PARAMETERS IV: CLINICAL CARE AND TREATMENT

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Outline for this session

- Care continuum as modeled
 - HIV testing
 - ART initiation (linkage to care with ART)
 - Dynamic engagement in care via ART on/off status
 - Dynamic suppression status
 - On ART, fully suppressed
 - On ART, partially suppressed
 - (Not suppressed because not on ART or undiagnosed)

HIV testing

HIV testing: concept

- HIV testing is critical prevention tool
 - Behavior changes post-diagnosis
 - Necessary for receipt of HIV care (and PrEP in PrEP models)
- Model conceptualized 2 types of testers
 - Never testers 6.5% of sample didn't have HIV test by age 40
 - Regular testers: memoryless exponential process, per mean time between tests
 - Susie Cassells (UCSB) has R21 extending to more types
- Also incorporated 21-day window period until Ab detection
 - HIV+ positive persons thus test HIV-ne

HIV testing: assessment

62. Have you ever been tested for HIV? 1 O Yes If yes, then #63, #64, #65, #66 0 No HIVtest_ever Binary_prefernot.
63. In what month and year did you have your most recent HIV test?
Month: January =1 February =2 March =3 April =4 May =5 June =6 July =7 August =8 September =9 October =10 November =11 December =12
64. Year: HIVtest_year HIVtest_result.
65. What was the result of your most recent HIV test? O Negative Positive If positive, then hide #67, #70, #71, #72, #73. If positive, then show #68, #69 Indeterminant/Inconclusive O Didn't get the results of my last HIV test

HIV testing: model metric

Mean time since last test: (today – date of last test)

BMSM: 301 days

WMSM: 315 days

 Assuming testing is memoryless exponential with rate parameter equal to above means, Bernoulli daily (-> weekly) testing probability is the inverse of these.

ART initiation (linkage to care with ART)

ART linkage/initiation

- Linkage to care defined in many ways, but typically about the duration from diagnosis until first HIV clinical visit
- The model has no direct representation of clinical visits
 - It does have ART, which is a primary purpose of HIV care
 - Linkage to care to data empirically available in some CDC reports
 - But not time to ART initiation. Assumed instant initiation of therapy once in care
- Following HIV diagnosis, MSM initiated treatment:
 - BMSM rate = 0.924/week (10.8 weeks on avg.)
 - WMSM rate = 1.271/week (7.9 weeks on avg.)

Dynamic engagement in care

Dynamic engagement in care: model concept

- Persons on ART can fall out of care
 - Viral load returns to 4.5 log₁₀ set-point linearly over 3 months
- Persons can re-engage in care and resume adherence process
- Rates based on cross-sectional CDC care continuum sources
 - Volumes can be said about the veracity of these estimates

Dynamic suppression status on ART

- Once on ART, may or not be fully adherent
 - Like "care" is modeled as ART on/off status, adherence is modeled in terms of VL suppression levels
 - Fully suppressed = 1.5 log₁₀ = 50 copies/mL = LLD
 - Partially suppressed = (Not suppressed can occur if on ART or undiagnosed)
 - (unsuppressed occurs if not diagnosed or not on ART)
 - Can cycle between full/partial states
 - Rate parameter back-calculation complicated and used CDC paper on:
 - Viral suppression at last test
 - "Durable" viral suppression at all test for 12 months

Dynamic suppression status on ART

Parameter	Black MSM	White MSM
Proportion of those initiating ART who	0.614	0.651
achieved full suppression		
Per-time step probability of falling out of	0.0102	0.0071
suppression		
Per-time step probability of re-achieving	0.00066	0.00291
suppression		

Implementing in the model

Testing

param.mard:

```
nction (nwstats, last.neg.test.B.int = 301, mean.test.B.int = 301, last.neg.test.W.int = 315, mean.test.W.int = 315, testing.pattern = "interval", test.window.int = 21, tt.traj.B.prob = c(0.077, 0, 0.356,
```

test.mard:

```
if (testing.pattern == "interval") {
    tsinceIntst <- at - dat$attr$last.neg.test
    tsinceIntst[is.na(tsinceIntst)] <- at - dat$attr$arrival.time[is.na(tsinceIntst)]
    tst.B <- which(active == 1 & race == "B" & tt.traj !=
        "NN" & (diag.status == 0 | is.na(diag.status)) &
        tsinceIntst >= mean.test.B.int)
    tst.W <- which(active == 1 & race == "W" & tt.traj !=
        "NN" & (diag.status == 0 | is.na(diag.status)) &
        tsinceIntst >= mean.test.W.int)
}
tst.pos.B <- tst.B[status[tst.B] == 1 & inf.time[tst.B] <=
        at - twind.int]
tst.neg.B <- setdiff(tst.B, tst.pos.B)
tst.pos.W <- tst.W[status[tst.W] == 1 & inf.time[tst.W] <=</pre>
```

Treatment initiation, halting, re-initiation

param.mard:

```
test.window.int = 21, tt.traj.B.prob = c(0.077, 0, 0.356, 0.567), tt.traj.W.prob = c(0.052, 0, 0.331, 0.617), tx.init.B.prob = 0.092, tx.init.W.prob = 0.127, tx.halt.B.prob = 0.0102, tx.halt.W.prob = 0.0071, tx.reinit.B.prob = 0.00066, tx.reinit.W.prob = 0.00291,
```

tx.mard: who initiates, halts, and re-starts treatment, e.g.:

```
tx.ieiiit.w.prob <= datsparamstx.reiiit.w.prob
tx.init.elig.B <- which(active == 1 & race == "B" & status ==
    1 & tx.status == 0 & diag.status == 1 & tt.traj %in%
    c("YP", "YF") & cum.time.on.tx == 0 & stage != "D")
tx.init.B <- tx.init.elig.B[rbinom(length(tx.init.elig.B),
    1, tx.init.B.prob) == 1]
tx.init.elig.W <- which(active == 1 & race == "W" & status ==</pre>
```