This expression is directly comparable to (4-58), written for a point outside the ellipsoid for which

$$
\int_{\beta}^{1}=\int_{1}^{1}=0
$$

so that $E, T$, and $Q$ vanish, and $D=D(1)=1$ as above:

$$
\begin{equation*}
V=\frac{G M}{r}\left[1-\frac{2}{5} \frac{R^{2}}{r^{2}} S_{1} P_{2}(\cos \theta)+\frac{12}{35} \frac{R^{4}}{r^{4}} P_{1} P_{4}(\cos \theta)\right] . \tag{6-14}
\end{equation*}
$$

Here we have introduced $G M / r$ in a similar way as in (6-1) and expressed $r$ in metric units, which explains the occurrence of $R$ to make dimensions right. Comparing the coefficients of $P_{4}(\cos \theta)$ in $(6-13)$ and $(6-14)$ thus yields

$$
-\frac{a^{4}}{r^{4}} J_{4}=\frac{12}{35} \frac{R^{4}}{r^{4}} P_{1}
$$

or

$$
P_{1}=-\frac{35}{12}\left(\frac{a}{R}\right)^{4} J_{4} .
$$

Since $J_{4}=O\left(f^{2}\right)$, we may put $(a / R)^{4}=1$ without losing accuracy, obtaining

$$
\begin{equation*}
P_{1}=-\frac{35}{12} J_{4} . \tag{6-15}
\end{equation*}
$$

By means of (6-12) and (6-15) we are thus able to express $J_{4}$ as follows:

$$
\begin{equation*}
J_{4}=-\frac{4}{5} f^{2}+\frac{4}{7} f m-\frac{32}{35} \kappa_{1} . \tag{6-16}
\end{equation*}
$$

This fundamental equation links the spherical-harmonic coefficient $J_{4}$ with the geometric parameters $f$ and $\kappa_{1}$ and with $m$. A useful check is provided by the fact that for the ellipsoid with $\kappa_{1}=0,(6-16)$ reduces to eq. (2-119) of (Heiskanen and Moritz, 1967).

In fact, for the level ellipsoid, $\kappa_{1}=0$ by (6-6), but we have retained $\kappa_{1} \neq 0$ because we shall consider, besides the level ellipsoid, also the case of the real earth.

### 6.2 Level Ellipsoid and Equilibrium Figures

As we have remarked, the basic spheroidal equations such as $(6-1),(6-3)$, or ( $6-16$ ) hold for the level ellipsoid (and the real earth) as well as for equilibrium figures, since they have been derived without presupposing hydrostatic equilibrium.

Now there comes a surprise: it is possible to find two corresponding mass distributions, one for equilibrium and the other for the level ellipsoid, for which the values of the ellipticity ( $4-48$ ),

$$
\begin{equation*}
e=f-\frac{5}{42} f^{2}+\frac{4}{7} \kappa \tag{6-17}
\end{equation*}
$$

are equal for each $\beta$. This is possible because the coefficients $A_{0}(\beta)$ and $A_{2}(\beta)$ in $(6-1)$, as given by (4-63) and (4-64), do not depend on the deviation $\kappa$, provided they are expressed in terms of $e$ instead of $f$; compare also (4-56) and (4-59)! This is an essential advantage of describing the elliptical shape of the internal surfaces of constant density by the "artificial" parameter (6-17) instead of the flattening $f$ itself.

In other terms, we may take

$$
\begin{gather*}
A_{0}^{E}(\beta) \equiv A_{0}^{H}(\beta)  \tag{6-18}\\
A_{2}^{E}(\beta) \equiv A_{2}^{H}(\beta) \tag{6-19}
\end{gather*}
$$

the superscript $E$ denoting the ellipsoid. Of course, $A_{4}^{E}$ will be different from $A_{4}^{H}=0$, and ( $6-19$ ) vanishes because of the equilibrium conditions (4-69).

Thus using now subscripts to denote the ellipsoidal and the hydrostatic cases,

$$
\begin{equation*}
e_{E}(\beta)=e_{H}(\beta) \tag{6-20}
\end{equation*}
$$

but since

$$
\begin{equation*}
\kappa_{E} \neq \kappa_{H} \tag{6-21}
\end{equation*}
$$

also

$$
\begin{equation*}
f_{E} \neq f_{H} \tag{6-22}
\end{equation*}
$$

except in second-order terms.
Thus we have (Moritz, 1973, p. 31) the
Theorem
To each mass distribution in hydrostatic equilibrium there corresponds a mass distribution for the equipotential ellipsoid in such a way that the density $\rho$ is the same function of $\beta$ and. that the values of the ellipticity $e$ are the same for any two surfaces corresponding to the same value of $\beta$.

In the sequel we shall always assume that $e=e_{E}$ is selected in this way. We then obtain an ellipsoidal mass configuration which deviates very little from an equilibrium configuration.

We shall also omit the index $E$ for ellipsoidal quantities.
On this assumption, (6-19) vanishes as we have seen, and (6-1) reduces to

$$
\begin{equation*}
W(\beta, \theta)=W_{0}(\beta)+W_{4}(\beta) P_{4}(\cos \theta) \tag{6-23}
\end{equation*}
$$

where, by (4-63) and (4-68) with $A_{2}=0$ :

$$
\begin{align*}
W_{0}(\beta) & =\frac{G M}{R} \beta^{2}\left[D\left(1+\frac{1}{3} \mu+\frac{4}{45} e^{2}+\frac{4}{45} e \mu\right)+\frac{3}{2} E-\right. \\
& \left.-\frac{4}{25} e S+\frac{8}{75} e T\right],  \tag{6-24}\\
W_{4}(\beta) & =\frac{G M}{R} \beta^{2} \cdot \frac{8}{35}\left[\left(\frac{3}{2} e^{2}-4 \kappa\right) D-3 e S+\frac{3}{2} P+\frac{4}{3} Q\right] \tag{6-25}
\end{align*}
$$

A further simplification of $W_{4}$ is obtained by subtracting the hydrostatic value

$$
\begin{equation*}
W_{4}^{H}(\beta)=\frac{G M}{R} \beta^{2} \cdot \frac{8}{35}\left[\left(\frac{3}{2} e^{2}-4 \kappa_{H}\right) D-3 e S+\frac{3}{2} P_{H}+\frac{4}{3} Q_{H}\right] \equiv 0, \tag{6-26}
\end{equation*}
$$

noting that $D$ and $S$ are equal in both cases. Thus we get

$$
\begin{equation*}
W_{4}(\beta)=\frac{G M}{R} \beta^{2} \cdot \frac{32}{105}\left[-3\left(\kappa-\kappa_{H}\right) D+\frac{9}{8}\left(P-P_{H}\right)+\left(Q-Q_{H}\right)\right], \tag{6-27}
\end{equation*}
$$

where, by (4-56),

$$
\begin{align*}
\frac{9}{8}\left(P-P_{H}\right) & =\beta^{-7} \int_{0}^{\beta} \delta \frac{d}{d \beta}\left[\left(\kappa-\kappa_{H}\right) \beta^{7}\right] d \beta  \tag{6-28}\\
Q-Q_{H} & =\beta^{2} \int_{\beta}^{1} \delta \frac{d}{d \beta}\left[\left(\kappa-\kappa_{H}\right) \beta^{-2}\right] d \beta \tag{6-29}
\end{align*}
$$

### 6.3 Equipotential Surfaces and Surfaces of Constant Density

Denote a surface of constant density, $\rho=\rho_{1}$, by $S_{1}$ and a corresponding surface of constant potential, $W=W_{1}$, by $S_{2}$. Let the surface $S_{1}$ be characterized by a value $\beta_{1}$ such that

$$
\begin{equation*}
\rho\left(\beta_{1}\right)=\rho_{1} ; \tag{6-30}
\end{equation*}
$$

then the constant $W_{1}$ will be determined by

$$
\begin{equation*}
W_{0}\left(\beta_{1}\right)=W_{1}, \tag{6-31}
\end{equation*}
$$

the function $W_{0}(\beta)$ being expressed by $(6-24)$. Thus a surface $S_{2}$ is made to correspond to each surface $S_{1}$ (Fig. 6.1).


FIGURE 6.1: A surface of constant density, $S_{1}$, and the corresponding surface of constant potential, $S_{2}$

