

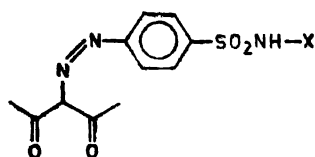
TABLE 2—3,5-DIMETHYL-4-(SUBSTITUTED SULFONAMIDO-BENZENE AZO, 4-SULFOPHENYL AZO AND 4-SULFONAPHTHYL AZO)PYRAZOLE

Sl. no.	X	R	M.p. °C
1	HOOCCH <sub>2</sub> -	R <sub>a</sub>	217
2	H <sub>3</sub> C <sub>2</sub> -	R <sub>a</sub>	194
3	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	R <sub>a</sub>	206
4	HOOCCH <sub>2</sub> -	R <sub>b</sub>	103
5	H <sub>3</sub> C <sub>2</sub> -	R <sub>b</sub>	160
6	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	R <sub>b</sub>	111
7	HOOCCH <sub>2</sub> -	R <sub>c</sub>	280
8	H <sub>3</sub> C <sub>2</sub> -	R <sub>c</sub>	166
9	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	R <sub>c</sub>	222
10	H <sub>3</sub> C <sub>2</sub> -	R <sub>d</sub>	166
11	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	R <sub>d</sub>	211
12	4-SO <sub>3</sub> H-C <sub>6</sub> H <sub>4</sub> -	R <sub>a</sub>	292
13	4-SO <sub>3</sub> H-C <sub>10</sub> H <sub>6</sub> -	R <sub>a</sub>	135
14	4-SO <sub>3</sub> H-C <sub>6</sub> H <sub>4</sub> -	R <sub>b</sub>	267
15	4-SO <sub>3</sub> H-C <sub>10</sub> H <sub>6</sub> -	R <sub>b</sub>	171
16	4-SO <sub>3</sub> H-C <sub>6</sub> H <sub>4</sub> -	R <sub>c</sub>	205
17	4-SO <sub>3</sub> H-C <sub>10</sub> H <sub>6</sub> -	R <sub>c</sub>	272
18	4-SO <sub>3</sub> H-C <sub>10</sub> H <sub>6</sub> -	R <sub>d</sub>	286

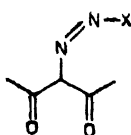
ture<sup>5</sup>. Azo dicarbonyl compounds were treated with the appropriate carbhydrazides using the procedure mentioned above, to obtain hydrazones (Table 1) and azo pyrazoles (Table 2), respectively.

**Screening for antibacterial activity:** The compounds in Table 2 have been evaluated for their antibacterial activity. The test organisms employed were *Bacillus subtilis*, *Staphylococcus aureus*, *Salmonella typhi*, *Escherichia coli*, and *Pseudomonas aeruginosa*. The antibacterial activity of the compounds were tested by ditch plate technique<sup>6</sup>, using the concentration levels 2 mg and 3 mg per ml.

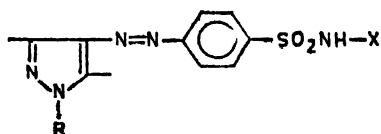
Compounds 1, 3 and 4 showed complete inhibition of *B. subtilis*. Compounds 12 and 16 showed complete inhibition of *S. aureus*. Compounds 12 and 16 showed complete inhibition of *S. typhi*, whereas compound 18 showed only partial inhibition. Compounds 14 and 16 showed complete inhibition of *P. aeruginosa*. The other compounds showed no activity on the organisms under study. None of the compounds studied showed any activity against *E. coli*. These results were identical for both the concentrations of the compounds studied.



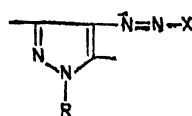
(Ia)



(Ib)

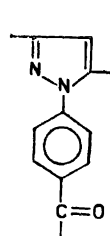
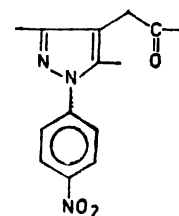
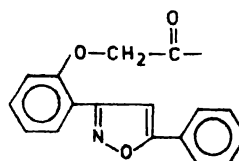
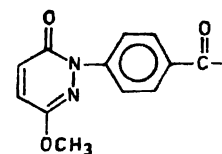


(IIa)

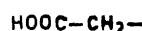
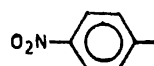


(IIb)

Substituents R = R<sub>a</sub>, R<sub>b</sub>, R<sub>c</sub>, R<sub>d</sub>.

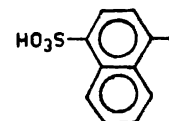
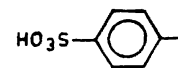
(R<sub>a</sub>)(R<sub>b</sub>)(R<sub>c</sub>)(R<sub>d</sub>)

X =



(Ia, IIa)

X =



(Ib, IIb)

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#### Reaction of Nitrosyl Chloride with Steroidal Olefins

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NITROSYL chloride gas was used for chloronitrosation of steroidal alcohols<sup>1</sup> and oximes<sup>2,3</sup>. The work of Tanabe *et al.*<sup>4</sup> which dealt with the