

# Identification and management of persons with chronic viral hepatitis in Europe

Country sessions

**BELGIUM**

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BASL

# Burden of disease: hepatitis B and C

- **In Belgium:**
- Passive surveillance through mandatory notification of hep B (Royal Decree 1.3.1971) and C cases (as of 2009 no longer in all regions).
- Few epidemiological data

# 1. Hepatitis C

# Burden of disease: hepatitis C

- **Cross sectional study in Belgium in 2003:**

HCV Ab+ : 0.12% (0.87% + in 1993 serum)

HBsAg+: 0.66% (0.70% + in 1993 serum)

- Saliva test send to n=6000 => 30,6% responded => 3/10 000 people in Flanders represented
- At risk population probably underrepresented

S. Quolin. Eur J Epidemiol 2007; 22:195 - 202

# Burden of disease: hepatitis C

S. Demaeght. Acta Gastroenterologica Belgica 2008;71: 19-23

- **HOSPITALS : Cross sectional study in Belgium in 2003**
- Prospective observatory in Belgium in 2003-2004, new consecutive anti-HCV+ patients in 9 hospitals (2° and 3° health care centers)  
N= 318      55% Men      median age 45y [11-87y]
- Results:
  - 87% PCR HCV +
  - 66% abN transaminases
  - Identified Risk factors:
    - IDU: 27% (underrepresented)
    - Blood transfusion: 23%
    - Invasive medical procedures: 11%
    - Unkown: 23%
  - Stage: 43% F0F1; 35% F2; 22% F3F4
  - Genotypes: G1: 59%; G3: 19%; G4: 14%
  - QOL: 61 ± 31 (0-100)

# Burden of disease: hepatitis C

- **HCV in IDU**
- IDU recruited at tt centers all over B\*: n = 147 antiHCV+
- 70% of all IDU are antiHCV + => 67% is PCR +
- G1: 38%; G2: 2%; **G3: 49%**; G4: 9%
  
- IDU recruited at 2 Flemish Centres (Antw/Limburg)° n = 155
- G3a remains Steady state; G1a becomes predominant => harmful on long term

G. Robaey. *Acta Gastroenterologica Belgica* 2002; 65:99-100

Micalessi\* *J Med Virol* 2008; 80: 640-645 . C. Matei ° *J Viral Hepat* 2008; 15:399-408.

# Burden of disease: hepatitis C

- **HCV/HIV coinfection: Belgian study**
- 9 centres, open, non controlled, prospective trial between 9/2001-dec/2003
- 10-15% of all HIV+ pts are antiHCV +
- N = 37

56% have  $\geq$  F3

65% receive HAART

SVR after 52w pegIFNa2b/Riba: G1/4: 13% G2/3: 47%

30% stopped tt because of side effects

# Screening programme HCV

## Implementation

- **Since 1990**: screening of blood and blood products/tissues for HCV
- **Universal precautions** in healthcare setting
- Screening of **pregnant** women
- Opportunistic screening by GP and gastroenterologists
- National screenings days HCV: 2001 and 2004, sponsored by medical company SP



# Treatment strategies in B: HCV

- **For G1, G4, G5 and G6:**

- Liver biopsy required
- 48w peg IFN alfa 2a or 2b + ribavirin
- At 12w: PCR should be  $\geq 2\log$  ↓
- Price:  $1000\text{€} \times 11 + 700\text{€} \times 11 = \pm 18\,700\text{€}/48\text{w}$
- RIZIV/INAMI pays for it: 98%
- The patient pays  $\pm 35$  euro/month =  $385\text{€}/48\text{w}$
- FU: w2-4-6-8, then every 4-6w, EOT, SVR  
if F3-F4: US +  $\alpha$ FP every 6months

# Treatment strategies: HCV

- **For G2, G3:**

- No liver biopsy required
- 24w peg IFN alfa 2a or 2b + ribavirin
- Price:  $1000\text{€} \times 6 + 700\text{€} \times 6 = \pm 10\,200\text{€}/24\text{w}$
- RIZIV/INAMI pays for it: 98%
- The patient pays  $\pm 35$  euro/month =  $210\text{€}/24\text{w}$
- FU: w2-4-6-8, then every 4-6w, EOT, SVR  
if F3-F4: US +  $\alpha$ FP every 6months

# Treatment strategies: HCV/HIV coinfection

- **For all genotypes:**

- Liver biopsy required
- 48w peg IFN alfa 2a or 2b + ribavirin
- Price:  $1000\text{€} \times 11 + 700\text{€} \times 11 = \pm 18\,700\text{€}/48\text{w}$
- RIZIV/INAMI pays for it: 98%
- The patient pays  $\pm 35$  euro/month =  $385\text{€}/48\text{w}$
- FU: w2-4-6-8, then every 4-6w, EOT, SVR  
if F3-F4: US +  $\alpha$ FP every 6months

## Treatment strategies:

### Health economic model HCV in Belgium

- 29% of all liver transplants due to HCV in B
- Immediate treatment of HCV with mild HCV F1
  - => more expensive
  - => less complications
  - => higher % cured

For G1, 4, 5, 6: 23.000 €/QALY

For G2, 3: 4.600 €/QALY

=> cost/effective if < 50 000€ /QALY

# **2. Hepatitis B**

# Burden of disease: hepatitis B

- **BASL registry 12m 3/2008-2/2009:** report all HBsAg+ pts at consultation
- N = 1421 pts                      26 centres                      71% prevalent
- Mean age: 42y                      67% male
- 52% Caucasian                      25% black Africans
- **Risk factors:**                      14% transfusion;                      9% IDU;                      6% surgery;  
    38% sexual behavior; 33% familial transmission
- 92/1421 = **12% coinfect** (HDV 26; HCV 28; HIV 32)
- **Liver biopsy** in 641pts:    F0F1:40%; F2: 24%; F3: 19%; F4: 17%
- **Phase:**                      Immune tolerant: 0.7%;                      HBeAg+: 17%;  
    HBeAg-: 29%;                      inactive carrier: 44%;  
    9% not classified

VHPB meeting "identification and management of chronic viral hepatitis in Europe" 18-19 march 2010, Budapest, Hungary.

# Burden of disease: hepatitis B

- **KCE registry 6m 1/2009-6/2009**: report all HBsAg+ pts at consultation + data of 2006 if available
- Objectives:
  - To estimate number of pts visiting a specialist for HBV
  - To document the distribution of different HBV stages
  - To document QOL per disease stage (EQ-5D CRF)
  - To document the expenses for health insurance per disease stage
- 18 centres                      n= 544 pts                      mean age: 46y                      47% men
- 51% European                      9% Turkey                      22% Africa                      18% Asia
- 2% had HCC                      11% were transplanted                      14% had cirrhosis
- Immune tolerants: 4%                      34% inactive carriers                      20% HBeAg+
- 40% HBeAg-                      2% HBsAg-

# Burden of disease: hepatitis B

- **KCE registry 6m 1/2009-6/2009**: report all HBsAg+ pts at consultation + data of 2006 if available
- **QOL**
  - HCC and decompensated cirrhosis: 0.67 and 0.7
  - LTx: 0.82
  - Rest: 0.8

Cost effectiveness model: has still to be done



# Screening programme HBV

## Implementation

- **Since 1972**: screening of blood and blood products/tissues for HBV
- **Universal precautions** in healthcare setting
- Screening of **pregnant women**
- Opportunistic screening by GP and gastroenterologists

# Screening/**prevention** programme HBV

## Results

- **Pregnant women**: if HBsAg+ => newborn should receive vaccination AND immunoglobulins within 24h
- Systematic vaccination for **infants**: since 1999
- Systematic vaccination for **adolescents**: since 1999
- **Reimbursement** for **HBV vaccine** since 1980ies for at risk population (hemophiliac, HD, family, pre-and post Tx, handicapped, ...)

# Treatment strategies: HBV

Since jan 2010: finally good reimbursement criteria

- **Liver biopsy** required: inflammation and/or fibrosis; **ALT** at 2 time points elevated
- **Viral load HBV DNA:** > 2000 IU/ml
- **Which products:**
  - Lamivudine in 1st line
  - Adefovir in 2<sup>nd</sup> line
  - Entecavir in 1st and 2<sup>nd</sup> line
  - Tenofovir in 1st and 2<sup>nd</sup> line
- **FU:** every 3 to 6m under therapy; if F3-F4: US every 6months

# Drugs available in Belgium

Drug	Cost per year	in B	TT duration
<b>Peg-interferon <math>\alpha</math>2a</b>	11.468 €	1°line	48w
<b>Lamivudine</b>	749 €	1° line	?
<b>Adefovir</b>	6.159 €	2° line	?
<b>Tenofovir</b>	4.992 €	1° +2° line	?
<b>Entecavir</b>	5.244 €	1° + 2°line	?

# Conclusions:

## Screening for HCV and HBV in Belgium

- No coordinated screening at federal/regional level
- As specialist => try to stimulate/alert general practitioners to screen the at risk population
- Centres for IDU, STI clinics, ... => screen for HIV, HBV and HCV

# Conclusions:

## Prevention for HCV and HBV in Belgium

- No coordinated prevention at federal/regional level
- Centres for IDU => vaccinate sometimes for HAV/HBV
- Systematic HBV vaccination: advised for all babies and children (11-12y) since 1999
- Reimbursement of HBV vaccine in some risk groups since 1980ies

# The end!

