# **Nuclear Astrophysics**

Lecture 11 Thurs. Jan. 19, 2012 Prof. Shawn Bishop, Office 2013, Ex. 12437 shawn.bishop@ph.tum.de

# REACTION RESONANCE PHENOMENON

V(r)Simple "toy" model to consider: 3-D SWE, s-wave scattering  $(\ell=0)$ and potential just a constant  $-V_0$ for  $r \leq R_0$ E $R_0$ rThe general radial SWE:  $\frac{d^2u}{dr^2} + \frac{2\mu}{\hbar^2}[E+V_0]u = 0$  $-V_0$ Or:  $\frac{d^2u}{dm^2} + k^2u = 0$   $k^2 = \frac{2\mu}{\hbar^2}[E + V_0]$ For  $r \leq R_0$ ,  $E - V_0 > 0 \Rightarrow u_{in}(r) = Ae^{ikr} + Be^{-ikr}$  $= A(\cos kr + i\sin kr) + B(\cos kr - i\sin kr)$ 

Recall:  $R(r) = \frac{u(r)}{r}$ 

To satisfy this at the origin, we require A+B=0

Therefore,  $u_{in}(r) = (A - B)i\sin kr = 2Ai\sin kr$ 

Outside potential, for  $r > R_0$ 

$$u_o(r) = Ce^{iKr} + De^{-iKr} \qquad K^2 = \frac{2\mu}{\hbar^2}E$$

 $= (C+D)\cos Kr + i(C-D)\sin Kr$ 

 $= E\sin Kr + F\cos Kr$ 

The constants E and F are normalization constants, and we can write them in any form we want (as long as the form is always a constant). Factor them both as follows:

$$E = G \cos \delta_0$$
 and  $F = G \sin \delta_0$ 

Then:

$$u_o(r) = G(\cos \delta_0 \sin Kr + \sin \delta_0 \cos Kr)$$
$$= G \sin(Kr + \delta_0)$$

At the boundary of the potential,  $r = R_0$ , we require continuity of wave functions:

$$A' \sin kR_0 = G \sin(KR_0 + \delta_0)$$
 Value matching

 $A'k\cos kR_0=GK\cos(KR_0+\delta_0)$  Derivative matching

Take the ratio of these two equations:

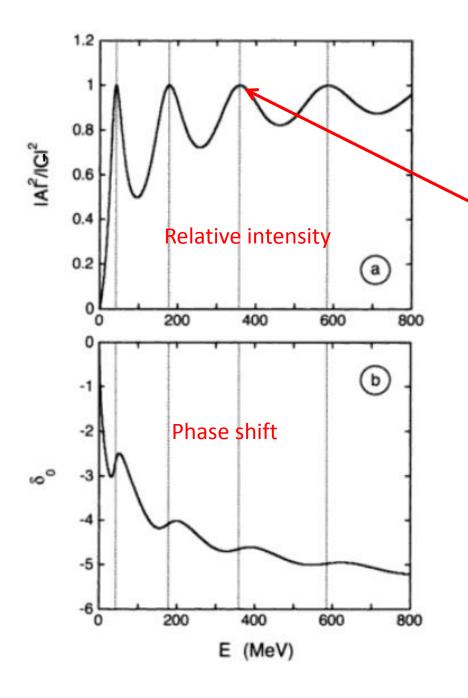
A' = 2Ai

We have for the phase shift of the outer wave function:  $\Rightarrow \delta_0 = -KR_0 + \arctan\left[\frac{K}{k} \tan kR_0\right]$ 

Now, what about the relative intensity of the outer wave function within the interior of the potential  $|A|^2/|G|^2$ ? Take the modulus-squared of both equations above and add the results:

$$\left|\frac{A^2}{G^2}\right| = \frac{K^2}{K^2 + [k^2 - K^2]\cos^2 kR_0}$$

$$= \frac{E}{E + V_0 \cos^2\left(\frac{\sqrt{2\mu(E+V_0)}}{\hbar}R_0\right)}$$



Results for  $V_0=-100~{\rm MeV}$ ,  $R_0=3~{\rm fm}$ and an energy E corresponding to:  $2\mu/\hbar^2=0.0484~{\rm MeV}^{-1}{\rm fm}^{-1}$ 

These structures are resonances. The amplitude matching between the incident (external) wave function and the internal wave function are optimized at these values of incident energy.

$$\left. \frac{A^2}{G^2} \right| = \frac{E}{E + V_0 \cos^2\left(\frac{\sqrt{2\mu(E+V_0)}}{\hbar}R_0\right)}$$

Maximized when the cosine term is zero; when argument is  $(n+1/2)\pi$  .

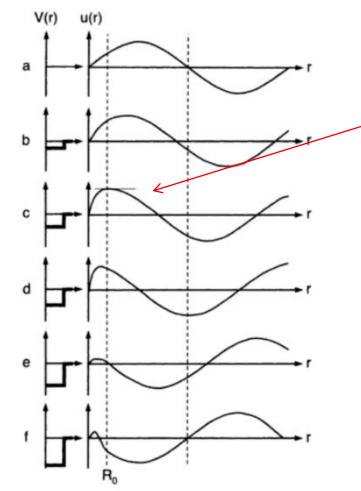
$$\Rightarrow E_n = \frac{\hbar^2}{2\mu} \frac{\pi^2}{R_0^2} \left( n + \frac{1}{2} \right) - V_0$$

These are Resonance Energies

When the incident beam energy is at, or close to, one of the system's resonance energies, the probability of forming a compound state in the potential is maximized.

Also note: the condition 
$$\cos kR_0 = \cos\left(rac{\sqrt{2\mu(E+V_0)}}{\hbar}R_0
ight) = 0$$

tells us something about the derivative matching equations on page 20: this cosine term is exactly the same as the internal wave function derivative condition being set to zero.



This suggests that resonances occur when the outer wave function and internal wave functions match, such that, the slope of wave function at the boundary radius is zero. Also note: the boundary and derivative matching equations on page 5 are equivalent to the following:

$$R\left(\frac{1}{u_{in}(r)}\frac{du_{in}(r)}{dr}\right)_{r=R_0} = R\left(\frac{1}{u_o(r)}\frac{du_o(r)}{dr}\right)_{r=R_0}$$

$$f_{\ell} \equiv R\left(\frac{1}{u_{\ell}(r)}\frac{du_{\ell}}{dr}\right)_{r=R_0}$$

This is called the *logarithmic derivative at the boundary*. Our boundary/derivative matching conditions, then, for any partial waves (not just s-waves) can be summarized succinctly as:

$$f_{\ell}^{in} = f_{\ell}^{o}$$

When  $f_{\ell}^{in} = 0$  the system should be at a resonance.

Recall from page 17, Lecture 7, the elastic cross section was found to be:

$$\sigma_{el} = \frac{4\pi}{k^2} \sum_{\ell=0}^{\infty} (2\ell+1) \sin^2 \delta_\ell = \frac{\pi}{k^2} \sum_{\ell=0}^{\infty} (2\ell+1) \left| 1 - e^{2i\delta_\ell} \right|^2$$

And the reaction cross section was found to be, on page 19, Lecture 7,

$$\sigma_{re} = \frac{\pi}{k^2} \sum_{\ell=0}^{\infty} (2\ell+1) \left( 1 - \left| e^{2i\delta_{\ell}} \right|^2 \right)$$

Recall from page 15, of Lec. 7, that the total outgoing scattered wave function was shown to be:

$$\psi_{\ell} = \frac{1}{2kr} \sum_{\ell=0}^{\infty} (2\ell+1)i^{\ell+1} \left[ e^{-i(kr-\ell\pi/2)} - e^{2i\delta_{\ell}} e^{i(kr-\ell\pi/2)} \right] P_{\ell}(\cos\theta) \equiv \frac{u_{\ell}(r)}{r}$$

Let's deal with  $\ell = 0$  only:  $\sigma_{el,\ell=0} = \frac{\pi}{k^2} \left| 1 - e^{2i\delta_0} \right|^2$ 

$$\sigma_{re,\ell=0} = \frac{\pi}{k^2} \left( 1 - \left| e^{2i\delta_0} \right|^2 \right) \quad \psi_0 = \frac{i}{2kr} \left( e^{-ikr} - e^{2i\delta_0} e^{ikr} \right) = \frac{u_0(r)}{r}$$

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These cross sections must be related back to the phase shift. Let's determine what the logarithmic derivative is for this outgoing wave function.

$$\frac{f_0}{R_0} = \left(\frac{1}{u_o}\frac{du_o}{dr}\right)_{r=R_0} = \frac{-ike^{-ikR_0} - ike^{2i\delta_0}e^{ikR_0}}{e^{-ikR_0} - e^{2i\delta_0}e^{ikR_0}}$$

After some algebraic work:  $e^{2i\delta_0} = {f_0 + ikR_0\over f_0 - ikR_0} \, e^{-2ikR_0}$ 

Behold! We can now drop this into the elastic and reaction cross section equations (I=0) at the bottom of previous page:

$$\sigma_{el,0} = \frac{\pi}{k^2} \left| 1 - \frac{f_0 + ikR_0}{f_0 - ikR_0} e^{-2ikR_0} \right|^2 = \frac{\pi}{k^2} \left| e^{2ikR_0} - \frac{f_0 + ikR_0}{f_0 - ikR_0} \right|^2$$
$$= \frac{\pi}{k^2} \left| -\frac{2ikR_0}{f_0 - ikR_0} + e^{2ikR_0} - 1 \right|^2 = \frac{\pi}{k^2} \left| A_{res} + A_{pot} \right|^2$$
$$A_{res} = -\frac{2ikR_0}{f_0 - ikR_0} \qquad A_{pot} = e^{2ikR_0} - 1$$

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Now, look at the form for the resonance scattering amplitude:

 $A_{res} = -\frac{2ikR_0}{f_0 - ikR_0}$ 

It has its maximum when  $f_0 = 0$ ; when the logarithmic derivative is zero.

For the reaction cross section:

$$\sigma_{re,0} = \frac{\pi}{k^2} \left( 1 - \left| e^{2i\delta_0} \right|^2 \right) = \frac{\pi}{k^2} \left( 1 - \left| \frac{f_0 + ikR_0}{f_0 - ikR_0} e^{-2ikR_0} \right|^2 \right)$$

Because the wave functions can be *complex*, the logarithmic derivative can be, in general, a complex function:  $f_0 = g + ih = \text{Re}f_0 + i \text{Im}f_0$ 

Subbing this in to the above equation, and two more steps of complex algebra will result in the following reaction cross section formula

$$\sigma_{re,0} = \frac{\pi}{k^2} \frac{-4kR_0 \operatorname{Im} f_0}{(\operatorname{Re} f_0)^2 + (\operatorname{Im} f_0 - kR_0)^2}$$

# Summary Thus Far:

Elastic cross section in terms of logarithmic derivative, and boundary of potential:

$$\sigma_{el,0} = \frac{\pi}{k^2} \left| -\frac{2ikR_0}{f_0 - ikR_0} + e^{2ikR_0} - 1 \right|^2 = \frac{\pi}{k^2} \left| A_{res} + A_{pot} \right|^2$$
$$A_{res} = -\frac{2ikR_0}{f_0 - ikR_0} \qquad \qquad A_{pot} = e^{2ikR_0} - 1$$

Reaction cross section in terms of logarithmic derivative and boundary of potential:

$$\sigma_{re,0} = \frac{\pi}{k^2} \frac{-4kR_0 \operatorname{Im} f_0}{(\operatorname{Re} f_0)^2 + (\operatorname{Im} f_0 - kR_0)^2}$$

What have we done? We've derived "formal" expressions for the elastic and reaction cross sections in terms of the logarithmic derivative, and in doing so, we have eliminated the rather mathematically "ugly" phase shift  $\delta_0$ 

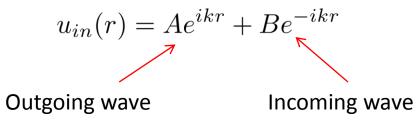
But we've traded this problem for another: in avoiding the phase shift, we've introduced the logarithmic derivative  $f_0$ . What have we "won" by doing this. How do we calculate  $f_0$ ?

We calculate according to its definition, 
$$\frac{f_{in}}{R_0} = \left(\frac{1}{u_{in}}\frac{du_{in}}{dr}\right)_{r=R_0}$$

And there are ways to build model wave functions for  $u_{in}(r)$ . Further, by using the *internal* wave function we continue to stay away from direct usage of  $\delta_{\ell}$ 

Let's do an example to see where the physics comes in to all of this mathematical formality.

As before, take as our (highly simplified) internal nuclear wave function to be of a form:



We require this structure of wave function because we are now allowing reactions to occur. Reactions involve beam particles penetrating into the nuclear volume (Incoming wave), and then leaving this volume through some reaction channel (outgoing wave).

We cannot expect these waves (incoming & outgoing) to be in phase with each other.
The incident beam particle may be absorbed in the nuclear volume (i.e. A (p,γ) reaction), so the outgoing wave amplitude *must* be smaller amplitude than incident wave

Let us therefore try the following *ansatz*:  $A = B e^{2i\xi} e^{-2q}$ 

With  $\xi$  and q both real numbers.

So take as our model internal wave function:  $u_{in}(r) = B(e^{2i\xi} e^{-2q} e^{ikr} + e^{-ikr})$ 

So we have:

$$u_{in}(r) = B(e^{2i\xi} e^{-2q} e^{ikr} + e^{-ikr})$$

$$= \frac{B}{2} \left[ e^{-i(kR_0 + \xi + iq)} + e^{i(kR_0 + \xi + iq)} \right] \times 2e^{(i\xi - q)}$$

$$= 2Be^{(i\xi-q)}\cos(kr + \xi + iq)$$

Exercise for you: Using this form for the internal wave function, determine the logarithmic derivative to be:

$$f_0 = R_0 \left(\frac{1}{u_{in}(r)} \frac{du_{in}(r)}{dr}\right)_{r=R_0} = -kR_0 \tan(kR_0 + \xi + iq)$$

#### This result must be zero for a resonance!

Now, think physics. The parameter q represents attenuation of the outgoing internal wave function due to absorption. (note it appeared as an exponential, similar to the exponential penetrability factor in our discussions on charged particle cross sections and the S-factor).

In *low energy* reactions, absorption is *weak* and elastic scattering is huge.

We therefore consider  $|q|\ll 1$  , and we demand that the log. derivative be zero for resonance

$$f_0(E_{\lambda}, q) = -kR_0 \tan(kR_0 + \xi + iq) = 0$$

Expand this in a Taylor Series around the resonance energy  $E_{\lambda}$  and around q=0

$$f_0 \approx f_0(E_\lambda, q) + (E - E_\lambda) \left(\frac{\partial f_0}{\partial E}\right)_{E_\lambda, q=0} + q \left(\frac{\partial f_0}{\partial q}\right)_{E_\lambda, q=0}$$

The first term is zero, by definition of resonance condition.

At resonance energy, from above, the tangent function is zero and:  $\frac{d}{dx}(\tan x) = \cos^{-2} x$ 

Therefore,

$$\left(\frac{\partial f_0}{\partial q}\right)_{E_{\lambda},q=0} = 1$$

$$\Rightarrow f_0 \approx (E - E_{\lambda}) \left(\frac{\partial f}{\partial E}\right)_{E_{\lambda}, q=0} - iqkR_0 = \operatorname{Re} f_0 + i\operatorname{Im} f_0$$

Referring back to pages 10-12, and using the previous results:

$$\sigma_{re,0} = \frac{\pi}{k^2} \frac{-4kR_0 \operatorname{Im} f_0}{(\operatorname{Re} f_0)^2 + (\operatorname{Im} f_0 - kR_0)^2}$$

$$= \frac{\pi}{k^2} \frac{4qk^2 R_0^2}{(E - E_\lambda)^2 \left(\frac{\partial f_0}{\partial E}\right)_{E_\lambda, q=0}^2 + (qkR_0 + kR_0)^2}$$

$$= \frac{\pi}{k^2} \frac{4qk^2 R_0^2}{(E - E_\lambda)^2 \left(\frac{\partial f_0}{\partial E}\right)_{E_\lambda, q=0}^2 + (qkR_0 + kR_0)^2}$$

$$= \frac{\pi}{k^2} \frac{(2kR_0)(2qkR_0)/(\partial f_0/\partial E)^2_{E_{\lambda},q=0}}{(E-E_{\lambda})^2 + (qkR_0 + kR_0)^2/(\partial f_0/\partial E)^2_{E_{\lambda},q=0}}$$

Continuing..

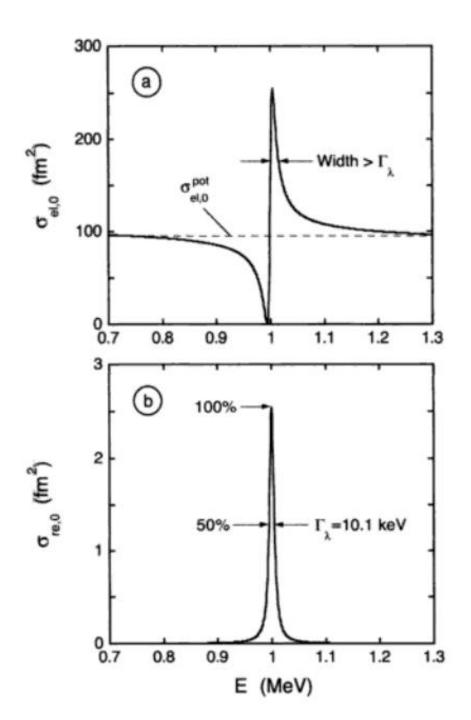
$$\sigma_{re,0} = \frac{\pi}{k^2} \frac{(2kR_0)(2qkR_0)/(\partial f_0/\partial E)_{E_\lambda,q=0}^2}{(E-E_\lambda)^2 + (qkR_0 + kR_0)^2/(\partial f_0/\partial E)_{E_\lambda,q=0}^2}$$
$$= \frac{\pi}{k^2} \frac{\Gamma_{\lambda e}\Gamma_{\lambda r}}{(E-E_\lambda)^2 + \Gamma_\lambda^2/4}$$

$$\begin{split} \Gamma_{\lambda e} &\equiv -\frac{2kR_0}{(\partial f_0/\partial E)_{E_{\lambda},q=0}} \qquad \Gamma_{\lambda r} \equiv -\frac{2qkR_0}{(\partial f_0/\partial E)_{E_{\lambda},q=0}} \\ \Gamma_{\lambda} &= \Gamma_{\lambda e} + \Gamma_{\lambda r} \end{split}$$

$$\sigma_{el,0} = \frac{\pi}{k^2} \left| -\frac{2ikR_0}{f_0 - ikR_0} + e^{2ikR_0} - 1 \right|^2$$

$$= \frac{\pi}{k^2} \left| -\frac{i\Gamma_{\lambda e}}{(E - E_{\lambda}) + i\Gamma_{\lambda}/2} + e^{2ikR_0} - 1 \right|^2$$

$$= \frac{\pi}{k^2} \left[ 2 - 2\cos(2kR_0) + \frac{\Gamma_{\lambda e}^2 - \Gamma_{\lambda e}\Gamma_{\lambda}\cos(2kR_0) + 2\Gamma_{\lambda e}(E - E_{\lambda})\sin(2kR_0)}{(E - E_{\lambda})^2 + \Gamma_{\lambda}^2/4} \right]$$
Interference term: changes sign above and below resonance energy  $E_{\lambda}$ 



Elastic resonant cross section showing the interference effect

Reaction resonant cross section. The width of the curve at 50% of maximum is the *total* width  $\Gamma_\lambda$ 

### **Resonant Reaction Rate:**

$$r_{12} = \left(\frac{8}{\pi\mu}\right)^{1/2} \frac{N_1 N_2}{1 + \delta_{12}} \tau^{-3/2} \int_0^\infty E\sigma(v) \exp\left(-\frac{E}{\tau}\right) dE$$
$$\sigma = \frac{\pi}{k^2} \frac{\Gamma_a \Gamma_b}{(E - E_r)^2 + \Gamma^2/4} = \frac{\pi\hbar^2}{2\mu E} \frac{\Gamma_a \Gamma_b}{(E - E_r)^2 + \Gamma^2/4}$$

Assume that both widths are small (total width is narrow) and they are constant (very weak energy dependence). Sub the above cross section into the integral, treating the  $\Gamma$  terms as constants:

$$r_{12} = \left(\frac{8}{\pi\mu}\right)^{1/2} \frac{\pi\hbar^2}{2\mu} \frac{N_1 N_2}{1+\delta_{12}} \tau^{-3/2} e^{-E_r/\tau} \frac{\Gamma_a \Gamma_b}{\Gamma} 2 \int_0^\infty \frac{\Gamma/2}{(E-E_r)^2 + \Gamma^2/4} dE$$

Resonant Reaction Rate (Single Resonance):

$$r_{12} = \left(\frac{2\pi}{\mu\tau}\right)^{3/2} \hbar^2 \frac{N_1 N_2}{1+\delta_{12}} \frac{2J_r + 1}{(2J_1 + 1)(2J_2 + 1)} \frac{\Gamma_a \Gamma_b}{\Gamma} e^{-E_r/\tau}$$

We have added the spin-statistical factors (now) to account for the fact that our theory up to this point has only considered spin zero particles (in entrance channel and exit channel) in the reaction.

More generally, for particles with spin, we have to multiply the cross section by the spin-statistical factor to account for the different permutations of spin alignments that are allowed.

 $J_r$  is the spin (intrinsic) of the resonance.

J1 "" beam particle.J2 "" target particle.

We define  $\omega\gamma\equiv rac{2J_r+1}{(2J_1+1)(2J_2+1)}rac{\Gamma_a\Gamma_b}{\Gamma}$  as the resonance strength.

In the event that several resonances can contribute to the rate, then we have:

$$r_{12} = \left(\frac{2\pi}{\mu\tau}\right)^{3/2} \hbar^2 \frac{N_1 N_2}{1+\delta_{12}} \sum_i (\omega\gamma)_i e^{-E_{r_i}/\tau}$$

Where the sum extends over all resonances that can contribute to the rate, and where each resonance has its own resonance energy and resonance strength.