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Controlled Comparison of the Efficacy of Fourteen Preparations in the Relief of Postoperative Pain

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Summary

Thirteen analgesic drugs, four of them at two dose levels, four analgesics in combination with antagonist or neuroleptic agents, and saline have been evaluated simultaneously in the relief of postoperative pain. The method of assessment was designed to favour drugs which provided freedom from pain with minimum depression of consciousness. Only levorphanol 2 mg proved significantly superior to pethidine 100 mg, which was used as the standard reference drug. Oxycodone 10 mg, pentazocine 20 mg, and the morphine 10 mg and cyclizine 50 mg combination were the most successful of the remaining drugs. None of the drug combinations was significantly better than the analgesic drug given alone.

Introduction

There are several possible approaches to the symptomatic relief of severe pain—namely, interruption of afferent nerve pathways, either chemically or surgically; “dissociation” of the patient from his pain, which may be achieved by such psychological devices as suggestion or hypnosis, or pharmacologically by agents such as chlorpromazine; and use of centrally acting analgesics, which may also produce mild sedation and dissociate patients from their pain. Only the latter, which involves the systemic use of drugs with opiate-like actions, offers ease of administration compatible with everyday clinical practice. It is therefore of importance that the relative merits of the numerous available drugs of this class should be established. Though the literature relating to these substances is abundant reports have usually encompassed only a few agents, and lack of standardized methodology makes the compiling of an overall assessment difficult.

The study reported here, which occupied four years, was undertaken to answer this need.

Method and Material

The method has been previously described and presented with a critical assessment of its sensitivity and validity (Loan *et al.*, 1968). Patients who had undergone upper or lower abdominal surgery were admitted to the trial when the nursing staff considered that they required their first postoperative analgesic. Pain severity was assessed before and after treatment on four criteria—namely, patient's and observer's subjective estimates on a five-point scale and, with upper abdominal wounds, measurements of vital capacity and peak expiratory flow rate. Pain relief was inferred from changes in these criteria and also estimated by the patient's direct retrospective comparison with his pretreatment state. Treatment in any individual patient was classified as a success only when all these criteria showed improvement, and this “overall” assessment was used in the final comparison of drugs.

The study was carried out in the same environment by two observers who were successive full-time research workers. Drugs were administered and assessments made under double-blind conditions.

The preparations, doses, and numbers of patients given each drug are listed in Table I, and mean ages and weights of patients are shown in Table II. Ideally, each drug should be included at several dose levels so that the basis of comparison would be dose/response curves, thus eliminating differences due solely to dosage. In practice such an approach would have required a vastly greater number of subjects and would have extended the time far beyond the four years taken to complete this study, which was mainly concerned with com-

TABLE I—Details of Drugs and Dosages used.

	Dose (mg)	No. of Patients
Morphine	10	80
Diamorphine	15	20
Papaveretum	5	20
Oxycodone	20	20
Dihydrocodeine	10	20
Levorphanol	50	40
Phenazocine	2	40
Pentazocine	2	20
Methadone	20	20
Dextromoramide	10	40
Pethidine	5	20
Phenoperidine	100	40
Fentanyl	1	20
Pethidine/levallorphan	2	40
Morphine/cyclizine	0.1	20
Fentanyl/droperidol	0.2	40
Phenoperidine/droperidol	100/1.25	20
	10/50	20
	0.2/5	40
	2/5	40

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TABLE II—Numbers of Patients in Treatment Groups with Operation Sites, Mean Ages, and Weights

Treatment	No. of Patients			Mean Weight ± S.E. (kg)	Mean Age ± S.E. (years)
	Upper Abdominal	Lower Abdominal	Total		
Morphine 10 mg	40	40	80	64 ± 2.8	48 ± 4.5
Morphine 15 mg	10	10	20	63 ± 3.6	54 ± 3.5
Diamorphine 5 mg	20	20	40	65 ± 3.4	49 ± 3.6
Papaveretum 20 mg	10	10	20	62 ± 3.7	49 ± 3.7
Oxycodone 10 mg	10	10	20	64 ± 4.4	40 ± 4.4
Dihydrocodeine 50 mg	10	10	20	64 ± 2.3	44 ± 2.5
Levorphanol 2 mg	20	20	40	62 ± 2.6	49 ± 2.9
Phenazocine 2 mg	10	10	20	62 ± 2.9	48 ± 2.7
Pentazocine 20 mg	10	10	20	65 ± 1.8	52 ± 3.2
Pentazocine 40 mg	20	20	40	64 ± 2.7	45 ± 3.2
Methadone 10 mg	20	20	40	62 ± 2.4	48 ± 2.3
Dextromoramide 5 mg	10	10	20	62 ± 2.4	57 ± 3.4
Pethidine 100 mg	20	20	40	63 ± 1.6	48 ± 2.6
Phenoperidine 1 mg	10	10	20	64 ± 4.2	49 ± 4.4
Phenoperidine 2 mg	20	20	40	65 ± 3.8	46 ± 4.1
Fentanyl 0.1 mg	10	10	20	66 ± 4.1	43 ± 4.4
Fentanyl 0.2 mg	20	20	40	66 ± 3.8	49 ± 3.9
"Pethilorfan 100"	10	10	20	69 ± 4.7	50 ± 3.5
"Cyclimorph 10"	10	10	20	67 ± 2.3	51 ± 3.7
Fentanyl 0.2 mg/ droperidol 5 mg	20	20	40	65 ± 3.7	46 ± 4.1
Phenoperidine 2 mg/ droperidol 5 mg	20	20	40	66 ± 3.6	46 ± 4.1
Saline	40	40	80	65 ± 2.9	51 ± 4.9

TABLE IV—Rank Order of Efficacy of Analgesia expressed as Percentage of Patients obtaining "Successful" Relief as defined in the Text

Incidence of "Success"	
≥75	Levorphanol 2 mg
70-74	"Cyclimorph 10", pentazocine 20 mg, oxycodone 10 mg
65-69	Methadone 10 mg, papaveretum 20 mg
60-64	Diamorphine 5 mg, morphine 10 mg, pentazocine 40 mg
55-59	Dextromoramide 5 mg, morphine 15 mg
50-54	Fentanyl 0.2 mg, pethidine 100 mg
45-49	Dihydrocodeine 50 mg, "pethilorfan 100"
≤44	Phenazocine 2 mg, saline

TABLE V—Rank Order of Drugs in respect of Analgesic Potency as judged by Patient's Estimate alone

	Groups in Alphabetical Order
Over 80% pain relief ..	Diamorphine 5 mg, papaveretum 20 mg, oxycodone 10 mg, levorphanol 2 mg, pethidine 100 mg, methadone 10 mg, "Cyclimorph 10"
75-80% pain relief ..	Morphine 10 mg, morphine 15 mg, pentazocine 20 mg, phenoperidine 2 mg, "Pethilorfan 100"
70-74% pain relief ..	Dextromoramide 5 mg
50-69% pain relief ..	Dihydrocodeine 50 mg, fentanyl 0.2 mg, phenazocine 2 mg
Less than 50% pain relief	Saline

TABLE III—Numbers of Patients showing Improvement in the Recorded Criteria after Administration of the Analgesic

Treatment	Upper Abdominal							Lower Abdominal*					Total					Total Overall Success (%)
	No. in Group	O.E.	P.E.	P.R.E.	V.C.	PEFR	Overall	No. in Group	O.E.	P.E.	P.R.E.	Overall	No. in Group	O.E.	P.E.	P.R.E.	Overall	
Morphine 10 mg	40	30	25	32	28	30	17	40	36	34	34	33	80	66	59	66	50	62.5
Morphine 15 mg	10	9	8	8	5	10	3	10	9	8	9	8	20	18	16	17	11	55
Diamorphine 5 mg	20	19	17	16	14	14	11	20	19	17	16	13	40	38	34	32	24	60
Papaveretum 20 mg	10	9	8	8	8	7	4	10	8	9	10	9	20	17	17	18	13	65
Oxycodone 10 mg	10	9	9	8	7	9	6	10	10	8	10	8	20	19	17	18	14	70
Dihydrocodeine 50 mg	10	6	5	6	5	6	5	10	7	5	5	4	20	13	11	10	9	45
Levorphanol 2 mg	20	19	18	15	19	8	15	20	17	15	17	15	40	36	33	32	30	75
Phenazocine 2 mg	10	7	5	4	5	6	1	10	9	8	6	7	20	16	13	10	8	40
Pentazocine 20 mg	10	9	7	6	9	9	5	10	10	8	8	9	20	19	15	14	14	70
Pentazocine 40 mg	20	17	14	8	15	16	11	20	18	15	17	14	40	35	29	25	25	62.5
Methadone 10 mg	20	17	18	17	16	16	10	20	20	16	17	16	40	37	34	34	26	65
Dextromoramide 5 mg	10	7	8	8	9	9	5	10	9	6	7	6	20	16	14	13	11	55
Pethidine 100 mg	20	19	18	15	7	9	6	20	18	15	13	15	40	37	33	28	21	52.5
Phenoperidine 1 mg	10	7	6	5	5	8	4	10	5	4	6	4	20	12	10	11	8	40
Phenoperidine 2 mg	20	15	13	15	13	14	9	20	19	18	17	17	40	34	31	32	26	65
Fentanyl 0.1 mg	10	7	5	5	7	6	4	10	6	5	6	4	20	13	10	11	8	40
Fentanyl 0.2 mg	20	15	12	13	12	13	6	20	18	14	15	14	40	33	26	28	20	50
"Pethilorfan 100"	10	9	9	8	6	5	3	10	10	7	9	6	20	19	16	17	9	45
"Cyclimorph 10"	10	9	8	9	7	7	5	10	10	9	9	9	20	19	17	18	14	70
Fentanyl 0.2 mg/ droperidol 5 mg	20	18	12	18	15	14	9	20	16	16	12	12	40	34	28	30	21	52.5
Phenoperidine 2 mg/ droperidol 5 mg	20	18	16	16	13	17	10	20	19	18	18	16	40	37	34	34	26	65
Saline	40	18	9	11	21	31	5	40	23	18	21	9	80	41	27	32	14	17.5

*Vital capacity and peak expiratory flow rate were not recorded in the groups having lower abdominal operations.
O.E. = Observer estimate. P.E. = Patient estimate. P.R.E. = Patient's retrospective estimate. V.C. = Vital capacity. P.E.F.R. = Peak expiratory flow rate.

parisons at one dose level. The doses chosen were those which appeared from the literature to be near-equipotent or to be equitoxic in terms of respiratory depression, or which were those in which the drug is at least widely used clinically. The newer preparations—fentanyl (0.1 and 0.2 mg), phenoperidine (1 and 2 mg), and pentazocine (20 and 40 mg)—were each studied at two dose levels, as there was little evidence in the literature at that time of their relative potencies. Morphine was also included at doses of 10 and 15 mg, as both of these are frequently used in clinical practice.

Results

The findings are presented in Table III. Pethidine 100 mg and saline were selected as standard and placebo treatments respectively and the remaining agents classified as being superior to pethidine 100 mg, equivalent to pethidine 100 mg, or inferior to pethidine 100 mg, and superior to saline or equivalent to saline.

All drugs were better than saline (minimum value $P < 0.025$) but only levorphanol 2 mg proved significantly superior to pethidine 100 mg ($P < 0.05$).

The commonly used doses of these drugs placed in rank order with respect to incidence of successful postoperative pain relief according to the criteria described above are given in Table IV. This table ignores differences of less than 5%.

Discussion

The objective of this investigation was the selection of the most successful out of the many currently available analgesic drugs. The difficulty of making such a choice on the basis of existing literature has already been mentioned, arising from differences in methodology. The logical approach was the inclusion of all of the agents in question in one trial carried out under standard conditions. This imposed limits on the number of patients receiving each drug and consequently reduced the chance of demonstrating statistically significant differences between treatments. Nevertheless, it also implied that such distinctions as were found would probably be of clinical as opposed to purely statistical importance.

It must also, of course, be remembered that analgesic potency represents only one aspect of a drug's performance; the incidence and severity of side effects being at least equally

important in determining its clinical value, and these have already been reported in respect of the drugs included in this trial (Loan *et al.*, 1969; Morrison *et al.*, 1969; Dundee *et al.*, 1970).

Before discussing individual drugs it is worth considering the rank order of efficacy of the analgesics studied (Table IV) compared with a similar grading of efficacy based solely on the patient's estimation of pain relief (Table V). This distinguishes less clearly between the most effective drugs but suggests that either respiratory depression or oversedation placed pethidine and pethilorfan in a less favourable position (Table IV). We were surprised to find that the very popular pethidine showed up so poorly in the evaluation shown in Table IV compared with that in Table V. This was due to the fact that patients receiving it were so heavily sedated that they were unable to perform the tests of respiratory function. (The method of assessment in Table IV is, of course, designed to favour agents producing freedom from pain with minimum impairment of consciousness.) It is also interesting to note that while morphine 10 mg and pethidine 100 mg will produce about the same incidence of drowsiness in patients before operation (Dundee *et al.*, 1965) unpublished findings have shown that when given postoperatively pethidine 100 mg was associated with a pronounced depression of consciousness in 55% of cases as compared with 37% in the case of morphine 10 mg. This tendency for pethidine to cause cerebral depression in the postoperative period has previously been commented on by Bromage (1955) and Masson (1962).

Levorphanol, at a dose of 2 mg, clearly emerged as the most effective agent according to the criteria adopted in this trial. It has previously been shown to be a powerful analgesic (Glazebrook, 1952; Hunt and Foldes, 1953) with a lower incidence of drowsiness than morphine (Brown, 1954), and might be expected to provide the pain relief with minimal depression of consciousness sought in this trial.

Oxycodone 10 mg, pentazocine 20 mg, and the morphine 10 mg and cyclizine 50 mg (Cyclimorph 10) combination were the most successful of the remaining drugs though not significantly better than the standard pethidine 100 mg.

There is little published work on the analgesic effectiveness of oxycodone but it has been suggested that the pectinate has a potency similar to that of morphine (Eddy *et al.*, 1957; Brittain, 1959), and this relationship has been to some extent confirmed in this study. In a study of side effects of various analgesics we found this to be fairly non-toxic (Dundee *et al.*, 1970).

Pentazocine 20 mg compared favourably with most of the established potent analgesics, confirming reports of its efficacy against postoperative pain (Keats and Telford, 1964; Conaghan *et al.*, 1966). While early claims that this drug was totally free from addiction potential have not been fully supported the magnitude of this risk can be judged from the fact that, after some four years of widespread use in Britain, it is not subject to control under the dangerous drugs legislation.

Diamorphine 5 mg did not in this trial display any unique properties, and the results tend to support the view of Lasagna (1964) of the essential similarity between diamorphine and morphine. It must be remembered, however, that diamorphine in solution is quite rapidly broken down to monacetylmorphine and morphine (Storey, 1944), and that where a necessarily prolonged study is carried out under blind conditions it is impossible to be certain that the agent used is in all cases pure. In clinical practice, of course, where the identity of the drug need not be concealed, the current widespread use of solutions freshly prepared from diamorphine powder should overcome this difficulty.

Studies in animals suggested that the analgesic effectiveness of fentanyl in terms of milligramme potency would be 200-400 times that of morphine. The results in this trial, however, where fentanyl 0.2 mg proved equianalgesic to morphine 10 mg, would suggest that milligramme potency of fentanyl is of the order of 50 times that of morphine. This potency

ratio is in keeping with the respiratory depressant ratio found by Stephen (1967)—namely, that 0.15 mg of fentanyl produced a similar degree of respiratory depression to that of morphine 10 mg. Finch and de Kornfeld (1967) reported comparable results from a trial in patients with acute pain—namely, that the analgesic potency of fentanyl 0.2 mg was similar to that of morphine 10 mg. This drug serves once again to show the irrelevancy of milligramme potency in the pharmacology of an analgesic drug. The fact that by weight fentanyl is 50 times as potent as morphine, or 500 times as potent as pethidine, is of little value when, at acceptable levels of toxicity, it is no more effective in relief of pain and is appreciably more toxic (Dundee *et al.*, 1970).

As in the case of fentanyl there is a wealth of data from animal experiments on which to base an estimate of comparative analgesic performance ratio for phenoperidine and morphine. In milligramme potency this ratio would appear to assess phenoperidine as being from 15 to 75 times as potent as morphine. Again, however, the present studies suggest that these ratios are not applicable to its analgesic effectiveness in man and indicate that the truer ratio lies in the region of 5:1. Search of the literature has failed to produce any other report of the analgesic effectiveness of this drug.

While classified in this paper as a single agent, papaveretum is, of course, a mixture of morphine with three opium alkaloids—codeine, narcotine, and papaverine. Papaveretum 20 mg contains, in fact, morphine alkaloid equivalent to that contained in morphine sulphate 13.3 mg (Loan *et al.*, 1966). Many unconfirmed claims have been made regarding the superior analgesic potency of papaveretum as opposed to morphine alone (Gray, 1911; Schall, 1917), but the findings of this study do not support this view.

Four drugs were studied at more than one dose level—namely, morphine at 10 and 15 mg, pentazocine at 20 and 40 mg, fentanyl at 0.1 and 0.2 mg, and phenoperidine at 1 and 2 mg. It is noteworthy that in the cases of phenoperidine and fentanyl the difference in analgesic potency between a clinical dose and one considerably lower was reflected in improved performance, whereas with morphine and pentazocine an increase in dosage above the usual levels produced, if anything, a slight reduction in the desired effect. This supports the concept of a non-linear dose response curve for analgesic drugs and the importance of selecting the optimum dose of each agent.

Four analgesics were studied in combination with other drugs—namely, phenoperidine 2 mg and fentanyl 0.2 mg with dehydrobenzperidol 5 mg, morphine 10 mg with cyclizine 50 mg, and pethidine 100 mg with levallorphan 1.25 mg (Pethilorphan 100). Only in the case of the morphine-cyclizine preparation was there any appreciable improvement in analgesic potency, though this did not reach levels of statistical significance. In the absence of any indication that cyclizine has inherent analgesic properties it is possible that reduction in side effects might have resulted in greater patient comfort. The addition of dehydrobenzperidol 5 mg to fentanyl 0.2 mg had minimal effect on analgesic potency. There is some difference of opinion in the literature regarding the analgesic status of this combination of drugs. Foldes *et al.* (1964) and Fox *et al.* (1967) have suggested that dehydrobenzperidol either has some analgesic effect itself or is capable of enhancing the effects of the opiate analgesics. In contrast, Morrison (1970) found that droperidol given alone increased the subject's sensitivity to experimentally-induced pain, and, when given in the combinations commonly employed in neuroleptic anaesthesia it appeared to reduce the analgesic action of phenoperidine and fentanyl. With the number of subjects in each treatment group in this trial, however, it is likely that a clinically significant difference in potency would be shown by a statistically significant difference.

The combination of levallorphan with pethidine was associated with a fall in analgesic potency of the latter, and this

mixture has little to commend it in clinical practice (Telford and Keats, 1961; Campbell *et al.*, 1965).

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MEDICAL MEMORANDA

Cervical and Mediastinal Fibrosis Presenting with Thyroid Swelling

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A woman presenting with a thyroid swelling was found to be suffering from cervical and mediastinal fibrosis. The aetiology and association with other conditions are discussed.

Case History

Twenty years before the present admission the patient had been investigated for thyrotoxicosis, and a small hard thyroid swelling had been noted. Now 52 years old she complained of hoarseness of the voice, a feeling of pressure on the throat, and occasional dysphagia and dyspnoea. On examination she was found to have a large non-toxic nodular goitre with retrosternal extension confirmed by X-ray examination. Results of blood tests, including protein-bound iodine and thyroid antibody tests, were all normal, and thyroid scan showed only two small areas of functioning tissue. Indirect laryngoscopy showed a paralysed right vocal cord.

At operation on 13 March 1969 initial appearances suggested thyroid carcinoma, but it was found that the right lobe could be enucleated from a bed of dense fibrous tissue which extended into the mediastinum. Most of this and the left lobe was removed and the resultant fixed-walled cavity drained. Histological examination showed a simple goitre in which the capsular tissues were infiltrated with inflammatory cells including many plasma cells, and there were some similar scattered deposits in the gland. Lymph nodes showed reactive hyperplasia only.

One month after discharge the patient complained of increasing dyspnoea, and examination showed venous congestion of the right side of the neck, and mild ptosis of the right lid. A hard mass could be felt extending from neck to sternum. Tomograms showed a large swelling extending into the mediastinum and compressing the trachea, but the barium swallow was normal. Sedimentation rate was raised from 13 to 57 mm, there was an increase in the

gammaglobulin protein fraction, but no gastric or thyroid autoantibodies could be detected.

At bronchoscopy the right vocal cord was still paralysed and the trachea was rigid and compressed from the right side, the mucosa being congested down to the carina, beyond which it was normal. At thoracotomy the apex of the lung was found to be adherent to a hard pale mass extending from the thoracic inlet and engulfing the superior vena cava and lung surface. A line of cleavage was made between lung and mediastinum, further surgery being considered too hazardous. Histological examination showed dense fibrous tissue with prominent vessels and patchy inflammatory cell infiltration. As no thyroid tissue was present a diagnosis of mediastinal fibrosis was made.

Postoperatively the patient was started on prednisolone 30 mg daily with some improvement in venous pressure in the neck. Over the following year she remained in reasonable health with only slight dyspnoea and hoarseness. When last seen venous pressure was only slightly raised and she was taking prednisolone 15 mg daily.

Comment

Mediastinal fibrosis associated with and antedating retroperitoneal fibrosis has been reported (Tubbs, 1946; Calne *et al.*, 1966) but cervical fibrosis has not been mentioned. The present patient had no clinical evidence of the retroperitoneal condition. Renal function was normal but intravenous pyelography showed minor clubbing of the calices on the right. At hysterectomy two years previously no abnormal retroperitoneal tissues had been observed. There was also no history of methysergide therapy.

The simple goitre was probably incidental to the fibrosis, but as the microscopical appearances of mediastinal fibrosis have been compared with those of Riedel's thyroiditis and other fibrosing conditions (Barrett, 1958) and a common inflammatory aetiology has been suggested I wonder if this is a way along which Riedel's thyroiditis could have developed.

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