

ORAL PRESENTATION

Open Access

O314. Efficacy and safety of peginterferon alfa-2a + RBV in cHCV/HIV- vs cHCV-infected patients: interim analysis of a multicenter German cohort

A Baumgarten¹, T Lutz², P Kreckel³, E Wellmann⁴, U Alshuth⁴, S Mauss⁵, J Rockstroh^{6*}

From Tenth International Congress on Drug Therapy in HIV Infection
 Glasgow, UK. 7-11 November 2010

Purpose of the study

Eradicating cHCV is necessary for the subsequent management of patients with HIV and every HIV/HCV co-infected patient should be considered for treatment. We describe differences between cHCV mono-infected and cHCV/HIV co-infected patients in baseline factors and outcome of cHCV-treatment with peginterferon alfa 2a + RBV in the worldwide largest cHCV cohort.

Methods

Noninterventional prospective multicenter German cohort, started January 2008 and still recruiting. Interim analysis of cHCV patients, stratified for cHCV mono-infection and cHCV/HIV co-infection. The results are

based on a cross-sectional analysis of all available data in April 2010.

Results

This interim analysis included 5.390 patients, who received HCV-treatment. 397 were cHCV/HIV co-infected (CI) and 4.993 cHCV mono-infected (MI). Main baseline- characteristics: 85.9% were GT1/4/5/6 patients in the CI-Group and 63.4% in the MI-Group, age was 41.0 (CI), 42.0 (MI) yrs, 89.7 (CI), 62.9 (MI)% were male, BMI was 22.8 (CI), 24.9 (MI) kg/m², naïve/relapse/non-responder/re-infection: 86.4/3.8/4.5/5.3(CI), 88.0/6.1/5.3/0.6 (MI) %, source of infection (>1 answer possible): iv drug use 25.2(CI), 44.9(MI) %, sexual trans-

	cHCV/HIV			cHCV		
	Overall	GT 1/4/5/6	GT 2/3	Overall	GT 1/4/5/6	GT 2/3
RVR % (n)	37.1 (111/299)	33.6 (86/256)	59.6 (25/42)	46 (1773/3858)	30.4 (789/2597)	78.1 (976/1250)
EVR % (n)	79.3 (207/261)	78.6 (173/220)	82.9 (34/41)	84.3 (3182/3775)	81.4 (2021/2484)	90.0 (1155/1284)

Figure 1

⁶Universitätsklinikum Bonn, Medizinische Klinik Poliklinik 1, Bonn, Germany
 Full list of author information is available at the end of the article

mission 60.7 (CI), 4.1(MI)%, other 8.0 (CI), 24.0 (MI), unknown 13.1(CI), 33.0 (MI)%. 86.4 % of the co-infected patients received antiretroviral HIV-treatment (ART), 66.2 % of them had an HIV-RNA level below 50 copies/mL, median CD4-cells/ μ count was 502. From those patients who finished treatment, 52.9% of the CI-Group and 67.7% of the MI-Group completed the planned course. Reasons for discontinuation (>1 answer possible) were non-response (59.3% in CI, 45.5 % in MI) and patient request (24.7% in CI, 14.5% in MI). Other reasons were tolerability (11.1% in CI, 12.0% in MI) and compliance issues (12.3% in CI, 10.5% in MI). Treatment response rates, stratified by genotypes, were already available regarding RVR and EVR (see Figure 1)

Conclusions

In this preliminary analysis HCV/HIV co-infected patients seemed to respond similar according to RVR and EVR in GT1/4/5/6 as HCV-mono infected patients on HCV-treatment. Treatment discontinuation due to non-response and patient request was much more common in the co-infected group. Other reasons for discontinuation like tolerability and compliance are equal in both arms. A more detailed analysis, in particular the influence of the HIV-ART on HCV-therapy outcome, may help interpreting these data. An updated analysis will be presented.

Author details

¹Privat praxis Dupke/Carganico/Baumgarten, Berlin, Germany. ²Infektiologikum Frankfurt, Frankfurt, Germany. ³Praxis Koeppel/Kreckel, Berlin, Germany. ⁴Roche Pharma AG, Grenzach-Wyhlen, Germany. ⁵Center for HIV and Hepatogastroenterology, Düsseldorf, Germany. ⁶Universitätsklinikum Bonn, Medizinische Klinik Poliklinik 1, Bonn, Germany.

Published: 8 November 2010

doi:10.1186/1758-2652-13-S4-O32

Cite this article as: Baumgarten *et al.*: O314. Efficacy and safety of peginterferon alfa-2a + RBV in cHCV/HIV- vs cHCV-infected patients: interim analysis of a multicenter German cohort. *Journal of the International AIDS Society* 2010 **13**(Suppl 4):O32.

**Submit your next manuscript to BioMed Central
and take full advantage of:**

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

