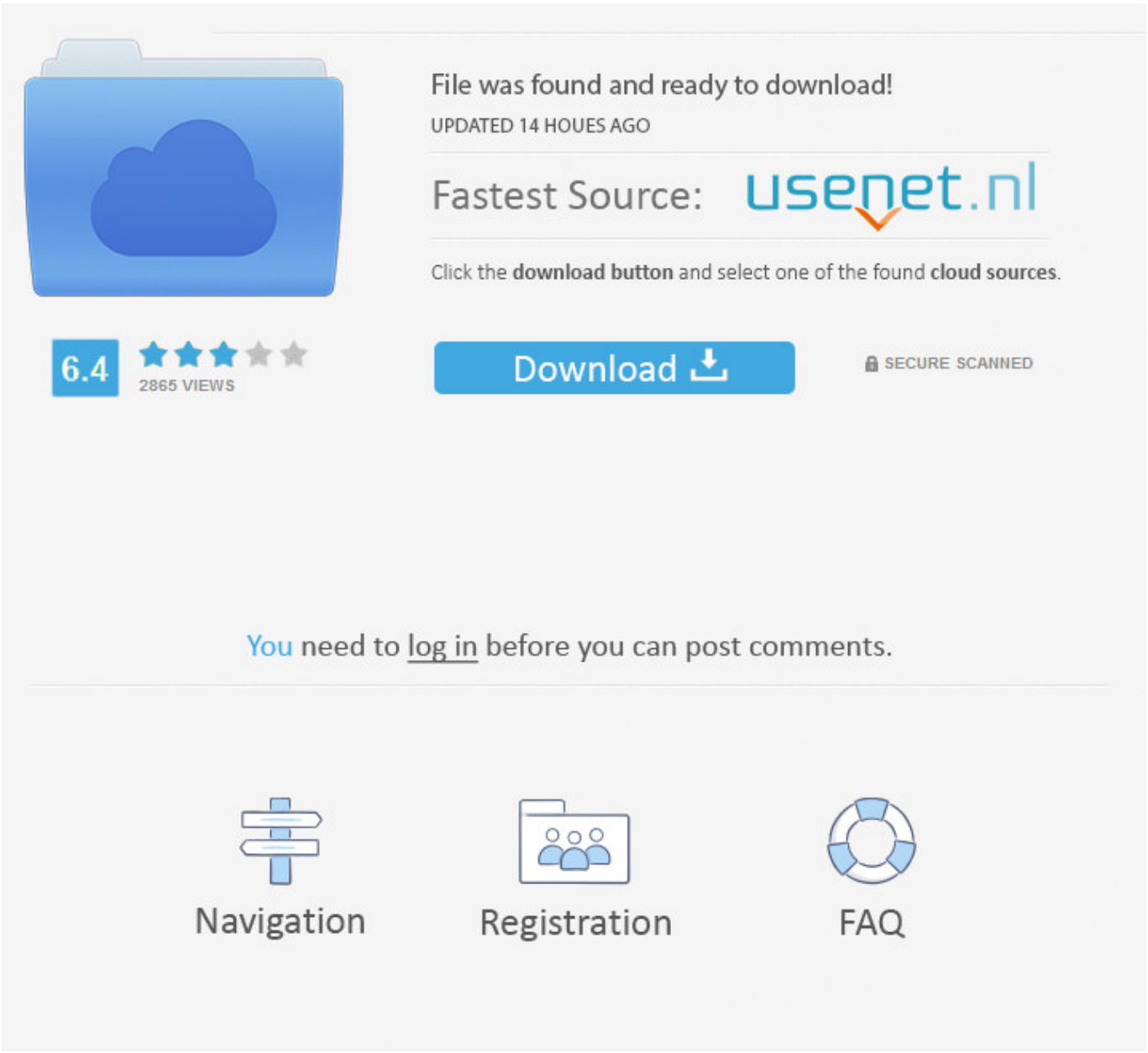

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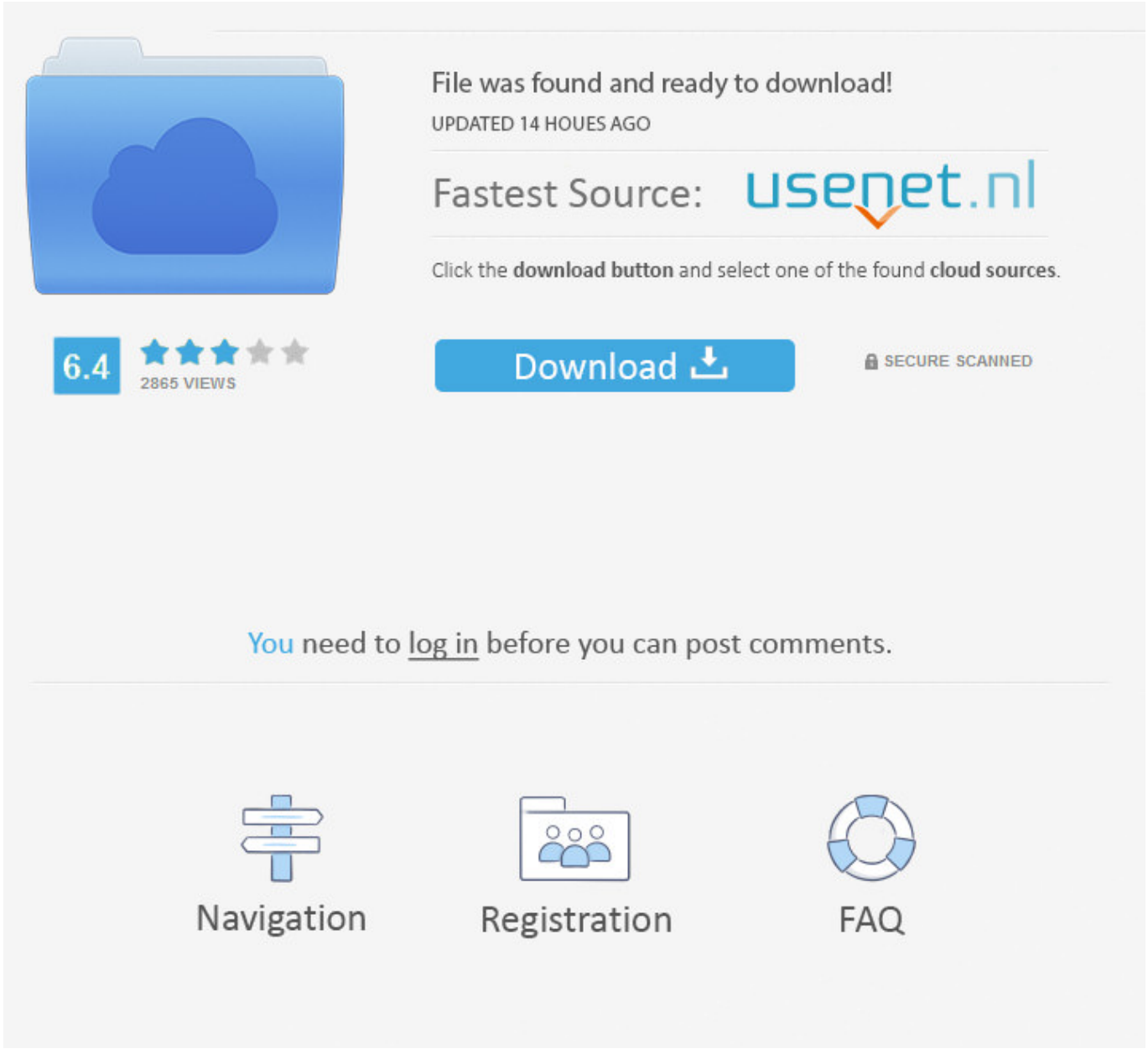
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Patients were randomly assigned to receive the NS5A inhibitor ledipasvir and the nucleotide polymerase inhibitor sofosbuvir in a once-daily, fixed-dose combination tablet for 12 weeks, ledipasvir–sofosbuvir plus ribavirin for 12 weeks, ledipasvir–sofosbuvir for 24 weeks, or ledipasvir–sofosbuvir plus ribavirin for 24 weeks.. The rates of sustained virologic response were high in all treatment groups: 94% (95% confidence interval [CI], 87 to 97) in the group that received 12 weeks of ledipasvir–sofosbuvir; 96% (95% CI, 91 to 99) in the group that received 12 weeks of ledipasvir–sofosbuvir and ribavirin; 99% (95% CI, 95 to 100) in the group that received 24 weeks of ledipasvir–sofosbuvir; and 99% (95% CI, 95 to 100) in the group that received 24 weeks of ledipasvir–sofosbuvir and ribavirin.. Among the estimated 170 million people in the world who have chronic hepatitis C virus (HCV) infection, approximately 60% have the genotype 1 strain of the virus.. The only interferon-free option currently approved for HCV genotype 1 infection is 24 weeks of sofosbuvir and ribavirin for patients who are ineligible to receive interferon because of absolute or relative contraindications.. Muir, M D , Mark Sulkowski, M D , and Paul Kwo, M D , for the ION-2 Investigators N Engl J Med 2014; 370:1483-1493 DOI: 10.. 1056/NEJMoa1316366 open through April 19, 2014 Methods We conducted a phase 3, randomized, open-label study involving patients infected with HCV genotype 1 who had not had a sustained virologic response after treatment with peginterferon and ribavirin, with or without a protease inhibitor.. Original Article Ledipasvir and Sofosbuvir for Previously Treated HCV Genotype 1 Infection Nezam Afdhal, M. 34bbb28f04
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