HIV/AIDS surges in Eastern Europe — Asia-Pacific next?

Twenty years after physicians first spotted the immune deficiency disease that became known as AIDS, the state of the worldwide AIDS epidemic is stirring both dismay and hope, judging from the latest AIDS Epidemic Update released on 28 November by the Joint United Nations Programme on HIV/AIDS (UNAIDS) and WHO.

In Eastern Europe and Central Asia, HIV infections are rising faster than anywhere in the world — an estimated one million people were living with HIV/AIDS there in 2001, up from 700 000 the year before — and “a huge epidemic may be imminent” in parts of the former Soviet Union, the UNAIDS/WHO report says. In Asia and the Pacific, low national incidence rates hide “serious” local epidemics that could break into the general population, particularly in China, where as many as one million people may already be infected. In the US, Western Europe, Canada and Australia, complacency is replacing the safer-sex ethic “promoted so successfully for much of the 1980s and 1990s”, posing the threat of resurgent epidemics.

Worldwide, an estimated five million people were newly infected with HIV in 2001 (vs 5.3 million in 2000) and an estimated 40 million people (vs 36.1 million) are believed to be living with the virus — 70% of them in sub-Saharan Africa. Urban men and women in Zambia report fewer sexual contacts and more condom use. HIV prevalence rates among South African adolescents have dropped sharply since 1998. And in Uganda, “the first African country to have subdued a major HIV/AIDS epidemic”, prevalence rates among pregnant women in urban areas have fallen for the past eight years running, from a peak of 29.5% in 1992 to 11.25% in 2000.

“Where there’s the will and commitment from all different sectors — from the political to the religious sectors — it’s possible to reverse the epidemic,” says Dr Jesus Maria Garcia Calleja, a UNAIDS epidemiologist in Geneva, Switzerland. “It’s not a question only of money. It’s a question of the country getting involved in the fight against HIV/AIDS.”

Massive prevention efforts could avert “a much larger and more generalized epidemic” in Eastern Europe and Central Asia, the UNAIDS/WHO report says, because the outbreak there is still at an early stage. Although diagnosed AIDS cases have risen dramatically over the past two years in the Russian Federation, the epidemic is still centred among IV drug users, not just in Russia but also in Ukraine and several Central Asian countries. However, a high proportion of IV drug users are sexually active young men, raising the prospect that HIV infection might explode into the broader population.

“What is needed is “a comprehensive response to reduce risky sexual and drug-injecting behaviour among young people, and to tackle the socioeconomic and other factors that promote the spread of the virus,” the report says. It points to signs that governments in the region are planning such an effort.

“The coming year can be the turning point in the fight against this global epidemic,” says WHO’s Brundtland. But she warns that “it will be a long fight.” Dr Peter Piot, UNAIDS executive director, cautions: “It will get worse before it gets better.”

Bruce Agnew, Bethesda, Maryland, USA

Haemoglobin variant gives strong protection against malaria

A genetic variation of haemoglobin, the oxygen-carrying protein found in red blood cells, can almost completely abolish the risk of falciparum malaria, the most lethal form of the disease, according to a study in West Africa reported by African and Italian researchers in the 15 November issue of Nature. Having two copies of the gene for the variant, called haemoglobin C, provides “almost complete protection” against the disease, according to malaria researcher and study team leader Dr Mario Coluzzi of the University of Rome La Sapienza.

Moreover, haemoglobin C comes with few unhealthy strings attached. Mild anaemia and, in some adults, gallstones are the only adverse effects. “Compared to all other red blood cell mutations that protect against malaria, haemoglobin C is not associated with serious health problems,” says Dr David Modiano, lead author of the study.

Scientists have known for years that another version of the haemoglobin gene, haemoglobin S, shields against malaria by crippling the red blood cells that malaria parasites need to survive. The abnormal haemoglobin distorts red blood cells into a sickle shape, and these misshapen cells along with any parasites they contain are destroyed by the spleen. But with haemoglobin S, malaria protection comes...
at a cost. People with two copies of the haemoglobin S gene often develop potentially lethal sickle-cell anaemia.

Earlier work by other research teams, including an epidemiological study in Mali and laboratory experiments on parasite proliferation, had suggested a link between haemoglobin C and malaria resistance, but the results were inconclusive. The Rome team confirmed the protective effect of haemoglobin C by studying the blood of 4348 children of the Mossi ethnic group in Ouagadougou, Burkina Faso. The majority of the participants were healthy, but 835 of them had malaria caused by the parasite Plasmodium falciparum. The scientists determined which forms of the haemoglobin gene each child had inherited.

When the researchers compared the genetic data between the healthy and sick children, they found far fewer haemoglobin C genes among the sick patients. Statistical analysis predicted that 14 of the sick children would have two copies of the haemoglobin C gene if the variant offered no protection against the disease. Instead, only one sick child had a double dose of the gene. Based on their findings, the scientists calculate that having a haemoglobin C gene paired with a normal haemoglobin gene reduces the risk of malaria by 29% compared to having two normal haemoglobin genes. Having two copies of the haemoglobin C gene reduces the risk by a huge 93%. That is slightly more protective than the 73% risk reduction experienced by people having a normal haemoglobin gene paired with a haemoglobin S gene (HbAS genotype), and certainly more protective than the approximately 67% protection afforded by having two copies of the haemoglobin S gene (HbSS), although the lethality of that genotype clearly offsets its protective effect.

Modiano and his colleagues are currently conducting a study of more than 5000 people, children and adults, in Burkina Faso to find out whether haemoglobin C merely keeps the disease in check or if it totally blocks infection. How the haemoglobin could accomplish either task remains a mystery. The abnormal protein might produce slight changes in the way antigens are exposed on the surface of red blood cells, says Modiano. Determining how haemoglobin C fends off malaria might guide the development of vaccines and better treatments that safely mimic the mechanism of action involved. ■

Charlene Crabb, Paris, France

Anti-inflammatory drugs slash Alzheimer risk

Extended use of nonsteroidal anti-inflammatory drugs (NSAIDs), such as ibuprofen, may reduce the risk of developing Alzheimer disease by as much as 80%, according to a seven-year prospective study published on 22 November in the New England Journal of Medicine (NEJM). If these results are confirmed in prevention trials, the drugs could have an enormous public health impact, say experts.

“Given the global spread of this disease, with about 11 million sufferers of Alzheimer dementia worldwide,” says WHO mental health expert Dr José Bertolote, “any evidence such as this clearly has the potential to be of public health significance. We have to be hopeful, however, that these early findings will live up to the expectations they raise.”

The new study, conducted by researchers at Erasmus University Medical School in Rotterdam, the Netherlands, followed nearly 7000 people over an average of just under seven years. All subjects were aged 55 or older and free of dementia at the beginning of the study. The researchers evaluated the mental and neurological health of the participants at the beginning, middle and end of the study. Throughout the study, the researchers used computerized pharmacy records to keep track of the subjects’ NSAID use. (NSAIDs were available only by prescription in the Netherlands during most of the study.)

Analysis revealed that study participants who used NSAIDs for two or more years were 80% less likely to develop Alzheimer disease compared to people who didn’t use the drugs. Ibuprofen, naproxen, and diclofenac were the most popular NSAIDs, together accounting for about 83% of the total number of NSAIDs taken by participants in the study.

This isn’t the first study to find a link between NSAIDs and Alzheimer risk. “There have been at least 20 epidemiological studies looking at NSAIDs and Alzheimer disease,” Dr Peter P. Zandi of Johns Hopkins University in Baltimore, Maryland, USA, told the Bulletin. But this latest study is especially strong because it was prospective, involved a large population group, included a long follow-up period, and gathered data from pharmacy records rather than relying on the subjects’ memory of past NSAID use. Zandi and his colleague at Johns Hopkins, Dr John Breitner, wrote in an accompanying editorial in the NEJM.

It is unclear how NSAIDs might decrease Alzheimer risk. Inflammation is believed to be involved in Alzheimer. For this reason, “we think that inhibition of inflammation [by these drugs] may play a role,” Dr Bruno Stricker, senior author of the study, told the Bulletin. But that is only one hypothesis. A study recently published in the 8 November issue of Nature suggested that NSAIDs might lessen Alzheimer risk by reducing brain levels of amyloid-β, a protein that accumulates in the brains of people with the disease. The exact mechanism by which these drugs might achieve this result is not entirely clear, but blockage of an enzyme involved in the production of amyloid-β is one possibility.

For now, the ability of NSAIDs to prevent Alzheimer disease awaits verification from clinical trials under way. “Widespread use [of NSAIDs for this purpose] should be discouraged until prospective double-blind randomized trials have confirmed this and have assessed the ideal dose and benefit/risk ratio,” Stricker told the Bulletin.

But if the benefits of NSAIDs are validated in further trials, the public health gains would be huge. “If a particular drug can be proven to prevent Alzheimer disease, it would have a tremendous public health impact,” Dr Neil Buckholtz, Chief of the Dementias of Aging Branch of the National Institute on Aging in Bethesda, Maryland, USA, told the Bulletin. “People have estimated that if you delayed the onset of Alzheimer by five years, you would decrease the number of people with the disease by half and cut the cost too.”

Those numbers could quickly add up. WHO estimates that by the year 2025 more than 22 million people worldwide will have Alzheimer disease. ■

Christie Aschwanden,
Nederland, Colorado, USA