

## 11 DILUTION PURIFICATION

Dilution purification (also known as *self purification*) refers to discharge of partially treated sewage into a body of water such as a stream or river. If the body is large and is adequately oxygenated, the sewage's BOD may be satisfied without putrefaction. Other conditions which must be monitored besides BOD are oxygen content and suspended solids.

Equation 8.20 can be used to calculate the final concentration of BOD, oxygen, and sediment when the two flows are mixed. Dilution requirements may be expressed in terms of ratios (e.g., 23 stream volumes per discharge volume) or absolute flow quantities (e.g., 4 to 7 cfs per 1000 population).

$$C_1 Q_1 + C_2 Q_2 = C_f (Q_1 + Q_2) \quad 8.20$$

### Example 8.3

Wastewater ( $DO = 0.9$  mg/l, 6 MGD) is discharged into a  $50^\circ\text{F}$  stream flowing at 40 cfs. Assuming the stream is saturated with oxygen, what is the oxygen content of the stream immediately after mixing?

From appendix B, the saturated oxygen content at  $50^\circ\text{F}$  ( $10^\circ\text{C}$ ) is 11.3 mg/l.

$$(6 \text{ MGD}) \left( 1.547 \frac{\text{cfs}}{\text{MGD}} \right) = 9.28 \text{ cfs}$$

$$C = \frac{(0.9)(9.28) + (11.3)(40)}{9.28 + 40} = 9.34 \text{ mg/l}$$

The *oxygen deficit* is the difference between actual and saturated oxygen concentrations. Since reoxygenation and deoxygenation of a polluted river occur simultaneously, an oxygen deficit will occur only if the reoxygenation rate is less than the deoxygenation rate. If the oxygen content goes to zero, anaerobic decomposition and putrefaction will occur.

The oxygen deficit at any time  $t$  is given by the *Streeter-Phelps equation*:

$$D_t = DO_{\text{sat}} - DO_t$$

$$= \frac{K_D BOD_u}{K_R - K_D} (10^{-K_D t} - 10^{-K_R t})$$

$$+ D_o (10^{-K_R t}) \quad 8.21$$

$D_t$  is the dissolved oxygen deficit,  $t$  is in days, and  $BOD_u$  is the ultimate carbonaceous BOD of the stream immediately after mixing.  $K_D$  and  $K_R$  are the deoxygenation and reoxygenation rate constants respectively, and  $D_o$  is the dissolved oxygen deficit immediately after mixing.<sup>9</sup>

$K_R$  can be approximated by equation 8.22 if field test data is not available.<sup>10</sup>

$$K_{R, 20^\circ\text{C}} \approx \frac{3.3v}{d^{1.33}} \quad 8.22$$

$K_R$  for different temperatures is given by equation 8.23. Typical values of  $K_R$  are given in table 8.12.

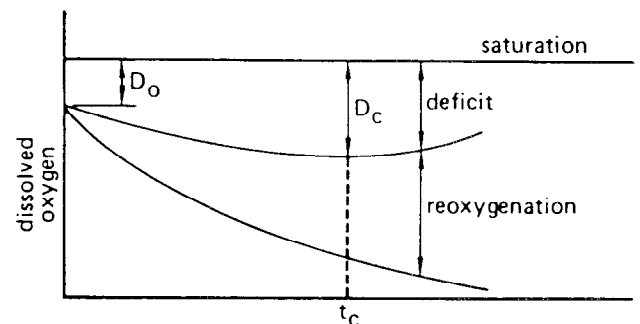
$$K_{R T} = (1.016)^{T-20} K_{R, 20^\circ\text{C}} \quad 8.23$$

**Table 8.12**

Typical Reoxygenation Constants (base 10, 1/days)

white water	0.5 and above
swiftly flowing	0.3 to 0.5
large streams	0.15 to 0.3
large lakes	0.10 to 0.15
sluggish streams	0.10 to 0.15
small ponds	0.05 to 0.10

Equations 8.6 and 8.21 can be plotted simultaneously as shown in figure 8.5. The plot of equation 8.21 is known as the *oxygen sag curve*. The difference between the two curves is the effect of reoxygenation.



**Figure 8.5** The Oxygen Sag Curve

<sup>9</sup>  $K_R$  and  $K_D$  may be written as  $K_1$  and  $K_2$  by other authors

<sup>10</sup> Equation 8.22 is the O'Connor and Dobbins formula for natural streams.