#### Hepatitis C Virus Infection in Massachusetts: A tale of two epidemics

Daniel Church, MPH Massachusetts Department of Public Health Bureau of Infectious Disease

### **Goals of presentation**

- Provide an overview of viral hepatitis, with a focus on hepatitis C virus (HCV)
- Discuss HCV epidemiology, transmission, testing and treatment
- Describe surveillance for viral hepatitis and recent trends in in Massachusetts
- Discuss role of public health nurses

# **Viral hepatitis**

- Hepatitis A Virus (HAV, fecal-oral transmission, vaccine available)
- Hepatitis B Virus (HBV, blood-borne, vaccine available)
- Hepatitis C Virus (HCV, blood-borne)
- Hepatitis D Virus (blood-borne, only causes problems for people infected with HBV)
- □ Hepatitis E Virus (fecal-oral, occurs rarely in U.S.)

# **Disease burden in the U.S.**

\*all numbers shown are estimates

Outcome	HAV	HBV	HCV	HIV
New Infections	21,000	38,000	(17,000)	47,500
Chronic Infections	NA	0.8-1.4 million	2.7-3.9 million	1.1 million
Deaths/ year	~80	3,000	15,100	12,700
Percent aware of infection status	NA	35%	25-50%	82%

IOM, 2010; CDC 2006-2008; CDC, 2013; Ly, et al, 2012

#### Hepatitis A epidemiology in Massachusetts

- □ Number of cases in sharp decline since 2006
- Statewide and localized outbreaks detected in past among MSM, IDU and homeless
  - Adult HAV vaccination programs implemented in past to respond to epidemiological findings
- □ HAV vaccine recommended for all children at age of one year since 2006.

# Hepatitis A

Reported confirmed acute HAV infections in Massachusetts by year, 1998-2012



#### Hepatitis B epidemiology in Massachusetts

- □ The number of reported cases of acute hepatitis B reported to MDPH has been decreasing since 2005
  - Birth dose of HBV vaccine recommended
  - Mandatory vaccination requirements in schools
- Testing among pregnant women and follow-up with infected mothers to reduce transmission to the infant and household contacts (MDPH Perinatal Hepatitis B Prevention Program)
- Cases only represent those people who have been screened, tested and reported to MDPH

#### **Acute HBV infection**

Confirmed acute HBV cases reported in Massachusetts by year, 2002-2011



\*Data as of 2/27/13 and are subject to change Source: MDPH Office of Integrated Surveillance and Informatics Services

#### **Chronic hepatitis B infection**

Chronic HBV cases by year and case status in Massachusetts, 2002-2011



Source: MDPH Office of Integrated Surveillance and Informatics Services

# Hepatitis C – the virus

- □ RNA virus
- Enters the bloodstream, goes into liver cells to reproduce, causing inflammation
- □ Mutates rapidly, evades the immune system
- Causes chronic infection in 75%-85% of infected individuals
- □ Viral infectivity:
  - Up to 63 days in a syringe barrel
  - Up to 21 days in H2O in a plastic container
  - Up to 14 days on inanimate faces

#### HCV – the virus

- □ 6 genotypes
- □ Most Americans have genotype 1
  - Genotype 1 remains the most challenging to treat
- A person can be infected or re-infected with more than 1 genotype
- □ There is no vaccine for HCV

#### What are the two HCV epidemics?

Reported HCV cases in Massachusetts: 2002 and 2009



# **HCV in the United States**

- Estimated 2.7-3.9 million people have chronic HCV in the United States
- □ Current incidence not well established
- Prevalence highest in groups with risk factors that include:
  - Current or former injection drug use
  - People who received blood products before June 1992

#### Long term hemodialysis

IOM, 2010;

CDC Disease Burden from Viral Hepatitis A, B, C in the United States updated 9/13/11

# **HCV prevalence**

#### □ NHANES

- General population: 1.6%
- Males: 2.1%
- Born between 1945-1965
- Non-Hispanic blacks: 3.0%
- Transfusion before 1992: 5.8%
- □ Injection Drug Users (IDUs):
  - **70% 90% (Alter, 1998; Hagan, 2008)**
- □ Incarcerated:
  - 12% 35% (Boutwell, et al, 2005)

#### **HCV** transmission

- Bloodborne pathogen
- Asymptomatic still potentially infectious
- □ Most people infected through:
  - Injection drug use (sharing drug injection equipment)
  - Blood transfusions/clotting factors/organ transplants prior to 1992
  - Chronic hemodialysis
  - Sexual transmission inefficient but does occur
  - Vertical transmission 4-7% of births to infected mothers (20% in HIV/HCV co-infected)

#### **Possible transmission risks**

- Occupational exposures
  - Risk from needlestick:
    - □ HIV=3/1000 HCV=2/100 HBV=3/10
  - Prevalence of HCV in health care workers is the same as the general population
- □ Sharing personal/household items with blood
- Intranasal drug use
- □ Tattoo/body piercing: nonsterile practices

# HCV – injection drug users (IDU)

- □ IDU accounts for 68% of all new infections (CDC)
- As many as 32% of IDUs are infected with HCV within 1 year of first injecting; 53% within 5 years (Hagan, et al, 2008)
- □ Sharing of syringes, cookers, cottons, rinse water, etc. from injection drug use is the greatest risk for HCV transmission
- □ HCV infection CAN be prevented among injection drug users
  - Access to sterile injection equipment and multi-component prevention programs is critical

#### **Sexual transmission of HCV**

- □ Occurs, but efficiency is low
- Low incidence (0.01-0.13%) among monogamous long-term partners, or one per 190,000 sexual contacts (Terrault, et al, 2013)
- May account for 15-20% of acute and chronic infections in the United States (CDC)
- Increased transmission among HIV+ MSM (CDC, 2011)

# Natural history of hepatitis C



Annual age-adjusted mortality rates from hepatitis B and hepatitis C virus and HIV infections listed as causes of death in the United States between 1999 and 2007



Ly K et al. Ann Intern Med 2012;156:271-278

#### Mortality among HCV cases in Massachusetts

#### Lijewski, et al, 2012

Timing of mortality among known HCV cases in Massachusetts, 1992-2009



76,122 HCV diagnoses were reported to the MDPH between 1992 and 2009, 8,499 of these reported HCV cases died and are represented in the figure. Data as of 1/11/2011.

### Why should we screen for HCV?

#### Public & personal health

- Those infected may transmit to others
- ESLD due to HCV great burden on health care system
- 18,000 deaths/year by 2020, 35,000 deaths/year by 2030
- □ We can do something about it
  - Over half (70-80%) of those with chronic HCV can be cured

### **CDC risk-based HCV screening recommendations (1998)**

- Ever injected illicit drugs
- Received a transfusion or blood products before July 1992
- Received clotting factor prior to 1988
- Children >18 months
   born to HCV-positive
   women

- Ever on hemodialysis
- □ HIV-positive
- Healthcare, emergency, public safety workers after needlestick/mucosal exposures to HCVpositive blood

How successful are current screening recommendations?

#### Only 25-50% of those infected with HCV are aware of their diagnosis

Kwiatkowski Addiction 2002 Wasley et al. J Clin Virol 2006 Hagan et al. Pub Health Rep 2006 Volk et al. Hepatology 2009

### **Changes to HCV screening recommendations (2012)**

- □ Move to focus on age-based screening
  - 2/3 of HCV cases among "baby-boomer" population
- Recommendation: One-time HCV screening for all people born between 1945-1965
  - Alcohol use screening and treatment for HCV+
- □ Risk-based screening still important

Treatment regimens	HCV Genotype	Treatment Regimen	Success (SVR*) rate	Side-effects
And coming very soon: <u>More DAAs:</u> - More protease inhibitors - NS5A inhibitors - Nucleotide polymerase inhibitors (e.g., Sofosbuvir) <u>Interferon-free regimens?</u> <u>Human pharmacogenetics</u> - IL28B polymorphisms	1	<ul> <li>Peginterferon</li> <li>Ribavirin</li> <li>12 week "boost" with a <u>Direct Acting</u> <u>Antiviral (DAA):</u> NS3 protease inhibitor (Telaprevir or Boceprevir)</li> <li>Duration: 24-48 weeks depending on early response</li> </ul>	62-80%	<ul> <li>Flu-like sx</li> <li>Mood changes</li> <li>Pancytopenia</li> <li>Autoimmunity</li> <li>Hemolytic anemia</li> <li>Teratogenicity</li> <li>For protease inhibitor regimens:</li> <li>More severe anemia</li> <li>Pruritus and rash</li> <li>Dysguesia</li> </ul>
	2 and 3	<ul> <li>Peginterferon</li> <li>Ribavirin</li> <li>Duration: 24 weeks</li> </ul>	78-82%	

\* Sustained Virologic Response = negative HCV viral load 24 weeks after treatment

#### Harm reduction

- HAV and HBV vaccination
- Alcohol cessation
- Avoidance of hepatotoxic medications or OTC products
- Hepatitis C education
- Counseling about transmission
- Referral to psychiatric or addiction treatment when appropriate
- Referral to hepatology for cirrhotics

# **HCV harm reduction for IDUs**

- To reduce spread of HCV, IDUs should:
- Be provided information on drug treatment options
- Be informed about existing needle exchange programs and pharmacy access
- □ Have access to harm reduction education
  - Clean works
  - Safe injection practices
  - Overdose prevention
  - Opioid replacement therapy

# Hepatitis C surveillance

- Estimated ~100,000 people in Massachusetts exposed to HCV in the past
- Number of cases relatively stable since 2002 with 8-10,000 newly diagnosed cases reported to MDPH annually
- Cases only represent those people who have been screened, tested and reported to MDPH
- Confirmed and probable case definitions according to CDC classification
- Data entered into and managed in MAVEN

#### Why do we do surveillance?

#### Detect potential outbreaks



#### Why do we do surveillance?

- □ Interrupt transmission
  - Provide harm reduction messages
- Identify at-risk populations and emerging issues
- □ Target areas for services

#### **Reported cases of HCV infection in Massachusetts: 2000-2011**





# HCV among youth in Massachusetts 2007-2011

- Starting in 2007 an increase of newly diagnosed HCV infection has been noted among youth ages 15-25
- Between 2002 and 2011, an increase of 62 to 132 cases per 100,000 population was reported in this age group
- Data suggest that the increase is due to youth injecting drugs (mostly heroin)
- Other jurisdictions have also seen this trend (CT, HI, KY, ME, MN, NY, PA and others)

#### MMWR: Age distribution of newly reported confirmed cases of hepatitis C virus infection ----Massachusetts, 2002 and 2009



\* N = 6,281; excludes 35 cases with missing age or sex information. + N = 3,904; excludes 346 cases with missing age or sex information.



Data as of January 23, 2013 and are subject to change

#### Confirmed and probable cases of HCV infection by age and gender in Massachusetts, 2011



### Role of Public Health Nurses

- Investigate suspect cases of acute HBV and HCV infection
  - Work with medical providers and cases as possible to gather information indicated in MAVEN
    - MDPH epidemiologists will contact you and provide guidance tip sheets in MAVEN forthcoming
  - Key issues: Determine acute status (when was the case first exposed?), risk history (was this health care acquired?), and provide education to case
- Be a resource for their communities for viral hepatitis information and referrals

#### Programs in Massachusetts

- □ No direct viral hepatitis funding from state legislature
- Integration of HCV services (prevention education, screening, testing) has been fully implemented with all HIV prevention and screening programs (34 programs)
  - Requirement of completing case report form recently implemented for those conducting point-of-care testing, including rapid HCV tests
- Integration of HCV medical management into HIV case management services (5 programs)
- For program location and contact information: <u>http://www.mass.gov/eohhs/docs/dph/aids/res</u> <u>ources-guide.pdf</u>

#### New report available



Shifting Epidemics: HIV and Hepatitis C Infection among Injection Drug Users in Massachusetts

Fifth in a Series of Reports on the Status of the HIV/AIDS Epidemic in Massachusetts

2012

Deval L. Patrick Governor Timothy P. Murray Lieutenant Governor

http://www.mass.gov/eohhs/docs/dph/aids/shifting-epidemics-report.pdf

#### Massachusetts Viral Hepatitis Coalition

- Conducts advocacy and education on viral hepatitis in Massachusetts
- Members include medical and social service providers, consumers, pharmaceutical companies and other community partners
- Quarterly meetings with some sub-committees
- For more information and to get involved contact Katie Boos (<u>kboos@aac.org</u>)

### **Resources: Provider Education**

- □ CDC <u>http://www.cdc.gov/hepatitis/</u>
- Hepatitis Web Study: <u>http://depts.washington.edu/hepstudy/</u>
- National Training Center for Integrated Hepatitis, HIV and STD Prevention Services

www.knowhepatitis.org

- Treatment Action Group <u>www.treatmentactiongroup.org/hepatitis</u>
- Caring Ambassadors Program: Hepatitis C <u>http://www.hepcchallenge.org/index.htm</u>

### **Resources: Patient Education**

- □ CDC <u>http://www.cdc.gov/hepatitis/</u>
  - "Know Hepatitis" campaign
- □ Harm Reduction Coalition <u>http://harmreduction.org/</u>
- Hepatitis C Support Project

http://www.hcvadvocate.org/

- Treatment Action Group <u>www.treatmentactiongroup.org/hepatitis</u>
- Caring Ambassadors Program: Hepatitis C <u>http://www.hepcchallenge.org/index.htm</u>

# **Resources: Policy**

□ US Department of Health and Human Services Viral Hepatitis Action Plan (2011)

http://www.hhs.gov/ash/initiatives/hepatitis/actionpl an\_viralhepatitis2011.pdf

 Institute of Medicine Report on Hepatitis and Liver Cancer (2010)

http://www.cdc.gov/hepatitis/IOMnews.htm

- National Viral Hepatitis Roundtable <u>www.nvhr.org</u>
- National Alliance of State and Territorial AIDS Directors (NASTAD) <u>www.nastad.org</u>

## Questions?

Dan Church, MPH

William A Hinton State Laboratory

305 South St. Jamaica Plain, MA 02130

617-983-6830

Daniel.church@state.ma.us