

HIV transmission links between past and newly diagnosed infections: a molecular epidemiology study

Tsz Ho Kwan, Ngai Sze Wong, Grace Chung Yan Lui, Kenny Chi Wai Chan, Owen Tak Yin Tsang, Wai Shing Leung, Kai Man Ho, Man Po Lee, Wilson Lam, Sze Nga Chan, Denise Pui Chung Chan, Sabrina Wai Chi To, Wing Cheong Yam, Shui Shan Lee

Abstract

Background Understanding the driving forces of HIV infection is crucial for developing intervention strategies. We aimed to infer these forces from genetic linkages between past and newly diagnosed cases by a molecular approach.

Methods We recruited newly diagnosed HIV patients from all three HIV specialist clinics in Hong Kong, China, between 2016 and 2018 to collect their HIV sequences with corresponding behavioural and clinical data. We retrieved archived HIV sequences from 1994 to 2012 separately. We pooled and analysed sequences by constructing a network with 1.5% TN93 distance threshold and measured node degree centrality. We did univariate analyses to identify factors associated with genetic links with archived sequences among newly diagnosed patients, and used a generalised linear model to differentiate archived sequences having links with new cases adjusting for diagnosis year.

Findings 2778 partial *pol* sequences were amalgamated for network construction, of which 426 were newly diagnosed. The network contained 1908 (69%) nodes and 23 305 edges. Some 703 (3%) edges linked 192 archived samples with 135 new cases. Of 239 newly diagnosed patients with genetic connections, those linked with past infections were more likely to be subtype B (odds ratio 3.31; $p < 0.0001$) and be 30 years or younger (2.26; $p = 0.0065$). Men who have sex with men (MSM) linked with past infections were less likely to be engaged in chemsex (0.52; $p = 0.039$) or users of a mainland Chinese gay dating mobile application (0.39; $p = 0.035$). Generalised linear model showed that people with past infections linked to new cases were younger ($p = 0.0002$), had a higher degree centrality ($p < 0.0001$), were less likely to be subtype CRF01_AE ($p = 0.0019$), and infected through heterosexual contact ($p < 0.0001$) or injection drug use ($p = 0.0002$) compared with MSM contact.

Interpretation HIV transmission chains linking new with past infections in Hong Kong were driven by younger MSM. Chemsex, emergence in recent years, only clustered in and contributed to some new infections. Formation of transmission linkages over time signified the importance of targeting new and established infections in developing prevention strategies.

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Contributors

THK contributed to data analysis and interpretation, and drafting of the Abstract. NSW and DPCC contributed to data analysis. GCYL, KCWC, OTYT, WSL, KMH, MPL, WL, SNC, SWCT, and WCY contributed to the acquisition of data. SSL conceptualised the study, contributed to the study design and data interpretation, and critically amended the Abstract. All authors have seen and approved the final version of the Abstract for publication.

Declaration of interests

We declare no competing interests.

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Jockey Club School of Public Health and Primary Care (T H Kwan BSc) and Stanley Ho Centre for Emerging Infectious Diseases (N S Wong PhD, D P C Chan PhD, S S Lee FRCP), The Chinese University of Hong Kong, Shatin, Hong Kong Special Administrative Region, China; Department of Medicine and Therapeutics, Prince of Wales Hospital, Shatin, Hong Kong Special Administrative Region, China (G C Y Lui FRCP); Special Preventive Programme, Centre for Health Protection, Department of Health, Kowloon Bay, Hong Kong Special Administrative Region, China (K C W Chan FRCP); Department of Medicine and Geriatrics, Princess Margaret Hospital, Lai Chi Kok, Hong Kong Special Administrative Region, China (O T Y Tsang FRCP, W S Leung FHKAM, K M Ho MNurs); Department of Medicine, Queen Elizabeth Hospital, Yau Ma Tei, Hong Kong Special Administrative Region, China (M P Lee FRCP, W Lam FHKAM, S N Chan FHKCMN); and Department of Microbiology, University of Hong Kong, Pokfulam, Hong Kong Special Administrative Region, China (S W C To PhD, W C Yam PhD)

Correspondence to: Prof Shui Shan Lee, Postgraduate Education Centre, Prince of Wales Hospital, Shatin, Hong Kong Special Administrative Region, China
sslee@cuhk.edu.hk