

HIV/HCV co-infection across Europe

L Peters¹, K Laut¹, C Leen², K Falconer³, T Trofimora⁴, D Paduto⁵, J Gatell⁶, A Rauch⁷, K Lacombe⁸, A Mocroft⁹, JD Lundgren¹, JK Rockstroh¹⁰ for EuroSIDA in EuroCoord

¹CHIP, Dept. of Infectious Diseases, Rigshospitalet, University of Copenhagen, Denmark. ²Western General Hospital, Edinburgh, UK. ³Karolinska Hospital, Stockholm, Sweden. ⁴Novgorod Centre for AIDS, Novgorod, Russia. ⁵Regional AIDS Centre, Svetlogorsk, Belarus. ⁶Hospital Clinic Universitari de Barcelona, Barcelona, Spain. ⁷Department of Infectious Diseases, Inselspital, Bern University Hospital, University of Bern, Switzerland. ⁸Hospital Saint Antoine, Paris, France. ⁹Dept Infection and Population Health, University College London, London UK. ¹⁰University of Bonn, Germany

Objectives

A large observational study is needed to monitor the uptake, efficacy, safety and clinical outcomes of direct-acting antivirals (DAAs) in unselected HIV/HCV co-infected patients in a real-life heterogeneous setting. In 2014, the EuroSIDA study began enrolment of a new cohort of 5,000 anti-HCV positive HIV patients from all European regions (South, Central West, North, Central East, East) plus Argentina (**Figure 1**).

The objective of this study is to describe the characteristics of patients co-infected with HIV and hepatitis C (HCV) prior to the introduction of effective DAAs.

Methods

Descriptive statistics were used to summarise characteristics of patients with HIV/HCV at baseline (latest of anti-HCV+, study recruitment or 1/1/2013).

Results

There were considerable differences among the 4608 persons included to date followed in different regions (**Table1**). Compared with patients from other regions, patients from East were more likely to be injection drug users, younger, had lower CD4 and higher CD4 nadir, were less likely to be on cART, and among those on cART, less likely to have a viral load<500 copies/ml. The median CD4 cell count at starting cART (available for 3452) was 205 cells/mm³ (IQR 95-315), and was ≤350 cells/mm³ for 80.5%, ranging from 73.3% in South to 88.2% in East.

Table 2 describes the HCV characteristics of the cohort. Information on HCV genotype was available for 2534 (55.0%); GT1 (n=1360, 53.7%) was the most common. Of 3283 (71.3%) with HCV-RNA information, 2010 (61.2%) were HCV-RNA-positive, and 1287/2866 (44.6%) with quantified HCV-RNA had HCV RNA ≥500,000 IU/ml. 757 (16.4%) had ever been exposed to anti-HCV therapy (DAA, interferon), but only 35 (0.8%) to DAA. Of the 4069 (88.3%) with information on fibrosis, 3444 (84.6%) had F0/F1 fibrosis, and 125 (3.1%), 104 (2.6%) and 396 (9.7%) had F2, F3 or F4 fibrosis respectively. ≥F2 fibrosis was significantly more common in South (17.6%) and East (16.1%).

Discussion

This large cohort of HIV/HCV co-infected persons was largely naïve to DAA and had a low proportion of advanced fibrosis; persons on cART had good HIV virological control although cART was started at a low CD4 count. All patients will be followed up for at least five years or until the time of death or loss to follow up, and the cohort will be important for investigating outcomes associated with DAAs when introduced into a heterogeneous clinic population.

Figure 1

Countries participating in EuroSIDA

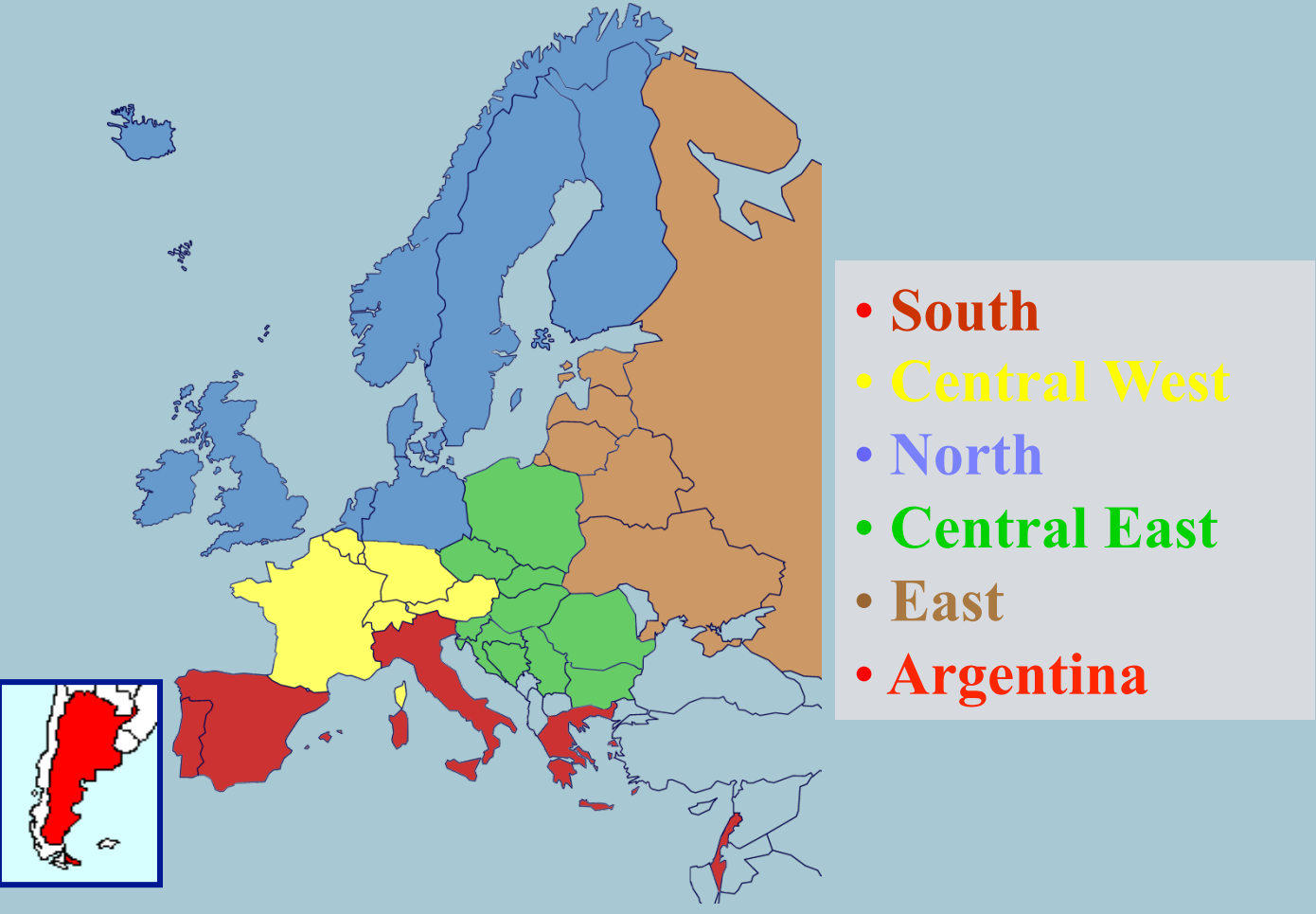


Table 1

Baseline characteristics of 4608 HIV/HCV co-infected individuals							
	All	S	W	N	CE	E	Argentina
N (%)	4608	1183 (25.7)	573 (12.4)	624 (13.5)	854 (18.5)	1266 (27.5)	108 (2.3)
Male (%)	69.9	72.5	74.4	74.7	76.5	63.0	65.7
Injection drug use (%)	61.6	67.2	47.3	47.9	63.7	70.1	38.9
HBV co-infected (%)	5.4	4.1	7.3	4.7	6.0	5.7	6.5
On cART (%)	81.2	82.4	79.1	85.4	87.0	74.2	93.5
On cART, VL<500 (%)	89.5	96.8	95.8	95.7	91.1	74.8	89.4
CD4 (median cells/mm ³)	50.8	588	588	520	532	412	451
Nadir CD4 (median cells/mm ³)	16.4	165	146	152	175	179	139
Age (median years)	4.5	50	51	50	40	36	46
Baseline (median mm/y)	10/13	10/13	10/13	07/14	10/13	10/13	10/13

Table 2

Baseline characteristics of 4608 HIV/HCV co-infected individuals							
		S	W	N	CE	E	Argentina
N (%)	4608	1183 (25.7)	573 (12.4)	624 (13.5)	854 (18.5)	1266 (27.5)	108 (2.3)
HCV-RNA negative (%)	38.8	33.6	43.2	43.6	37.5	41.1	38.0
HCV-RNA >500,000 IU/ml (%)	44.6	50.5	39.3	43.6	40.2	45.6	35.9
HCV genotype 1 (%)	53.7	53.9	57.8	58.2	41.3	57.6	64.2
HCV genotype 2 (%)	2.8	1.7	2.7	6.8	1.1	2.5	3.9
HCV genotype 3 (%)	29.0	26.5	24.7	25.6	32.6	39.9	23.1
HCV genotype 4 (%)	14.4	17.9	14.5	8.7	25.0	0	3.9
≥F2 fibrosis (%)	15.4	17.6	13.0	15.6	13.9	16.1	6.7
Prior HCV treatment (%)	16.4	20.4	26.2	14.9	15.7	9.1	22.2
Prior DAA treatment (%)	0.8	1.4	1.2	1.0	0.7	0	0

The EuroSIDA Study Group

The multi-centre study group, EuroSIDA (national coordinators in parentheses).

Argentina: M Lasso, Hospital JM Ramos Mejia, Buenos Aires. Austria: (N Vetter), Pulmologisches Zentrum der Stadt Wien, Vienna. R Zangerle, Medical University Innsbruck, Innsbruck. Belarus: (I Karпов), A Vassilenko, Belarus State Medical University, Minsk. VM Mitsura, Gomel State Medical University, Gomel. D Paduto, Regional AIDS Centre, Svetlogorsk. Belgium: (N Clumeck), S De Wit, M Delforge, Saint-Pierre Hospital, Brussels. E Florenco, Institute of Tropical Medicine, Antwerp. L Vandecasteele, University Ziekenhuis Gent, Gent. Bosnia-Herzegovina: (V Hadzimehmedovic), Klinicki Centar Univerziteta Sarajeva, Sarajevo. Bulgaria: (K Kostov), Infectious Diseases Hospital, Sofia. Croatia: (J Begovic), University Hospital of Infectious Diseases, Zagreb. Czech Republic: (L Machala), D Jlich, Faculty Hospital Bulovka, Prague. D Sedlacek, Charles University Hospital, Pilsen. Denmark: G Kronborg, T Benfield, Hvidovre Hospital, Copenhagen. J Gerstoft, T Kaldesen, A-B E Hansen, Rigshospitalet, Copenhagen. C Pedersen, NP Møller, Odense University Hospital, Odense. L Ostergaard, Skejby Hospital, Århus. U B Dingsø, Roskilde Hospital, Roskilde. L N Nielsen, Hillerød Hospital, Hillerød. Estonia: (K Zimar), West-Tallinn Central Hospital, Tallinn. Jelena Smrd, Nakhosakond Saelek, Kottla-Järve. Finland: (M Ristola), I Aho, Helsinki University Central Hospital, Helsinki. France: (C Katlama), Hôpital de la Pitié-Salpêtrière, Paris. J-P Vaire, Hôtel-Dieu, Paris. P-M Girard, Hôpital Saint-Antoine, Paris. L Cotte, Hôpital de la Croix Rousse, Lyon. C Pradier, E Fontas, Hôpital de la Pitié, Nice. F Dabis, D Neau, Unité INSERM, Bordeaux. C Duvalier, Hôpital Necker-Enfants Malades, Paris. Germany: (J Rockstroh), Universitäts Klinik Bonn. R Schmidt, Medizinische Hochschule Hannover. J van Lunzen, O Degen, University Medical Center Hamburg-Eppendorf. Infectious Diseases Unit, Hamburg. HJ Stellbrink, IPM Study Center, Hamburg. C Stefan, JW Goethe University Hospital, Frankfurt. J Bogner, Medizinische Poliklinik, Munich. G. Felsenberg, Universität Köln, Cologne. Georgia: (N Chikharishvili), Infectious Diseases, AIDS & Clinical Immunology Research Center, Tbilisi. Greece: (J Kormlidis), P Gargalianos, G Xylomenos, P Lourdas, Athens General Hospital. H Sambatakou, Ipokration General Hospital, Athens. Hungary: (D Barhegyi), Szent László Hospital, Budapest. Iceland: (M Gottfredsson), Landspítali University Hospital, Reykjavik. Ireland: (F Mulcahy), St. James's Hospital, Dublin. Israel: (I Nisli), D Turner, M Burke, Ichilov Hospital, Tel Aviv. E Shahar, G Hasson, Raminan Medical Center, Haifa. H Elina, M Hazzou, Hadassah University Hospital, Jerusalem. ZM Shoshan, AIDS Center (Neve Or), Jerusalem. Italy: (A D'Amico Montforte), Istituto Di Clinica Malattie Infettive e Tropicale, Milan. R Esposito, I Mazzeo, C Mussini, Università Modena, Modena. F Mazzotta, A Gabbuti, Ospedale S Maria Annunziata, Firenze. V Vullo, M Lichtner, Ospedale di Roma la Sapienza, Rome. M Zaccarelli, A Antonini, R Acciarini, G D'Onofrio, Istituto Nazionale Malattie Infettive Lazzaro Spallanzani, Rome. A Lazzaro, A Castagna, N Gianotti, Ospedale San Raffaele, Milan. M Galli, A Ridolfi, Osp. L. Sacco, Milan. Latvia: (B Rozentale), Infectology Centre of Latvia, Riga. Lithuania: (V Uzdavicius), Vilnius University Hospital Santariskiu Klinikos, Vilnius. R Matulionyte, Center of Infectious Diseases, Vilnius University Hospital Santariskiu Klinikos, Vilnius. Luxembourg: (T Staub), R Hemmer, Centre Hospitalier, Luxembourg. Netherlands: (P Rees), Academisch Medisch Centrum bij de Universiteit van Amsterdam, Amsterdam. Norway: (V Ormaasen), A Haueland, J Bruun, Ullevål Hospital, Oslo. Poland: (B Kryszewski), J Gasiorowski, M Ingot, Medical University, Wrocław. A Horban, E Bakowska, Centrum Diagnostyki i Terapii AIDS, Warszawa. A Grzeszczuk, R Flisak, Medical University, Białystok. M Parczewski, M Pryka, K Maciejewska, Medical University, Szczecin. M Benikow, E Mulanska, Otorok Diagnostyki i Terapii AIDS, Chorzów. J Smatcz, M Gensberg, Medical University, Gdańsk. E Jablonowska, E Malolepsza, K Wojcik, Wojewódzki Szpital Specjalistyczny, Łódź. I Moser-Liesewska, Poznan University of Medical Sciences, Poznan. Portugal: (M Doraana), L Caldeira, Hospital Santa Maria, Lisbon. K Mansinho, Hospital de Egas Moniz, Lisbon. F Maltzer, Hospital Curry Cabral, Lisbon. Romania: (R Radoi), C Oprea, Spitalul de Boli Infectioase si Tropicale, Dr. Victor Babes, Bucuresti. Russia: (A Raikmanova), Medical Academy Borok Hospital, St Petersburg. A Raikmanova, St Petersburg AIDS Centre, St Petersburg. T Trofimova, Novgorod Centre for AIDS, Novgorod. I Khromova, Centre for HIV/AIDS & Infectious Diseases, Kaliningrad. E Kuznetsova, Nizhny Novgorod Scientific and Research Institute, Nizhny Novgorod. Serbia: (D Jevlovic), The Institute for Infectious and Tropical Diseases, Belgrade. Slovakia: A Shumra, D Stanekova, Drier Hospital, Bratislava. Slovenia: (J Tomcovic), University Clinical Centre Ljubljana, Ljubljana. Spain: (JM Gatell), JM Miró, Hospital Clinic Universitari de Barcelona, Barcelona. S Moreno, J M Rodriguez, Hospital Ramon y Cajal, Madrid. B Cotel, A Joui, R Paredes, C Tural, J Puig, J Bravo, Hospital Germans Trias i Pujol, Badalona. P Domingo, M Gutierrez, G Mateo, MA Sambasat, Hospital Sant Pau, Barcelona. JM Laporte, Hospital Universitario de Alava, Vitoria-Gasteiz. Sweden: (A Blighul), Verneis-Söderström Hospital, Stockholm. L Farnham, Mairno University Hospital, Mairno. K Falconer, A Thaimie, A Sonnenberg, Karolinska University Hospital, Stockholm. Switzerland: (B Ledergerber), R Weber, University Hospital Zurich. M Cavassini, University Hospital Lausanne. A Calmy, University Hospital Geneva. H Furrer, University Hospital Bern. M Battegay, L Eizi, University Hospital Basel. P Schmidt, Cantonal Hospital St. Gallen. Ukraine: (E Kravchenko), N Chentsova (deceased), Kiev Centre for AIDS, Kiev. V Frolov, G Krut'ya, I Baskakov, Luhansk State Medical University, Luhansk. A Kuznetsova, Kharkov State Medical University, Kharkov. G Kyselova, Crimean Republican AIDS centre, Simferopol. United Kingdom: (B Gazzard), St. Stephen's Clinic, Chelsea and Westminster Hospital, London. AM Johnson, E Simons, S Edwards, Mortimer Market Centre, London. A Phillips, A Mocroft, Royal Free and University College Medical School, London. Royal Free Campus, London. C O'Brien, Royal Free Hospital, London. J Weber, C Soutter, Imperial College School of Medicine at St. Mary's, London. A Clarke, Royal Sussex County Hospital, Brighton. C Leen, Western General Hospital, Edinburgh. The following centers have previously contributed data to EuroSIDA: Bernhard Nocht Institut für Tropenmedizin, Hamburg, Germany; 1st I.K.A Hospital of Athens, Athens, Greece; Ospedale Runiti, Divisione Malattie Infettive, Bergamo, Italy; Ospedale di Bolzano, Divisione Malattie Infettive, Bolzano, Italy; Ospedale Cetruglio, II Divisione Malattie Infettive, Napoli, Italy; Hospital Carlos III, Departamento de Enfermedades Infecciosas, Madrid, Spain; Odessa Region AIDS Center, Odessa, Ukraine.

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