DC-SIGN Polymorphisms Associate with Risk of Hepatitis C Virus Infection Among Men who Have Sex with Men but not Among Injecting Drug Users

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We aimed to identify whether genetic polymorphisms within L-SIGN or DC-SIGN correlate with hepatitis C virus (HCV) susceptibility. A men who have sex with men (MSM) and an injecting drug users (IDU) cohort of HCV cases and multiple-exposed uninfected controls were genotyped for numerous L-SIGN and DC-SIGN polymorphisms. DC-SIGN single nucleotide polymorphisms (SNPs) −139, −871, and −939 correlated with HCV acquisition in the MSM cohort only. When the same SNPs were introduced into a transcription activity assay they demonstrated a reduction in expression with predicted alteration in binding of transcription factors. DC-SIGN promoter SNPs correlated with risk of HCV acquisition via sexual but not IDU exposure, likely through modulation of mRNA expression levels.

Keywords. Hepatitis C virus; lectin; DC-SIGN; single nucleotide polymorphism; sexual transmission.

Hepatitis C virus (HCV) represents a major global health burden, with 350,000 people dying annually from HCV-related liver disease [1]. Intravenous drug use is now the major transmission route. Nevertheless, since 2000, sexual transmission has been reported frequently among HIV-infected men who have sex with men (MSM) and is associated with high-risk sexual behavior. Interestingly, some individuals remain uninfected despite practicing high-risk behavior(s). Studies have shown that ultimately 10%–20% of injecting drug users (IDU) do not seroconvert, suggesting a biological reason why some individuals are less prone to contract HCV [2].

DC-SIGN (dendritic cell specific ICAM-grabbing nonintegrin, CD209) and L-SIGN (DC-SIGN related, CD209L) are C-type lectins, which have been suggested to play a role in HCV transmission and infection [3]. DC-SIGN is a calcium-dependent cell-surface lectin of dendritic cells (DCs) [4]. DCs are localized in skin and mucosal tissues and may serve as a replication reservoir for HCV [4, 5]. L-SIGN is mainly expressed on liver and lymph node sinusoidal endothelial cells. It shares 77% amino acid identity with DC-SIGN and it has been shown to capture several viruses, including HCV [3]. Whereas the DC-SIGN neck region on exon 4 is highly conserved (7 repeats in the majority of individuals) the L-SIGN neck region is very variable [6]. This repeat region has been suggested to affect disease susceptibility and outcome for HIV-1 infection [7–9].

The objective of this study was to analyze the frequency of previously reported genetic variations in DC/L-SIGN genes in individuals from 2 well-defined cohorts at risk of HCV infection who either seroconverted or remained uninfected. We identified 3 DC-SIGN SNPs that were associated with HCV susceptibility through high-risk sexual exposure but not with IDU. Furthermore, we assessed whether these SNPs in the DC-SIGN promoter affect its activity.

METHODS

Study Populations

MSM Cohort (MOSAIC)

Sixty-two HIV-1 infected MSM participating in the MSM Observational Study of Acute Infection with Hepatitis C (MOSAIC) cohort were included. Risk behavior data was available from behavioral questionnaires collected at 6-month intervals and MOSAIC Risk Scores were calculated [10]. Participants were categorized as multiple exposed uninfected (MEU, n = 30) or multiple exposed infected (MEI, n = 32) based on reported behavioral risk factors at inclusion or any of the follow-up visits, which have been shown to be associated with increased risk of acquiring HCV sexually [10, 11]. The MOSAIC study was approved by the Institutional Review Board of the Academic Medical Center under assigned study numbers NL26485.018.09 and NL48572.018.14.

IDU Cohort (Amsterdam Cohort Studies)

Sixty-two participants from the Amsterdam Cohort Studies (ACS) among IDUs were selected, who started injecting drugs intravenously before 1990, a period of high HCV...
incidence (up to 27.5/100 person years) [12]. The ACS among IDUs was an open prospective cohort study recruiting drug users between 1985 and 2016 investigating the epidemiology, natural history, and pathogenesis of HIV-1 infection and other blood-borne and/or sexually transmitted diseases. Participants who injected more than 2 years and remained HCV seronegative during follow-up (n = 40) were classified as MEU whereas 22 MEI seroconverted for HCV during follow-up. Total duration of injecting drugs and follow-up was similar for MEU and MEI (Supplementary Table S1). During follow-up no statistical difference was found between MEI and MEU when comparing needle-sharing events. The ACS study was approved by the Institutional Review Board of the Academic Medical Center under assigned study numbers MEC 07/182 and MEC 09/040.

DNA Isolation and Genotyping
DNA was isolated from 200 µL participant serum utilizing the QIAamp DNA blood mini kit (Qiagen). The number of repeat domains within the L-SIGN repeat region was determined for each subject by polymerase chain reaction (PCR). PCR reactions contained 5 µL of template DNA, 400 nM forward primer, 400 nM reverse primer, 2.5 mM MgCl₂, 0.2 mM dNTPs, 0.1 mg/mL bovine serum albumin (BSA), 1.25 units FastStart Taq DNA polymerase in a total volume of 25 µL 1 × Faststart PCR buffer.

L-SIGN SNP rs2277998 was assessed using the Ready-to-use hot start reaction mix for High Resolution Melting (HRM) curve analysis using the LightCycler® 480 (Roche). The reaction contained 2.0 µL DNA template, 2.5 mM MgCl₂, 8 ng α-casein, 450 nM forward primer (Biologio) and 450 nM reverse primer (Biologio) in a total volume of 20 µL 1 × HRM master mix.

To assay reported DC-SIGN SNPs in the promoter region at positions −939 (rs735240), −871 (rs735239), −336 (rs4804803), and −139 (rs2287886), a DNA fragment covering approximately 1000 bp upstream of the ATG translation start site was amplified with 2 primer sets. The amplicons were sequenced in both directions with the same primers using Big Dye Terminator directions with the same primers using Big Dye Terminator

RESULTS

DC-SIGN −139GG, −871GG, and −939AA Are Associated With Reduced HCV Susceptibility in MSM
Patient characteristics are summarized in Supplementary Table S1. In the MSM cohort, 3 DC-SIGN SNPs were significantly associated with HCV infection (Table 1). The −139GG was found more frequently in MEU (63.3% in MEU compared to 37.5% in MEI). Additionally, −871GG (36.7% in MEU compared to 12.5% in MEI) and the −939AA (53.3% in MEU compared to 21.9% in MEI) were found more often in MEU, indicating that −139GG, −871GG, and −939AA genotypes protect against HCV acquisition (OR, 0.35; P = .045; OR, 0.23 P = .027; and OR, 0.23 P = .009, respectively). The −336 SNP was not significantly associated with HCV susceptibility.
As a statistically significant difference was found in the baseline Mosaic Risk Score between MEU and MEI, a sensitivity analysis was done, including only participants with a MOSAIC Risk Score ≥ 2. The association became stronger for all 3 SNPs (−139, −871, and −939), with strong statistical significance for SNP −871 and SNP −939 (Supplementary Table S3). In the ACS IDU cohort, no significant associations were found between SNPs and HCV susceptibility.

**No Associations Between L-SIGN Polymorphisms and HCV Susceptibility**

No association with HCV susceptibility was found for L-SIGN SNP rs2277998. In addition, the L-SIGN repeat distribution between MEI and MEU was similar for both cohorts (Supplementary Table S4). No significant difference in zygosity for the L-SIGN repeat was found between MEI and MEU (OR, 0.982 P = .961) (Supplementary Table S5).

**DC-SIGN SNPs Affect Promoter Activity**

We tested the effect of the promoter variants within the DC-SIGN promoter on transcription activity by using luciferase promoter constructs (Supplementary Figure S1). The −139G caused a 2.6-fold reduction (P < .001), the −871G a 3.3-fold reduction (P < .001), and the −939A a 1.4-fold reduction (P = .086) (Figure 1A). These data suggest that the DC-SIGN promoter variants affect transcription levels and thereby protein and cell surface expression patterns. We investigated whether the observed decrease in DC-SIGN promoter activity for specific SNPs could be due to alterations in TF binding sites, by a in silico comparison.
of predicted TF binding sites of promoter variants (Figure 1B). The variants at the −139, −871, and −939 sites do affect multiple predicted TF binding sites, with some putative sites lost (GR, C/EBP, PR-B, PR-A, HOXD9, HOXD10) and some TF binding sites gained (GR-Alpha, AP-2Alpha). This would indicate that the SNPs identified within the DC-SIGN promoter region can modulate activity through differential binding of transcription factors.

**DISCUSSION**

Here we investigated whether polymorphisms in DC-SIGN and L-SIGN correlated with susceptibility to HCV infection in 2 well-defined cohorts consisting of individuals at high risk of HCV infection through sexual or intravenous exposure. We selected polymorphisms based on what has been reported within the literature for HCV as well as other infectious agents. In the MSM cohort we identified an association of HCV susceptibility with 3 DC-SIGN SNPs. These SNPs were not associated with HCV susceptibility in the IDU cohort. No effects were found for the DC-SIGN −336 SNP, the L-SIGN SNP rs2277998, and repeat polymorphism in either cohort.

We studied 4 SNPs in the DC-SIGN promoter region, of which 3 (−139, −871, and −939) were found to correlate with HCV susceptibility in MSM, with −139G showing the strongest effect. Although the same SNPs have previously been associated with other infectious diseases, this is the first time SNPs have been reported to be associated with susceptibility to sexual transmission of HCV. Interestingly, the −139G SNP has also been reported to protect against sexual transmission of HIV-1 [14].

It has been published previously that the combination of −139G and −939A in the DC-SIGN promoter region significantly reduces DC-SIGN expression on immature DCs compared to −139A and −939G [15]. We now show that the −139G and −871G SNP independently cause a reduction in promoter activity, while the −939A variant failed to reach statistical significance (P = .085). The DC-SIGN promoter encodes multiple TF binding sites, which are in silico predicted to be affected by the −139, −871, and −939 variants. This strongly suggests that the decreased promoter activity observed in vitro is (at least partly) caused by a reduction in TF binding, which will require further testing.

As our study was small and HCV exposure may have been lower in the uninfected study groups, our observations clearly need to be confirmed in larger cohorts. However, the functional data supports the associations of the SNPs with protection against HCV acquisition. Collectively, our data suggest that DC-SIGN plays a role in HCV acquisition via sexual and not intravenous exposure. This effect appears to be mediated by reduced DC-SIGN expression, suggesting that DC-SIGN on DCs plays a role in sexual transmission of HCV, similar to its role in HIV infection [4]. We hypothesize that DCs transfer HCV to the liver through DC-SIGN; individuals with the protective genotypes will have lower DC-SIGN expression, resulting in a reduced susceptibility to sexual acquisition of HCV. Alternatively, DC-SIGN expression on DCs at mucosal surfaces may influence HCV antigen capture and induction of localized immune responses and modulate mucosal protection against HCV acquisition, which does not play a role in intravenous exposure. Further studies into the exact mechanism behind DC-SIGN affecting HCV infection susceptibility are warranted to better understand how DC-SIGN expression levels might influence immune responses, as well as mechanisms of transmission.

**Supplementary Data**

Supplementary materials are available at *The Journal of Infectious Diseases* online. Consisting of data provided by the authors to benefit the reader, the posted materials are not copyedited and are the sole responsibility of the authors, so questions or comments should be addressed to the corresponding author.

**Notes**

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References
1. World Health Organization. Data and statistics. World Health Statistics 2015. www.who.int/gho/publications/world_health_statistics/2015. Accessed 1 March 2016.
2. Sutton AJ, Gay NJ, Edmunds WJ, Hope VD, Gill ON, Hickman M. Modelling the force of infection for hepatitis B and hepatitis C in injecting drug users in England and Wales. BMC Infect Dis 2006; 6:93.
3. Cormier EG, Durso RJ, Tsamis F, et al. L-SIGN (CD209L) and DC-SIGN (CD209) mediate transinfection of liver cells by hepatitis C virus. Proc Natl Acad Sci U S A 2004; 101:14067–72.
4. Geijtenbeek TB, Kwon DS, Torensma R, et al. DC-SIGN, a dendritic cell-specific HIV-1-binding protein that enhances trans-infection of T cells. Cell 2000; 100:587–97.
5. Goutagny N, Fatmi A, De Ledinghen V, et al. Evidence of viral replication in circulating dendritic cells during hepatitis C virus infection. J Infect Dis 2003; 187:1951–8.
6. Barreiro LB, Patin E, Neyrolles O, Cann HM, Gicquel B, Quintana-Murci L. The heritage of pathogenic pressures and ancient demography in the human innate-immunity CD209/CD209L region. Am J Hum Genet 2005; 77:869–86.
7. Wichukchinda N, Kitamura Y, Rojanawiwat A, et al. The polymorphisms in DC-SIGNR affect susceptibility to HIV type 1 infection. AIDS Res Hum Retroviruses 2007; 23:686–92.
8. Chaudhary O, Bala M, Singh J, Hazarika A, Kumar R, Luthra K. The DC-SIGNR 7/5 genotype is associated with high dendritic cell counts and their subsets in patients infected with HIV-1. J Clin Immunol 2013; 33:788–97.
9. Liu H, Carrington M, Wang C, et al. Repeat-region polymorphisms in the gene for the dendritic cell-specific intercellular adhesion molecule-3-grabbing nonintegrin-related molecule: effects on HIV-1 susceptibility. J Infect Dis 2006; 193:698–702.
10. Newsum AM, Stolte IG, van der Meer JT, et al. Development and validation of the HCV-MOSAIC risk score to assist testing for acute hepatitis C virus (HCV) infection in HIV-infected men who have sex with men (MSM). Euro Surveill 2017; 22 pii: 30540.
11. Vanhommerig JW, Lambers FA, Schinkel J, et al. Risk factors for sexual transmission of hepatitis C virus among human immunodeficiency virus-infected men who have sex with men: a case-control study. Open Forum Infect Dis 2015; 2:ofv115.
12. van den Berg CH, Smit C, Bakker M, et al. Major decline of hepatitis C virus incidence rate over two decades in a cohort of drug users. Eur J Epidemiol 2007; 22:183–93.
13. Farré D, Roset R, Huerta M, et al. Identification of patterns in biological sequences at the ALGGEN server: PROMO and MALGEN. Nucleic Acids Res 2003; 31:3651–3.
14. Kagoné TS, Bisseye C, Méda N, et al. A variant of DC-SIGN gene promoter associated with resistance to HIV-1 in serodiscordant couples in Burkina Faso. Asian Pac J Trop Med 2014; 7(Suppl 1):S93–6.
15. Mezger M, Steffens M, Semmler C, et al. Investigation of promoter variations in dendritic cell-specific ICAM3-grabbing non-integrin (DC-SIGN) (CD209) and their relevance for human cytomegalovirus reactivation and disease after allogeneic stem-cell transplantation. Clin Microbiol Infect 2008; 14:228–34.