HIV-1 molecular epidemiology evidence and transmission patterns in the Middle East and North Africa

Ghina Mumtaz,¹ Nahla Hilmi,² Francisca Ayodeji Akala,² Iris Semini,³ Gabriele Riedner,⁴ David Wilson,⁵ Laith J Abu-Raddad^{1,6,7}

¹Infectious Disease Epidemiology Group, Weill Cornell Medical College — Qatar, Cornell University, Qatar Foundation, Education City, Doha, Qatar

²Human Development Sector, Middle East and North Africa Region, World Bank, Washington, DC, USA

³Joint United Nations Programme on HIV/AIDS Regional Support Team, Middle East and North Africa, Cairo, Egypt

⁴Regional Office of the Eastern Mediterranean, World Health Organization, Cairo, Egypt

⁵Global HIV/AIDS Program, World Bank, Washington, DC, USA

⁶Department of Public Health, Weill Cornell Medical College, Cornell University, New York, New York, USA

⁷Vaccine and Infectious Disease Institute, Fred Hutchinson Cancer Research Center, Seattle, Washington, DC, USA

Correspondence to

Dr Laith J Abu-Raddad, Infectious Disease Epidemiology Group, Weill Cornell Medical College — Qatar, Qatar Foundation, Education City, P.O. Box 24144, Doha, Qatar; lja2002@qatar-med.cornell.edu

Accepted 24 August 2010
Published Online First
29 October 2010

This paper is freely available online under the BMJ Journals unlocked scheme, see <http://sti.bmj.com/site/about/unlocked.xhtml>

ABSTRACT

The distribution of HIV-1 subtypes in a population tracks the spread and evolution of the epidemic. This study is a systematic review of all available evidence on HIV-1 molecular epidemiology and subtype distribution in the Middle East and North Africa. Sources of data included Medline and various institutional documents and databases. In several countries, a diverse distribution of HIV-1 subtypes was observed principally reflecting travel-related exogenous exposures. A trend for a dominant HIV-1 subtype was observed in a few other settings and was often linked to HIV transmission within specific high-risk groups such as subtype A and CRF35_AD among injecting drug users and subtype C among commercial sex networks. Multiple exogenous introductions of HIV-1 variants seemed common to all countries, as observed from the high diversity in subtypes, or the high genetic divergence among any specific subtype even if predominant. In several countries though, epidemic-type clustering of specific subtypes suggests established or nascent HIV epidemics among classic core risk groups for HIV infection. HIV prevention efforts in MENA must be prioritized for these high-risk groups.

HIV-1 is characterised by a high genetic variability and includes three groups, with the main group responsible for the HIV pandemic (group M) being further divided into several subtypes (A–K) and a number of mosaic strains known as circulating recombinant forms (CRF).¹ With the exception of sub-Saharan Africa, where the largest diversity of subtypes and CRF has been observed, HIV-1 subtypes show a specific geographical distribution on the global map.^{2,3} Subtype A is the predominant variant in central and eastern Africa and in countries of the former Soviet Union. Subtype B is predominant in western and central Europe, the Americas, Australia, and is commonly found in several countries of Southeast Asia.² Subtype C is the main genetic form in southern Africa and India, the two countries where more than 80% of the global HIV cases occur. Finally, CRF are the most predominant forms in Southeast Asia (CRF01_AE) and in west and central Africa (CRF02_AG).²

Molecular epidemiology investigation can be a powerful tool in developing sensitive diagnostic tools, in the management of individual infections, and most importantly in tracking transmission patterns and the spread and evolution of the epidemic.⁴ Molecular epidemiology research in the Middle East and North Africa (MENA) region is at

an early stage of development and has yet to contribute to informed HIV prevention policy and programming. A recent review on the epidemiology of HIV in MENA indicates that in this region where HIV spread in the general population is very limited, there are emerging epidemics in high-risk populations including injecting drug users (IDU), men who have sex with men (MSM), and to a lesser extent female sex workers (FSW) in a few countries.⁵ Exogenous HIV exposures among nationals linked to travel abroad also appear to be the dominant observed HIV transmission pattern in a few MENA countries.⁵ However, there remain many gaps in the understanding of HIV transmission pathways in MENA.

The objective of this work was to review for the first time all evidence on HIV virus diversity and subtype distribution in MENA countries to complement our knowledge of HIV epidemiology and to understand transmission patterns.

METHODS

This work was part of a comprehensive systematic review of all available data about HIV in MENA, which was complemented by another detailed search focused specifically on published HIV molecular epidemiology literature in MENA. The countries included in our definition of MENA are: Afghanistan, Algeria, Bahrain, Djibouti, Egypt, Iran, Iraq, Jordan, Kuwait, Lebanon, Libya, Morocco, Oman, Pakistan, Palestine, Qatar, Saudi Arabia, Somalia, Sudan, Syria, Tunisia, United Arab Emirates and Yemen. Considering geographical proximity and similarity in the sociocultural context, data were occasionally included on Mauritania, Israel and Turkey.

The main source of data for this investigation was the MENA HIV/AIDS Epidemiology Synthesis Project whose mandate was to collect and synthesise all available data on HIV, sexually transmitted infections (STI) and sexual behaviour in MENA.⁶ Details of this project and of the search strategy used in this comprehensive systematic review were described in a previous publication.⁵ In brief, we undertook a literature search of Medline (PubMed) using a strategy with both free text and MeSH headings and with no time limitation. The main search criteria in relevance to the current work were (('HIV Seropositivity' OR 'HIV' OR 'HIV Infections') AND ('Middle East' OR 'Islam' OR 'Arabs' OR 'Arab World' OR 'Africa, Northern' OR 'Mauritania' OR 'Sudan' OR 'Somalia' OR

'Djibouti' OR 'Pakistan')). We also searched for relevant data in country-level reports and databases, governmental and non-governmental organisations studies and publications, international organisations' reports and databases, as well as other institutional reports related to HIV and STI in MENA.⁵

The above generic search was complemented by a specific search of the Medline database on all HIV-1 molecular data in MENA. The MeSH search criteria used were (('HIV-1/classification' OR 'HIV-1/genetics') AND ('Middle East' OR 'Islam' OR 'Iran' OR 'Arabs' OR 'Arab World' OR 'Africa, Northern' OR 'Mauritania' OR 'Sudan' OR 'Somalia' OR 'Djibouti' OR 'Pakistan')). All relevant studies identified in this search were already covered by the wide-umbrella search of the Synthesis Project above. However, this search was undertaken and respective citations reviewed for further reassurance that no relevant data were missed.

Finally, we searched the Los Alamos HIV sequence database, which gathers global HIV sequence data stratified by geographical location, but without any information on data source and study methodology.⁷ Data from all MENA countries were extracted from this online database and compared with results of published studies.

RESULTS

Status of the evidence

As part of the comprehensive Medline search that was undertaken for the Synthesis Project, a total of 1092 citations on HIV in MENA were retrieved as of 29 May 2010, out of which 49

citations were specific to the molecular epidemiology search. After excluding non-relevant studies, there remained 24 studies that examined the nature of HIV-1 subtypes and suggested transmission pathways in the population (table 1). HIV-1 subtype data were also extracted from the Los Alamos HIV database for all MENA countries and suggested trends similar to those observed in published studies (table 2).

All the reviewed studies used standard and comparable methodologies^{4 31} in their molecular analyses, consisting of the amplification of specific genomic regions (mostly *env*, *gag* and/or *pol*) using PCR, followed by sequencing of the PCR products and phylogenetic tree analysis. It appears unlikely that there is bias in comparing findings between studies and over time due to differences in the methodology of molecular analyses.

Settings with variability of subtypes

A number of countries in MENA display a very diverse distribution of HIV-1 subtypes. This is the case of Lebanon where a complex HIV-1 subtype distribution pattern and high levels of intrasubtype diversity were found representing the travel history of its nationals, particularly to Europe and the Americas where subtype B (44%) is acquired, and to Africa and the United Arab Emirates where most of the non-B subtypes, especially subtype A (40%), are acquired.¹⁸ An analogous trend of high strain diversity including many CRF was observed in Saudi Arabia.^{24–26} However, the identification of a subcluster with a new CRF in this country (CRF43_02G) could indicate a local transmission network.²⁴ HIV-1 subtypes in Yemen also displayed a complex distribution, which appeared to be a mix of

Table 1 Distribution of HIV-1 subtypes in MENA as extracted from various studies

Country	Study	Sample size	Year*	Genes	Subtype (%)
Afghanistan	Sanders-Buell ⁸	4 IDU	2005–6	<i>Pol, Env</i>	CRF35_AD (100)
Algeria	Bouzeghoub ⁹	134 Isolates	NA†	<i>Rt, Prot, Env</i>	B (56), CRF02_AG (12.7), CRF06_cpx (4), Inter-CRF recombinants (9.7)
Djibouti	Maslin ¹⁰	34 Isolates	2002	<i>Pol, Env</i>	C (73), CRF02_AG (18), D (6), A (3)
	Lasky ¹¹	33 French military personnel in Djibouti	NA†	<i>Env</i>	C (48), B (33), A (15), E (3)
Iran	Sarrami-Farooshani ¹²	46 IDU and 15 haemophiliacs in Teheran	2003	<i>Gag, Env</i>	IDU: A (100) Haemophiliacs: B (100)
	Naderi ¹³	12 IDU in Mashhad	2005	<i>Gag, Env</i>	A (100)
	Soheili ¹⁴	13 Isolates, mostly IDU	NA†	<i>RT, Prot, Env</i>	CRF35_AD (100)
	Mousavi ¹⁵	39 Isolates, mostly IDU	2005–6	<i>Pol</i>	CRF35_AD (100)
	Hamkar ¹⁶	42 Isolates, 36% IDU	2009	<i>Pol</i>	A/D recombinants (48), B (43), A (5), CRF01_AE (5)
Israel/Palestine	Gehring ¹⁷	92 Isolates, 45% MSM	NA†	<i>Env</i>	B (98), A (1), D (1)
Lebanon	Pieniazek ¹⁸	25 Isolates	1996	<i>Prot, Gag, Env</i>	A (44), B (40), C (4), D (4), G (4),
Libya	de Oliveira ¹⁹	44 Children	1998	<i>Gag</i>	CRF02_AG (100)
Morocco	Elharti ²⁰	200 Isolates	NA†	NA†	B (93.5), A (1%), F (0.5%)
	Abid ²¹	14 Isolates	NA†	<i>Env, Tat</i>	B (78.6%), A (21.4%)
	Bakhouch ²²	98 Isolates	2004–7	<i>Rt, Prot</i>	B (74.6%), CRF02_AG (15.5), CRF01_AE (4.2), G (2.8), C (1.4), F2 (1.4%)
Pakistan	Khan ²³	34 IDU in Karachi	2006	<i>Gag, Env</i>	A (100)
Saudi Arabia	Yamaguchi ²⁴ Badreddine ²⁵	56 Saudi nationals in Riyadh	2003–4	<i>Gag, Pol, Env</i>	C (39.3), B (17.8), Intersubtype recombinants (16.1), CRF43_02G (10.7), CRF25_cpx (7.1), D (3.6), A (1.8), CRF02_AG (1.8)
	Alzahrani ²⁶	12 Isolates in Dammam	2004–7	<i>Gag, Env</i>	C (58%), B (17%), A (8%), D (8%), G (8%)
Sudan	Hierholzer ²⁷	30 Blood donors	1998–9	<i>Pol, Env</i>	D (50), C (30), recombinants (10), A (6.7), B (3.3)
Tunisia	Ben Halima ²⁸	25 Isolates	1999	<i>Env</i>	B (95), CRF02_AG (5)
Turkey	Yilmaz ²⁹	27 Isolates	NA†	<i>Env</i>	B (70.4), A (14.8), F1 (7.4), C (3.7), D (3.7)
Yemen	Saad ³⁰	19 Isolates	2000–2	<i>Pol</i>	B (47.3), C (31.6), D (10.5), A (5.3), URF (5.3)

*Year of data collection.

†Not available in original document.

CRF, circulating recombinant form; IDU, injecting drug user; MENA, Middle East and North Africa; MSM, men who have sex with men.

Table 2 Distribution of HIV-1 subtypes in MENA as extracted from the Los Alamos HIV database⁷

Country	No of isolates	Subtype (%) [*]
Afghanistan	4	35_AD (100)
Algeria	478	B (56.3), CRF02_AG (18.0), CRF06_cpx (17.2), G (2.7), 206 (2.7), D (1.3), other (1.9)
Djibouti	92	C (64.1), CRF02_AG (26.1), D (5.4), A (3.3), CRF01_AE (1.1)
Egypt	24	B (91.7), O (4.2), CRF01_AE (4.2)
Iran	277	A (65), B (32.9), C (2.2)
Israel/Palestine	301	C (85.4), B (8.6), CRF02_AG (1.7), A (1.3), other (3.0)
Kuwait	2	C (100)
Lebanon	59	A (40.7), B (35.6), D (6.8), U (5.1), C (3.4), G (3.4), CRF02_AG (1.7), AD (1.7), AG (1.7)
Libya	68	CRF02_AG (100)
Morocco	2	B (100)
Saudi Arabia	169	C (36.7), G (24.3), B (18.3), D (4.7), CRF43_02G (4.7), A (4.1), CRF02_AG (3.6), CRF25_cpx (2.4), other (1.2)
Somalia	24	C (87.5), AC (12.5)
Sudan	40	D (55), C (30), A (7.5), B (5), U (2.5)
Tunisia	24	B (95.8), CRF02_AG (4.2)
Yemen	25	B (48), C (28), D (16), A (4), U (4)

^{*}Frequency in the HIV database and not necessarily in the respective population. MENA, Middle East and North Africa.

genetic variants closely related to those found over a wide geographical range, including Europe, North America, Asia and Africa.³⁰

Settings with dominance of specific subtypes

In a few countries, a trend of a dominant HIV-1 subtype was observed and linked to HIV transmission chains within specific high-risk groups such as among IDU in Afghanistan, Iran and Pakistan (table 1). In Iran, all viral sequences identified among IDU in two different cities formed a single cluster within subtype A and were found to be related, although not necessarily directly, to Ugandan/Kenyan isolates.^{12 13 32} Interestingly, there was a lack of strong phylogenetic correlation between the sequences from Iran and those from other Middle Eastern countries, apart from Afghanistan, where subtype A is common.^{13 32} The confined variability in the genes suggests the relatively recent nature of the epidemic in Iran.^{12 13 32} Similarly, subtype A was the only clade observed among a group of IDU in Karachi, Pakistan, where the isolated viruses formed a tight monophyletic group indicating a founder effect.^{23 33} In more recent studies among IDU in Iran,^{14–16} CRF35_AD was identified; this subtype had also recently been identified among a small group of IDU in Kabul, Afghanistan.⁸

Clustering within other subtypes was also observed in some MENA countries. In Djibouti, 73% of infections in 2002 were of subtype C, which had previously been described among FSW in the horn of Africa.¹⁰ Subtype C was found in 48% of French military personnel infected with HIV who served in Djibouti.¹¹ Similarly, subtype C accounted for 87.5% of HIV-1 cases in Somalia (table 2). Sudan was the only country in MENA where subtype D was observed at a high frequency of 50%.²⁷

Subtype B was found to be the prevalent subtype in a number of MENA countries. In Algeria subtype B was found to be predominant, particularly in the north of the country, accounting for 56% of samples studied.⁹ There is, however, a high diversity of recombinant subtypes in the southern part of

Algeria, bordering with sub-Saharan Africa.⁹ Subtype B also accounted for 98% of the cases among Caucasian Israeli and Palestinian isolates.¹⁷ In Tunisia, 95% of HIV-1 infections were found to be of subtype B and had probable origins in western Europe.²⁸ Subtype B was historically also found to be the dominant strain in Morocco, accounting for 93.5% of infections in the 1990s and reflecting transmission pathways of western European origins.²⁰ However, more recent data indicate that although subtype B remains the predominant strain, there is an increasing diversity of HIV-1 variants, with over 25% of the infections being of non-B subtypes, including several CRF.²² In one study from Morocco, 21.4% of the infections were of subtype A and clustered with strains of Kenyan origin.²¹

DISCUSSION

General overview

It is not straightforward to draw firm conclusions from available studies, especially as many are based on an analysis of a small number of isolates, which in some settings may be reflecting transmission patterns from the past rather than current ongoing transmission (table 1). However, the gathered molecular data on HIV-1 subtype distribution in the region, substantiated by other epidemiological evidence on disease distribution, HIV dynamics among high-risk populations and risk networks in respective countries (G Mumtaz, 2010, unpublished data),^{5 6} point towards two major patterns of HIV acquisition and transmission in MENA countries: one of random exogenous exposures related to travel history, which is present all over MENA (figure 1A and 1B), and another of local epidemic-type clustering related to concentrated HIV epidemics in specific high-risk groups in a number of MENA countries (figure 1B).

Patterns of exogenous HIV exposures

In a number of MENA countries, as was observed in Lebanon, Saudi Arabia and Yemen, sequence analyses indicated multiple distinct subtypes in the isolated strains, suggesting multiple travel-related introductions rather than the endemic transmission of genetically diverse HIV-1 subtypes (figure 1A). In some other settings such as in northern Algeria, Palestine and Tunisia, although one subtype is predominant, the intrasubtype diversity suggests that HIV transmission appears to be strongly affected by multiple introductions, but from confined geographical sites. In both kinds of settings, HIV dynamics, based strictly on molecular evidence, do not appear to be dominated by a specific considerable epidemic in one or multiple high-risk groups (figure 1A).

The high diversity in subtypes and/or the high genetic divergence among any specific subtype suggest that HIV infections in these countries are driven by travel-related exogenous exposures among nationals or HIV acquisition by their sexual partners upon their return.⁵ Data from case notification reports confirm this postulated trend.³⁴ For example in Lebanon, 45.4% of notified cases up to 2004 were linked to travel abroad,³⁵ as were half of reported AIDS cases in Yemen.³⁴ In these settings, local spread is usually limited to immediate sexual partners and does not lead to an epidemic.³⁶ In one study in Saudi Arabia, a strong phylogenetic relationship between HIV-1 strains was observed among several husband–wife pairs, indicating heterosexual transmission between spouses.²⁵ Indeed, 97% of HIV-infected women in Saudi Arabia who acquired HIV sexually acquired the infection from their husbands.³⁷ This trend of travel-associated infections has also been observed in other countries outside of MENA, such as in Cuba where the epidemic is characterised by a high diversity of subtypes mainly due to the

Figure 1 Schematic representation of countries with dominance of travel-related exogenous HIV exposures (A), and dominance of endogenous epidemic-type HIV transmissions in addition to the contribution of travel-related exogenous HIV exposures (B). The rate and number of exogenous exposures varies between Middle East and North Africa countries.

return of Cuban military troops from Angola, the large numbers of Cuban aid workers serving in sub-Saharan countries, and the recent development of tourist resorts in the country.^{36 38}

The high population mobility in MENA including migration, tourism, studying and working abroad, and military deployment among others explain in large part the observed pattern of exogenous exposures and subsequent subtype diversity,⁶ as well as explaining the observed Africanisation of subtypes in some settings.^{6 9 22 25} Southern Algeria and Morocco are both important migration hubs for migrants from sub-Saharan African countries towards Europe and this is reflected in the high diversity of non-B subtypes and CRF in these regions.^{9 22} In Turkey, non-B subtype infections are also thought to be mainly caused by immigrants or residents from Africa, the Balkans and the Middle East.²⁹ The high prevalence of subtype C in Saudi Arabia is expected in view of the country's ties with neighbouring Horn of Africa countries.²⁵ The very high diversity of subtypes observed in Lebanon reflects the frequent population movements and return of expatriates such as from West Africa.¹⁸ The high level of emigration to western Europe explains the predominance of subtype B in Tunisia, which is thought to have been initially introduced mainly by young male IDU or individuals with haemophilia infected in Europe.²⁸

Patterns of local epidemic-type clustering

On the other hand, the dominance of specific subtypes with limited sequence variability in Afghanistan, Djibouti, Iran, Libya, Pakistan and Somalia in addition possibly to one subcluster in Saudi Arabia, suggests that HIV acquisitions in some countries are occurring in transmission chains that have been propagating locally for at least a number of years. This suggests active high-risk networks where HIV is being transmitted via typical founder effect, which translates into local epidemic-type clustering in one or multiple risk groups (figure 1B).^{8 10 12 13 23 32 33} This trend was documented in Russia, for example, where injecting drug use represented the main mode of HIV transmission for the longest time and where subtype A was the predominant clade driving the epidemic among IDU.³⁹

This conjecture is supported in Iran and Pakistan as examples, by robust evidence for concentrated epidemics among IDU.⁵ By the end of 2007, IDU accounted for at least 67% of HIV notified cases in Iran.⁴⁰ In Pakistan, the contribution of IDU to the total notified cases has been rising, and IDU accounted for the largest proportion of HIV infections in 2008.⁴¹ The similarities between

HIV-1 subtypes in Afghanistan, Iran and apparently Pakistan suggest that strains have been circulating between these countries, probably through the return of Afghani refugees, and may indicate that social networks of drug producers, refiners, distributors and IDU transcend national borders.⁸

The local epidemic-type clustering is also supported by epidemiological evidence from Djibouti and from some parts of Somalia, where there is evidence of concentrated HIV epidemics in commercial heterosexual sex networks,⁵ with rates of HIV up to 70% being reported among street-based FSW in Djibouti.^{42 43} Clustering within subtype C in HIV-1 infections in Djibouti and Somalia is in accordance with previous studies in the Horn of Africa, which have identified subtype C to be the dominant clade, accounting for up to 98.5% of infections among FSW in Ethiopia.^{44 45} Similarly, in Sudan the epidemic appears to be dominated by East African viruses also circulating in neighbouring countries.²⁷ Libya is yet another example where clustering of subtypes was observed, but in the context of a long-standing nosocomial transmission scenario.¹⁹

Transmission links between epidemics

One of the insightful applications of HIV molecular research is that it can indicate whether there is a link between two epidemics occurring simultaneously in two different risk groups in one country. This was reported in many regions of the world such as in Thailand, where in the first decade of the epidemic, HIV-1 subtypes were segregated by risk group, with IDU being infected with subtype B and patients infected through the heterosexual route being infected with CRF01_AE.^{46 47}

Similarly in Iran, molecular epidemiology investigations indicated that the HIV epidemic among IDU is independent of the epidemic among individuals with haemophilia as seen by the different subtype clustering in each transmission network: IDU were found to be exclusively infected with subtype A, which has been possibly introduced to Iranian IDU by a pilgrim visiting Mashhad, Iran's holiest city,³² whereas individuals with haemophilia were exclusively infected by subtype B.¹² In Israel/Palestine, whereas subtype B was the predominant clade, subtype C was exclusively observed among the Ethiopian Israeli population, indicating two distinct HIV transmission pathways in the population that possibly reflect the geographical site of virus acquisition.¹⁷

Recent phylogenetic analyses have also linked epidemics in MENA such as the emerging epidemic among MSM in Pakistan

Key messages

- ▶ HIV molecular epidemiology investigation is a powerful tool for tracking transmission patterns and the spread and evolution of the epidemic in a population.
- ▶ This is the first review of all available data on HIV-1 subtype distribution in MENA and implications in terms of transmission patterns.
- ▶ Molecular evidence shows two main patterns: the dominance of exogenous exposures in several countries and the dominance of local epidemic-type clustering among high-risk groups in others.
- ▶ The virus subtype distribution in MENA reflects the transmission pathways of virus introduction into this region from the other parts of the world.

and that which emerged among IDU earlier in this decade in this country (S. Ali, personal communication, 2010).

Recommendations for future research and interventions

With the weak HIV surveillance systems in this region, it is useful to expand HIV molecular research, especially as some of the available studies have been conducted a decade ago. Overall in MENA, only a very small number of HIV cases have ever been subtyped. Despite the high cost of molecular analyses, identifying the subtypes and studying the phylogenetic clustering of at least small subsamples of notified HIV cases and HIV-seropositive cases in surveyed populations is pertinent. It can pinpoint the most affected risk groups and the prevalent modes of transmission in the population, and help establish some reliable information on the subtype distributions. However, the priority in MENA countries that are still developing their HIV surveillance infrastructure would be to invest in integrated biobehavioural surveillance using state of the art sampling methodologies for hard-to-reach populations. This kind of surveillance is essential in order to facilitate the prioritisation of HIV response to where needs are and to avert the possibility of HIV efforts being dissipated on non-consequential programmes.

The molecular evidence gathered in this review, supported by other epidemiological evidence (G Mumtaz, 2010, unpublished data),^{5,6} indicates the absence of a sustainable general population HIV epidemic in MENA, with the exception possibly of southern Sudan. As such, any current HIV prevention efforts focused on the general population are misplaced. Whereas it is challenging to identify, prevent or even control travel-related exogenous exposures, the number of infections arising from these random introductions remains relatively small. Instead, surveillance and prevention efforts should be prioritised for high-risk groups including IDU, MSM and FSW, in whom low intensity and concentrated epidemics are emerging or are already established. This would prevent further HIV transmission or the emergence of localised and hidden HIV epidemics among these population groups.

Funding The MENA HIV/AIDS Epidemiology Synthesis Project was funded through a joint partnership of the World Bank, the MENA Regional Support Team of the Joint United Nations Programme on HIV/AIDS (UNAIDS) and the Eastern Mediterranean Regional Office (EMRO) of the WHO. LJA and GM are also grateful for the Qatar National Research Fund for supporting this work (NPRP 08-068-3-024).

Competing interests None.

Contributors LJA conceived and led the design and conduct of this study; co-led the literature reviews and collection, analysis and interpretation of the data; and contributed to the drafting of the manuscript. GM led the drafting of the manuscript

and co-led the literature reviews and collection, analysis and interpretation of the data. NH contributed to the literature reviews and assisted in the general conduct of the research. FAA, IS, GR and DW contributed to the interpretation of the results, contributed to the drafting of the manuscript, and provided data at the regional and country level. All authors were involved in the finalisation of the manuscript and approved the final version.

Provenance and peer review Not commissioned; externally peer reviewed.

REFERENCES

1. Woodman Z, Williamson C. HIV molecular epidemiology: transmission and adaptation to human populations. *Curr Opin HIV AIDS* 2009;**4**:247–52.
2. Buonaguro L, Tornesello ML, Buonaguro FM. Human immunodeficiency virus type 1 subtype distribution in the worldwide epidemic: pathogenetic and therapeutic implications. *J Virol* 2007;**81**:10209–19.
3. Papathanasopoulos MA, Hunt GM, Tiemessen CT. Evolution and diversity of HIV-1 in Africa — a review. *Virus Genes* 2003;**26**:151–63.
4. Tatt ID, Barlow KL, Nicoll A, et al. The public health significance of HIV-1 subtypes. *AIDS* 2001;**15**(Suppl 5):S59–71.
5. Abu-Raddad LJ, Hilmi N, Mumtaz G, et al. Epidemiology of HIV infection in the Middle East and North Africa. *AIDS* 2010;**24**(Suppl 2):S5–23.
6. Abu-Raddad L, Akala FA, Semini I, et al. *Characterizing the HIV/AIDS epidemic in the Middle East and North Africa: time for strategic action. Middle East and North Africa HIV/AIDS Epidemiology Synthesis Project*. Washington DC, USA: World Bank/UNAIDS/WHO Publication, 2010.
7. Los Alamos HIV database, National Institute of Allergy and Infectious Diseases (NIAID). <http://www.hiv.lanl.gov/content/index> (accessed Feb 2010).
8. Sanders-Buell E, Saad MD, Abed AM, et al. A nascent HIV type 1 epidemic among injecting drug users in Kabul, Afghanistan is dominated by complex AD recombinant strain, CRF35_AD. *AIDS Res Hum Retroviruses* 2007;**23**:834–9.
9. Bouzeghoub S, Jauvin V, Recordon-Pinson P, et al. High diversity of HIV type 1 in Algeria. *AIDS Res Hum Retroviruses* 2006;**22**:367–72.
10. Maslin J, Rogier C, Berger F, et al. Epidemiology and genetic characterization of HIV-1 isolates in the general population of Djibouti (Horn of Africa). *J Acquir Immune Defic Syndr* 2005;**39**:129–32.
11. Lasky M, Perret JL, Peeters M, et al. Presence of multiple non-B subtypes and divergent subtype B strains of HIV-1 in individuals infected after overseas deployment. *AIDS* 1997;**11**:43–51.
12. Sarrami-Forooshani R, Das SR, Sabahi F, et al. Molecular analysis and phylogenetic characterization of HIV in Iran. *J Med Virol* 2006;**78**:853–63.
13. Naderi HR, Tagliamonte M, Tornesello ML, et al. Molecular and phylogenetic analysis of HIV-1 variants circulating among injecting drug users in Mashhad, Iran. *Infect Agent Cancer* 2006;**1**:4.
14. Soheili ZS, Ataiee Z, Tootian S, et al. Presence of HIV-1 CRF35_AD in Iran. *AIDS Res Hum Retroviruses* 2009;**25**:123–4.
15. Mousavi SM, Hamkar R, Gouya MM, et al. Surveillance of HIV drug resistance transmission in Iran: experience gained from a pilot study. *Arch Virol* 2010;**155**:329–34.
16. Hamkar R, Mohraz M, Lorestani S, et al. Assessing subtype and drug-resistance associated mutations among antiretroviral-treated HIV-infected patients. *AIDS* 2010;**24**(Suppl 2):S85–91.
17. Gehring S, Maayan S, Ruppach H, et al. Molecular epidemiology of HIV in Israel. *J Acquir Immune Defic Syndr Hum Retroviral* 1997;**15**:296–303.
18. Pieniazek D, Baggs J, Hu DJ, et al. Introduction of HIV-2 and multiple HIV-1 subtypes to Lebanon. *Emerg Infect Dis* 1998;**4**:649–56.
19. de Oliveira T, Pybus OG, Rambaut A, et al. Molecular epidemiology: HIV-1 and HCV sequences from Libyan outbreak. *Nature* 2006;**444**:836–7.
20. Elharti E, Elouad R, Amzazi S, et al. HIV-1 diversity in Morocco. *AIDS* 1997;**11**:1781–3.
21. Abid M, Luo CC, Sekkat S, et al. Characterization of the V3 region of HIV type 1 isolates from Morocco. *AIDS Res Hum Retroviruses* 1998;**14**:1387–9.
22. Bakhouch K, Oulad-Lahcen A, Bensghir R, et al. The prevalence of resistance-associated mutations to protease and reverse transcriptase inhibitors in treatment-naïve (HIV1)-infected individuals in Casablanca, Morocco. *J Infect Dev Ctries* 2009;**3**:380–91.
23. Khan S, Rai MA, Khanani MR, et al. HIV-1 subtype A infection in a community of intravenous drug users in Pakistan. *BMC Infect Dis* 2006;**6**:164.
24. Yamaguchi J, Badreddine S, Swanson P, et al. Identification of new CRF43_02G and CRF25_cpx in Saudi Arabia based on full genome sequence analysis of six HIV type 1 isolates. *AIDS Res Hum Retroviruses* 2008;**24**:1327–35.
25. Badreddine S, Smith K, van Zyl H, et al. Identification and characterization of HIV type 1 subtypes present in the Kingdom of Saudi Arabia: high level of genetic diversity found. *AIDS Res Hum Retroviruses* 2007;**23**:667–74.
26. Alzahrani AJ. Analysis of HIV subtypes and the phylogenetic tree in HIV-positive samples from Saudi Arabia. *Saudi Med J* 2008;**29**:1394–6.
27. Hierholzer M, Graham RR, El Khidir I, et al. HIV type 1 strains from East and West Africa are intermixed in Sudan. *AIDS Res Hum Retroviruses* 2002;**18**:1163–6.
28. Ben Halima M, Pasquier C, Slim A, et al. First molecular characterization of HIV-1 Tunisian strains. *J Acquir Immune Defic Syndr* 2001;**28**:94–6.
29. Yilmaz G, Midilli K, Turkoglu S, et al. Genetic subtypes of human immunodeficiency virus type 1 (HIV-1) in Istanbul, Turkey. *Int J Infect Dis* 2006;**10**:286–90.
30. Saad MD, Al-Jaify A, Graham RR, et al. HIV type 1 strains common in Europe, Africa, and Asia cocirculate in Yemen. *AIDS Res Hum Retroviruses* 2005;**21**:644–8.

31. **Thomson MM**, Najera R. Molecular epidemiology of HIV-1 variants in the global AIDS pandemic: an update. *AIDS Rev* 2005;**7**:210–24.
32. **Tagliamonte M**, Naderi HR, Tornesello ML, *et al.* HIV type 1 subtype A epidemic in injecting drug user (IDU) communities in Iran. *AIDS Res Hum Retroviruses* 2007;**23**:1569–74.
33. **Rai MA**, Nerurkar VR, Khoja S, *et al.* Evidence for a “Founder Effect” among HIV-infected injection drug users (IDUs) in Pakistan. *BMC Infect Dis* 2010;**10**:7.
34. **WHO, UNICEF, UNAIDS, Yemen.** *Epidemiological facts sheets on HIV/AIDS and sexually transmitted infections*. Geneva, Switzerland: UNAIDS, 2006.
35. **Karam W**, Aftimos G, Jurjus A, *et al.* Prevalence of sexually transmitted infections in Lebanese women as revealed by pap smear cytology: a cross sectional study from 2002–2006. Cairo, Egypt. *WHO/EMRO* 2007.
36. **Perrin L**, Kaiser L, Yerly S. Travel and the spread of HIV-1 genetic variants. *Lancet Infect Dis* 2003;**3**:22–7.
37. **Alrajhi AA**, Halim MA, Al-Abdely HM. Mode of transmission of HIV-1 in Saudi Arabia. *AIDS* 2004;**18**:1478–80.
38. **Cuevas MT**, Ruibal I, Villahermosa ML, *et al.* High HIV-1 genetic diversity in Cuba. *AIDS* 2002;**16**:1643–53.
39. **Thomson MM**, Vinogradova A, Delgado E, *et al.* Molecular epidemiology of HIV-1 in St Petersburg, Russia: predominance of subtype A, former Soviet Union variant, and identification of intrasubtype subclusters. *J Acquir Immune Defic Syndr* 2009;**51**:332–9.
40. **Iran Ministry of Health.** *AIDS/HIV surveillance report, fourth quarter*. Teheran: Ministry of Health, 2007.
41. **Pakistan Ministry of Health.** *AIDS/HIV surveillance report, third quarter*. Islamabad, Pakistan, 2008.
42. **Marcelin AG**, Grandadam M, Flandre P, *et al.* Comparative study of heterosexual transmission of HIV-1, HSV-2 and KSHV in Djibouti. 8th Retrovir Oppor Infect (abstract no. 585). Foundation for Retrovirology and Human Health. Chicago, Illinois, USA. 4–8 February 2001.
43. **Marcelin AG**, Grandadam M, Flandre P, *et al.* Kaposi's sarcoma herpesvirus and HIV-1 seroprevalences in prostitutes in Djibouti. *J Med Virol* 2002;**68**:164–7.
44. **Hussein M**, Abebe A, Pollakis G, *et al.* HIV-1 subtype C in commercial sex workers in Addis Ababa, Ethiopia. *J Acquir Immune Defic Syndr* 2000;**23**:120–7.
45. **Zewde A**, Bahiru S, Sanders E, *et al.* HIV-1 seroprevalence and subtypes in police recruits from Afar regional state, Ethiopia. *Ethiop Med J* 2002;**40**(Suppl 1):1–10.
46. **Ou CY**, Takebe Y, Weniger BG, *et al.* Independent introduction of two major HIV-1 genotypes into distinct high-risk populations in Thailand. *Lancet* 1993;**341**:1171–4.
47. **Arroyo MA**, Phanuphak N, Krasaesub S, *et al.* HIV type 1 molecular epidemiology among high-risk clients attending the Thai Red Cross Anonymous Clinic in Bangkok, Thailand. *AIDS Res Hum Retroviruses* 2010;**26**:5–12.