A keystone of Australia’s pandemic planning has been the introduction of a vaccination program as quickly, safely and effectively as possible after the pandemic strain of influenza is recognised. Since 2003 the Australian Government has been putting in place sleeper contracts to ensure that when a pandemic strain of influenza emerged Australia would be well-placed to acquire a vaccine.

In April 2009, a novel strain of influenza A (H1N1) was identified. It quickly became apparent that recent seasonal influenza vaccination was unlikely to provide protection against this strain and a vaccine covering it would be required. On 28 May 2009 the Australian Government placed an order for enough pandemic vaccine to vaccinate approximately 10 million individuals (20 million doses).

This paper describes the pre-pandemic planning for vaccine roll-out, controversies encountered during the roll-out and initial progress of the roll-out in New South Wales (NSW).

Pre-pandemic planning
NSW pandemic plans, written in preparation for a more severe pandemic, consider the use of mass vaccination clinics an effective method of distributing vaccines to the population. Consequently guidelines were in development to assist area health services (AHSs) in delivering vaccination clinics. The pandemic vaccine was always likely to be supplied in multi-dose vials due to the increased speed of production. Needles and syringes to administer a vaccine were stockpiled by the Australian Government.

Progress during the pandemic
Decisions regarding the supply of the pandemic vaccine were made early in the pandemic, one month after the World Health Organization declared an emergency and well before the disease was established in Australia. It was correctly anticipated that worldwide demand for the pandemic vaccine would outstrip the production capacity of the vaccine manufacturer. This was despite the pandemic being characterised as moderate and the World Health Organization stating that the overwhelming majority of cases recover with no treatment. By August 2009 it became clear that a vaccine would likely be available only after the first wave of the pandemic had subsided and the rate of admissions for critical care and deaths had decreased.

Controversies in rolling out this program
Guillain-Barré syndrome
In 1976 the United States of America implemented a large-scale vaccination program for a swine-origin influenza virus. Following the roll-out, and particularly in the first 5 weeks after vaccination, a number of people developed Guillain-Barré syndrome (at a rate of around one case per 100 000 vaccinated). This incidence has not been found in other influenza vaccination programs; nevertheless concerns remained that the 2009 H1N1 vaccination program might lead to an increase in cases of Guillain-Barré syndrome in the community. In response to these concerns the
Australian Technical Advisory Group on Immunisation released an information sheet about Guillain-Barré syndrome and the vaccine, stating that recently there had not been an increased association between Guillain-Barré syndrome and influenza vaccination and that vaccination may, in fact, be protective against the development of Guillain-Barré syndrome.9

Use of a multi-dose vial

While it has long been documented that multi-dose vials would be used for a pandemic influenza vaccination program, the use of the vials in a mild pandemic elicited concern among health professionals. Several instances of transmission of infectious diseases, from dates prior to the adoption of modern infection control practices, were cited as a cause for concern by infectious disease specialists and subsequently reported in the media.10 While it is preferrable to use a single-dose pre-filled syringe, due to the potential infection control issues, the worldwide demand for the vaccine was such that taking delivery only in that form would have created a significant delay in vaccine availability.

The inclusion of thiomersal, a preservative containing small amounts of mercury, in the multi-dose vial has also elicited some controversy. While thiomersal has been removed from most childhood vaccines as a precaution during a time when it was theorised that it may cause harmful effects,11 the World Health Organization’s Global Advisory Committee on Vaccine Safety has concluded that there is ‘…currently no evidence of mercury toxicity in infants, children, or adults exposed to thiomersal in vaccines’.12

Vaccine program development and roll-out in NSW, 2009

The moderate nature of the pandemic and the fact that a vaccine was only available after the rate of infections was subsiding decreased the need for an urgent roll-out of the vaccine. Within NSW, representative primary care groups and AHSs were consulted regarding preferred methods for vaccine roll-out. Mass vaccination clinics were no longer considered to be appropriate. Rather, a more usual general practitioner (GP) and Aboriginal Health Service-based model for delivery of the vaccine was thought to be more accessible and appropriate.

All AHSs developed plans to deliver the vaccine to healthcare workers, to consider opportunistic vaccination of vulnerable patients and to provide assistance in areas where GP and Aboriginal medical services were less accessible.

Once the vaccine was approved, a dedicated webpage with training packages for the safe use of multi-dose vials and a general introduction to the program was uploaded to the NSW Health website.13 Links to important national documents and webpages were included.

The program commenced for people aged 10 years and over on 30 September 2009. This involved the distribution of hundreds of thousands of doses of the vaccine (n = 664 860)
within a 5-day period across NSW. Figure 1 shows the number of doses delivered between 23 September and 23 October 2009.

A number of enquiries (24) were received at the NSW Department of Health’s generic email account. The majority were enquiring about gaining access to the vaccine (Figure 2).

A computer-aided telephone interview as part of the NSW Health Survey will monitor vaccine uptake within NSW and NSW Health is working with the Therapeutic Goods Administration to monitor any adverse events following H1N1 vaccination. Triage-text searching of emergency department presentations for persons presenting with possible adverse reactions is also ongoing.

Future planning
In December 2009 the pandemic vaccine was made available to children aged from 6 months to 10 years.14 It has been announced that the 2010 annual influenza vaccination will contain the pandemic strain, leading to a need for clear communications to medical practitioners regarding the roles of the pandemic vaccine and the seasonal influenza vaccine.15

Future pandemics may be more severe than that caused by the current H1N1 virus, resulting in a completely different approach to a national vaccination program. This possibility emphasises the importance of building a degree of flexibility into any pandemic response plan.

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References


