HIV Testing in New Jersey:
A 2010 Status Report

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New Jersey’s HIV Epidemic

• The face of the NJ HIV epidemic is a bit different from urban centers like San Francisco and New York City
  – Although NJ is a high prevalence state, the face of HIV is made up increasingly of women and minorities.
  – The NJ epidemic is characterized by urban pockets of infection, drug abuse and poverty
  – Historically,
    • 5th in the US in cumulative reported AIDS cases
    • 3rd in the US in cumulative reported pediatric AIDS cases
    • 1st in the proportion of women with AIDS among its cumulative reported AIDS cases.
    • 34,915 persons living with HIV or AIDS (12/31/08)
HIV cases among IVDU

- Historically, 41% of HIV cases in New Jersey are among IVDU
- Recently, between 8-10% cases have been associated with IVDU
HIV AIDS in NJ

Reported AIDS Cases by Race/Ethnicity, Cumulative through 2005, New Jersey
N = 48,431

- White, not Hispanic (26.2%)
- Black, not Hispanic (55.9%)
- Hispanic (17.0%)
- Asian/Pacific Islander (0.1%)
- American Indian/Alaska Native (0.1%)
- Unknown/Other (0.7%)
HIV and Drug Use

• Drug users are at high-risk population for HIV infection/transmission
  – NOT ONLY through actual “use” → studies have shown a decrease in prevalence and incidence after successful implementation of syringe needle exchange programs
  – BUT MOSTLY through high-risk sexual activities (rates of STDs have not decreased)
NJ HIV

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Rapid Testing is Widely Available in NJ

County and Municipal Statistics

Prevalence Rate:
Persons Living with HIV/AIDS per 100,000 population
- 0.0 - 199.9
- 200.0 - 399.9
- 400.0 - 1199.9

Statewide Summary:
Prevalence of Persons Living with HIV/AIDS

Persons Living with HIV/AIDS: 34,470
Total Population, Estimate 07/01/06: 8,724,560
Prevalence Rate per 100,000 pop.: 395.1

Cases not on map
- County Unknown: 63
- Incarcerated at Diagnosis: 1,907

Note: Top number indicates number of persons living with HIV/AIDS (HIV Positive Infection or AIDS) as of 12/31/2007. Not included in this number are cases of Perinatal HIV Exposure that are not confirmed HIV Positive. Bottom number in parentheses indicates prevalence rate of persons living with HIV/AIDS per 100,000 population (July 1, 2006 estimate).
New Jersey
Rapid HIV Testing

- Counseling sites - mid-1980s
- Testing added - late 1980s
- Rapid testing added - late 2003
- Currently:
  - 73 laboratory licenses
  - Over 90 CTS testing sites:
    - Hospitals/EDs (23), STD clinics, CBOs, Health Depts., Mobile Vans, Prisons, Faith-based Initiatives, Methadone treatment programs, substance abuse programs
- 2006 ASTHO Vision Award
Centralized Approach to Rapid HIV Testing

- laboratory oversight
- uniform procedures, policies
- quality assurance
- proficiency testing
- training
- certification
- support with testing discordants
- logistics
- licensing
- inventory
- testing data for rapid HIV program
- monitoring, site visits, compliance and competency reports
- 2 MD, 1 PhD, 1 manager, 6 techs
Goal for Today

1. Rapid Testing Algorithms (RTA) – “Rapid-Rapid” – The first 25,000!
2. “Pooled Screening” - Testing for very early infection in antibody negative blood – Newark ER’s
3. Developments - i.e impending licensure of 4th generation POC tests
LIMITATIONS OF RAPID TESTING

• The Rapid HIV Test Measures
  – Antibodies to HIV

• It DOES NOT Measure?
  – HIV virus/RNA or DNA

• How Sensitive Is It?
  – At least as sensitive as more complex EIA technology used in hospitals and laboratories
  – In some cases it is more sensitive than the Western blot, the ‘Gold Standard’ for validation.
Western blot – “The Gold Standard”

• Every HIV Positive currently requires a confirmation
  – Getting old...
    • Complex test - prone to QA issues.
    • Cost issues
    • Supply and Manufacturing issues
    • Indeterminate HIV-1 *Western Blots*:
      – Need for additional *testing*
      – Differences in Western blot performance

  – ...... and delays in rendering a final interpretation
  – Sometimes we focus too much on the issue of specificity and not enough on the consequences of delay.
Problem - confirmatory process

• People refuse confirmatory tests
  – 7.1% of positives could not be confirmed because specimens not collected

• Many don’t return to get their final results
  – New Jersey: 25 – 30% fail to return for a second testing-related visit.
  – Los Angeles: 35-40% fail to return
  – Other urban environments – similar story sometimes even worse

• Bottom line:
  – ONLY ~ 70% actually get their confirmed + result!!

• Impact ➔ Linkage to Care is:
  – Delayed
Why Rapid Verification?

NJ Statewide Data - 2004

- Problem
  - Preliminary Positive clients fail to return for results (21.8%)
  - NAP succeeds ONLY 20% of the time in locating these clients
- Solution
  - Confirmatory testing on-site, same day

Disposition of Confirmed HIV+ Clients

<table>
<thead>
<tr>
<th>Number</th>
<th>Confirmed HIV+</th>
<th>Result returned to client</th>
<th>Did Not Receive Results</th>
<th>Referred to NAP</th>
<th>Found by NAP</th>
</tr>
</thead>
<tbody>
<tr>
<td>326</td>
<td>244</td>
<td>82</td>
<td>47</td>
<td>11</td>
<td></td>
</tr>
</tbody>
</table>
Why? What can we do?

- Its difficult for some clients to work up the courage → if there is any delay it is an excuse for procrastination
- Identifying an infected individual and linking them to care immediately → increases the likelihood for treatment
- Consistently identifying truly infected individuals improves overall program credibility with providers.

- Goal: Test and Link to Care in a single visit
The solution: Rapid-Rapid Testing

- **Widespread use worldwide** ~ fifteen years
  - Resource-poor
  - High prevalence environments
- **What was not tested was low prevalence environments**
- **Los Angeles/San Francisco CDC pilot studies** “rapid-rapid” – RTA 2008-9
- **New Jersey implementation** – 2009
- **New York implementation** – upstate - 2009
Three Test RTA vs. Two Test RTA

• **Definitive (3 Test RTA)**
  - Does program credibility hang on ‘getting it completely right’?
  - A 3 test RTA will allow you to resolve more discordant events; BUT
  - **Downside:** QC costs and potential operator errors for seldom used tests

• **Efficient (2 Test RTA)**
  - If we identify 98% of infected clients in a single visit, and successfully connect them to healthcare ➔ way ahead.
  - Less to remember, less to forget in a two-test algorithm
  - **Downside:** A very small number will not be resolvable at the time of initial testing.
  - **Key:** What happens to the problem cases – NJ is centralized ➔ laboratorian/physician interaction early.
Validation of a Testing Algorithm

“Validation is the process of demonstrating that an analytical procedure is suitable for its intended use” - CBER

• The use of other rapid tests to confirm a rapid HIV test is not new or novel - WHO recommends this approach for countries in which the prevalence of HIV exceeds 10% for a number of years

• What was new and novel was using this approach in sites with 2% or lower prevalence
Validation of an RTA – 2004-8

• Goal – To satisfy ourselves that a second, independent rapid HIV test could reliably identify false positive HIV tests
  - 2004 – Using residual serum confirmed all Western blot positive sera obtained in the previous year and available at the Public Health Labs
  - 2005-8:
    • Using residual sera and plasma samples to confirm that a second independent rapid HIV test could reliably identify false positive HIV tests
Rapid confirmation trial

July 1, 2004 through April 19, 2005

- 15,923 OraQuick tests statewide
- 363 prelim positive samples to state lab for confirmatory testing
  - 355 Western Blot positive
  - 8 Western Blot negative
- A second rapid test – Unigold identified all 8 false positive rapids and agreed with all 355 HIV + diagnoses
Rapid-Rapid Implementation

- DEPLOYMENT PLAN:
  - December, 2008: 3 pilot sites began the ‘roll-out’
  - Higher prevalence first, lower prevalence later
  - Policies, Procedures, Counseling Messages and Forms were completed for the entire system available before training
  - Side-step the issue of confirmation

- EXPECTATIONS:
  - Doesn’t eliminate Western blot confirmation, BUT allow immediate linkage to care reliably without a western blot!
  - Less than 1 in 100 would later be removed from care because of a failure to confirm

- UNKNOWNS: What will be the real world performance of a rapid test in a confirmatory setting?
  - Does reducing the delay really improve the linkage to care?
  - False Pos frequency? Would there be False Neg’s
Status of Rapid-Rapid in New Jersey

February, 2010
Issues

• Number of Rapid-Rapid Sites
• Distribution of sites throughout the state
• Training
• Real-world performance of the ‘second rapid’
• Costs of the ‘second rapid’
• Does it increase the linkage to care?
Implementation of Rapid-Rapid

Rapid-Rapid Pgm

- Prelim Pos
- UniGold Confirmed
Various Venues for Rapid-Rapid Program

- 21 Primary Sites Operational in NJ
  - 42 Satellite Sites
    - FQHC’s: 4
    - Hospital ER’s: 7
    - CBO’s: 16
    - Mobile Van Initiatives: 15
- >140 individuals Trained
Diversity of sites using an RTA

County and Municipal Statistics

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Notes: Top number indicates number of persons living with HIV/AIDS (HIV Positive Infection or AIDS) as of December 31, 2007. Not included in this number are cases of Vertical HIV Exposure that are not confirmed HIV Positive. Bottom number in parenthesis indicates prevalence rate of persons living with HIV/AIDS per 100,000 population (July 1, 2007, estimated).
<table>
<thead>
<tr>
<th></th>
<th>Tests</th>
<th>PCT</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Rapid Test 1</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>StatPak</td>
<td>19,830</td>
<td>77.4%</td>
</tr>
<tr>
<td>Oraquick Oral</td>
<td>3,005</td>
<td>11.7%</td>
</tr>
<tr>
<td>Oraquick Finger Stick</td>
<td>2,778</td>
<td>10.8%</td>
</tr>
<tr>
<td><strong>Rapid Test 2</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unigold</td>
<td>213</td>
<td></td>
</tr>
</tbody>
</table>

**RTA Total Tested**: 25,623
## Outcomes

<table>
<thead>
<tr>
<th>Date</th>
<th>Result Description</th>
<th>Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>2/28/2010</td>
<td>Preliminary Positive</td>
<td>213</td>
</tr>
<tr>
<td></td>
<td>Unigold Verified</td>
<td>194</td>
</tr>
<tr>
<td></td>
<td>Same-Day Connect to Care</td>
<td>146</td>
</tr>
<tr>
<td></td>
<td>Verified &amp; Linked Same Day</td>
<td>75%</td>
</tr>
</tbody>
</table>

![Bar chart showing Linkage to Care Rapid-Rapid NJ](chart.png)
Discordant Results
NJ Rapid Testing Program
Rapid AND Rapid-Rapid

Rapid HIV Discordants

Tests

Discordants

Dec-05  Dec-06  Dec-07  Dec-08  Dec-09

0  10  20  30  40  50  60  70  80  90  100

0  10 000  20 000  30 000  40 000  50 000  60 000  70 000  80 000  90 000  100 000

Tests

Discordants

Tests  Discordants
Discordant Issues

RTA Verification
- UniGold Confirmed: 193
- Discordant: 15

Discordant Characterization
- False Pos: 3
- False Neg: 2

RTA Program Specificity: 99.94%
## Rapid-Rapid Summary

**February, 2010**

<table>
<thead>
<tr>
<th>WB Results</th>
<th>1st Rapid Positive</th>
<th>2nd Rapid Positive</th>
<th>2nd Rapid Negative</th>
<th>Notes: Percentages calculated excluding those who refused WB</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total WB results</td>
<td>197</td>
<td>186</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Pct WB POS</td>
<td>95.4%</td>
<td>99.5%</td>
<td>27.3%</td>
<td></td>
</tr>
<tr>
<td>Pct WB Ind</td>
<td>0.0%</td>
<td>0.0%</td>
<td>9.1%</td>
<td></td>
</tr>
<tr>
<td>Pct WB Neg</td>
<td>4.1%</td>
<td>0.5%</td>
<td>80.0%</td>
<td></td>
</tr>
<tr>
<td>Pct Refused WB</td>
<td>7.0%</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Linkage to Care
Linkage to Care
Rapid Test + W. Blot Algorithm: 2005-2006

• 644 cases - rapid test confirmed by W. Blot
  – 425 (66%) had at least one CD4 count or VL in eHARS database indicating ‘in treatment’
  – Based on the date of the 1st lab result, 253 of the 425 (60%) got into treatment within 3 mos
  – Of the 172 (41%) who got into treatment after 3 months, the median time to treatment was 524 days (range 93 – 1416 days)
  – > one-third of the 644 cases did not have evidence of a CD4 count or viral load test result

• Getting results 3.6x more likely to be in RX
Who Gets Linked to Care - RTA

- 75% of ‘verified’ HIV positives receive appts on the same day
- 26% DID NOT receive appts on the same day!!
- Site Specific Issues - Ongoing
- How to improve linkage
- 15% More then traditional testing!!
Linkage to Care with an RTA – Where matters!

- 62 RTA positives identified in the first six months of RTA program: 76.7% - same day appointments for treatment
  - Academic medical centers (1) and FQHCs (4) identified 33 HIV positive individuals using an RTA
  - 82% received immediate appt
  - **97% were in care at six months, 1 lost to care**
- Health Departments (2) and CBOs identified 29 infections
  - 16 (55%) appts. were made on same day
  - **19 (47%) were in care at 6 months, 10 (34.4%) lost to care**

- Efforts to better connect screened, infected clients to providers is needed in non-traditional healthcare settings
Facts

• Roughly 7% of clients in NJ refuse Western blot testing altogether. These have been linked to care on the basis of a second rapid, BUT NOT in its absence.
• Clients who must return to receive Western blot results fail to do so 25% of the time.
• → ~32%
• Partly because of resistance by providers to take on patients prior to a conclusive ‘diagnosis’ – Western blot has become a roadblock to entering care.
• If we run two rapid HIV tests from different manufacturers AND they both are positive, 99.5% will confirm WHEN a Western blot is completed.
• If the two rapids disagree:
  – 27.3% of the time the Western blot will be POSITIVE,
  – 9.1% of the time it will be INDETERMINATE
  – 80% of the time it will be completely NEGATIVE
• Using a ‘Rapid-Rapid’ less than 1:100 will later be pulled out of care. OBVIOUS QUESTION: Why not refer on the basis of a second rapid?
Facts

1. The result of the second rapid is “credible verification”.
2. Clients exposed to the results of the second rapid have little reason to procrastinate about linking to care.
3. The cost of a second rapid is between $7-15. The cost of a Western blot is between $70 - $250.
The Obvious Question

• Given the expense, the limited sensitivity, the complexity AND the inability to move the Western blot into the POC environment, do we need to consider the proposition that the Western blot has outlived its usefulness in a screening arena?
Testing for Early Infection

“NAT Testing of Pooled Negatives”
Viremia During Early HIV Infection

- **Ramp-up Viremia Doubling Time**: 21.5 hrs
- **Peak Viremia**: $10^6 - 10^8$ gEq/mL
- **Viral set-point**: $10^2 - 10^5$ gEq/mL
- **WINDOW**
  - Antibody – 22 Days
  - Antigen – 16 Days
  - Pooled NAT – 14 Days
  - Individual NAT – 11 Days
Natural History - HIV Infection

Couthino et al., Bulletin of Mathematical Biology 2001
Why worry about the ‘30 day window’?

• HIV transmission
  – 50% occurs PRIOR to appearance of antibodies!

How do we EFFECTIVELY narrow the ‘HIV window’

  – Implement alternative technologies to identify
    • Other serologic markers that appear earlier
    • Utilize nucleic amplification technologies to look for the virus
Future Developments

AHI Study in Emergency Rooms

- High prevalence area – central ward Newark
- ? How frequently do individuals present in the ER with evidence of acute HIV infection?
  - AHI: Appearance of HIV virus associated with burst of infectivity. No antibody present. Possibly p24 Ag is present
- University Hospital ER and St. Michael’s ER
Acute HIV Infection

- HIV virus is associated with burst of infectivity.
- No detectable antibody present for ~ 22 days
- Possibly p24 Ag present

- ?: If we screen with a POCT product detecting p24 Ag can we:
  - Detect most or all of the AHI episodes?
  - Is that good enough?
  - Does the immediate result offset the delay issues?
Technology trade-offs

**POOLED NAT**
- ~ 14 days
  - Sensitive
  - labor intensive
  - Expensive
  - Often referred
  - Reporting delays: Days → Weeks

**P24 Ag Detect**
- ~ 16 days
- Originally a lab-based technology
- Approaching licensure is a POCT like rapid HIV
  - ? Pricing
  - ? How to utilize
Ongoing Efforts

• Expanding Rapid-Rapid – focusing on the ‘linkage to care’
• Determining whether p24 Ag detection is an acceptable alternative to NAT in a screening environment.
THANKS!

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