# Recent detectable viral loads among adults living with HIV in the Asia-Pacific between 2015 to 2020



Maylin Palatino<sup>1</sup>, Dhanushi Rupasinghe<sup>2</sup>, Alvina Widhani<sup>3</sup>, Ngoc Chi<sup>4</sup>, I Ketut Agus Somia<sup>5</sup>, Nagalingeswaran Kumarasamy<sup>6</sup>, Suwimon Khusuwan<sup>7</sup>, Vohith Khol<sup>8</sup>, Man Po Lee<sup>9</sup>, Sasisopin Kiertiburanakul<sup>10</sup>, Thach Ngoc Pham<sup>11</sup>, Sanjay Pujari<sup>12</sup>, Anchalee Avihingsanon<sup>13</sup>, Jun Yong Choi<sup>14</sup>, Romanee Chaiwarith<sup>15</sup>, Yu-Jiun Chan<sup>16</sup>, Iskandar Azwa<sup>17</sup>, Rossana Ditangco<sup>18</sup>, Rohidas T Borse<sup>19</sup>, Oon Tek NG<sup>20</sup>, Junko Tanuma<sup>21</sup>, Fujie Zhang<sup>22</sup>, Yasmin Gani<sup>23</sup>, Jeremy Ross<sup>24</sup>, and Awachana Jiamsakul<sup>2</sup>, on behalf of the International Epidemiology Databases to Evaluate AIDS (IeDEA) Asia-Pacific

<sup>1</sup>University of Maryland, USA, <sup>2</sup>The Kirby Institute, UNSW Sydney, Australia, <sup>3</sup>Faculty of Medicine Universitas Indonesia - Dr. Cipto Mangunkusumo General Hospital, Jakarta, Indonesia, <sup>4</sup>Bach Mai Hospital, Vietnam, <sup>5</sup>Faculty of Medicine Udayana University - Prof. Dr. I.G.N.G. Ngoerah Hospital, Indonesia, <sup>6</sup>CART CRS, Voluntary Health Services, India, <sup>7</sup>Chiangrai Prachanukroh Hospital, Thailand
<sup>8</sup>National Center for HIV/AIDS, Dermatology & STDs, Cambodia, <sup>9</sup>Queen Elizabeth Hospital, Hong Kong SAR, <sup>10</sup>Mahidol University, Thailand, <sup>11</sup>National Hospital for Tropical Diseases, Vietnam, <sup>12</sup>Institute of Infectious Diseases, India, <sup>13</sup>Faculty of Medicine, Chulalongkorn University, Thailand, <sup>14</sup>Yonsei University College of Medicine, South Korea, <sup>15</sup>Chiang Mai University, Thailand, <sup>16</sup>Taipei Veterans General Hospital, Taiwan, <sup>17</sup>University of Malaya, Malaysia, <sup>18</sup>Research Institute for Tropical Medicine, Philippines, <sup>19</sup>BJ Government Medical College and Sassoon General Hospital, India, <sup>20</sup>Tan Tock Seng Hospital, Singapore, <sup>21</sup>National Center for Global Health and Medicine, Japan, <sup>22</sup>Capital Medical University, China, <sup>23</sup>Hospital Sungai Buloh, Malaysia, <sup>24</sup>TREAT Asia, amfAR - The Foundation for AIDS Research, Bangkok, Thailand

#### BACKGROUND

#### RESULTS

A proportion of people living with HIV (PLWH) continue to have detectable viral load (VL) while on robust antiretroviral therapy (ART) regimens and risk onward HIV transmission. We aimed to estimate the proportion and factors associated with having a detectable VL between 2015 and 2020 in two Asia-Pacific HIV observational cohorts.

Figure 1. Asia-Pacific countries and territories included in TAHOD and TAHOD-LITE cohorts



### METHODS

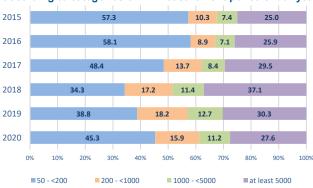
This analysis included available data from adult PLWH enrolled in two prospective observational cohort studies of IeDEA Asia-Pacific: the TREAT Asia HIV Observational Database (TAHOD) and TAHOD Low Intensity Transfer (TAHOD-LITE) cohorts (Figure 1). Eligible participants were on ART for  $\geq$ 1 year, in follow-up between 2015 and 2020, and with  $\geq$ 1 VL measurement during the follow-up period.

Detectable VL was defined as having  $\geq 1$  VL measurement  $\geq 50$  copies/mL during 2015-2020. The proportion with detectable VL during the follow-up period and the distribution of VL measurements per year were obtained. The proportions with viral blips, i.e., detectable VL measurements followed by a VL  $\leq 50$  copies/mL on retesting within 6 months, were likewise computed. Factors associated with detectable VL were analyzed using repeated measures logistic regression. Of the 20,765 PLWH included, the majority were male (64%) and had heterosexual contact as their mode of HIV exposure (72%). The median age was 36 (Table 1).

Characteristics	Total PLWH	PLWH who never had detectable VL	PLWH who ever had detectable VL (≥50 copies/mL)
Total, N (%)	20,765 (100)	14,727 (71)	6,038 (29
Sex, N (%) Males Females	13,376 (64) 7,389 (36)	9,290 (63) 5,437 (37)	4,086 (68 1,952 (32
Age (years) at ART initiation, median (IQR)	36 (30-43)	36 (30-43)	36 (30-42
HIV Exposure, N (%) Heterosexual contact MSM Injecting drug use Other/Unknown	15,012 (72) 2,985 (14) 755 (4) 2,013 (10)	10,454 (71) 2,322 (16) 612 (4) 1,339 (9)	4,558 (75 663 (11 143 (2 674 (11
CD4 cell count (cells/µl) at ART initiation, median (IQR)	168 (59-287)	174 (54-299)	158 (67-262
Viral Load (log <sub>10</sub> (copies/mL)) at ART initiation, median (lQR)	2 (2-3)	2 (2-3)	2 (2-2
Hepatitis B surface antigen, N (%) Negative Ever positive Not reported	13,682 (66) 1,451 (7) 5,632 (27)	10,388 (70) 1,138 (8) 3,201 (22)	3,294 (55 313 (5 2,431 (40
Hepatitis C antibody, N (%) Negative Ever positive Not reported	12,802 (62) 1,418 (7) 6,545 (31)	10,405 (68) 1,131 (8) 3,551 (24)	2,757 (45 287(5 2,994 (50
World Bank country income, N (%) Lower Middle Upper Middle High	10,215 (49) 7,514 (36) 3,036 (15)	6,035 (41) 6,410 (44) 2,282 (16)	4,180 (69 1,104 (18 754 (13

During the study period, 6,038 (29%) had  $\geq$ 1 detectable VL. Of these, 2,881 (48%) had VL measurements between 50 and <200 copies/mL, 684 (11%) between 200 and <1000, 495 (8%) between 1000 and <5000, and 1,978 (33%)  $\geq$ 5000 copies/mL. The highest and lowest proportions with VL measurements  $\geq$ 5,000 copies/mL were in years 2018 (37%) and 2015 (25%), respectively (Figure 2). The proportions with viral blips varied over time: 2015 (10%), 2016 (11%), 2017 (13%), 2018 (17%), 2019 (21%), and 2020 (14%).





The factors associated with decreased odds of detectable VL included female sex compared to males; older age compared to  $\leq$ 30 years; male-male sex and injecting drug use compared to heterosexual contact as mode of HIV exposure; non-nucleoside reverse transcriptase inhibitor-based ART (NNRTI) compared to integrase strand transfer inhibitor (INSTI)-based regimens; hepatitis B or hepatitis C co-infection, higher CD4 count compared to CD4 <200 cells/µL; and higher country income compared to lower-middle. Detectable VL was more likely among those on protease inhibitor-based ART (Figure 3).

## Figure 3. Factors associated with having detectable viral load (≥50 copies/mL) using repeated measures

Covariate		Adj OR (95% CI)
Sex		
Female vs Male	-	0.84 (0.78, 0.90)
Time-updated age (years)	1	
31-40 vs at most 30	-#-	0.68 (0.60, 0.76)
41-50 vs at most 30		0.56 (0.49, 0.63)
at least 51 vs at most 30	+	0.43 (0.38, 0.49)
HIV mode of exposure	1	
MSM vs heterosexual contact		0.86 (0.75, 0.97)
Injecting drug use vs heterosexual contact		0.65 (0.52, 0.83)
Other/Unknown vs heterosexual contact	-	0.58 (0.52, 0.65)
Time-updated ARV Regimen		
PI-based vs INSTI-based		
NNRTI-based vs INSTI-based	-	0.47 (0.43, 0.52)
Other/No ART vs INSTI-based		0.99 (0.78, 1.25)
HBV surface antigen		
Ever positive vs negative		0.80 (0.70, 0.92)
HCV antibody		
Ever positive vs negative		0.81 (0.69, 0.97)
Time-updated CD4 (cells/microliter)	1	
200-350 vs <200	+	0.34 (0.30, 0.38)
351-500 vs <200		0.20 (0.18, 0.23)
>500 vs <200		0.17 (0.15, 0.18)
World Bank country income		
Upper Middle vs lower Middle		0.22 (0.20, 0.24)
High vs lower Middle		0.36 (0.32, 0.40)
Time-updated calendar year		
2016 vs 2015	-	0.81 (0.76, 0.86)
2017 vs 2015		0.60 (0.56, 0.64)
2018 vs 2015		0.39 (0.36, 0.42)
2019 vs 2015		0.37 (0.34, 0.40)
2020 vs 2015	<b>.</b>	0.34 (0.31, 0.38)
	0.2 0.5 0.8 1	
	Adjusted odds r	atio

### CONCLUSIONS

Almost one third of PLWH in our analysis had detectable VL between 2015 and 2020, of these, a third had VL measurements ≥5,000 copies/mL, indicating the need for strengthening life-long adherence counselling and follow-up, particularly among those with increased odds of detectable VL.

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The TREAT Asia HIV Observational Database: V Khol\*, V Ouk, C Pov, National Center for HIV/AIDS, Dermatology & STDs, Phnom Penh, Cambodia; FJ Zhang\*, HX Zhao, N Han, Beijing Ditan Hospital, Capital Medical University, Beijing, China; MP Lee\*, PCK Li, TS Kwong, TH Li, Queen Elizabeth Hospital, Hong Kong SAR; N Kumarasamy\*, C Ezhilarasi, Chennai Antiviral Research and Treatment Clinical Research Site (CART CRS), VHS-Infectious Diseases Medical Centre, VHS, Chennai, India; S Pujari\*, K Joshi, S Gaikwad, A Chitalikar, Institute of Infectious Diseases, Pune, India; RT Borse\*, V Mave, I Marbaniang, S Nimkar, BJ Government Medical College and Sassoon General Hospital, Pune, India; IKA Somia\*, TP Merati, AAS Sawitri, F Yuliana, Faculty of Medicine Udayana University - Prof.Dr.I.G.N.G. Ngoerah Hospital, Bali, Indonesia; E Yunihastuti\*, A Widhani, S Maria, TH Karjadi, Faculty of Medicine Universits Indonesis. Or. Cipto Mangunkusumo General Hospital, Jakara, Indonesia; I Tanuma\*, S Oka, T Nishijima, National Center for Global Health and Medicine, Tokyo, Japan; YC Loh\*, Na S, JM Kim, Division of Infectious Diseases, Department of Internal Medicine, Yonea; VI Gani\*, NB Rudi, Hospital Suloh, Surgai Bluoh, Surgai Bluoh, Surgai Bluoh, Surgai Bluoh, Surgai Suloh, Xalaysia; I Zawa\*, A Kamarutzaman, SF Syed Omar, S Ponnampalavanar, University Malaya Medical Centre, Kuala Lumpur, Malaysia; R Ditangoc\*, MK Pane; Nesvea: A Institute for Tropical Medicine, Muntinlupa City, Philippines; YJ Cha\*, HP Chen, PC Wu, E Ke, Taipei Veterans General Hospital, Taipei, Taipei







