Outcomes for Indigenous and non-Indigenous patients who access treatment for hepatitis C in the Top End of the Northern Territory

TO THE EDITOR: Chronic hepatitis C virus (HCV) infection affects over 225,000 Australians and is a leading cause of the need for liver transplantation and of liver-related death, but curative treatments are available. Ethnicity is a major determinant of treatment responsiveness, with the lowest sustained virological response (SVR) rates reported in African patients, and the highest in Asian patients. Much of this difference is accounted for by racial differences in polymorphisms in the interleukin-28B (IL28B) gene; however, it is unknown how common these polymorphisms are in Indigenous Australians, and no studies have been published about hepatitis C treatment outcomes among Indigenous Australians.

The hepatitis C treatment service for the Top End of the Northern Territory is run from a community-based sexual health clinic in Darwin. As clinicians working at this service, our perception was that Indigenous people rarely accessed the service or received treatment for HCV infection. Further, we were concerned that — due to social, cultural and linguistic barriers — Indigenous people who accessed the service may be less likely to commence treatment and to successfully complete treatment and achieve an SVR.

During the period 1 January 2006 to 31 December 2010, 243 patients were seen on at least two occasions for assessment of HCV infection; all were adults and 22 (9%) were Indigenous. During the audit period, HCV infection was treated with pegylated interferon-α plus ribavirin for 24–48 weeks. There were no significant differences in the proportion of patients who went on to commence and complete treatment, and to achieve an SVR, between Indigenous and non-Indigenous patients (Box). Of five Indigenous patients tested for IL28B genotype, all had the favourable CC polymorphism at the rs12979860 locus. Compared with the unfavourable TT and CT polymorphisms, the CC polymorphism at this locus is associated with at least a twofold higher chance of achieving a cure of HCV with interferon treatment, due to enhanced host immune responsiveness to interferon.

In conclusion, Indigenous people in the NT who access hepatitis C treatment services have a similar chance of achieving a cure . . . to non-Indigenous people.

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 Davis et al

Indigenous compared with non-Indigenous patients who attended a hepatitis C treatment service on at least two occasions for assessment of HCV infection*

<table>
<thead>
<tr>
<th></th>
<th>Indigenous (n = 22)</th>
<th>Non-Indigenous (n = 221)</th>
<th>P</th>
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</thead>
<tbody>
<tr>
<td>Age, median (interquartile range)</td>
<td>41.0 (36.2–45.0)</td>
<td>47.3 (39.3–52.1)</td>
<td>0.14</td>
</tr>
<tr>
<td>Men</td>
<td>15/22 (68% [45%–86%])</td>
<td>144/221 (65% [58%–71%])</td>
<td>0.78</td>
</tr>
<tr>
<td>HCV genotype 1</td>
<td>10/18 (56% [31%–78%])</td>
<td>87/173 (50% [43%–58%])</td>
<td>0.67</td>
</tr>
<tr>
<td>Commenced HCV treatment</td>
<td>11/22 (50% [28%–72%])</td>
<td>99/221 (45% [38%–52%])</td>
<td>0.58</td>
</tr>
<tr>
<td>Completed HCV treatment</td>
<td>9/11 (82% [48%–98%])</td>
<td>80/99 (81% [72%–88%])</td>
<td>0.94</td>
</tr>
<tr>
<td>Achieved sustained virological response†</td>
<td>4/8 (50% [16%–84%])</td>
<td>54/88 (61% [50%–72%])</td>
<td>0.53</td>
</tr>
</tbody>
</table>

HCV = hepatitis C virus. * Data are number/denominator (% [95% CI]) unless otherwise indicated. † Denominators represent those patients for whom 6-month post-treatment blood test results were available.

MJA 199 (1) · 8 July 2013

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doi: 10.5694/mja13.10083

Competing interests: No relevant disclosures.


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