



A · M · E · R · I · C · A · N  
A N T H R O P O L O G I C A L  
A S S O C I A T I O N

---

Folk Zoological Life-Forms: Their Universality and Growth

Author(s): Cecil H. Brown

Reviewed work(s):

Source: *American Anthropologist*, New Series, Vol. 81, No. 4 (Dec., 1979), pp. 791-817

Published by: [Wiley-Blackwell](#) on behalf of the [American Anthropological Association](#)

Stable URL: <http://www.jstor.org/stable/676586>

Accessed: 14/11/2012 18:51

---

Your use of the JSTOR archive indicates your acceptance of the Terms & Conditions of Use, available at <http://www.jstor.org/page/info/about/policies/terms.jsp>

JSTOR is a not-for-profit service that helps scholars, researchers, and students discover, use, and build upon a wide range of content in a trusted digital archive. We use information technology and tools to increase productivity and facilitate new forms of scholarship. For more information about JSTOR, please contact [support@jstor.org](mailto:support@jstor.org).



Wiley-Blackwell and American Anthropological Association are collaborating with JSTOR to digitize, preserve and extend access to *American Anthropologist*.

<http://www.jstor.org>

# Folk Zoological Life-Forms: Their Universality and Growth

CECIL H. BROWN  
Northern Illinois University

*Folk zoological life-form terms, like folk botanical life-form terms (Brown 1977a), are added to languages in a highly regular manner. Life-forms of the triad FISH, BIRD, and SNAKE are lexically encoded first, although in no particular order, followed by WUG (e.g., American English bug) and then MAMMAL. Four general principles of naming-behavior underlie these regularities: (1) criteria clustering; (2) conjunctivity (including binary opposition); (3) dimension salience; and (4) marking. In addition, size of folk zoological life-form vocabularies is positively correlated with societal complexity. This is caused by the decay of folk biological taxonomies as societies become more complex. [cognitive anthropology, ethnobiology, folk classification, language universals, language change]*

THIS IS THE SECOND of two papers treating the universality and growth of folk biological life-forms. The first paper showed that five botanical life-forms, "tree," "grerb" (small herbaceous plant), "vine," "bush," and "grass," are added to languages in a highly regular manner (Brown 1977a). These uniformities were explained through reference to general principles of naming-behavior. The present study similarly shows that five folk zoological life-forms are lexically encoded in a highly regular order. General principles of naming-behavior also underlie zoological life-form encoding.

The order in which biological life-forms are added to languages is inferred from synchronic implicational universals. For example, if a language has a label for "grerb," it will also have a label for "tree," although not necessarily vice versa. Thus the occurrence of a "grerb" term universally implies the occurrence of a "tree" term (Brown 1977a). Similarly, if a language has a word for "mammal," it will also have a word for "wug" (e.g., American English *bug*) although not necessarily vice versa. Thus "mammal" universally implies "wug." For such relationships to have been realized in fact, life-form

CECIL H. BROWN's research interests lie primarily in cognitive anthropology and linguistics. His published works in these fields include articles on kinship semantics, ethnobiology, ethnoanatomy, color classification, language universals, language change, and the historical/comparative linguistics of Mesoamerican languages. He has also written a book, *Wittgensteinian Linguistics* (1974), dealing with overlapping issues in contemporary linguistics and philosophy. Following fieldwork among Huastec Mayan speakers of northern Veracruz, Mexico, in 1971 he earned a Ph.D. in anthropology from Tulane University. Brown is currently an associate professor of anthropology at Northern Illinois University where he has been since 1970. His principal nonacademic interest is warm weather.



Copyright © 1979 by the American Anthropological Association  
0002-7294/79/040791-27\$3.20/1



criteria for affiliation with that rank nor are they always particularly large and heterogeneous. These criteria, however, served as an important guide in an initial phase of research involving a survey of published and unpublished reports of firsthand investigation of native zoological taxonomies (see discussion of Type A language cases below). The principal finding of the survey is that among those zoological classes in languages that are unambiguously identified as life-forms by Berlin's nomenclatural criteria, five categories consistently occur: FISH, BIRD, SNAKE, WUG, and MAMMAL. No other zoological life-forms are as pervasive as these five. The critical features associated with these regularly occurring folk zoological life-form taxa are as follows:

- FISH** Creature possessing fins, gills, and a streamlined body, adapted to an aquatic environment. (This life-form is occasionally extended to other aquatic animals lacking some or all of these features, e.g., whales, aquatic crustaceans. In such cases true fish usually constitute the focal members of the class [cf. Hunn 1977:250]. )
- BIRD** Creature possessing feathers, wings, and a bill or beak, adapted to flying. (This life-form is occasionally extended to bats.)
- SNAKE** Featherless, furless, elongated creature adapted to crawling, usually lacking appendages. (This life-form in its greatest extension includes worms, snakes, lizards, and, occasionally, other elongated creatures such as reptilelike insects.)
- WUG** Small creature usually other than those included in FISH, BIRD, and SNAKE. (This life-form always encompasses bugs, i.e., insects and other very small creatures such as spiders, and frequently is extended to worms. Occasionally, the category also includes other small creatures such as snails and crabs and, in addition, creatures such as lizards, tortoises, and frogs if these are small.)
- MAMMAL** Large creature other than those included in FISH, BIRD, and SNAKE. (Typically this life-form is restricted to mammals. Occasionally, however, it is extended to other large animals such as iguanas and crocodiles and, in addition, to such creatures as tortoises and frogs if these are large.) (Note: *Animal* is used more commonly than *mammal* as a name for this life-form by speakers of American English. Since *animal* is used also as a unique beginner to refer to creatures in general, to avoid ambiguity of reference, it is not employed as a life-form gloss here.)

The five classes defined above stand apart from most other categories meeting Berlin's life-form criteria in that inclusion within them is based solely on *the form of the whole animal* or, in other words, on gross morphology. Membership in nonpervasive or nonuniversal life-form categories is frequently found to be based on criteria other than gross morphology. These include animal habitat (e.g., house vs. forest), edibility (e.g., poisonous vs. nonpoisonous, tabooed vs. nontabooed), feeding habits (e.g., meat-eating vs. plant-eating vs. insect-eating), relationship to human beings (e.g., dangerous vs. harmless, domesticated vs. wild), locomotion (e.g., flying vs. crawling vs. trotting vs. burrowing), occurrence in environment (e.g., common vs. rare), and so on. The finding that universal zoological life-forms are based on gross morphology is identical to the finding for universal folk botanical life-forms, which are based solely on the form of the whole plant (Brown 1977a:320).

Labeled zoological categories that meet the definitional criteria of universal life-forms

(involving gross morphology) occasionally do not meet the nomenclatural criteria for life-forms as outlined by Berlin. In addition, in some instances these classes are not especially large or heterogeneous. In this study definitional criteria take precedence over other criteria (nomenclature, heterogeneity), and such categories are formally treated as life-forms.<sup>1</sup>

#### LIFE-FORMS AS BINARY OPPOSITIONS

Although universal life-forms are typically encoded through use of a single label, such categories can be lexically recognized in other ways. For example, the life-form BIRD is frequently encoded through binary opposition. Languages doing so will lack a label for "birds in general" but will name one or both of the contrasting classes "large bird"/"small bird." I have encountered examples of the use of binary opposition in encoding FISH, SNAKE, and MAMMAL in addition to BIRD (see Table I).

Encoding by binary contrast is a common feature of language. Indeed, dimensions are apparently always encoded through binary opposition before they are assigned single labels. For example, the dimensions denoted by the contrasts sharp/dull and hard/soft differ from dimensions denoted by deep/shallow and wide/narrow in that they lack non-productive labels. Productive labels for the former two dimensions are, of course, derived from words encoding the contrasts they designate: sharpness or dullness and hardness or softness. The developmental priority of binary opposition is also reflected by the fact that nonproductive names for dimensional concepts are frequently derived as well from words encoding contrasts: depth (from deep) and width (from wide).

For botanical life-forms, "tree" and "grerb" emerge when the binary opposition "large plant"/"small plant" is lexically encoded (Brown 1977a, 1977b). Languages having "tree" and "grerb" will often lack a term for plants in general. However, the contrast itself signals recognition of plants as a unified grouping of living things set off from animals. A common procedure for deriving labels for the botanical unique beginner lends support to this argument. Frequently, words for "plant," like labels for dimensional concepts (e.g., sharpness, depth), are derived from terms encoding associated contrasts. Berlin (1972:78-80), for example, cites languages in which labels for "tree" and "grerb" are used as well in reference to "plant," and argues that the nonunique beginner usage is developmentally prior.

Shoshoni offers an example in which a term for a zoological life-form is derived from one of the labels for an associated contrast. Hage and Miller (1976) report that the Shoshoni lexically distinguish "large bird"/"small bird." The "large bird" term, *kwinaa*, is three-way polysemous since it refers also more specifically to "eagle" and more generally to BIRD. The latter usage is clearly the more recent of the term's three referential applications. The original meaning of *kwinaa* almost certainly was "eagle." This term, for example, is found in Northern Paiute, in which it is restricted to "eagle" (Fowler and Leland 1967). In Shoshoni, *kwinaa* expanded in reference from eagle to large birds in general. This type of lexical change is discussed in detail in Berlin (1972) and in Brown (1979). With the lexical encoding of the "large bird"/"small bird" distinction, BIRD emerged as a life-form category. A single label for this life-form was subsequently acquired by promoting to life-form status one of the two words encoding the contrast.

Ethnobiological oppositions of the types discussed above occasionally involve overt marking (see Brown 1977a:334-335). This results when a label for one oppositional pole is derived from its counterpart. Encoding of adjectival oppositions sometimes entails overt marking. In Spanish, for example, the opposition "deep"/"shallow" is encoded through use of *profundo* and *poco profundo*, respectively. The latter label is an example of overt marking whereby the term for the unmarked pole ("deep") is used as part of a

compound label for the marked pole ("shallow"). *Poco profundo* translates literally into "little deep."

Several possible examples of overtly marked "grerb" were listed in the botanical life-form paper (Brown 1977a:335) and several additional cases in Mayan languages have been reported (Brown 1979). These polynomial "grerb" terms consist of a label for "tree" and a modifier (e.g., Kanjobal Mayan † *eʔ* "tree"/*mok'an* † *eʔ* "grerb," literally "short tree"). I have encountered five possible examples of overt marking pertaining to zoological life-forms: Gimi *umi* "large mammal"/*atumi* "small mammal," Ndumba *fai* "large mammal"/*faahi* "small mammal," Cayapa *piñi* "snake"/*tupiñi* "worm," Tarahumara *si-nó-w-i* "snake"/*no-w-i* "worm," and Natick *askook* "snake"/*oohk* "worm" (literal translations of overtly marked items could not be determined from original sources).

The Cayapa, Tarahumara, and Natick examples are direct evidence that "snake" and "worm" terms arise through lexical encoding of a "large elongated animal"/"small elongated animal" opposition. In this study the life-form SNAKE or "elongated animal" is judged present when a language demonstrates a label for "snake" or "worm" or labels for both. These two categories are then treated as a binary contrast. It is frequently the case that when only one pole of this opposition, i.e., "snake" or "large elongated animal," is lexically encoded, worms or small elongated animals are included in the small-creature class WUG. SNAKE, of course, is also lexically encoded through use of a single term that designates both snakes and worms and, on occasion, other elongated creatures as well.

#### LANGUAGE CASES

This article surveys the zoological life-form inventories of 112 languages. Data were assembled from two major sources: (a) published and unpublished accounts of firsthand investigation of native zoological classification and (b) dictionaries. Languages are referred to as Type A or Type B cases according to the kind of source used in collecting data pertaining to them, respectively (a) and (b) above. Type A cases were investigated by the author, were obtained from ethnozoological reports (monographs and articles) and from dictionaries with special sections dealing with zoological classification, or were supplied to the author by individuals who gathered data firsthand. Type B cases were collected from dictionaries. Dictionaries with less than 3,000 entries were used only if they proved especially thorough with respect to animal glosses. In some instances Type A cases were supplemented by dictionary materials.

Table I indicates the approximate ranges of zoological life-forms possessed by languages. For example, "[SNAKE(snake); WUG(worm + bug)]" preceding cases 102 and 103 (respectively Delaware and Mongolian) indicates that the SNAKE life-forms in these languages encompass snakes alone, and that their WUG life-forms include both worms and bugs. As another example, "[SNAKE(snake/worm); WUG(bug)]" preceding a number of cases (55-67) indicates that SNAKE is encoded through binary opposition ("large elongated animal"/"small elongated animal") and that WUG is restricted to bugs. Details of FISH, BIRD, and MAMMAL are indicated only when these differ in some important respect from equivalent classes in English: e.g., "BIRD(large bird/small bird)," which notes that this life-form is recognized through a large/small contrast. In addition, superordinate and subordinate distinctions are indicated: e.g., "FISH(fish/small fish)" means that a language has a general term for "fish" and a separate term for "small fish," the former taxonomically preceding the latter.

Most decisions concerning the presence or absence of zoological life-forms in languages involved unequivocal information. A brief discussion will illustrate the nature of some of the less straightforward cases. Fowler and Leland (1967) describe a category for Northern

TABLE 1. STAGE AFFILIATIONS OF 112 TYPE A AND B LANGUAGE CASES WITH ASSOCIATED SOCIETAL COMPLEXITY SCORES.

STAGE AFFILIATION	COMPLEXITY SCORE
<i>Stage 1 (one item)</i>	
BIRD	
1. Pintupi <sup>b</sup>	—
<i>Stage 2 (two items)</i>	
FISH/BIRD	
2. Chrau <sup>a1</sup>	—
3. Yana <sup>b</sup>	2
FISH/SNAKE	
[SNAKE (snake/worm)]:	
4. Muyuw <sup>b</sup>	—
5. Tausug <sup>b</sup>	—
[FISH (large fish/small fish); SNAKE (large snake/worm)]:	
6. Bontok Ignorot <sup>b</sup>	—
BIRD/SNAKE	
[SNAKE (snake[/worm?])]:	
7. Smith River <sup>a1</sup>	—
8. Karok <sup>a1</sup>	—
[SNAKE (snake/worm)]:	
9. Crow <sup>b</sup>	2
[SNAKE (worm)]:	
10. Northern Paiute <sup>a1</sup>	1
11. Diegueño <sup>b</sup>	1
<i>Stage 3 (three items)</i>	
FISH/BIRD/SNAKE	
[SNAKE (snake + eel[/worm?])]:	
12. Nungubuyu <sup>a1</sup>	—
[SNAKE (snake + worm + eel)]:	
13. Nukuoro <sup>b</sup>	—
[SNAKE (snake + worm/worm)]:	
14. Mixtec <sup>b</sup>	—
[SNAKE (snake + worm + other reptile/snake)]:	
15. Anglo-Saxon <sup>b</sup>	—
[SNAKE (snake + lizard/snake/worm)]:	
16. Hawaiian <sup>b</sup>	5
[SNAKE (snake/worm)]:	
17. Cayapa <sup>b</sup>	1
18. Classical Nahuatl <sup>b</sup>	—
19. Huastec <sup>a3</sup>	5
20. Igbo <sup>b</sup>	—
21. Jacaltec <sup>b</sup>	—
22. Kekchi <sup>b</sup>	—
23. Maidu <sup>b</sup>	3
24. Marshallese <sup>b</sup>	—
25. Palauan <sup>b</sup>	4
26. Pangasinan <sup>b</sup>	—
27. Tarahumara <sup>b</sup>	2
28. Tewa <sup>a1</sup>	1
29. Tifal <sup>b</sup>	—
30. Yareba <sup>b</sup>	—

Table I continued

STAGE AFFILIATION	COMPLEXITY SCORE
[SNAKE (snake + worm)]:	
31. Totonac <sup>b</sup>	7
[SNAKE (snake)]:	
32. Luiseño <sup>b</sup>	1
[SNAKE (worm)]:	
33. Kusaiean <sup>b</sup>	4
34. Mokilese <sup>a1</sup>	—
[BIRD (bird/large bird/small bird); SNAKE (snake/worm)]:	
35. Shoshoni <sup>a1, b</sup>	1
[BIRD (bird/large bird); SNAKE (snake/worm)]:	
36. Achumawi <sup>b</sup>	—
37. Quichua <sup>b</sup>	—
[BIRD (bird/small bird); SNAKE (snake/worm)]:	
38. Huitoto Muinane <sup>b</sup>	—
39. Natick <sup>b</sup>	—
[FISH (fish/small fish); BIRD (large bird/small bird); SNAKE (snake/worm)]:	
40. Aguaruna <sup>a1, b</sup>	—
FISH/SNAKE/WUG	
[SNAKE (snake/worm); WUG (bug + bird)]:	
41. Tequistlatec <sup>b</sup>	—
42. Zapotec <sup>b</sup>	—
<i>Stage 4 (four items)</i>	
FISH/BIRD/SNAKE/WUG	
[SNAKE (snake + worm/snake); WUG (bug)]:	
43. Maranao <sup>b</sup>	—
[SNAKE (snake + worm/worm); WUG (bug)]:	
44. Zoque <sup>b</sup>	—
[SNAKE (snake + worm/snake/worm); WUG (bug)]:	
45. Indonesian <sup>a3</sup>	20.5
[SNAKE (snake + worm/worm); WUG (bug + bird)]:	
46. Huave <sup>b</sup>	—
[SNAKE (snake + other reptile); WUG (worm + bug)]:	
47. Osage <sup>b</sup>	—
[SNAKE (snake + lizard/worm); WUG (worm + bug)]:	
48. Pashto <sup>a3, b</sup>	—
[SNAKE (snake/worm); WUG (worm + bug)]:	
49. Choctaw <sup>b</sup>	3
50. Congo <sup>b</sup>	6
51. Kiowa <sup>b</sup>	—
52. Tibetan <sup>b</sup>	7
53. Yoruba <sup>b</sup>	7
[SNAKE (snake/worm); WUG (worm + bug + small reptile)]:	
54. Cornish <sup>b</sup>	—
[SNAKE (snake/worm); WUG (bug)]:	
55. Bikol <sup>b</sup>	—
56. Biloxi <sup>b</sup>	—
57. Efik <sup>b</sup>	—
58. Fijian <sup>b</sup>	4
59. Finnish <sup>b</sup>	47.5
60. Iroquois <sup>b</sup>	3
61. Kayan <sup>a4</sup>	—



Table I continued

STAGE AFFILIATION	COMPLEXITY SCORE
62. Lake Miwok <sup>b</sup>	—
63. Mexicano <sup>b</sup>	7
64. Tagalog <sup>b</sup>	7
65. Tzotzil <sup>b</sup>	—
66. Wappo <sup>b</sup>	1
67. Yapese <sup>b</sup>	5
[SNAKE (snake [/worm?]); WUG (bug + crab)];	
68. Thai <sup>a1</sup>	13.7
[SNAKE (snake/large snake/worm); WUG (worm + bug)];	
69. Japanese <sup>b</sup>	41.5
[SNAKE (snake); WUG (worm + bug)];	
70. Amharic <sup>b</sup>	7
71. Atakapa <sup>b</sup>	—
[SNAKE (snake); WUG (worm + bug + snake)];	
72. Western Apache <sup>a2</sup>	1
[BIRD (bird/small bird); SNAKE (snake + worm/snake); WUG (bug)];	
73. Pahlavi <sup>b</sup>	—
[BIRD (bird/small bird); SNAKE (snake + other reptile/worm); WUG (bug)];	
74. Ocaina <sup>b</sup>	—
[BIRD (bird/small bird); SNAKE (snake + other reptile/snake/worm); WUG (bug)];	
75. Chamorro <sup>b</sup>	—
[BIRD (bird/small bird); SNAKE (snake/worm); WUG (bug)];	
76. Arabic <sup>b</sup>	7
[BIRD (bird/small bird); SNAKE (snake); WUG (worm + bug)];	
77. Navaho <sup>a1, b</sup>	1
[BIRD (bird/large bird); SNAKE (snake/large snake/worm); WUG (bug)];	
78. Galla <sup>b</sup>	2
[BIRD (large bird/small bird); SNAKE (snake/worm); WUG (bug)];	
79. Dyola <sup>b</sup>	—
[BIRD (large bird/small bird); SNAKE (snake); WUG ( <i>literally</i> , "small creature")];	
80. Kikuyu <sup>b</sup>	2
[FISH (fish + worm); SNAKE (snake); WUG (bug)];	
81. Papago-Pima <sup>a1, 2</sup>	1
FISH/BIRD/SNAKE/MAMMAL	
[SNAKE (snake/worm)];	
82. Tzeltal <sup>a1, 4</sup>	—
[BIRD (bird/small bird); SNAKE (snake [/worm?])];	
83. Yurok <sup>a1</sup>	1
[SNAKE (snake + worm + lizard + reptilelike insect + fish); MAMMAL (large mammal/small mammal)];	
84. Kyaka Enga <sup>a4</sup>	—
BIRD/SNAKE/WUG/MAMMAL	
[SNAKE (snake + worm + lizard); WUG (bug)];	

Table I continued

STAGE AFFILIATION	COMPLEXITY SCORE
MAMMAL (large mammal/small mammal): 85. Gimi <sup>a1</sup>	—
[SNAKE (snake + worm + lizard); WUG (bug + snail + frog + snake + worm + lizard); MAMMAL (large mammal/small mammal): 86. Ndumba <sup>a4</sup>	—
<i>Stage 5 (five items)</i>	
FISH/BIRD/SNAKE/WUG/MAMMAL	
[SNAKE (small snake + large worm/snake + large worm/small worm); WUG (bug)]: 87. Chorti <sup>a1</sup>	—
[SNAKE (snake/worm); WUG (worm + bug + reptile)]: 88. Tiruray <sup>b</sup>	—
[SNAKE (snake/worm); WUG (worm + bug)]: 89. Czech <sup>a4, b</sup>	65.5
90. Hungarian <sup>b</sup>	36.8
91. Pali <sup>b</sup>	—
92. Welsh <sup>b</sup>	—
[SNAKE (snake); WUG (bug + small nonsnake reptile + toad + tortoise [+ worm?]); MAMMAL (mammal + large nonsnake reptile): 93. Azande <sup>a1</sup>	5
[SNAKE (snake); WUG (bug + small nonsnake reptile + small amphibian + crustacean [+ worm?]); MAMMAL (mammal + large nonsnake reptile): 94. Hill Pandaram <sup>a1</sup>	—
[SNAKE (snake/worm); WUG (bug)]: 95. Albanian <sup>b</sup>	—
96. Armenian <sup>b</sup>	—
97. American English <sup>a3</sup>	109.4
98. Irish <sup>b</sup>	—
99. Italian <sup>b</sup>	41.3
100. Serbo-Croatian <sup>b</sup>	—
[SNAKE (snake/worm); WUG (bug + crayfish)]: 101. Turkish <sup>b</sup>	23.9
[SNAKE (snake); WUG (worm + bug)]: 102. Delaware <sup>a1, b</sup>	1
103. Mongolian <sup>b</sup>	5
[SNAKE (snake); WUG (worm + bug + small nonsnake reptile)]: 104. Chinese <sup>a3, b</sup>	13.0
[SNAKE (snake); WUG ( <i>literally</i> , "small creature")]: 105. Zulu <sup>b</sup>	7
[BIRD (bird/small bird); SNAKE (snake + worm); WUG (bug)]: 106. Mixe <sup>b</sup>	—
[BIRD (bird/small bird); SNAKE (snake/worm); WUG (worm + bug)]: 107. Mende <sup>a2</sup>	5
[SNAKE (snake + lizard); WUG (worm + bug); MAMMAL (large mammal/small mammal): 108. Fore <sup>a1</sup>	—
[SNAKE (snake + lizard/snake/large snake/small snake/worm);	

Table I continued

STAGE AFFILIATION	COMPLEXITY SCORE
WUG (bug); MAMMAL (large mammal/small mammal + frog): 109. Kalam <sup>a1</sup>	—
[SNAKE (snake + worm + lizard + fish); WUG (bug + frog); MAMMAL (large mammal + cassowary + eel/small mammal)]: 110. Rofaifo <sup>a1,4</sup>	—
[BIRD (large bird/"tree" bird/"water" bird); SNAKE (snake); WUG (worm + bug); MAMMAL (large mammal/small mammal + small residual bird)]: 111. Puget Salish <sup>b</sup>	—
[BIRD (large bird/small bird); SNAKE (snake); WUG (worm + bug + snake + toad); MAMMAL (large mammal/small mammal)]: 112. Ojibwa <sup>a1,b</sup>	1
Type A cases: a1, monograph or article source a2, dictionary (special treatment) a3, collected by author a4, personal communication	
Type B cases: b, dictionary source	

Paiute (10) which includes ants, spiders, ticks, beetles, lizards, and snakes. Flying insects are excluded. In addition, the category's label, *nuyúad̄i*; translates literally as "crawlers." Thus, *nuyúad̄i* actually refers to small creatures having a particular means of locomotion and consequently does not meet the definitional criteria of a WUG life-form (involving gross morphology). Similarly, the word *kúj*, glossed by "louse; bacteria; bug; flea; germ; parasite" in Abo et al.'s (1976) dictionary of Marshallese (24), appears to refer to small creatures that infest the bodies of people and animals, rather than to small creatures in general, i.e., WUG.

Other examples involve positive rather than negative decisions. Chrau (2) is judged as having a BIRD life-form. The term involved is used in reference to all birds (large and small) with the exception of birds of prey. Similarly, Puget Salish (111) is judged as possessing a FISH life-form even though the category is not extended to all fish. The label involved is a general term for all sea-going fish (salmon, trout, etc.) that spawn in fresh water.

Other decisions involved codification of a category as one or another of the five universal life-forms. Tequistlatec (41) and Zapotec (42), for example, both lump together birds and bugs (including both flying and nonflying insects) in labeled classes. I have identified those categories as WUG rather than as BIRD, for reasons to be set forth presently.

#### ZOOLOGICAL LIFE-FORM ENCODING SEQUENCE

There are 32 (2<sup>5</sup>) logically possible combinations of the 5 universal zoological life-forms that can occur in languages.<sup>2</sup> The 112 cases of Table I, however, demonstrate only 10 of these patterns. There are, therefore, severe constraints upon the types of animal life-form classes that may occur together in the lexicons of language. These constraints are realized as implicational universals.

With only four exceptions (cases 41, 42, 85, and 86), if a language has a WUG life-form, it will also possess FISH, BIRD, and SNAKE classes. However, languages having

one, two, or all of the latter three life-forms do not necessarily have WUG. Thus, the presence of WUG implies the presence of FISH, BIRD, and SNAKE. (The occurrence of WUG *always* implies at least two of the three life-forms FISH, BIRD, and SNAKE.) Similarly, with only three exceptions (cases 82, 83, and 84), if a language has MAMMAL, it will have WUG, although not necessarily vice versa. Thus, MAMMAL implies WUG.

If these synchronic implicational relationships are interpreted diachronically, then FISH, BIRD, and SNAKE are added to lexicons before WUG, and WUG before MAMMAL. This encoding sequence is summarized in Figure 1, which shows that languages at Stage 1 in zoological life-form growth encode one of the triad FISH, BIRD, and SNAKE, but in no particular order; those at Stage 2 encode two; and those at Stage 3 encode three. Stage 4 languages add WUG and Stage 5 languages add MAMMAL.

Thirty-four of the 112 languages are Type A cases (identified by use of a superscript *a* in Table I). Type A languages are affiliated with four of the five growth stages. The numbers of Type A cases per stage are as follows: Stage 2 (4), Stage 3 (6), Stage 4 (12), and Stage 5 (12).

Of the language cases surveyed 105 (93.75%) conform to the encoding sequence while only seven (6.25%) do not. Four of these exceptions, Tequistlatec (41), Zapotec (42), Gimi (85), and Ndumba (86) have not encoded all three of the FISH, BIRD, and SNAKE triad before adding WUG. The remaining exceptions, Tzeltal (82), Yurok (83), and Kyaka Enga (84), have the initial triad and MAMMAL but lack WUG.

Tequistlatec and Zapotec have added WUG after encoding only FISH and SNAKE (BIRD is lacking). Though these languages are exceptions to the encoding sequence, they do not constitute serious anomalies with respect to principles of naming-behavior underlying regularities in zoological life-form categorization to be discussed presently.

Gimi and Ndumba have added WUG after encoding only BIRD and SNAKE (FISH is missing). Speakers of these two languages live in the Eastern Highlands of New Guinea. Glick, who describes Gimi ethnozoological classification, reports that his informants were only familiar with eels and tinned fish (1964:280). Terence E. Hays (personal communication) similarly notes that eels are the only native fish known to Ndumba speakers. Lack of fish in areas in which Gimi and Ndumba are spoken undoubtedly is related to the absence of FISH life-forms in these languages. This ecological factor may also account for the manner in which fish are classified in two other New Guinea Highlands languages. Both Kyaka Enga (84) and Rofaifo (110) have labeled FISH classes, but these are respectively included in highly heterogeneous SNAKE life-forms (see Table I). Apparently, native exemplars of fish are so few in number that speakers of these languages do not feel justified in maintaining their FISH categories as unaffiliated groupings.<sup>3</sup> The inclusion of fish in SNAKE is understandable since these animals share the feature "elongation" with other creatures such as snakes, worms, lizards, and so on. In this connection, it is interesting that the desert dwelling Papago-Pima (81) lump together fish and worms in a single labeled class.

Tzeltal, one of the three deviant cases having all life-forms except WUG, may not in fact constitute an exception to the encoding sequence. The word *čanbalam*, according to Hunn (1977), is used by Tzeltal speakers as if it were a MAMMAL label. The same word also functions as a unique beginner label referring to animals in general, except human beings. Thus, it would appear that *čanbalam* is polysemous. Hunn (1977:135), however, does not claim that the word has two distinct referential applications. Rather, it seems that the most typical or representative *čanbalam* "animal" is found in the mammal grouping. Thus, the term merely appears to function as a MAMMAL label when the referential cutoff point for any Tzeltal speaker using it is something short of animals in general. As Hunn (1977:135) puts it, "The cut-off point in a given case will depend on the momentary perception of the balance between the core [i.e. the mammal focus] and

the extended range of the term." If Tzeltal lacks MAMMAL, it is simply an unexceptional Stage 3 case.

#### GALTON'S PROBLEM AND GROWTH STAGES

Galton's problem could be raised in response to the findings reported here. In other words, the implicational relationships for folk zoological life-forms may be an artifact of the "relatedness" of the languages surveyed rather than the product of universal underlying principles of naming-behavior. Geographically and/or genetically related languages may be affiliated with the same stage of life-form growth. If so, many of the logically possible combinations of life-forms do not occur in languages because of areal diffusion of only a limited number of patterns and/or because only a few combinations were originally represented in the lexicons of parent languages.

Table II organizes the 112 languages surveyed according to genetic relationship and roughly by geography, and indicates zoological life-form stage affiliation. Genetically related languages of the sample are not regularly associated with the same life-form stage. For example, well-represented language groupings (e.g., Indo-European, Mayan, Niger-Congo, Central New Guinea, and Uto-Aztecan) are affiliated with at least three of the total five stages. Austronesian, the family represented by the most languages (18), shows association with all but one of the five stages. Geographically contiguous languages also tend not to share the same stage of life-form growth. For example, 17 Western U.S. languages (Achumawi, Crow, Diegueño, Karok, Lake Miwok, Luiseño, Maidu, Navaho, Northern Paiute, Papago-Pima, Shoshoni, Smith River, Tewa, Wappo, Western Apache, Yana, and Yurok) range from Stage 2 to Stage 4. Thus, neither diffusion nor genetic relationship appears to have figured significantly in the observed distribution of life-form inventories.

TABLE II. CLASSIFICATION OF LANGUAGES WITH STAGE AFFILIATIONS.

---

*Salish*: Puget Salish (Stage 5).

*Macro-Algonkian*:

*Algonkian*: Natick (Stage 3), Delaware, Ojibwa (Stage 5).

*Other*: Yurok (Stage 4).

*Gulf\**:

*Siouan*: Crow (Stage 2), Biloxi, Osage (Stage 4).

*Other*: Atakapa, Choctaw (Stage 4).

*Iroquoian*: Iroquois (Stage 4).

*Athapaskan*: Smith River (Stage 2), Navaho, Western Apache (Stage 4).

*Penutian*: Maidu (Stage 3), Lake Miwok (Stage 4).

*Yukian*: Wappo (Stage 4).

*Hokan*: Diegueño, Yana, Karok (Stage 2), Achumawi, Tequistlatec (Stage 3).

*Aztec-Tanoan*:

*Kiowa-Tanoan*: Tewa (Stage 3), Kiowa (Stage 4).

*Uto-Aztecan*:

*Aztecan*: Classical Nahuatl (Stage 3), Mexicano (Stage 4).

*Sonoran*: Tarahumara (Stage 3), Papago-Pima (Stage 4).

*Takic*: Luiseño (Stage 3).

*Numic*: Northern Paiute (Stage 2), Shoshoni (Stage 3).

*Mesoamerican\*\**:

*Otomanguean*: Mixtec, Zapotec (Stage 3).

*Mayan*: Huastec, Jacaltec, Kekchi (Stage 3), Tzeltal, Tzotzil (Stage 4), Chorti (Stage 5).

*Zoquean*: Zoque (Stage 4), Mixe (Stage 5).

*Others*: Totonac (Stage 3), Huave (Stage 4).

Table II continued

*Macro-Chibchan*: Cayapa (Stage 3).

*Macro-Carib*:

*Witotoan*: Huitoto Muinane (Stage 3), Ocaina (Stage 4).

*Andean-Equatorial*: Aguaruna, Quichua (Stage 3).

*Austronesian*:

*Eastern Oceanic*:

*Polynesian*: Hawaiian, Nukuoro (Stage 3).

*Other*: Fijian (Stage 4).

*Micronesian*: Kusaian, Marshallese, Mokilese (Stage 3), Yapese (Stage 4).

*Northwest Austronesian*: Bontok Igorot, Tausug (Stage 2), Palauan, Pangasinan (Stage 3), Bikol, Chamorro, Maranao, Tagalog (Stage 4), Tiruray (Stage 5).

*West Indonesian*: Indonesian, Kayan (Stage 4).

*Kam-Tai*: Thai (Stage 4).

*Mon-Khmer*: Chrau (Stage 2).

*Central New Guinea*:

*East New Guinea Highlands*:

*East Central*: Gimi (Stage 4), Fore, Rofaifo (Siane) (Stage 5).

*Eastern*: Ndumba (Tairora) (Stage 4).

*West Central*: Kyaka Enga (Stage 4).

*Karam*: Kalam (Stage 5).

*Southeast New Guinea*: Yareba (Stage 3).

*Central and South New Guinea*: Tifal (Stage 3).

*Unclassified New Guinea language (Austronesian?)*: Muyuw (Stage 2).

*Australian Macro-Phylum*: Pintupi (Stage 1), Nunggubuyu (Stage 3).

*Sino-Tibetan*: Tibetan (Stage 4), Chinese (Stage 5).

*Japanese*: Japanese (Stage 4).

*Ural-Altaiic*:

*Altaic*: Mongolian, Turkish (Stage 5).

*Uralic*: Finnish (Stage 4), Hungarian (Stage 5).

*Dravidian*: Hill Pandaram (Stage 5).

*Indo-European*:

*Indo-Iranian*: Pahlavi, Pashto (Stage 4), Pali (Stage 5).

*Slavic*: Czech, Serbo-Croatian (Stage 5).

*Italic*: Italian (Stage 5).

*Celtic*: Cornish (Stage 4), Irish, Welsh (Stage 5).

*Germanic*: Anglo-Saxon (Stage 3), American English (Stage 5).

*Others*: Albanian, Armenian (Stage 5).

*Afroasiatic*:

*Semitic*: Amharic, Arabic (Stage 4).

*Cushitic*: Galla (Stage 4).

*Niger-Congo*:

*Benue-Congo*:

*Bantu Proper*: Congo, Kikuyu (Stage 4), Zulu (Stage 5).

*Cross River*: Efik (Stage 4).

*Kwa*: Igbo (Stage 3), Yoruba (Stage 4).

*West Atlantic*: Dyola (Stage 4).

*Adamawa-Eastern*: Azande (Stage 5).

*Mande*: Mende (Stage 5).

---

\*Springer and Witkowski (1978)

\*\*Witkowski and Brown (1978a)

This is not to propose that diffusion and genetic relationship never influence life-form growth. For example, while genetically related languages frequently possess more biological life-forms than their parent language, they rarely, if ever, possess fewer (cf.

Brown 1979). Mayan languages of the sample (Chorti, Huastec, Jacalteco, Kekchi, Tzeltal, and Tzotzil), for instance, are associated with Stages 3-5 (having three to five zoological life-forms). Lexical reconstruction indicates that Proto-Mayan probably had both a FISH and a SNAKE life-form and, possibly a BIRD life-form, thus suggesting that the language was associated with Stage 3 in life-form growth. Mayan daughter languages have at least as many life-form terms (three) (and sometimes four or five) as the estimation for Proto-Mayan.

There are instances in which zoological life-form classes have diffused. For example, Tequistlatec (41) and Zapotec (42), both spoken in the southeastern corner of the Mexican state of Oaxaca, share unusual zoological life-form categories that are identical in referential range (though not in lexical designation). These languages are exceptions to the encoding sequence since both lack BIRD but nonetheless encode WUG. WUG life-forms in these languages encompass bugs and birds, a category range occurring in only one other language surveyed. This sharing of an unusual category is not attributable to genetic relationship. Zapotec is a member of the Zapotecan grouping, which is one of eight families of the Otomanguean stock located entirely in Mesoamerica. On the other hand, Tequistlatec is related to Yuman languages spoken in California and Arizona and thus is probably intrusive to Mesoamerica. Consequently, that Zapotec and Tequistlatec share intimate details of zoological life-form classification is almost certainly caused by diffusion.

Huave (46), the other language in which WUG encompasses birds, is spoken in an area bordering on Tequistlatec and Zapotec territory in southeast Oaxaca. Huave has a labeled BIRD taxon, but this class is included in WUG. While Huave is distantly related to Zapotec (Witkowski and Brown 1978a), the relationship is so remote that diffusion is the more plausible explanation of identical WUG ranges in these two languages.

A number of New Guinea languages encode MAMMAL through binary opposition. The distribution of this feature is also caused by areal diffusion. All six East New Guinea Highlands languages surveyed, Kyaka Enga (84), Gimi (85), Ndumba (86), Fore (108), Kalam (109), and Rofaifo (110), lexically encode "large mammal"/"small mammal" through use of dissimilar sets of two terms. While these languages are genetically related (Voegelin and Voegelin 1977:257-259), it is unlikely that the contrast is of great antiquity and, thus, traceable to a parent language, since MAMMAL tends to emerge very late in zoological life-form growth. Independent development of the large/small contrast for MAMMAL by these languages also seems unlikely since this distinction otherwise appears only rarely and sporadically in the sample of 112 cases (see only Puget Salish [111] and Ojibwa [112]).

#### SOCIETAL COMPLEXITY AND GROWTH STAGES

The botanical life-form study (Brown 1977a:328-332) reported a positive association between size of life-form lexicon and societal complexity. A similar correlation exists between the number of zoological life-forms and societal complexity. Languages with three or fewer zoological life-forms (Stages 1-3) are usually spoken by peoples living in small-scale societies lacking the complex political integration, social stratification, and technological elaboration of those who speak languages having four or more zoological life-form terms (Stages 4 and 5).

Measures of societal complexity employing a wide range of index variables (e.g., Freeman 1957; Marsh 1967; Naroll 1956) all correlate strongly with each other (Schaefer 1969). Marsh's (1967:338-347) scale indexes societal complexity in terms of size and integration of political units and degree of social stratification. A primary scale ranging from 0 (low) to 7 (high) applies to societal units other than contemporary national societies.

National societies range from 8.6 (for Portuguese Guinea) to 109.4 (for the United States).

Table I lists societal complexity scores for the 49 societies (languages) of the set of 112 found in Marsh's index. The association between societal complexity and size of zoological life-form lexicons (given as life-form stages) is presented in Table III. The correlation coefficient, gamma, is .66 ( $p < .02$ ,  $N = 49$ ).

Positive correlations between number of botanical and zoological life-forms and societal complexity are linked to taxonomic devolution. Folk biological taxonomies tend to decay "from the bottom up" (from more specific classes to less specific classes) as people urbanize and become increasingly separated from direct reliance and dependence on the world of plants and animals (Berlin 1972; Brown 1977a; Witkowski and Brown 1978b). Taxonomic decay involves both loss of knowledge of biological referents of terms for generic and specific classes and loss of knowledge of the terms themselves. As a consequence, more general terms, such as life-form names, become increasingly useful and salient (cf. Dougherty 1978a), so much so that they tend to increase in number. Thus, the growth of ethnobiological life-forms ironically indexes an overall lack of interest in and interaction with the natural world.

For folk zoological taxonomy, an increase in societal complexity may not have always resulted in a uniform devolution of systems. Some parts of zoological taxonomies may tend to decay less extensively than others. For example, possibly those portions of animal taxonomies dealing with the classification of mammals have not devolved to the degree that other parts have. For most of humankind's existence, mammals have constituted important, if not the most important, animals in its environment. As societies shifted from hunting and gathering to food-producing economies, mammals, unlike creatures such as fish, birds, and snakes, acquired new significance as domesticated animals. The late emergence of MAMMAL in zoological life-form growth may be partly caused by high levels of interest in mammals characterizing both small-scale and large-scale societies. Since specific and generic mammal terms maintain their usefulness and salience, broad zoological categories encompassing all mammals are less useful than otherwise.

#### EXPLANATION OF LIFE-FORM ENCODING REGULARITIES

While MAMMAL's late encoding may be traced to relative interest and cultural importance, most of the zoological life-form encoding regularities are underlain by more general principles of naming-behavior. These include: (1) criteria clustering; (2) conjunctivity (including binary opposition); (3) dimension salience; and (4) marking (see Brown 1977a, 1977b, 1979; Brown and Witkowski 1979; Witkowski and Brown 1977, 1978b).

TABLE III. ASSOCIATION BETWEEN SOCIETAL COMPLEXITY AND SIZE OF ZOOLOGICAL LIFE-FORM LEXICON (GIVEN AS LIFE-FORM STAGE).

Societal complexity	Life-form stage		
	1-3	4	5
High (above 7)	0	4	6
Medium (4-7)	5	9	4
Low (0-3)	10	9	2
gamma = .66	p < .02		N = 49



*Criteria clustering* occurs when certain defining features of natural objects "bundle" so that the presence of any one feature is highly predictive of the presence of other features. This leads to the expectancy of features being present together. In this way "natural breaks" occur in the physical world that are often followed in the classification of objects (Bruner, Goodnow, and Austin 1956:47; Hunn 1977:46).

Criteria clustering figures prominently in the encoding priority of FISH, BIRD, and SNAKE. Bruner, Goodnow, and Austin (1956:47), for instance, illustrate this principle by citing the example of birds, animals possessing feathers, wings, and a bill or beak. A creature's possession of feathers is highly predictive of wings and a bill or beak, so much so that an expectancy of all these features being present together is built up. This expectancy can lead to the lexical encoding of BIRD. Similarly, FISH and SNAKE have respective sets of defining features showing high levels of mutual predictability. The occurrence of gills predicts fins and a streamlined body, and greatly elongated animals usually lack appendages as well as feathers and fur. In brief, FISH, BIRD, and SNAKE constitute salient discontinuities in nature and thus are natural candidates for lexical recognition. (See Hunn [1976, 1977] for a detailed consideration of the influence of discontinuities in nature upon folk classification.)

The late encoding of WUG and MAMMAL (see Figure 1) is in part a function of their *relative indistinctiveness* as natural discontinuities vis-à-vis the distinctiveness of FISH, BIRD, and SNAKE. Each of these two groupings is distinctly heterogeneous, demonstrating little criteria clustering. For example, while most mammals have four appendages used for locomotion and/or object manipulation, so do many other animals including such common creatures as lizards, salamanders, frogs, and turtles. Consequently, possession of four appendages is not particularly predictive of other animal characteristics. Exemplars of WUG have even less in common than creatures included in MAMMAL. WUG, for example, encompasses animals having legs and lacking them, having wings and lacking them, having segments and lacking them, and so on.

*Binary opposition* and *dimension salience* underlie the encoding of WUG and MAMMAL. Encoding through binary opposition is a common feature of language. Oppositional characteristics of dimensional concepts such as height, width, depth, etc., are usually encoded by two terms producing such familiar adjectival contrasts as tall/short, wide/narrow, deep/shallow, and so on. Sometimes nondimensional substantive concepts are also encoded through binary opposition. This entails lexical recognition of how related objects such as plants, elongated animals, birds, etc., contrast along some dimension or dimensions, e.g., "tree"/"grerb" (large plant/small plant), "snake"/"worm," "large bird"/"small bird," and so on.

Binary opposition is one way in which the principle of *conjunctivity* is realized in natural language categorization (Witkowski and Brown 1978b). The recognition of dimensional concepts such as height, width, and depth through binary opposition is conjunctive naming-behavior. An alternative way of encoding a dimension would be to name its midsection with a single term and its two extremes with another. This never occurs in dimensional naming, however, because it would violate conjunctivity. The category combining dimensional extremes would be disjunctive, and such categories are rare in human naming systems. Binary opposition represents the only way that does not contravene conjunctivity of partitioning a dimension through use of two terms. Thus, the principle of binary opposition in dimensional naming is the surface result of conjunctivity considerations.

Conjunctivity also underlies encoding nondimensional concepts through binary opposition, e.g., "tree"/"grerb." Languages could, for instance, categorize botanical objects through attention to size by assigning a name to all midsized plants and another name to all very large and very small plants. To treat plants in such a way would violate

conjunctivity and does not occur. Combining very large and very small plants (minus midsized plants) would be disjunctive. The “tree”/“grerb” or “large plant”/“small plant” contrast is the only way that is conjunctive of categorizing plants by size using two terms. Consequently, oppositional classification of objects in terms of underlying dimensions is due also to conjunctivity constraints.

In folk biological classification suprageneric categories based on binary opposition always involve size. The importance of size in suprageneric biological classification illustrates the principle of *dimension salience*. Highly salient dimensions pertain to large and varied sets of objects. Dimensions are not particularly salient if they only apply to a small number of different objects. Since all biological organisms vary by size, there is a strong tendency to incorporate this dimension into suprageneric classificatory strategies involving plants and animals.

The addition of WUG and MAMMAL to languages constitutes use of size in distinguishing a “residual” group of animals through binary opposition.<sup>4</sup> Animals included in these categories are residual since they are the creatures that are left over after the highly distinctive discontinuities FISH, BIRD, and SNAKE are lexically encoded. Thus addition of WUG and MAMMAL encodes the contrast “small residual animal”/“large residual animal.”

A classic example of the WUG/MAMMAL contrast is described by Evans-Pritchard (1963) for the Azande (93). He reports the following extensions for the Azande MAMMAL and WUG life-forms, respectively: “Reptiles, except the snakes, tend to be described as *anya* . . . if they are large and as *agbiro* . . . if they are small” (1963:139, emphasis in original). Thus Azande MAMMAL (*anya*) includes such creatures as iguanas in addition to mammals. Evans-Pritchard also notes that Azande WUG (*agbiro*) includes toads and tortoises in addition to bugs and small nonsnake reptiles.

MAMMAL and WUG in Hill Pandaram (94) parallel Azande life-forms. Morris (1976:349–350) describes Hill Pandaram MAMMAL as follows:

The category *mrgam* includes not only all known species of mammals . . . but several of the large species of reptiles. It excludes however snakes . . . and smaller reptiles and amphibia which are normally referred to, if categorized at all, as *puchi* [emphasis in original].

Morris (1976:348) describes *puchi* (WUG) as “a residual category . . . which includes insects, crustaceans and several other creatures.”

Inclusion of large creatures that are not mammals in MAMMAL and small creatures that are not worms or bugs in WUG offers evidence that these categories are based on residualness and size. Bulmer and Tyler (1968), for example, report that Kalam (109) speakers include frogs in their “small mammal” class (*as*). These authors (1968:352) speculate on why frogs and small mammals are lumped together. From the comparative perspective of this paper, frogs and small mammals are residual creatures sharing large size relative to Kalam WUG (*joŋ*) and small size relative to Kalam “large mammal” (*kmn*). The Rofaifo (110), who also make the “large mammal”/“small mammal” distinction, group bugs and frogs together in WUG. By so doing, Rofaifo (Siane) speakers have drawn the line separating big residual animals from little residual animals at a place along a continuum of animals ranked by size different from the point at which Kalam speakers have drawn it.

Rofaifo (Siane) speakers also include fresh water eels and cassowaries in their “large mammal” category. Since eels are not found in the Rofaifo environment and are known only through trade (Peter Dwyer, personal communication), they are treated as large residual creatures in suprageneric classification. Cassowaries, probably because they are large and flightless, are excluded from BIRD life-forms in Rofaifo (Siane) and in two related New Guinea languages, Fore (108) and Kalam (109). While Fore and Kalam have

elected to treat residual cassowaries as if they belong to an unaffiliated class of cassowaries, Rofaifo (Siane) speakers have incorporated these creatures into their named grouping of the largest residual animals.

Another language that encodes MAMMAL through binary contrast includes certain birds in a residual animal class. Puget Salish (111) speakers partition the universe of birds in a complex fashion. Three suprageneric categories are encoded, "large bird" (mainly birds of prey), "tree birds," and "waterfowl" (good swimmers, not shorebirds). There exists a residue of small birds that do not fit into any one of these three classes. In Puget Salish the latter creatures are included in "small mammal."

A number of languages surveyed, in addition to those already mentioned, include small residual creatures that are neither worms nor bugs in WUG. The WUG life-form of Chinese (104) encompasses small reptiles other than snakes; that of Cornish (54), various small reptiles; that of Ndumba (86), snails, frogs, and lizards; that of Ojibwa (112), toads; that of Thai (68), crabs; and that of Turkish (101), crayfish. According to the *Oxford English Dictionary* the English WUG term, *insect*, once had similar extensions. Part of the *OED* entry for *insect* reads, ". . . formerly (and still by the uneducated) applied still more widely, e.g., to earthworms, snails, and even some small vertebrates, as frogs and tortoises."

Nomenclature is sometimes indicative of the roles of size and residualness in the derivations of WUG and MAMMAL. For example, languages having zoological unique beginners such as the English *animal* will occasionally encode WUG through use of compound labels translating literally as "little animal." Five languages of those surveyed have such labels: Zapotec (42), Zoque (44), Efik (57), Fijian (58), and Mixe (106).

Unique beginner terms are frequently polysemous, referring to WUG or MAMMAL in addition to creatures in general. This type of polysemy is indicative of the residualness of organisms included in these two life-forms. An ordinary way of identifying residual creatures *before* emergence of WUG and MAMMAL terms is by use of a zoological unique beginner term, if present in a language. For example, any bug might be referred to simply as "a creature." When WUG or MAMMAL develops, a unique beginner term may, through extension of its common use as a label for residuals, develop quite naturally as a label for such a class, and in this manner acquire a polysemous application.

Such a development, for example, would account for the Bikol (55) use of *háyoḡ* as a label for both "creature" (unique beginner) and WUG. Cognates of this word found in other Philippine (northwest Austronesian) languages function exclusively as zoological unique beginners. Consequently, it is likely that *háyoḡ* was originally used by Bikol speakers only as a label for animals in general and subsequently developed WUG as an additional referent. American English *animal* is an example of a zoological unique beginner label that has taken on the additional referent MAMMAL. Among the 112 languages surveyed, Bikol (55), Tzotzil (65), Chomorro (75), and Tiruray (88) have polysemous terms referring to animals in general and, more specifically, to small residual animals (WUG); and Tzeltal (82), Chorti (87), Czech (89), American English (97), Delaware (102), and Mixe (106) have unique beginner terms used to designate large residual animals (MAMMAL) as well. In some of these cases, however, the less inclusive application (WUG or MAMMAL) may have preceded developmentally the more inclusive application (zoological unique beginner).

Though the tendency to add FISH, BIRD, and SNAKE before WUG and MAMMAL is obviously strong, it is not inconceivable that one or both of the latter two life-forms may occasionally arise before all three of the initial triad have been encoded. In such case, given principles underlying the emergence of WUG and MAMMAL, it is likely that animals that are associated with life-forms normally encoded before WUG and MAMMAL but have not been encoded in fact, would be treated as additional residual

creatures categorized by size. This has apparently occurred in two languages of the sample, Tequistlatec (41) and Zapotec (42).

These languages have added WUG to their life-form inventories while encoding only FISH and SNAKE of the initial triad. In both cases, birds are lumped with bugs in WUG. This classificatory treatment accords with principles underlying the emergence of WUG. In Tequistlatec and Zapotec, birds have been treated as zoological residuals sharing the feature "relative smallness" with other residual creatures. (Zapotec employs a compound label for WUG, *mani huiini?*, which translates literally as "little animal," while Tequistlatec uses a unitary lexeme, *taga*, for the life-form.)

Neighboring Tequistlatec and Zapotec share WUG life-forms encompassing birds and bugs as a result of diffusion (see earlier discussion). A third neighboring language, Huave (46), has the same extension but differs in that it has labeled BIRD. Animals included in BIRD are, in turn, included in WUG. Huave has adopted, probably through borrowing, the WUG range found in Tequistlatec and Zapotec while nonetheless recognizing BIRD as a distinct discontinuity.

The Huave example indicates that the tendency to focus on size in making suprageneric distinctions is apparently so strong that lack of residualness will sometimes be overlooked in assigning creatures to "residual" categories, especially to WUG. Seventeen other languages of those surveyed demonstrate this phenomenon. The Ndumba (86) SNAKE class, which encompasses snakes, worms, and lizards, is immediately included in WUG, as are SNAKE classes (restricted to snakes) in Western Apache (72) and Ojibwa (112). In Tiruray (88), all elongated animals included in the labeled classes "snake" and "worm" are included also in WUG. Finally, 13 languages label "worm" but nonetheless include worms in their WUG life-forms: Pashto (48), Choctaw (49), Congo (50), Kiowa (51), Tibetan (52), Yoruba (53), Cornish (54), Japanese (69), Czech (89), Hungarian (90), Pali (91), Welsh (92), and Mende (107).

The encoding priority of WUG vis-à-vis MAMMAL fits into the framework of *marking* developed over the years by Jakobson (1941), Greenberg (1966, 1969, 1975), and others. Marking frequently finds expression in antonyms such as deep/shallow and wide/narrow (Greenberg 1966:53). In such pairs one item will often be marked and the other unmarked. An unmarked item is more salient, tending to occur more frequently in ordinary language use than its marked opposite. This is especially noticeable in neutralized linguistic environments. For example, queries concerned with depth more often use the unmarked form "deep" than the marked form "shallow." Thus, people customarily ask "How deep is the river?" not "How shallow is the river?" In addition, marked items tend to be phonologically more complex than unmarked items, indicating that the unmarked forms are older and, hence, developmentally prior to marked items. (See Greenberg [1975] and Brown and Witkowski [1979] for additional features of marking relationships.)

The same pole of a dimension underlying an adjectival contrast will often be unmarked in all languages recognizing the dimension. Greenberg (1966:53), for example, shows that "deep" is universally unmarked, while its counterpart, "shallow," is universally marked. This result extends to both botanical and zoological life-forms encoded through binary opposition. In botanical classification "tree" or "large plant" is unmarked (since it is developmentally prior), whereas its counterpart, "grerb" or "small plant," is marked (Brown 1977a:334). Similarly, in zoological classification, WUG or "small residual animal" is unmarked, and its counterpart, MAMMAL, or "large residual animal," is marked.

Fitting these life-form encoding uniformities into the framework of marking shows that they do not constitute exceptional linguistic regularities. It is therefore possible that explanations eventually brought to bear upon marking uniformities in the adjectival com-

ponent of language will extend to marking in the biological life-form lexicon. On the other hand, these parallel regularities may not be due in all cases to the same underlying factors. For example, the unmarked status of WUG vis-à-vis MAMMAL is probably attributable to special considerations that are not linked to marking in other parts of the vocabulary.

I argued earlier that the growth of zoological life-form lexicons indexes an increasing indifference toward and a decreasing involvement with the animal world as societies become more complex. Peoples in all societies, both simple and complex, demonstrate relatively high levels of interest in large residual animals. MAMMAL, therefore, tends to be the last holdout against pressures for adding all five universal life-forms to vocabularies. Consequently, WUG, along with FISH, BIRD, and SNAKE, regularly precedes MAMMAL in life-form encoding and, hence, is the unmarked component of the WUG/MAMMAL opposition.

#### CHILD ACQUISITION OF ZOOLOGICAL LIFE-FORMS

Paul K. Chase (1978) reports that American English-speaking children acquire folk zoological life-forms in the same order in which these are added to languages (see Figure 1). Chase investigated the zoological life-form vocabularies of children ranging in age from about three to ten. This investigation involved sorting tasks (using pictures of animals pasted on cards) and traditional elicitation techniques. Chase found that children from three to five years old unambiguously recognize only two zoological life-forms of the universal set of five, FISH and SNAKE. Between the ages of five and seven, a third life-form, BIRD, is acquired, and shortly thereafter WUG. WUG for children younger than eight is occasionally extended to creatures such as crabs, lizards, and turtles in addition to bugs. After age eight, MAMMAL is learned. Principles of naming-behavior underlying zoological life-form growth almost certainly account also for parallel patterns in child acquisition of suprageneric animal classes.<sup>5</sup>

#### NOTES

*Acknowledgments.* The following people contributed to this project in important ways: Mark Flotow, Kuo Huang Han, M. Jamil Hanifi, Donn V. Hart, Robert M. Laughlin, Robert H. McDaniel, Jay Miller, Ronald Provencher, Brian Stross, Liana Suryo-Atmojo, and Paul R. Turner. The author is very grateful for their assistance. Ralph N. H. Bulmer, Michael Burton, Peter Dwyer, Terence E. Hays, Albert Heinrich, Eugene Hunn, and Stanley R. Witkowski are especially thanked for commenting in great detail on an earlier draft of this paper. It is impossible to say how many of the better ideas expressed here were formulated during many long and profitable discussions with my colleague Stanley R. Witkowski. Certainly, without his intellectual stimulation this paper would not have been written.

<sup>1</sup> Labeled classes meeting definitional criteria of universal life-forms but not meeting nomenclatural and other criteria of that rank have possibly devolved from full-fledged life-form categories, or, just as likely, are in the process of consolidating into such taxa.

<sup>2</sup> Only nine of these possible combinations are predicted by the zoological life-form encoding sequence (see Figure 1).

<sup>3</sup> Peter Dwyer (personal communication) reports that older male Rofaifo (Siane) speakers always include *laiva* (FISH) in the taxon *hoiafa* (SNAKE). In formal contexts, however, younger men accord *laiva* unaffiliated status. Only two types of fish other than eels are known to the Rofaifo. Ralph N. H. Bulmer (personal communication) recorded only five fish species in the Kyaka Enga domain.

<sup>4</sup> The role of "residualness" in biological classification was first recognized and discussed by Hunn (1976, 1977).

<sup>5</sup> Dougherty (1978b) similarly describes parallels between botanical life-form acquisition by American English-speaking children and botanical life-form growth (Brown 1977a).

## REFERENCES CITED

- Abo, Takaji, et al.  
1976 *Marshallese-English Dictionary*. Honolulu: University of Hawaii.
- Berlin, Brent  
1972 *Speculations on the Growth of Ethnobotanical Nomenclature*. *Language in Society* 1: 51-86.  
1976 *The Concept of Rank in Ethnobiological Classification: Some Evidence from Aguaruna Folk Botany*. *American Ethnologist* 3:381-399.
- Berlin, Brent, Dennis E. Breedlove, and Peter H. Raven  
1973 *General Principles of Classification and Nomenclature in Folk Biology*. *American Anthropologist* 75:214-242.  
1974 *Principles of Tzeltal Plant Classification: An Introduction to the Botanical Ethnography of a Mayan-speaking People of Highland Chiapas*. New York: Academic Press.
- Brown, Cecil H.  
1977a *Folk Botanical Life-Forms: Their Universality and Growth*. *American Anthropologist* 79:317-342.  
1977b *Lexical Universals and the Human Language Faculty*. In *Georgetown University Round Table on Languages and Linguistics 1977*. M. Saviile-Troike, ed. pp. 75-91. Washington, D.C.: Georgetown University Press.  
1979 *Growth and Development of Folk Botanical Life-Forms in the Mayan Language Family*. *American Ethnologist* 6:366-385.
- Brown, Cecil H., and Stanley R. Witkowski  
1979 *Language Universals*. In *Explaining Human Behavior*. David Levinson and Martin Malone, principal authors. New Haven: HRAF Press (in press).
- Bruner, J. S., J. J. Goodnow, and G. A. Austin  
1956 *A Study of Thinking*. New York: Wiley.
- Bulmer, R. N. H., and M. J. Tyler  
1968 *Karam Classification of Frogs*. *Journal of the Polynesian Society* 77:333-385.
- Chase, Paul K.  
1978 *Acquisition of Folk Zoological Life-Forms by American Children*. Manuscript. Department of Anthropology, Northern Illinois University.
- Dougherty, J. W. D.  
1978a *Salience and Relativity in Classification*. *American Ethnologist* 5:66-80.  
1978b *Learning Names for Plants and Plants for Names*. Manuscript. Department of Anthropology, University of Illinois, Urbana.
- Evans-Pritchard, E. E.  
1963 *Notes on Some Animals in Zandeland*, *Man* 63:139-142.
- Fowler, Catherine S., and Joy Leland  
1967 *Some Northern Paiute Native Categories*. *Ethnology* 6:381-404.
- Freeman, L. C.  
1957 *An Empirical Test of Folk-Urbanism*. Ann Arbor: University Microfilms.
- Glick, Leonard B.  
1964 *Categories and Relations in Gimi Natural Science*. *American Anthropologist* 66 (Number 4, Part 2, special publication): 273-280.
- Greenberg, Joseph H.  
1966 *Language Universals with Special Reference to Feature Hierarchies*. The Hague: Mouton.  
1969 *Language Universals: A Research Frontier*. *Science* 166:473-478.  
1975 *Research on Language Universals*. *Annual Review of Anthropology* 4:75-94.
- Hage, Per, and Wick R. Miller  
1976 "Eagle" = "Bird": A Note on the Structure and Evolution of Shoshoni Ethnoornithological Nomenclature. *American Ethnologist* 3:481-488.

- Hunn, Eugene S.  
 1976 Toward a Perceptual Model of Folk Biological Classification. *American Ethnologist* 3: 508-524.  
 1977 Tzeltal Folk Zoology: The Classification of Discontinuities in Nature. New York: Academic Press.
- Jakobson, Roman  
 1941 *Kindersprache, Aphasie, und allgemeine Lautgesetze*. Uppsala, Sweden: Almqvist and Wiksell.
- Marsh, Robert M.  
 1967 *Comparative Sociology: A Codification of Cross-societal Analysis*. New York: Harcourt, Brace & World.
- Morris, Brian  
 1976 Whither the Savage Mind? Notes on the Natural Taxonomies of a Hunting and Gathering People. *Man* 11:542-557.
- Naroll, Raoul S.  
 1956 A Preliminary Index of Social Development. *American Anthropologist* 58:687-715.
- Schaefer, James M.  
 1969 A Comparison of Three Measures of Social Complexity. *American Anthropologist* 71: 706-708.
- Springer, James W., and Stanley R. Witkowski  
 1978 A Reassessment of Southeastern Linguistics and Archaeology. *Bulletin of the Southeastern Archaeological Conference for 1978* (in press).
- Voegelin, C. F., and F. M. Voegelin  
 1977 *Classification and Index of the World's Languages*. New York: American Elsevier.
- Witkowski, Stanley R., and Cecil H. Brown  
 1977 An Explanation of Color Nomenclature Universals. *American Anthropologist* 79:50-57.  
 1978a Mesoamerican: A Proposed Language Phylum. *American Anthropologist* 80:942-944.  
 1978b Lexical Universals. *Annual Review of Anthropology* 7:427-451.

*Submitted 16 June 1978*

*Accepted 20 October 1978*

*Revisions received 11 November 1978*

#### APPENDIX: LANGUAGE CASE SOURCES

- Achumawi*. David L. Olmsted (1966), *Achumawi Dictionary*. Berkeley: University of California Press.
- Aguaruna*. Per Hage and Wick R. Miller (1976), "Eagle" = "Bird": A Note on the Structure and Evolution of Shoshoni Ethnoornithological Nomenclature. *American Ethnologist* 3:481-488 (specifically the personal communication from Brent Berlin quoted on pp. 484-485); Mildred L. Larson (1966), *Vocabulario Aguaruna de Amazonas*. Perú: Instituto Lingüístico de Verano.
- Albanian*. Stuart E. Mann (1957), *An English-Albanian Dictionary*. Cambridge: Cambridge University Press.
- American English*. Investigated by author.
- Amharic*. Wolf Leslau (1976), *Concise Amharic Dictionary: Amharic-English, English-Amharic*. Berkeley: University of California Press.
- Anglo-Saxon*. Joseph Bosworth (1888), *A Compendius Anglo-Saxon and English Dictionary*. London: Reeves and Turner; Carl Darling Buck (1949), *A Dictionary of Selected Synonyms in the Principal Indo-European Languages*. Chicago: University of Chicago Press; Gregory K. Jember (1975), *English-Old English, Old English-English Dictionary*. Boulder, Colo.: Westview Press.
- Arabic*. Socrates Spiro (1974), *An English-Arabic Dictionary of the Colloquial Arabic of Egypt*. Beirut: Librairie du Liban.
- Armenian*. Adour H. Yacoubian (1944), *English-Armenian and Armenian-English Concise Dictionary*. Los Angeles: Armenian Archives Press.

*Atakapa.* Albert S. Gatschet and John R. Swanton (1932), *A Dictionary of the Atakapa Language*. Bureau of American Ethnology, Bulletin 108.

*Azande.* E. E. Evans-Pritchard (1963), *Notes on Some Animals in Zandeland*. *Man* 63:139-142.

*Bikol.* Malcolm W. Mintz (1971), *Bikol Dictionary, PALI Language Texts: Philippines*. Honolulu: University Press of Hawaii.

*Biloxi.* James Owen Dorsey and John R. Swanton (1912), *A Dictionary of the Biloxi and Ofo Languages*. Bureau of American Ethnology, Bulletin 47.

*Bontok Igorot.* Walter Clayton Clapp (1908), *A Vocabulary of the Igorot Languages as Spoken by the Bontok Igorots*. Bureau of Science, Division of Ethnology Publications, Manila: Bureau of Printing.

*Cayapa.* John N. Lindscoog and Carrie A. Lindscoog (1964), *Vocabulario Cayapa*. Quito: Instituto Lingüístico de Verano.

*Chamorro.* Donald M. Topping, Pedro M. Ogo, and Bernadita C. Dungca (1975), *Chamorro-English Dictionary, PALI Language Texts: Micronesia*. Honolulu: University Press of Hawaii.

*Chinese.* Investigated by author; Janey Chen (1970), *A Practical English-Chinese Pronouncing Dictionary*. Rutland, VT: Charles E. Tuttle Co.; R. H. Mathews (1960), *Mathews' Chinese-English Dictionary*, Cambridge, Mass.: Harvard University Press.

*Choctaw.* Cyrus Byington (1915), *A Dictionary of the Choctaw Language*. John R. Swanton and Henry S. Halbert, eds. Bureau of American Ethnology, Bulletin 46.

*Chorti.* Charles Wisdom (1949), *Materials on the Chortí Language*. Microfilm Collection of Manuscripts in Middle American Cultural Anthropology, Number 28. Chicago: University of Chicago Library.

*Chrau.* Dorothy Thomas (1966), *Chrau Zoology: An Ethnolinguistic Study*. In *Papers on Four Vietnamese Languages*. David D. Thomas, ed. pp. 1-14. Auckland: Linguistic Society of New Zealand.

*Classical Nahuatl.* Alonso de Molina and Antonia de Spinosa (1966), *Vocabulario Nahuatl-Castellano, Castellano-Nahuatl (segunda edición: abreviada y modernizada, primera edición 1571)*. México: Ediciones Colofon.

*Congo.* Henry Craven and John Barfield (1971), *English-Congo and Congo-English Dictionary*. Freeport, N.Y.: Books for Libraries Press.

*Cornish.* Fred W. P. Jago (1887), *An English-Cornish Dictionary*. London: Simpkin, Marshall; R. Morton Nance (1971), *A Cornish-English Dictionary*, Cornish Language Board; R. Morton Nance (1973), *An English-Cornish Dictionary*, Cornish Language Board.

*Crow.* Robert H. Lowie (1960), *Crow Word Lists*. Berkeley: University of California Press.

*Czech.* Supplied to author by Mark Flotow; Alois Čermák (1963), *English-Czech, Czech-English Dictionary*. New York: Saphrograph.

*Delaware.* Jay Miller (1975), *Delaware Alternative Classifications*. *Anthropological Linguistics* 17:434-444; David Zeisberger (1887), *Zeisberger's Indian Dictionary*. Cambridge: John Wilson and Son.

*Diegueño.* Ted Couro and Christina Hutcheson (1973), *Dictionary of Mesa Grande Diegueño: ṽlṽpay Aa-English/English-ṽlṽpay Aa*. Banning, Calif.: Malki Museum Press.

*Dyola.* Le R. P. Wintz, ed. (1909), *Dictionnaire Français-Dyola et Dyola-Français*. Elinkine (Casamance): Mission Catholique.

*Efik.* Hugh Goldie (1964), *Dictionary of the Efik Language*. Farnborough, England: Gregg Press.

*Fijian.* A. Capell (1973), *A New Fijian Dictionary*. Suva, Fiji: Government Printer.

*Finnish.* Raija Hurme (1973), *English-Finnish General Dictionary*. Werner Söderströ Osakeyhtiö: Maritta Pesonen.

*Fore.* J. M. Diamond (1966), *Zoological Classification System of a Primitive People*. *Science* 151:1,102-1,104.

*Galla.* E. C. Foot (1913), *A Galla-English, English-Galla Dictionary*. Cambridge, England: University Press.

*Gimi.* Leonard B. Glick (1964), *Categories and Relations in Gimi Natural Science*. *American Anthropologist* 66 (Number 4, Part 2, special publication): 273-280.

*Hawaiian.* Mary Kawena Pukui and Samuel H. Elbert (1957), *Hawaiian-English Dictionary*.



Honolulu: University Press of Hawaii; Mary Kawena Pukui and Samuel H. Elbert (1964), *English-Hawaiian Dictionary*. Honolulu: University Press of Hawaii.

*Hill Pandaram*. Brian Morris (1976), *Whither the Savage Mind? Notes on the Natural Taxonomies of a Hunting and Gathering People*. *Man* 11:542-557.

*Huastec*. Investigated by author.

*Huave*. Jorge A. Suárez (1975), *Estudios Huaves*. México, D. F.: Instituto Nacional de Antropología e Historia; Milton Warkentin and Clara Warkentin (1952), *Vocabulario Huave*. México, D. F.: Instituto de Lingüístico de Verano.

*Huitoto Muinane*. Eugene E. Minor and Dorothy Hendrich de Minor (1971), *Vocabulario Huitoto Muinane*. Yarinacocha, Perú: Instituto Lingüístico de Verano.

*Hungarian*. L. Országh (1957), *Hungarian-English Dictionary*. Brooklyn: K. P. Schick; L. Országh (1957), *A Dictionary of the English and Hungarian Languages*. Brooklyn: K. P. Schick.

*Igbo*. Northcote W. Thomas (1913), *Anthropological Report on the Ibo-Speaking Peoples of Nigeria, Part II. English-Ibo and Ibo-English Dictionary*. London: Harrison and Sons; Beatrice F. Welmers and William E. Welmers (1968), *Igbo: A Learner's Dictionary*. Los Angeles: University of California, Los Angeles, and the United States Peace Corps.

*Indonesian*. Investigated by author.

*Irish*. Tomás de Bhaldraithe (1959), *English-Irish Dictionary*. Oifig An tSoláthair: Baile Átha Cliath; Patrick S. Dinneen (1927), *An Irish-English Dictionary*. Dublin: Irish Texts Society.

*Iroquois*. David Zeisberger (1887), *Zeisberger's Indian Dictionary*. Cambridge: John Wilson and Son.

*Italian*. Mario Hazon (1961), *Garzanti Comprehensive Italian-English, English-Italian Dictionary*. New York: McGraw Hill Book Company, Inc.

*Jacalteco*. Christopher Day (1977), *Diccionario Jacalteco-Español: The Vocabulary of a Mayan Language of Guatemala*. Unpublished manuscript. Department of Anthropology, University of Rochester.

*Japanese*. Andrew Nathaniel Nelson (1962), *The Modern Reader's Japanese-English Character Dictionary*. Rutland, VT: Charles E. Tuttle Co.; M. Takahashi (1973), *Romanized English-Japanese Dictionary*. Tokyo: Taiseido Shobo.

*Kalam*. Ralph N. H. Bulmer (1974), *Folk Biology in the New Guinea Highlands*. *Social Science Information* 13(4/5): 9-28; Ralph N. H. Bulmer and J. I. Menzies (1972-73), *Karam Classification of Marsupials and Rodents, Parts I and II*. *Journal of the Polynesian Society* 81:472-499 and 82:86-107; Ralph N. H. Bulmer, J. I. Menzies, and F. Parker (1975), *Kalam Classification of Reptiles and Fishes*. *Journal of the Polynesian Society* 84:267-308; Ralph N. H. Bulmer and M. J. Tyler (1968), *Karam Classification of Frogs*. *Journal of the Polynesian Society* 77:333-385.

*Karok*. Jane O. Bright and William Bright (1965), *Semantic Structures in Northwestern California and the Sapir-Whorf Hypothesis*. *American Anthropologist* 67:249-258.

*Kayan*. Supplied to author by Robert H. McDaniel.

*Kekchi*. Guillermo Sedat S. (1955), *Nuevo Diccionario de las Lenguas K'ekchi' y Española*, Chamelco. Alta Verapaz, Guatemala, C.A.: Instituto Indigenista Nacional.

*Kikuyu*. A. Ruffell Barlow and T. G. Benson (1975), *English-Kikuyu Dictionary*. Oxford, England: Clarendon Press.

*Kiowa*. John P. Harrington (1928), *Vocabulary of the Kiowa Language*, Bureau of American Ethnology, Bulletin 84.

*Kusaiean*. Kee-Dong Lee (1976), *Kusaiean-English Dictionary*. Honolulu: University Press of Hawaii.

*Kyaka Enga*. Supplied to author by Ralph N. H. Bulmer.

*Lake Miwok*. Catherine A. Callaghan (1965), *Lake Miwok Dictionary*. Berkeley: University of California Press.

*Luiseno*. William Bright (1968), *A Luiseno Dictionary*. Berkeley: University of California Press.

*Maidu*. William F. Shipley (1963), *Maidu Texts and Dictionary*. Berkeley: University of California Press.

*Maranao*. Batua A. Macaraya, Hadji Pambaya Bayabao, and David L. Hamm (1975), *The Dansalan College Maranao-English Dictionary*. Marawi City, Philippines: Dansalan Research Center, Dansalan College; Howard P. McKaughan and Batua A. Macaraya (1967), *A Maranao Dictionary*. Honolulu: University Press of Hawaii.

*Marshallese.* Takaji Abo, Byron W. Bender, Alfred Capelle, and Tony DeBrum (1976), *Marshallese-English Dictionary*. Honolulu: University Press of Hawaii.

*Mende.* Gordon Innes (1969), *A Mende-English Dictionary*. Cambridge, England: University Press; F. W. H. Migeod (1908), *The Mende Language*. London: Kegan Paul, Trench, Trübner and Co.

*Mexicano.* Forrest Brewer and Jean G. Brewer (1971), *Vocabulario Mexicano de Tetelcingo*. Morelos, México, D.F.: Instituto Lingüístico de Verano.

*Mixe.* Alvin Schoenhals and Louise C. Schoenhals (1965), *Vocabulario Mixe de Totontepec*. México: Instituto Lingüístico de Verano.

*Mixtec.* Anne Dyk and Betty Stoudt (1973), *Vocabulario Mixteco de San Miguel el Grande*. México, D.F.: Instituto Lingüístico de Verano; Brenda J. Pensinger (1974), *Diccionario Mixteco-Español, Español-Mixteco*. México, D.F.: Instituto Lingüístico de Verano.

*Mokilese.* Barbara Demory (1972), *A Preliminary Survey of the Ethnobiology of a Micronesian Atoll: Mokil or "A Rat is a Hairy Lizard."* Unpublished manuscript. Department of Anthropology, University of California, Berkeley.

*Mongolian.* John G. Hangin (1970), *A Concise English-Mongolian Dictionary*. The Hague: Mouton.

*Muyuw.* Daphne Lithgow and David Lithgow (1974), *Dictionaries of Papua New Guinea Volume 1, Muyuw Language*. Ukarumpa, Papua New Guinea: Summer Institute of Linguistics.

*Natick.* James Hammond Trumbull (1903), *Natick Dictionary*. Bureau of American Ethnology, Bulletin 25.

*Navaho.* Berard Haile (1951), *A Stem Vocabulary of the Navaho Language*. St. Michaels, Ariz.: St. Michaels Press; Norma Perchonock and Oswald Werner (1969), *Navaho Systems of Classification: Some Implications for Ethnoscience*. *Ethnology* 8:229-242; Oswald Werner, Allan Manning, and Kenneth Y. Begishe (n.d.), *A Taxonomic View of the Traditional Navajo Universe*. Unpublished manuscript. Department of Anthropology, Northwestern University; Leland C. Wyman (1964), *Navaho Indian Ethnoentomology*, University of New Mexico Publications in Anthropology, Number 12. Albuquerque: University of New Mexico.

*Ndumba.* Supplied to author by Terence E. Hays.

*Northern Paiute.* Catherine S. Fowler and Joy Leland (1967), *Some Northern Paiute Native Categories*. *Ethnology* 6:381-404.

*Nukuoro.* Vern Caroll and Tobias Soulik (1973), *Nukuoro Lexicon, PALI Language Texts: Polynesia*. Honolulu: University Press of Hawaii.

*Nungubuyu.* Jeffrey Heath (1978), *Linguistic Approaches to Nungubuyu Ethnobotany and Ethnobotany*. In *Australian Aboriginal Concepts*. L. R. Hiatt, ed. pp. 40-55. Canberra: Australian Institute of Aboriginal Studies.

*Ocaina.* Ilo M. Leach (1969), *Vocabulario Ocaina*. Perú: Instituto Lingüístico de Verano.

*Ojibwa.* R. R. Bishop Baraga (1973), *A Dictionary of the Otchipwe Language* (originally published in 1878 by Ross and Haines). Minneapolis: Ross and Haines; Mary B. Black (1969), *Eliciting Folk Taxonomies in Ojibwa*. In *Cognitive Anthropology*. Stephen A. Tyler, ed. pp. 165-189. New York: Holt, Rinehart and Winston.

*Osage.* Francis La Flesche (1932), *A Dictionary of the Osage Language*. Bureau of American Ethnology, Bulletin 109.

*Pahlavi.* D. N. MacKenzie (1971), *A Concise Pahlavi Dictionary*. London: Oxford University Press.

*Palauan.* Edwin G. McManus (1977), *Palauan-English Dictionary*. Honolulu: University Press of Hawaii.

*Pali.* A. P. Buddhadatta Mahatera (1955), *English-Pali Dictionary*. Pali Text Society.

*Pangasinan.* Richard A. Benton (1971), *Pangasinan Dictionary*. Honolulu: University Press of Hawaii.

*Papago-Pima.* Joseph B. Casagrade and Kenneth L. Hale (1967), *Semantic Relationships in Papago Folk-Definitions*. In *Studies in Southwestern Ethnolinguistics: Meaning and History in the Languages of the American Southwest*. Dell H. Hymes, ed. pp. 165-193. The Hague: Mouton; Dean Saxton and Lucille Saxton (1969), *Dictionary: Papago and Pima to English and English to Papago and Pima*. Tucson: University of Arizona Press.

*Pashto.* Investigated by author; Henry Walter Bellew (1901), *A Dictionary of the Pukshto or Pukshto Language*. Lahore: Rai Sahib M. Gulab Singh and Sons.

- Pintupi*. K. C. Hansen and L. E. Hansen (1974), *Pintupi Dictionary*. Darwin, Australia: Summer Institute of Linguistics.
- Puget Salish*. Thom Hess (1976), *Dictionary of Puget Salish*. Seattle: University of Washington Press.
- Quichua*. Carolyn Orr and Betsy Wrisley (1965), *Vocabulario Quichua del Oriente del Ecuador*. Quito, Ecuador: Instituto Lingüístico de Verano; Guillermo Escobar Risco (1951), *Vocabulario y Phrasis en la Lengua General de Los Indios del Perú, Llamada Quichua*. Lima: Instituto de Historia de la Facultad de Letras.
- Rofaifo*. Supplied to author by Peter D. Dwyer; Peter D. Dwyer (1976), *An Analysis of Rofaifo Mammal Taxonomy*. *American Ethnologist* 3:425-445.
- Serbo-Croatian*. Morton Benson (1971), *Serbocroatian-English Dictionary*. Philadelphia: University of Pennsylvania Press; Svetomir Ristić and Živojin Simić (1959), *An English-Serbocroatian Dictionary*. Beograd: Prosveta.
- Shoshoni*. Per Hage and Wick R. Miller (1976), "Eagle" = "Bird": A Note on the Structure and Evolution of Shoshoni Ethnoornithological Nomenclature. *American Ethnologist* 3:481-488; Wick R. Miller (1972), *Newe Natekwinapphe: Shoshoni Stories and Dictionary*. University of Utah Anthropological Papers, Number 94. Salt Lake City: University of Utah Press.
- Smith River*. Jane O. Bright and William Bright (1965), *Semantic Structures in Northwestern California and the Sapir-Whorf Hypothesis*. *American Anthropologist* 67:249-258.
- Tagalog*. Leo James English (1965), *English Tagalog Dictionary, Republic of the Philippines: Department of Education*; Maria Odulio de Guzman (1966), *An English-Tagalog and Tagalog-English Dictionary*. Manila: G.O.T.
- Tarahumara*. K. Simon Hilton (1959), *Tarahumara y Español*. México, D.F.: Instituto Lingüístico de Verano; Andrés Lionnet (1972), *Los Elementos de la Lengua Tarahumara*. México: Universidad Nacional Autónoma de México.
- Tausug*. Irene Hassan, Nurhadan Italud, Seymour Ashley, and Lois Ashley (1975), *Tausug-English Dictionary*. Manila, Philippines: Summer Institute of Linguistics.
- Tequistlatec*. Paul Turner and Shirley Turner (1971), *Dictionary: Chontal to Spanish-English, Spanish to Chontal*. Tucson: University of Arizona Press.
- Tewa*. Junius Henderson and John Peabody Harrington (1914), *Ethnozoology of the Tewa Indians*. Bureau of American Ethnology, Bulletin 56.
- Thai*. S. J. Tambiah (1969), *Animals Are Good to Think and Good to Prohibit*. *Ethnology* 8:423-459.
- Tibetan*. H. A. Jäschke (1881), *A Tibetan-English Dictionary*. New York: Frederick Ungar.
- Tifal*. Phyllis M. Healey and Walter Steinkrans (1972), *A Preliminary Vocabulary of Tifal with Grammar Notes, Language Data Microfiche AP2*. Santa Ana, California: Summer Institute of Linguistics.
- Tiruray*. Stuart A. Schlegel (1971), *Tiruray-English Lexicon*. Berkeley: University of California Press.
- Totonac*. Herman Pedro Aschmann (1973), *Diccionario Totonaco de Papantla*. Veracruz, México, D.F.: Instituto Lingüístico de Verano; Aileen A. Reid and Ruth G. Bishop (1974), *Diccionario Totonaco de Xicotepc de Juárez*. Puebla, México, D.F.: Instituto Lingüístico de Verano.
- Turkish*. H. C. Hony (1947), *A Turkish-English Dictionary*. London: Oxford University Press; Fahir Iz and H. C. Hony (1952), *An English-Turkish Dictionary*. London: Oxford University Press.
- Tzeltal*. Supplied to author by Brian Stross; Eugene S. Hunn (1975), *The Tenejapa Tzeltal Version of the Animal Kingdom*. *Anthropological Quarterly* 48:14-30; Eugene S. Hunn (1977), *Tzeltal Folk Zoology: The Classification of Discontinuities in Nature*. New York: Academic Press.
- Tzotzil*. Robert M. Laughlin (1975), *The Great Tzotzil Dictionary of San Lorenzo Zinacantán*. Smithsonian Contributions to Anthropology, No. 19.
- Wappo*. Jesse O. Sawyer (1965), *English-Wappo Vocabulary*. Berkeley: University of California Press.
- Welsh*. Thomas Jones (1811), *An English and Welsh Dictionary*. Denbigh, Wales: Thomas Gee; Jac L. Williams (1973), *Geiriadur Terman—Dictionary of Terms*. Cardiff: University of Wales Press.
- Western Apache*. Staff of the White Mountain Apache Culture Center (1972), *Western Apache Dictionary, White Mountain Apache Tribe*.

*Yana.* Edward Sapir and Morris Swadesh (1960), *Yana Dictionary*. Mary R. Haas, ed. Berkeley: University of California Press.

*Yapese.* John Thayer Jensen (1977), *Yapese-English Dictionary*. Honolulu: University Press of Hawaii.

*Yareba.* Harry Weimer and Natalia Weimer (1974), *Dictionaries of Papua New Guinea Volume 2*. Ukarumpa, Papua New Guinea: Summer Institute of Linguistics.

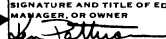
*Yoruba.* Samuel Crowther (1913), *A Dictionary of the Yoruba Language*. London: Oxford University Press.

*Yurok.* Jane O. Bright and William Bright (1965), *Semantic Structures in Northwestern California and the Sapir-Whorf Hypothesis*. *American Anthropologist* 67:249-258.

*Zapotec.* Velma Pickett (1973), *Vocabulario Zapoteco del Istmo*. México, D.F.: Instituto Lingüístico de Verano.

*Zoque.* Raul de la Grasserie (1898), *Langu Zoque et Langu Mixe*. Paris: Bibliothèque Linguistique Américaine, 22; Roy W. Harrison and Margarita B. Harrison (1948), *Diccionario Español-Zoque y Zoque-Español*. México, D.F.: Instituto Lingüístico de Verano.

*Zulu.* C. M. Doke, D. McK. Malcolm, J. M. A. Sikakana (1958), *English-Zulu Dictionary*. Johannesburg: Witwatersrand University Press; C. M. Doke and B. W. Vilakazi (1953), *Zulu-English Dictionary*. Johannesburg: Witwatersrand University Press.

U.S. POSTAL SERVICE STATEMENT OF OWNERSHIP, MANAGEMENT AND CIRCULATION (Required by 39 U.S.C. 3685)		
1. TITLE OF PUBLICATION AMERICAN ANTHROPOLOGIST		2. DATE OF FILING 10/1/79
3. FREQUENCY OF ISSUE QUARTERLY		4. ANNUAL SUBSCRIPTION PRICE \$40.00
5. LOCATION OF KNOWN OFFICE OF PUBLICATION (Street, City, County, State and ZIP Code) (Not printers) 1703 NEW HAMPSHIRE AVENUE, NW WASHINGTON, DC 20009		
6. NAMES AND COMPLETE ADDRESSES OF PUBLISHER, EDITOR, AND MANAGING EDITOR		
PUBLISHER (Name and Address) AMERICAN ANTHROPOLOGICAL ASSOCIATION, SEE #4		
EDITOR (Name and Address) D. L. OLMSTED, SEE #4		
MANAGING EDITOR (Name and Address) KEN PATTERSON, SEE #4		
7. OWNER (If owned by a corporation, its name and address must be stated and also immediately thereunder the names and addresses of stockholders owning or holding 1 percent or more of total amount of stock. If not owned by a corporation, the names and addresses of the individual owners must be given. If owned by a partnership or other unincorporated firm, its name and address, as well as that of each individual must be given. If the publication is published by a nonprofit organization, its name and address must be stated.)		
NAME	ADDRESS	
AMERICAN ANTHROPOLOGICAL ASSOCIATION	SAME AS #4	
8. KNOWN BONDHOLDERS, MORTGAGEES, AND OTHER SECURITY HOLDERS OWNING OR HOLDING 1 PERCENT OR MORE OF TOTAL AMOUNT OF BONDS, MORTGAGES OR OTHER SECURITIES (If there are none, so state)		
NAME	ADDRESS	
NONE		
9. FOR COMPLETION BY NONPROFIT ORGANIZATIONS AUTHORIZED TO MAIL AT SPECIAL RATES (Section 132.122, PSN) The purpose, function, and nonprofit status of this organization and the exempt status for Federal income tax purposes (Check one)		
<input checked="" type="checkbox"/> HAVE NOT CHANGED DURING PRECEDING 12 MONTHS <input type="checkbox"/> HAVE CHANGED DURING PRECEDING 12 MONTHS (If changed, publisher must submit explanation of change with this statement.)		
10. EXTENT AND NATURE OF CIRCULATION	AVERAGE NO. COPIES EACH ISSUE DURING PRECEDING 12 MONTHS	ACTUAL NO. COPIES OF SINGLE ISSUE PUBLISHED NEAREST TO FILING DATE
A. TOTAL NO. COPIES PRINTED (Net Press Run)	12,500	12,500
B. PAID CIRCULATION 1. SALES THROUGH DEALERS AND CARRIERS, STREET VENDORS AND COUNTER SALES	0	0
2. MAIL SUBSCRIPTIONS	11,296	11,301
C. TOTAL PAID CIRCULATION (Sum of 10B1 and 10B2)	11,296	11,301
D. FREE DISTRIBUTION BY MAIL, CARRIER OR OTHER MEANS SAMPLES, COMPLIMENTARY, AND OTHER FREE COPIES	20	20
E. TOTAL DISTRIBUTION (Sum of C and D)	11,316	11,321
F. COPIES NOT DISTRIBUTED 1. OFFICE USE, LEFT OVER, UNACCOUNTED, SPOILED AFTER PRINTING	1,184	1,179
2. RETURNS FROM NEWS AGENTS	0	0
G. TOTAL (Sum of E, F1 and 2—should equal net press run shown in A)	12,500	12,500
11. I certify that the statements made by me above are correct and complete.	SIGNATURE AND TITLE OF EDITOR, PUBLISHER, BUSINESS MANAGER, OR OWNER  MANAGING EDITOR	