



SENTINEL GENERAL PRACTICES

The NSW Public Health Bulletin has recently highlighted the usefulness of sentinel general practices for monitoring non-notifiable infectious diseases such as influenza¹.

Another recent review article shows general practice surveillance need not be confined to infectious diseases, but can cover other conditions such as asthma and injuries and other risk factors like hypertension and smoking².

Sentinel general practices have operated in the UK³, other parts of Europe^{4,5} and parts of the US⁶ for several years. They have also been running successfully in South Australia⁷ and more recently Queensland⁸.

The National Health and Medical Research Council recommended at its 97th session in 1986 "that each State and Territory health authority be encouraged to produce sentinel morbidity survey similar to the survey in South Australia"⁹.

Although there have been attempts to establish general practitioner sentinel networks in NSW since 1975¹⁰, long-term general practice surveillance has generally been unsuccessful in this State, partly because of a lack of the necessary administrative support. The network of new Public Health Units (PHUs) could be the catalyst in helping set up the schemes throughout NSW.

A network of six sentinel general practices covering five disease entities (Table 1) was established in the Illawarra in June 1990, and has been producing regular reports (Figure 1). Between 500 and 600 patients are seen in participating practices each week. The general practitioners (GPs) also receive regular feedback of the results, and found these sufficiently useful to have asked that the network be expanded to 12 practices (a surveillance base of 1000 to 1300 patients weekly) from October 1990.

The number of conditions under surveillance is likely to be increased to a (maximum) of 12 over a one- to two-year period.

Experience from the Illawarra may be of use to other PHUs in setting up similar networks.

SUCCESSFUL IMPLEMENTATION

Successful implementation depends on an understanding of the strengths and limitations of sentinel general practice surveillance. It is not intended to provide complete prevalence data or a comprehensive data set.

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Sentinel General Practices

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TABLE 1

| DISEASE AND CLINICAL DEFINITIONS | | DISEASE RECORDING |
|----------------------------------|--------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Number | Disease | Clinical Criteria |
| 01 | Influenza | Fever, chills, headache, myalgia, coryza, sore throat, cough and prostration (six out of eight required); nasopharyngeal aspirate for index cases only (send patient to Path. OPD, Wollongong Hospital). |
| 02 | Measles | Fever, coryza, cough, florid rash beginning on the face, conjunctivitis, Koplick's spots. May be modified in vaccinated people. |
| 03 | Lower RTI | Fever, new or changed cough, production of new or changed sputum (no pathology initially). |
| 04 | Asthma | Recurrent wheeze, cough, dyspnoea (existing and new cases where symptoms/treatment is primary reason for consultation). |
| 05 | Infectious Mononucleosis | Fever, sore throat, often exudative pharyngitis or tonsillitis and/or lymphadenopathy; morbilliform rash; splenomegaly or persistence of exudative pharyngitis (for more than 3/7 in spite of antibiotic therapy); FBC (lymphocytosis more than 50 per cent with more than 10 per cent atypical cells); ± positive Monospot. |
| 06 | Free Comment | Space is provided for your free comments. This may be an interesting case which you think is worthy of reporting to us if there is a cluster/epidemic occurring. |

But it does show changes in incidence (in terms of general practice attendance), allow the GP to make more confident diagnoses, and facilitate an early response to changing incidence or distribution of particular diseases.

Important initial steps are:

- Enlist the support of local GPs. (The sub faculty of the RACGP, or postgraduate education group is an obvious route.)
- Emphasise the value to them of active participation (such as early feedback on changing disease patterns, greater certainty in diagnosis and prescribing and reduced need for confirmatory laboratory tests).
- Form a management committee with majority representation from local GPs.
- Allow the management committee to select participating practices. A good geographic and sociodemographic spread is essential — there should be a minimum of 1 per cent of possible general practices in the area, but more is desirable.
- Allow the management committee to select initial list of diseases and conditions. Keep it short — no more than four or five. There should be one or two relatively common conditions (to maintain interest). Others should reflect local priorities and interests.

- Allow a space on the form for free comment. This will provide early warning of increased incidence of other diseases, such as infantile gastroenteritis, or anything unusual.
- Ensure regular feedback to the GPs.
- Develop an outer circle of interested practices to allow interchange of participating practices over time.
- Have regular meetings of the management committee to fine-tune the network.

CONCLUSIONS

Most people continue to regard the GP as their main source of primary health care and advice¹.

Links formed between PHUs and GPs in setting up a successful sentinel general practice network will be valuable in securing their co-operation in other public health initiatives such as improved immunisation and notification rates.

The Illawarra sentinel general practice network now consists of 12 practices and 14 GPs — about 10 of whom are active at any one time. The network has already proved its value several times:

- It was able to ascertain rapidly the extent of spread of a measles outbreak at Woonona.
- It was able to reassure authorities that a patient with Legionella was an isolated case, and that there was no major upsurge in lower respiratory tract infection at the time.
- It also revealed that despite reports in the media of a 'late flu epidemic' there had been no rise in influenza cases monitored by the network.

There is a proposal for a national network of sentinel general practices — Australian Sentinel Practice Research Network (ASPERN)². The research committee of the NSW Faculty of the Royal Australian College of General Practitioners has been contacted about NSW participation.

With infrastructure and support provided by the PHUs, a NSW network of sentinel general practices would not only provide additional useful public health data for this State, but would complement the proposed national network. Most unit directors appear to favour such a network. Dr Michael Levy from the Epidemiology Branch will co-ordinate efforts to implement this approach in 1991.

Dr David Jeffs

Director — Illawarra Public Health Unit

Dr Rod McMahon

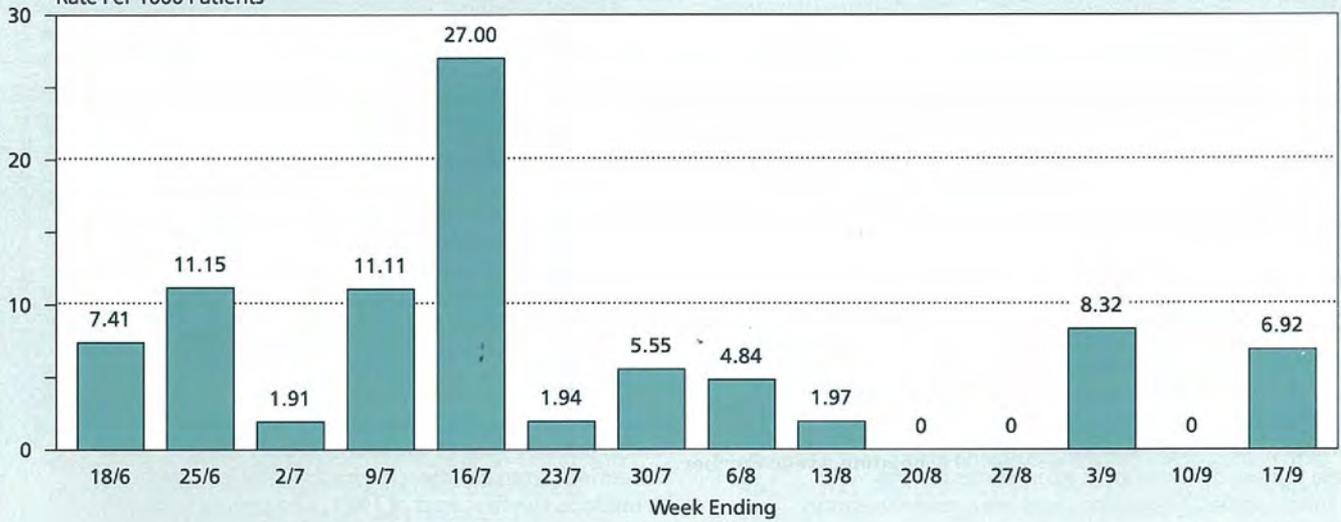
Chairman — Illawarra Institute of General Practitioners

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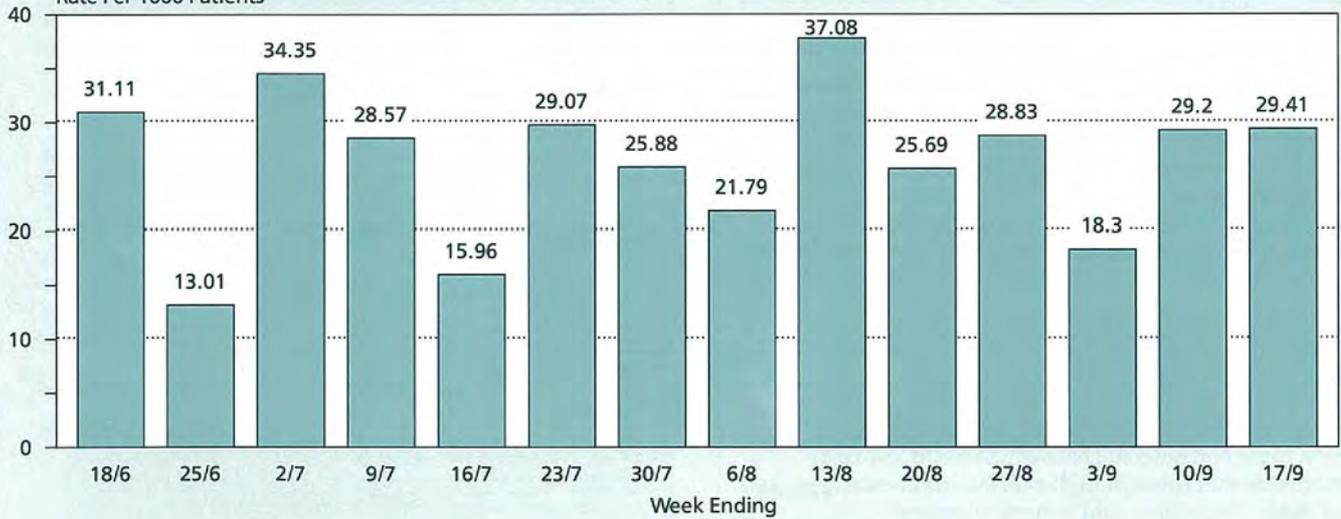
FIGURE 1

SAMPLE WEEKLY RETURNS FROM ILLAWARRA SENTINEL GENERAL PRACTICES

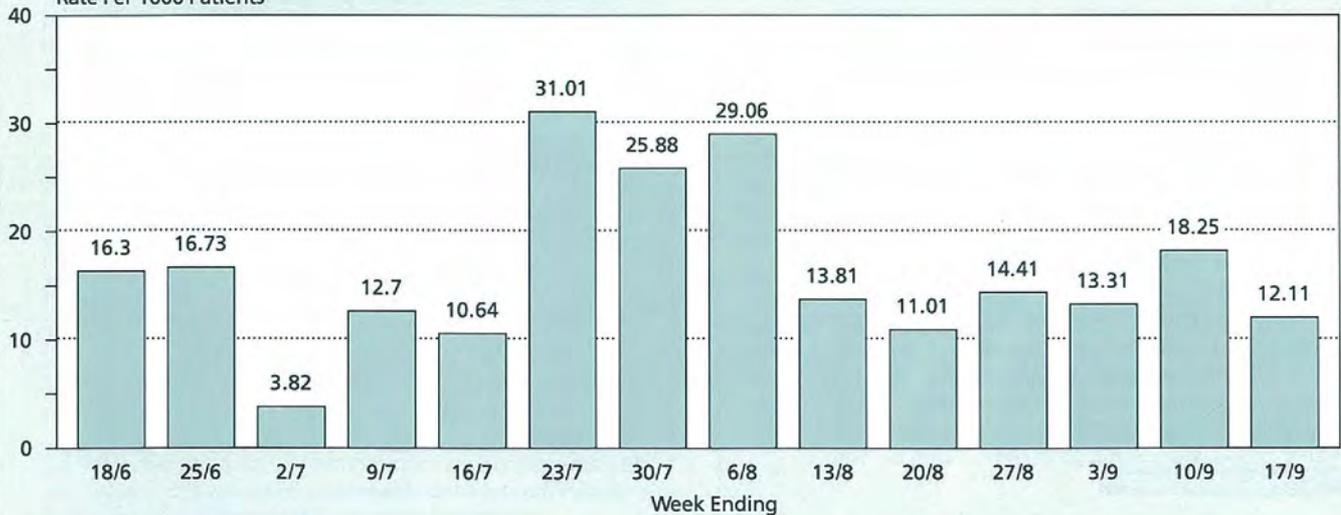
Occurrences of Influenza
Rate Per 1000 Patients



Occurrences of Lower RTI
Rate Per 1000 Patients



Occurrences of Asthma
Rate Per 1000 Patients



STUDY OF HEPATITIS B IN SCHOOLCHILDREN

Horizontal (non-sexual, perinatal or parenteral) transmission of hepatitis B infection may sometimes occur in schools even in countries with low endemicity but where there are small numbers of carriers among the students. Horizontal transmission is documented in households where one person is a carrier but the degree of infectivity of individual carriers, even those who are HBe Ag-positive, may vary widely.

Horizontal transmission of hepatitis B infection has been reported among schoolchildren in New Zealand. But there is little evidence of horizontal transmission between high- and low-carrier rate groups in Australia so routine vaccination of infants in the general population is not advocated. The Australian studies have involved a mixed-race community in Brewarrina, NSW¹, and schoolchildren in the Kimberley region of Western Australia². A third study, soon to be published, was carried out in rural NSW³ and the National Health and Medical Research Council is supporting a Darwin survey, the results of which have not yet been made public. No attempt had been made to survey schoolchildren in the major urban population centres to help formulate vaccination policy for the whole of Australia.

Children of any race who acquire hepatitis B infection very early (at less than five years of age) probably have a higher risk of becoming chronic carriers of the virus and of developing serious long-term sequelae. Review of data from highly endemic areas has emphasised the importance of documentation of the levels of horizontal transmission among young children⁴. The mode of transmission in these circumstances has not been clarified but is suggested to be by saliva or blood from skin abrasions.

Early in 1990 it was decided to study horizontal transmission of hepatitis B in Sydney schoolchildren. The best study design would have been one in which a cohort of low-risk children was followed from birth through pre-school and primary school, comparing infection rates and outcome between those in low-risk environments and those in high-risk environments. Such a study would have had to involve several thousand children and would have been time-consuming and costly.

A reasonable starting plan for a survey of horizontal transmission was therefore a cross-sectional serological study sampling children in the last year of primary school (aged about 12 years). These children have had a long period of school and in some cases pre-school to be exposed to horizontal risks. A plan was developed to sample a group of 750 low-risk children (children of ethnic groups with a carriage rate of less than 5 per cent) in 'low-risk' schools (schools with fewer than 5 per cent children from ethnic groups with more than 10 per cent carriage rate); and, 750 from 'high-risk' schools (more than 20 per cent of children from ethnic groups with more than 10 per cent carriage rate). A similar number of high-risk children (children of ethnic groups with a carriage rate of more than 10 per cent) would be sampled from the same 'high-risk' schools. The Director General of School Education gave permission for the studies.

The Sydney survey began in August 1990, at the beginning of the third school term. A group of schools has been chosen for the control arm of the study, mainly from the northern metropolitan area. All children taking part are volunteers whose parents have given written informed consent. Nurses and doctors from The Children's Hospital, Camperdown, collect venous blood from the children during school hours. There has been excellent co-operation from the schools and a good participation rate from the children. Some demographic data are being collected as well as information about the schools' lifestyles (such as types of games). The sera are being tested for hepatitis B markers and parents will be given their child's results, which will be confidential. Counselling will be available for the families of chronic carriers. Schools will not be given individual results but will be informed of the overall results of the survey.

Collection of sera from the control group has been completed this year in the fourth school term. Collections from children in the 'high-risk' schools will be made in the first part of 1991. The project is jointly funded and supported by the NSW Health Department and The Children's Hospital. Results will be available promptly and will be helpful in evaluating whether universal hepatitis B vaccination is needed.

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FACT SHEETS FOR NSW

With this issue of the Public Health Bulletin readers will find the first of a series of NSW Health Indicator Fact Sheets being produced by the Epidemiology and Health Services Evaluation Branch.

Fact Sheets will provide information on priority health goals in a format which is easy to use and understand. Fact Sheet No 90/1 is about the main cause of death in NSW — ischaemic heart disease — and includes time trends, age and sex differences and comparisons between health areas and regions.

Send your comments on the Fact Sheets to Judy Jones, Health Department Epidemiology Branch, Locked Bag, PO Box 961, North Sydney, 2059. The facsimile number is: (02) 391 9232.

HIV TESTING IN METHADONE PATIENTS

Intravenous drug users (IVDUs) with human immunodeficiency virus (HIV) infection are an important potential source of HIV infection through needle sharing and/or sexual contact. There is however a lack of data about HIV prevalence and incidence data for IVDUs in NSW.

The Epidemiology and Health Services Evaluation Branch, with the NSW Methadone Program, recently explored the feasibility of improving data on the HIV status of IVDUs by collecting patient HIV testing data from methadone prescribers in NSW.

The NSW Methadone Program is managed by the Directorate of the Drug Offensive (DODO) in conjunction with the Pharmaceutical Services Section of the NSW Health Department, Area Health Services and Health Regions. The Health Department is responsible for granting approval to doctors to prescribe methadone. Each prescriber's approval is reviewed every six months.

Patients receive treatment at public clinics, from approved private medical practitioners or in the prison methadone program. There are similar numbers of public and private patients in the program. Treatment is available in the Sydney metropolitan area, at several regional clinics and from prescribers at rural centres. The patients represent a sizeable (more than 4000 patients¹) and accessible group of intravenous drug users and former users. While they are not necessarily representative of IVDUs in general, they may comprise up to one-third of the estimated 12,000 IVDUs in NSW².

HIV-RELATED POLICIES

The program has documented policies on HIV testing among patients. These include: counselling, confirmatory testing of those with a positive HIV test and, when patients are being assessed for entry to the program, HIV-related risk assessment and HIV screening when requested by the patient³. There is no policy on regular follow-up screening.

EXISTING DATA ON HIV STATUS

Entry and exit forms

Prescribers are required to submit to the Health Department a detailed entry assessment form and a termination form for each patient. They also update selected patient details, such as dosage, every six months for renewal of authority to prescribe methadone. These data are collated by the Department. Neither the termination form nor the six-monthly prescription update includes information on HIV status.

Although the entry form has a three-part question on HIV testing and status, the data it yields are inadequate for surveillance purposes. About half the prescribers do not complete the questions (completion of all items is not enforced) and the information is based on patient self-report so its reliability is uncertain. A 'Review of Methadone Treatment' form which is also due every six months includes questions on HIV status but compliance by doctors in completing this form is poor.

HIV testing practices

There is a variety of screening practices, which range from arranging HIV tests at the methadone clinic for virtually all patients to always referring patients elsewhere (such as clinics specialising in AIDS/HIV) for HIV-test-related action. Neither the Health Department nor DODO has collected systematic data on individual prescribers' or clinics' HIV screening practices.

STUDY OBJECTIVES

Our chief objective was to develop and trial a new procedure for directly obtaining accurate, timely HIV surveillance data from methadone prescribers. The main data of interest were:

- HIV sero-conversion prevalence rates among methadone patients; and,
- HIV sero-conversion incidence rates among methadone patients.

The data would also provide:

- Baseline data on HIV testing status (when and where previously tested) for new patients;
- Information on HIV testing practices of methadone prescribers; and,
- Opportunities to explore relationships between HIV status and selected patient characteristics⁴.

METHODS

Ten private prescribers and three public clinics agreed to complete four items of information on patients who had had an HIV test. Prescribers were selected for this study because they were likely to be interested and/or co-operative. Participation was voluntary and of those approached, only one refused to take part. Some prescribers offered to take part without being approached. We discussed the study with the individual prescribers to confirm their co-operation and commitment.

We supplied specially designed data collection forms on which the prescribers recorded, for each patient who had an HIV test, the month and year of the most recent test, where the test took place, the result and whether the result was verified by the prescriber or based solely on the patient's self-report. Patients were identified only by their unique identifying number⁵. Names and addresses were not included. We requested information on the following patients:

- Those on the program at September 30, 1989;
- Those who entered the program in October, November or December 1989; and,
- Those who had an HIV test in October, November or December 1989.

RESULTS

Prescriber response

Of the 13 prescribers who agreed to participate, 10 supplied the information. These 10 prescribers were treating around one-quarter of the patients in the program. Their patients were similar to other patients on the program in terms of sex, age, marital status and duration of narcotic use and dependence.

Most prescribers had to be followed up, some many times, before any forms were returned. The main reasons given for slow completion of forms were a lack of time (especially for those without secretarial support) and having to identify patients by number rather than by name.

HIV testing practices

The private prescribers in this study had incorporated routine HIV testing into their entry procedures and were about to begin or had already started regular re-testing for patients in treatment. In the public clinic, procedures ranged from almost complete testing at the clinic to complete referral. Private patients, therefore,

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HIV Testing in Methadone Patients

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were much more likely to have been tested and those who were tested during September-November 1989 were usually tested around the same time either by the prescriber or at a clinic with which that prescriber was associated.

Data quality

Data on HIV testing collected directly from prescribers may be more accurate, complete and/or reliable than that available on the treatment assessment form. Prescribers verified the HIV test results of most (91 per cent) of their patients. For the 70 per cent of patients whose most recent test was in September 1989 or later, this was usually because the prescribers had arranged the test.

As with the total population of methadone patients, HIV-related information for patients in this study was usually missing from their entry assessment forms (for example, HIV testing status had not been recorded for 64 per cent of the patients).

HIV test results

The prescribers indicated that 467 (35 per cent) of their patients had had an HIV test. Of these, 9 (2 per cent) were HIV-positive. All positive results were verified by the prescriber. This low HIV-positive rate is consistent with four recent studies of IVDU in western Sydney, in each of which fewer than 1 per cent of patients were found to be HIV-positive^{6,7}.

Most patients (95 per cent) had been tested in 1989. Almost half (49 per cent) were tested between October and December 1989, usually by their methadone prescriber. Patients whose most recent HIV test was before October 1989 were similar to those who had not been tested in age, sex, marital status, employment status, partner's drug use, duration of drug use, duration of drug dependence and time spent in methadone programs. Those with children were more likely to have had an HIV test.

CONCLUSIONS

- 1 Prevalence and incidence data on HIV testing and status of patients in the Methadone Program can be obtained directly from methadone prescribers.
- 2 Considerable follow-up was needed to obtain voluntarily supplied data from this group of cooperative prescribers. If a system of compulsory supply of HIV testing data were to be introduced, the important issue of how to enforce this would need to be addressed. For example, prescribers tend to comply with the six-monthly prescription update because Departmental authority for them to continue to prescribe methadone is contingent upon it, whereas they leave many sections of the treatment assessment form unanswered because no such consequence applies.
- 3 The volume and quality of the data available from each prescriber will be related to the HIV testing practices of that prescriber. Those who do not directly arrange testing for their patients have difficulty supplying accurate, verified data, even of the most basic kind (date, place, result of

HIV test), since the confidentiality of HIV test results is protected by legislation. Accurate, updated information on individual prescribers' HIV testing practices is needed.

- 4 The Methadone Program's policy on HIV testing allows prescribers the full range from no testing or referrals whatsoever to the equivalent of comprehensive regular screening. The policy needs clarification.
- 5 Where prescribers have access to HIV testing information about their clients (for example, because they arrange regular HIV testing), the data supplied are likely to be more timely, accurate and complete than has been available from entry assessment forms.
- 6 The accuracy and comprehensiveness of data collected would be improved if prescribers were encouraged to arrange HIV testing, for example by making resources available to assist them with testing, counselling and/or data transcription.
- 7 HIV surveillance data obtained from prescribers can be successfully linked with patient characteristics on the Methadone Stats Unit database via the patients' unique identifying numbers.
- 8 Introduction of uniform and reliable HIV antibody screening procedures in the Methadone Program would assist efforts to gain a clearer picture of the prevalence and incidence of HIV in the patient population.

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The assistance of the Directorate of the Drug Offensive, in particular Stuart Riley of the NSW Methadone Program, and of the participating prescribers and clinics is gratefully acknowledged.

EDITORIAL NOTE

Following on from this study, a panel of NSW AIDS and drug researchers, clinicians and administrators was formed to draft strategies to improve the monitoring of HIV infection in IVDUs.

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2. Sandland, R, *Estimation of the Number of Heroin Users in NSW using Police Arrest Data: Development of a Statistical Model*. Sydney: NSW Drug and Alcohol Authority, April 1986.
3. Directorate of the Drug Offensive, *Policies and Procedures for the Methadone Treatment of Opioid Dependence in NSW*. Sydney: NSW Department of Health (undated).
4. The Methadone Stats Unit (Directorate of the Drug Offensive) supplied a de-identified database of patients in the methadone program (current to October 1989) which was derived from completed treatment admission forms.
5. The Department of Health assigns a unique identifying number to each patient who starts on the methadone program. Patients retain this number through different entries, exits and prescribers.
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NSW HEALTH INDICATORS

Health Areas and Regions

This Fact Sheet is part of a series produced by the Epidemiology and Health Services Evaluation Branch and the Information Centre of the NSW Health Department to provide information about important health indicators in the State. Data on deaths are derived from the official mortality statistics for NSW based on deaths reported to the Registry of Births, Deaths and Marriages. Data on hospital admissions are based on the Hospital Inpatient Statistics Data Collection supplied to the Department by all hospitals in NSW. Disease classifications are from the International Classification of Diseases, 9th Revision.

HEART DISEASE 1990

1 - DEATHS DUE TO ISCHAEMIC HEART DISEASE

Heart disease is the leading cause of death in NSW. In 1988, 27 per cent of all deaths were due to ischaemic heart disease. The main risk factors for heart disease have been well-documented — elevated blood cholesterol levels, smoking, elevated blood pressure, overweight and lack of exercise. These have been targeted for both primary and secondary prevention.

- The ICD-9 codes 410-414 cover ischaemic heart disease (IHD), comprising acute myocardial infarction, other acute and sub-acute forms of heart disease, old myocardial infarction, angina pectoris and other forms of chronic ischaemic heart disease. Ischaemic heart disease results from the build-up of fatty and scar-like deposits (atherosclerosis) which cause narrowing of the coronary arteries.
- The main cause of death due to ischaemic heart disease is acute myocardial infarction (AMI). (In myocardial infarction, there is permanent damage to the heart muscle.) Studies carried out by the U. of Newcastle (Dobson et al, *Am J of Epid*, 1988, 128) have shown that although definite AMIs are usually coded into their appropriate ICD-9 category (410), probable AMIs may be coded in any of the five categories. For this reason, deaths in all five categories have been included.
- The IHD mortality rate in Australia has reduced considerably in the past 15 years, but it is still higher than more than half of industrialised countries studied by the WHO in 1988. In NSW, the declining death rate due to heart disease which began in the 1960s has continued into the 1980s. Between 1978 and 1988, the mortality rate fell by almost 15 per cent. Overall, the decline has been greater for males so that the differential between males and females has decreased over time.

- The greatest percentage decrease in deaths between 1984 and 1988 has been in the younger age groups. This may be related to a number of factors, such as lifestyle changes, earlier detection and more effective treatment resulting in prolonged life. For older females, there was very little decrease in the death rate between 1984 and 1988.

DEATHS IN NSW 1978-1988† ISCHAEMIC HEART DISEASE

| YEAR | NUMBER | RATE* |
|-------|--------|-------|
| 1978 | 12,480 | 247.1 |
| 1979 | 11,455 | 224.5 |
| 1980 | 11,523 | 223.1 |
| 1981 | 11,731 | 224.1 |
| 1982 | 12,084 | 227.8 |
| 1983 | 11,769 | 219.9 |
| 1984 | 11,559 | 213.9 |
| 1985 | 11,991 | 219.4 |
| 1986 | 11,649 | 210.6 |
| 1987 | 11,936 | 212.9 |
| 1988# | 11,996 | 210.4 |

† ICD-9 Codes 410-414

* Crude death rate per 100,000 population

Provisional figures only

- In four health areas (Central Sydney, Hunter, South-West Sydney and Western Sydney) the standardised mortality ratio (SMR) for males was significantly higher than would have been predicted on State figures. In the latter two of these, the female SMR was also significantly elevated.

DEATH RATE† IN NEW SOUTH WALES, 1984-1988, BY AGE AND SEX ISCHAEMIC HEART DISEASE

| Age Group | Sex | 1984 | 1985 | 1986 | 1987 | 1988* | % Decr. |
|-----------|-----|--------|--------|--------|--------|--------|---------|
| 25-44 | M | 22.8 | 18.9 | 19.2 | 13.9 | 19.5 | 14.5 |
| | F | 5.1 | 4.2 | 4.0 | 3.6 | 3.0 | 41.2 |
| 45-64 | M | 345.2 | 323.9 | 298.9 | 297.6 | 290.4 | 15.8 |
| | F | 104.7 | 110.7 | 108.7 | 88.6 | 84.7 | 19.1 |
| 65+ | M | 1931.7 | 1990.5 | 1821.8 | 1855.2 | 1769.9 | 8.4 |
| | F | 1308.4 | 1349.3 | 1302.8 | 1306.1 | 1297.0 | 0.9 |

† Death rate per 100,000 population

* Provisional figures only

Note: The large percentage decrease for females aged 25-44 years should be interpreted with caution because of the small numbers

**OBSERVED (O) AND EXPECTED (E)* DEATHS DUE TO ISCHAEMIC HEART DISEASE
HEALTH AREAS & REGIONS, 1984-1988, BY AGE AND SEX**

| Area/ Region | Age: | 25-44 | | 45-64 | | 65† | | Total | | Standardised Mortality Ratio*† | |
|------------------|------|------------|------------|-------------|-------------|--------------|--------------|--------------|--------------|-----------------------------------|--------------|
| | Sex: | M | F | M | F | M | F | M | F | M | F |
| Sth. Sydney | O | 56 | 11 | 738 | 225 | 2132 | 2342 | 2927 | 2579 | 89.0 ↓ | 94.4 ↓ |
| | E | 73 | 15 | 867 | 280 | 2346 | 2436 | 3288 | 2731 | | |
| E. Sydney | O | 56 | 12 | 650 | 176 | 1764 | 1890 | 2470 | 2078 | 99.6 | 99.5 |
| | E | 56 | 10 | 586 | 179 | 1837 | 1901 | 2480 | 2090 | | |
| Cen. Sydney | O | 36 | 9 | 526 | 145 | 1546 | 1813 | 2109 | 1967 | 107.3 ↑ | 105.8 |
| | E | 45 | 8 | 489 | 141 | 1430 | 1707 | 1965 | 1859 | | |
| S-W Sydney | O | 96 | 24 | 859 | 304 | 1634 | 1621 | 2589 | 1950 | 110.5 ↑ | 119.5 ↑ |
| | E | 83 | 17 | 790 | 247 | 1468 | 1368 | 2343 | 1632 | | |
| W. Sydney | O | 126 | 15 | 862 | 293 | 1788 | 2013 | 2776 | 2321 | 113.2 ↑ | 122.4 ↑ |
| | E | 87 | 19 | 790 | 238 | 1575 | 1639 | 2452 | 1896 | | |
| Wentworth | O | 37 | 7 | 297 | 76 | 686 | 677 | 1020 | 760 | 105.8 | 107.2 |
| | E | 39 | 7 | 259 | 81 | 665 | 620 | 964 | 709 | | |
| Nth. Sydney | O | 61 | 11 | 832 | 238 | 3131 | 3672 | 4026 | 3922 | 86.4 ↓ | 87.1 ↓ |
| | E | 109 | 23 | 1147 | 376 | 3405 | 4100 | 4661 | 4501 | | |
| Cen. Coast | O | 17 | 7 | 335 | 120 | 1316 | 982 | 1668 | 1109 | 98.2 | 93.6 |
| | E | 26 | 5 | 320 | 116 | 1351 | 1064 | 1698 | 1185 | | |
| Hunter | O | 79 | 18 | 822 | 275 | 2195 | 1994 | 3097 | 2287 | 107.6 ↑ | 103.0 |
| | E | 66 | 12 | 733 | 234 | 2080 | 1972 | 2879 | 2219 | | |
| Cen. Western | O | 29 | 7 | 259 | 79 | 782 | 673 | 1071 | 759 | 105.7 | 100.5 |
| | E | 23 | 4 | 248 | 76 | 744 | 674 | 1013 | 755 | | |
| Sth. Eastern | O | 27 | 7 | 289 | 101 | 851 | 714 | 1167 | 822 | 103.5 | 103.8 |
| | E | 26 | 5 | 294 | 89 | 810 | 698 | 1128 | 792 | | |
| Illawarra | O | 53 | 11 | 526 | 164 | 1198 | 1051 | 1779 | 1226 | 103.5 | 107.7 |
| | E | 40 | 8 | 488 | 149 | 1192 | 979 | 1719 | 1138 | | |
| North Coast | O | 37 | 7 | 529 | 163 | 1728 | 1244 | 2294 | 1414 | 94.4 | 85.6 ↓ |
| | E | 43 | 8 | 534 | 175 | 1853 | 1467 | 2429 | 1652 | | |
| New England | O | 44 | 6 | 418 | 130 | 1130 | 961 | 1593 | 1097 | 104.9 | 102.2 |
| | E | 33 | 7 | 376 | 118 | 1109 | 950 | 1519 | 1074 | | |
| Orana/FW | O | 29 | 5 | 249 | 76 | 631 | 509 | 910 | 590 | 109.6 | 102.0 |
| | E | 20 | 3 | 216 | 62 | 592 | 512 | 830 | 579 | | |
| Sth. West | O | 20 | 6 | 314 | 108 | 968 | 857 | 1302 | 971 | 91.1 | 93.3 |
| | E | 34 | 7 | 366 | 111 | 1028 | 923 | 1429 | 1041 | | |
| NSW Total | | 803 | 163 | 8505 | 2673 | 23480 | 23013 | 32798 | 25852 | 100.0 | 100.0 |

* Expected number of deaths for age group and sex based on State death rate
 * Standardised mortality ratio, that is, Observed deaths divided by Expected deaths X 100
 † Whether there were significantly more deaths (↑) or fewer deaths (↓) than for the State

INFECTIOUS DISEASES

MORE UP-TO-DATE INFORMATION

This month we report data up to November 30, 1990. The information includes notifications processed up to December 1. In future issues we will include provisional data for the month preceding the Bulletin issue — the January 1991 edition will contain notification data for December 1990. In 1991 we will aim to reduce substantially transmission times from the doctor/laboratory to PHU and PHU to Epidemiology Branch in an effort to make the data in the Bulletin as up-to-date as possible.

HIV AND AIDS

HIV data are included for the first time in routine reporting of infectious diseases.

The NSW Health Department reports that 9263 people have been infected with the human immunodeficiency virus (HIV) in NSW. This figure is based on information from the three State laboratories that perform confirmatory testing for HIV infection. The laboratories are at St Vincents, Westmead and Prince of Wales hospitals.

The data are incomplete. St Vincents Hospital, which reported 82 per cent of the positive tests in NSW up to June 1989, has provided data only to September 1989 (6905 cases), and Westmead Hospital has provided data through to August 31, 1990 (1940 cases). The data from Prince of Wales are to November 30, 1990 (418 cases). Based on available data it is estimated the overall figures for NSW would be about 9800 cases at November 30, 1990. This is consistent with projections made by the National Centre for Epidemiology and Population Health at the Australian National University.

Information on the transmission category of these 9263 people infected with HIV will be published in the NSW Public Health Bulletin in January 1991.

By November 14, 1468 cases of AIDS had been reported in NSW. Of these, 906 people have died.

Mechanisms are being put in place to ensure that by February 1991 complete NSW data on AIDS and HIV will be published monthly in the NSW Public Health Bulletin. These data will also subsequently be provided to the National Centre for HIV Epidemiology and Clinical Research, the agency responsible for national AIDS/HIV reporting.

NOTIFICATION DELAY

The response of the Public Health Unit to an infectious disease notification depends on timely notification by doctors. Figure 2 indicates the extent of delays in reporting notifications. Only 2 per cent of notifications are received within two days of the onset of the disease, and only 10.2 per cent are received within one week of onset. Fifty per cent of notifications are received within 16 days of onset. These data indicate the need for faster methods of case reporting to PHUs. We suggest that medical practitioners and laboratory staff be encouraged to telephone notifications where feasible. We look forward to seeing your creative solutions to this problem.

Key features of the conditions notified (as occurring in October) are as follows:

TABLE 2

INFECTIOUS DISEASE NOTIFICATIONS, NSW
Notifications to the end of November 1990

| CONDITION | Number of Cases Notified | | | | | |
|-----------------------------|--------------------------|--------------|------------|--------------|--------------|------------|
| | Period | | | Cumulative | | |
| | October 1989 | October 1990 | Nov. 1990* | October 1989 | October 1990 | Nov. 1990* |
| AIDS | 30 | 32 | - | 256 | 295 | 295 |
| Amoebiasis | 1 | - | - | 7 | 9 | 9 |
| Ancylostomiasis | - | - | - | - | - | - |
| Anthrax | - | - | - | - | - | - |
| Arboviral infection (NOS) | 1 | - | - | 1 | 1 | 1 |
| Brucellosis | - | - | - | - | 5 | - |
| Campylobacter infection | 81 | 154 | 60 | 1434 | 1500 | 1560 |
| Chancroid | - | - | - | - | - | - |
| Chlamydia infection (NOS) | 8 | 21 | 1 | 35 | 361 | 362 |
| Cholera | - | - | - | - | 1 | 1 |
| Congenital rubella syndrome | - | - | - | - | - | - |
| Diphtheria | - | - | - | - | - | - |
| Donovanosis | - | - | - | - | - | - |
| Encephalitis (NOS) | - | - | - | 1 | 1 | 1 |
| Food poisoning (NOS) | - | 1 | 1 | 7 | 22 | 23 |
| Genital herpes | 34 | 75 | 1 | 565 | 857 | 858 |
| Giardiasis | 25 | 25 | 3 | 562 | 508 | 511 |
| Gonococcal ophthalmia neo. | - | - | - | 1 | - | - |
| Gonorrhoea | 27 | 20 | 9 | 490 | 338 | 347 |
| Hepatitis A | 3 | 2 | - | 56 | 24 | 24 |
| Hepatitis B | 47 | 29 | 1 | 392 | 370 | 371 |
| Hepatitis C | N/A | 8 | - | N/A | 24 | 24 |
| Hepatitis unspecified | 5 | - | - | 17 | 3 | 3 |
| HIV | N/A | N/A | N/A | N/A | N/A | **9263 |
| Hydatid disease | - | - | - | 1 | 2 | 2 |
| Infantile diarrhoea (NOS) | 69 | 10 | 4 | 409 | 119 | 123 |
| Legionnaires' disease | 1 | - | - | 48 | 23 | 23 |
| Leprosy | 1 | - | - | 10 | 5 | 5 |
| Leptospirosis | 4 | 3 | 2 | 47 | 39 | 41 |
| Lymphogranuloma venereum | - | - | - | - | - | - |
| Malaria | 5 | 7 | - | 69 | 142 | 142 |
| Measles | 21 | 21 | 14 | 42 | 102 | 116 |
| Meningococcal infection | 2 | 7 | 2 | 46 | 86 | 88 |
| Non specific urethritis | 101 | 90 | 1 | 1388 | 1265 | 1266 |
| Ornithosis | - | - | - | 4 | - | - |
| Pertussis | 37 | 1 | - | 114 | 125 | 125 |
| Plague | - | - | - | - | - | - |
| Poliomyelitis | - | - | - | - | - | - |
| Q fever | 10 | 7 | - | 91 | 124 | 124 |
| Rabies | - | - | - | - | - | - |
| Ross River fever | 5 | 2 | - | 383 | 246 | 246 |
| Rubella | 3 | 1 | - | 3 | 4 | 4 |
| Salmonella infection | 32 | 68 | 39 | 954 | 1179 | 1218 |
| Shigella infection | 5 | 6 | - | 62 | 114 | 114 |
| Syphilis | 27 | 18 | 12 | 267 | 289 | 301 |
| Tetanus | - | - | 1 | - | - | 1 |
| Trachoma | - | - | - | - | 1 | 1 |
| Tuberculosis | 41 | 20 | 5 | 351 | 406 | 411 |
| Typhoid & paratyphoid | 3 | 3 | 1 | 21 | 32 | 33 |
| Typhus | - | - | - | - | - | - |
| Vibrio infection (NOS) | - | - | 13 | 14 | 9 | 22 |
| Viral haemorrhagic fevers | - | - | - | - | - | - |
| Yellow fever | - | - | - | - | - | - |
| Yersinia infection | 5 | 12 | 1 | 71 | 117 | 118 |

* Preliminary data only

** Cumulative data from St Vincents 1984-September 1989, Westmead 1984-August 1990, Prince of Wales 1984-November 1990

Continued on page 59 ▶

FIGURE 2

**NOTIFICATION OF DISEASES
TIMELINESS**

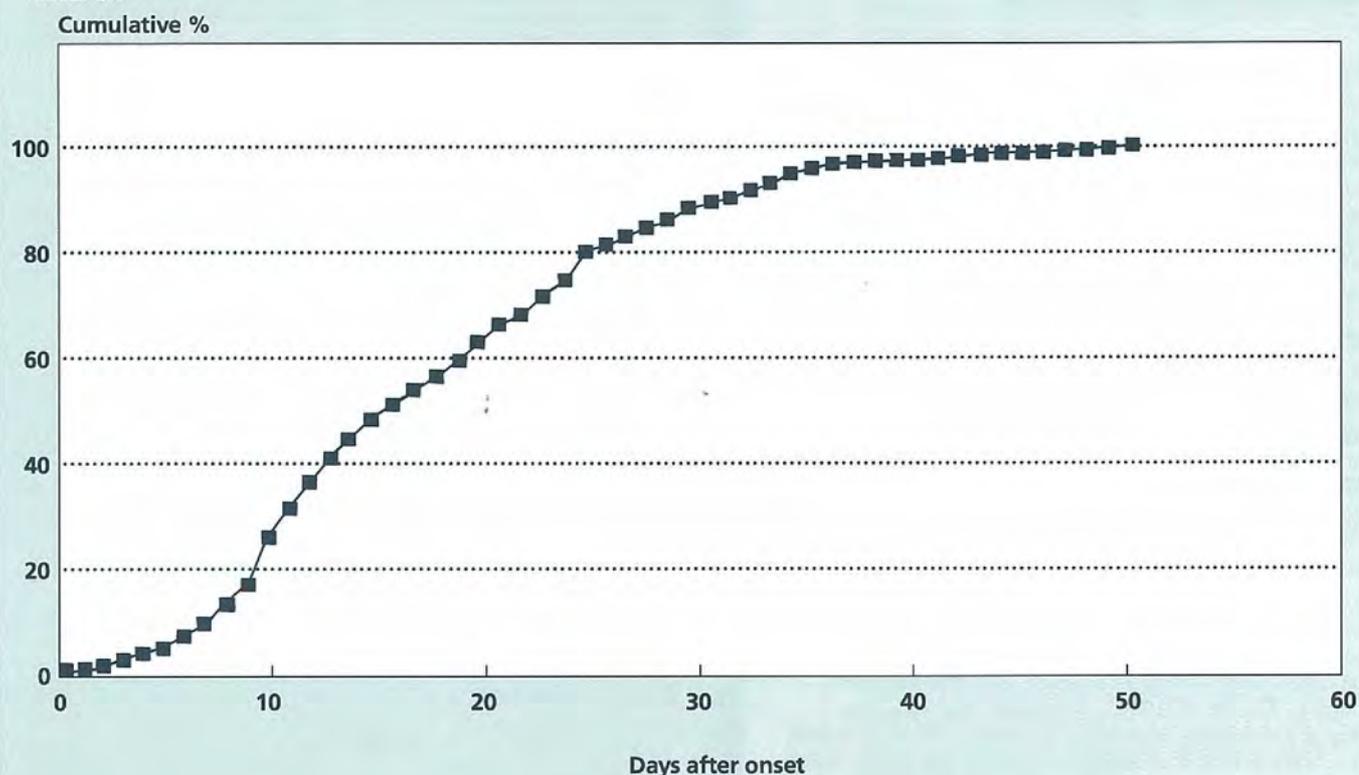


TABLE 3

**INFECTIOUS DISEASE NOTIFICATIONS*
BY HEALTH AREA & REGION, NSW,
JANUARY 1 TO NOVEMBER 30, 1990**

| HEALTH AREA/REGION | DOCTOR NOTIFICATIONS | RATE** PER 100,000 | LABORATORY NOTIFICATIONS | RATE** PER 100,000 | TOTAL NOTIFICATIONS | RATE** PER 100,000 |
|---------------------------|----------------------|--------------------|--------------------------|--------------------|---------------------|--------------------|
| Central Sydney Area | 284 | 85.2 | 115 | 34.5 | 399 | 119.7 |
| Eastern Sydney Area | 713 | 220.6 | 1833 | 567.1 | 2546 | 787.7 |
| Southern Sydney Area | 326 | 61.9 | 269 | 51.1 | 595 | 113.1 |
| South Western Sydney Area | 468 | 76.3 | 240 | 39.2 | 708 | 115.5 |
| Western Sydney Area | 265 | 45.0 | 251 | 42.6 | 516 | 87.6 |
| Wentworth Area | 170 | 63.7 | 295 | 110.5 | 465 | 174.1 |
| Northern Sydney Area | 347 | 47.8 | 260 | 35.8 | 607 | 83.6 |
| Central Coast Area | 102 | 47.2 | 30 | 13.9 | 132 | 61.1 |
| Illawarra Region | 204 | 68.7 | 76 | 25.6 | 280 | 94.3 |
| Hunter Region | 271 | 56.9 | 83 | 17.4 | 354 | 74.3 |
| North Coast Region | 290 | 85.0 | 682 | 199.8 | 972 | 284.8 |
| New England Region | 192 | 78.0 | 379 | 154.1 | 571 | 232.1 |
| Orana & Far West | 181 | 130.1 | 86 | 61.8 | 267 | 191.9 |
| Central West Region | 64 | 38.9 | 71 | 43.1 | 135 | 82.0 |
| South West Region | 49 | 19.5 | 59 | 23.5 | 108 | 43.1 |
| South East Region | 36 | 18.9 | 39 | 20.5 | 75 | 39.4 |
| Unknown | 33 | 0.6 | 60 | 1.1 | 93 | 1.6 |
| Total† | 3995 | 70.1 | 4828 | 84.7 | 8823 | 154.8 |

† Notifications on interstate and overseas residents visiting NSW accounted for an additional 100 cases

* Excludes HIV

** Rate per 100,000 population

Notification Delay

► Continued from page 57

- Pertussis notifications continue to be lower than the comparable period in 1989.
- Measles notifications do not reflect the current situation of medium to large outbreaks throughout at least nine of the 16 Health Areas/Regions. We encourage PHU staff to elicit formal notifications from medical practitioners.
- Tuberculosis notifications for the year are 13 per cent higher than for the comparable period of 1989 — only 6.7 per cent of notifications originated from the country Regions. The NSW Health Department recently reviewed State tuberculosis services. A report on this will be published in a future issue of the Bulletin.
- Meningococcal infections continue to be notified and are 85 per cent higher than 1989 notification rates. New England Region reports a rate of 6.8 cases per 100,000 population a year. Further discussion of this will appear in the January Bulletin.
- Ross River notifications are low and reflect expected seasonal trends. Arbovirus surveillance through chickens and mosquito trapping will continue in 1991. No activity has been detected in the sentinel chicken flocks.
- AIDS figures appearing in previous Public Health Bulletins have been revised upwards following active surveillance by Epidemiology and Health Services Evaluation Branch, Public Health Medicine Registrars and AIDS co-ordinators. This effort revealed 122 previously unreported cases of AIDS, and of these 117 people remain alive. Finding these extra cases will enhance the NSW AIDS services budget by more than \$3.5 million.

INFLUENZA SURVEILLANCE

For the period June 4 to October 28, 1990, the proportions of all general practitioner (GP) consultations for both influenza-like illness and the International Classification of Health Problems in Primary Care, 2nd edition [ICHPPC-2] defined influenza are shown in Figure 3. The proportions are expressed as the number of cases per 100 consultations.

During the study period the Epidemiology Branch received information from 55,663 GP consultations. The proportion of GP consultations relating to influenza-like illness was 3.4 per cent, while the proportion fulfilling ICHPPC-2 criteria was 1.8 per cent.

Of the people presenting with influenza-like illness who had been immunised this year against influenza, 54 per cent were below the NHMRC recommended age of 65 years.

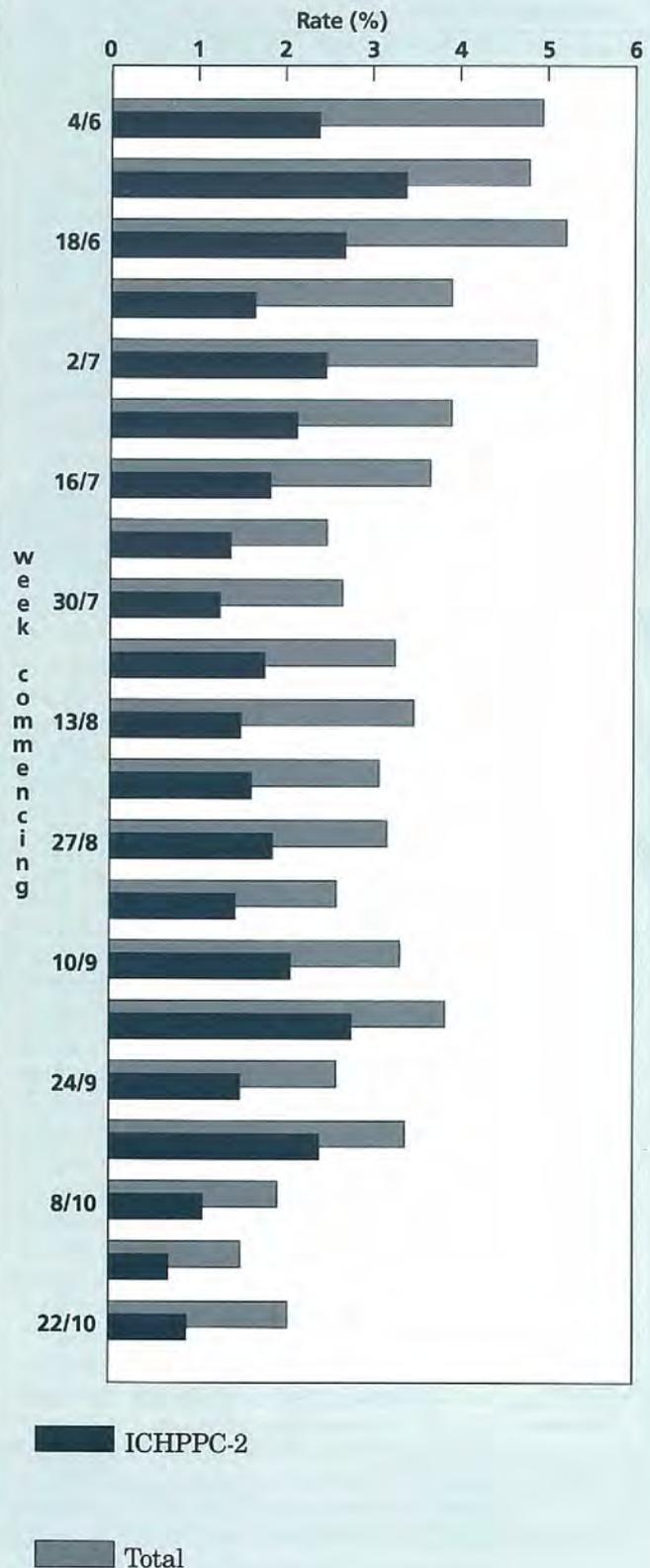
The project was established so any influenza outbreak could be monitored. The study showed that 1990 was not an epidemic year in NSW for influenza. This was in contrast to the predictions, based on the 1989/1990 experience of the northern hemisphere.

The influenza surveillance project provided Epidemiology Branch with experience in sentinel surveillance. Further co-operative projects with the Royal Australian College of General Practitioners are planned for 1991.

*Infectious Diseases Section
Epidemiology and Health Services Evaluation Branch
NSW Health Department.*

FIGURE 3

RATE OF INFLUENZA-LIKE ILLNESS
June 4 to October 28, 1990



rate influenza/100 consultations

TABLE 4

**INFECTIOUS DISEASE NOTIFICATIONS,
BY HEALTH AREA & REGION, NSW,
FOR MONTH OF OCTOBER 1990**

| CONDITION | CSA | ESA | SSA | SWS | WSA | WEN | NSA | CCA | ILL | HUN | NCR | NER | OFR | CWR | SWR | SER | IS | U/K | TOTAL |
|------------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|-----|-------|
| AIDS | 6 | 13 | - | 2 | 2 | - | 6 | - | 1 | - | 2 | - | - | - | - | - | 1 | 1 | 32 |
| Campylobacter inf. | 6 | - | 37 | 8 | 20 | 29 | 12 | 2 | 7 | 4 | 6 | 10 | 1 | 6 | 1 | 1 | 4 | - | 154 |
| Chlamydia inf. | - | 11 | - | - | - | - | - | - | - | - | 6 | 3 | - | - | - | - | - | 1 | 21 |
| Food poisoning (NOS) | - | - | - | - | 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 |
| Genital herpes | 1 | 60 | - | 2 | - | - | - | - | - | - | 8 | 2 | - | 2 | - | - | - | - | 75 |
| Giardiasis | - | - | 4 | - | 1 | - | - | 2 | - | - | 17 | - | - | - | - | - | 1 | - | 25 |
| Gonorrhoea | - | 12 | 1 | 1 | 1 | 1 | - | - | 1 | - | 1 | 2 | - | - | - | - | - | - | 20 |
| Hepatitis A | - | - | - | - | - | - | - | - | - | - | - | 1 | - | - | 1 | - | - | - | 2 |
| Hepatitis B | 1 | 5 | - | 2 | 1 | 1 | 9 | - | 1 | - | 2 | 3 | 2 | - | 1 | 1 | - | - | 29 |
| Hepatitis C | - | - | - | - | 2 | - | 6 | - | - | - | - | - | - | - | - | - | - | - | 8 |
| Infantile diarr. (NOS) | - | - | - | - | - | - | - | - | 3 | - | 7 | - | - | - | - | - | - | - | 10 |
| Leptospirosis | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 | - | - | 1 | 1 | 3 |
| Malaria | - | - | 1 | - | 1 | - | 3 | - | 2 | - | - | - | - | - | 1 | - | - | - | 7 |
| Measles | - | - | 1 | - | - | - | 2 | - | 1 | - | 12 | 2 | - | - | - | 3 | - | - | 21 |
| Meningococcal inf. | 1 | - | 1 | - | - | - | - | - | - | 4 | 1 | - | - | - | - | - | - | - | 7 |
| Nonspecific urethritis | - | 77 | - | 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 90 |
| Pertussis | - | - | - | - | - | 1 | - | - | - | - | - | - | - | - | - | - | - | - | 1 |
| Q Fever | - | - | - | - | - | - | - | - | - | - | 1 | 6 | - | - | - | - | - | - | 7 |
| Ross River virus | - | - | - | 1 | - | - | - | - | - | - | 1 | - | - | - | - | - | - | - | 2 |
| Rubella | - | - | - | - | - | - | - | - | - | - | - | 1 | - | - | - | - | - | - | 1 |
| Salmonella inf. | 2 | 5 | 8 | 7 | 8 | 8 | 14 | 2 | 1 | 1 | 6 | 2 | 1 | 1 | 1 | 1 | - | - | 68 |
| Shigella inf. | - | 1 | - | - | 2 | 1 | - | - | - | - | 2 | - | - | - | - | - | - | - | 6 |
| Syphilis | 2 | 7 | 2 | 2 | - | - | 2 | - | - | - | 1 | - | 2 | - | - | - | - | - | 18 |
| Tuberculosis | 2 | 1 | 3 | 2 | 2 | - | 4 | - | - | 4 | 1 | - | - | - | - | 1 | - | - | 20 |
| Typhoid & paratyphoid | - | - | - | - | - | - | - | - | - | 2 | - | - | - | - | - | 1 | - | - | 3 |
| Yersinia inf. | 4 | 1 | - | 1 | 2 | - | 2 | - | - | - | - | 1 | - | - | - | 1 | - | - | 12 |

TABLE 5

**INFECTIOUS DISEASE NOTIFICATIONS,
BY HEALTH AREA & REGION, NSW,
1 January, 1990 to October 31, 1990**

| CONDITION | CSA | ESA | SSA | SWS | WSA | WEN | NSA | CCA | ILL | HUN | NCR | NER | OFR | CWR | SWR | SER | IS | OS | U/K | TOTAL |
|-------------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|----|-----|-------|
| AIDS | 58 | 117 | 18 | 7 | 12 | 7 | 40 | 2 | 5 | 9 | 6 | 3 | - | 1 | 2 | 1 | 1 | - | 6 | 295 |
| Aeromonas Hydroph. | - | - | - | - | - | - | - | - | - | - | - | 2 | - | - | - | - | - | - | - | 2 |
| Amoebiasis | - | 2 | - | 1 | - | 1 | - | - | - | 2 | 3 | - | - | - | - | - | - | - | - | 9 |
| Arboviral inf. (NOS) | - | - | - | - | - | - | - | - | - | - | 1 | - | - | - | - | - | - | - | - | 1 |
| Brucellosis | - | - | - | - | - | - | - | - | - | - | 3 | 2 | - | - | - | - | - | - | - | 5 |
| Campylobacter inf. | 89 | 71 | 267 | 138 | 164 | 211 | 153 | 31 | 22 | 33 | 96 | 127 | 16 | 20 | 6 | 8 | 33 | 2 | 13 | 1500 |
| Chlamydia inf. | 1 | 197 | 2 | 5 | 3 | - | 1 | - | 25 | 10 | 60 | 45 | 3 | - | 2 | 1 | - | - | 6 | 361 |
| Cholera | - | 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 |
| Encephalitis (NOS) | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 | - | - | - | - | - | 1 |
| Food Poisoning (NOS) | 1 | - | - | - | 6 | 1 | 3 | - | - | - | - | - | - | 1 | 10 | - | - | - | - | 22 |
| Genital herpes | 2 | 626 | - | 16 | 8 | 20 | 3 | 3 | 15 | 21 | 77 | 45 | 8 | 7 | 1 | - | 3 | - | 2 | 857 |
| Giardiasis | 14 | 19 | 51 | 18 | 25 | 26 | 35 | 28 | - | 27 | 208 | 30 | 7 | 7 | - | 2 | 2 | - | 9 | 508 |
| Gonorrhoea | 11 | 199 | 6 | 16 | 10 | 4 | 1 | 4 | 2 | 11 | 28 | 21 | 16 | 4 | 1 | 1 | - | - | 3 | 338 |
| Hepatitis A | 1 | 2 | - | - | 4 | 3 | 5 | - | - | 1 | 1 | 3 | - | - | 3 | 1 | - | - | - | 24 |
| Hepatitis B | 8 | 90 | 8 | 64 | 18 | 7 | 22 | 6 | 6 | 4 | 34 | 33 | 56 | 3 | 5 | 4 | 2 | - | - | 370 |
| Hepatitis C | 2 | - | - | - | 4 | - | 13 | 1 | - | - | 2 | 2 | - | - | - | - | - | - | - | 24 |
| Hepatitis Unspecified | - | - | - | - | - | - | - | - | - | 1 | - | 1 | - | 1 | - | - | - | - | - | 3 |
| Hydatid disease | - | - | - | - | - | - | - | - | - | - | 1 | - | - | 1 | - | - | - | - | - | 2 |
| Infantile diarr. (NOS) | - | - | - | 5 | 4 | 16 | - | - | 15 | 1 | 67 | 9 | 2 | - | - | - | - | - | - | 119 |
| Influenza Type A | - | - | - | - | 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 |
| Legionnaires' dis. | - | 1 | 4 | 3 | 2 | - | 4 | - | 1 | 3 | 1 | - | - | 1 | - | 1 | 2 | - | - | 23 |
| Leprosy | 1 | 1 | - | 2 | - | - | 1 | - | - | - | - | - | - | - | - | - | - | - | - | 5 |
| Leptospirosis | - | 1 | 1 | - | 3 | - | - | - | 4 | 3 | 6 | 5 | - | 2 | 4 | 3 | 3 | - | 4 | 39 |
| Malaria | 9 | 18 | 3 | 5 | 11 | 3 | 41 | 3 | 6 | 7 | 4 | 4 | 1 | 4 | 7 | 1 | 1 | 1 | 13 | 142 |
| Measles | - | - | 18 | 3 | 3 | 1 | 3 | 2 | 1 | 6 | 52 | 6 | - | - | 1 | 5 | - | - | 1 | 102 |
| Meningococcal inf. | 5 | 1 | 10 | 8 | 12 | 4 | 3 | 1 | - | 9 | 12 | 14 | 4 | 2 | - | - | - | - | 1 | 86 |
| Mumps | - | - | - | - | - | - | - | - | - | - | - | 1 | - | - | - | - | - | - | - | 1 |
| Nonspecific urethritis | 1 | 910 | 2 | 143 | 3 | 1 | 2 | 1 | 85 | 82 | 14 | 8 | 4 | - | 1 | 2 | 1 | - | 5 | 1265 |
| Pertussis | 15 | 2 | 8 | 12 | 8 | 20 | 9 | 10 | - | 4 | 10 | 18 | 4 | 3 | - | 1 | 1 | - | - | 125 |
| Q Fever | - | 3 | - | - | 4 | - | 1 | 1 | 2 | 1 | 27 | 30 | 7 | 39 | 4 | 1 | 2 | - | 2 | 124 |
| Ross River virus | 1 | 4 | 1 | 1 | - | 1 | 2 | 1 | 5 | 26 | 96 | 46 | 14 | 7 | 23 | - | - | - | 5 | 246 |
| Rubella | - | - | - | - | - | - | - | - | - | - | 2 | 2 | - | - | - | - | - | - | - | 4 |
| Salmonella inf. | 75 | 61 | 100 | 143 | 128 | 93 | 162 | 28 | 51 | 49 | 88 | 66 | 35 | 18 | 28 | 24 | 18 | - | 12 | 1179 |
| Shigella inf. | 4 | 16 | 3 | 8 | 5 | 3 | 13 | 2 | 3 | 4 | 20 | 11 | 8 | 5 | 1 | - | 6 | 1 | 1 | 114 |
| Syphilis | 16 | 109 | 14 | 26 | 1 | - | 5 | - | 6 | 9 | 14 | 11 | 73 | - | 2 | - | - | - | 3 | 289 |
| Trachoma | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 | - | - | - | 1 |
| Tuberculosis | 57 | 71 | 56 | 46 | 51 | 9 | 45 | 5 | 12 | 21 | 3 | 10 | 3 | 3 | 5 | 3 | 2 | - | 4 | 406 |
| Typhoid & paratyphoid | 1 | 6 | 1 | 1 | 2 | 2 | 3 | 1 | 3 | 3 | 6 | - | - | - | - | 1 | 1 | - | 1 | 32 |
| Vibrio Parahaemolyticus | - | - | 2 | 1 | - | 1 | - | - | - | 2 | - | - | - | - | - | - | - | - | - | 6 |
| Vibrio SPP | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 1 |
| Vibrio Vulnificus | - | - | - | - | 1 | - | - | - | - | - | - | - | - | 1 | - | - | - | - | - | 2 |
| Yersinia inf. | 22 | 8 | 11 | 13 | 8 | 4 | 21 | 1 | 1 | 1 | 16 | 5 | 2 | - | - | 1 | 2 | - | 1 | 117 |

Abbreviations used in this Bulletin:

CSA Central Sydney Health Area, ESA Eastern Sydney Health Area, SSA Southern Sydney Health Area, SWS South Western Sydney Health Area, WSA Western Sydney Health Area, WEN Wentworth Health Area, NSA Northern Sydney Health Area, CCA Central Coast Health Area, ILL Illawarra Health Area, HUN Hunter Health Area, NCR North Coast Health Region, NER New England Health Region, OFR Orana & Far West Health Region, CWR Central West Health Region, SWR South West Health Region, SER South East Health Region, IS Interstate, U/K Unknown, OS Overseas, NOS Not Otherwise Stated

Please note that the data contained in this Bulletin are provisional and subject to change because of late reports or changes in case classification. Data are tabulated where possible by area of residence and by the disease onset date and not simply the date of notification or receipt of such notification.

NEWS AND COMMENT

PUBLIC HEALTH UNIT MEETING

A meeting of Public Health Unit staff was held on November 30 at Rozelle Hospital. The following key points emerged:

ENVIRONMENTAL HEALTH GROUP

The group made several recommendations: that more communication occur between PHUs — directly and through the Bulletin; that all PHUs have a directory of information resources based on the health surveyors policy manual, 24-hour access to computerised databases, and protocols for dealing with health hazards posed by environmental toxins; and that training in environmental health be reviewed.

The group at its next meeting will discuss linkages between organisations concerned with environmental health, prepare a list of environmental health issues, and develop a plan for developing an environmental health strategy.

It was suggested that existing plans for dealing with environmental health hazards and lists of members of relevant committees be made available to PHUs.

That the Fire Brigade has information on the storage of chemicals suggested that the Environmental Health network could improve its links with organisations dealing with environmental pollution.

INFECTIOUS DISEASES GROUP REPORT

The group endorsed the new list of notifiable diseases proposed by the NSW Infectious Disease Advisory Committee and the use of CDC case definitions and made recommendations about information flow from laboratories to PHUs, information content of notification forms, and frequency of meetings.

A general practitioners survey will be conducted in seven PHUs to test notification forms.

The computerised infectious diseases data system is now available. PHUs should start entering data on the system by January 1, 1991.

COUNTRY PUBLIC HEALTH UNIT MEETING REPORT

The meeting looked at practical issues, aiming to identify differences and commonalities. Issues discussed included a fish kill strategy, pesticides in the cotton industry, overseas travel and linking public health medicine training with PHUs.

Changes to the role of environmental health officer were fully supported and the development of links with food inspectors encouraged.

The cost of travel — in money and time — discouraged quarterly PHU meetings, so annual meetings should be considered. One person should represent all regions on each special interest group.

TRAVEL HEALTH ADVICE

Travel health advice consumes substantial PHU resources. David Jeffs and others agreed to advance options for handling this load to the infectious diseases group.

HEALTH PROMOTION

The health promotion budget has doubled since 1986 to more than \$18 million. Sixty per cent of this is distributed to Areas/Regions. In the past two years national and State policies have been developed, with priorities given to nutrition, older people, injuries, cancers, and cardiovascular disease.

Michael Ward indicated that the Department has been instructed to develop specific health goals. PHU and health promotion staff can collaborate to set realistic goals and to develop action and evaluation plans.

GAZETTEL OF MEDICAL OFFICERS OF HEALTH (MOHs)

The list of MOHs for each Area/Region will be gazetted shortly.

THE LEGIONELLA COMMITTEE REPORT

Tony Burns outlined the functions of the committee and the recommendations of the report.

AIDS NOTIFICATIONS

A recent search for unreported cases of AIDS netted 122 new notifications. These notifications attract \$3.5 million of Commonwealth funding for AIDS services.

To improve AIDS surveillance, each Area/Region was asked to provide an HIV/AIDS surveillance contact person, who should liaise with Tim Sladden of the Epidemiology Branch and regularly contact specialist AIDS treatment centres, base hospitals, sexual health services and other doctors who treat people with AIDS.

There was debate about the AIDS notification mechanism. A handout to assist in AIDS surveillance has been produced, including notification forms, a summary of AIDS notification procedure and CDC definitions.

HEALTH INDICATOR FACT SHEETS

Health Indicator Fact Sheets will be produced from available morbidity and mortality figures. The monthly sheets will aim to be brief and useful to planners, evaluators and program staff. The first appears with this Bulletin issue.

IMMUNISATION

The Health and Education Departments should co-ordinate efforts to improve information on and immunisation levels of schoolchildren. This collaboration will be the one area of focus of the Health Department review of immunisation to be completed by May 1991.

SENTINEL PRACTICE IN THE ILLAWARRA

David Jeffs presented an overview of the Illawarra GP Sentinel Surveillance System (see article on Page 51). When concern was raised about Legionnaires' disease in the Illawarra, the sentinel network was useful in assessing general practice attendances for respiratory tract infections and unusual symptoms. Other PHUs should consider the value of establishing small sentinel surveillance networks.

NEWS AND COMMENT

THE ILLAWARRA LEAD STUDY

Blood lead levels were monitored among 170 children in Illawarra primary schools. The study, which was expensive and difficult to perform, involved concerned companies.

PUBLIC HEALTH BULLETIN

The Public Health Bulletin has been delayed by technical and editorial problems. From January, the target will be for Bulletins to appear every second week of the month. It is important to have a rapid turnaround and for State data to be published locally, before publication in the national forum.

The Bulletin offers an important mode of communication between PHUs. It could include reports of meetings, program successes and failures, debates on issues and ideas.

If production problems continued, a plainer format could be considered. But with the appointment of a new sub editor, timely production is expected.

PUBLIC HEALTH TRAINING

There are six Public Health Medicine Registrars, with four more and two Health Services Management Registrars joining the scheme in 1991. Approval has been given to the selection of three non-medical public health trainees.

A group has been formed to develop a plan for future public health training in NSW.

Public health educational sessions should be timed to allow access to staff outside central administration. Short courses in aspects of public health should be developed.

TUBERCULOSIS

The Department has approved the reorganisation of tuberculosis services into the public health system. Margaret Thomas has moved into the Epidemiology Branch. CEOs will be asked to integrate tuberculosis sisters into the PHUs.

The Department is developing a plan to eradicate tuberculosis. South Australia has set as a target the year 2017. A tuberculosis advisory committee has been formed, consisting of the Deputy Chief Health Officer, representatives from the Thoracic Society, Chest Association and two PHUs.

ASTHMA

In the hours after a thunderstorm on November 1 a large number of people attended Tamworth Casualty with asthma. By the end of the week, 96 cases had been to casualty. Attendances in nearby towns were more than 10 times the rate in other areas.

Stephen Corbett outlined the results of an investigation of the outbreak of asthma in Tamworth. While outbreaks have been investigated elsewhere, this appears to be the first thorough study of a non-point source outbreak.

HIV INJURY SURVEILLANCE

The Department plans to introduce surveillance of health care workers exposed to HIV-infected blood and body fluids. The real risk, number of people exposed and outcomes are unknown in NSW. Surveillance will be hospital-based and linked with zidovudine prophylaxis protocols.

PHUs will be asked to distribute forms and information to all hospitals. Follow-up initially will be arranged centrally. A hospital infection control seminar is planned for next year.

FUTURE MEETINGS

Directors of PHUs will meet in the morning to discuss administrative issues and other interested parties will join in the afternoon for broader discussion. Meetings should be timed to coincide with short courses or special interest group meetings. The next meeting is scheduled for March 7 at the Rozelle Conference Centre. Agenda items should be sent to George Rubin.

IN TOUCH ARTICLE DECEMBER ISSUE

The Public Health Association of Australasia requested an article on the Public Health Units for its December issue. Watch out for it!

Health in New South Wales: Current Indicators is now available. To obtain a copy please call Ashley Curtis on (02) 391 9223 or write to the editor.

PUBLIC HEALTH EDITORIAL STAFF

The Bulletin's editorial advisory panel is as follows:

Dr Sue Morey, Chief Health Officer, Department of Health; Professor Stephen Leeder, Professor of Community Medicine, University of Sydney; Professor Geoffrey Berry, Professor of Epidemiology & Biostatistics, University of Sydney; Dr Robert Reznik, Acting Director, Department of Community Medicine, Royal Prince Alfred Hospital; Professor Ian Webster, Professor of Community Medicine, University of NSW; Dr Christine Bennett, Acting Associate Director, Service Development, Department of Health; Dr Michael Frommer, Epidemiologist, Epidemiology & Health Services Evaluation Branch; Ms Jane Hall, Research Officer, Department of Community Medicine, Westmead Hospital; and Mr Michael Ward, Manager, Health Promotions Unit, Department of Health.

The editor is Dr George Rubin, Director, Epidemiology and Health Services Evaluation Branch, Department of Health, NSW.

Design and Production — Health Public Affairs Unit, Department of Health, NSW.

Please send your articles, news, comments or letters to Dr George Rubin — Locked Bag 961, North Sydney NSW 2059 or Fax (02) 391 9293.

Suggestions for improving the reporting of infectious diseases are most welcome.