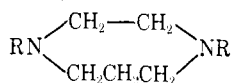


TABLE I
SYMMETRICAL N,N'-DIALKYLHOMOPIPERAZINES



R	Yield, ^a %	Bp (mm) or mp, °C	Formula	Analyses
<i>n</i> -C ₁₀ H ₂₁		150-152 (3)	C ₂₃ H ₃₂ N ₂	C, H, N
<i>n</i> -C ₁₂ H ₂₅	34	10	C ₂₉ H ₄₀ N ₂	H, N; C ^b
<i>n</i> -C ₁₄ H ₂₉	45	32-33	C ₃₃ H ₄₈ N ₂	C, H, N
<i>n</i> -C ₁₆ H ₃₃	72	41-42	C ₃₇ H ₇₆ N ₂	C, H, N
<i>n</i> -C ₁₈ H ₃₇	74	47-48	C ₄₁ H ₈₄ N ₂	C, H, N

^a Yield calculated after one recrystallization. ^b C: calcd, 79.74; found, 79.04.

0.11 mole of the alkyl bromide in 50 ml of EtOH for 3 hr. After cooling, the resultant salt was filtered and washed (cold EtOH). The salt was suspended in 50% EtOH and a slight excess of 20% KOH solution was added. Upon warming, the free amine separated as a colorless oil, which was separated by extracting three times with C₆H₆. The C₆H₆ extracts were distilled *in vacuo* to remove EtOH and H₂O. The residue was taken up in warm C₆H₆ and filtered through hard paper and the solvent again was removed *in vacuo*. The residue was taken up in Me₂CO and recrystallized from this solvent. The dialkylhomopiperazines are white, waxy solids or colorless oils, insoluble in H₂O and only slightly soluble in EtOH and EtOAc. The yields obtained varied from 35 to 75%.

N,N'-Dialkyl-N,N'-dimethylhomopiperazinium dimethosulfates (Table II) were prepared by dissolving 0.0025 mole of the dialkylhomopiperazine in EtOH (10 ml), adding 0.006 mole of

TABLE II
SYMMETRICAL N,N'-DIALKYL-N,N'-DIMETHYLHOMOPIPERAZINIUM
DIMETHOSULFATES AND DIMETHIOLIDES

R	Dimethosulfates		Dimethioides	
	Yield, ^a %	Mp, °C dec	Formula ^b	Mp, °C Formula ^b
<i>n</i> -C ₁₀ H ₂₁	55	260-262	C ₂₉ H ₆₄ N ₂ O ₈ S ₂ ^c	214-216 C ₂₇ H ₃₈ I ₂ N ₂
<i>n</i> -C ₁₂ H ₂₅	69	263	C ₃₃ H ₇₂ N ₂ O ₈ S ₂	206-207 C ₃₁ H ₆₆ I ₂ N ₂
<i>n</i> -C ₁₄ H ₂₉	83	258-259	C ₃₇ H ₈₀ N ₂ O ₈ S ₂	214 C ₃₅ H ₇₄ I ₂ N ₂
<i>n</i> -C ₁₆ H ₃₃	88	261-265	C ₄₁ H ₈₈ N ₂ O ₈ S ₂	218-222 C ₃₉ H ₈₂ I ₂ N ₂
<i>n</i> -C ₁₈ H ₃₇	86	250-252	C ₄₅ H ₉₆ N ₂ O ₈ S ₂	195-197 C ₄₃ H ₉₀ I ₂ N ₂

^a Yield calculated after one recrystallization. ^b All compounds were analyzed for C, H, N. ^c C: calcd, 54.68; found, 54.18.

redistilled Me₂SO₄, and refluxing for 3 hr. The reaction mixture was chilled and filtered and the product was recrystallized from EtOH and dried *in vacuo*. The yields obtained varied from 55 to 88%. The dimethosulfates are white, waxy solids insoluble in Et₂O and only slightly soluble in cold EtOH and EtOAc.

N,N'-Dialkyl-N,N'-dimethylhomopiperazinium dimethioides (Table II) were prepared by dissolving 0.0025 mole of the dialkylhomopiperazine in EtOH (10 ml), adding 0.0075 mole of MeI, and refluxing for 3 hr. The product was filtered from the chilled reaction mixture, recrystallized twice from EtOH, and dried *in vacuo*. These bis-quaternary salts are light yellow in color, insoluble in H₂O, and only slightly soluble in EtOH, Me₂CO, and EtOAc. The yields obtained were nearly quantitative.

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Book Reviews

Medical Research: A Series of Monographs. Volume 2. Drugs Affecting the Central Nervous System. Edited by ALFRED BURGER. Marcel Dekker, Inc., New York, N. Y. 1968. xv + 437 pp. 16 × 24 cm. \$19.75.

This is the second volume of a series dealing with the subject of neuropharmacology. The first volume considered those families of drugs acting specifically upon the peripheral nervous system; this present volume surveys those drugs which act directly upon the central nervous system. In keeping with the style of the first volume, the reviews and monographs collected represent discussions of fundamental rather than of applied pharmacology. The eight chapters of this volume range from general and comprehensive reviews of classes of central nervous system drug effects, to the specific and detailed minutiae of narrow chemical systems having CNS action.

At the general review end of the spectrum, there is a compact and complete chapter on the subject of narcosis and anesthesia by Larsen, Van Dyke, and Chenoweth of The Dow Chemical Company, wherein the treatment of a diverse group of chemicals is unified through the consideration of common mechanisms. Domino, Hudson, and Zografis discuss the frequently reviewed field of phenothiazines from the point of view of biochemical and pharmacological modes of action. A compilation of the many chemical types that may be grouped under the title psychotomimetic agents has been made by Hofmann, and both their medical and their paramedical uses are discussed. A broad chemical review by Donahoe and Kimura compares the wide range of families of agents effecting skeletal muscle relaxation through central nervous system action. This chapter serves as a useful companion piece to the chapter in Volume 1 that dis-

cusses similar pharmacological consequences through action in the area of the myoneural junction. A general review of the diverse types of drugs effective in the treatment of mental depression has been assembled by Biel.

The remaining chapters deal with minor topics in fine detail. Sternbach, Randall, Banziger, and Lehr of Hoffmann-La Roche present a thorough structure-activity relationship study of the 1,4-benzodiazepine analogs of Librium and Valium; however, their conclusions derive from animal screening data, and there is no extrapolation to human effectiveness. Similarly, Janssen and Van der Eycken present an overwhelming amount of detail, with archival thoroughness, on a series of compounds chemically related to, and pharmacologically more potent than, morphine. Lastly, Abood reviews the family of atropine-like glycolate esters, with special emphasis on the interrelationship between their structures and their mechanism of action. Issue may be taken with the employment of the term "psychotomimetic" in his title of this review of highly active anticholinergic drugs, for this usage conflicts with the generally accepted definition given by Hofmann.

All in all, this volume is an excellent reference work and will be of value to all scientists in this field. The printing is clear, the chemical formulas are free from typographical error to an unprecedented degree, and the subject and author indexes are extensive and complete. The appearance of the next volume of this set, concerning pharmacological testing methods, should complete a trilogy indispensable to anyone active in this area of research.

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