Individual psychophysical functions for 28 odorants*

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Individual scales of odor intensity were obtained for 28 different chemical compounds by the method of magnitude estimation. Eleven Ss participated in an experiment with 196 olfactory stimuli which differed in both quality and intensity. It was found (1) that power functions described the relationship between partial vapor pressure of the odorants and their subjective odor intensity for all Ss, (2) that all exponents were less than one but varied greatly between Ss, (3) that consistent intraindividual differences in the exponents of different odorants exist, and (4) that these are attributable to perceptual differences rather than to response bias.

Early psychophysical studies of olfaction were concerned primarily with the problem of odor quality, but, with the advent of direct psychophysical methods, more attention has recently been paid to the relationship between stimulus intensity and the corresponding subjective intensity. It has thus been shown that the perceived intensity is a negatively accelerated function of physical intensity. The relationship is practically always described by a power function $R = cS^n$, where the value of n is less than one. The exponents obtained in different studies have been found to range between 0.13 for octanol and 0.72 for eugenol.

In an early study of odor intensity with the use of a magnitude-estimation method, an exponent of 0.2 was obtained for benzaldehyde in air by Reese and Stevens (see Stevens, 1957, p. 166). Jones (1958a, b) studied systematically nine different organic compounds, which were presented to Ss in test tubes containing the odorants in liquid form, and he found that the exponents of the power function ranged from 0.42 for *n*-butanol to 0.58 for pyridine. Exponents of approximately the same magnitude were also obtained by Reese and Stevens (1960) for the odors of heptane and coffee diluted in air. In a

number of experiments with liquid odorants by Engen and his associates, the exponent of the power function was found to range from a value as low as 0.13 for octanol to 0.57 for amyl acetate (Engen, 1961, 1965; Engen & Lindström, 1963; Engen, Cain, & Rovee, 1968; Cain & Engen, 1970; see also Cain, 1969). Engen (1961) has pointed out that occasional deviations observed from the power function may be due to changes in perceived quality that sometimes accompany changes in the intensity of an odorant.

Results of 10 investigations are summarized in Table 1. The table is based on the scaling of perceived odor intensity of different chemical compounds and serves as a review of direct scaling experiments in this area. It is difficult to make comparisons among the different studies since, unfortunately, chemical formulas, as well as information about the purity of the odorants, are usually insufficient or lacking. Another complication is the type of diluent (air, gas, or liquid) used. An interesting finding made by Cain (1969) was that air-diluted compounds tend to give higher exponents than do the same compounds presented in liquid form, although relative values of the exponents are independent of the method of dilution and presentation of the stimulus. Furthermore, comparisons of the exponents are difficult to make, because sometimes partial vapor pressure and sometimes concentration in volume percent is used as the physical measure of intensity. Obviously, the same values of the parameter of the power function cannot be expected for two physical scales that are not linearly related. However, as the molecular weight of the odorant approaches the molecular weight of the diluent, the relative values of the scales approach a linear relationship. This is the

reason that the exponent for octanol is almost the same in both units, but the exponent for propanol is not.

The subject of the present investigation is the individual differences in the exponent of the power function. A brief review of this problem is presented by Ekman, Hosman, Lindman, Ljungberg, and Åkesson (1968) for several psychophysical continua. An early study of individual functions was performed by Künnäpas (1958), who studied apparent length of lines and verified the power function for all his Ss. J. C. Stevens and Mack (1959) noted that, despite individual differences in the slopes of psychophysical functions, the power function described the individual as well as the group scales of apparent force obtained by both magnitude estimation and magnitude production (see also S. S. Stevens, 1960). Similar results were obtained for sensory continua such as loudness of 1,000-Hz tone (J. C. Stevens & Guirao, 1964), brightness of white light (Marks & J. C. Stevens, 1966), and saltiness of sodium chloride and sweetness of sucrose (Ekman & Åkesson, 1965). Among the scaling experiments concerned with olfactory intensity, one by Jones and Marcus (1961) has reported individual data. A median exponent of 0.56 ($Q_1 = 0.47$ and $Q_3 = 0.93$) was found for the odor of benzene.

In the present case, individual scales of perceived odor intensity were obtained for a large sample of odorants with a procedure permitting direct intraindividual comparisons of these scales. A magnitude-estimation method was used, with one standard as a reference for all variables independent of the intensity and quality.

THE EXPERIMENT

Stimuli

In the present experiment 28 different chemical compounds, the purest available commercially, were used as odorants. They were chosen with regard to both chemical and psychological characteristics according to five criteria: (1) None of the chemical compounds should exceed the rating 2 ("moderate") in Sax's (1966) system of toxic hazards regarding inhalation. (2) A large variation in molecular weight and vapor pressure (at 20°C) was desired, as well as a variation in chemical structure of the chemical compounds. (3) Different characteristics of solubility in polar and nonpolar substances were to be represented. (4) The different compounds should produce qualitatively different odors. (5) Both pleasant and unpleasant odors should be included in the stimulus sample.

Each of the 28 odorants was diluted into seven steps in a geometric series according

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to the formula $(100/2^n)\%$, where n ranged from 0 to 6, such that the concentrations ranged from 100% to 1.56% in equal log steps. The percentages represent milliliters of odorant per 4 ml of the solution. For three of the compounds, guaiacol, menthol, and phenyl acetic acid, which are crystals in pure form, the 100% solution was replaced by 50%, 40%, and 28%, respectively, the highest liquid concentrations available for these particular odorants. Diethyl phthalate was used as diluent, except for *n*-heptane, nonane, and *n*-octane, which were diluted in silicon oil. It was assumed that all solutions would deviate negligibly from ideal behavior when diluted in diethyl phthalate. Therefore, the partial vapor pressures of the odorous compounds in mm Hg at 20°C were calculated from the molefraction according to Raoult's law. With regard to *n*-heptane, direct vapor pressure measurements were performed on the solution of *n*-heptane in silicon oil. Since the same vapor pressure was found for *n*-heptane in silicon oil as in vacuum, it can be concluded that the silicon oil has a negligible vapor pressure. On the assumption that Raoult's law is applicable, the molecular weight of silicon oil was calculated from the data obtained from the solution of *n*-heptane in silicon oil. This value was then used in the calculations of the vapor pressure for nonane and *n*-octane diluted in silicon oil according to Raoult's law. All chemical compounds used in the experiment are presented in Table 2.

The experiment was performed in a well-ventilated room of 30 m³, with a fan evacuating 1.5 m³ of air per minute. The stimuli were presented by a technique used earlier by Engen (1965). The prepared solutions were kept at a temperature of ca 20°C in 15 x 100 mm glass tubes with a cork wrapped in aluminum foil. Attached to each cork was a glass rod with a piece of cotton wrapped around its lower end. The Ss sniffed the odors from the cotton, which was contained in the tube and partially immersed in the liquid odorant when not in use.

Procedure

The method of *magnitude estimation* with a modulus was used. A standard acetone at 100% concentration (vapor pressure of 175 mm Hg at 20°C) was presented before a block of seven variable stimuli. The Ss were asked to call this standard 100, to memorize its perceived intensity, and then to estimate the perceived intensity of subsequent stimulus variables in relation to it. The same standard was used throughout the

Table 1
Summary of Results Concerning Power Functions for Different Odorants

Odorant	Diluent	Scaling Method	Number of Ss	Exponent	Reference
Acetone	Air	Magnitude estimation	15	.71	Cain (1969)
Amyl acetate	Benzyl benzoate	Magnitude estimation	4	.33	Engen (1961)
Amyl acetate	Benzyl benzoate	Magnitude estimation	4	.20	Engen (1961)
Amyl acetate	Benzyl benzoate	Magnitude estimation	13	.49 ^c	Engen & Lindstrom (1963)
Amyl acetate	Benzyl benzoate	Ratio estimation	13	.42	Engen & Lindstrom (1963)
Amyl alcohol	Air	Magnitude estimation	15	.56	Cain (1969)
Benzaldehyde	Air	Magnitude estimation	30	.20	Stevens (1957) and personal communication
Benzene	Mineral oil	Magnitude estimation	36	.55	Jones (1958b)
Benzene	—	Magnitude estimation	52	.56 ^b	Jones & Marcus (1961)
<i>n</i> -Butanol ^a	Mineral oil	Magnitude estimation	36	.42	Jones (1958a)
<i>n</i> -Butanol	Diethyl phthalate	Magnitude estimation	12	.23	Cain (1969)
<i>n</i> -Butanol	Air	Magnitude estimation	15	.64	Cain (1969)
<i>sec</i> -Butanol ^a	Mineral oil	Magnitude estimation	36	.57	Jones (1958a)
<i>iso</i> -Butanol ^a	Mineral oil	Magnitude estimation	36	.56	Jones (1958a)
Coffee	Air	Magnitude estimation	12	.55	Reese & Stevens (1960)
Cyclohexane ^a	Mineral oil	Magnitude estimation	36	.44	Jones (1958a)
Dimethyl disulphide	Air	Finger span matching	8	.40	Berglund, Berglund, & Lindvall (In preparation) ¹
Dimethyl monosulphide	Air	Finger span matching	8	.41	Berglund, Berglund, & Lindvall (In preparation) ¹
Ethyl acetate ^a	Mineral oil	Magnitude estimation	36	.53	Jones (1958a)
Eugenol	Air	Magnitude estimation	20	.72	Mitchell & Gregson (1968)
Geraniol	Diethyl phthalate	Magnitude estimation	12	.20	Cain (1969)
Geraniol	Air	Magnitude estimation	15	.29	Cain (1969)
Heptane ^a	Mineral oil	Magnitude estimation	52	.55	Jones (1958b)
Heptane	Air	Magnitude estimation	18	.60	Reese & Stevens (1960)
Heptanol	Diethyl phthalate	Magnitude estimation	12	.15	Cain (1969)
Hexanol	Diethyl phthalate	Magnitude estimation	12	.16	Cain (1969)
Octane ^a	Mineral oil	Magnitude estimation	52	.55	Jones (1958a)
Octanol	Diethyl phthalate	Magnitude estimation	11	.15	Cain (1969)
Octanol	Diethyl phthalate	Magnitude estimation	11	.13	Engen (1965)
Propanol	Diethyl phthalate	Magnitude estimation	11	.42	Engen (1965)

Table 1 (Continued)

Odorant	Diluent	Scaling Method	Number of Ss	Exponent	Reference
Propanol	Diethyl phthalate	Magnitude estimation	12	.38	Cain (1969)
Propanol	Diethyl phthalate	Magnitude estimation	12	.36	Cain (1969)
Propanol	Diethyl phthalate	Magnitude estimation	12	.52	Cain (1969)
Propanol	Air	Magnitude estimation	15	.69	Cain (1969)
Pyridine ^a	Mineral oil	Magnitude estimation	36	.58	Jones (1958a)

^aVapor pressure is used as physical scale of intensity. ^bMedian of 52 individually obtained exponents.

^cMean of obtained exponents for different standards.

experiment. It was chosen to represent the upper end of the subjective intensity continuum for the whole set of 28 odorants in order to obtain estimates in submultiples, since some Ss have been found not to handle both multiples and fractions properly at the same time (Svenson & Åkesson, 1966; see also Stevens, 1956). Only part of the total set of 196 stimuli was presented in any one session in order to minimize adaptation and cross-adaptation effects due to extensive exposure to a 100% acetone standard.

The comparison odorants, varying in both quality and intensity, were presented in a different irregular order for each S, in which each individual stimulus appeared four times. The stimulus order was drawn at random but modified such that for each compound the lowest, the third lowest, the fifth lowest, and the highest concentration was placed next to the standard at least once in the whole random set. This restriction was made in order to balance out a possible change in the memory or state of adaptation due to the standard that might be present within the blocks.

Each of the 196 different stimuli was presented four times to each S in sessions comprising 70-77 comparison stimuli. Eleven such sessions, usually two each day, were run per S. A session lasted about 20 min, with intervals of 15 sec between stimuli and a rest of 20 min between sessions.

Subjects

Eleven Ss, five men and six women, participated in the experiment. All but E, G, H, and K were psychology students, but none of them had any previous experience in olfactory experiments.

RESULTS

The individual exponents of the odorants represent the main results of this study and are presented in Table 3. Individual scales of odor intensity for each of the 28 odorants were computed by

averaging the four estimates for each stimulus. A straight line representing a power function seems to fit the data fairly well in all 308 individual graphs, when these arithmetic means are plotted in log-log coordinates against the stimulus concentration measured as partial vapor pressure.² Of course, it may be possible to fit other types of mathematical functions to some of the data plots because of scatter of points, but there seem to be no systematic trends over individuals or odorants indicating that a different function would be better.

A comparison of the individual scales suggests that there is a positive correlation over Ss in the range of numbers assigned to the different odorants. This effect is even more evident in the results for all Ss and odorants. Whether this apparent correlation may be related to interindividual differences in olfactory sensitivity or ability in handling numbers cannot be decided on the basis of present data.

As indicated, some of the Ss apparently did not perceive any differences in intensity for some of the odorants, but with three exceptions this phenomenon is limited to certain odorants for certain Ss. That is, the results of all Ss for diethyl monosulphide, n-heptanal, and phenyl acetic acid vary little in perceived intensity within the stimulus range investigated. The stimulus range for these three odorants probably covers only an extremely small part of the possible dynamic range. It would be useful to determine whether the absolute thresholds for these odorants are low compared to the lowest intensity used in the present experiment.

Table 3 also shows that the size of the individual exponents of two of the Ss, D and I, deviate remarkably from those of the other Ss. With a few exceptions, the exponents for D and I are less than 0.10, and, in addition, their numerical estimates are generally high in magnitude relative to the standard. The very narrow numerical range used by these two high estimators

could be related to uncertainty and problems of discrimination, but, interestingly, there does not seem to be any obvious difference between them and the rest of the Ss as far as the ability to rank-order stimulus intensities is concerned.

In order to evaluate the interindividual differences in the relative size of exponents over the different odorants, a product-moment correlation was calculated on the rank orders of exponents for paired Ss because of presence of ties. In computing these coefficients, the empty cells of Table 3 were regarded as containing zero exponents. Despite the great number of tied ranks, these coefficients range from 0.12 to 0.65. Even S D (0.14-0.65) and S I (0.23-0.65), with their narrow response ranges, correlated as well with each other and the rest as did any of the other Ss.

In addition, Kendall's W was calculated for the whole group, and the value of 0.44 obtained corresponds to an average rank coefficient of 0.38 and is significant at the 0.1% level, according to a chi-square test (cf. Ferguson, 1959, p. 188). There is no indication that valuable information about the exponents for different odorants will get distorted or lost when the results are summarized in terms of group data.

The exponents of the average scales of perceived intensity obtained by computing the arithmetic means of the 11 individual scales for each of the 28 odorants are presented in the last column of Table 3. Straight lines fitted to these functions by eye seem to fit very well in all except two cases (diethyl monosulphide and n-heptane).² This indicates that the power function governs the relationship between perceived intensity and partial vapor pressure for different odorants. The arithmetic means of the individual exponents are also presented in Table 3. The exponents obtained through these two different procedures show a rank-order coefficient of correlation of 0.70. There is no evidence of any systematic difference between group functions and individual functions, even though the latter are based on only four estimates per stimulus. Finally, it should be noted that the exponents obtained for all the 28 odorants are less than one for all individuals.

DISCUSSION

Individual Functions

The finding that the psychophysical power function yields a satisfactory fit to nearly all individual data for the 28 different odorants is noteworthy, since published data on the psychophysical function for the individual Ss is limited, especially in the area of olfaction. This kind of information is particularly

Table 2
Chemical Compounds Used as Stimuli in the Experiment

Chemical compound	Purity*	Molecular weight	Density at 20° C	Vapor pressure in mm Hg at 20° C	Solubility in g/100 ml H ₂ O at ca. 20° C	Odor	Stimulus range (S _{max} /S _{min}) in vapor pressure
Acetone CH ₃ ·CO·CH ₃	pro analysi	58.08	0.792	175	∞	Mildly pungent, somewhat aromatic	24.33
n-Amyl acetate CH ₃ ·COO(CH ₂) ₄ ·CH ₃		130.19	0.879	3.27	0.18	Apples, bananas	48.09
Anethole CH ₃ ·CH=CH·C ₆ H ₄ ·O·CH ₃	purum	148.21	0.985	0.03	0	Anethole	49.18
Benzaldehyde C ₆ H ₅ ·CHO	puriss.	106.12	1.050	0.7	0.33	Bitter almonds	33.14
Benzyl acetate CH ₃ ·COOCH ₂ ·C ₆ H ₅	purum	150.18	1.057	0.07	0	Jasmine, pears	46.05
1-Butanol CH ₃ ·(CH ₂) ₃ ·CH ₂ OH	pro analysi	74.12	0.810	4.3	7.91	Weak fusel oil	30.11
n-Butyric acid CH ₃ ·CH ₂ ·CH ₂ ·COOH	puriss.	88.11	0.959	0.7	∞	Pungent, rancid	30.38
Diethyl monosulphide CH ₃ ·CH ₂ ·S·CH ₂ ·CH ₃	purum	90.18	0.837	48	0.313	Unpleasant	35.29
Diethyl phthalate C ₆ H ₄ (COOC ₂ H ₅) ₂	puriss.	222.24	1.123	0	0.7	None	—
Ethanol CH ₃ ·CH ₂ OH	99.5%	46.07	0.790	41	∞	Ethereal odor	19.84
Ethyl acetate CH ₃ ·COOC ₂ H ₅	pro analysi	88.11	0.901	71	8.6	Fruity	31.86
Ethyl methyl ketone C ₂ H ₅ ·CO·CH ₃	pro analysi	72.11	0.805	71.2	35.3	Acetone-like	29.48
Eugenol CH ₃ ·CH ₂ ·C ₆ H ₃ (O·CH ₃) ₂ ·OH	purum	164.21	1.066	7.1	0	Glove	50.02
Furfural O·CH=CH·C ₄ ·CHO	pro analysi	96.09	1.159	1.1	8.3	Almond odor	27.36
Geranial (CH ₃) ₂ C=CH·CH ₂ ·CH ₂ ·C(CH ₃)=CH·CHO	purum	152.24	0.887	0.03	0	Lemon, sweet rose	55.56
Guaiacol CH ₃ ·O·C ₆ H ₄ ·OH	puriss.	124.14	1.129	0.06	1.16	Agreeable aromatic	45.92
n-Heptanal CH ₃ ·(CH ₂) ₅ ·CHO	pro analysi	114.19	0.850	1.9	0	Penetrating fruity	43.76
n-Heptane CH ₃ ·(CH ₂) ₅ ·CH ₃	pro analysi	100.21	0.684	34	0.0052	Gasoline	19.30
1-Heptanol CH ₃ ·(CH ₂) ₅ ·CH ₂ OH	puriss.	116.21	0.822	0.15	0.09	—	46.01
Menthol CH ₃ ·CH·CH ₂ ·CH(OH)·CH(CH(CH ₃) ₂)·CH ₂ ·CH ₂	puriss.	156.27	0.890	0.003	0.04	Peppermint	60.94
Methyl salicylate HO·C ₆ H ₄ ·COOCH ₃	purum	152.15	1.184	0.009	0.074	Strong winter-green	41.86
Nonane CH ₃ ·CH ₂ ·CH ₂ ·CH ₂ ·CH ₂ ·CH ₂ ·CH ₂ ·CH ₂ ·CH ₃	pract.	128.25	0.718	3.22	0	Candied ginger	47.1
n-Octane CH ₃ ·(CH ₂) ₆ ·CH ₃	pract.	114.22	0.703	10	0.0015	—	17.4
1-Octanol CH ₃ ·(CH ₂) ₆ ·CH ₂ OH	puriss.	130.23	0.825	0.01	—	Penetrating aromatic	51.26
1-Pentanol CH ₃ ·(CH ₂) ₃ ·CH ₂ OH	pro analysi	88.15	0.817	2.8	0	Characteristic odor	35.32
Phenyl acetic acid C ₆ H ₅ ·CH ₂ ·COOH	puriss.	136.15	1.228	0.0001	1.66	—	52.63
2-Phenyl ethyl alcohol C ₆ H ₅ ·CH ₂ ·CH ₂ OH	purum	122.17	1.024	0.002	1.6	Faint rose	38.99
Pyridine N·CH=CH·CH=CH	pro analysi	79.10	0.982	14	∞	Disagreeable	26.64
Silicon oil DC 702		337.00	1.070	0	—	None	—
iso-Valeric acid (CH ₃) ₂ ·CH·CH ₂ ·COOH	puriss.	102.14	0.931	0.4	4.2	Disagreeable, rancid cheese	35.89

* Pro analysi is the highest purity grade for analytical work. Puriss. is of min 99% purity; purum is of min 98% purity, and pract. is of high but varying purity. All purity grades of the chemicals are guaranteed against containing any additional odor except the characteristic one. From *The Handbook of Chemistry & Physics*, 1969.

invaluable in comparing psychophysical functions in general. For example, the finding that intraindividual differences in the exponent for qualitatively different odorants are less than interindividual differences in the exponent for any particular odor might suggest the operation of nonsensory factors (Jones & Marcus, 1961).

Since characteristic exponents of the power functions are found to relate psychophysical magnitude to stimulus magnitude, the exponent may be described as "modality bound" (S. S. Stevens, 1961). Of course, individual differences in the exponent have been found for different continua as well as for qualitative variation within one sense modality (Ekman &

Sjöberg, 1965; Ekman, Hosman, & Lindström, 1965; Ekman & Åkesson, 1965; S. S. Stevens, 1968; Cain, 1969). Two possible interpretations of the interindividual variation in the exponent of the psychophysical function have been proposed; it may reflect *genuine perceptual differences* or some kind of *response bias* related to the S's conception of numerical

Table 3
Individual and Mean Exponents of the Power Functions for the 28 Odorants. Cells with a bar indicate that the S could not discriminate odor intensity within the range investigated for that particular odorant.

Odorant	Subject											Mean	SD	Range	Group Exponent
	A	B	C	D	E	F	G	H	I	J	K				
Acetone	0.94	0.63	0.59	0.12	0.79	0.40	0.49	1.02	0.14	0.73	0.13	0.54	0.32	0.12-1.02	.34
n-Amyl acetate	0.10	0.18	0.24	0.07	0.08	0.04	0.23	0.11	0.05	0.26	0.06	0.13	0.08	0.04-0.26	.11
Anethole	0.17	0.21	0.14	0.07	0.21	0.14	0.09	0.22	0.04	0.32	0.14	0.16	0.08	0.04-0.32	.15
Benzaldehyde	0.27	0.30	0.17	0.13	0.38	0.09	0.24	0.24	0.07	0.11	0.09	0.19	0.10	0.07-0.38	.16
Benzyl acetate	--	0.10	0.23	0.05	0.35	0.22	0.25	0.36	0.07	0.32	0.17	0.21	0.11	0.05-0.36	.15
l-Butanol	0.37	0.70	0.27	0.04	0.72	0.16	0.33	0.30	0.03	0.32	0.20	0.31	0.23	0.03-0.72	.22
n-Butyric acid	0.41	0.39	0.07	0.04	0.33	0.12	0.18	0.41	0.07	0.26	0.11	0.22	0.15	0.04-0.41	.16
Dimethyl monosulphide	--	--	0.16	--	0.07	0.05	--	0.03	--	--	--	(0.08)	(0.06)	(0.03-0.16)	--
Ethanol	--	0.40	0.23	0.11	1.00	0.19	--	0.27	0.09	--	0.12	0.30	0.30	0.09-1.00	.07
Ethyl acetate	0.43	0.33	0.07	0.08	0.29	0.17	0.19	0.25	0.10	0.27	0.10	0.21	0.12	0.07-0.43	.19
Ethyl methyl ketone	0.65	0.42	0.47	0.10	0.37	0.25	0.24	0.48	0.08	0.45	0.15	0.33	0.18	0.08-0.65	.27
Eugenol	0.29	0.34	0.19	0.08	0.69	0.22	0.21	0.21	0.09	0.45	0.15	0.27	0.18	0.08-0.69	.19
Furfurole	0.15	0.70	0.24	0.04	0.29	0.13	0.25	0.49	0.07	0.47	0.25	0.28	0.20	0.04-0.70	.23
Geranial	0.23	0.39	0.16	0.08	0.15	0.05	0.21	0.12	0.04	0.39	0.10	0.17	0.12	0.04-0.39	.13
Guaiacol	0.32	0.40	0.30	0.09	0.37	0.24	0.08	0.23	0.04	0.24	0.20	0.23	0.12	0.04-0.40	.17
n-Heptanal	--	--	--	--	0.05	--	--	0.18	--	0.06	--	(0.10)	(0.07)	(0.05-0.18)	--
n-Heptane	0.48	0.55	--	0.10	--	0.14	0.11	0.43	0.08	0.77	0.08	0.30	0.26	0.08-0.77	.19
l-Heptanol	0.12	0.24	0.20	0.04	0.28	0.10	--	0.19	--	0.31	0.04	0.71	0.10	0.04-0.31	.09
Menthol	0.36	0.64	0.13	--	--	0.21	0.15	0.31	0.04	0.20	--	0.26	0.19	0.04-0.64	.11
Methyl salicylate	0.28	0.31	0.18	0.04	0.26	0.25	--	0.15	0.04	0.28	0.16	0.20	0.10	0.04-0.31	.12
Nonane	0.27	0.73	0.08	0.04	0.44	0.13	0.22	0.38	--	0.48	0.24	0.30	0.21	0.04-0.73	.13
n-Octane	0.16	0.57	0.30	0.06	0.16	0.11	--	0.28	0.08	0.61	0.10	0.24	0.20	0.06-0.61	.15
l-Octanol	0.31	0.38	0.03	0.06	0.17	--	--	0.11	0.02	0.09	0.07	0.14	0.13	0.02-0.38	.09
l-Pentanol	--	0.23	0.15	0.07	0.50	0.07	0.26	0.28	0.05	0.26	0.14	0.20	0.14	0.05-0.50	.15
Penyl acetic acid	--	--	0.05	--	0.20	--	0.07	0.12	--	0.06	--	(0.10)	(0.06)	(0.05-0.20)	.08
2-Phenyl ethyl alcohol	--	0.35	--	0.03	0.08	0.08	0.16	0.36	--	0.23	0.21	0.19	0.12	0.03-0.36	.06
Pyridine	0.30	0.70	0.25	0.17	0.25	0.22	0.06	0.23	0.08	0.25	0.15	0.24	0.17	0.06-0.70	.27
iso-Valeric acid	0.21	0.29	0.24	0.04	0.22	0.19	0.12	0.33	0.10	0.30	0.22	0.21	0.09	0.04-0.33	.16

quantity. Jones and Marcus (1961) determined individual exponents and performed an analysis of variance for three sense modalities including odor, and they concluded that interindividual differences largely reflected differences in handling numbers. In later experiments with a large number of psychologically very different continua, Ekman et al (1968) found that correlations (ca 0.40) between subjective ranges were low compared to the reliability coefficients characteristic of the individual scales (ca 0.96). Although the present paper was not designed to deal directly with the issue, the data obtained are relevant. Of particular interest is the fact that the interindividual variability, with regard to the exponent in the present study (Table 3), is of the same order of magnitude as the variability found for other sensory continua (Ekman et al, 1968).

The Power Function for Different Odorants

The exponents obtained for the different compounds in the present study seem to reveal genuine perceptual differences. The experimental design permits the interpretation that the numerical response range used by the Ss is not limited to one particular odorant, which would be the case in a more conventional design when each odorant is scaled in separate experiments. The scaling of an odorant is performed in the context

of the whole set of odorants, and the tendency noted above for each S to use the same range of numbers in separate experiments is avoided (Jones & Marcus, 1961; Ekman et al, 1968).

It should be noted, though, that for the S there was presumably only one total range of perceptual intensity for the whole set of 196 stimuli, and, therefore, the physical ranges of individual odorants scaled could not logically have affected the specific response range chosen for each odorant. As a result, it can be assumed that the variation in the exponents is characteristic of the odorants rather than of response bias. The relatively high rank-order correlations obtained for pairs of Ss and the fact that the variation in stimulus range (S_{max}/S_{min}) in vapor pressure (Table 2) accounts for less than 25% of the total variance also support this interpretation.

For smaller samples of odorants, the relative value of the exponents have been shown to correspond to the physical property of water solubility. Cain (1969) has obtained high positive rank-order coefficients of correlation between these variables, both with regard to his own data and to data from Jones (1958a). That finding is not supported on the basis of the present larger sample of odorants (Table 2). The rank-order coefficient of correlation is only +0.27 and +0.25 for group exponents and averaged individual exponents, respectively.

Some of the present odorants have also been scaled in earlier studies, and the results are not always in agreement. However, it should be borne in mind that the comparison of group results from small samples of individuals is limited, owing to the relatively large individual differences in the exponent. For example, the exponents obtained earlier for amyl acetate (e.g., Cain & Engen, 1970) are higher than our group exponents. On the other hand, exponents obtained for other odorants, for example, acetone, butanol, and octanol, agree with those obtained here, despite differences in experimental design (e.g., Cain, 1969). The task of the Ss in the present experiment must be more difficult than that in the typical scaling experiment, for they work relatively long hours and, therefore, motivational effects may be more likely to play a role (S. S. Stevens & Greenbaum, 1966). Finally, effects of adaptation and cross-adaptation are often expected in studies of olfaction, but they probably did not affect the present results. If they had, the exponents should have been higher rather than lower than those obtained in other studies, because the effect of adaptation seems to increase the exponent in olfaction (Cain & Engen, 1970) as in other modalities (J. C. Stevens & S. S. Stevens, 1963).

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NOTES

1. Berglund, B., Berglund, U., & Lindvall, T. Perceived odor intensity of two organic sulfides and mixtures thereof. In preparation.
2. Graphs showing both individual and average psychophysical functions are available in a preliminary report of the data and may be obtained by writing the authors at Psychological Laboratories, University of Stockholm, Box 6801, S-113 86 Stockholm, Sweden.

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