who were previously HIV-negative or unaware of their HIV status.

RPR was reactive or equivocal in 20 patients out of 89 tested (8 men and 3 women were not examined for syphilis). All of them were MSM. When taking into account all available data (past medical history, clinical presentation along with previous and current syphilis test results), an ongoing Treponema pallidum infection was
ultimately diagnosed in 11 out of 85 men tested (12.9%). Only 2 of them (both HIV-negative) had the symptomatic disease (secondary syphilis), 4 men (two were HIV-positive) had the first-time diagnosis of latent syphilis, and 5 patients (all were HIV-positive) had recurrent latent syphilis. In addition, 14 individuals (including 11 HIV-positive) with no active disease were treated for syphilis in the past.

The presence of HBsAg was found in 3 out of 100 subjects (all were MSM, two co-infected with HIV). They all knew about their HBV infection before the episode of hepatitis A occurred. Anti-HCV positivity was shown in 3 out of 99 persons tested. All were HIV-infected MSM, and 2 of them were aware of their HCV infection. HCV-RNA was negative in the third patient.

Importantly, in 17.2% (16/93) of these men, a previously unrecognized episode of STI was diagnosed.

The need for periodic screening for syphilis in HIV-positive patients and those with syphilis for HIV-infection is well documented and recommended [7]. Such a paradigm was also recently confirmed by two groups from Poland [8, 9].

Based on the results of our study, we postulate that at least 23.9% (17/71) of inpatients concerned these inpatients. Secondly, no tests for other STI. Some previous observations suggested the validity of such an approach in Poland, and the epidemic of 2017 confirmed this [3, 10]. It is very important for STI, which may be asymptomatic, including syphilis and HIV infection. Their recognition allows to screen the analysis. However, in the study period, about 70% of all hepatitis A cases diagnosed in our centre concerned these inpatients. Secondly, no tests for syphilis were performed in some persons, which might result in an underestimation of this disease. Only in single cases were both HIV and HCV assays not carried out. Thirdly, while writing this Letter, no molecular testing results were available to prove linkage with outbreaks in other European countries.

In conclusion, the hepatitis A outbreak in the Greater Poland region in 2017 concerned mainly MSM. Men with hepatitis A should always be screened for other STI, including syphilis and HIV infection.

### Conflict of interest

The authors declare no conflict of interest.

| Table 1. Peak levels of hepatic biochemistry parameters in study participants (n = 100) |
| Parameter | Median (Q1–Q3) |
|-----------|----------------|
| Alanine aminotransferase (ALAT) [U/l] | 8287 (1423–4141) |
| Reference range: 4–45 |
| Aspartate aminotransferase (AspAT) [U/l] | 1427 (475–3020) |
| Reference range: 4–35 |
| Total bilirubin [μmol/l] | 167 (114–200) |
| Reference range: 3–17 |
| γ-Glutamyltransferase (GGT) [U/l] | 248 (160.5–344) |
| Reference range: 5–55 |
| Alkaline phosphatase (ALP) [U/l] | 209 (165–265) |
| Reference range: 46–116 |

Q1, Q3 – quartiles 1 and 3.

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