

Zooplankton

TZMAX

55. TZMAX is the maximum ingestion rate for zooplankton (1/day). The zooplankton compartment includes the groups Cladocera, Copepoda, and Rotatoria which are classified as either herbivores or as carnivores.

56. Two types of feeding behavior exist: filter feeding and grasping feeding. Daphnia and some copepods are filter feeders. They collect particulate matter, including algae and detritus, by sieving lake water through the fine meshes of their filtering apparatus (Jorgensen 1975). Algae are swept into the feeding appendages to the mouth region where they are ingested as boluses containing many cells. Filter-feeding zooplankton make up the greater proportion of the zooplankton community and have been studied in greater detail.

57. The filtering rate per animal decreases as food concentration **increases**; above a critical concentration of food, the feeding rate is independent of food concentration.

58. Factors that influence food consumption by filter-feeding zooplankton include (a) animal density, size, sex, reproductive state, nutritional or physiological state as well as (b) the type, quality, concentration, and particle size of food. Other factors include water quality and temperature.

59. A second type of feeding behavior, raptorial or grasping feeding, is exhibited by most copepods and some cladocerans. They pursue prey and grasp large particles, including algae and detritus. Apparently, some copepods can switch feeding modes.

60. Several experiments have been able to demonstrate a maximum grazing rate allowing for long-term acclimation to food concentration above the incipient limiting level. Values for TZMAX range from 0.045 to 3.44 l/day.

61. Dissolved organic matter (DOM) is another potential source of food for zooplankters, although this feeding transfer is not modeled in CE-QUAL-R1. values for maximum ingestion rates for zooplankton are given in Table 12.

Table 12
Maximum ingestion rates for zooplankton (l/day)

| <u>PREDATOR</u> | <u>VALUE</u> | <u>FOOD SOURCE</u> | <u>REFERENCE</u> |
|-------------------------------|--------------|--------------------------|--|
| Bosmina | 0.01 | detritus | Bogdan and McNaught 1975 |
| Brachionus rubens | 3.438 | Chlorella vulgaris | Pilarska 1977 |
| Cladocerans | 0.15 | detritus | Bogdan and McNaught 1975 |
| Copepods | 0.10 | detritus | Bogdan and McNaught 1975 |
| Daphnia | 0.01 | detritus | Bogdan and McNaught 1975 |
| Daphnia magna | 0.251 | Saccharomyces cervisiae | McMahon and Rigler 1965 |
| Daphnia magna | 0.452 | Tetrahymena pyriformis | McMahon and Rigler 1965 |
| Daphnia magna | 0.301 | Chlorella vulgaris | McMahon and Rigler 1965 |
| Daphnia magna | 0.045 | Escherichia coli | McMahon and Rigler 1965 |
| Daphnia magna | 0.760 | Chiarella vulgaris | Kersting and Van De Leeuw-Leeqwater 1976 |
| Daphnia magna | 0.350 | Saccharomyces cerevisiae | Rigler 1961 |
| Daphnia magna | 1.. | Chiarella vulgaris | Ryther 1954 |
| Daphnia magna | 2.2 | Navicula pelliculosa | Ryther 1954 |
| Uaphnia magna | 2.3 | Scenedesmus quadricauda | Ryther 1954 |
| Daphnia pulex | 0.120 | Chlorococcwn sp. | Monokov and Sorokin 1961 |
| Daphnia rosea | 0.900 | Rhodotaru1a glutinis | Burns and Rigler 1967 |
| Diaptomus | 0.47 | detritus | Bogdan and McNaught 1975 |
| IN SITU EXPERIMENTS | | | |
| Heart Lake, Canada | 0.801 | Various | Haney 1973 |
| Lake Vechten, The Netherlands | 0.24 | Various | Gulati 1978 |
| Lake Krasnoye, USSR | 1.20 | various | Andronikova 1978 |

TZMORT

62. TZMORT is the maximum nonpredatory mortality rate for zooplankton (1/day). Nonpredatory mortality rate may be obtained by measuring total mortality and predatory mortality and subtracting to obtain the difference (a direct approach is to measure mortality rate and eliminate predators altogether). Nonpredatory mortality may be influenced by oxygen concentration, temperature, diet, age, and population density. Nonpredatory mortality rates are normally less than 1 percent per day. Values for maximum nonpredatory mortality rate are given in Table 13.

Table 13
Zooplankton mortality rates (1/day)

| <u>SPECIES</u> | <u>TZHORT</u> | <u>REFERENCE</u> |
|--------------------------------------|---------------|--------------------------|
| <i>Calanus helgolandicus</i> | 0.003-0.048 | Paffenhofer 1976 |
| <i>Calanus nelgolandicus</i> | 0.024 | Mullin and Brooks 1970 |
| Carnivorous zooplankton | 0.002-0.013 | Petipa et al. 1970 |
| <i>Ceriodaphnia reticulata</i> | 0.0016 | Clark and Carter 1974 |
| Copepod nauplii | 0.006-0.017 | Petipa et al. 1970 |
| <i>Daphnia galeata</i> | 0.017 | Hall 1964 |
| <i>Daphnia pulex</i> | 0.012 | Craddock 1976 |
| <i>Daphnia pulex</i> | 0.018-0.027 | Frank et al. 1957 |
| <i>Daphnia retrocurva</i> | 0.001 | Clark and Carter 1974 |
| <i>Daphnia rosea</i> | 0.001-0.007 | Dodson 1972 |
| <i>Daphnia rosea</i> | 0.001 | Clark and Carter 1974 |
| <i>Daphnia</i> spp. | 0.002 | Wright 1965 |
| <i>Diaptomus clavipes</i> | 0.004-0.155 | Gehrs and Robertson 1975 |
| <i>Diaphanosoma leuchtenbergiana</i> | 0.001 | Clark and Carter 1974 |
| Omnivorous zooplankton | 0.010-0.013 | Petipa et al. 1970 |
| <i>Paracalanus</i> sp. | 0.003-0.006 | Petipa et al. 1970 |
| <i>Rhincalanus nasutus</i> | 0.006-0.015 | Mullin and Brooks 1970 |
| <i>Sirnocephalus serrulatus</i> | 0.003 | Hall et al. 1970 |

ZEFFIC

63. ZEFFIC, the zooplankton assimilation efficiency (A/G) {dimensionless}, is the proportion of food consumed (G) to food assimilated (A), i.e., food actually absorbed from an individual's digestive system. The assimilation efficiency is used to modify consumption and to determine the quantity of energy entering an individual or population.

64. Of the factors affecting assimilation efficiency, the most significant is food type. For herbivores-detritivores, the range in ZEFFIC is wide because these animals often consume foods of varying energy content and digestibility. Among the carnivores, for which food type varies little, A/G ranges between 0.80 and 0.95. Values for zooplankton assimilation efficiency are given in Table 14.

Table 14
Zooplankton assimilation efficiency coefficients (dimensionless)

| <u>SPECIES</u> | <u>ZEFFIC</u> | <u>REFERENCE</u> |
|---------------------------------|---------------|------------------------------------|
| <i>Acartia clausi</i> | 0.66-0.73 | Penchen'-Finenko 1977 |
| <i>Bosmina coregoni</i> | 0.09-0.77 | Semenova 1914 |
| <i>Bosmina longirostris</i> | 0.32-0.31 | Gute'l'mackher 1977 |
| <i>calanus firmarchicus</i> | 0.45-0.96 | Marshall and Orr 1956 |
| <i>Calamoezia lucase</i> | 0.63-0.67 | Green 1975 |
| <i>Ceriodaphnia reticulata</i> | 0.106 | Czeczuga & Bobiatynska-Ksok 1970 |
| <i>ceriodaphnia reticulata</i> | 0.47-0.73 | Czeczuga , Bobiatynska-Ksok 1970 |
| <i>Cyclops strennus</i> | 0.50 | Schindler 1971 |
| <i>cyclops vicimus</i> | 0.80 | Monakov 1972 |
| <i>Daphnia longispina</i> | 0.10-0.25 | Monakov , Sorokin 1961 |
| <i>Daphnia longispina</i> | 0.42 | Honakov 1972 |
| <i>Daphnia magna</i> | 0.60-0.84 | Schindler 1968 |
| <i>Daphnia pulex</i> | 0.14-0.31 | Richman 1958 |
| <i>Daphnia schodleri</i> | 0.60-0.90 | Hayward , Gallup 1976 |
| <i>Daphnia sp.</i> | 0.08-0.25 | Cohn 1958 |
| <i>Diaptomus graciloides</i> | 0.81 | Penchen'-Finenko 1977 |
| <i>Diaptomus graciloides</i> | 0.45-0.50 | Klekowski & Shushkina 1966 |
| <i>Diaptomus sicioides</i> | 0.40-0.83 | Comita 1972 |
| <i>Diaptomus oregonensis</i> | 0.71 | Richman 1964 |
| <i>Eury cercus lameillatic</i> | 0.07-0.32 | Smirnov 1962 |
| <i>Holopedium gibberum</i> | 0.10-0.47 | Gute'l'mackher 1977 |
| <i>Leptodora kindtii</i> | 0.40 | Cummins et al. 1969 |
| <i>Leptodora kindtii</i> | 0.87 | Hi1ibricht-I1kowska & Karabin 1970 |
| <i>Macro cyclops albidus</i> | 0.45-0.50 | Klekowski & Shushkina 1966 |
| <i>Mesocyclops albidus</i> | 0.20-0.75 | Klekowski & Shushkina 1966 |
| <i>Polyphemus pediculus</i> | 0.42 | Monakov 1972 |
| <i>Sida crystallima</i> | 0.11-0.99 | Monakov 1912 |
| <i>Simocephalus esplno5us</i> | 0.46 | Sorokin 1969 |
| <i>Simocephalus vetulus</i> | 0.31-0.72 | Klekowski 1970 |
| <i>Simocephalus vetulus</i> | 0.31-0.72 | Ivanova & Klekowski 1912 |
| 10 herbivores | 0.476 | Comita 1972 |

PREF1, PREF2, PREF3

65. All zooplankters are selective feeders resulting from a combination of (a) an organism's mechanical limitations in capturing and processing food items of varying size and configuration, (b) the chemical composition of the food items, and (c) feeding behavior. Food preference is demonstrated if an organism consumes a food item in a proportion different from the food item's relative contribution to the total of all available foods in the environment. If all foods occur at the same concentration, then the preference factors equal the fractions of ingestion contributed by each food compartment. Seasonal abundance of phytoplankton, bacteria, and detritus may be the main factor determining the percent composition of these components in the diets of many zooplankters.

66. Filamentous bluegreen algae are generally not considered to be as assimilable as are other algal species. They are seldom found in the guts of zooplankton, because they either are not eaten or are actively rejected. Most species of green algae and diatoms are filtered at about the same rate and digested. However, it is not necessarily the taxonomic position of the alga that makes it suitable or unsuitable as food, but rather the attributes of each algal species such as size, shape, and toxicity.

67. Although ample evidence exists to show that detritus is consumed by zooplankton, no evidence exists to show that it is consumed preferentially; rather, detritus is ingested in proportion to its composition in the environment. When detritus is included as a food source in a grazing formulation, it should be given equal ranking with other suitable foods. It should be noted that bacteria that colonize detritus constitute an important source of protein in the diet.

68. Filter feeders discriminate among particles on the basis of size, shape, and texture. There are upper and lower limits to the sizes of particles that can be managed by zooplankton feeding appendages. Particles of 0.8μ and larger can be retained; an upper limit is related to the size of the animal. Algae that clog the filtering appendages are rejected from them by a claw on the lower abdomen.

69. Raptorial feeders can seize large prey and tear it apart before eating (Ambler and Frost 1974, Brandl and Fernando 1975), but there are limits to the size of prey they capture.

70. PREF1 is the preference factor of zooplankton for the ALGAE1 compartment, PREF2 is the preference factor of zooplankton for the ALGAE2 compartment, and PREF3 is the preference factor of zooplankton for the detritus compartment. The food preference factors are dimensionless; the total of the three factors must equal 1. Values for these preference factors are given in Table 15.

Table 15
Food preference factors of zooplankton (dimensionless)

| PREDATOR | PREF | PREY | REFERENCE |
|-------------|------|-----------------|--------------------------|
| Bosmina | 0.33 | nannoplankton | Bogdan and McNaught 1975 |
| Bosmina | 0.33 | netplankton | Bogdan and McNaught 1975 |
| Cladocerans | 0.30 | nannoplankton | Bogdan and McNaught 1975 |
| Cladocerans | 0.30 | netplankton | Bogdan and McNaught 1975 |
| Cladocerans | 0.20 | bluegreen algae | Bogdan and McNaught 1975 |
| copepods | 0.45 | nannoplankton | Bogdan and McNaught 1975 |
| copepods | 0.15 | netplankton | Bogdan and McNaught 1975 |
| copepods | 0.20 | bluegreen algae | Bogdan and McNaught 1975 |
| Daphnia | 0.33 | nannoplankton | Bogdan and McNaught 1975 |
| Daphnia | 0.17 | netplankton | Bogdan and McNaught 1975 |
| Diaptomus | 0.40 | nannoplankton | Bogdan and McNaught 1975 |
| Diaptomus | 0.17 | netplankton | Bogdan and McNaught 1975 |

TZRESP

71. TZRESP is the maximum zooplankton respiration rate (1/day). Respiration is the sum of all physical and chemical processes by which organisms oxidize organic matter to produce energy. Respiration rates of aquatic invertebrates usually are estimated directly by monitoring oxygen consumption. By multiplying oxygen consumed times an oxycaloric coefficient {i.e., 4.83 cal/ml O₂ (Winberg et al. 1934» and the energy-to-carbon relation for aquatic invertebrates (*i.e.*, 10.98 cal/mg C (Salonen et al. 1976»), the amount of carbon metabolized can be determined and converted to biomass.

72. Conover (1960) has indicated that carnivores have higher respiration rates than herbivores. Values for maximum zooplankton respiration rates are given in Table 16.

Table 16
Zooplankton maximum respiration rates (1j day)

| <u>SPECIES</u> | <u>TZRESP</u> | <u>REFERENCE</u> |
|--------------------------------|---------------|--|
| <i>Bosmina coregoni</i> | 0.170 | Manuilova 1958 |
| <i>Bosmina longirostris</i> | 0.185 | Sushchenya 1958 |
| <i>Ceriodaphnia reticulata</i> | 0.18-.50 | Gophen 1976 |
| Copepoda | 0.075-.204 | Bishop, 1968 |
| Copepod adults | 0.043-.131 | Williams 1982 |
| Copepod copepodites | 0.054-.171 | Williams 1982 |
| Copepod nauplii | 0.165-.695 | Williams 1982 |
| Copepod total | 0.056-.183 | Williams 1982 |
| <i>Daphnia ashland!!</i> | 0.447-.74 | Duval and Geen 1976 |
| <i>Daphnia clavipes</i> | 0.117-.165 | Comita 1968 |
| <i>Daphnia cululata</i> | 0.161 | Manui10va 1958 |
| <i>Daphnia galeata</i> | 0.13-.772 | LaRow et al. 1975 |
| <i>Daphnia hyalina</i> | 0.179 | Blazka 1966 |
| <i>Daphnia longispina</i> | 0.121-.135 | Tezuka 1971 |
| <i>Daphnia longispina</i> | 0.16 | Manuilova 1958 |
| <i>Daphnia longispina</i> | 0.146 | Shushkina and Pecen' 1964 |
| <i>Daphnia magna</i> | 0.085-.175 | Kersting and Van De Leeuw-Leegwater 1976 |
| <i>Daphnia magna</i> | 0.014 | Sushchenya 1958 |
| <i>Daphnia oregonensis</i> | 0.194 | Richman 1964 |
| <i>Daphnia pulex</i> | 0.582 | Buikema 1972 |
| <i>Daphnia pulex</i> | 0.18-.19 | Tezuka 1971 |
| <i>Daphnia septopus</i> | 0.008-.18 | Comita 1968 |
| <i>Daphnia siciloides</i> | 0.006-.52 | Comita 1968 |
| <i>Diaphanosoma brachyurum</i> | 0.272 | Sushchenya 1958 |
| <i>Diaptomus kenai</i> | 0.272-.448 | Duval and Geen 1976 |
| <i>Leptodora kindtii</i> | 0.471 | Moshiri et al. 1969 |
| <i>Leptodora kindtii</i> | 0.125 | Hillbricht-Ilkowska and Karabin 1970 |
| <i>Simocephalus vetulus</i> | 0.131 | Sushchenya 1958 |
| <i>Simocephalus vetulus</i> | 0.154 | Manuilova 1958 |
| <i>Simocephalus vetulus</i> | 0.096-.201 | Ivanova and Klekowski 1972 |
| Total zooplankton | 0.063-.210 | Williams 1982 |

ZS2P

73. ZS2P is the zooplankton half-saturation coefficient for grazing on algae and detritus (mg/L). It has been found that zooplankton exhibit reduced feeding rates at high food concentrations; the relationship between feeding rate and food concentration has been reported to be curvilinear by a number of investigators (Burns and Rigler 1967, Parsons et al. 1967, McQueen 1970, Frost 1972, Mnakov 1972, Gaudy 1974, and Chisholm et al. 1975).

74. The most realistic calculation of zooplankton grazing rate is based on their rate of removal of biomass of food (Mullin 1963); therefore, it is important that investigators report results in terms of biovolume or biomass instead of cell number. The method most used to determine ingestion rate is to count prey in controls and experimental chambers after feeding zooplankton. Values for zooplankton HSC are given in Table 17.

Table 17
Zooplankton half-saturation coefficients (mg/L)

| SPECIES | ZS2P | REFERENCE |
|-----------------------|----------|-----------------------|
| Bosmina coregoni | 4.0 | Scavia and Eadie 1976 |
| Daphnia magna | 9.6-15.0 | Scavia and Eadie 1976 |
| Daphnia rosea | 0.16 | Scavia and Eadie 1976 |
| Diaptomus oregonensis | 1.6 | Scavia and Eadie 1976 |

ZOOT1, ZOOT2, ZOOT), ZOOT4

75. values for zooplankton temperature coefficients are given in Table 18.

- a. ZOOT1 is the lower temperature bound at which metabolism continues to occur. It is generally DOC.

- h. ZOOT2 is the lowest temperature at which processes are occurring near the maximum rate (0 C) .**
- c. ZOOT3 is the upper temperature bounding the range of maximum rates (QC).**
- d. ZOOT4 is the upper lethal temperature (QC).**

Table 18
Zooplankton temperature coefficients (DC)

| SPECIES | ZOOT1 | ZOOT2 | ZOOT3 | ZOOT4 | REFERENCE |
|--------------------------------|-------|-------|-------|-------|--|
| Calamoezia lusasi | NA' | 20 | 24 | NA | Green 1975 |
| Ceriodaphnia reticulata | NA | 2. | 27 | NA | Gophen 1976 |
| Daphnia galeata | NA | 20 | 24 | NA | Burns 1969 |
| Daphnia longispina | NA | 11 | 18 | NA | Nauwerck 1959 |
| Daphnia magna | NA | 24 | 2. | 35 | McMahon 1965 |
| Daphnia magna | NA | 25 | NA | NA | Burns 1969 |
| Daphnia middendorffiana | NA | 24 | 25 | NA | Kryutchkova and Kondratyuk 1966 |
| Daphnia pulex | NA | 20 | 24 | NA | Burns 1969 |
| Daphnia pulex | NA | 20 | 24 | NA | Geller 1975 |
| Daphnia pulex | NA | NA | 25 | NA | Geller 1975 |
| Daphnia rosea | NA | 20 | 24 | NA | Burns & Rigler 1967 |
| Daphnia rosea | NA | 14 | 15 | NA | Kibby 1971 |
| Daphnia schedleri | NA | 20 | 22 | NA | Burns 1969 |
| Daphnia schedleri | NA | 20 | 24 | NA | Hayward & Gallup 1976 |
| Diaptomus sp. | NA | 1. | 18 | NA | Nauwerck 1959 |

* NA not available.

76. As with the phytoplankton, zooplankton are able to adapt to the ambient temperature with time. This is demonstrable throughout the different regions of the United States and at different times of the year. Zooplankton found in temperate regions of the United States are exposed to lower average temperatures throughout the year and consequently have lower temperature factors (i.e., ZOOT1, ZOOT2, ZOOT3, and ZOOT4) than those found in more southern regions. Again, these values are unavailable from the literature but have been estimated by Leidy and Ploskey (1980) based upon acclimation temperatures (Table 19).

Table 19

Acclimation temperature, upper and lower lethal temperature, and the temperature range for a constant maximum grazing rate for zooplankton exposed to rapid temperature stress (OCJ
lfrom Leidy and Ploskey 1980)

| Acc 1. Temp. | ZOOT1 | ZOOT2 | ZOOT3 | ZOOT4 |
|-----------------|---------------|-------|-------|-------|
| 5 | 0 | 5 | 6 | 25 |
| 10 | 0 | 10 | 12 | 30 |
| 15 | 2 | 15 | 18 | 33 |
| 20 | 5 | 20 | 24 | 33 |
| 25 | 7 | 25 | 30 | 34 |
| 29 | 10 | 29 | 34 | 34 |
| 30 | 10 | 30 | 34 | 34 |
| 31 | 12 | 31 | 34 | 34 |
| 34 | 15 | 34 | 34 | 34 |
| 35 | Lethal | | | |

Table 20
Daily ration of benthic organisms (from Leidy and Ploskey 1980)
Cl/day)

| <u>SPECIES</u> | <u>FOOD</u> | <u>RATION</u> | <u>REFERENCE</u> |
|----------------------------------|----------------------------------|---------------|------------------------|
| NEMATODA | | | |
| <i>Aphelenchus avenae</i> | fungal mycelia | 0.26 | Soyza 1973 |
| <i>Plectus palustris</i> | <i>Acinetobacter</i> sp. | 6.50 | Duncan et al. 1974 |
| MOLLUSCA | | | |
| <i>Dreisena polymorpha</i> | bacteria | 0.01-.12 | SOrokin 1966 |
| <i>Goniobala clavaeformis</i> | aufwucks | 0.01-.24 | Malone and Nelson 1969 |
| ARTHROPODA | | | |
| <i>Hyalella & zteca</i> | sediments | 0.17- 1.03 | Hargrave 1970 |
| <i>Pontogammarus robustoides</i> | <i>Cladophora</i> sp. | 0.007-.98 | Xititsyna 1975 |
| <i>Pontogammarus robustoides</i> | <i>Tubifex</i> sp. | 0.187-1.63 | Xititsyna 1975 |
| PODOCOPA | | | |
| <i>Chaoborus flavicans</i> | natural phytoplankton population | 0.036-.114 | Kajak and Dusoqe 1970 |
| <i>Berpetocypris reptans</i> | <i>Spirogyra</i> sp. | 1.28 | Yakovleva 1969 |
| <i>Herpetocypris reptans</i> | <i>Zygnema</i> sp. | 0.93 | Yakovleva 1969 |
| <i>Herpetocypris reptans</i> | <i>Mougeotia</i> sp. | 0.93 | Yakovleva 1969 |
| <i>Herpetocypris reptans</i> | <i>Chironomus plwnosus</i> | 0 .66 | Yakovleva 1969 |
| <i>Herpetocypris reptans</i> | <i>Asellus aquaticus</i> | 0.66 | Yakovleva 1969 |
| <i>Herpetocypris reptans</i> | fish fry | 1.09 | Yakovleva 1969 |
| <i>Procladius choreus</i> | Chironomidae | 0.007-.11 | Kajak and DU60ge 1970 |
| EPHEMEROPTERA | | | |
| <i>Stenonema pulche11wn</i> | <i>Navicula minima</i> | 0.234 | Trama 1972 |
| PLECOPTERA | | | |
| <i>Acroneuria californica</i> | <i>Hydropsyche</i> sp. | 0.002-.087 | Heiman and Knight 1975 |

Benthos

TBMAX

77. TBMAX is the maximum ingestion rate for benthos (1/day) and is measured at food densities above the incipient limiting food concentration. The food source for this compartment is organic sediment; its dominant members for most reservoir benthic communities are the aquatic oligochaetes and Chironomidae. Filter feeders, predators, deposit feeders, and surface grazers are all represented in most benthic communities.

78. Daily rations (an approximation of the daily grazing rate) of some benthic species compiled by Leidy and Ploskey (1980) are listed in Table 20. Other values for maximum ingestion rate are given in Table 21.

Table 21
Benthos maximum ingestion rates (1/day)

| <u>SPECIES</u> | <u>TBMAX</u> | <u>REFERENCE</u> |
|----------------------------|--------------|------------------------|
| Acroneuria californica | 0.002-.09 | Heiman and Knight 1975 |
| Asellus aquaticus | 0.25 | Prus 1972 |
| Carnivores | 0.0282 | Bigelow et al 1977 |
| Chaoborus flavicans | 0.036-.114 | Kajak and Dusoge 1970 |
| Deposit feeder | 0.111 | Gordon 1966 |
| Hyalella azteca | 0.11-1.3 | Hargrave 1970 |
| Omnivores | 0.043 | Bigelow et al. 1977 |
| Pontagamrnarus robustiodes | 0.074-.98 | Kititsyna 1975 |
| Procladius choreus | 0.07-.11 | Kajak and Dusoge 1970 |
| Selective deposit feeder | 0.05 | Bigelow et al. 1977 |
| Stenonema pulchellum | 0.21-.23 | Trama 1972 |

TBMORT

79. TBMORT is the nonpredatory mortality rate for benthos (1/day). Leidy and Ploskey (1980), in their review of the literature, show most benthos nonpredatory mortality rates to be between 0.001 and 0.02/day.

BEFFIC

80. BEFFIC is the assimilation efficiency for benthos (dimensionless). The assimilation efficiency is multiplied by the ingestion rate to obtain an assimilation rate. Values for benthos assimilation efficiency are given in Table 22.

Table 22
Benthos assimilation efficiencies (dimensionless)

| <u>SPECIES</u> | <u>VALUE</u> | <u>REFERENCE</u> |
|---------------------------------|--------------|-----------------------------|
| <i>Anatopina dijari</i> | 0.30 | Teal 1957 |
| <i>Asellus aquaticus</i> | 0.30 | Klekowski 1970 |
| <i>Asellus aquaticus</i> | 0.26-0.44 | Prus 1971 |
| <i>Bandsiola crotchii</i> | 0.31-0.40 | Winterbourn 1974 |
| <i>Calopsectra dives</i> | 0.20 | Teal 1957 |
| Carnivores | 0.20-0.97 | Lawton 1970 |
| <i>Gammarus pseudolimnaeus</i> | 0.10-0.20 | Barlocher and Kendrick 1975 |
| <i>Gammarus pseudolimnaeus</i> | 0.42-0.75 | Barlocher and Kendrick 1975 |
| <i>Gammarus pseudolimnaeus</i> | 0.10 | Marchant and Hynes 1981 |
| <i>Gammarus pulex</i> | 0.30-0.40 | Nilsson 1974 |
| <i>Glossosoma nigrior</i> | 0.17-0.32 | Cummins 1973 |
| <i>Hedriodiscus</i> | 0.59 | Stockner 1971 |
| <i>Hyalella azeteca</i> | 0.05-0.80 | Hargrave 1970 |
| <i>Hydrophilus triangularis</i> | 0.55 | Hallmark and Ward 1972 |
| <i>Lepidostoma</i> | 0.07-0.12 | Grafius 1973 |
| <i>Lestes sponsa</i> | 0.36 | Klekowski et al. 1970 |
| <i>Lethocerus americanus</i> | 0.07 | Guthrie and Brust 1969 |
| <i>Limnodrilus hoffmeisteri</i> | 0.5 | Teal 1957 |
| Most invertebrates | 0.5 | Monakov 1972 |
| <i>Potamopyrgus jenkinsi</i> | 0.04 | Heywood and Edwards 1962 |
| <i>Potomophylax cingulatus</i> | 0.10-0.30 | Otto 1974 |
| <i>Pteronarcys scotti</i> | 0.11 | McDiffett 1970 |
| <i>Pyrrhosoma</i> | 0.77-0.91 | Lawton 1970 |
| <i>Simulium</i> | 0.57 | McCullough 1975 |
| <i>Stenonema</i> | 0.52 | Trama 1957 |
| <i>Tricorythodes minutus</i> | 0.07-0.55 | McCullough 1975 |
| <i>Tubifex tubifex</i> | 0.5 | Ivlev 1939 |

BS2SED

81. BS2SED is the half-saturation coefficient for benthos feeding on organic sediment (g/m^2). Leidy and Ploskey (1980), after a thorough review of the literature, wrote that they were unable to find a single reference that documented, in units convertible to carbon, the change in benthic grazing as a function of food concentration. In addition, the value of the coefficient depends on the depth of the sediment being modeled, which is itself a variable. The authors of the present report recommend using values slightly smaller than half the initial condition for the sediment, which is reported in g/m^2 .

TBRESP

82. TBRESP is the maximum respiration rate for benthos (l/day). Respiration rates are estimated directly by monitoring benthic oxygen consumption by manometric, chemical, or polarographic methods. Values for the respiration rate for benthos are given in Table 23.

Table 23
Maximum respiration rates for benthos II/day)

| <u>SPECIES</u> | <u>TBRESP</u> | <u>TEMP °C</u> | <u>REFERENCE</u> |
|-------------------------|---------------|----------------|-----------------------|
| Acartia | 0.129-.215 | NA" | Williams 1982 |
| Ancylus fluvialis | 0.035-.049 | " | Berg 1952 |
| Baetes sp. | 0.47-.72 | 10 | Fox et al. 1937 |
| Bithynia tentaculata | 0.020 | 13 | Berg & Ockelmann 1959 |
| Bithynia leachi | 0.031 | 13 | Berg & Ockelmann 1959 |
| Chironomus anthracinus | 0.005 | 11 | Berg et al. 1962 |
| Chironomus strenzkei | 0.12-.14 | 30 | Platzer-Schultz 1970 |
| Chloeon dipterum | 0.16-.46 | 10-16 | Fox and Simmonds 1933 |
| Coenis sp. | 0.075 | 10 | Fox et al. 1935 |
| Corethra flavicans | 0.002 | 11 | Berg et al. 1962 |
| Corycaeus | 0.051-.270 | NA | Williams 1982 |
| Echyonurus venosus | 0.17-.34 | 10 | Fox et al. 1935 |
| Ephemera simulans | 0.063 | 20 | Olson and Rueger 1968 |
| Ephemera vulgaris | 0.072-.19 | 10 | Fox et al. 1935 |
| Ephemera damica | 0.095-.21 | 10 | Fox et al. 1935 |
| Ephemerella ignita | 0.24 | 10 | Fox et al. 1935 |
| Erpobdella oculata | 0.034 | 20 | Mann 1956 |
| Erpobdella testacea | 0.052 | 20 | Mann 1956 |
| Gammarus pulex | 0.10-.12 | NA | Fox and Simmonds 1933 |
| Gastropoda, Veliger | 0.107 | NA | Williams 1982 |
| Glossiphonia complanata | 0.044 | 20 | Mann 1956 |
| Helobdella stagnalis | 0.052 | 20 | Mann 1956 |
| Ilyodrilus hammoniensis | 0.0009 | 11 | Berg et al. 1962 |
| Larvaceans | 0.014-.043 | NA | Williams 1982 |
| Lumbricillus rivalis | 0.006 | 11 | Berg et al. 1962 |
| Lymnaea aricularia | 0.016 | 13 | Berg & Ockelmann 1959 |
| Lymnaea palustris | 0.027 | 13 | Berg & Ockelmann 1959 |
| Lymnaea peregrina | 0.023 | 13 | Berg & Ockelmann 1959 |
| Many groups | 0.0001-.04 | NA | Olson and Rueger 1968 |
| Myxas glutinosa | 0.026 | 13 | Berg & Ockelmann 1959 |
| Oligotrichs | 0.257 | NA | Williams 1982 |
| Physa fontinalis | 0.041 | 13 | Berg & Ockelmann 1959 |
| piscicola geometra | 0.088 | 20 | Mann 1956 |
| Procladius sp. | 0.002 | 11 | Berg et al. 1962 |
| Tintinnids | 0.245 | NA | Williams 1982 |
| Tubifex barbatus | 0.005 | 11 | Berg et al. 1962 |
| Tubifex tubifex | 0.001 | 11 | Berg et al. 1962 |
| Valvata piscinalis | 0.041 | 13 | Berg & Ockelmann 1959 |

* NA = not available.

BENT1, BENT2, BENT3, BENT4

83. Values for benthos temperature coefficients are given in Table 24.

- a. BENT1 is the lower temperature bound at which metabolism continues to occur; it is usually 0 °e.
- b. BENT2 is the lowest temperature at which processes are occurring near the maximum rate.
- c. BENT3 is the upper temperature bounding the range of maximum rates.
- d. BENT4 is the upper lethal temperature.

Table 24

Temperature coefficients for benthos metabolism (OC)

| SPECIES | BENT1 | BENT2 | BENT3 | BENT4 | REFERENCE |
|--------------------------------|-------|-------|-------|-------|-----------------------|
| <i>Asellus aquaticus</i> | 0 | 15 | NA* | NA | Moore 1975 |
| <i>Gammarus pulex</i> | 0 | 18 | NA | NA | Moore 1975 |
| <i>Gammarus pseudolimnaeus</i> | 0 | 20 | NA | NA | Marchant & Hynes 1981 |

* NA not available.

Fish

84. CE-QUAL-R1 has three fish compartments for simulating piscivorous, planktivorous, and benthic-feeding assemblages in a reservoir. Since many fish species are omnivorous, however, the weighting procedure for computing composite compartment rates is different from other compartments. A report by Leidy and Jenkins (1977) provides all the information necessary to compute the required composite rate coefficients.

85. In the model, the piscivorous fish (compartment 1) feed only on the other two fish compartments. Fish in the second compartment feed on detritus, zooplankton, and the two algal groups; fish in the third compartment feed on

sediment and benthos.

TFMAX

86. **TFMAX,1** is the maximum ingestion rate (1/day) for the piscivorous fish compartment. The composite rate for the compartment should be computed based on the mean annual standing crop estimate. Ingestion rates vary as a function not only of species, but also of other factors such as condition or age class; the ingestion rate should reflect these factors by using, for example, average age class estimates.

87. **TFMAX,2** is the maximum ingestion rate for planktivorous fish (1/day). The planktivorous fish consume zooplankton, algae, and detritus.

88. **TFMAX,3** is the maximum ingestion rate for benthic fish (1/day). Benthic-feeding fish ingest both benthos and organic sediment.

89. In general, a TFMAX coefficient of 0.01 represents maintenance without growth; 0.04 to 0.05 represents optimum growth efficiency (Leidy and Jenkins 1977).

FS2BEN, FS2Z00, FS2FSH

90. To adjust the ingestion rate of fish due to the available food supply, the fishery model uses half-saturation constants; these represent the amount of food present that results in fish ingestion at half the maximum growth rate. It has been suggested that the half-saturation constant be considered to be 5 percent of fish wet body weight consumed per day at 20°C (Leidy and Jenkins 1977). Five percent of the body weight consumed per day corresponds closely with the food intake rate for optimum efficiency in growth (4 to 5 percent for many species). User's of CE-QUAL-RI should refer to Leidy and Jenkins (1977) because

of the difficulty in estimating half-saturation coefficients.
Estimates of fish half-saturation coefficients are given in
Table 25.

- a. FS2BEN is the benthic-feeding fishes' (FISH3) half-saturation coefficient for benthos and sediment grazing (mg/L).
- b. FS2Z00 is the planktivorous fishes' (FISH2) half-saturation coefficient for zooplankton, detritus, and algae (mg/L).
- c. FS2FSH is the piscivorous fishes' (FISH1) half-saturation coefficient for feeding on FISH3 and FISH2 (mg/L).

Table 25

Estimated half-saturation coefficients for fish growth (mg/L)
 (from Leidy and Jenkins 1977)

| <u>SPECIES</u> | <u>FOOD TYPE</u> | <u>VALUE</u> | <u>REFERENCE</u> |
|--------------------|------------------|--------------|---------------------------|
| Largemouth bass | minnows | .6 | Thompson 1941 |
| Smallmouth bass | minnows | 7.2 | Williams 1959 |
| Muskellunge | minnows | 5.6 | Gammon 1963 |
| Reticulate sculpin | midge larvae | 000 | Davis and Warren 1965 |
| Sockeye salmon | mixed diet | 3.9-7.9 | Brett et al. 1969 |
| Channel catfish | mixed diet | 3.1 | Andrews and Stickney 1972 |

F2ALG, F2DET, F2Z00, F3BEN, F3SED

91. Preference factors for fish compartments 2 and 3
are as follows:

- a. F2ALG is the preference of FISH2 for algae (dimensionless).
- b. F2DET is the preference of FISH2 for detritus (dimensionless).
- c. F2Z00 is the preference of FISH2 for zooplankton (dimensionless).
- d. F3BEN is the preference of FISH3 for benthos (dimensionless).
- e. F3SED is the preference of FISH3 for sediment (dimensionless).

Information relating to fish preference factors is supplied in Leidy and Jenkins (1977) and is reprinted here in Table 26 below. Unfortunately, the different fish foods are expressed as fractions of the total diet rather than as quantities (i.e. grams) consumed, making preference factors difficult to estimate from this information.

Table 26
Fish food expressed as a fraction of the diet
(from Leidy and Jenkins 1977)

| SPECIES | PLANT | DETRITUS | ZOOPL | BENTHOS | FISH |
|---------------------------|-------|----------|-------|---------|------|
| Gizzard shad | 0.10 | 0.80 | 0.05 | 0.05 | |
| Threadfin shad (young) | 0.30 | 0.50 | 0.10 | 0.10 | |
| Threadfin shad (old) | 0.30 | 0.05 | 0.15 | 0.55 | 0.10 |
| Rainbow trout | 0.05 | | 0.60 | 0.15 | |
| Brook trout | | | 0.90 | 0.05 | |
| Carp | 0.30 | 0.40 | 0.20 | 0.10 | |
| Minnows | 0.20 | | 0.20 | 0.60 | |
| Carpsuckers | 0.15 | 0.65 | 0.05 | 0.15 | |
| Suckers | 0.15 | 0.65 | 0.05 | 0.15 | |
| Hogsuckers | | 0.80 | 0.05 | 0.15 | |
| Buffalofish | 0.05 | 0.40 | 0.05 | 0.15 | |
| Redhorse | | | 1.00 | | |
| Bullhead | 0.10 | 0.25 | 0.50 | | 0.15 |
| Catfish | 0.27 | 0.10 | | | 0.80 |
| Madtoms | | | 0.55 | | 0.18 |
| Silversides | | | 0.20 | 0.80 | |
| Temperate bass | | | 0.20 | 0.10 | 0.70 |
| Sunfish | 0.10 | 0.05 | 0.65 | | 0.05 |
| Black bass | | | 0.08 | | 0.86 |
| Crappie | 0.05 | 0.05 | 0.20 | 0.15 | 0.55 |
| Perch | | | 0.20 | 0.20 | 0.60 |
| Freshwater drum | | 0.08 | 0.58 | | 0.34 |

92. An example is given for calculating preference factors for the third fish compartment when actual quantities consumed are known. Suppose a particular species of fish consumes 2.9 out of an available 16.0 g of benthos and 0.26 g out of an available 120.0 g of sediment. The preference factor (P_i) for the i th food category equals

$$P_i = (E_i/A_i)/\sum E_i/A_i \quad (22)$$

where

E_i = the amount of the i th food consumed

A_i = the amount of the i th food available

For the above examples the preference factors would be

$$P(\text{benthos}) = (2.0/16.0)/0.127166 = 0.983$$

$$P(\text{sediment}) = (0.26/120.0)/0.127166 = 0.017$$

FSHT1, FSHT2, FSHT3, FSHT4

93. Upper and lower temperature tolerances for fish ingestion are presented as follows:

- a. FSHT1 is the lower temperature boundary, usually 0 °C, at which metabolism continues.
- b. FSHT2 is the lowest temperature at which processes are occurring at the maximum rates.
- c. FSHT3 is the upper temperature bounding the range of maximum rates.
- d. FSHT4 is the upper lethal temperature.

94. For most warmwater species, upper and lower temperature tolerances are similar, the lower limit being reached at DoC and the upper limit between 33 and 37°C; the optimum temperature is about 27°C. Coldwater species such as salmonids reach a lower temperature limit at DoC, but the upper limit is near 25°C; the optimum temperature is about 14°C. Temperature tolerance values and the various acclimation temperatures (ACCL), where available, are given in Table 27.

Table 27
Temperature coefficients for fish ingestion (Oe)
(from Leidy and Jenkins 1977)

| SPECIES | ACCL | FSHT1 | FSHT2 | FSHT3 | FSHT4 | REFERENCE |
|-------------------|------|-------|-------|-------|----------------------------|------------------------|
| Pickerel | | ● | | 24 | 34.4 | Leidy and Jenkins 1977 |
| Minnows | | ● | 27 | 33.4 | Leidy and Jenkins 1977 | |
| Catfish | | ● | 3. | 37.1 | Leidy and Jenkins 1977 | |
| Sunfish | | 2.5 | 27.5 | 35.7 | Leidy and Jenkins 1977 | |
| Black bass | | 1.6 | 27 | 36.5 | Leidy and Jenkins 1977 | |
| Crappie | | | 23 | 32.5 | Leidy and Jenkins 1977 | |
| Yellow perch | | ● | 24.2 | 30.9 | Leidy and Jenkins 1977 | |
| Yellow perch | | | | 29 | Schneider 1973 | |
| Fingerling salmon | | | 15 | | Brett et al. 1969 | |
| Bluntnose minnow | 5 | | | 26.0 | Hart 1947 | |
| Bluntnose minnow | " | | | 2B.3 | Hart 1947 | |
| Bluntnose minnow | 15 | 1.. | | 30.6 | Hart 1947 | |
| Bluntnose minnow | 2. | 4.2 | | 31.7 | Hart 1947 | |
| Bluntnose minnow | 25 | 7.5 | | 33.3 | Hart 1947 | |
| Flathead minnow | " | | | 2B.2 | Hart 1947 | |
| Flathead minnow | 2. | 1.5 | | 31.7 | Hart 1952 | |
| Flathead minnow | 3. | 10.5 | | 33.2 | Hart 1952 | |
| Creek chub | 5 | | | 24.7 | Hart 1952 | |
| Creek chub | " | | | 27.3 | Hart 1952 | |
| Creek chub | 15 | | | 29.3 | Hart 1952 | |
| Creek chub | 2. | .7 | | 30.3 | Hart 1952 | |
| Creek chub | 25 | 4.5 | | 30.3 | Hart 1952 | |
| Chub | " | | | 27.1 | Black 1953 | |
| Finescaled sucker | 14 | | | 26.9 | Black 1953 | |
| White sucker | 25 | | | 31.2 | Brett 1944 | |
| White sucker | 5 | | | 26.3 | Hart 1947 | |
| White sucker | " | | | 27.7 | Hart 1947 | |
| White sucker | 15 | | | 29.3 | Hart 1947 | |
| White sucker | 2. | 2.5 | | 29.3 | Hart 1947 | |
| White sucker | 25 | 6.0 | 27 | 29.3 | Hart 1947 | |
| White sucker | | | | | McCormick and Mischuk 1973 | |
| Brown bullhead | 5 | | | 27.B | Hart 1952 | |
| Brown bullhead | " | | | 29.0 | Hart 1952 | |
| Brown bullhead | 15 | | | 31.0 | Hart 1952 | |
| Brown bullhead | 2. | | | 32.5 | Hart 1952 | |
| Brown bullhead | 25 | | | 33.8 | Hart 1952 | |
| Brown bullhead | 3. | | | 34.8 | Hart 1952 | |
| Brown bullhead | 34 | | | 34.8 | Hart 1952 | |
| Black bullhead | 23 | | | 35 | Black 1953 | |
| Channel catfish | 25 | | | 35.5 | Allen and Strawn 1968 | |
| Channel catfish | 35 | | | 38 | Allen and Strawn 1968 | |
| Channel catfish | | | 18 | | Andrews and Stickney 1972 | |
| Channel catfish | 15 | ●● | | 30.3 | Hart 1952 | |
| Channel catfish | 2. | 2.5 | | 32.B | Hart 1952 | |
| Channel catfish | 25 | 6.. | | 33.5 | Hart 1952 | |
| Bluegill | 15 | 2.5 | | 30.7 | Hart 1952 | |
| Bluegill | 2. | 5.. | | 31.5 | Hart 1952 | |
| Bluegill | 25 | 7.5 | | | Hart 1952 | |

Table 27 (concluded)

| SPECIES | ACCL | PSHT1 | FSHT2 | FSHT3 | PSHT4 | REFERENCE |
|-------------------------|------|-------|-------|-------|-------|--------------------------|
| Bluegill | 30 | 11.1 | | | 33.B | Hart 1952 |
| Bluegill | | | 22 | | 33.B | McComish 1971 |
| Longear sunfish | 25 | | | | 35.6 | Neill et al. 1966 |
| Longear sunfish | 30 | | | | 36.B | Neill et al. 1966 |
| Longear sunfish | 35 | | | | 37.5 | Neill et al. 1966 |
| Pumkinseed | 25 | | | | 24.5 | Brett 1944 |
| Smallmouth bass | 35 | 1.. | 26.3 | | 35.0 | Horning and Pearson 1973 |
| Small mouth bass | | | 28.3 | | | Peck 1965 |
| Largemouth bass | | | 27.5 | 30 | | Strawn 1961 |
| Largemouth bass | | | 25 | | | Niimi and Beamish 1974 |
| Largemouth bass | 20 | 5.5 | | | 32.5 | Hart 1952 |
| Largemouth bass | 25 | | | | 34.5 | Hart 1952 |
| Largemouth bass | 30 | 11.8 | | | 36.4 | Hart 1952 |
| Yellow perch | 5 | | | | 21.3 | Hart 1947 |
| Yellow perch | 10 | 1.1 | | | 25.0 | Hart 1947 |
| Yellow perch | 15 | | | | 27.7 | Hart 1947 |
| Yellow perch | 25 | 3.7 | | | 29.7 | Hart 1947 |
| Yellow perch-juvenile | 24 | | 20 | 23.3 | | McCauley and Read 1973 |
| Yellow perch-adult | 24 | | 17.6 | 20.1 | | McCauley and Read 1973 |
| Yellow perch | 8 | | 18.6 | | | Ferguson 1958 |
| Yellow perch | 10 | | 19.3 | | | Ferguson 1958 |
| Yellow perch | 15 | | 23.0 | | | Ferguson 195B |
| Yellow perch | 20 | | 23.1 | | | Ferguson 1958 |
| Yellow perch | 25 | | 24.5 | | | Ferguson 1958 |
| Yellow perch | 30 | | 26.7 | | | Ferguson 1958 |
| Sockeye salmon-fry | 5 | 0 | | | 22.2 | Brett 1952 |
| Sockeye salmon-fry | 10 | 3.1 | | | 23.4 | Brett 1952 |
| Sockeye salmon-fry | 15 | 4.1 | | | 24.4 | Brett 1952 |
| Sockeye salmon-fry | 20 | 4.7 | | | 24.8 | Brett 1952 |
| Sockeye salmon-juvenile | 15 | | 15 | 17 | | Brett et al. 1969 |
| Coho salmon | 5 | 0.2 | | | 20.9 | Brett 1952 |
| Coho salmon | 10 | 1.7 | | | 23.7 | Brett 1952 |
| Coho salmon | 15 | 3.5 | | | 24.3 | Brett 1952 |
| Coho salmon | 20 | 4.5 | | | 25.0 | Brett 1952 |
| Chinook salmon | | | 18.4 | | | Olson and Foster 1955 |
| Northern pike | 25 | | | | 32 | Scott 1964 |
| Lake trout | | | 11.7 | | | McCauley and Tait 1970 |
| Lake trout | | | 8 | 10.9 | | Rawson 1961 |
| Rainbow trout | 18 | | 17 | 20 | | McCauley and Pond 1971 |
| Brook trout | 5 | | | | 23.7 | Fry et al. 1946 |
| Brook trout | 10 | | | | 24.4 | Fry et al. 1946 |
| Brook trout | 15 | | | | 25.0 | Fry et al. 1946 |
| Brook trout | 20 | | | | 25.3 | Fry et al. 1946 |
| Brook trout | 25 | 0.5 | | | 25.3 | Fry et al. 1946 |
| Brook trout | | | 14 | 19 | | Graham 1949 |

FEFFIC

95. FEFFIC, the assimilation efficiency for fish (dimensionless), ranges from 0.66 to 0.98; a value of 0.80 is realistic for most fish (Leidy and Jenkins 1977). The assimilation efficiency is multiplied by the ingestion rate to obtain an assimilation rate. Values for fish assimilation efficiency are given in Table 28.

Table 28
Assimilation efficiencies of fish (dimensionless)

| <u>SPECIES</u> | <u>FEFFIC</u> | <u>REFERENCE</u> |
|------------------------|---------------|--------------------------|
| Bleak | 0.80 | Mann 1965 |
| Blueback herring | 0.80 | Burbridge 1974 |
| Bluegill | 0.80 | Pierce and Wissing 1974 |
| Bluegill | 0.97 | Gerking 1955 |
| Carnivorous fish | 0.80 | Wingerg 1956 |
| Carp | 0.74 | Ivlev 1939a |
| Carp | 0.95 | Kobashi and Deguchi 1971 |
| Cichlasama bimaculatum | 0.69-0.89 | Warren and Davis 1967 |
| Cutthroat trout | 0.84-0.86 | Krokhin 1959 |
| Ctenopharyngodon | 0.14 | Fisher 1970 |
| Dace | 0.79 | Mann 1965 |
| Goldfish | 0.71-0.86 | Davies 1964 |
| Green sunfish | 0.94 | Gerking 1952a |
| Longear sunfish | 0.94-0.97 | Gerking 1952a |
| Northern pike | 0.72 | Johnson 1966 |
| Perea fluviatilis | 0.35 | Klekowski et al. 1970 |
| Perch | 0.79 | Mann 1965 |
| Reticulate sculpin | 0.74-0.84 | Davis and Warren 1965 |
| Roach | 0.78 | Mann 1965 |
| White bass | 0.66-0.69 | Wissing 1974 |

TFMORT

96. TFMORT is the nonpredatory mortality rate for fish (1/day). Mortality rate is that fraction of fish biomass that is converted to detritus by death. Nonpredatory mortality rates can be highly variable depending on species, age, exploitation rate, and numerous environmental variables.

The average rate calculated by Leidy and Jenkins (1977) is 0.001 for exploited populations.

97. Ricker (1945) has reviewed techniques for calculating various mortality rates (total, instantaneous, conditional, natural, and fishing). Values for nonpredatory mortality are given in Table 29.

Table 29
Fish nonpredatory mortality rates (l/day)

| <u>SPECIES</u> | <u>TFMORT</u> | <u>REFERENCE</u> |
|-----------------|---------------|----------------------------|
| American shad | 0.002 | Walburg 1961 |
| Bluegill | 0.002 | Patriarche 1968 |
| Bluegill | 0.0002 | Gerking 1952b |
| Bluegill | 0.001 | Ricker 1945 |
| Brook trout | 0.001 | Latta 1962 |
| Brook trout | 0.003-.004 | Alexander and Shetter 1961 |
| Brook trout | 0.56-1.34 | Hatch and Webster 1961 |
| Brown bullhead | 0.001 | McCammon and Seeley 1961 |
| Brown bullhead | 0.001 | Rawstran 1967 |
| Channel catfish | 0.001 | Ricker 1958 |
| Cutthroat trout | 0.001-.002 | Hansen 1971 |
| Cutthroat trout | 0.001 | Ball and Cope 1961 |
| Freshwater drum | 0.001 | Butler 1965 |
| Largemouth bass | 0.00037 | Mraz and Threinen 1955 |
| Longnose sucker | 0.002 | Geen et al. 1966 |
| Northern pike | 0.002 | Groebner 1960 |
| Northern pike | 0.002 | Johnson and Peterson 1955 |
| Rock bass | 0.002 | Ricker 1947 |
| Walleye | 0.001 | Olson 1957 |
| White catfish | 0.001 | McCammon and Seeley 1961 |

TFRESP

98. TFRESP is the fish respiration rate (*l/day*). There are three types of respiration that can be defined: (a) standard respiration--oxygen consumed in the absence of measurable movement (i.e., nonactive respiration, basal of resting metabolism), (b) routine respiration--rate of

oxygen consumption of fish showing normal activity, and (c) active respiration--maximum rate of oxygen consumption under continuous forced active respiration. It would appear that the best estimates of the rate of respiration for normal active fish are values for routine metabolism (i.e., type 2 above) (Winberg 1956). Values for fish respiration rate are given in Table 30.

Table 30
Fish maximum respiration rates (l/day)

| SPECIES | TFRESP | TYPE | REFERENCE |
|--------------------------|------------|----------|---------------------|
| Brown bullhead | 0.001 | routine | Beamish 1964 |
| Brook trout | 0.003 | routine | Beamish 1964 |
| Carp | 0.001 | routine | Beamish 1964 |
| Lake trout | 0.001 | standard | Gibson and Fry 1954 |
| Rainbow trout | 0.002 | standard | Florke et al. 1954 |
| Salvelinus fontinalis | 0.006-.024 | standard | Madsen et al. 1977 |
| Salvelinus fontinalis | 0.019-.101 | active | Madsen et al. 1977 |
| Sockeye salmon | 0.002 | standard | Brett 1944 |
| White sucker | 0.002 | routine | Beamish 1964 |

Other Coefficients

TDSETL

99. TDSETL is the detrital settling velocity (m/day). Detrital settling velocities vary from 0.001 to over 200 m/day depending on the detrital characteristics and reservoir hydrodynamics. Settling rates should be obtained from quiescent settling chamber studies because advective and turbulent forces in the mixed layer that can reduce settling in a reservoir are modeled separately. For most studies, settling velocities are in the range of 0.05 to 1.0 m/day.

Much higher values are often reported for fecal pellets, as shown in Table 20; however, such high settling coefficients may be questionable because they produce unrealistically low detritus values in the modeling studies. Values for detritus settling velocities are given in Table 31.

Table 31
Detritus settling velocities (m/day)

| <u>SOURCE</u> | <u>TDSETL</u> | <u>REFERENCE</u> |
|----------------------------------|---------------|------------------------------|
| Ceratium balticum | 9.0 | Apstein 1910 |
| Chaetoceros borealis | 5.0 | Apstein 1910 |
| Chaetoceros didymus | 0.85 | Eppley et al. 1967b |
| Cricosphaera carterae | 1.70 | Eppley et al. 1967b |
| Ditylum brightwellii | 2.0 | Apstein 1910 |
| Fecal pellets: | | |
| Acartia clausii | 116.0 | Smayda 1971 |
| Fecal pellets: | | |
| Euphausia krohnii | 240.0 | Fowler and Small 1972 |
| Fecal pellets: | | |
| Euphausia pacifica | 43.0 | Osterberg et al. 1963 |
| Fecal pellets: | | |
| Pontella meadii | 54.0-88.0 | Turner 1977 |
| Phaeodactylum tricornutum | 0.02-.04 | Riley 1943 |
| Rhizosolenia herbetata | 0.22 | Eppley et al. 1967b |
| Stephanopyxis tunis | 2.1 | Eppley et al. 1967b |
| Tabellaria flocculosa | 0.46-1.5 | Smayda 1971 |
| Thalassiosira psuedonana | 0.85 | Hecky and Kilham 1974 |

DETT1, DETT2

100. DETT1 is the lower temperature boundary at which decomposition continues to occur. It is usually 0 °C.

101. DETT2 is the temperature at which decomposition occurs near the maximum rate. Temperature coefficients for decomposition are given in Table 32.

Table 32
Temperature coefficients for decomposition (°C)

| <u>SUBSTRATE OR SITE</u> | <u>DETT1</u> | <u>DETT2</u> | <u>REFERENCE</u> |
|---|--------------|--------------|-----------------------|
| Pseudomonas fluorescens: natural substrate | 0 | 25-30 | Tison and Pope 1980 |
| E. coli: natural substrate | 0 | 37 | Tison and Pope 1980 |
| Glucose: Lake George, New York | 0 | 25 | Tison et al. 1980 |
| Glucose | 0 | 20-30 | Bott 1975 |
| Glucose: Lake Wingra, Wis. | | 25-30 | Boylen and Brock 1973 |

TDOMDK

102. TDOMDK is the dissolved organic matter (DOM) decay rate (1/day). DOM in natural waters is the organic substrate for heterotrophic metabolism. The composition of natural DOM is highly variable and little understood, but its sources are generally grouped into (a) excretion from phytoplankton and macrophytes, (b) decomposition of phytoplankton and macrophytes, (c) excretion by animals, and (d) allochthonous drainage (e.g., humic compounds from upstream sources).

103. Aquatic bacteria appear to be chiefly responsible for the removal of DOM compounds from the water; they are the major agents for bacterial mineralization of organic solutes in fresh water (Wright 1975), using organic matter as an energy source. Various methods have been tested to determine the decay rate of DOM in water. Modification of the basic Parson and Strickland (1963) technique have been developed to quantify the kinetics.

104. DOM decomposition rates have also been represented by filtered carbonaceous biochemical oxygen demand (BOD) decay rates. If sufficient oxygen is available, the

aerobic biological decomposition of organics will continue until all the DOM is consumed. In the standard test for BOD, a sample is diluted with water containing a known amount of oxygen. The loss of oxygen after the sample has been incubated for 5 days at 20°C is known as the 5-day BOD. The value of the first-order decay rate is generally about 0.05 to 0.20 per day.

105. The BOD test suffers from several serious deficiencies. The test has no stoichiometric validity, for example: the arbitrary S-day period usually doesn't correspond to the point where all the organic matter is consumed.

106. Contributing to the errors involved in measuring decay rates of DOM is the extensive variability in the composition and stage of decomposition of DOM. Allochthonous inputs of DOH are likely to be more refractory than autochthonous inputs, and as a result, decomposition rates will be slower and decay may be incomplete; therefore, the length of time the organic matter is available for decomposition is important. In addition, as particles sink out of the euphotic zone, both dissolved and detrital organic substrates may be limited to more resistant fractions thereby arresting attached microbial growth. Therefore, the rate of DOM decomposition may be lower in the hypolimnion of a stratified reservoir.

107. Oxygen consumption rate (mg O₂/L/hr) can be transformed into a mineralization rate of organic carbon (mg C/L/hr) by application of a conversion factor of 0.29 (Seepers 1981). Values for DOM decay rate are given in Table 33.

Table 33
DOM decay rates (1jday)

| <u>COMPOUND</u> | <u>TDOMDK</u> | <u>REFERENCE</u> |
|-----------------|---------------|-------------------------|
| Acetate | 0.2 | Wright 1975 |
| Amino acids | 0.64 | Williams et al. 1976 |
| Glucose | 0.24 | Williams et al. 1976 |
| Glucose | 0.32-.50 | Toerien and Cavari 1982 |
| Glucose | 0.111 | Wright 1975 |
| Glutamate | 0.11-.625 | Carney and Colwell 1976 |
| Glycine | 0.312-.45 | Vaccaro 1969 |
| Glycine | 0.048 | Vaccaro 1969 |
| Glycolate | 0.024-.432 | Wright 1975 |
| Glycolate | 0.012-.25 | Wright 1975 |
| Glycolic acid | 0.004 | Tanaka et al. 1974 |

TNH3DK

108. TNH3DK is the ammonia decay rate (i.e., the rate at which ammonia is oxidized to nitrite) (1/day). Ammonia is generated by heterotrophic bacteria as the primary end product of decomposition of organic matter, either directly from proteins or from other nitrogenous organic compounds. Although ammonia is a major excretion product, this nitrogen source is minor in comparison to decomposition.

109. Nitrification is the biological conversion of organic and inorganic N compounds from a reduced state to a more oxidized state (Alexander 1965). The nitrifying bacteria capable of oxidation of NH₄⁺ to NO₂⁻ are largely confined to the species Nitrosomonas, bacteria which are mesophilic (1-37 °C).

110. Nitrification rate can be determined by a number of different techniques. Courchaine (1968) has plotted nitrogenous BOD on a logarithmic scale and determined the decay rate from the slope of the line. Thomann et al. (1971) used a finite-difference approximation to solve a

set of simultaneous linear equations.

III. Laboratory measurements for the ammonia decay rate can produce results that differ from what might be measured *in situ*. Several environmental factors influence the rate of nitrification, including pH, temperature, suspended particulate concentration, hydraulic parameters and benthos.

112. Nitrification can be measured as a one- or two-step process. In the one-step method, only the end product of the entire reaction, nitrate, is measured. In the two-step method, (a) nitrite accumulation is measured as ammonia is oxidized to nitrite and (b) nitrate accumulation is measured as nitrite is oxidized to nitrate. Oxidation of ammonia to nitrite is the rate-limiting step in the total reaction; therefore, experiments that measure the rate of the total reaction (i.e., the one-step method) can be used to estimate this parameter. Ammonia oxidation rates are given in Table 34.

Table 34
Ammonia oxidation rates (1/day)

| SITE | TNH3DK | REFERENCE |
|----------------------------|------------|------------------|
| Wastewater treatment plant | 0.05-0.30 | Wild et al. 1971 |
| Grand River, Ill. | 0.80 | Bansal 1976 |
| Grasmere Lake, U.K. | 0.001-.013 | Hall 1982 |
| Truckee River, Nev. | 0.09-1.30 | Bansal 1976 |
| Upper Mohawk River, N.Y. | 0.23-0.40 | Bansal 1976 |
| Middle Mohawk River | 0.30 | Bansal 1976 |
| Lower Mohawk River | 0.30 | Bansal 1976 |
| Ohio River | 0.25 | Bansal 1976 |
| Big Blue River, Neb. | 0.17-0.25 | Bansal 1976 |
| Flint River, Mich. | 0.76-0.95 | Bansal 1976 |

TN02DK

113. TN02DK is the decay rate of nitrite to nitrate (1/day).

TDETDK

114. TDETDK is the detritus decay rate (1/day). Detritus as defined by Wetzel et al. (1972) consists of organic carbon lost from an organism by nonpredatory means (including egestion, excretion, secretion, etc.) from any trophic level component, or input from sources external to the ecosystem that enter and cycle in the system (i.e., allochthonous organic carbon). For CE-QUAL-RI, this should be considered to be particulate material only.

115. The rate of detritus decay can be determined by measuring the use of oxygen during decomposition, with results expressed as a first-order decay coefficient (k base $e = \text{mg oxygen used/mg day}$). Many workers have measured rates of oxygen uptake by detritus, suggesting that oxygen uptake is related to the organic matter available for decomposition. Odum and de la Cruz (1967) and Fenchel (1970), for example, demonstrated an inverse relation between detritus particle size and oxygen consumption. Oxygen uptake is an integrative measure of all oxidative processes occurring in the sample, both chemical and biological: reducing substances are usually rapidly oxidized; respiration of the organisms associated with detritus is primarily bacterial, although algae, protozoa, and fungi may also contribute. Measurement of the oxygen uptake reflects the metabolism of communities of microorganisms involved in the decomposition of natural substances.

116. As a detrital particle decomposes with time, there is a decline in oxygen uptake accompanied by succession of communities of microorganisms; this decline occurs

as the matter changes from labile to refractory; refractory matter often accumulates in the sediment. Rates of decay are generally high initially and slow down as the material becomes refractory; the rate is influenced by temperature, detrital composition, and age of the detritus. Macrophyte communities are the primary source of detritus in most systems. Submersed and floating macrophytes generally decay more rapidly than the highly lignified emergent species. Particulate organic matter of dead bluegreen algae decomposes much faster than that derived from green algae diatoms and desmids. Particulate organic matter (POM) is especially resistant (Gunnison and Alexander 1975). As detritus decays, there is a decrease in the C:N ratio as a result of a buildup of microbial protein (Mann 1972). A 1-g sample of detritus at 20°C consumes about 1 mg oxygen/hr (Hargrave 1972).

117. Plant litter consists of a variety of compounds (*i.e.*, sugars, hemicellulose, lignin, waxes) which decay at different rates. The decay curves initially tend to follow the exponential decay functions of the more readily degradable fractions, particularly aquatic macrophytes, which account for a large proportion of the weight of plant litter; therefore, the majority of the litter's weight loss occurs in the first year. Over the long term, the decay rates change, especially for deciduous leaf litter which has a larger proportion of decay-resistant material than do aquatic macrophytes and therefore decays at a much slower rate.

118. Decay rates can also be measured by suspending a nylon mesh bag of detrital material in situ or under controlled conditions and determining weight loss with time. This actually measures weight loss due to enzymatic decomposition by bacteria and fungi, solution of soluble sub-

stances, and loss of fragments through the container pores.

119. Decay rates have also been determined by measuring the mineralization rates of carbon, nitrogen, and phosphorus (Otuski and Hanya 1972). Decomposition of detritus generated from planktonic communities of surface lake water occurs at rates on the order of 10 percent per day (Saunders 1972), based upon radioactive carbon tracer studies.

120. Consideration should be given to the primary or expected sources of detritus. Decomposition rates for allochthonous detrital sources are generally lower than for autochthonous sources to reflect the more refractory nature of allochthonous material after its transport through the upper portions of the reservoir. While a one-dimensional model like CE-QUAL-RI assumes instantaneous dispersal of inflow constituents, much of the decomposition in the prototype reservoir system occurs in the headwater area. The labile fraction of autochthonous detritus produced in the pelagic zones of the lower reservoir will decompose more rapidly in the water column and should have a higher decomposition rate than allochthonous detritus. However, in a stratified reservoir the POM in the hypolimnion may not be exchanged with the epilimnetic waters. The POM becomes more refractory with time, and rates of decomposition decrease.

121. Microbial decomposition of detritus can be represented by three stages: a very quick solution of soluble organic components, a relatively rapid decomposition of labile organic constituents, and slow decomposition of refractory organic constituents. Detritus decay rates are given in Table 35.

Table 35
Detritus decay rates (1/day)

| <u>DETTRITUS SOURCE</u> | <u>TDETDK</u> | <u>REFERENCE</u> |
|-----------------------------------|---------------|-------------------------|
| Beech | 0.001-.004 | Hanlon 1982 |
| Cladophera glornerata | 0.007 | Piecznska 1972 |
| Dead green algae | 0.016-.076 | Otsuki and Hanya 1972 |
| Dead mixed algae | 0.007-.111 | Jewell and McCarty 1971 |
| Dead mixed algae | 0.007-.06 | Fitzgerald 1964 |
| <i>Gloeotrichia</i> echinulata | 0.001-.007 | Piecznska 1972 |
| Isoetes lacustris | 0.003-.015 | Hanlon 1982 |
| Leaf packs | 0.005-.017 | Sedell et al. 1975 |
| Osier | 0.001-.005 | Hanlon 1982 |
| Potamogeton crispus | 0.002-.004 | Rogers and Breen 1982 |
| Potomogeton perfoliatlls | 0.002-.007 | Hanlon 1982 |

TCOLDK

122. TCOLDK is the coliform decay rate (1/day). Estimates of coliform die-off rates may be obtained in the laboratory or in situ. In situ, where there are no flow regime data, or where flows are of a transient nature, a commonly used method is to add a slug of a conservative tracer substance (a dye, rare element, or radioisotope) to steady-state discharge. The discharge plume is sampled, dilution is estimated from the concentration of tracer, and the decay rate is estimated from the dilution-corrected coliform counts. This technique gives misleading results in cases where the tracer is diluted by water heavily contaminated with the same discharge. Since the tracer was introduced as a slug, there is no way to know how many of the surviving coliforms originated in the tracer-dosed effluent and how many came from pre- or post-dosing effluent. This problem is reduced where the flow regime is sufficiently stable (Zison et al. 1978).

123. There are two approaches to estimating die-off rates. Frost and Streeter (1924) were able to estimate the die-off rate using seasonal averages of coliform counts from a downstream station, by assuming plug flow in the river. Errors in the rates determined by this approach are attributable to (a) dilution and to longitudinal mixing that produced overestimates and (b) unconsidered sources of coliforms that produced underestimates.

124. In a second approach, a mathematical model of the flow and mixing in the system is used to correct the measurements for the effects of dilution. In this manner Marais (1974) analyzed coliform die-off in wastewater maturation ponds as a first-order decay reaction in a series of completely mixed steady-state reactors. Errors in the decay rates determined in this way are primarily attributable to the reliability of the system model.

125. Table 36 gives decay rates for coliform and fecal streptococcus. In Table 37 from Mitchell and Chamberlain (1978), the median die-off value was 0.040/hr for freshwater coliform. In general, the die-off follows first-order decay kinetics, although a significant increase in coliform levels is commonly observed in the first several miles downstream from the outfall.

126. Factors affecting coliform decay rate include sedimentation, solar radiation, nutrient deficiencies, predation, algae, bacterial toxins, and physiochemical factors.

Table 36
Coliform and fecal streptococcus decay rates (1/ day)

| <u>SPECIES</u> | <u>TCOLDK</u> | <u>REFERENCE</u> |
|---------------------------|---------------|--------------------------------------|
| Fecal coliform | 0.048-.096 | Evans et ala 1968 |
| Fecal streptococci | 0.063 | Evans et ala 1968 |
| Fecal streptococci | 0.004-.013 | Geldreich et al. 1968 |
| Total coliform | 4.48-5.52 | Kittrell and Furfari 1963 |
| Total coliform | 0.199-.696 | Klock 1971 |
| Total coliform | 1.99 | Marais 1974 |
| Total coliform | 0.168-1.56 | Geldreich et ala 1968 |
| Total coliform | 0.009-.028 | Klock 1971 |
| Total coliform | 0.021-.038 | Evans et al. 1968 |
| Total coliform | 0.045-.049 | Frost and Streeter 1924 |
| Total coliform | 0.024-.105 | Hoskins et al. 1927 |
| Total coliform | 0.48-2.04 | Mitchell and Chamberlain 1978 |

Table 37
Freshwater die-off rates of coliform bacteria measured in situ (l/da.)
 from Mitchean Chamber ain 97

| SITE | TEMP/SEASON | RATE | REFERENCE |
|-------------------------------|-------------|------------|------------------------------|
| Ohio River | Summer 20°C | 1.175 | Frost and Streeter 1924 |
| Ohio River | Winter 5°C | 1.08 | Frost and Streeter 1924 |
| Upper Illinois River | June-Sept. | 2.04 | Hoskins et al. 1927 |
| Upper Illinois River | OCT.-Hay | 2.52 | Hoskins et al. 1927 |
| Upper Illinois River | Dec. Mar. | 0.576 | Hoskins et al. 1927 |
| Upper Illinois River | Apr.-Nov. | 1.032 | Hoskins et al. 1927 |
| Lower Illinois River | June-Sept. | 2.04 | Hoskins et al. 1927 |
| Lower Illinois River | Oct.-May | 0.888 | Hoskins et al. 1927 |
| Lower Illinois River | Dec.-Mar. | 0.624 | Hoskins et al. 1927 |
| Lower Illinois River | Apr. - Nov. | 0.696 | Hoskins et al. 1927 |
| Shallow turbulent stream | Summer | 15.12 | Kittrell and Koschitzky 1947 |
| Missouri River | Winter | 0.48 | Kittrell and Furfari 1963 |
| Tennessee River (.Knoxville) | Summer | 1.03 | Kittrell and Furfari 1963 |
| Tennessee River (Chattanooga) | Summer | 1.32 | Kittrell and Furfari 1963 |
| Sacramento River, Calif. | Summer | 1.752 | Kittrell and Furfari 1963 |
| Cumberland River, Md. | Summer | 5.52 | Kittrell and Furfari 1963 |
| Groundwater stream | 10°C | 0.504 | Wuhrmann 1912 |
| Leaf River, Miss. | NA | 0.408 | Mahloch 1974 |
| Wastewater lagoon | 7.9-2S.S.C | 0.199-.696 | Klock 1971 |
| Maturation ponds | N | 1.99 | Marais 1974 |
| Maturation ponds | 19°C | 1.68 | Marais 1974 |
| Oxidation ponds | 20°C | 2.59 | Marais 1974 |

TSEDDK

127. T5EDDK is the organic sediment decomposition rate (1/day). While sediment consists primarily of settled organic detritus, the decomposition rate should reflect the changing nature of the detritus as it reaches the sediment; i.e., it becomes more refractory since the labile portion of the organic detritus decomposes as it settles through the water column. In addition, since the initial value for sediment is in 91m^2 , the thickness of the sediment layer, along with TSEDDK, will affect the amount of predicted decomposition. Thus, if high initial values are used for sediment, TSEDDK may have to be lowered since only the top few centimeters of sediment are usually involved in aerobic decomposition. Hargrave (1969) found the following relationship between the rate of oxygen consumption by sediments ($\text{ml O}_2/\text{m}^2/\text{hr}$) and the temperature ($T, ^\circ\text{C}$):

$$\ln (\text{O}_2 \text{ consumption rate}) = 1.74 * \ln(T) - 1.30 \quad (23)$$

At 6° C this would be $214.3 \text{ mg O}_2/\text{m}^2/\text{day}$, assuming a constant rate for the day and the conversion formula found in the CE-QUAL-R1 User's Manual (Environmental Laboratory 1982, p. 188). At 25° C the rate would be $2567 \text{ mg/m}^2/\text{day}$. The amount of sediment (in mg/m^2) times the value for TSEDDK times 1.4 (i.e., the stoichiometric equivalent of oxygen uptake to sediment decay) should be near the $6-25^\circ\text{C}$ range.

DOMT1, DOMT2

128. DOMT1, the critical low temperature for DOM decay, is usually 0°e .

129. DOMT2 is the optimum temperature for DaM decay (ee). Temperature coefficients for DaM decay are given in Table 38.

Table 38
Temperature coefficients for DOM decay (Oe)

| SUBSTRATE | DOMT1 | DOMT2 | REFERENCE |
|----------------------------|-------|-------|-------------------------|
| Glucose | 5.0 | 35.5 | Toerien and Cavari 1982 |
| Glucose: Lake George, N.Y. | 0 | 25 | Tison et al. 1980 |
| Glucose | 0 | 20-30 | Batt 1975 |
| Glucose: Lake Wingra, Wis. | 0 | 25-30 | Boylen and Brock 1973 |

NH3T1, NH3T2

130. Researchers have generally found temperature to affect nitrification rates, especially in the range of 10 to 35°C.

- a. NH3T1 is the lower temperature boundary at which ammonium nitrification continues. It is generally 0 °C.
- b. NH3T2 is the optimum temperature for oxidation of NH3-N. The optimum temperature for nitrification is generally accepted to be between 25 and 30°C.

Temperature factors for ammonia oxidation are given in Table 39.

Table 39
Temperature coefficients for ammonia oxidation (Oe)

| SPECIES OR SITE | NH3T1 | NH3T2 | REFERENCE |
|----------------------------|-------|-------|---------------------|
| Nitrosomonas | 5 | 30 | Knowles et al. 1965 |
| Wastewater treatment plant | 5 | 25 | Wild et al. 1971 |
| Ann Arbor, Michigan | 2 | 20 | Borchardt 1966 |

N02T1, N02T2

131. N02T1 is the lower temperature boundary at which nitrate nitrification occurs (QC).

132. N02T2 is the lowest temperature (OC) at which the oxidation of nitrite to nitrate occurs near the maximum rate.

TSSETL

133. TSSETL is the suspended solids settling velocity (rn/day). The settling rate is dependent on the type of particle, grain size, density, temperature, viscosity, and turbulence. Most of the larger particles entering a reservoir settle very quickly and should not be included in the inflow. Lane (1938) gives figures of 0.86 to 860.0 m/day for particle diameters of 0.002 to 0.1 mm. Particles found in the main body of a reservoir are usually at the lower end of this scale.

Q1OCOL

134. CE-QUAL-R1 uses a 010 formulation to modify the coliform die-off rate as a function of temperature. All other rates are modified by temperature through the RMULT function in CE-QUAL-R1. The 010 coefficient is usually 1.04.

PART III: RECOMMENDATIONS

135. This report provides information about, and values for, many of the coefficients needed for use of the version of the model CE-QUAL-R1 described in the User's Manual (Environmental Laboratory 1982).

136. Research on processes described in **this report is likely to provide more information needed to refine the equations used in the model.** Future versions of the model may therefore require additional coefficients.

137. This report may be updated to provide information about, and values for, any additional coefficients needed for use of future versions of the model.

138. Application, calibration, and verification of the model to a variety of sites *is likely to identify coefficient values that are best suited to the model.* These values may be included in updates to this report.