available to me, and I would further hope that the manufacturers, while taking due and proper notice of the cases which have been brought to our attention, will not fail to recall the silent majority who have benefited, and continue so to do, from this drug.—I am, etc.,

M. D. CASHMAN.

St. Annes-on-Sea,
Lancs.

SIR,—A number of reports of hepatic dysfunction associated with the administration of iprindole (Prondol) have appeared in your columns (7 February, p. 367; 25 April, p. 238; and 7 November, p. 368).

Iprindole has been available to hospitals from November 1967 and to general practitioners from August 1968. Since its introduction it has been widely prescribed and commented upon as being a useful anti-depressant compound with a low incidence of anticholinergic side effects.^{1,2}

To date, some 44 cases of liver dysfunction have been reported with the use of this drug, associated with the usage of 63 million tablets over the three-year period. The reports vary from raised transaminases and bile in the urine on the one hand to overt clinical jaundice on the other; patients concerned returned to normal on withdrawal of the drug.

I would, therefore, like to re-emphasize the statement in our technical literature where the side effects, precautions, and contraindications to be considered when this drug is used are enumerated. Attention is drawn to the fact that jaundice has been reported, and that in such cases further treatment with the drug is contraindicated—as it is with patients who have had a previous history of hepatic disorder.—I am, etc.,

D. J. RICHARDS,
Medical Director,
John Wyeth and Brother Ltd.

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Accident and Emergency Services

SIR,—My letter concerning accident and emergency services (14 November, p. 429) might be construed as suggesting that the casualty department at St. Mary's Hospital (Harrow Road) is not efficient. I wish to correct any such impression by stating that it is an excellent department giving a very high standard of care to patients, and good training for potential surgeons.—I am, etc.,

NIGEL H. HARRIS.

St. Mary's Hospital (Harrow Road),
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Vasectomy

SIR,—As my name was mentioned in Mrs. Helen Wolfers's article on psychological aspects of vasectomy (31 October, p. 297) may I make it clear that, while I was interested in her observations, I do not agree with all conclusions. For example "...

submission to vasectomy may itself be indicative of personality disorder. . . ." Not so. A man whose family is complete is doing the sensible thing in having vasectomy. In interviewing three thousand candidates for vasectomy my colleagues and I have found that an outstanding characteristic of the men is that they are considerate and thoughtful, men who do not want to risk unwanted pregnancies or to subject their wife to sterilization or years of pills.

Further, "... a contraceptive method with harmful side effects . . ." could not, in my experience, be fairly applied to vasectomy. "... It is reasonable to assume that histories of pre-existing marital, sexual, or psychological instability should be taken as a contraindication to vasectomy. . . ." Not so, but as a contraindication to having (further) children. The couple has asked for a small operation that would spare them work, anxiety, and expense. Any child would have, as well as the possibility of genetic defect, the disadvantage of being brought up by parents with instability.—I am, etc.,

J. K. MONRO.

Simon Trust Clinic,
Swindon, Wilts.

Retaining Device for Nasogastric Tube

SIR,—The standard method of retaining a nasogastric tube in position is to fix it to the patient's nose with a piece of zinc-oxide strapping. This simple but crude method has the disadvantages that it is uncomfortable for the patient and deters easy adjustment of the tube's position. Further it cannot be assumed that the position will remain correct throughout the period of its utilization as kinking may occur. Change of position of the patient, retching, and alterations of gastric tone may subsequently prevent the tube coming into contact with the stomach contents. A "nil" gastric aspiration result recorded repeatedly on the balance chart is the give-away of the situation, and indicates the necessity for adjusting the tube's position.

An alternative means of retaining a nasogastric tube in position is a flange 4 cm. by 6 cm. made of German silver bearing at right angles to it a fenestrated blade with

slots of graduated size to accommodate the several gauges of nasogastric tubing (Fig.). The flange of the device is stuck to the patient's forehead with a double-sided plaster; the tube correctly positioned is threaded through the fenestrated blade, which acting with a cleat action grips it and holds it in position without occluding its lumen which remains clear for intermittent aspiration and open drainage. Disposable bags suitable for open drainage management of nasogastric tube are available.

Adjustment of the tube's position is readily accomplished by liberating it from the cleat until the new correct position is obtained then cleating the tube again. These devices can remain in position for several days and are well tolerated by patients who quickly become unaware of their presence.—I am, etc.,

LAURENCE F. TINKLER.

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Bristol.

Thyroid Dysfunction

SIR,—Your timely leading article (11 July, p. 61) on the protean symptomatology of myxoedema draws attention to how easily one can miss the diagnosis of early hypothyroidism. The importance of diagnosing early thyroid failure, however, cannot be overemphasized.

For the past eight months biochemical investigations (measurement of thyroid function—T-3 uptake) have been done on autistic children at the radiotherapy department of the Royal Berkshire Hospital. The study was based on 62 autistic children diagnosed on the nine points Creak scale. Out of these 62 autistic children, 45 were found to be hypothyroid and two were hyperthyroid.

The hypothesis I put forward is that some children diagnosed as autistic are, in fact, suffering from the thyroid dysfunction. Some biochemical disturbance is taking place in the hypothalamic-pituitary-thyroid-adrenal axis manifested clinically as an autistic syndrome or childhood psychosis. The syndrome presents different clinical pictures of hypothyroidism (myxoedematous features) and hyperthyroidism. The autistic manifestations of thyroid failure have occurred as a result of delay in diagnosing early hypothyroidism or hyperthyroidism.

Autism can be a feature of thyroid dysfunction and thyroid disturbance can produce childhood psychosis or autistic syndrome. It is therefore suggested that all children with autistic features should have routine thyroid function tests done.—I am, etc.,

A. ALI KHAN.

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Warneford Hospital,
Oxford.

Complication of Hysterotomy

SIR,—I wish to report an unusual, yet important, complication of hysterotomy. This I believe will be of general medical interest, especially for those who are involved with undertaking termination of pregnancy.

A 33-year-old housewife was admitted to hospital for abdominal termination of pregnancy and sterilization. The indications were parity, proctocolitis, and acute anxiety.

She was para 3+0. The second was a twin pregnancy. All four children were alive and well. Her last menstrual period was on 6 July 1968. The uterine size was compatible with the period of gestation. On 22 October 1968 hysterotomy and Pomeroy sterilization was performed by an experienced member of our team. A single fetus was removed from the uterine cavity, and the postoperative period was satisfactory apart from mild urinary tract infection.

At the follow-up clinic on 27 December 1968 the patient presented with a uterus the size of a 24-week gestation. She was obviously pregnant and she was booked for a hospital delivery. On 1 February 1969 she was admitted in premature labour and had a spontaneous delivery of a live infant weighing 3 lb. (1.4 kg.). The puerperium was normal. The infant continued to thrive.

Although it cannot be proved, there is strong circumstantial evidence that this was a twin pregnancy and that only one gestation sac was removed at the time of hysterotomy. The other fetus must have continued to thrive. As far as I am aware a similar case has never been previously reported. The medicolegal implications of such a case are obvious, and I would suggest that everyone undertaking such a procedure should search for a second gestation sac and make appropriate notes at the time of operation.—I am, etc.,

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West Middlesex Hospital,
Isleworth, Middx.

L-dopa for Postencephalitic Parkinsonism

SIR,—In recent months there have been several trials on the efficacy of L-dopa in the treatment of Parkinson's disease. Few of these have referred to postencephalitic Parkinsonism and it has been noted that patients in this group do not tolerate L-dopa as well as patients in whom the disorder is of a different origin.¹

L-dopa is now freely available to hospital and general practice. Fortunately, most manufacturers temper this free availability with a note of caution, such as: "Long term safety and efficacy have yet to be established."

We wish to present our own findings in a nine-week trial of this preparation in twelve patients with postencephalitic Parkinsonism who have manifested signs of their condition for between 10 and 49 years. There were six women and six men, and their ages ranged from 50 to 70 years. Treatment with traditional therapy (benzhexol and orphedadrine) was continued.

The Table shows the dosages employed (these were given in divided doses three times daily with meals), the duration of treatment, the effects on tremor, rigidity, and writing ability, and also the side effects encountered during the trial. The side effects which may be seen with this drug are reviewed elsewhere,² but those most frequently seen in the present study were nausea, vomiting, psychological disturbances, and involuntary movements. Among the last were facial grimacing, tongue protrusion, and foot stamping; and eight out of twelve patients exhibited some kind of emotional upset, including euphoria, depression, aggressive and antisocial behaviour, hallucinations, and delusions.

Of the twelve patients who commenced treatment, therapy was withdrawn in eight patients because of side effects, seven showed no response at all, three showed substantial improvement,

Patient No.	Age	Stabilized Daily Dosage of L-dopa (mg.)	Duration of Treatment (Days)	Effect on			Side Effects
				T	R	W	
1 ..	54	1,500	63*	○	○	○	Depression; nausea; grimacing
2 ..	60	750	63*	+	○	+	Depression; uncooperative; nausea
3 ..	53	1,500	28	○	+	○	Exhibitionism; more oculogyric crises; involuntary foot stamping; hemiballismus; nausea
4 ..	56	750	8	○	○	○	Mouth twitching; screaming and weeping attacks; respiratory panting
5 ..	64	1,500	36	○	○	○	Nausea and vomiting
6 ..	57	1,500	63*	○	○	○	Involuntary movements of tongue; withdrawn
7 ..	70	1,500	28	++	++	+	Aggressive and uncontrollable; exhibitionism
8 ..	69	4,500	51	○	++	++	None—patient died of acute bronchopneumonia
9 ..	58	1,500	32	+	++	++	Nausea and vomiting; euphoria; thrombocytopenia (Platelets < 10,000/cu. mm.)
10 ..	50	1,500	32	○	○	○	Involuntary protrusion of tongue; respiratory panting; intermittent penile erection
11 ..	58	1,500	33	○	○	○	Involuntary protrusion of tongue
12 ..	61	2,000	63*	+	+	+	Hallucinations and delusions

○ = No improvement.
+ = Moderate improvement.
++ = Substantial improvement.
* = Treatment continues.

T = tremor.
R = rigidity.
W = writing ability.

and two showed moderate improvement. It is noteworthy that in all three patients who benefited conspicuously, the drug had to be discontinued because of side effects. Four patients remain on treatment.

Some of the side effects of this drug will obviously necessitate withdrawal, but the less troublesome ones may be controlled by appropriate adjustment of dosage and the addition of dopa-decarboxylase inhibitors;³ these were not used in the present trial.

It is confirmed that not all types of Parkinson's disease show equal benefit on treatment with L-dopa. Full assessment of the disability and the planning of treatment should be undertaken in hospital and dosage be cautiously increased to the limits of tolerance. In summary, the use of L-dopa in the treatment of our patients with postencephalitic Parkinsonism has been

attended by the frequent occurrence of unpleasant side effects. The poor tolerance shown by these patients calls for further study.

We wish to thank Professor Stanley Alstead for help and advice in carrying out this investigation.

—We are, etc.,

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JANET M. CORNELIUS.

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- 2 Krasner, N., in *British Encyclopaedia of Medical Practice, Medical Progress 1970/71—Clinical Pharmacology and Therapeutics* (Treatment of Parkinson's Disease). In press.
- 3 Tissot, R., *Presse Médicale*, 1969, **77**, 617.

Depression and Oral Contraception

SIR,—Dr. Brenda N. Herzberg and her collaborators are to be congratulated on their careful and painstaking attempt to elucidate the difficult problem of depression occurring in women taking oral contraceptives (17 October, p. 142). Some possible sources of error are listed, but my main difficulty in accepting their conclusions lies in the fact that their controls are unmatched.

Broadly speaking, women requesting contraception fall into two groups: those who want virtually 100% efficiency and those who are satisfied with something less. Excluding certain periods of time associated with massive adverse publicity (for example, late 1969 to mid 1970), into the first group will go the majority of women who have enough or too many children, are unmarried, have had abortions, have suffered from severe physical or psychiatric disease, are either very young or very old (reproductively speaking), and those with unstable family relationships, financial problems, or housing difficulties. To these women a pregnancy would often be a catastrophe, and they therefore choose the

reliable pill. Into the second group go the remainder, and, again excluding those frightened off the pill, will be mainly women delaying or spacing their family. To them a pregnancy may be ill-timed, inconvenient, or unwanted—rarely a catastrophe.

In other words, the reason for the development of depression and for the choice of the oral contraceptive method may well be the same. To say that one group can act as controls for the other without taking this factor into account seems to me unrealistic. The difference between these two self-selected groups is quite enough to explain a difference of 6% in the incidence of severe depression—indeed I would have expected it to be higher than 6% and it may well be that the freedom from the fear of pregnancy conferred by the pill is the reason for this low figure.

Your leading article (p. 127) is relevant. Not only should the doctor look for symptoms of depression, but should also look for causes other than the pill. Not only should the doctor consider changing or abandoning the oral contraceptive, but should also con-