Reply to: “Treatment of veterans with hepatitis C in the United States Department of Veterans Affairs”

To the Editor:
I would like to thank Dr. Ross.

(1) Dr. Ross does not state how many veterans with HCV are currently receiving care at the Department of Veterans Affairs (VA). In 2008, VHA clinicians cared for over 147,000 veterans with chronic HCV [1]. Treating 4500 patients with HCV in 20 months is only 225 patients per month. The VA is currently treating less than 2% of infected veterans per year with boceprevir and telaprevir. It will take more than fifty years for the VA to treat all of their HCV infected patients. Evidence based care of an infectious disease is cure of the infection not the development of integrated models to address comorbidities. If 98% of patients with a curable infection are not treated each year, the VA’s response is inadequate.

(2) The VA does a better job with the human immunodeficiency virus (HIV) treating 78% of veterans [2]. The number of patients on antiviral therapy clearly indicates that HIV is a high priority for the VA while HCV treatment is not.

(3) Telaprevir is not available as a non-formulary drug at the Louisville VA. Boceprevir is on the formulary there.

(4) More than 1800 patients with HCV antibodies have been identified at the Louisville VA over 19 years. They had multiple physicians providing care.

(5) $100 million for antiviral therapy over 20 months is $5 million per month. This is clearly inadequate to treat 147,000 veterans with hepatitis C. This is why legislation should be passed so that all veterans with HCV immediately pre-qualify for their choice of Medicaid or Medicare. They could then obtain antiviral therapy in the private sector instead of waiting for the VA to treat 2% of them each year. Now, many are trapped in the VA system while their curable infection progresses to liver cancer, liver failure and death.

Conflict of interest
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References

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Use of TNFα antagonists in refractory AIH: Revealing the unforeseen

To the Editor:
We read with considerable interest the paper by Weiler-Normann et al. in the Journal of Hepatology [1], which reported promising results regarding the use of infliximab as a therapeutic option in difficult-to-treat patients with autoimmune hepatitis (AIH). Although the exact role of tumor necrosis factor α (TNFα) in the pathogenesis of AIH has not been elucidated yet, very recently, it has been shown in a mouse model of fatal AIH that TNFα is essential in the induction of AIH through upregulation of hepatic CCL20 expression, which allows migration of dysregulated splenic T cells [2]. As a consequence, the efficacy of anti-TNFα therapy in AIH could have a pathophysiological basis, taking also into account that TNFα is produced in large amounts in the liver, in the context of AIH, by macrophages, CD8+ T cells and possibly Th17 lymphocytes [3]. However, it is already known from the use of anti-TNFα treatment in various autoimmune diseases that anti-TNFα can also be immunogenic, with development of either autoantibodies or true autoimmune diseases, making infliximab a two-edged sword [4].

The induction of AIH is one of the examples of the latter “therapeutic paradox” during anti-TNFα treatment. In fact, the hepatic flare reported in the second patient of the study of Weiler-Normann et al. [1] could have been such an effect, especially if it was combined with an IgG increase. Here, we are reporting an additional case of a 30-year old female patient admitted to our department because of infliximab induced AIH, in an attempt to further emphasize the “two-sided” face of anti-TNFα treatment. Our patient had a history of refractory psoriasis treated with infliximab (5 mg/kg at week 0, 2, 6 and then every 8 weeks by intravenous infusion) and presented to our department because of infliximab induced AIH, in an attempt to give a simplified score of 7, confirming the diagnosis of definite AIH [5]. Apart from infliximab withdrawal, the patient was treated after an informed consent, according to our experience and