TESTING & DIAC	GNOSIS					
First Author,	Study	Decima	Laatian	Danulatian.	Sample Indigenous	
Publication Year	Period	Design	Location	Population	n (%)	Key Cascade Findings
DuBois 1996	1994	cross-sectional	CAN: Montreal	Urban	99 (100)	Proportion ever tested: 48.5% Odds of testing among men, vs European
			NZ: Auckland /	STD clinic		Maori: 0.6; Pacific islander: 0.3; Other: 1.1 Odds of testing among women, vs European
Connor 1997	1991-1992	cross-sectional	Christchurch	attenders	1363 (16.1)	Maori: 0.6; Pacific islander: 0.2; Other: 1.7
			AUS: Anangu			
Miller 1998	1994-1996	cohort (population)	Pitjantjatjara	Adults	592 (100)	Tests per year: 1994: 83> 1996: 592 (24% of pop)
						with SCI: 31%
14 4000	1000		US: Western	Co-morbid (SCI)	070 (400)	without SCI: 60%
Krause 1999	1996	cross-sectional	Mountain Region		672 (100)	Significant difference (p<0.05)
Martin 2002	1992-2000	cohort	CAN: BC	Substance use	2345 (100)	Proportion tested while in drug Tx: 67%
Mak 2003	1997-2002	cohort (population)	AUS: Western	Perinatal	1903 (66.0)	Model: Aboriginal ethnicity not significantly associated with HIV testing
Denny 2003	1998-2000	cross-sectional (BRFSS)	US: National	Adults	4854 (2.2)	Proportion never tested: 50.5% (47.6%-53.4%) Significantly lower than non-AIAN
Defility 2003	1990-2000	cross-sectional	OO. National	Addits	4034 (2.2)	Proportion opting out of prenatal testing: 55 (2.2%)
Wang 2005	2000	(population)	CAN: Alberta	Perinatal	2549 (6.6)	Model: First Nations more likely to opt out AOR: 1.91; 95%CI: 1.42-2.58
Foley 2005	2002-2004	intervention	US: Southwest	Substance use	134 (100)	Proportion tested in drug Tx: 105/134 (78.4%)
Wardman 2006	2003	cross-sectional	CAN: BC	Engaged	219 (100)	Proportion ever tested: 60.4%
77 di di 11 di 12 di 00	2000	or odd oddiona.	07 11 11 20		2.0 (.00)	Proportion tested in past year: 79/218 (36%)
Lapidus 2006	2001	cross-sectional	US: Portland	Adults	222 (100)	Proportion ever tested: 180/218 (82.6%)
'		cross-sectional			` '	Proportion tested in past year: 31 (77.5%)
Sanchez 2006	2003-2005	(NHBSS)	US: National	MSM	40 (0.4)	Proportion ever tested: 38 (95%)
			CAN: Northern	newly diagnosed		Proportion with a late diagnosis (CD4 <200): 49 (38%)
Plitt 2009	1998-2003	cohort (NAHIVP)	Alberta	PLWHA	216 (41.1)	No different than non-Indigenous p=0.20
Long 2010	2003-2008	cohort (population)	CAN: Alberta	Co-morbid (TB)	97 (12.5)	Proportion tested in routine screening in TB care: 86 (89%)
		cross-sectional			520493	Proportion tested in past year: 13.2%
Orchard 2010	2001	(APS)	CAN: National	Off-reserve	(100)	Proportion ever tested: 41.7%
Worthington 2010;						Proportion over tested: 210/412 (E10/1)
Mill 2008 & 2011; Prentice 2011	2004-2005	orose contional	CAN: National	Youth	412 (100)	Proportion ever tested: 210/413 (51%) Proportion with late diagnosis: 6/26 (23.1%)
Prentice 2011	2004-2005	cross-sectional	CAN: National	TOULTI	413 (100)	Proportion ever tested: 43.0%
		cross-sectional				Men: 40.3%; Women 46.8%
Barnes 2010	2004-2008	(NHIS)	US: National	Adults	793 (0.6)	Model: Al/AN more likely than other ethnicities to be tested for HIV
		,				Estimated number of undiagnosed
						1200 (25.8%; 95%CI: 600-1800)
0 "" 0040	0000	mathmatical			1000 (0.4)	60.4 per 100,000 people (95%Cl: 25.1-95.6)
Campsmith 2010	2006	modeling	US: National	General	4600 (0.4)	Model: Al/AN more likely to be undiagnosed vs. white (1.52; 95%Cl: 1.43-1.64)
Hodge 2010	?	cross-sectional	US: California	General	457 (100)	Proportion ever tested: 171/423 (40.4%)
0	0000 0040	cross-sectional	ALIO, National	Health care	40444 (4.0)	Testing rate per 100 encounters: 5.57 (95%Cl: 3.97-7.16)
Sawleshwarkar 2011	2000-2010	(BEACH) cross-sectional	AUS: National	providers	13114 (1.3)	Model: Indigenous more likely to be tested (AOR:1.74; 95%CI:1.26-2.41)
Sondag 2011	2007-2008	(BRFSS)	US: Montana	Adult	? (100)	Proportion ever tested: 39%
Takahashi 2011	2007-2000	cross-sectional	US: S. California	Adult	179 (100)	Proportion ever tested: 53 %
rananaoni 2011	2000	or odd oddiona.	AUS: Western	, taut	()	Model: Aboriginal people more likely to get STI/BBV testing according to
Kwan 2012	2007-2010	pre-post test	Australia	Perinatal	390 (14.3)	guidelines (AOR: 7.45, 95%CI: 5.53–10.05)
		cross-sectional				Proportion tested in past 6 months: 8% (95%CI: 0-20%)
Bauer 2012	2009-2010	(Trans Pulse)	CAN: Ontario	Trans people	35 (8.1)	Proportion never tested: 15% (95%CI: 3-36%)
						Proportion tested in past year: 66.7% (95%CI: 57.9–75.4%)
D	0000		ALIO: 0. 1	Outest	400 (00 1)	Proportion never/unsure if tested: 20.7% (95%CI: 13.2-28.3%)
Paquette 2013	2008	cross-sectional	AUS: Sydney	Substance use	120 (20.4)	No difference compared to non-Indigenous
Ward 2013	2007-2008	cross-sectional	AUS: NSW	Youth	293 (100)	Proportion tested for STIs incl. HIV in past year: 58%
Pearson 2013	2005-2007	cross-sectional	US: 7 cities	MSM	174 (100)	Proportion tested in past 3 months: 67 (45.6%)
Sionean 2014	2010	cross-sectional (NHBSS)	US: National	Heterosexual	56 (0.6)	Proportion tested in past year: 14/56 (25%); (men: 25%; women: 25) Proportion ever tested: 42/56 (75%); (men: 67%; women: 81)
Long 2014	2003-2012	cohort (population)	CAN: Alberta	Co-morbid (TB)	182 (12.5)	Proportion tested in routine screening in TB care:167/182 (91.8%)
LUNY 20 14	2000-2012	conort (population)	AUS: Western	SO-MOIDIG (TD)	102 (12.0)	Proportion tested in routine screening in 18 care: 167/162 (\$1.6%)
Williams 2015	2010	cross-sectional	Australia	Youth	244 (100)	Proportion ever tested: 46%
					\/	Late diagnosis (Stage 3 AIDS at diagnosis):
						AI/AN: 15 (34.1%)
Nwangwu-lke 2015	2012	cohort (NHSS)	US: National	Women	731 (0.3)	Native Hawaiian: 4 (40%)
						Proportion tested in past year: 101 (6.5%)
						Model: Odds of recent HIV testing higher for AI/AN 50-54 (AOR:2.47;
Fand 2045	2010	cross-sectional	LIC. National	Olden edults	4540 (4.4)	95%CI:1.09–5.58) and 60-64 (AOR:2.99; 95%CI:1.50–5.97) vs white people
Ford 2015	2010	(BRFSS)	US: National	Older adults	1542 (1.1) 15260 STI	the same age. No differences among 55-59 year olds.
Ward 2016	2010-2014	trial (STRIVE)	AUS: 65 remote communities	Young people	tests (100)	Proportion tested within 30 days of positive STI test: 4858/15260 (31.8%)
Anastario 2017	?	cross-sectional	US: Fort Peck	Substance use	51 (100)	Proportion tested within 30 days of positive \$11 test: 4636/13260 (\$1.6%) Proportion tested in past year: 66%
,	•	cross-sectional	JULI GUN	Substanto use	31 (100)	
Cahn 2017	2006-2010	(NSFG)	US: National	Men	923 (14.8)	Proportion tested in past year: 11.5%
-		` '			, -,	

LINKAGE TO C	ARE					
First Author, Publication Year	Study period	Study design	Location	Population	Sample Indigenous n (%)	- Key cascade findings
Diamond 2001	1989-1998	chart review	US: Seattle / Denver / LA	PLWHA	151 (1.3)	Presented late (AIDS diagnosis): 58 (38.0%) Significantly less likely than non-Indigenous p=0.07
Plitt 2009		cohort (NAHIVP)	CAN: Northern Alberta	newly diagnosed PLWHA	216 (41.1)	Median time to care (days): 38 (95%CI: 30-44) Model: Aboriginal ethnicity marginally significant correlate of longer times to care (AHR: 0.82; 95% CI 0.68–1.01)
Prentice 2011	2004-2005	cross-sectional	CAN: National	Youth	413 (100)	Never accessed care following diagnosis: 13/25 (52%) Range of time to care: within 3 months to after 2 years
Dickson 2012	2005-2010	cohort (AIDS Epidemiology Group)	NZ: National	PLWHA	65 (8.6)	Presented late (CD4 <350 cells/mL or AIDS-defining event): 30/65 (46.1%) MSM: 20/49 (40.8%); hetero: 10/16 (62.5%) Model: Maori MSM were more likely to present to care with 'advanced HIV disease' compared with European settler MSM (RR: 2.1, 95%CI: 1.4–3.2)
Connel 2014	?	chart review	US: Oklahoma	APHA	112 (100)	Presented late (CD4 <200/µL): 27 (24%)
Nwangwu-Ike 2015	2012	cohort (NHSS)	US: National	Women	731 (0.3)	>=1 lab test within 3 mo of diagnosis Al/AN: 4 (100%) Native Hawaiian: 4 (80.0%)
Kronfli 2017	2013-2015	cross-sectional (CHIWOS)	CAN: BC / Ontario / Quebec	Women	288 (20.2)	Model: Indigenous women more likely to have never accessed care (aOR: 4.30; 95%Cl: 1.31-14.13) and to have delayed linkage to care (aOR: 2.04; 95%Cl: 1.42-2.92), vs. Caucasian women.
Mitsch 2017	2011-2013	cross-sectional (NHSS)	US: 28 jurisdictions	APHA	1233 (100)	>=1 lab test within 1mo of diagnosis among those diagnosed in 2013: 43/58 (74.1%)

RETENTION IN	N CARE					
					Sample	-
First Author,	Study				Indigenous	
Publication Year	period	Study design	Location	Population	n (%)	Key Cascade Findings
						Received medical care in the past 4 months: 24 (86%)
Duran 2000	1997-1998	cross-sectional	US: Western	rural APHA	28 (100)	Avg. number of clinic visits: 4
Diamand 2004	1000 1000	alaant nasiiassi	US: Seattle /	DLWIIA	454 (4.2)	Mean annual outpatient visits:13
Diamond 2001	1989-1998	chart review	Denver / LA	PLWHA	151 (1.3)	Significantly lower than non-Indigneous p=0.001
		cross-sectional (HIV				Ever recieved VL monitoring: 23/25 (92.0%)
Grierson 2004	2001-2002	Futures Survey)	NZ: National	APHA	25 (100)	Avg. frequency of VL monitoring past year: 3.1
			CAN: Northern	ART naïve		No viral load monitoring in 6-months after baseline: 28/172 (16%)
Martin 2010	1999-2005	cohort (NAHIVP)	Alberta	PLWHA	172 (37.3)	Significantly higher than non-Indigenous (p=0.0059)
					•	Odds of missing at least one appointment
						Aboriginal: AOR: 1.2; 95%CI: 0.7-2.1
Erickson 2015	? - 2013	chart review	CAN: Winnipeg	PLWHA	218 (38.7)	Aboriginal PWUD: AOR: 3.33; 95%CI: 1.39–7.97
						Laboratory monitoring (CD4): 2.76 tests/year
						Significantly lower than non-Indigenous p=0.025
						Laboratory monitoring (VL): 2.53 tests/year
Templeton 2015	1999-2013	cohort	AUS: National	PLWHA	42 1.9)	Significantly lower than non-Indigenous p<0.001
· ·					,	>=1 lab tests in 2011
						AI/AN: 65 (47.1%)
						Native Hawaiian: 47 (51.6%)
						Model: Al/AN women less likely than white to have 1+ lab test (PR: 0.74;
						95%CI:0.62-0.88)
						>=2 lab tests at least 3 months apart
						AI/AN: 46 (33.3%)
						Native Hawaiian: 34 (37.4%)
Nwangwu-Ike						Model: Al/AN women less likely than white to have 2+ lab tests (PR: 0.70;
2015	2011	cohort (NHSS)	US: National	Women	731 (0.3)	95%CI:0.55-0.89)
			CAN: BC /		(5.5)	Model: No significant differences between Indigenous vs. Caucasian
		cross-sectional	Ontario /			women in terms of having been linked to care but not having received HIV
Kronfli 2017	2013-2015	(CHIWOS)	Quebec	Women	288 (20.2)	medical care in the past year.
		cross-sectional	US: 28			
Mitsch 2017	2011-2013	(NHSS)	jurisdictions	APHA	1233 (100)	>=2 CD4 or VL tests >=3 months apart in 2012: 551/1175 (46.9%)
						Continuous care (>=2 VL result >=90 days apart/yr): 103/175 (58.9%) Discontinuous care (>=1 yr with only 1 VL result): 55/175 (31.4%)
						Gap in care (a yr without a VL result): 17/175 (9.7%)
Rachlis 2017	2000-2012	cohort (OHTN)	CAN: 10 clinics		175 (9.5)	Model: Indigenous more likely to experience discontinuous care (aOR: 1.91; 95% CI: 1.28–2.84) vs. White.
Nacillis 2017	2003-2012	CONOIL (OLLIN)	OAN. TO CHITICS	I FANITY	110 (0.0)	1.01, 0070 OI. 1.20-2.04) VS. WIIILE.

ART INITIATIO	ON					
					Sample	
First Author,	Study				Indigenous	
Publication Year	period	Study design	Location	Population	n (%)	Key Cascade Finding
						Ever on ART: 7 (70%)
						Not different than non-Indigenous p=0.540
				PLWHA who		Duration on ART (median days): 97 (30-1416)
Goldstone 2000	1997	chart review	CAN: Vancouver	died	10 (16.1)	Significantly lower than non-Indigenous p=0.014
			US: Seattle /			on ART: 56%
Diamond 2001	1989-1998	chart review	Denver / LA	PLWHA	151 (1.3)	Not different than non-Indigenous p=0.8
						Ever on ART: 81/149 (54.4%)
						Significantly less than non-Indigenous p=0.001
						Model: Aboriginal people significantly less likely than other
				PLWHA who		ethnicities to have ever received ART (AOR: 0.60; 95%CI:
Wood 2003	1995-2001	cohort	CAN: BC	died	149 (12.0)	0.41–0.87; p=0.008)
		cross-sectional (HIV				Never been on ART: 7/25 (28.0%)
Grierson 2004	2001-2002	Futures Survey)	NZ: National	APHA	25 (100)	Currently on ART: 16/24 (66.7%)
				ART naïve		AIDS diagnosis at baseline: 20 (14%)
Miller 2006	1996-2002	cohort (BC DTP)	CAN: BC	PLWHA	146 (16.4)	Not different than non-Indigenous p=0.361
						On ART at 2 years: 29.2%
						Model: Aboriginal participants initiated ART at a significantly
Wood 2006	1996-2003	cohort (BART)	CAN: Vancouver	Substance use	149 (47.8)	slower rate ARH: 0.37; 95%CI: 0.15-0.93; p=0.035
						Ever been on ART: 146/185 (78.9%)
_					,,,	Ever been on HAART: 129/185 (70.1%)
Gorgos 2006		chart review	US: Phoenix	APHA	235 (100)	Currently on HAART: 107/185 (59.1%)
			ALIC: Mastana			
Nauraan 2007	2002	arana anational	AUS: Western	ADLIA	20 (400)	On ART: 11/00 (EE 00/.)
Newman 2007	2003	cross-sectional	Australia US: Navajo	APHA	20 (100)	On ART: 11/20 (55.0%)
Duran 2010	2003-2007	program evaluation	Nation	Case mgmt	46 (100)	On ART: 75%
Prentice 2011		cross-sectional	CAN: National	Youth	413 (100)	Ever on ART: 6/25 (24.0%)
Frende 2011	2004-2003	CIOSS-SECTIONAL	CAN. National	Touti	413 (100)	on ART: 51/80 (64%)
						First Nations: 60.0%; Metis or Inuit: 70.0%
Monette 2009 &				off-reserve		men: 70%; women: 61%
Beswick 2013	2006	cross-sectional (PSHP)	CAN: Ontario	APHA	80 (100)	lgbtq: 70%; hetero: 60%
DC3WIOR Z010	2000	cross sectional (i Crii)	Of IIV. Officiallo	Co-morbid	00 (100)	195tq. 7070, Notoro. 0070
Cain 2013	_	cross-sectional	CAN: National	(depression)	72 (100)	Currently not on ART: 28/72 (39%)
Connel 2014		chart review	US: Oklahoma	APHA	112 (100)	On ART at baseline: 35 (31%)
Conner 2014		Chartieview	OS. Oklarionia	ALIIA	112 (100)	On ART at baseline. 33 (3170)
Templeton 2015	1999-2013	cohort (prospective)	AUS: National	PLWHA	42 (1.9)	ART naive at enrolment: 14/42 (33.3%)
. 31110101011 2010	1000 2010	σσ.ιστε (ρισοροσανο)		. = ******	(1.0)	Currently not on ART: 20%
Jaworsky 2016	2007-2011	cross-sectional (OHTN)	CAN: Toronto	PLWHA	70 (9.5)	Not different than Caucasian p=0.15
Klakowicz 2016	2007-2012	pre-post (VNHS)	CAN: Vancouver		323 (59.2)	Ever been on ART while a patient: 280/323 (87%)
		/			\ - /	Ever been on ART at baseline: 234/326 (71.8%)
						On ART in past 6 months: Increased from 51-94%
						Model: No difference in likelihood of Indigenous vs non-
					/	Indigenous participants being dispensed ART in the
Milloy 2016	2005-2014	cohort (ACCESS)	CAN: Vancouver	Substance use	326 (38.6)	previous 180 days (aOR: 1.32; 95%CI: 0.92-1.89).

					Sample	_
First Author, Publication Year	Study period	Study design	Location	Population	Indigenous n (%)	Key Cascade Finding
					()	Received ART >=75% of the time: 23/81 (28.4%)
				PLWHA who		Model: No differences in time to ART discontinuation were
Wood 2003	1995-2001	cohort	CAN: BC	died	149 (12.0)	observed by ethnicity (ARH:1.19; 95%CI:0.87-1.61)
		sectional (HIV			•	
		Futures				Avg. doses taken per day: 2.4 (Range 0-6; Median: 2)
Grierson 2004	2001-2002		NZ: National	APHA	25 (100)	Ever took break from Tx: 8/16 (50.0%)
						5 day ratio: 33 (83%)
						SOAR: 22 (55%)
Ka'opua 2004	?	cross-sectional	US: Hawaii	PLWHA	40 (50.0)	No difference for either compared to White
	-	cohort (BC		ART naïve	(0010)	Adherent in first year on ART: 75-100%
Miller 2006	1996-2002	,	CAN: BC	PLWHA	146 (16.4)	Significantly lower than non-Indigenous p=0.001
		cohort (BC		ART naïve		Adherence <95%: 58/88 (66%)
Lima 2006	1996-1999	,	CAN: BC	PLWHA	88 (14.1)	No difference compared to non-Indigenous (p=0.2995)
		program	USA: Navajo			,
Duran 2010	2003-2007		Nation	Case mgmt	46 (100)	No missed doses in p4w: 24/34 (71%)
Iralu 2010		cross-sectional		rural APHA	36 (100)	Excellent adherence: 19/29 (61.3%)
	20012000	0.000 000.0.10.	00.000		00 (100)	% doses taken in past week (mean, SE): 91% (3)
				PLWHA on		% doses taken in past month (mean, SE): 94% (2)
Gunther 2014	2010	cross-sectional	CAN: Edmonton	ART	23 (35.4)	No difference compared to non-Indigenous
		cohort study			7	Tx interruption >=90 days: 46 (21.8%)
Samji 2014	2007-2010	,	CAN: BC	PLWHA	211 (27.5)	Model: Indigenous ethnicity not associated with Tx interruption
•		,				Tx interruption >=90 days: 242/414 (58.5%)
		cohort	CAN: BC /	ART naïve		Model: Higher risk of Tx interruption among Aboriginal vs non-
Samji 2015	2000-2011	(CANOC)	Ontario / Quebec	PLWHA	414 (5.4)	Aboriginal (aHR: 1.67, 95%CI: 1.27–2.20)
		,			, ,	Adherence >=95% at baseline: 107/326 (32.8%), remained
						similar over study period
		cohort				Model: No difference between Indigenous and non-Indigenous
Milloy 2016	2005-2014	(ACCESS)	CAN: Vancouver	Substance use	326 (38.6)	in adherence in past six months (aOR: 0.90; 95%CI: 0.73-1.10
•		,				Adherence >=95% (pharmacy refill), mean proportion (SD)
						No IDU history: Men: 71.3 (5.4); Women: 59.5 (7.1)
						With IDU history: Men: 77.1 (1.6); Women: 57.0 (2.3)
		cohort				Model: Indigenous people less likely to have optimal adherence
Puskas 2017	2000-2014	(HOMER)	CAN: BC	PLWHA	589 (13.0)	compared to non-Indigenous people (p<0.001)
		, ,			, ,	Tx interruption >=90 days: 617/1022 (60.4%)
		cohort (BC				Higher risk of Tx interruption among Aboriginal vs. non-
Wang 2017	1996-2015	DTP)	CAN: BC	PLWHA	1022 (12.6)	Aboriginal (p<0.001)

VIRAL SUPPRES	30.014				Sample	
First Author	Ctudy					_
First Author, Publication Year	Study period	Study design	Location	Population	Indigenous n (%)	Key Cascade Finding
T abrication Tour	portou	Otady doolgii	Location	PLWHA	11 (70)	noy babbaab i manig
Goldstone 2000	1997	chart review	CAN: Vancouver	who died	10 (16.1)	Median (IQR) viral load last reported: 32,000 (22,000 - 54,000)
		cross-sectional				
		(HIV Futures				
Grierson 2004	2001-2002	Survey)	NZ: National	APHA	25 (100)	Mean copies/mL (range): 558 (<500 to 2.5 million)
				ADT noïve		Median (IQR) viral load at baseline: 71,950 (25,000-151,000)
Miller 2006	1996-2002	cohort (BC DTP)	CAN: BC	ART naïve PLWHA	146 (16.4)	Remain unsuppressed: 24% Model: Aboriginal ethnicity not associated with suppression.
Willier 2000	1990-2002	conort (BC DTI)	CAN. BO	ART naïve	140 (10.4)	Median (IQR) viral load at baseline: 89,350 (21, 750 – 185,000)
Lima 2006	1996-1999	cohort (BC DTP)	CAN: BC	PLWHA	88 (14.1)	Significantly lower compared to non-Indigneous (p<0.001)
	(cohort	AUS: Western			Proportion suppressed prior to delivery (<50 copies/mL): 11 (48%
Gilles 2007	1991-2005	(retrospective)	Australia	Perinatal	16 (39.0)	No difference compared to non-Indigenous (p=0.183)
Iralu 2010	2004-2006	cross-sectional	US: Southwest	rural APHA	36 (100)	Median log viral load: 2.3 (2.3, 7.3)
						Median (IQR) copies/mL at baseline: 105,000 (38,000-405,000)
						Did not suppress within 6mo: 65/172 (37.8%)
			CAN: Northern	ART naïve		Model: Odds of suppression (vs non-Aboriginal non-IDU) Aboriginal IDU: OR:0.33, 95%CI:0.19-0.60, p=0.0002
Martin 2010	1999-2005	cohort (NAHIVP)	Alberta	PLWHA	172 (37.3)	Aboriginal non-IDU: OR:0.38, 95%CI:0.21-0.67, p=0.0009
		/	CAN: Northern	ART naïve	\/	Median (IQR) copies/mL at baseline: 100,000 (22,000-390,000)
Martin 2011	1999-2005	cohort (NAHIVP)	Alberta	PLWHA	194 (35.4)	No difference compared to non-Indigenous (p=0.46)
						Median (IQR) copies/mL
						Aboriginal IDU: 50 (50-44,000)
						Aboriginal non-IDU: 50 (50-71)
						Significantly higher (p=0.019) >400 copies/mL
						Aboriginal IDU: 7 (33%)
		cross-sectional	CAN: Northern			Aboriginal non-IDU: 2 (15%)
Martin 2013	2006-2007	(NAHIVP)	Alberta	PLWHA	34 (35.4)	Marginally higher (p=0.058)
						Median (IQR) copies/mL at baseline: 71,100 (19,700-151,000)
ART-CC 2013	1998-2009	cohort (HOMER)	CAN: BC	PLWHA	205 (0.4)	Significantly lower than non-Indigenous (p<0.001)
0 0011	4007 0040		CAN- DO	DLAMIA	FFO (O.4)	Proportion unsuppressed: 78 (14%)
Cescon 2014	1997-2010	cohort (BC DTP)	CAN: BC	PLWHA	558 (9.1)	Significantly higher than non-Indigenous (p<0.001) Mean (range) copies/mL: 137,807 (43–750,000)
Connel 2014	?	chart review	US: Oklahoma	APHA	112 (100)	Proportion suppressed at entry into care: 20/112 (17.9%)
000. 2011			oor omanoma	7.1.7.7	()	Median (IQR) copies/mL at baseline: 32,000 (8600-98000)
						No VL sup within 6 mo of ART initiation: 24/87 (27.5)
						Model: Aboriginal less likely to suppress within 6mo of initiating ART
Lafabura 2014	2006 2012	achort (NIALII) (D)	CAN: Northern	ART naïve	07 (07 0)	vs. Canadian-born non-Aboriginal (AOR:0.44; 95%CI: 0.20-0.96;
Lefebvre 2014	2006-2012	cohort (NAHIVP)	Alberta	PLWHA	87 (27.0)	p=0.04). No difference in odds of virological failure by ethnicity. Proportion suppresed (<400 copies/ml 12mo post-ART): 60.0%
						Proportion suppresed (<400 copies/ml 24mo post-ART): 81.0%
Templeton 2015	1999-2013	cohort (prospective)	AUS: National	PLWHA	42 (1.9)	No difference compared to non-Indigenous p=0.635
•				Substance	, ,	Proportion never been suppressed: 39/152 (25.7%)
Erickson 2015	? - 2013	chart review	CAN: Winnipeg	use	218 (38.7)	Significantly more likely than non-Indigenous p=0.008
						Proportion suppressed (<=200 copies/mL)
						Al/AN: 41 (29.7%)
		cohort (National				Native Hawaiian: 36 (39.6%)
Nwangwu-lke 2015	2008-2012	HIV Surveillance System)	US: National	Women	731 (0.3)	Model: Al/AN less likely to be suppressed vs white (PR: 0.64; 95%CI: 0.49-0.83). No difference between Native Hawaiian and white.
imaigwa ike 2013		cross-sectional	CO. Hadional	TTOINGII	, 01 (0.0)	Proportion suppressed: 76%
Jaworsky 2016	2007-2011		CAN: Toronto	PLWHA	70 (9.5)	No difference compared to non-Indigenous p=0.88
						Median (IQR) copies/mL at baseline: 196 (35-16531)
Klakowicz 2016	2007-2012	pre-post (VNHS)	CAN: Vancouver	PLWHA	323 (59.2)	Proportion suppressed within 1 year of HAART: 128 (40%)
				04		Proportion suppressed (<50 copies/mL): Increased from 23-65%
Milloy 2016	2005 2014	cohort (ACCESS)	CAN: Vancouver	Substance	326 (38.6)	Model: No difference between Indigenous and non-Indigenous in terms of being undetectable (aOR: 1.15; 95%: 0.91-1.46)
Milloy 2016	2000-2014	cohort (ACCESS)	CAN: Vancouver	use	020 (00.0)	terms of being undetectable (aOrt. 1.10, 30%, 0.81-1.40)
Benoit 2017 (a)	2000-2012	cohort (CANOC)	Ontario / Quebec	PLWHA	497 (7.0)	Log10 copies/mL at baseline: 4.85 (4.38–5.00)
_ 5.1.0.1 = 0 11 (u)			3	. =	(1.0)	Suppression at 1yr after ART initiation: 54%
						Rebound at 1 yr after ART initiation: 13%
						Model: Indigenous less likely to achieve suppression vs. Caucasian
						(aHR=0.58, 95% CI 0.50-0.68, p<0.0001). Risk of rebound not
			CAN: BC /			significantly different between Indigenous and Caucasian (aHR=1.03,
Benoit 2017 (b)		cohort (CANOC)	Ontario / Quebec	PLWHA	497 (7.0)	95% CI 0.84, 1.27, P=0.75).
Mitsch 2017		cross-sectional (NHSS)	US: 28	APHA	1233 (100)	Proportion suppressed (<200 conice/ml): 530/4175 (45 49/)
Mitsch 2017	2012	(141 100)	jurisdictions	ALLIA	1200 (100)	Proportion suppressed (<200 copies/mL): 530/1175 (45.1%)