# Physics with neutrons 2 

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## ExERCISE 3.1

The potential

$$
U(r, \vartheta, \varphi)=-U_{0} \Theta(R-r)
$$

is called a hard sphere potential with radius $R .(\Theta(x)$ is the Heaviside step function, which is defined to be zero for $x<0$ and unity for $x \geq 0$.)

1. Calculate the differential and the total cross section of scattering from this potential.
2. Using small-angle neutron scattering, a biologist would like to measure the diameter of spherical micelles (aggregated "clusters" of molecules in a solvent). What is the form factor $F(Q R)$ (i.e. the $Q$-dependent part of the differential scattering cross section) of one such micelle under the assumption that it can be approximated by a homogeneous sphere with a radius of 200 nm ?
3. For small values of $Q R$, the form factor can be Taylor-expanded. What is the resulting behavior?
4. Plot the form factor (versus $Q R$ ) on a log-log scale. For large values of $Q R$, what is the behavior of $F(Q R)$ when one averages over the oscillations?
5. What happens (qualitatively) when the sphere is placed in a solvent? What happens when there are multiple spheres present?

Solution. The scattering amplitude is given by

$$
f(\vec{Q})=-\frac{m}{2 \pi \hbar^{2}} \int d^{3} r U(\vec{r}) \exp (-i \vec{Q} \cdot \vec{r})
$$

or, in spherical coordinates,

$$
f(Q)=-\frac{m}{2 \pi \hbar^{2}} \int d r d(\cos \vartheta) d \varphi r^{2} U(r) \exp (-i Q r \cos \vartheta)
$$

Since the potential is spherically symmetric, we can use $\vartheta$ as the angle between $\vec{Q}$ and $\vec{r}$ in the exponential because we can select the coordinate system freely, so that $\vec{Q}$ is along the $z$ axis. Inserting our given potential and doing the trivial $\varphi$ integration we get

$$
f(Q)=-\frac{m}{2 \pi \hbar^{2}}\left(-2 \pi U_{0}\right) \int_{0}^{\infty} d r r^{2} \Theta(R-r) \int_{-1}^{1} d(\cos \vartheta) \exp (-i Q r \cos \vartheta)
$$

The integration over $\cos \vartheta$ is also easy:

$$
f(Q)=\frac{m U_{0}}{\hbar^{2}} \int_{0}^{\infty} d r r^{2} \Theta(R-r) \frac{1}{-i Q r}(\exp (-i Q r)-\exp (i Q r)) .
$$

Now we replace the exponential representation of the sine and resolve the $\Theta$ function by adjusting the integration limits:

$$
\begin{gathered}
f(Q)=\frac{2 m U_{0}}{\hbar^{2}} \frac{1}{Q} \int_{0}^{R} d r r \sin (Q r) . \\
f(Q)=\frac{2 m U_{0}}{\hbar^{2}} \frac{1}{Q}\left[\frac{\sin Q r}{Q^{2}}-\frac{r \cos Q r}{Q}\right]_{0}^{R}=\frac{2 m U_{0}}{\hbar^{2}} \frac{\sin Q R-Q R \cos Q R}{Q^{3}} .
\end{gathered}
$$

The final step is to rewrite this a bit:

$$
f(Q)=\frac{m U_{0}}{2 \pi \hbar^{2}} \frac{4 \pi R^{3}}{3} \frac{3(\sin Q R-Q R \cos Q R)}{(Q R)^{3}}=\rho \cdot V_{s} \cdot 3 \frac{\sin Q R-Q R \cos Q R}{(Q R)^{3}},
$$

where $\rho=m U_{0} / 2 \pi \hbar^{2}$ is the "scattering length density" (SLD) and $V_{s}$ the volume of the sphere. Note that the SLD definition matches well with the Fermi pseudopotential used for scattering at single nuclei: the singular $b \delta(r)$ is replaced by an $\rho(r)$ extended over the sphere. This is sensible because in small angle scattering, we are looking at very large structures and so cannot resolve individual scattering centers inside the spheres any more.
The differential cross section $d \sigma / d \Omega$ is simply given by $|f(Q)|^{2}$.
The total cross section is obtained by integrating over all solid angle. For this, we need to express $d \Omega$ in terms of $Q$ :

$$
\begin{gathered}
Q=2 k \sin \frac{\theta}{2} \Longrightarrow \frac{d Q}{d \theta}=k \cos \frac{\theta}{2} \\
\Rightarrow d \Omega=2 \pi d \theta \sin \theta=2 \pi \frac{d Q}{k \cos \frac{\theta}{2}} \sin \theta=2 \pi d Q \frac{2 \sin \frac{\theta}{2}}{k}=2 \pi \frac{Q}{k^{2}} d Q .
\end{gathered}
$$

The integration now gives

$$
\sigma=\int_{\Omega} d \Omega \frac{d \sigma}{d \Omega}=\int_{0}^{2 k} d Q \frac{2 \pi Q}{k^{2}}|f(Q)|^{2} ;
$$

note that we integrate in $Q$ from 0 to $2 k$, which is the maximum momentum transfer (a backscattering process). Inserting $f(Q)$ gives
$\sigma=\int_{0}^{2 k} d Q \frac{2 \pi Q}{k^{2}} 9 \rho^{2} V^{2} \frac{(\sin Q R-Q R \cos Q R)^{2}}{(Q R)^{6}}=\frac{18 \pi \rho^{2} V^{2}}{(k R)^{2}} \int_{0}^{2 k R} d u \frac{\sin ^{2} u-2 u \sin u \cos u+u^{2} \cos ^{2} u}{u^{5}}$
with the substitution $u=k R$, and solving the integral we have

$$
\sigma=\frac{9 \pi \rho^{2} V^{2}}{2(k R)^{2}}\left[1-\frac{1}{(2 k R)^{2}}+\frac{\sin 4 k R}{(2 k R)^{3}}-\frac{\sin ^{2} 2 k R}{(2 k R)^{4}}\right]
$$

as the total scattering cross section.
The form factor for the micelle is the $Q$-dependent part of the differential cross section for the hard sphere potential with $R=200 \mathrm{~nm}$ :

$$
F(Q R)=\rho^{2} \cdot V_{s}^{2} \cdot 9 \frac{\sin ^{2} Q R-2 Q R \sin Q R \cos Q R+Q^{2} R^{2} \cos ^{2} Q R}{(Q R)^{6}} .
$$

For small values of $Q R$, the Taylor expansion of the form factor reads

$$
F(Q R) \approx 1-\frac{(Q R)^{2}}{5}
$$

This is called the "Guinier approximation", and it gives information about the micelle size. A plot of the form factor:


The behavior for large $Q R$, when averaging over oscillations, is $\propto(Q R)^{-4}$.
With the sphere placed in a solvent, the scattering length density $\rho$ is replaced by the SLD contrast between sphere and solvent $\rho_{\text {sphere }}-\rho_{\text {solvent }}$. By appropriate "contrast matching" with different solvents (e.g. different mixtures of $\mathrm{D}_{2} \mathrm{O}$ and $\mathrm{H}_{2} \mathrm{O}$ ), one can accurately determine $\rho_{\text {sphere }}$. For $N$ spheres, the form of $d \sigma / d \Omega$ stays the same (with a factor of $N$ ) only if the spheres are noninteracting. In reality, they are interacting (e.g. certain sphere-sphere distances are preferred), which leads to an additional factor in $d \sigma / d \Omega$ called the "structure factor".

## EXERCISE 3.2

When investigating an object with small-angle scattering, the Patterson function yields useful statistical information. It is defined by

$$
P(\mathbf{r})=\int \rho\left(\mathbf{r}_{1}\right) \rho\left(\mathbf{r}_{2}\right) d \mathbf{V}, \quad \text { with } \mathbf{r}=\mathbf{r}_{1}-\mathbf{r}_{2},
$$

where $\rho$ is the scattering-length density function of the object. The correlation function $\gamma(r)$ is the orientational average of the Patterson function $P(\mathbf{r})$, which is in two dimensions

$$
\begin{equation*}
\gamma(r)=\frac{1}{2 \pi} \int_{0}^{2 \pi} P(\mathbf{r}) \mathrm{d} \varphi . \tag{1}
\end{equation*}
$$

It answers the question: given that there is an atom of the particle at some place, what is the probability that the atoms in the distance $r$ are also situated inside the particle?

Numerically calculate the (two-dimensional) Patterson function and subsequently the characteristic function $\gamma_{0}(r)=\frac{\gamma(r)}{\gamma(0)}$ of the following objects (black area $\rho=1$, white area $\rho=0$ )


Why is $\gamma(r \geq D)=0$ when $D$ is the largest possible distance of two atoms inside the particle? What is the connection between Patterson function and scattering signal of the object?

Solution. The Patterson function corresponds to a convolution of the scattering length density $\rho(r)$ with $\rho(-r)$. This operation is called Autocorrelation and it describes the correlation of a function with itself translated by a distance $r^{\prime}$. In our case, the function corresponds to the particle shape. We get the following Patterson functions (Matlab function conv2)


Certain characteristics of the particle shape are still visible in this representation. In the next step we perform the radial integration around the centre to get the characteristic function of the particle


The characteristic function is important for interpreting scattering data, since the WienerKhinchin theorem states that the Fourier transform of the patterson function is equal the squared Fourier transform of the scattering length density, e.g. the scattering signal

$$
|\mathscr{F}(\rho)|^{2}=|\mathscr{F}(P)| .
$$

Consequently, we can calculate the radial averaged scattering signal by performing the radial Fourier transform of the characteristic

$$
I(Q)=4 \pi \int_{0}^{D} \gamma(r) \frac{\sin (Q r)}{Q} r \mathrm{~d} r .
$$



As a cross check we calculate the scattering signal in both ways, from the Patterson function and the squared Fourier transform of the scattering length density

## Fourier transform of the Patterson function



Squared Fourier transform of the scattering length density





