

ID Care Research Bibliography

Infectious Disease Research 2023 & Earlier • 155 studies

Care Safety

2023

Yuen MF, Heo J, **Nahass R**, Wong G et al. Preliminary safety and antiviral activity of AB-729 combination treatment with pegylated interferon alfa-2a in virally suppressed, HBeAg-negative subjects with chronic HBV infection. EASL 2023 Vienna Austria.

2022

Sulkowski MS, Agarwal K, Ma X, Nguyen TT, Schiff ER, Hann HWL, Dieterich D, **Nahass RG**. Efficacy and safety of vebicorvir administered with entecavir in treatment-naïve patients with chronic hepatitis B virus infection J Hepatology 2022.
<https://doi.org/10.1016/j.jhep.2022.05.027>

Yuen MF, Agarwal k, Ma X, Nguyen TT, Schiff ER, Hann HWL, Dieterich DT, **Nahass RG**. Safety and efficacy of vebicorvir administered in virologically-suppressed patients with chronic hepatitis B virus infection. <https://doi.org/10.1016/j.jhep.2022.04.005> (doi: 10.1016/j.jhep.2022.04.005)

2020

Fung S, Sulkowski M, Lalezari J, Schiff E, Dieterich D, Hassanein T, Kwo R, Elkhashab M, **Nahass R**. Antiviral activity and safety of the hepatitis B core inhibitor ABI-H0731 administered with a nucleos(t)ide reverse transcriptase inhibitor in patients with HBeAg-negative chronic hepatitis B. EASL 2020 London, England.

Impact Of Baseline Alanine Aminotransferase Levels On The Safety And Efficacy Of Remdesivir In Severe Covid-19 Patients, Aasld 2020

Jacobson IM, Ma X, Nguyen T, Schiff ER, Yuen MY, Hann HM, Sulkowski M, **Nahass R**, et al. Analysis Of Longer-Term Safety Profile Of The Hepatitis B Virus Core Inhibitor ABI-H0731 In An Open-Label Extension Study. AASLD 2020.

Yuen MF, Agarwal E, Ma X, Nguyen TT, Schiff E, Hann HW, Dieterich D, **Nahass R**, et al. Antiviral activity and safety of the hepatitis B core inhibitor ABI-H0731 administered with a nucleos(t)ide reverse transcriptase inhibitor in patients with HBeAg positive chronic hepatitis B infection in a long-term extension study. EASL 2020. Virtual

Janssen H, Hou J, Asselah T, Chan H, Zoulim F, Tanaka Y, Janczewska E, Efficacy and **Nahass RG**, Safety Results of the Phase 2 JNJ-56136379 JADE Study in Patients with Chronic Hepatitis B: Interim Week 24 Data. EASL 2020. Virtual

2019

Ma X, Lalezari J, Nguyen T, Bae H, Schiff ER, Fung S, Yuen MF, Hassanein T, Hann HW, Elkhashab M, Dieterich D, Sulkowski M, Kwo P, **Nahass RG**, et al. Interim Safety and Efficacy Results of the ABI-H0731 Phase 2a Program Exploring the Combination of ABI-H0731 with Nuc Therapy in Treatment-Naive and Treatment-Suppressed Chronic Hepatitis B Patients. EASL 2019, Amsterdam, Netherlands.

2018

Flamm SL, Cheng-Yuan P, Shibolet O, **Nahass R**, et al. Efficacy and Safety of Elbasvir (EBR)/Grazoprevir (GZR) in Hepatitis C Virus (HCV) GT1- and GT4-Infected Participants 65 Years and Older: An Integrated Analysis of Twelve Clinical Trials. Gastro 2018

2017

Wedemyer H, Wyles D, Reddy R, Luetkemeyer A, Jacobson I, Vierling JM, Gordon S, **Nahass RG**. Safety and efficacy of the fixed-dose combination regimen of MK-3682/grazoprevir/ruzasvir in cirrhotic or non- cirrhotic patients with chronic HCV GT1 infection who previously failed a direct-acting antiviral regimen (C-SURGE). EASL, Amsterdam 2017

Safety and Efficacy of Elbasvir (EBR)/Grazoprevir (GZR) in Hepatitis C Virus (HCV) GT1- and GT4-infected Participants 65 Years and Older: An Integrated Analysis of Twelve Clinical Trials. AASLD 2017

FlammSL, Peng CY, Shibolet O, **Nahass RG**, et al. Safety and efficacy of 12 weeks of Elbasvir (EBR)/Grazoprevir (GSR) in Hepatitis C Virus (HCV) GT2- and GT4- infected participants 65 years and older: an integrated analysis of twelve clinical trials. AASLD 2017.

2016

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2014

Everson GT, Tran TT, Towner WJ, Davis MN, Wyles D, **Nahass RG**, McNally J, Bainard DM et al. Safety and Efficacy of Treatment with the Interferon-free, Ribavirin-free combination of Sofosbuvir+GS-5816 for 12 Weeks in Treatment Naïve Patients with Genotype 1-6 HCV Infection. EASL, London UK 2014

Sulkowski M, Mallolas J, Pol S, Bouliere M, Gerstoft, Shibolet O, **Nahass R**, et al EFFICACY AND SAFETY OF THE ALL-ORAL REGIMEN, MK-5172/MK-8742 +/- RBV FOR 12 WEEKS IN GT1 HCV/HIV CO-INFECTED PATIENTS: THE C-WORTHY STUDY. EASL, London, UK. 2014

Sulkowski MS, Hezode C, Gerstoft J, Vierling JM, Mallolas J, Pol S, Kugelmas M, Murillo A, Weis N, **Nahass R**. Efficacy and safety of MK-5172 + MK-8742 ± ribavirin in HCV mono-infected and HIV/HCV co-infected treatment-naïve, non-cirrhotic patients with hepatitis C virus genotype 1 infection: The C-WORTHY study. 65th Meeting of the American Association for the Study of Liver Diseases, Boston MA, 2014

2013

High Efficacy and Safety of the All-Oral Combination Regimen, MK-5172/MK-8742 +/- RBV for 12 Weeks in HCV Genotype 1 Infected Patients: The C-WORTHY Study. AASLD 2013. Washington DC

K Patel, S Gordon, A Sheikh, Z Younes, J Ma, J McNally, DM Brainard, WT Symonds, JG McHutchison, **R Nahass**, EM Yoshida, H Reesink, D Nelson. Efficacy and Safety of Sofosbuvir in Patients According to Fibrosis Stage: An Analysis of Phase 3 Data. AASLD 2013. Washington, DC

Keyur Patel, Stuart C. Gordon, Aasim M. Sheikh, Ziad Younes, Julie Ma, John McNally, Diana M. Brainard, William T. Symonds, John G. McHutchison, **Nahass RG**, Eric M. Yoshida, Hendrik W. Reesink, Nelson DR. Efficacy and Safety of Sofosbuvir in Patients According to Fibrosis Stage: An Analysis of Phase 3 Data. 64th Meeting of the American Association for the Study of Liver Diseases, Washington DC, 2013.

Lawitz E, Vierling J, Murillo A, Kugelmas M, Gerstoft J, Winkle P, Balart L, Christensen P, Ghalib R, **Nahass RG**, Shaughnessy M, Sun X, Hwang P, Wahl J, Robertson M, Haber B. High safety and efficacy of all oral combination regimen, MK 5172/MK 8742 +/- RBV for 12 weeks in HCV infected patients the C Worthy study. 64th Meeting of the American Association for the Study of Liver Diseases, Washington DC, abstract 76, 2013.

Clinical Disease Management

2023

Brunetti L, Chapy H, **Nahass RG**, Moore R, Wassef A, Adler D, Yurkow E, Kagan L. Relationship between Body Composition and Serum Immunoglobulin Concentrations after Administration of Intravenous Immune Globulin—Preclinical and Clinical Evidence. *Pharmaceutics*. 2023; 15(2):510.
<https://doi.org/10.3390/pharmaceutics15020510>

Kohm K, Seneca K, Heineman D, Smith K, **Nahass RG**. Successful Treatment of *Cutibacterium acnes* Prosthetic Joint Infection with Single-Stage Exchange and Oral Antibiotics In Press OFID 2023

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San Filippo S, Crovetto B, Bucek J, **Nahass RG**, Milano M, MD, Brunetti L, PharmD, PhD, Comparative efficacy of early Covid-19 monoclonal antibody therapies: a retrospective analysis, *Open Forum Infectious Diseases*, 2022; ofac080, <https://doi.org/10.1093/ofid/ofac080>

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Nahass RG, Heineman D, Seneca KJ. Successful Treatment of *Cutibacterium acnes* (CA) Prosthetic Device Infection (PDI) with Oral Linezolid and Rifampin (LR). **Nahass RG**, Esquivel M, Smith K, Heinemann D, Seneca KH, ID Week 2021. Virtual

Seneca KH, Homer R, **Nahass RG**. Treating Hepatitis C Virus (HCV) in Young Active Drug Users is Possible. ID Week 2021. Virtual.

Crovetto B, SanFilippo S, Milano M, Bucek J, **Nahass R**, Brunetti L. Does time from COVID-19 symptom onset to administration of anti-spike protein monoclonal antibody predict response? ID Week 2021. Virtual

2020

J Goldman, D Lye, D Hui, K Marks, R Bruno, R Montejano, Spinner CD, Galli M, Ahn M, **Nahass RG**, et al Remdesivir for 5 or 10 days in patients with severe COVID-19. N Engl J Med (2020 May 27) DOI: 10.1056/NEJMoa2015301

Brunetti, L.; Diawara, O.; Tsai, A.; Firestein, B.L.; **Nahass, R.G.**; Poiani, G.; Schlesinger, N. Colchicine to Weather the Cytokine Storm in Hospitalized Patients with COVID-19. *J. Clin. Med.* 2020, 9, 2961

Hong TS, Gonzalez J, **Nahass RG**, Brunetti L. Impact of hydroxychloroquine on mortality in hospitalized patients with COVID-19: systemic review and meta-analysis. Pharmacy 2020

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Dalgard O, Litwin A, Dore G, Shibolet, Grebely J, **Nahass R**. Patient-reported outcomes among people receiving opioid agonist therapy and treatment for hepatitis C virus infection: results from CO-STAR. EASL 2020. London England.

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Sulkowski MS, Agarwal K, Fung S, yuen MF, Ma X, Lalezari J, Nguyen TT, Bae H, Schiff ER, ... **Nahass RG**. Continued Therapy with ABI-H0731 + NrtI Results in Sequential Reduction/Loss of HBV DNA, HBV RNA, HBeAg, HBcrAg and HBsAg in HBeAg-Positive Patients. AASLD 2019. Boston MA.

2018

Poster Presentation: Decreasing Clostridium difficile Infection – One Hospital's experience: SHEA 2018 conference, Portland, OR

2017

Wyles D, Wedemyer H, Ben-Ari Z, Gane EJ, Hansen J, Jacobsen IM, Laursen B, Lautkmeyer A, **Nahass RG** et al. Grazoprevir, ruzasvir, and uprifosbuvir for hepatitis C virus after NS5A treatment failure. *HEPATOLOGY* 2017;66:1794–1804)

Gane E, Metivier S, **Nahass RG**, et al. The Emergence of NS5B Resistance Associated Substitution S282T after Sofosbuvir-Based Treatment. *J Hepatology Communications* 2017;1:538–549.

2016

Miao B, Brunetti L, Bucek J, **Nahass RG**; Increased Cardiac Risk With Concomitant Levofloxacin (LVQ) and Amiodarone (AMIO) Therapy, Open Forum Infectious Diseases, Volume 3, Issue suppl_1, 1 December 2016, 1981, <https://doi.org/10.1093/ofid/ofw172.1529>

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Gane E, **Nahass RG**, Luketic V, et al. Efficacy of 12 or 18 weeks of grazoprevir plus elbasvir with ribavirin in treatment-naive, noncirrhotic HCV genotype 3-infected patients - J Hepatology 2015;62:S621

Foster GR, Pianko C, Cooper,A Brown A,Forton D, Nahass RG, Sofosbuvir + PegInterferon/Ribavirin for 12 weeks vs Sofosbuvir + Ribavirin for 16 or 24 weeks in Genotype 3 HCV infected patients and treatment-experienced cirrhotic pacientes with genotype 2 HCV: The BOSON study. Hepatology 2015;62:S259-S260

Pol S, Sulkowski M, Hassanein T, Gane E, Liu L, Mo H, Doeble B, Kanwar B, Brainard D, Subramanian G, Symonds W, Mchutchinson J, **Nahass RG**, Bennett m, Jacobson I. Sofosbuvir plus peginterferon and ribavirin in patients with genotype 1 HCV in whom prior therapy with direct-acting antivirals has failed. Hepatology (in Press).

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Nelson M, Rockstroh J, Mallolas J, Katlama C, **Nahass RG**, et al. High efficacy of grazoprevir/elbasvir among HCV GT1, GT4, or GT6 infected patients with HIV Co Infection. ID Week 2015 San Diego, CA

2014

Marcellin P, Manns MP, Janczewska E, Muir AJ, Wu X, Trenkle D, Pang P, Kanawar B, Dvory-Sobel H, Subramanian M, McHutchison J, **Nahass RG**, Gordon SC, Jacobson IM. Ledipasvir plus GS-9451 and PegIFN/RBV with high SVR12 rates in genotype 1 hepatitis C infection. NEJM 2014;370:1483-1493.

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Markowitz M, **Nahass RG**, et al. Switch from NNRTI plus TVD to STB Maintains HIV Suppression and is Well-Tolerated. American Conference for Treatment of HIV, Denver 2014

Nahass RG, Vierling JM, Safety of ABT-450/r/Ombitasvir + Dasabuvir With or Without Ribavirin in HCV Genotype 1-infected Patients, by Baseline Demographics. ID Week 2014, Philadelphia PA.

Gregory Everson G, Sims K, Thuluvath P, Schwartz H, Hassanein T, Lawitz E, Webster L, Bräu N, Desta T, Galati J, Ghalib R, Gitlin N, Han S, Hinestrosa F, Rodriguez-Torres M, **Nahass R**, et al. Daclatasvir in combination with asunaprevir and BMS-791325 for prior null responders with chronic HCV genotype 1 infection. 65th Meeting of the American Association for the Study of Liver Diseases, Boston MA, 2014

2013

Patel S, Segal R, Brunetti L, Kalabalik J, **Nahass RG**. Effectiveness of a pharmacist managed vancomycin dosing protocol in achieving therapeutic levels in a Community Medical Center. IDSA 2013. San Francisco CA

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Sukrut Dwivedi, Rohit Bhalla, Donald R. Hoover, Melvin P. Weinstein

Clinical Trials

2022

Janssen HLA, Hou J, Asselah T, Chan HLY, Zoulim F, Tanakan Y, Janczewska E, **Nahass RG**, et al. A Randomized Phase 2 Study (JADE) of the HBV Capsid Assembly Modulator NJ-56136379 With or Without a Nucleos(t)ide Analogue in Patients with Chronic Hepatitis B Infection. Gut 2023; 0:1-14. doi:10.1136/gutjnl-2022-328041In Press Gut

2020

Janssen H, Hou J, Asselah T, Chan H, Zoulim F, Tanaka Y, Janczewska E, **Nahass R**. Week 24 antiviral activity, safety and pharmacokinetics interim results of a phase 2 study of JNJ-56136379, a capsid assembly modulator, administered in combination with a nucleos(t)ide analogue in patients with chronic hepatitis B. EASL 2020, London, England.

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Jacobson IM, Lawitz E, Gane EJ, Willems BE, Ruane PJ, **Nahass RG**, et al. Efficacy of 8 Weeks of Sofosbuvir, Velpatasvir, and Voxilaprevir In Patients With Chronic HCV Infection: 2 Phase 3 Randomized Trials. Gastroenterology. 2017 Jul;153(1):113-122)

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Lawitz E, Reau N, Hinestrosa F, Rabinovitz M, Schiff E, Sheikh A, Younes Z, Herring R, Reddy R, Tran T, Bennett M, **Nahass R**, Yang J, An D, Dvory-Sobol H, Stamm L, Brainard D, McHutchison J, et al. Sofosbuvir, velpatasvir, and GS-9857 in patients with genotype 1 hepatitis C virus infection: an open-label, phase 2 trial, 2016 Jul 30. pii: S0016-5085(16)34835-1. doi: 10.1053/j.gastro.2016.07.039. [Epub ahead of print]

Lawitz E, Kowdley K, Curry M, Reau N, Nguyen M, Kwo P, Jacobson I, Tran T, **Nahass RG**. High efficacy of sofosbuvir/velpatasvir plus gs-9857 for 12 weeks in treatment-experienced genotype 1-6 hcv-infected patients, including those previously treated with direct-acting antivirals. EASL 2016 Barcelona Spain.

Nahass RG, et al. Sofosbuvir/Velpatasvir Plus GS-9857(100MG)for 6, 8, or 12 Weeks in Genotype 1-6 HCV-Infected Patients: An Integrated Analysis of Safety and Efficacy from Two Phase 2 Studies, ID Week 2016 New Orleans

Jacobson I, Asselah T, Nahass R, et al. A Randomized Phase 3 Trial of Sofosbuvir/Velpatasvir/Voxilaprevir for 8 Weeks Compared to Sofosbuvir/Velpatasvir for 12 Weeks in DAA-naïve Genotype 1-6 HCV-Infected Patients: The POLARIS-2 Study. AASLD, Boston 2016

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Foster GR, Pianko S, Cooper C, Brown A, Forton D, **Nahass RG**, et al. Sofosbuvir + PEGInterferon/Ribavirin for 12 weeks vs sofosbuvir + ribavirin for 16 or 24 weeks in genotype 3 HCV infected patients and treatment-experienced cirrhotic patients with genotype 2 HCV: The Boson Study. EASL, Vienna 2015.

Kwo P, Gitlin N, Nahass RG, et al. A phase 3, randomised, open-label study to evaluate the efficacy and safety of 12 and 8 weeks of simeprevir (smv) plus sofosbuvir (sof) in treatment-naïve and -experienced patients with chronic hcv genotype 1 infection without cirrhosis: optimist-1. EASL, Vienna, Austria 2015.

2014

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Epidemiology

2022

Dalgard O, Litwin AH, Shibolet O, Grebely J, **Nahass R**, et al. Health-related quality of life in people receiving opioid agonist treatment and treatment for hepatitis C virus infection, *Journal of Addictive Diseases* 2022. DOI: [10.1080/10550887.2022.2088978](https://doi.org/10.1080/10550887.2022.2088978)

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10/2022 Recurrence of Streptococcus agalactiae Bacteremia - Risk Factors and Complications. IDSA ID Week 2022

Serratia is a rare cause for Infective Endocarditis of the Tricuspid Valve. Samer I Talib, Nazar Raoof, Jonathan Kovacs, Natalia Raoof, Abdallah Khashan, Vasudev Daliparty, Srijana Poudel. *Open Forum Infectious Diseases* 9 (Supplement_2), ofac492. 1618, 2022

Castleman disease, a dilemma in HIV patients: A clinical presentation and diagnostic approach. Samer Talib, Nazar Raoof

Rachel Cowan, Shruti Varadarajan, **Abraham Wei**, Tanzila Salim, Michelle DallaPiazza. Microbial perils of the tropics: A case of cutaneous leishmaniasis in an immigrant from South America. *IDCases*. 2022 Dec 31;31:e01669.

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Bhatt PJ, Shiao S, Brunetti L, Xie Y, Solanki K, Khalid S, Mohayya S, Au PH, Pham C, Uprety P, **Nahass R**, Narayanan N. Risk Factors and Outcomes of Hospitalized Patients with Severe COVID-19 and Secondary Bloodstream Infections: A Multicenter, Case-Control Study. *Clin Infect Dis*. 2020 Nov 20:ciaa1748. doi: 10.1093/cid/ciaa1748. Epub ahead of print. PMID: 33216875; PMCID: PMC7717183.

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Parronchi CJ, Doshi V, **Nahass RG**. Histoplasmosis endophthalmitis – case report and review. Open Infect Dis J 2018;10:71-75.

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Akyar E, Mora N, Luke A, Layden J, Phillips R, Agyarko-Poku T, Owusu, Helena D **Nahass RG**. Risk Factors for Hepatitis C in Western Africa: An Observational Study in an STI Clinic. ID Week 2018. San Francisco

2017

Prevalence of Infection with Multiple Strains of Hepatitis C Virus (HCV) in Patients Enrolled in HCV Clinical Trials EASL Amsterdam 2017

Hepatitis C virus (HCV) reinfection and injection risk behavior following elbasvir/grazoprevir (EBR/GZR) treatment in patients on opioid agonist therapy (OAT): Co-STAR three-year follow-up study. DDW

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