Corrections to "Introduction to Environmental Biophysics", Second Printing, 2000.

Chapt	er 2
CIIGP	

P. 35	line 36	In prob. 1d, "sensible heat flux," should be "sensible heat flux
		density,".

P. 44 line 26 Change "...dependence
$$(0.01\%/C)$$
..." to "...dependence $(0.1\%/C)$..." . Chapter 3

Chapter 4

Chapter 5

Chapter 6

P. 78 last 2 lines Change "... dividing by
$$c_p$$
 ..." to "...dividing by $\hat{\rho} c_p$..." on page Change "...quantity k/c_p is ..." to "...quantity $k/\hat{\rho} c_p$..."

P. 79 Eq. 6.8 Add ") " so right term is "=
$$c_p (T_s - T_a)/r_H$$

Chapter 7

P. 92 Example 7.3 Change the number
$$40x10^{-6}$$
 m⁻² to $40x10^{+6}$ m⁻². in g_{vs} (adaxial)

P. 103 Eq. 7.30a Since units of
$$g_{Ha}$$
 usually are mol m⁻² s⁻¹, the constant of 240 should be 0.240. The constant of 240 means g_{Ha} in this Eq. is in mmol m⁻² s⁻¹.

Chapter 8

And the sentence that follows should read "where erfc is the complimentary error function, a function ..."

Chapter 9

P. 129 Eq. 9.1	All K's should be $K(\psi_m)$
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P. 130 Table 9.1
$$\theta_{-1500} = 0.21$$
 for silty clay loam (not 0.32)

P. 135 Example 9.1 Change "Solve Eq.(9.1) for
$$\theta$$
:" to "Solve Eq.(9.3) for θ :"

P. 140 Eq. 9.16 Insert a minus so
$$R_P = -\psi_{Lm}/E_{pmax}$$

P. 142 Eq. 9.21 All potentials are normalized so all
$$\psi$$
's should be ψ^* 's.

Chapter 10

P. 147 line 5
$$c = 3 \times 10^8 \text{ m/s}$$

P. 151 Ex. 10.1 PPFD=500 J/(
$$m^2$$
 s) x 1/(2.17 x 10⁵ J) = 2.3 mol/(m^2 s) = 2300 μ mol/(m^2 s)

P. 159 Eq. 10.5 For more clarity, denominator should be
$$\lambda^5$$
 [exp(hc/(k λ **T**)) – 1]

P. 160 Fig. 10.4 Add units of
$$W/m2/\mu m$$
 to right hand axis of graph

P. 164 Eq. 10.11 The "
$$T_a$$
" should be " T_a " indicating K temperature

Chapter 11

P. 171 line 29 Converting to hours gives
$$128^{\circ} \times 1 \text{ hr}/15^{\circ} = 8.56 \text{ hrs}$$
. The time of first twilight is $11.87 - 8.56 = 3.31 \text{ hrs}$ (local time). The daylength is $2 \times 8.56 = 17.1 \text{ hrs}$. The time of sunrise in PDT is $3.31 \text{ hrs} + 1 \text{ hr} = 4.31 \text{ hrs}$ (PDT).

P. 181 line 6
$$F_d = F_a = (1 - \cos \gamma)/2$$

Chapter 14

- P. 244 line 6 Replace "... a little above 20°C at..." with "...a little below 20°C at ..."
- P.245 Penman ref. Penman, H.L. (1948) Natural evaporation from open water, bare soil, and grass. Proc. R. Soc. London, A, 194:120-145.

Chapter 15

- P. 257 Eq.(15.10) Replace $\rho_{b,cpy}^*$ with $\rho_{b,cpy}^*(\psi)$ so that $\rho_{b,cpy} \approx \rho_{b,cpy}^*(\psi) (\rho_{b,cpy}^*(\psi) \rho_S) \exp(-2\sqrt{\alpha}K_{be}(\psi)L_t)$
- P. 260 line 14 Example 15.2 total incident PAR above canopy should be $Q_{ob} + Q_{od} = 2000 \ \mu mol \ photons \ m^{-2} \ s^{-1}$.
- P. 261 Example 15.2 At mid page the equation for $\overline{Q}_{sh} = \dots$ should be $\alpha \, \overline{Q}_{sh} = \dots$ so $\alpha \, \overline{Q}_{sh} = 0.8 \, (177 + 26) = 0.8 \, x \, 203$ The two equations for $\overline{Q}_{sl} = \dots$ should be $\alpha \, \overline{Q}_{sl} = \dots$ so $\alpha \, \overline{Q}_{sl} = \alpha (K_{be}(\psi)Q_{ob}) + \overline{Q}_{sh}$ and $\alpha \, \overline{Q}_{sl} = 0.8(0.652 \, x \, 1600 + 203) = 0.8 \, x \, 1246$ This is because these equations are for the PAR absorbed.
- P.261 last 2 lines Insert α in front of \overline{Q}_{sl} and \overline{Q}_{sh} so that on page The leaf assimilation rates can be obtained from Fig. 14.6 using α \overline{Q}_{sh} for shaded leaves and α \overline{Q}_{sl} for sunlit leaves.
- P.262 last Eq. on Insert α in front of Q, \overline{Q}_{sl} and \overline{Q}_{sh} so that Page $\overline{Q} = \frac{\alpha \overline{Q}_{sl} L_t^* + \alpha \overline{Q}_{sh} (L_t - L_t^*)}{L_t} = \frac{997 \times 1.32 + 162 \times 1.68}{3.0}$
- P.279 Table A1 Column headed with ρ should be changed to $\hat{\rho}$.
- P. 281 Table A3(continued) Units of slope of saturation vapor pressure versus temperature (triangle symbol at column head) should be PaC⁻¹ as in Table A3 on page 280.
- P. 281 Table A5

 Units on Gas constant should be J mol⁻¹ K⁻¹

 Units on Boltzmann constant should be J K⁻¹

 Units on Stefan-Boltzman constant should be W m⁻² K⁻⁴
- P.284 line 5 "Flick's diffusion law" should be "Ficks' diffusion law"