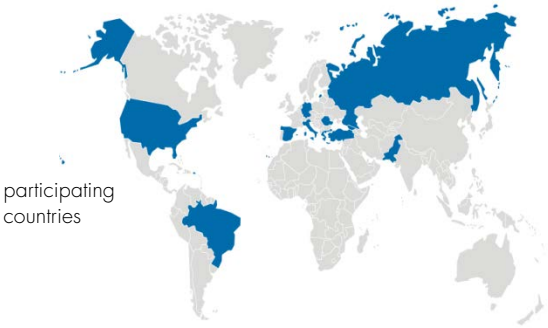
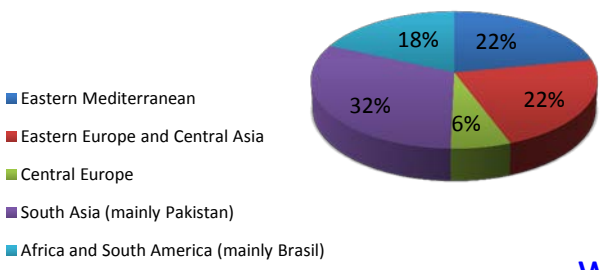
Hepatitis Delta International Network

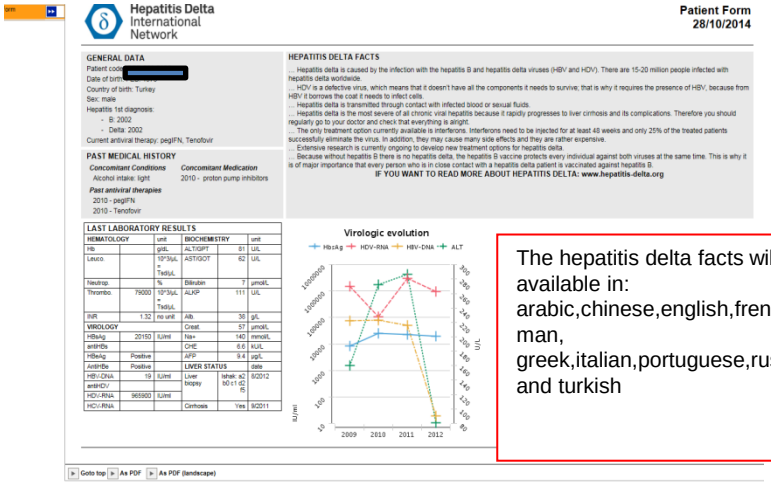


The primary aim of the HDIN registry is to establish a large global data base of patients chronically infected with HDV to better define the course of hepatitis delta and response to antiviral therapy in the context of different HDV and HBV genotypes and diverse host genetic and environmental backgrounds. A structured eCRF optimized for hepatitis delta was implemented and 12 centers world-wide are participating.
Over 800 patients are now documented in this registry.

Patients were divided according to the country of birth into Eastern Mediterranean (EM), Eastern Europe and Central Asia (EE/CA),Central and Southern Europe (CE), South Asian (SA mainly Pakistan) and Africa/South America (AF/SA)



THE HDIN-Patient Form : the Patient Form will be established automatically by the online system. This document will provide the **most important data regarding the patients´ clinical course, facts and advice about chronic hepatitis D.** This is supposed to give the patient a better understanding of the disease and to help him pay good attention to his health in order to prevent or at least delay sequel complications. Furthermore it will contain information to facilitate optimal treatment by any other doctors.


Hepatitis Delta International Network

Patient Form
28/10/2014

GENERAL DATA

HEPATITIS DELTA FACTS

PAST MEDICAL HISTORY

LAST LABORATORY RESULTS

Virologic evolution

The hepatitis delta facts will be available in: arabic,chinese,english,french,german, greek,italian,portuguese,russian and turkish

www.hepatitis-delta.org

ClinicalTrials.gov Identifier:
NCT02375906