

Vital Surveillances

Changing Proportions of HIV-1 Subtypes and Transmitted Drug Resistance Among Newly Diagnosed HIV/AIDS Individuals — China, 2015 and 2018

Jingjing Hao¹; Shan Zheng¹; Mengze Gan¹; Aobo Dong¹; Ruihua Kang¹; Miaomiao Li¹; Shuai Zhao¹; Jing Hu¹; Chang Song¹; Lingjie Liao¹; Yi Feng¹; Yiming Shao¹; Yuhua Ruan¹; Hui Xing^{1,†}

ABSTRACT

Introduction: With the expansion of human immunodeficiency virus (HIV) antiretroviral therapy (ART), HIV drug resistance is becoming more and more serious. This study describes the changing prevalence of HIV-1 subtypes and transmitted drug resistance (TDR) among newly diagnosed individuals in China, 2015 and 2018.

Methods: A total of 8,980 individuals in 2015 and 2018 from 31 provincial-level administrative divisions (PLADs) were enrolled in this study. Viral RNAs were amplified and sequenced using an in-house polymerase chain reaction (PCR) protocol. The Stanford HIV Drug Resistance Database (HIVdb) was used to predict susceptibility to 12 antiretroviral drugs.

Results: The prevalence of TDR was not significantly increased over time. The prevalence of TDR was 3.8% and 4.4% in 2015 and 2018, respectively ($P=0.13$). The prevalence of CRF55_01B increased from 2.3% in 2015 to 3.9% in 2018 ($P<0.001$). The drug resistance prevalence of non-nucleoside reverse transcriptase inhibitors (NNRTI) increased from 2.4% in 2015 to 3.3% in 2018 ($P<0.01$). The prevalence of E138 ($P<0.001$), H221 ($P=0.03$), and V179 ($P<0.001$) mutations increased from 0.30%, 0.09%, and 0.70% in 2015 to 1.10%, 0.30%, and 1.70% in 2018, respectively.

Conclusions: HIV drug resistance affects the effect of antiretroviral treatment, so the monitoring of HIV TDR should be strengthened to control the transmission of HIV drug resistance.

INTRODUCTION

The four main human immunodeficiency virus (HIV)-1 subtypes in China over the past 20 years include CRF07_BC, CRF01_AE, CRF08_BC, and subtype B. The prevalence of CRF07_BC and

CRF01_AE has increased since 2006 and they remain the dominant subtypes in China present in 31 provincial-level administrative divisions (PLADs) (1). In 2012, a study reported the HIV-1 CRF55_01B subtype, composed of CRF01_AE and subtype B, which possibly originated in men having sex with men (MSM) in Shenzhen (2). After its origin in Shenzhen, it spread rapidly to all cities in China (3). Various factors contribute to the spread and diversification of HIV-1 subtypes in China, including the emergence of more dating apps, the rapid development of the economy and means of transportation, and people moving to different parts of the country for employment.

From 2012, several first-line antiretroviral drugs were available to HIV patients for free, including TDF/AZT+3TC+EFV/NVP. In 2014, only HIV patients with a cluster of differentiation 4 (CD4) cell count of less than 500 cells/ μ L were eligible to receive ART. The Chinese government adjusted the standard for free ART in 2016 and recommended that all HIV patients receive ART (4). Adherence to ART is critical to achieving viral suppression; however, poor drug adherence or withdrawal from treatment can result in the development of drug resistance (5). Moreover, the problem of drug resistance can be exacerbated by the expansion of ART regimens. One possible solution would be conducting TDR surveys, which can offer an effective guide for structuring future first- and second-line ART regimens.

METHODS

The study design was conducted according to the WHO protocol for TDR in each region of China. Depending on the number of newly reported HIV/AIDS cases, China was divided into high prevalence regions, moderate prevalence regions and low prevalence regions. At least 300 patients were recruited in high prevalence regions, moderate

prevalence regions and low prevalence regions (6–7). The individuals (16 years old or older at the time of HIV-1 diagnosis) diagnosed with HIV-1 infection in 2015 and 2018 were enrolled in this study. All individuals provided written informed consent. Study variables included the following epidemiological and clinical characteristics: age, sex, education, HIV-1 transmission route, and HIV-1 subtype. Plasma from participants was collected and sent to the laboratory via cold chain transport.

Viral RNAs were extracted from 200 µL of plasma using the QIAasymphony platform. The pol fragment (HXB2 positions 2,253–3,312, 1,060 bp) was amplified and sequenced using an in-house PCR protocol. The nucleotide sequences were aligned separately using the HIV Align tool (https://mafft.cbrc.jp/alignment/server/add_fragments.html). The aligned sequences were manually adjusted using BioEdit (version 7.0.9.1, Borland, CA, USA). Sequences less than 1,000 nucleotides in length were excluded from the analysis. HIV-1 subtypes were determined based on a neighbor-joining tree with Mega [Mega 7.0: Molecular Evolutionary Genetics Analysis across computing platforms (Kumar S, Stecher G, and Tamura K 2016)]. The Stanford HIV Drug Resistance Database (HIVdb) (<https://HIV-1db.stanford.edu/HIV-1db/by-sequences>) genotypic resistance interpretation system was used to predict susceptibility to 12 antiretroviral drugs recommended for use by the WHO (EFV, NVP, ABC, AZT, 3TC, TDF, FTC, D4T, DDI, LPV/r, ATV/r, and DRV/r). This system assigns a drug penalty score to each mutation. Scores for individual mutations are greater than -15: negative numbers indicate increased susceptibility compared to wild type virus; 0 indicates no change to susceptibility; susceptible (0–9), potential low-level resistance (10–14), low-level resistance (15–29), intermediate resistance (30–59), and high-level resistance (≥ 60). Drug resistance is defined as 15 points or more.

SAS (version 9.4, SAS Institute Inc., Cary, NC, USA) was used for statistical analysis. The χ^2 test was used to analyze variations of 12 types of drugs and drug resistance mutations in 2015 and 2018. Univariate and multivariate logistic regression tests were used to analyze the factors associated with drug resistance, with $P < 0.05$ considered as statistically significant.

RESULTS

A total of 8,980 individuals were diagnosed with

HIV-1 in 2015 and 2018, with 4,704 diagnosed in 2015 and 4,276 in 2018. The characteristics of the individuals included in this study are listed in Table 1. The proportions of HIV-1 individuals aged 26–49 and ≥ 50 years old were 56.8% and 24.8%, respectively. Approximately 81.9% of HIV-1 individuals were men. About 40.6% of HIV-1 individuals graduated from high school or had higher education. The main pattern of transmission was heterosexual intercourse (52.5%).

The most common HIV-1 subtype was CRF07_BC (39.3%), followed by CRF01_AE (36.2%), CRF08_BC (8.9%), B (4.3%), and CRF55_01B (3.1%) (Table 1). The prevalence of CRF55_01B increased from 2.3% in 2015 to 3.9% in 2018 ($P < 0.001$) (Table 2).

The TDR prevalence was 3.80% in 2015 and 4.40% in 2018; overall TDR prevalence did not change significantly between 2015 and 2018. TDR prevalence of NNRTI, NRTI, and PI changed from 2.40%, 1.10%, and 0.20% in 2015, to 3.30%, 0.80%, and 0.07% in 2018, respectively. The increased prevalence of NNRTI drug resistance was from 2.40% in 2015 to 3.30% in 2018 ($P < 0.01$), whereas no increase in drug resistance was observed for NRTI and PI. Moreover, the prevalence of certain mutations, including E138, H221, and V179, significantly increased from 0.30%, 0.09%, and 0.70% in 2015 to 1.10%, 0.30%, and 1.70% in 2018, respectively (Table 3).

Multivariate logistic regression analysis showed that CRF08_BC [adjusted odds ratio (AOR)=1.51, 95% confidence interval (CI): 1.03–2.23], CRF55_01B (AOR=4.17, 95% CI: 2.77–6.27), and URF (AOR=1.69, 95% CI: 1.14–2.49) were independent factors associated with TDR (Table 4).

DISCUSSION

In China, CRF07_BC and CRF01_AE were still the two main subtypes. The proportion of the CRF55_01B subtype increased from 2015 to 2018. As a new recombinant subtype formed by the recombination of CRF01_AE and subtype B, CRF55_01B prevalence increased exponentially from 2005 to 2009 after its origin in Shenzhen. One of the factors contributing to its rapid transmission was the spread of the virus to different PLADs in 2007, as mediated by the rapid development of the Beijing-Guangzhou and Beijing-Kowloon railways (8). Moreover, even though this particular subtype

TABLE 1. General characteristics of newly diagnosed HIV individuals in 2015 and 2018, China.

Variable	2015		2018		Total	
	Number of cases	Proportion (%)	Number of cases	Proportion (%)	Number of cases	Proportion (%)
Total	4,704	100.0	4,276	100.0	8,980	100.0
Age (years)						
16–25	862	18.3	676	15.8	1,538	17.1
26–49	2,749	58.5	2,350	55.0	5,099	56.8
≥50	1,003	21.3	1,227	28.7	2,230	24.8
Unknown	90	1.9	23	0.5	113	1.3
Sex						
Male	3,763	80.0	3,595	84.1	7,358	81.9
Female	876	18.6	681	15.9	1,557	17.4
Unknown	65	1.4	0	0.0	65	0.7
Education						
Primary school	1,338	28.5	993	23.2	2,331	26.0
Junior high school	1,412	30.0	1,205	28.2	2,617	29.1
Senior high school or above	1,769	37.6	1,877	43.9	3,646	40.6
Unknown	185	3.9	201	4.7	386	4.3
Route of HIV transmission						
HET	2,533	53.9	2,181	51.0	4,714	52.5
MSM	1,855	39.4	1,906	44.6	3,761	41.9
IDU	115	2.4	53	1.2	168	1.9
Others	201	4.3	136	3.2	337	3.7
Subtype						
CRF01_AE	1,835	39.0	1,578	36.9	3,413	38.0
CRF07_BC	1,675	35.6	1,696	39.7	3,371	37.5
CRF08_BC	418	8.9	384	9.0	802	8.9
CRF55_01B	110	2.3	168	3.9	278	3.1
B	261	5.6	183	4.3	444	5.0
Others	405	8.6	267	6.2	672	7.5

Abbreviations: HIV=human immunodeficiency virus; HET=heterosexual; MSM=men who have sex with men; IDU=injecting drug users.

TABLE 2. Changes in HIV subtypes among newly diagnosed HIV individuals in 2015 and 2018, China.

Subtype	2015		2018		P
	Number of cases	Proportion (%)	Number of cases	Proportion (%)	
CRF01_AE	1,835	39.0	1,578	36.9	0.04
CRF07_BC	1,675	35.6	1,696	39.7	<0.0001
CRF08_BC	418	8.9	384	9.0	0.88
CRF55_01B	110	2.3	168	3.9	<0.0001
B	261	5.6	183	4.3	0.006
Others	405	8.6	267	6.2	<0.0001

Abbreviation: HIV=human immunodeficiency virus.

originated in homosexual males, it was transmitted among heterosexuals. Currently, in-depth studies on CRF55_01B are lacking, especially as it relates to the

underlying mechanism of drug resistance.

The TDR prevalence was 3.80% in 2015 and 4.40% in 2018, with no significant increase over time. A study

TABLE 3. Changes of HIV mutations among newly diagnosed HIV individuals in 2015 and 2018, China.

Variable	2015		2018		P
	Number of cases	Proportion (%)	Number of cases	Proportion (%)	
Total	178	3.80	189	4.40	0.13
NNRTI	113	2.40	142	3.30	0.01
A98G	5	0.10	7	0.20	0.46
E138A/G/K/Q	13	0.30	45	1.10	<0.001
G190A	13	0.30	9	0.20	0.53
H221Y	4	0.09	12	0.30	0.03
K101E	15	0.30	11	0.30	0.59
K103N/S	34	0.70	34	0.80	0.69
L100I	1	0.02	1	0.02	1.00
P225H	1	0.02	5	0.10	0.11
V106I/M	16	0.30	23	0.50	0.15
V108I	15	0.30	8	0.20	0.22
V179A/D	35	0.70	72	1.70	<0.001
Y181C	7	0.20	11	0.30	0.25
NRTI	52	1.10	33	0.80	0.10
D67N	3	0.06	4	0.09	0.72
K219Q	3	0.06	2	0.05	1.00
K65R	7	0.10	7	0.20	0.86
K70E/R	8	0.20	8	0.20	0.85
L210W	7	0.20	3	0.07	0.35
L74I	2	0.04	2	0.05	1.00
M184I/V	15	0.30	9	0.20	0.32
M41L	9	0.20	5	0.10	0.37
T69D/N	6	0.10	0	0.00	0.03
V75M/A	2	0.04	2	0.05	1.00
Y115F	1	0.02	1	0.02	1.00
PI	7	0.20	3	0.07	0.26
I50V/L	1	0.02	1	0.02	1.00
I54L/M	1	0.02	0	0.00	1.00
M46I/L	3	0.06	2	0.05	1.00
F53L	1	0.02	0	0.00	1.00
I47V	2	0.04	0	0.00	1.00
L90M	0	0.00	1	0.02	0.96
L33F	3	0.06	1	0.02	0.63

Abbreviations: HIV=human immunodeficiency virus; NNRTI=non-nucleoside reverse transcriptase inhibitor; NRTI=nucleoside reverse transcriptase inhibitor; PI=protease inhibitor.

that included TDR prevalence data from 2001 to 2017 in China estimated that TDR prevalence was 4.1% (9), which is consistent with the results of this study. Reports on drug resistance testing from the WHO

revealed that between 2014 and 2019, the median rate of transmission-resistant drugs was 4.1% in Southeast Asia, 6.0% in Sub-Saharan Africa, 9.1% in Latin America, 8.5% in Europe, and 14.2% in North America (10). Conversely, the prevalence of TDR was much higher in Europe and North America, which was consistent with the longer existence and variety of antiviral treatment.

The prevalence of transmitted NNRTI resistance increased from 2015 to 2018. According to the analysis of resistance mutations, the prevalence of the E138, H221, and V179 mutations increased. V179E is a nonpolymorphic mutation weakly selected by NVP and EFV, E138G is another nonpolymorphic accessory mutation occasionally in patients receiving NVP and EFV (11). Molecular monitoring results of newly diagnosed individuals in China showed that the most common mutations in patients' resistance to NNRTI were E138G and V179E (12). Moreover, E138K was found to be the most common mutation in HIV-1 patients resistant to NNRTI in the US, according to an analysis of TDR from 2014 to 2018 (10). The impact of the combination of E138 and V179 on both NNRTI susceptibility and virologic outcome in patients deserves investigation.

CRF55_01B was associated with drug resistance. Moreover, V179E often appears in CRF55_01B together with E138G and has low resistance to EFV and NVP (13). The rapid spread of the CRF55_01B subtype across China was mediated by an increase of plasma HIV RNA load and the relatively lower CD4 count (14). Moreover, these factors may prolong the asymptomatic phase and increase the risk of HIV transmission, which may soon lead to further increases and a potential epidemic. Finally, these factors may also help explain the recent surging dominance and continued expansion of this subtype among MSM in China.

This study was subject to several limitations. First, during the onsite questionnaire survey, the information was not filled in detail, such as ID number, so there were deviations in the data analysis. The second is that the two-year sequences were obtained by the same method, but not by the same team, so there will be certain differences in comparison. The experimental methods were the same, but the Stanford drug resistance discriminant version was different. This study retested for resistance against the latest version.

Although the overall prevalence of TDR was low in China, it is necessary to remain vigilant, especially of

TABLE 4. Factors associated with HIV drug resistance among newly diagnosed HIV individuals in 2015 and 2018, China.

Factors	Total	TDR (%)	OR (95% CI)	P	AOR (95% CI)	P
Total	8,980	367 (4.1)				
Age (years)						
16–25	2,230	78 (3.5)	1.00		1.00	
26–49	5,099	209 (4.1)	1.24 (0.95–1.61)	0.12	1.21 (0.93–1.58)	0.16
≥50	1,538	72 (4.7)	1.41 (1.02–1.95)	0.04	1.39 (1.00–1.94)	0.05
Unknown	113	8 (7.1)	2.11 (1.00–4.48)	0.05	2.05 (0.96–4.37)	0.06
Sex						
Male	7,358	299 (4.1)	1.00			
Female	1,557	65 (4.2)	1.03 (0.78–1.35)	0.84		
Unknown	65	3 (4.6)	1.12 (0.35–3.60)	0.84		
Education						
Primary school	2,331	90 (3.9)	1.00			
Junior high school	2,617	120 (4.6)	1.20 (0.91–1.58)	0.21		
Senior high school or above	3,646	147 (4.0)	1.05 (0.80–1.37)	0.74		
Unknown	386	10 (2.6)	0.66 (0.34–1.28)	0.22		
Route of HIV transmission						
HET	4,714	201 (4.3)	1.00			
MSM	3,761	140 (3.7)	0.87 (0.70–1.08)	0.21		
IDU	168	11 (6.5)	1.57 (0.84–2.95)	0.16		
Others	337	15 (4.5)	1.04 (0.61–1.78)	0.89		
Subtype						
CRF01_AE	3,413	135 (4.0)	1.00		1.00	
CRF07_BC	3,371	107 (3.2)	1.26 (0.97–1.63)	0.08	1.25 (0.97–1.62)	0.09
CRF08_BC	802	37 (4.6)	1.48 (1.01–2.16)	0.05	1.51 (1.03–2.23)	0.04
CRF55_01B	278	34 (12.2)	4.25 (2.83–6.39)	<0.001	4.17 (2.77–6.27)	<0.001
B	444	19 (4.3)	1.36 (0.83–2.25)	0.22	1.36 (0.83–2.24)	0.23
Others	672	35 (5.2)	1.67 (1.13–2.47)	0.01	1.69 (1.14–2.49)	0.01
Year						
2015	4,704	178 (3.8)	1.00			
2018	4,276	189 (4.4)	1.18 (0.95–1.45)	0.13		

Abbreviations: HIV=human immunodeficiency virus; TDR=transmitted drug resistance; OR=odds ratio; CI=confidence interval; AOR=adjusted odds ratio; HET=heterosexual; MSM=men who have sex with men; IDU=injecting drug users.

CRF55_01B. Strengthened surveillance of TDR can help reduce the transmission of TDR to the key target populations, especially MSM. In 2018, TDR reached a moderate epidemic level in Xinjiang Uygur Autonomous Region and other regions. Key areas should pay attention and take measures to control the spread of drug resistance. Reducing the prevalence of TDR is important for formulating future treatment and prevention guidelines. Surveillance and prevention of drug resistance should be a critical component of China's programmatic response to HIV to ensure the long-term efficacy and sustainability of ART and

achieve 95-95-95 targets.

Conflicts of interest: No conflicts of interest.

Acknowledgments: Provincial CDC staff.

Funding: The Ministry of Science and Technology of China (2017ZX10201101).

doi: 10.46234/ccdcw2021.251

Corresponding author: Xing Hui, xingh@chinaaids.cn.

¹ State Key Laboratory of Infectious Disease Prevention and Control (SKLID), National Center for AIDS/STD Control and Prevention (NCAIDS), Chinese Center for Disease Control and Prevention (China CDC), Collaborative Innovation Center for Diagnosis and Treatment of Infectious Diseases, Beijing, China.

Submitted: October 22, 2021; Accepted: November 19, 2021

REFERENCES

1. He X, Xing H, Ruan YH, Hong KX, Cheng CL, Hu YY, et al. A comprehensive mapping of HIV-1 genotypes in various risk groups and regions across China based on a nationwide molecular epidemiologic survey. *PLoS One* 2012;7(10):e47289. <http://dx.doi.org/10.1371/journal.pone.0047289>.
2. Han XX, An MH, Zhang WQ, Cai WP, Chen X, Takebe Y, et al. Genome sequences of a novel HIV-1 circulating recombinant form, CRF55_01B, identified in China. *Genome Announc* 2013;1(1):e00050 – 12. <http://dx.doi.org/10.1128/genomeA.00050-12>.
3. Liang BY, Wei QY, Yang Y, Yang Y, Liu J, Chu JM, et al. Identification of a novel HIV-1 CRF55_01B/B recombinant isolate in Guangxi, China. *AIDS Res Hum Retroviruses* 2020;36(5):434 – 9. <http://dx.doi.org/10.1089/aid.2019.0222>.
4. Chinese Center for Disease Control and Prevention. Notice of the General Office of the National Health and Family Planning Commission on adjusting the standard of free antiviral treatment for AIDS. 2016. http://www.chinaaids.cn/zlgh/hdjz6/201606/t20160615_131433.htm. [2016-6-15]. (In Chinese).
5. Benson C, Wang X, Dunn KJ, Li N, Mesana L, Lai J, et al. Antiretroviral adherence, drug resistance, and the impact of social determinants of health in HIV-1 patients in the US. *AIDS Behav* 2020;24(12):3562 – 73. <http://dx.doi.org/10.1007/s10461-020-02937-8>.
6. Kang RH, Liang SJ, Ma YL, Liang S, Xiao L, Zhang XH, et al. Pretreatment HIV drug resistance in adults initiating antiretroviral therapy in China, 2017. *Infect Dis Poverty* 2020;9(1):54. <http://dx.doi.org/10.1186/s40249-020-00668-5>.
7. World Health Organization. HIV drug resistance surveillance guidance, 2015 update. 2016. <https://apps.who.int/iris/handle/10665/204471>. [2020-2-5].
8. Gan MZ, Zheng S, Hao JJ, Ruan YH, Liao LJ, Shao YM, et al. The prevalence of CRF55_01B among HIV-1 strain and its connection with traffic development in China. *Emerg Microbes Infect* 2021;10(1):256 – 65. <http://dx.doi.org/10.1080/22221751.2021.1884004>.
9. Ye JR, Hao MQ, Xing H, Zhang FJ, Wu H, Lv W, et al. Transmitted HIV drug resistance among individuals with newly diagnosed HIV infection: a multicenter observational study. *AIDS* 2020;34(4):609 – 19. <http://dx.doi.org/10.1097/QAD.0000000000002468>.
10. McClung RP, Oster AM, Ocfemia MCB, Saduvala N, Heneine W, Johnson JA, et al. Transmitted drug resistance among HIV-1 diagnoses in the United States, 2014–2018. *Clin Infect Dis*, 2021ciab583. <http://dx.doi.org/10.1093/cid/ciab583>.
11. Liu YJ, Li HP, Wang XL, Han JW, Jia L, Li TY, et al. Natural presence of V179E and rising prevalence of E138G in HIV-1 reverse transcriptase in CRF55_01B viruses. *Infect Genet Evol* 2020;77: 104098. <http://dx.doi.org/10.1016/j.meegid.2019.104098>.
12. Zhang D, Zheng CL, Li HP, Li H, Liu YJ, Wang XL, et al. Molecular surveillance of HIV-1 newly diagnosed infections in Shenzhen, China from 2011 to 2018. *J Infect* 2021;83(1):76 – 83. <http://dx.doi.org/10.1016/j.jinf.2021.04.021>.
13. Dong AB, Liu L, Xiao L, Liang S, Li K, Hu J, et al. First detection of a circulating recombinant form of HIV-1 CRF01_AE/08_BC (CRF105_0108) with drug-resistant mutations in Sichuan, China. *AIDS Res Hum Retroviruses* 2020;36(7):625 – 30. <http://dx.doi.org/10.1089/AID.2020.0034>.
14. Wei L, Li H, Lv X, Zheng CL, Li GL, Yang ZR, et al. Impact of HIV-1 CRF55_01B infection on the evolution of CD4 count and plasma HIV RNA load in men who have sex with men prior to antiretroviral therapy. *Retrovirology* 2021;18(1):22. <http://dx.doi.org/10.1186/s12977-021-00567-z>.