

ANALYSIS OF A LOW ENRICHED URANIUM CENTRIFUGAL GAS CORE  
REACTOR

by

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## ABSTRACT

DARRIN LEER. Analysis of a Low Enriched Uranium Centrifugal Gas Core Reactor. (Under the direction of DR. DONALD JACOBS)

With the new NASA directive of returning to the Moon in 2024 as a stepping stone to missions to Mars and beyond, there is a renewed interest in developing nuclear thermal rockets (NTR) to reduce trip times. This thesis will focus on the analysis of a conceptual reactor design for use as an NTR. The low enriched uranium centrifugal gas core reactor (CGCR) is a low technology readiness level (TRL) concept that uses centrifuge technology to separate uranium gas from hydrogen propellant. There is also a new US directive for additional focus on research and development of reactors that utilize low enriched uranium (LEU) instead of high enriched uranium (HEU). The inclusion of a moderator in between the gas enables the use of LEU and a lower mass system compared to previous gas core concepts. In addition, the CGCR operates at lower temperatures than previous gas core concepts enabling higher uranium densities, which suggests that the centrifugal separation will aid in minimization of uranium entrainment. This research will cover a thorough analysis of neutronics, thermal transport, fluid dynamics, and comparison to alternative NTR designs utilizing computational methods, such as the Monte Carlo N-Particle transport code MCNP, and analysis software platforms Mathematica and MATLAB.

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## LIST OF ABBREVIATIONS

CGCR	Centrifugal Gas Core Reactor
GCR	Gas Core Reactor
HEU	High Enriched Uranium
LEU	Low Enriched Uranium
MCNP	Monte-Carlo N-Particle
NASA	National Aeronautics and Space Administration
NTP	Nuclear Thermal Propulsion
NTR	Nuclear Thermal Rocket
PBR	Particle Bed Reactor
SNTP	US Space Nuclear Thermal Propulsion
TRL	Technology Readiness Level
US	United States

## CHAPTER 1: INTRODUCTION

### 1.1 Motivation

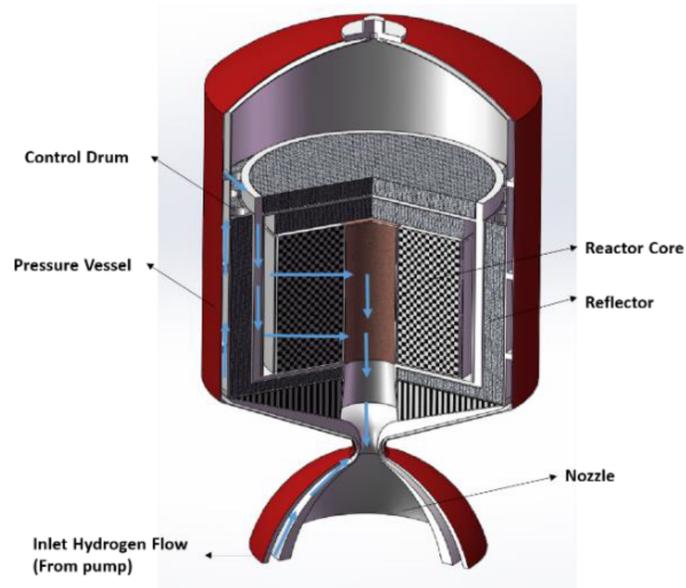
Long-term human space exploration missions come with many challenges and an increasing level of risk. One way to reduce that risk to make missions beyond the moon more feasible is to make the journey less long term. The best way to do that is to make our rockets faster by making advancements in propulsion technologies. With the new NASA directive of returning to the Moon in 2024 as a stepping stone to missions to Mars and beyond, there is a renewed interest in developing nuclear thermal rockets (NTR) to reduce trip times. This thesis will focus on the analysis of a conceptual reactor design for use as an NTR. The low enriched uranium centrifugal gas core reactor (CGCR) is a low technology readiness level (TRL) concept that uses centrifuge technology to separate uranium gas from hydrogen propellant. There is also a new US directive for additional focus on research and development of reactors that utilize low enriched uranium (LEU) of enrichment less than 20% instead of high enriched uranium (HEU). Some NASA studies suggest that in order to accomplish a round-trip mission to Mars, a high thrust engine with a specific impulse,  $I_{sp}$ , greater than 1300 seconds may be required.[4][5] Specific impulse is defined as the total impulse delivered per unit of propellant consumed, and is equivalent to the generated thrust divided by the mass flow rate.

Modern chemical rockets have high thrust, but still require an enormous amount of fuel, having an  $I_{sp} \sim 500s$ , making them less efficient for longer journeys. Nuclear thermal propulsion (NTP) is an alternative to chemical rockets that provide a boost to the  $I_{sp}$  numbers. Current NTP designs estimate  $I_{sp}$  double the most advanced chemical rockets at comparable thrust values. In addition to the resulting

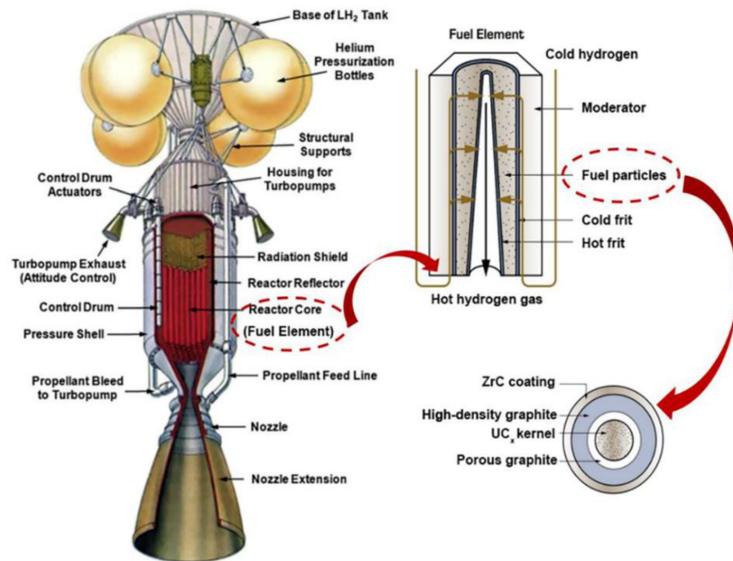
faster transit times, NTP engines could double as or be converted for electric power generation, as well as being expected to double or triple payload capacity compared to chemical propellants.

First generation NTP engines are limited to an  $I_{sp} \sim 900$ s, with designs for up to  $\sim 2000$ s. However, no significant work has been performed since the 1970s, due to changes in NASA policies and the spectre that comes with the words "nuclear" and "rocket". With this renewed initiative to continue work on NTP systems, new generation concepts are needed; particularly, ones that build on 1st Generation designs and experience.

One approach is to build on designs used in the US Space Nuclear Thermal Propulsion (SNTP) program. These designs used relatively large nuclear fuel elements in a moderator block where propellant would flow through the fuel elements. Nuclear fuel in the SNTP engine design was held in place using a "cold frit" and a "hot frit" to allow hydrogen to flow radially inward through the fuel and then exit axially at high temperature.[4] This can be seen in Figures 1.1 & 1.2.



(a)



(b)

Figure 1.1: Schematic view of a particle bed reactor (PBR) for propulsion. (a) Individual PBR core; (b) fuel elements-assembled PBR core [1]

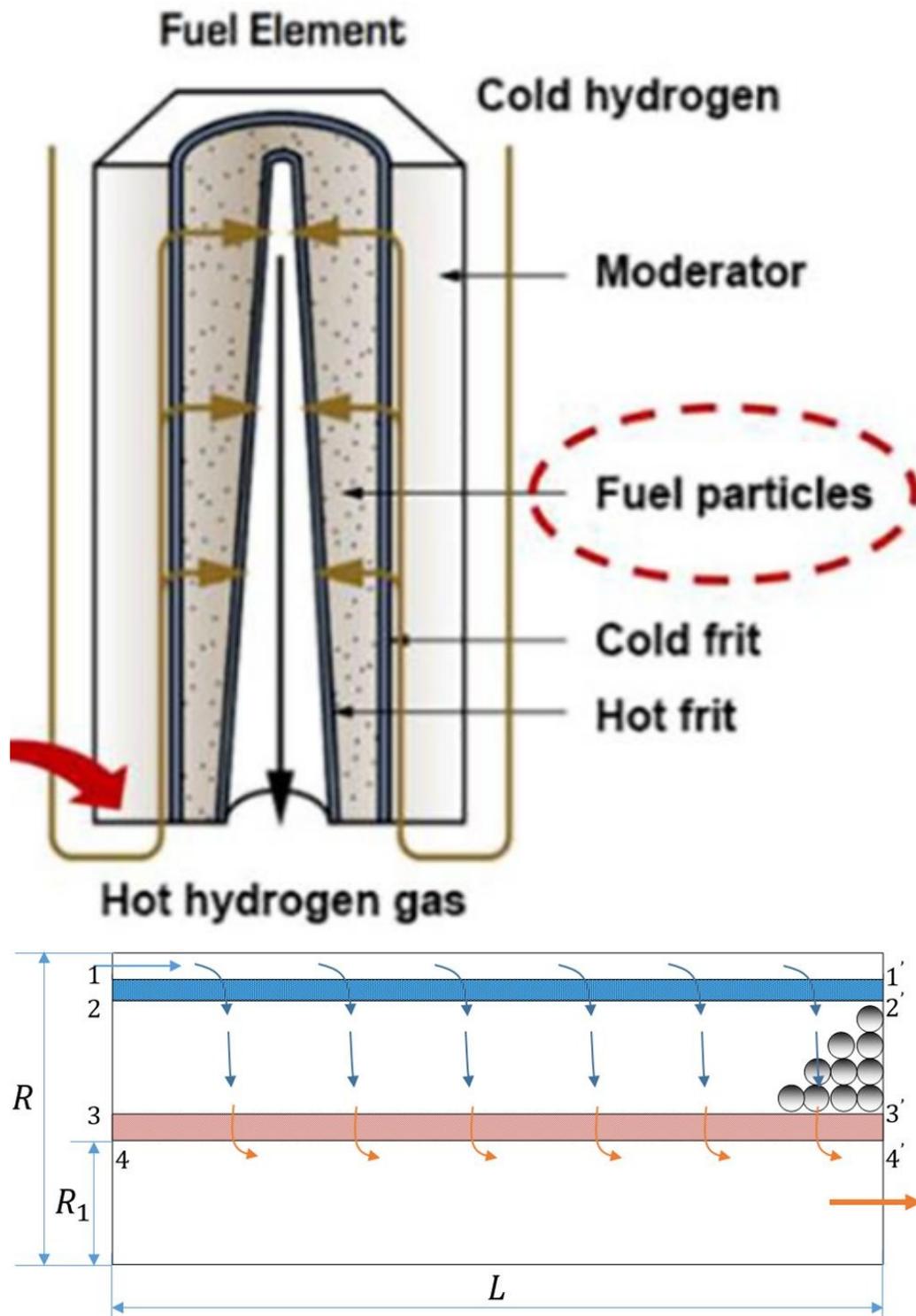


Figure 1.2: Schematic view of a particle bed reactor (PBR) fuel element and its simplification [1]

A similar concept for a 2nd Generation NTP system would utilize centrifugal force as a means of fuel containment instead of the "hot frit" element, allowing for portions of the fuel to operate beyond its melting point. This centrifuge concept could yield good separation between propellant and nuclear fuel. This particular design is the focus of this research and can be seen in Figures 1.3 & A.1.

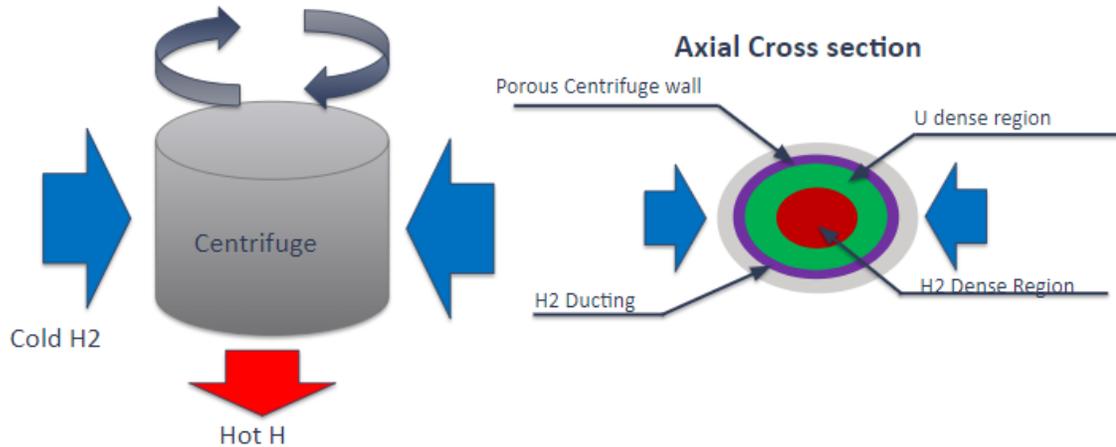


Figure 1.3: Rotating radial in-flow fuel element (no hot frit)[2]

The centrifugal gas core reactor can be explored in theory by initially looking at a model of the heat transfer, growth, and dynamics of radially-injected hydrogen bubbles in a rotating finite multi-phase fluid system under zero-gravity. As the CGCR is a nuclear reactor, it is necessary to determine its criticality, and having a proper density profile of nuclear fuel is highly important in this determination. This problem is coupled in nature: the density profile as a function of temperature, which is a function of nuclear heating, which is determined by the density profile of the nuclear fuel. Therefore, an iterative approach must be employed initially to determine the proper density profile.

## 1.2 Problem Statement

The design of this reactor fuel element, as seen in Figures 1.4 & 1.5, consists of coaxial annular cylinders of uranium metal housed by a porous cold frit (graphite), hydrogen coolant region, and a reflector/moderator/pressure vessel material, with a central hot hydrogen plenum. Rotating elements are rotating with angular velocity  $\Omega$ . Cold hydrogen will flow radially inward through the cold frit, through the uranium fuel layer, and then finally exiting axially at high temperature.

### Rotating Radial In-Flow Fuel Element

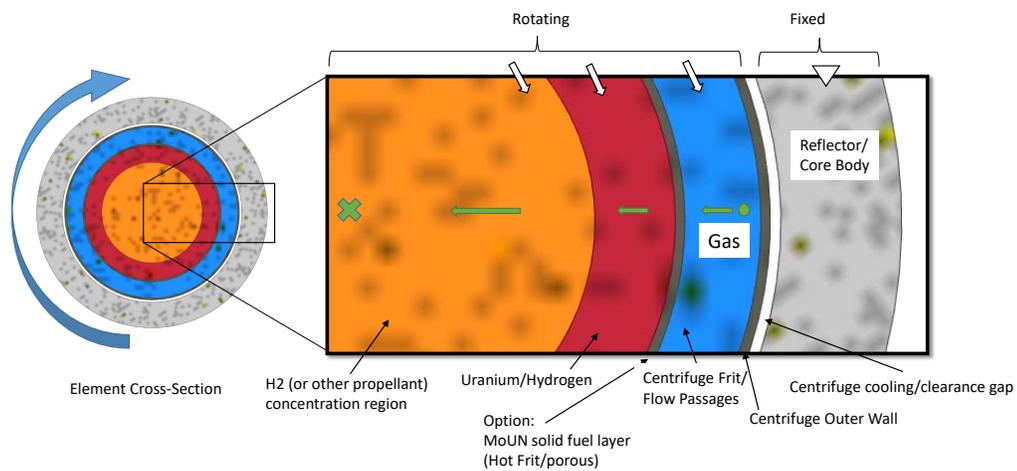


Figure 1.4: Rotating radial in-flow fuel element (no hot frit)[2]

## Temperature Map

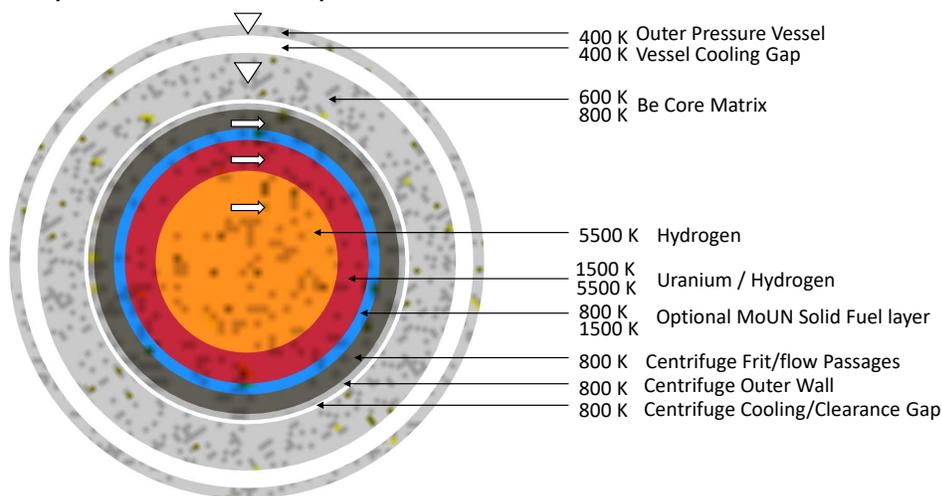


Figure 1.5: Temperature map of rotating radial in-flow fuel element (no hot frit)[2]

In this model, Figure 1.6, a hydrogen bubble is traveling radially inward through a layer of high density multi-phase fissioning uranium metal, passing into a central hydrogen plenum. The reactor is designed with a Beryllium pressure vessel of  $OD = 100cm$  and length  $110cm$ , a  $Li_7H$  reflector of thickness  $1cm$  and  $OD = 14.2cm$ , a hydrogen coolant region of thickness  $1mm$  and  $OD = 12.2cm$ , a porous graphite frit with  $H_2$  ducting of thickness  $1cm$  and  $OD = 12cm$ , LEU uranium region (19.75% enrichment) with  $OD = 10cm$ , hot dense hydrogen region  $OD = 6cm$ ; the non-Beryllium region is  $100cm$  in length. Where  $r_u$  is the outer radius of the uranium layer,  $r_{u_f}$  is the location of the interface between solid and liquid uranium,  $r_{u_v}$  is the location of the interface between liquid and vapor uranium, and  $r_p$  is the location of the interface between uranium and the hot hydrogen plenum.

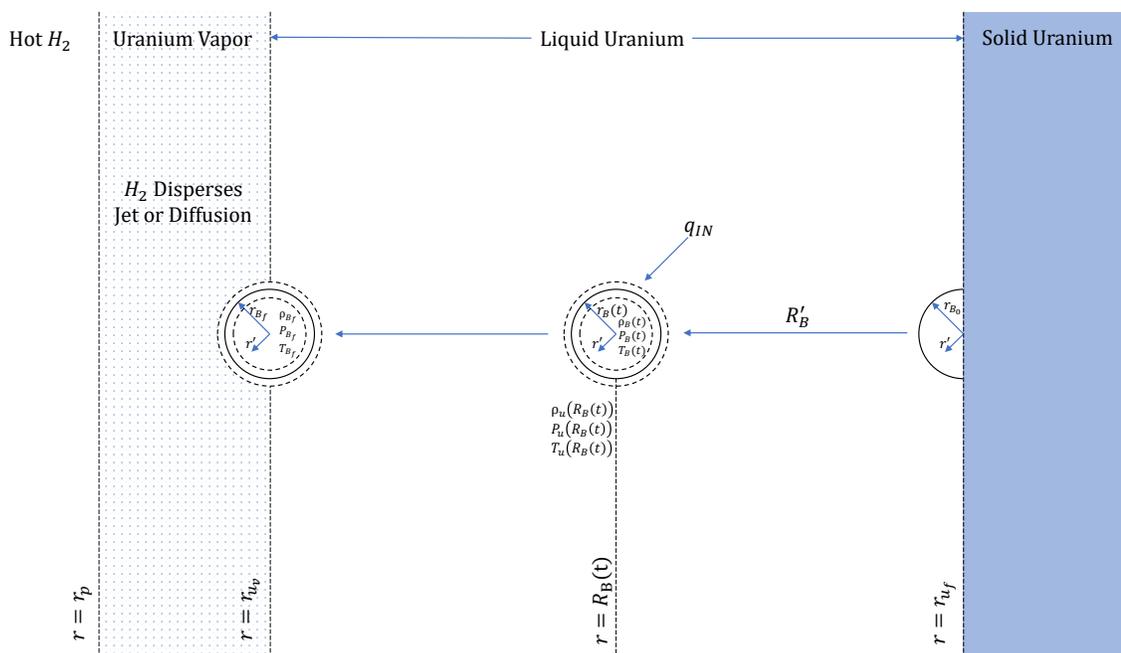


Figure 1.6: Hydrogen bubble traveling through uranium layer

## CHAPTER 2: MODEL: Theory and Methods

This chapter will demonstrate the building of a model framework that describes several coupled equations that will be solved numerically for a single hydrogen bubble traveling through the liquid uranium layer. In reality there will be more than one bubble traveling the same streamline through the uranium layer, as new bubbles will be formed as each bubble travels radially inward away from the site of injection; however, until a basic model of a single bubble is determined, bubble-bubble interactions will not be discussed. A more accurate model, for the purposes and use as an NTR, may in fact be a column or jet of hydrogen, but as an initial model, a single spherical bubble will be discussed. For this initial model, it is assumed that once the bubble reaches the uranium vapor, the bubble will disperse and become well-mixed within the uranium vapor. Molecular Hydrogen ( $H_2$ ) is considered, however dissociation may need to be explored in later models. To take advantage of the rotational symmetry of the cylindrical reactor, this model will be built in a cylindrical coordinate system. The desired model seeks to simultaneously calculate the uranium layer temperature, density, and pressure profiles; the bubble temperature, position, and radius; as well as solve for the nuclear fission effective multiplication factor, and the energy deposition rate. The variables will be defined as  $T_u(r)$ ,  $\rho_u(r)$ ,  $P_u(r)$ ,  $T_B(t)$ ,  $R_B(t)$ ,  $r_B(t)$ ,  $k_{eff}$ , and  $E_d$ , respectively. Fluid dynamics, heat transfer, and neutronics must be explored in order to determine the performance of this reactor concept.

## 2.1 Bubble Energy Balance

A good starting point for this model will begin by developing a governing equation that includes many of the above unknowns. Starting from the energy balance on the bubble, we have:

$$\begin{aligned} \text{a)} \quad & dt \int_{V_B(t)} \rho_B c_{pB} T_B dV = \oint_{S_B(t)} \mathbf{q} \cdot \hat{n} dA \\ \text{b)} \quad & c_{pB} \frac{d}{dt} (\rho_B T_B) V_B = q_{IN}(t) 4\pi r_B^2(t) + \frac{dP_B}{dt} V_B \end{aligned}$$

$$\text{Given that } q_{IN} = h[T_u(R_B(t)) - T_B(t)]$$

$$c_{pB} \frac{d}{dt} (\rho_B T_B) V_B = h[T_u(R_B(t)) - T_B(t)] 4\pi r_B^2(t) + \frac{dP_B}{dt} V_B \quad (2.1)$$

where  $h, c_{pB}, \rho_B, R_B, r_B, T_B, T_u(R_B(t))$  are the convective heat transfer coefficient, specific heat of the bubble, bubble density, bubble position relative to plenum center-line, bubble radius, bubble temperature, and uranium temperature at  $R_B(t)$ , respectively, and the bubble volume is  $V_B(t) = \frac{4}{3}\pi r_B^3(t)$  and the bubble surface area is  $S_B(t) = 4\pi r_B^2(t)$ . Equation (2.1) assumes that the hydrogen is well-mixed at all times ( $t$ ) within the bubble, the uranium temperature ( $T_u$ ) is uniform around bubble, and the bubble is spherical.

Assuming  $H_2$  behaves as a real gas:

$$\rho_B(t) = \frac{P_B(t)}{Z R_{H_2} T_B(t)} \quad (2.2)$$

$$R_{H_2} = \frac{\bar{R}}{MW_{H_2}} \quad (2.3)$$

where  $R_{H_2}$  is the specific gas constant of hydrogen,  $\bar{R}$  is the ideal gas constant, and  $Z$  is the compressibility factor of hydrogen.

If we take  $t = 0$  to be the instant that the bubble is injected into the uranium layer, then the initial conditions can be set as  $P_B(t = 0) = P_0$ ,  $T_B(t = 0) = T_0$ ,  $\rho_B(t = 0) = \frac{P_0}{Z R_{H_2} T_0}$ ,  $r_B(t) = r_{B0}$ , and  $R_B(t = 0) = r_{uf}$ , where  $r_{uf}$  is the location of

the interface between solid and liquid uranium. We now have multiple time-dependent unknowns:  $T_B(t)$ ,  $P_B(t)$ ,  $T_u(R_B(t))$ ,  $R_B(t)$ , and  $r_B(t)$ .

In order to tackle the above unknowns, we can start with the Young-Laplace equation:

$$P_B(t) - P_u(R_B(t)) = \frac{2\sigma_{st}}{r_B(t)} \quad (2.4)$$

where  $P_u(R_B(t))$  is the pressure in the uranium layer at  $R_B(t)$ , and  $\sigma_{st}$  is the surface tension coefficient for the uranium-hydrogen interface. At the melting point, the surface tension is approximately 1500 dynes/cm according to [6] and [7].

Equation (2.4) introduces a new unknown  $P_u(r)$ , which we can assume  $P_u(r)$  is calculable via the Navier-Stokes equations applied within the uranium layer, allowing us to determine  $r_B(t)$ .

## 2.2 U-Layer Heat Transfer

In order to find for temperature distribution  $T_u(r)$ , which is assumed to be steady-state, we will need to use the energy equation [8][9] applied within the uranium layer:

$$\rho_u c_{p_u} \left[ \frac{u_\theta T_{u,\theta}}{r} + u_r T_{u,r} \right] = K_u \left[ T_{u,rr} + \frac{T_{u,r}}{r} \right] + \dot{S}(r) \quad (2.5)$$

where  $\rho_u$ ,  $c_{p_u}$ ,  $K_u$  are the density, specific heat, and thermal conductivity of uranium, respectively;  $\dot{S}(r)$  is the volumetric heating term due to nuclear heating;  $u_\theta = u_\theta(r)$  is the  $\theta$ -velocity component in the uranium layer;  $u_r$  is the r-velocity component in the uranium layer, which we assume is negligible, and therefore:

$$u_r = 0 \quad (2.6)$$

If we assume the fluid cylinder has rotational symmetry, then  $T_u$  is symmetric:

$$\Rightarrow T_{u,\theta} = 0 \quad (2.7)$$

Equation (2.5) assumes  $K_u$  is fixed across u-layer, however since  $\Delta T_u$  is large, it may be worth exploring a variable  $K_u$  in further study via  $\Phi_q = -K\nabla T$ . In this model, we will also neglect axial variation in  $T_u$ . Using the above assumptions, equation (2.5) reduces to:

$$\frac{d^2 T_u}{dr^2} + \frac{1}{r} \frac{dT_u}{dr} + \frac{\dot{S}(r)}{K_u} = 0 \quad (2.8)$$

For any given  $\dot{S}(r)$ , (2.8) can be solved numerically, or analytically, if  $\frac{d\dot{S}(r)}{dr} \ll 1$ , by further assuming  $\dot{S}(r) \approx \text{constant} \equiv \dot{S}_0$ . Then (2.8) becomes:

$$T_u'' + \frac{1}{r} T_u'' + \frac{\dot{S}_0}{K_u} = 0 \quad (2.9)$$

### 2.3 U-Layer Fluid Dynamics

In order to determine  $r_B(t)$  in equation (2.4) we first need to find the pressure distribution in the U-Layer,  $P_u(r)$ .

a) Working in cylindrical coordinates, the Navier-Stokes equation for the azimuthal component  $\theta$  [9]:

$$\rho_u \left[ u_r u_{\theta,r} + u_z u_{\theta,z} + \frac{u_\theta}{r} u_{\theta,\theta} + \frac{u_r u_\theta}{r} \right] = -\frac{1}{r} \frac{\partial P_u}{\partial \theta} + \mu_u \left[ u_{\theta,rr} + \frac{u_{\theta,r}}{r} - \frac{u_\theta}{r^2} \right]$$

Assuming  $u_z$  is negligible,  $u_{,rr} = u_{,zz} = 0$ , symmetry of the velocity field, neglecting temperature-dependence of  $\mu_u$  (uranium viscosity), neglecting presence of hydrogen bubbles, and taking into account equation (2.6), the above equation reduces to:

$$\Rightarrow \frac{1}{r} \frac{d}{dr} \left( r \frac{du_\theta}{dr} \right) - \frac{u_\theta}{r^2} = 0 \quad (2.10)$$

Solving (2.10) analytically gives:

$$u_\theta(r) = r\Omega_0 \quad (2.11)$$

where  $\Omega_0$  is the angular velocity of the cylinder.

b) From the  $r$ -component of the Navier-Stokes equation:

$$-\rho_u \frac{u_\theta^2}{r} = -\frac{\partial P_u}{\partial r} \quad (2.12)$$

and since  $P_u =$  function of  $r$  only:

$$\Rightarrow \frac{\partial P_u}{\partial r} = \frac{dP_u}{dr} \quad (2.13)$$

As the  $u$ -layer consists of solid, liquid, and gas, densities of each region will be different:

$$\rho_u(r) = \begin{cases} \rho_{u_s}(r) = \text{Constant} & r_{u_f} \leq r \leq r_u \\ \rho_{u_l}(r) = 17270 - 1.4485(T_u(r) - 1408) & r_{u_v} \leq r < r_{u_f} \\ \rho_{u_v}(r) = \frac{P_u(r)}{R_u T_u(r)} & r_p \leq r < r_{u_v} \end{cases} \quad (2.14)$$

where  $R_u$  is the specific gas constant for uranium, and the density for liquid uranium comes from [10][11].

Using (2.11), (2.13), and the vapor region of (2.14), we can rearrange (2.12) in terms of  $\rho_{u_v}(r)$ :

$$\rho_{u_v}(r)r\Omega_0^2 = \frac{d}{dr}(\rho_{u_v}(r)R_u T_u(r))$$

Solving for  $\rho_{u_v}(r)$  gives:

$$\rho_{u_v}(r) = C \exp \left[ \frac{\Omega_0^2 r^2}{2R_u T_u(r)} \right], \quad r_p \leq r < r_{u_v} \quad (2.15)$$

Given the B.C.s of:

$$P_{u_v}(r_{u_v}) = P_{u_l}(r_{u_v})$$

$$\rho_{u_v}(r_{u_v}) = \frac{P_{u_v}(r_{u_v})}{R_u T_u(r_{u_v})}$$

we can solve for the constant in (2.15), giving:

$$\rho_{u_v}(r) = \frac{P_{u_v}(r_{u_v})}{R_u T_u(r_{u_v})} \exp \left[ \frac{\Omega_0^2}{2R_u} \left[ \frac{r^2}{T_u(r)} - \frac{r_{u_v}^2}{T_u(r_{u_v})} \right] \right], \quad r_p \leq r < r_{u_v} \quad (2.16)$$

Plugging (2.16) back into (2.14) and using (2.12) to solve for  $P_u(r)$ , we find:

$$\rho_u(r) = \begin{cases} \text{Constant} & r_{u_f} \leq r \leq r_u \\ 17270 - 1.4485(T_u(r) - 1408) & r_{u_v} \leq r < r_{u_f} \\ \frac{P_u(r_{u_v})}{R_u T_u(r_{u_v})} \exp \left[ \frac{\Omega_0^2}{2R_u} \left[ \frac{r^2}{T_u(r)} - \frac{r_{u_v}^2}{T_u(r_{u_v})} \right] \right] & r_p \leq r < r_{u_v} \end{cases} \quad (2.17)$$

$$P_u(r) = \begin{cases} \text{Constant} & r_{u_f} \leq r \leq r_u \\ \int \rho_u(r) r \Omega_0^2 dr & r_{u_v} \leq r < r_{u_f} \\ \rho_u(r) R_u T_u(r) & r_p \leq r < r_{u_v} \end{cases} \quad (2.18)$$

This analysis of rotating immiscible gases can generalize the above density and pressure functions to:

$$\rho_j(r, \Omega) = C_j \exp[\Omega^2 r^2 / 2R_j T_j(r)] \quad (2.19)$$

$$P_j(r, \Omega) = \rho_j(r, \Omega) R_j T_j(r) \quad (2.20)$$

where  $j$  represents differing gas regions as seen in Figure 2.1.

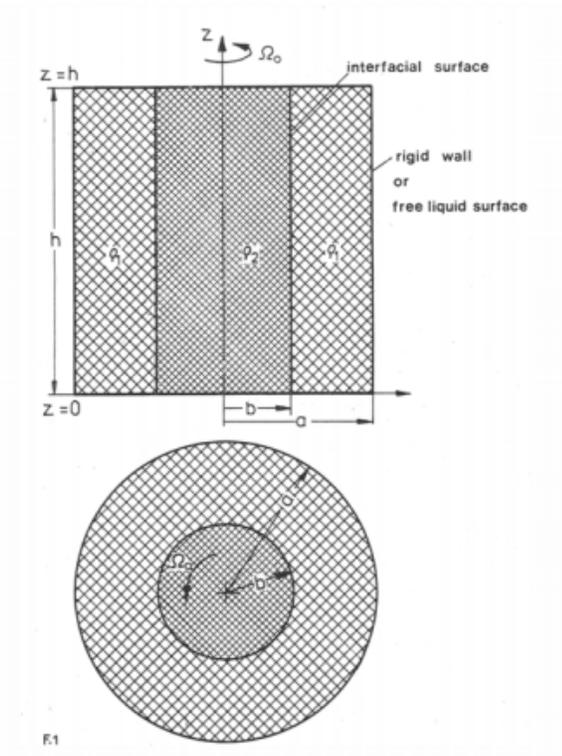


Figure 2.1: Rotating immiscible fluids [3]

When looking at the interaction between the gaseous uranium and the hydrogen plenum at the interface  $r = r_p$ ,

$$P_u(r_p) = P_{H_2}(r_p)$$

And thus equation (2.19) for each gas yields:

$$\rho_{u_v}(r, \Omega_0) = \rho_u(r_p) \exp \left[ \frac{\Omega_0^2}{2R_u T_u(r_p)} (r^2 - r_p^2) \right], \quad r_p \leq r < r_{u_v} \quad (2.21)$$

$$\rho_{H_2}(r, \Omega_0) = \frac{P_u(r_p)}{R_{H_2} T_{H_2}(r_p)} \exp \left[ \frac{\Omega_0^2}{2R_{H_2} T_{H_2}(r_p)} (r^2 - r_p^2) \right], \quad 0 \leq r \leq r_p \quad (2.22)$$

Given the acceleration potentials for each region from Bauer [3], which consist of pressure and centrifugal terms:

$$\Psi_u = \frac{P_u - P_0}{\rho_u} - \frac{1}{2}\Omega_0^2(r^2 - r_p^2) - \frac{1}{2}\Omega_0^2\frac{\rho_{H_2}}{\rho_u}r^2 \quad (2.23)$$

$$\Psi_{H_2} = \frac{P_{H_2} - P_0}{\rho_{H_2}} - \frac{1}{2}\Omega_0^2r^2 \quad (2.24)$$

At the interfacial surface  $r = r_p$ , the normal velocities of the U and  $H_2$  regions are equal, and therefore the free surface boundary condition is given by:

$$u_u = u_{H_2} \text{ at the interface } r = r_p$$

$$\begin{aligned} \rho_{H_2}(r_p)\frac{\partial\Psi_{H_2}}{\partial t} - \rho_u(r_p)\frac{\partial\Psi_u}{\partial t} + (\rho_{H_2}(r_p) - \rho_u(r_p))r_p\Omega_0^2u_{H_2} \\ + \frac{\sigma_{stp}}{r_p^2} \left( u_{H_2} + \frac{\partial^2 u_{H_2}}{\partial\theta^2} + r_p^2\frac{\partial^2 u_{H_2}}{\partial z^2} \right) = 0 \end{aligned} \quad (2.25)$$

where  $\sigma_{stp}$  is the interfacial tension at  $r = r_p$  between gases U and  $H_2$ . If we assume steady state and that  $u_{H_2,zz} = u_{H_2,\theta\theta} = 0$ , then solving for  $r_p$  yields:

$$r_p = \left[ \frac{\sigma_{stp}}{\Omega_0^2\rho_u(r_p) \left( 1 - \frac{\rho_{H_2}(r_p)}{\rho_u(r_p)} \right)} \right]^{\frac{1}{3}} \quad (2.26)$$

$$\frac{\rho_{H_2}}{\rho_u} \ll 1$$

$$r_p = \left[ \frac{\sigma_{stp}}{\Omega_0^2\rho_u(r_p)} \right]^{\frac{1}{3}} \quad (2.27)$$

This can also be stated in terms of the rotational Bond number  $\tilde{B}_0$ :

$$\tilde{B}_0 = \Omega_0^2 r_{uv}^3 \rho_u(r_p) / \sigma_{stp}$$

$$r_p = \frac{r_{uv}^3}{\tilde{B}_0} \quad (2.28)$$

## 2.4 Bubble Dynamics

Due to the pressure gradient within the uranium layer, a bubble injected radially inward will see a buoyant force, pushing the bubble inward from  $r = r_{u_f}$  to  $r = r_{u_v}$ . Given  $P_u(r)$ , we can calculate the instantaneous radial pressure force,  $F_{B_P}(t)$ .

$$\begin{aligned}
 F_{B_P}(t) &= - \oint_{S_B(t)} P_u \hat{n} \cdot \hat{e}_r dA \\
 &= - \int_{V_B(t)} \nabla \cdot (P_u(r) \hat{e}_r) dV \\
 &= - \int_{V_B(t)} \left[ \hat{e}_r \frac{\partial}{\partial r} + \frac{\hat{e}_\theta}{r} \frac{\partial}{\partial \theta} \right] \cdot (P_u(r) \hat{e}_r) 4\pi r^2 dr \\
 &= - \int_0^{r_B(t)} \left[ \frac{dP_u(r)}{dr} + \frac{P_u(r)}{r} \right] 4\pi r^2 dr
 \end{aligned}$$

As  $P_u(r)$  will be evaluated over  $r = R_B(t) + r'$ :

$$F_{B_P}(t) = - \int_0^{r_B(t)} \left[ \frac{dP_u(r)}{dr} \Big|_{r=R_B(t)+r'} + \frac{P_u(r)|_{r=R_B(t)+r'}}{R_B(t) + r'} \right] 4\pi r'^2 dr' \quad (2.29)$$

The bubble is also subject to a drag force  $F_{B_D}(t)$  and a centrifugal force  $F_{B_C}(t)$ :

$$F_{B_D}(t) = -6\pi\mu_u r_B(t) \frac{dR_B}{dt} \quad (2.30)$$

$$F_{B_C}(t) = M_B R_B(t) \Omega_0^2 \quad (2.31)$$

where  $M_B$  is the mass of the hydrogen bubble, which is a fixed quantity.

The general form of drag on a sphere described by Stoke's Law consists of two components: a normal force  $F_{B_n}$  due to pressure acting perpendicularly to the surface,

and a tangential force  $F_{B_t}$  due to shear stress.

$$\begin{aligned} F_{B_D} &= F_{B_n} + F_{B_t} \\ F_{B_n} &= -2\pi\mu_u r_B(t) \frac{dR_B}{dt} \\ F_{B_t} &= -4\pi\mu_u r_B(t) \frac{dR_B}{dt} \end{aligned}$$

If we neglect tangential drag, as the bubble sphere surface is gas and it is also assumed that there is no relative motion in the tangential direction between the bubble and the uranium layer ( $v_{\theta_u} = v_{\theta_B} = \Omega r$ ), (2.30) reduces to drag due to pressure:

$$\Rightarrow F_{B_D}(t) = -2\pi\mu_u r_B(t) \frac{dR_B}{dt} \quad (2.32)$$

Adding equations (2.29)-(2.32), we now have the total radial force on the bubble:

$$F_B(t) = F_{B_p}(t) + F_{B_D}(t) + F_{B_c}(t) \quad (2.33)$$

Now we can write Newton's second law governing the bubble's radial motion:

$$M_B \frac{d^2 R_B}{dt^2} = F_B(t) \quad (2.34)$$

$$M_B = \rho_0 V_B(t = 0)$$

$$M_B = \left( \frac{P_0}{Z R_{H_2} T_0} \right) \left( \frac{4}{3} \pi \right) r_{B_0}^3 \quad (2.35)$$

As (2.34) contains both  $R_B(t)$  and  $r_B(t)$ , we can revisit the Young-Laplace equation (2.4):

$$r_B(t) = \frac{2\sigma_{st}}{P_B(t) - P_u(R_B(t))} \quad (2.36)$$

The initial bubble radius  $r_{B_0}$  can be found from Blottner[12] and simultaneously

solving for  $\Omega_0$  by utilizing the Young-Laplace equation at the solid/liquid uranium interface, setting  $R_B(0) = r_{uf}$ , and  $P_B(R_B(0)) = P_{B_0}$ :

$$r_B = C * A$$

where  $C = 3.97$  and  $A$  is the Laplace constant:

$$A = \left[ \frac{\sigma_{st}}{g(\rho_l - \rho_g)} \right]^{1/2}$$

where  $g = R_B(t)\Omega_0^2$ ,  $\rho_l = \rho_u$ , and  $\rho_g = \rho_B$ .

$$\left\{ \begin{array}{l} r_{B_0} = C * \left[ \frac{\sigma_{st}}{r_{uf}\Omega_0^2(\rho_u(r_{uf}) - \rho_{B_0})} \right]^{1/2} \\ P_{B_0} - P_u(r_{uf}, \Omega_0) = \frac{2\sigma_{st}}{r_{B_0}} \end{array} \right. \quad (2.37a)$$

$$(2.37b)$$

$P_B(t)$  can be found via (2.36):

$$M_B = \left( \frac{P_B(t)}{ZR_{H_2}T_B(t)} \right) \left( \frac{4}{3}\pi \right) r_B^3(t)$$

$$P_B(t) = \frac{3M_B R_{H_2} Z T_B(t)}{4\pi r_B^3(t)} \quad (2.38)$$

Plugging (2.38) back into (2.36) yields:

$$r_B(t) = \frac{2\sigma_{st}}{\frac{C_0 Z T_B(t)}{r_B^3(t)} - P_u(R_B(t))} \quad (2.39)$$

where  $C_0 = \frac{3M_B R_{H_2}}{4\pi}$

Equation (2.39) describes the bubble growth, and requires both  $R_B(t)$  and  $T_B(t)$ .

Thus,  $r_B(t)$ ,  $R_B(t)$ , and  $T_B(t)$  are all coupled.

## 2.5 Final Model

Putting everything together, this model seeks to simultaneously calculate  $T_u(r)$ ,  $\rho_u(r)$ ,  $P_u(r)$ ,  $T_B(t)$ ,  $\rho_B(t)$ ,  $P_B(t)$ ,  $R'_B(t)$ ,  $R_B(t)$ , and  $r_B(t)$  given the following governing equations:

$$(2.1) \quad c_{pB} \frac{d}{dt}(\rho_B T_B) = h[T_u(R_B(t)) - T_B(t)]4\pi r_B^2(t)$$

$$(2.8) \quad \frac{d^2 T_u}{dr^2} + \frac{1}{r} \frac{dT_u}{dr} + \frac{\dot{S}(r)}{K_u} = 0$$

$$(2.10) \quad \frac{1}{r} \frac{d}{dr} \left( r \frac{du_\theta}{dr} \right) - \frac{u_\theta}{r^2} = 0$$

$$(2.12) \quad -\rho_u \frac{u_\theta^2}{r} = -\frac{\partial P_u}{\partial r}$$

$$(2.33) \quad F_B(t) = - \int_0^{r_B(t)} \left[ \frac{dP_u(r)}{dr} \Big|_{r=R_B(t)+r'} + \frac{P_u(r) \Big|_{r=R_B(t)+r'}}{R_B(t) + r'} \right] 4\pi r'^2 dr' \\ - 2\pi \mu_u r_B(t) \frac{dR_B}{dt} + M_B R_B(t) \Omega_0^2$$

$$(2.34) \quad M_B \frac{d^2 R_B}{dt^2} = F_B(t)$$

$$(2.39) \quad r_B(t) = \frac{2\sigma_{st}}{\frac{c_0 Z T_B(t)}{r_B^3(t)} - P_u(R_B(t))}$$

The governing equations lead to the following system of nonlinear ODEs:

$$\left\{ \begin{array}{l} R_B'' = \frac{F_B(t)}{M_B} \quad (2.40a) \\ P_B' = \frac{3h [T_u(R_B(t)) - T_B(t)]}{r_B(t) \left[ \frac{c_{pB}}{Z R_{H_2}} - 1 \right]} \quad (2.40b) \\ r_B' = \frac{-2\sigma_{st}}{[P_B(t) - P_u(R_B(t))]^2} \left[ P_B(t) - R_B' \frac{dP_u}{dR_B} \right] \quad (2.40c) \\ \rho_B' = \frac{-3C_0}{R_{H_2}} \frac{r_B'}{r_B(t)^4} \quad (2.40d) \\ T_B' = \frac{h [T_u(R_B(t)) - T_B(t)] 4\pi r_B(t)^2 + P_B' V_B(t)}{c_{pB} M_B} - \frac{\rho_B'}{\rho_B(t)} T_B(t) \quad (2.40e) \end{array} \right.$$

## 2.6 Nuclear & MCNP

As the CGCR is a nuclear reactor, it is necessary to adhere to the design parameter of being in a critical state. That is to say that the effective multiplication factor,  $k_{eff}$ , determined by the six factor formula is equal to 1:

$$k_{eff} = \frac{\text{neutrons produced by fission in one neutron generation}}{\text{number of neutrons lost through absorption in the preceding neutron generation}}$$

$$k_{eff} = \eta \cdot \varepsilon \cdot p \cdot f \cdot P_f \cdot P_t$$

where  $\eta, \varepsilon, p, f, P_f$ , and  $P_t$  represent the reproduction factor, fast fission factor, resonance escape probability, thermal utilization factor, fast non-leakage probability, and thermal non-leakage probability, respectively.

The reproduction factor is defined as the ratio of the number of fast neutrons produced by thermal fission to the number of thermal neutrons absorbed in the fuel, which determines the number of neutrons created in the new generation. The fast fission factor is defined as the ratio of the fast neutrons produced by fissions at all energies to the number of fast neutrons produced in thermal fission. The resonance escape probability is the probability that a neutron will be slowed to thermal energy and will escape resonance capture. The thermal utilization factor is the fraction of the thermal neutrons that are absorbed in the nuclear fuel. The fast non-leakage factor is defined as the ratio of the number of fast neutrons that do not leak from the reactor core during the slowing down process to the number of fast neutrons produced by fissions at all energies. And last but not least, the thermal non-leakage factor is defined as the ratio of the number of thermal neutrons that do not leak from the reactor core during the neutron diffusion process to the number of neutrons that reach thermal energies.

If  $k_{eff} > 1$ , then the reactor is supercritical and the number of neutrons is increasing

exponentially in time. If  $k_{eff} < 1$ , then the reactor is subcritical and the number of neutrons is decreasing and therefore the chain reaction will never be self-sustaining.

The other critical relation necessary in this problem is nuclear heating, or the nuclear energy deposition rate [13][14]:

$$E_d = \frac{N}{V\rho} \int_V \int_t \int_E H(E)\Phi(\vec{r}, E, t)dEdtdV \quad (2.41)$$

Where  $N$  and  $\rho$  are the atomic and mass densities, respectively,  $H(E)$  is the heating response, and  $\Phi(\vec{r}, E, t)$  is the particle flux. Energy deposition is in units of MeV/g. Energy deposition will be utilized for the nuclear heating term in (2.8) by scaling the raw F6 tally by the mass density of the material and by a scalar quantity  $C_i$  which has units [1/s] and is determined by the power normalization given by[13]:

$$Q_i = E_{d_i}\rho_i C_i [Watts/cm^3] \quad (2.42)$$

$$ReactorPower = Q_{tot} = \sum_{cells} Q_i V_i$$

The normalization factor can be determined by:

$$NormKCODE = \frac{\nu Q_{tot}}{(1.602 * 10^{-13})Q_{fis}k_{eff}} \quad (2.43)$$

where  $\nu$ , and  $Q_{fis}$  are the average number of neutrons per fission, and recoverable energy per fission in MeV/fission, respectively, and the units of (2.43) are (kcode source neutrons)/second. From (2.43):

$$E_d = NormKCODE * F_6 \quad (2.44)$$

Both  $k_{eff}$  and energy deposition can be calculated via radiation transport utilizing the Monte Carlo N-Particle transport package MCNP v.6.1 [14]. This package is

capable of Continuous Energy neutron and photon transport. Interaction probabilities are derived from experimentally obtained nuclear physics cross section data when available.[13] MCNP allows the problem domain to be spatially discretized for reaction rate tallying. For example, this would enable the end user to obtain an estimate for the local energy deposition inside a given volume embedded in the model. The KCODE mode will be used to solve for  $k_{eff}$  and the energy deposition rate in equation (2.41) will be determined by the F6 tally in MCNP.

The problem domain within MCNP is geometrically discretized into cells defined by Boolean combinations of spatial regions. Each cell must be defined spatially, as well by characteristic material, density, and particle importance in said cells and can be seen in the input file in Appendix B.1. The CGCR designed in MCNP can be seen in Figures 2.2 & 2.3. To be noted, the uranium region (in blue) has been subdivided into 1mm increments due to the varying density profile.

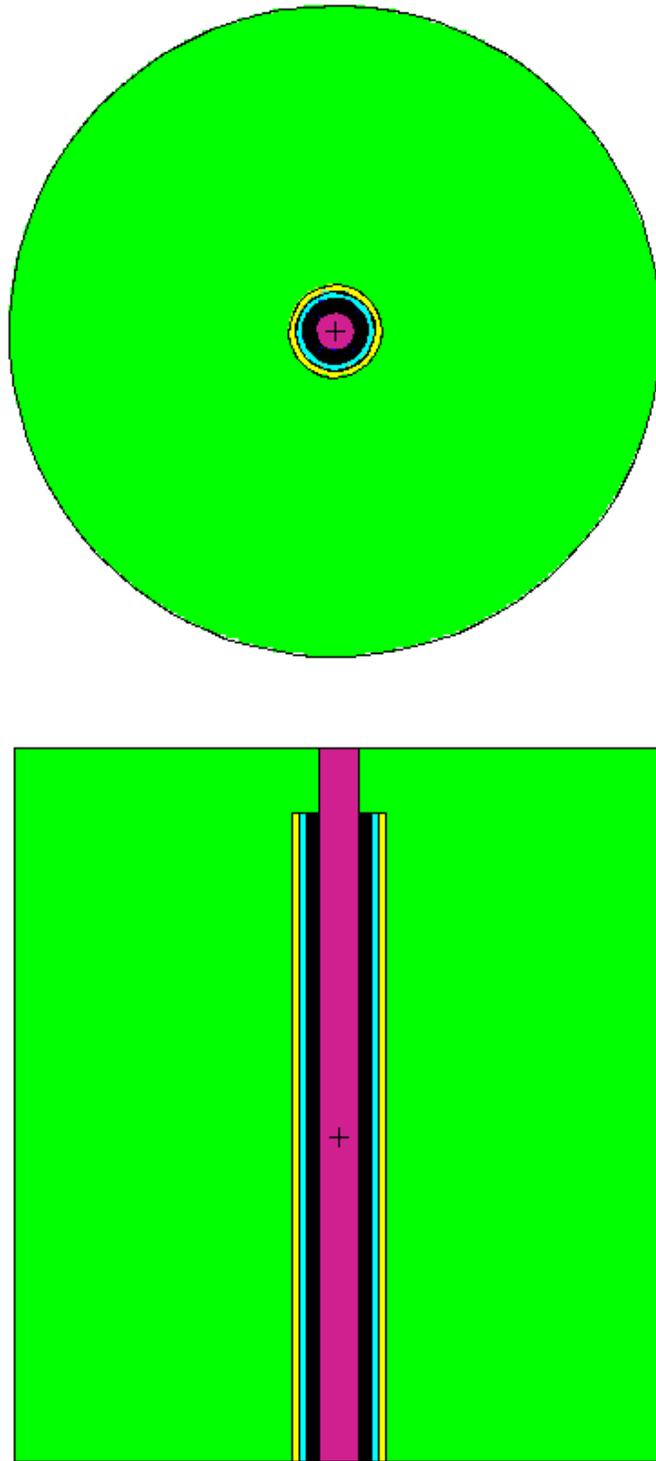


Figure 2.2: Reactor top & side cross-section centrifugal region, as designed in MCNP. Green = Beryllium, Yellow =  $Li_7H$ , Orange = cold  $H_2$ , Light Blue = Graphite frit, Blue = Uranium, Purple = hot  $H_2$

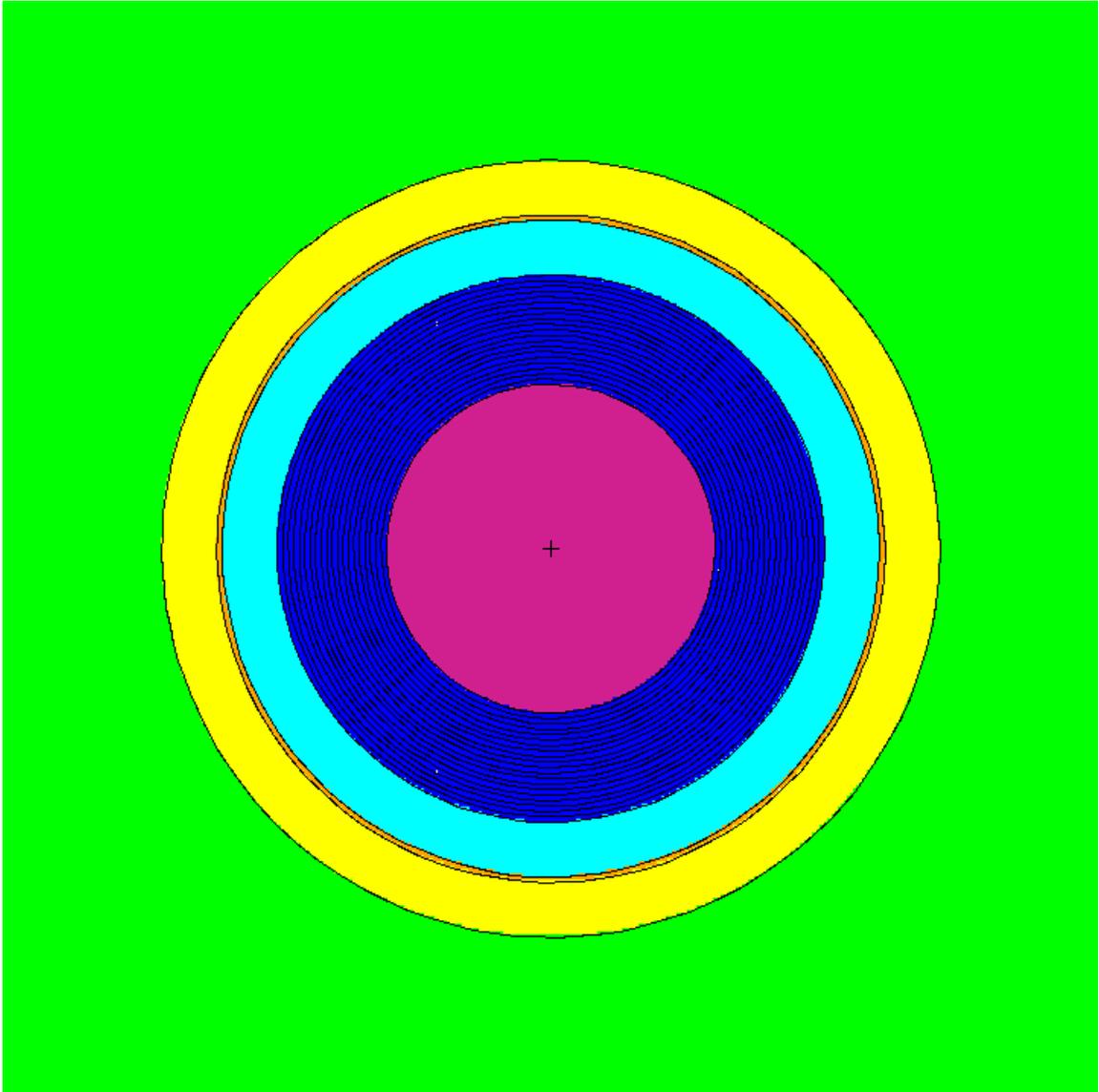


Figure 2.3: Reactor top cross-section centrifugal region, as designed in MCNP. Green = Beryllium, Yellow =  $Li_7H$ , Orange = cold  $H_2$ , Light Blue = Graphite frit, Blue = Uranium, Purple = hot  $H_2$

## CHAPTER 3: RESULTS

Utilizing analytical and numerical methods via Mathematica and MATLAB, as well as an initial reactor designed and analyzed in MCNP, a cyclic iteration method was utilized to determine temperature, density, and pressure profiles of the uranium layer  $(T_u(r), \rho_u(r), P_u(r))$ . The initial parameters were chosen to be as follows:

Table 3.1: Initial Parameters<sup>1</sup>

$T_u(r = r_u) = 800 \text{ K}$	$r_u = 5 \text{ cm}$	$K_u = 27 \text{ W/m K}$
$T_{ufus} = 1405.3 \text{ K}$	$r_p = 3 \text{ cm}$	$Q_{tot} = 10 \text{ MW}$
$T_{uwap} = 4404 \text{ K}$	$H = 100 \text{ cm}$	$Q_{fis} = 200 \text{ MeV/fission}$
$T_u(r = r_p) = 5500 \text{ K}$	$\mu_u = 6.5 \text{ cP}$	$\rho_{u_s} = 19100 \text{ kg/m}^3$
$T_{B_0} = T_{ufus}$	$\rho_{B_0} = 5 \text{ kg/m}^3$	$\rho_{B_{max}} = 10000 \text{ kg/m}^3$
$h = 1000 \text{ W/m}^2\text{K}$	$\dot{m}_{H_2} = 0.05 \text{ kg/s}$	$\sigma_{st} = 1500 \text{ dyne/cm}$
$c_{pB} = 18000 \text{ J/kg-K}$	$R_{H_2} = 4124 \text{ kg/mol}$	$R_u = 34.9328 \text{ kg/mol}$
$T_{crit_{H_2}} = 33.2 \text{ K}$	$P_{crit_{H_2}} = 12.797 \text{ atm}$	$P_0 = 30 \text{ MPa}$

The effective multiplication factor,  $k_{eff}$ , was found to have a value of  $0.884 \pm 0.001$ , and the raw energy deposition  $F_6$  was found to have a mean of  $7.254\text{e-}04 \text{ MeV/g} \pm 6.833\text{e-}04 \text{ MeV/g}$ . From the temperature profile and melting and boiling points of uranium in Figure 3.1, the locations of uranium phase change were determined to be  $r_{uf} \approx 0.0468m$  and  $r_{uv} \approx 0.0338m$ . From the density profile in Figure 3.2, the mass of the uranium in the finite cylinder of height  $H = 100cm$  was found to be  $69.114kg$ . For multiples of  $\Omega_0$  (5,10,15,20), density profiles were fed back into MCNP to find corresponding  $k_{eff} = 0.89812, 0.93220, 0.95964, 0.96092$ . This can be seen in Figures 3.3 & 3.4. From the density profile, the pressure profile was determined and is shown in Figure 3.5. The optimum angular velocity ( $\Omega_0$ ) for this setup was determined to be  $1605.36 \text{ Hz}$ . This was found by utilizing equation (2.37).

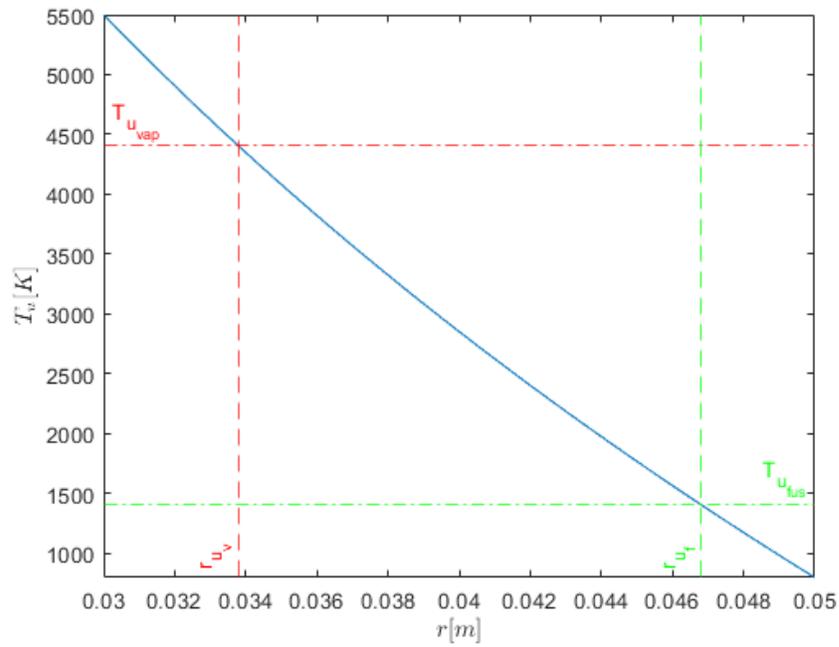


Figure 3.1: Radial temperature profile  $T_u$  within uranium layer. Melting and boiling temperatures of uranium are included to show the location of phase change.

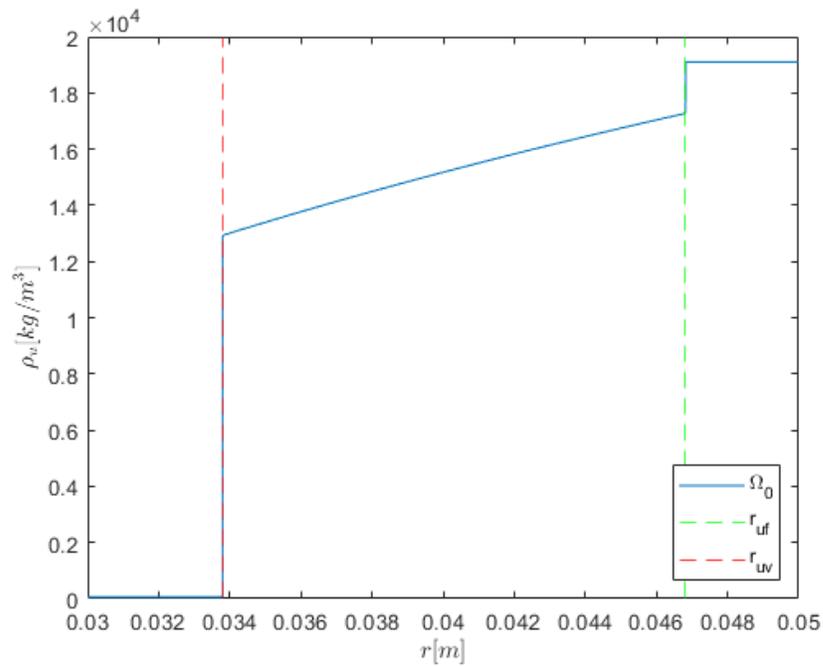


Figure 3.2: Radial density profile  $\rho_u$  for  $\Omega_0 \approx 1605$  Hz

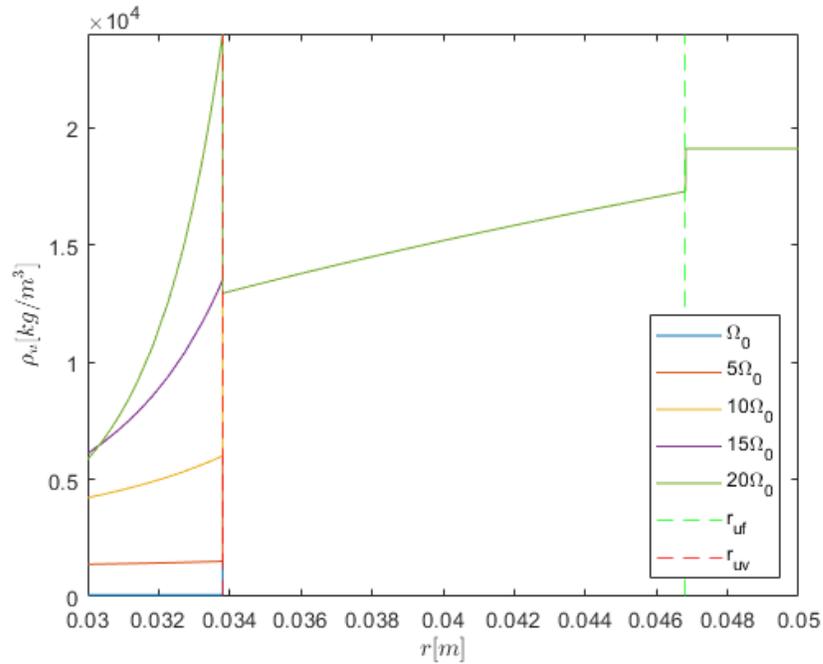


Figure 3.3: Radial density profile  $\rho_u$  for various angular velocities. Increasing angular velocity ( $\Omega$ ) results in an increased density shift radially outwards in the vapor region, as expected.

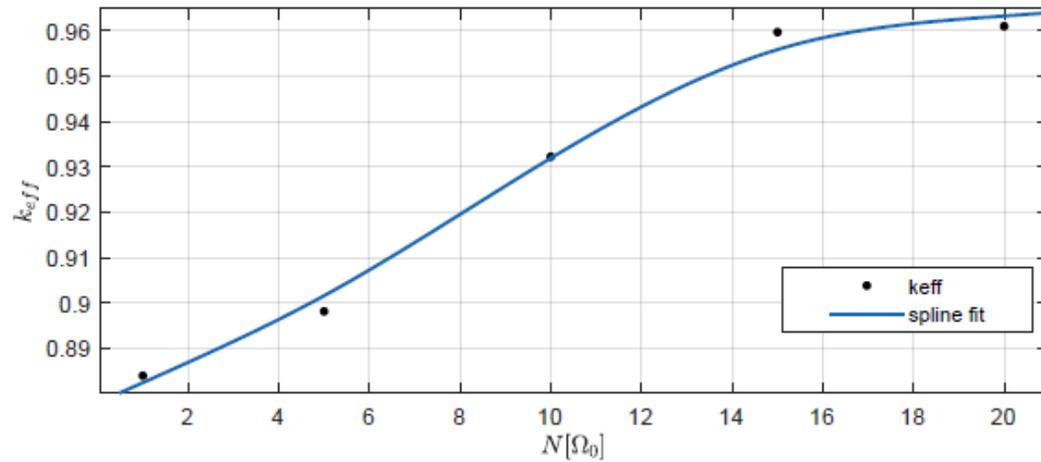


Figure 3.4:  $k_{eff}$  vs. various multiples of  $\Omega_0$ . This shows that  $k_{eff}$  increases with increasing angular velocity as more mass is allowed in uranium vapor layer.

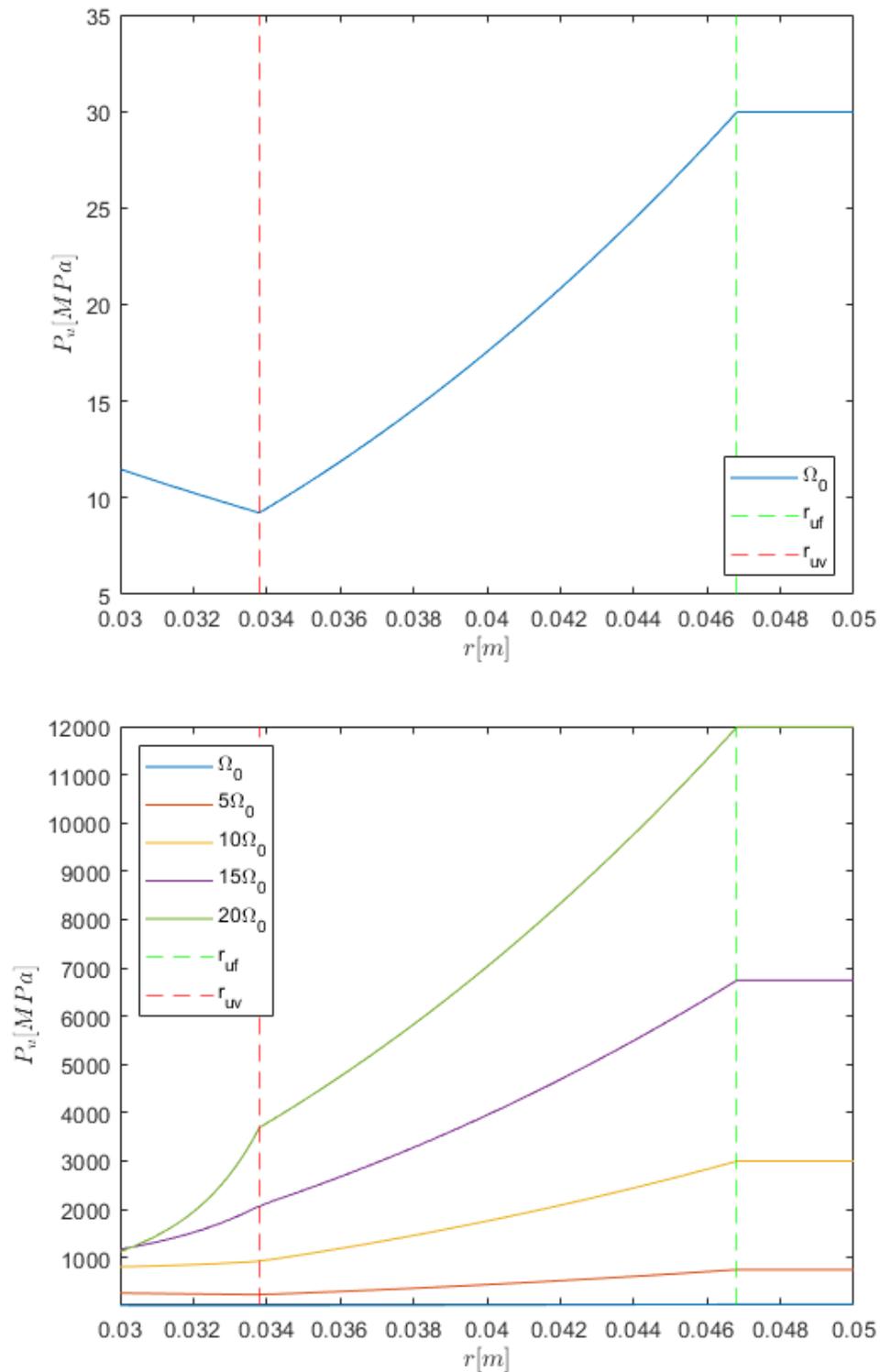


Figure 3.5: Top: Radial pressure profile  $P_u$  for  $\Omega_0 \approx 1605$  Hz. Bottom: Radial pressure profile  $P_u$  for various angular velocities. Increasing angular velocity ( $\Omega$ ) results in an increased pressure shift radially outwards in the vapor region, as well as increased pressure throughout the liquid region, as expected.

From the real gas equation (2.2) and Table 3.1,  $Z_0 = 1.0353$ . Given conservation of mass and Table 3.1, Initial velocity  $R'_B(0) = -\dot{m}_{H_2}/(2\pi H r_{uf} \rho_{B_0}) \approx -0.034\text{m/s}$ . Initial bubble radius  $r_{B_0}$  is calculated to  $\approx 101.3\mu\text{m}$ . As the bubble is injected radially inward, the solution to ODEs (2.40) suggest an immediate drop in bubble temperature and bubble radius as the bubble compresses to a minimum size  $r_{B_{min}} = 8.04\mu\text{m}$  limited by  $\rho_{B_{max}}$ , which was chosen to be on the order of metallic hydrogen assuming fusion doesn't occur. By the time the bubble traverses the liquid uranium layer and reaches  $r_{uv}$ ,  $t = 2.0881\text{ ms}$ ,  $R'_B = -3.0856\text{m/s}$ ,  $P_B = 834.795\text{ MPa}$ , and  $T_B = 18.3421\text{ K}$ . The trends can be seen in Figures 3.6-3.10.

Looking to the bubble evolution described by the governing equations (2.1)-(2.40) and the figures in the previous chapter, it can be noted the second term in equation (2.40e) is dominant until  $\rho_{B_{max}}$  is reached and  $\rho_B(t)' \Rightarrow 0$ ; once this occurs, the second term vanishes, causing  $T_B(t)'$  to become positive. Once the hydrogen bubble fully compresses, the centrifugal term in equation (2.40a) becomes dominant, thus decelerating the bubble. This therefore shows logical consistency between the bubble evolution equations and figures.

As an extension of this study, for comparison with the preliminary reference design in Figure A.1, this single fuel element design was also made into a lattice structure of 13 elements, as seen in Figure 3.11. This lattice resulted in an expected increased  $k_{eff} = 1.190 \pm 0.001$ .

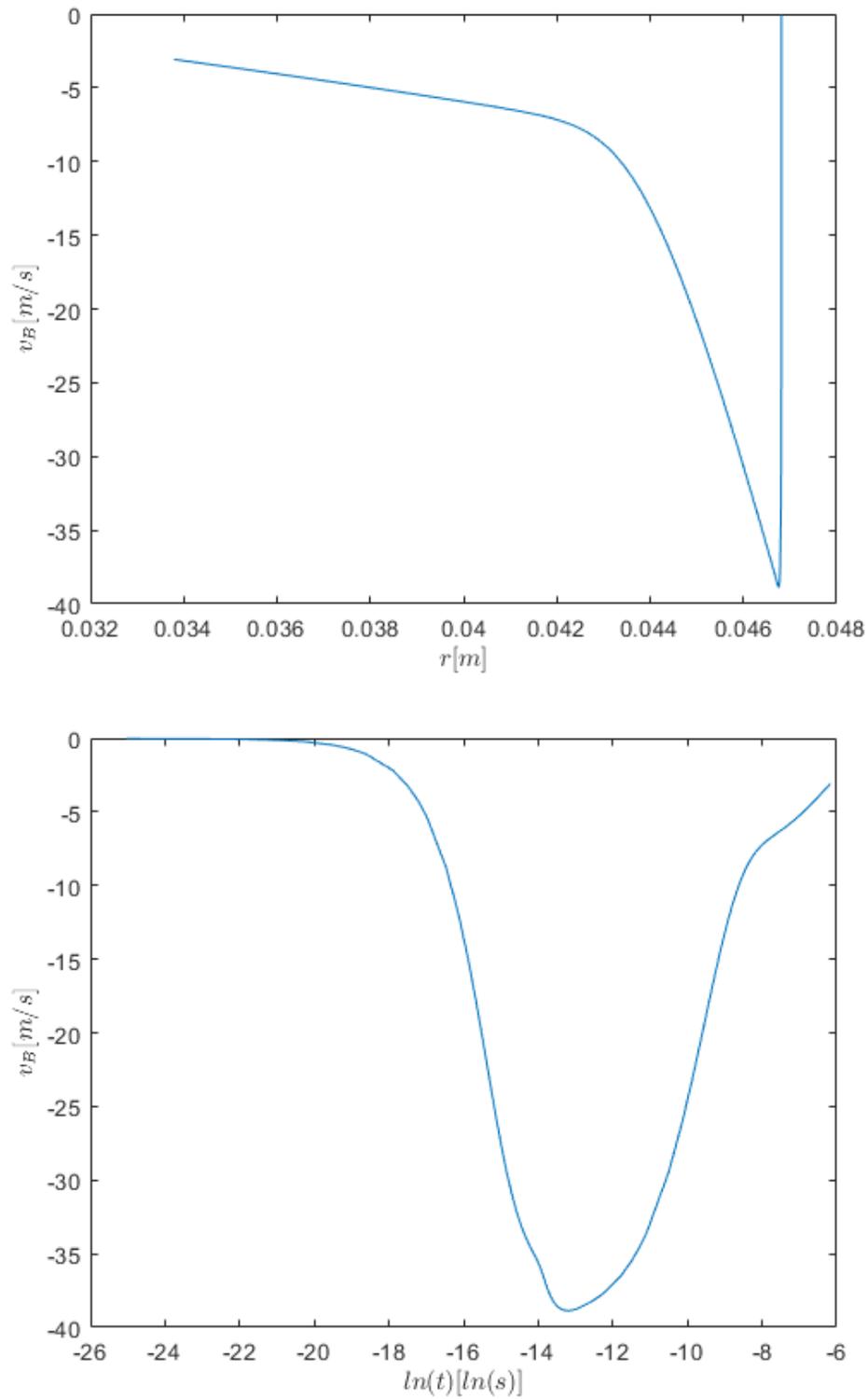


Figure 3.6: Top: Radial bubble velocity  $v_B$  vs. radial position  $r$  for  $\Omega_0 \approx 1605$  Hz. Bottom: Radial bubble velocity  $v_B$  vs.  $\ln(t)$  for  $\Omega_0 \approx 1605$  Hz. Indicates the bubble accelerates rapidly during compression and then decelerates once minimum radius is reached.

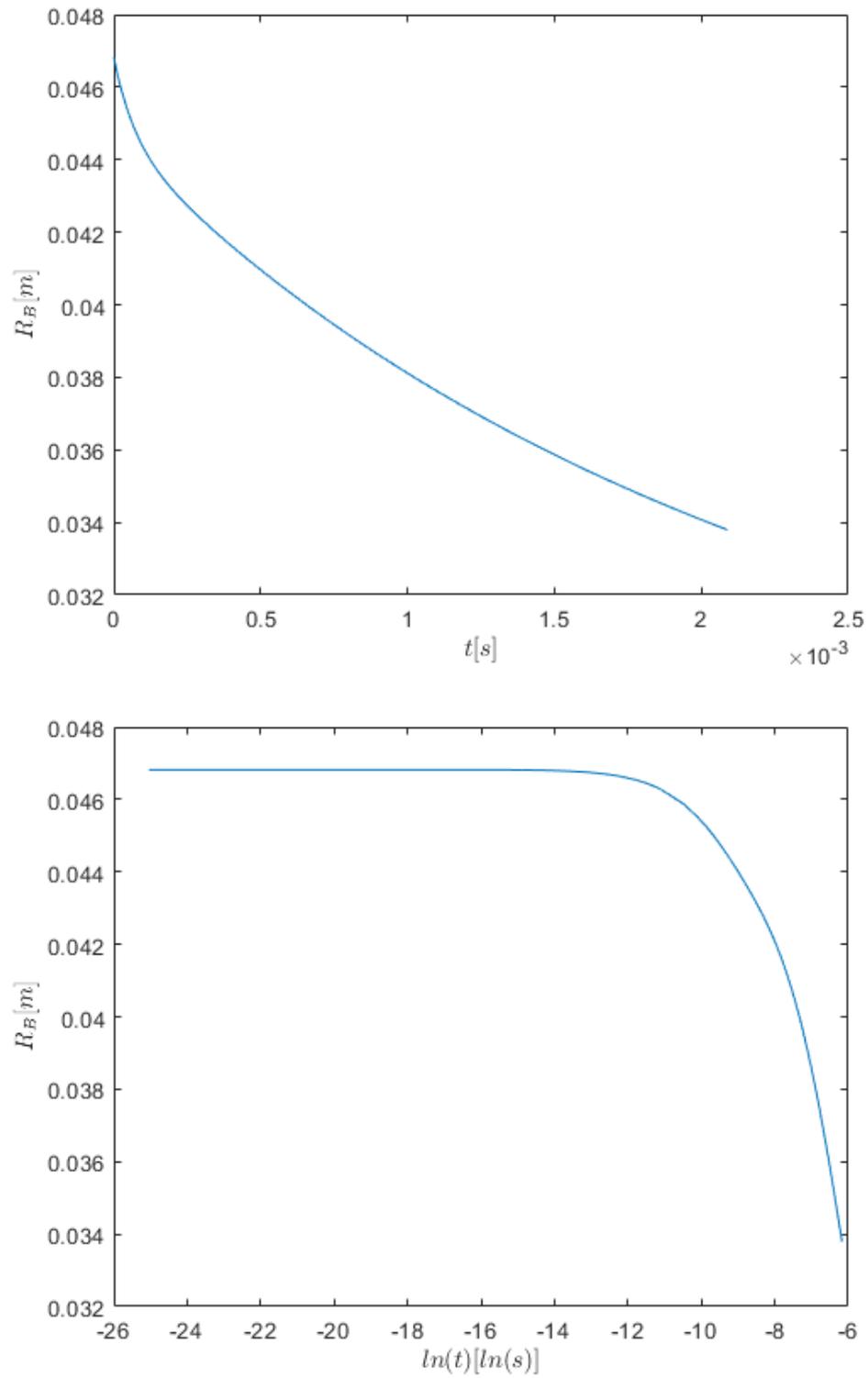


Figure 3.7: Top: Radial bubble position  $R_B$  vs.  $t$ , for  $\Omega_0 \approx 1605$  Hz.  
 Bottom: Radial bubble position  $R_B$  vs.  $\ln(t)$ , for  $\Omega_0 \approx 1605$  Hz.

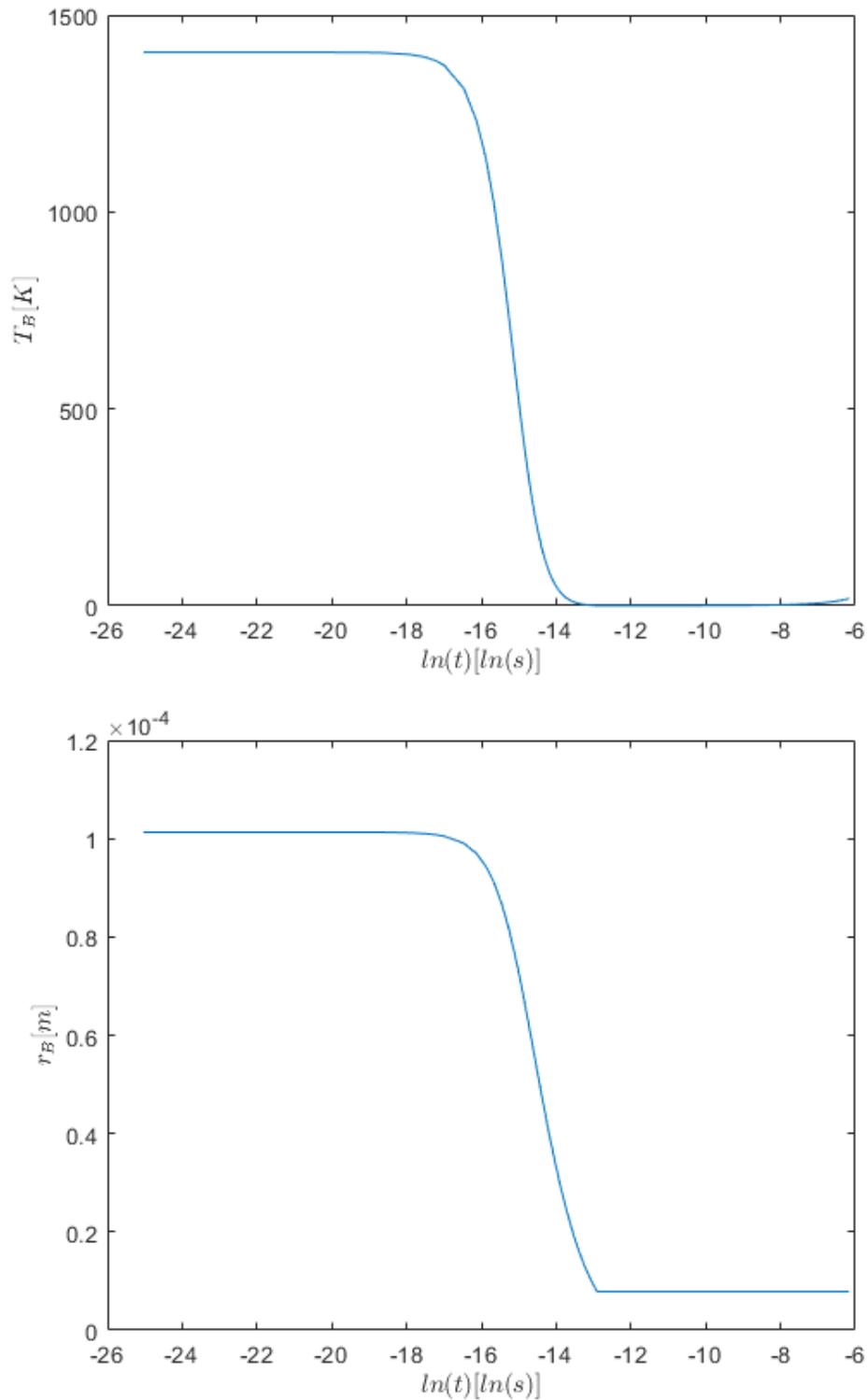


Figure 3.8: Top: Bubble temperature  $T_B$  vs.  $\ln(t)$ , for  $\Omega_0 \approx 1605$  Hz. Bubble initially compresses and super-cools to a minimum temperature  $T_{B_{min}} \approx 0.6591$  K, and then increases to 18.3421 K. Bottom: Bubble radius  $r_B$  vs.  $\ln(t)$ , for  $\Omega_0 \approx 1605$  Hz. Bubble initially compresses to a minimum radius  $r_{B_{min}}$  and then remains constant.

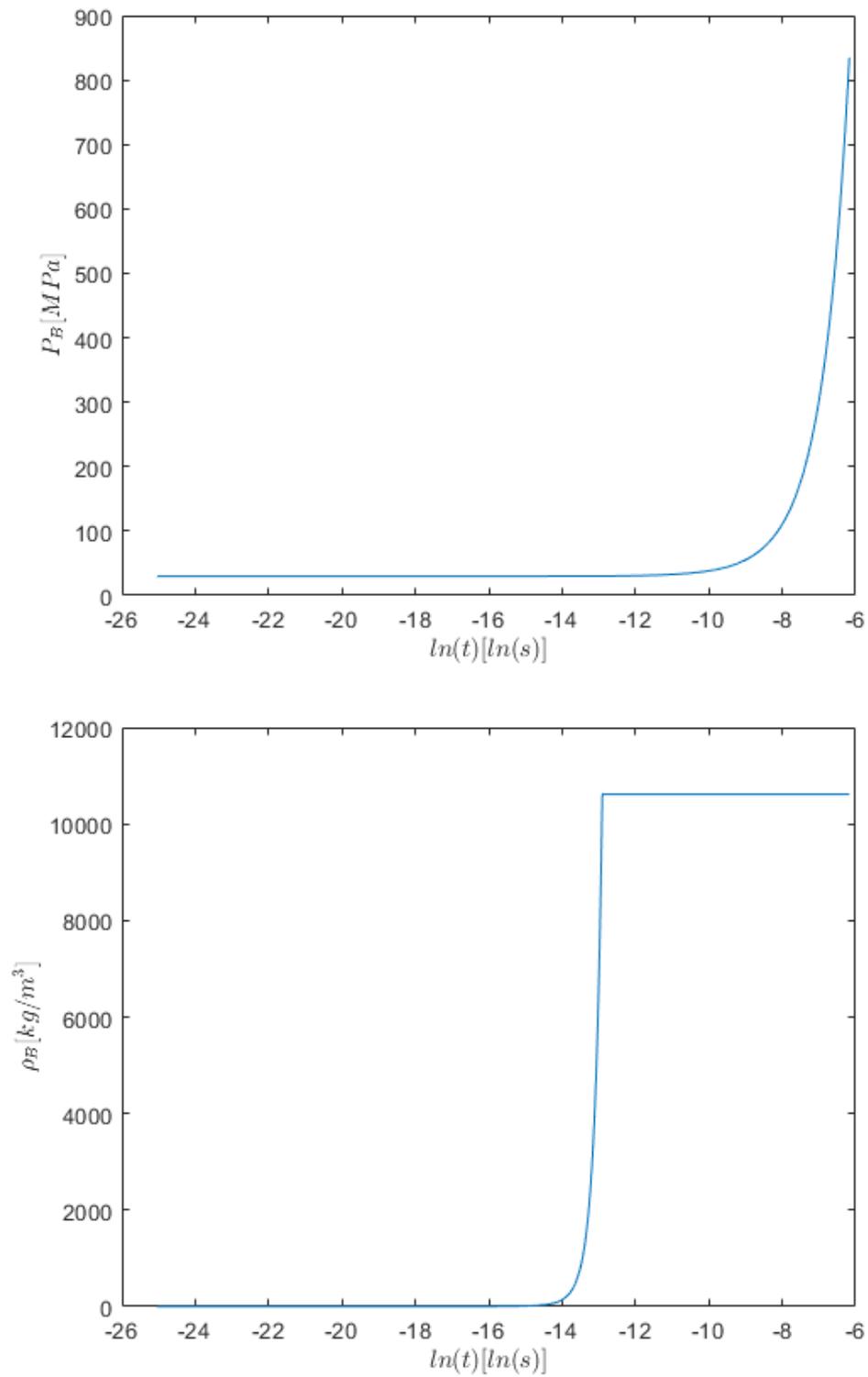


Figure 3.9: Top: Bubble pressure  $P_B$  vs.  $\ln(t)$ , for  $\Omega_0 \approx 1605$  Hz. Bubble pressure increases as it moves radially inward. Bottom: Bubble density  $\rho_B$  vs.  $\ln(t)$ , for  $\Omega_0 \approx 1605$  Hz. Bubble initially compresses to a maximum density  $\rho_{B_{max}}$  and then remains constant.

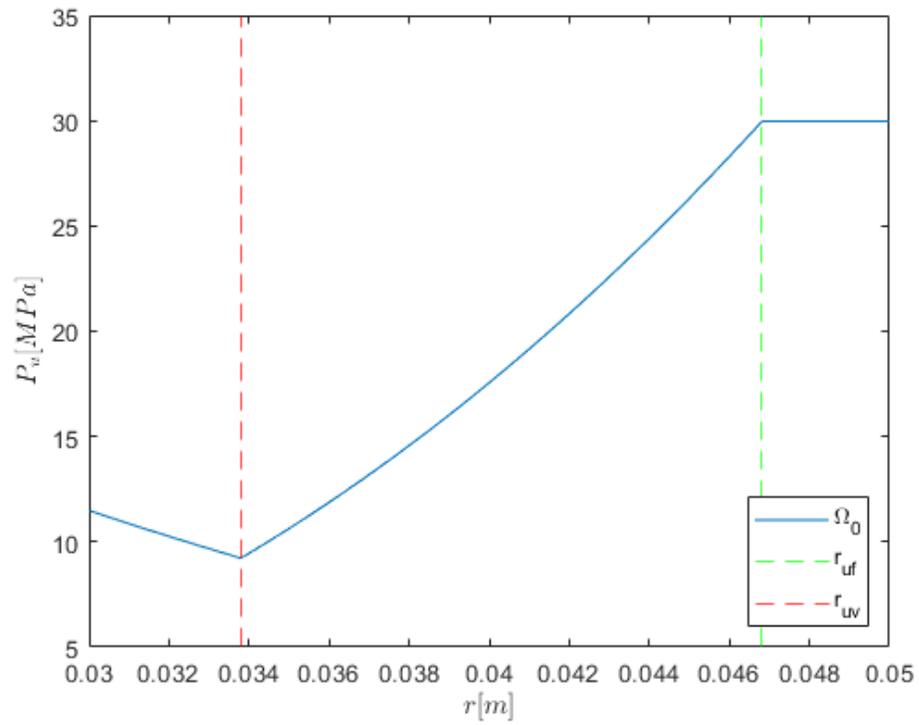
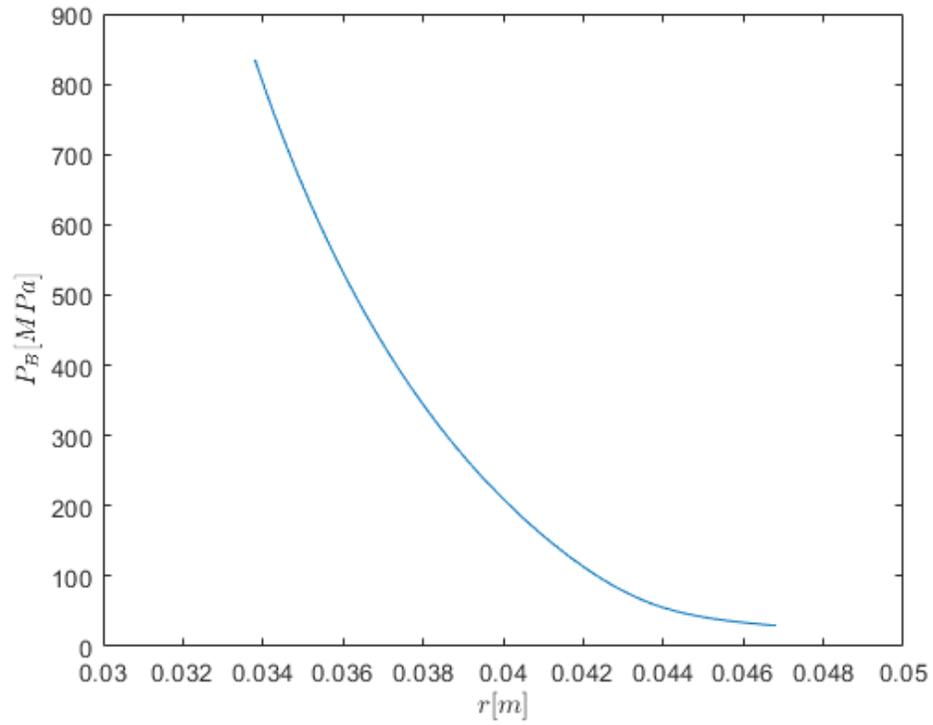


Figure 3.10: Bubble pressure  $P_B$  vs. radial position  $r$ , for  $\Omega_0 \approx 1605$  Hz.  
 Bottom: Uranium layer pressure  $P_u$  vs. radial position  $r$ , for  $\Omega_0 \approx 1605$  Hz.

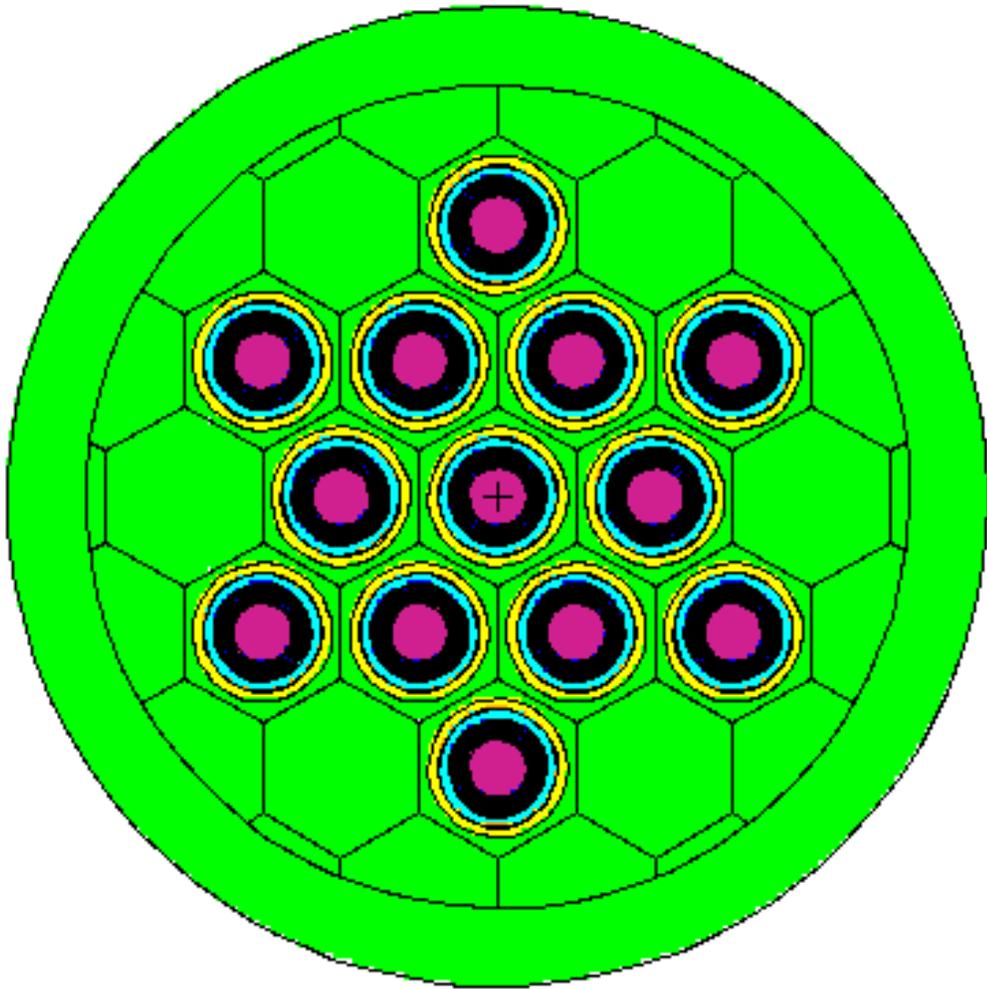


Figure 3.11: Reactor with 13 fuel elements

## CHAPTER 4: DISCUSSION AND FURTHER STUDY

Results in the previous chapter suggest that centrifugal separation of uranium and hydrogen will be beneficial. It can also be noted that in Figure 3.3 varying angular velocity will change the density at the liquid/gas uranium interface, as well as indicating that for a constant mass system,  $r_p$  will vary with varying angular velocity. However, when looking at the pressure profile in Figure 3.5 and the Young-Laplace relation (2.4), we can see that the angular velocity is limited by the initial input pressure provided by the NTRs propellant turbopump as well as current centrifugal technology. A turbopump is necessary to reduce the mass and thickness of the hydrogen pressure tank wall while achieving the required operational pressures for high performance and thrust of the rocket. Pressures above 30 MPa are currently outside of current turbopump capabilities, however, when looking at the trend of bubble pressure within the liquid uranium layer, the potential for utilizing this behavior in dense liquids to step up the pressure could be studied in the future. However, this also relies on the improvement of centrifugal technology; magnetic bearings currently are seeing rpm values of 500,000 (=8333 Hz). Minor variation in angular velocity may provide significant control over the dynamics of the uranium gas region; varying angular velocity would result in the change in plenum location  $r_p$ , as well as allow for the possibility of turning the mass flow rate off/on by increasing/decreasing  $P_u(ruf, \Omega)$  above/below the input hydrogen pressure. The results of the bubble evolution suggest that this technique may also be able to produce metastable metallic hydrogen, which is a more efficient propellant, that reaches the reaction chamber.

Further study may be required to numerically solve for equation (2.8) with varying  $E_d(r)$ . As the bulk density of uranium and hydrogen seemed negligibly different,

the uranium region was treated as purely uranium; however, further study may be required in regards to accurate mass fractions and porosity of the uranium layer. As the hydrogen bubble reaches the uranium vapor layer, two possible scenarios arise: either (1) the bubble disperses and becomes well-mixed in the uranium layer, in which a diffusion model may be necessary for future study, or (2) the bubble disperses into a column/finger spanning the thickness of the vapor layer, in which a jet model may be in order. Further optimization of design parameters could be done to minimize critical mass and maximize thermal coupling between nuclear fuel and propellant, for comparison to other reactors.

Although the design parameters for the model of a single fuel element resulted in a subcritical state ( $k_{eff} < 1$ ), an expanded lattice structure of 13 elements was tested with the designed density profiles resulting in a supercritical state ( $k_{eff} > 1$ ). A higher  $k_{eff}$  for a multi-element reactor is to be expected, as there is more fissionable material for neutrons from each element to interact with; however, both subcritical and supercritical states are not desired for a self-sustaining reactor. Therefore continued iteration and optimization with the lattice structure is necessary to find a convergence of the density profile and  $k_{eff}=1$ . Additional study of various moderator distributions may also prove beneficial to reactor performance. This could be achieved by seeding the hydrogen propellant with a moderator, by using a temporary "hot frit" that melts/vaporizes and mixes with the uranium fuel upon start-up, by using similar fuel pellets used in PBRs, or a combination thereof.

As this design is highly conceptual, many assumptions will need to be revisited in future models to address thermal properties that vary with temperature. Due to limited data on the thermal properties of high temperature (gaseous) uranium, future experimentation to extend the knowledge base of uranium thermal properties is required to advance the accuracy of this model. Additional study may also investigate the system of mostly gaseous uranium, however such a system may be unstable due

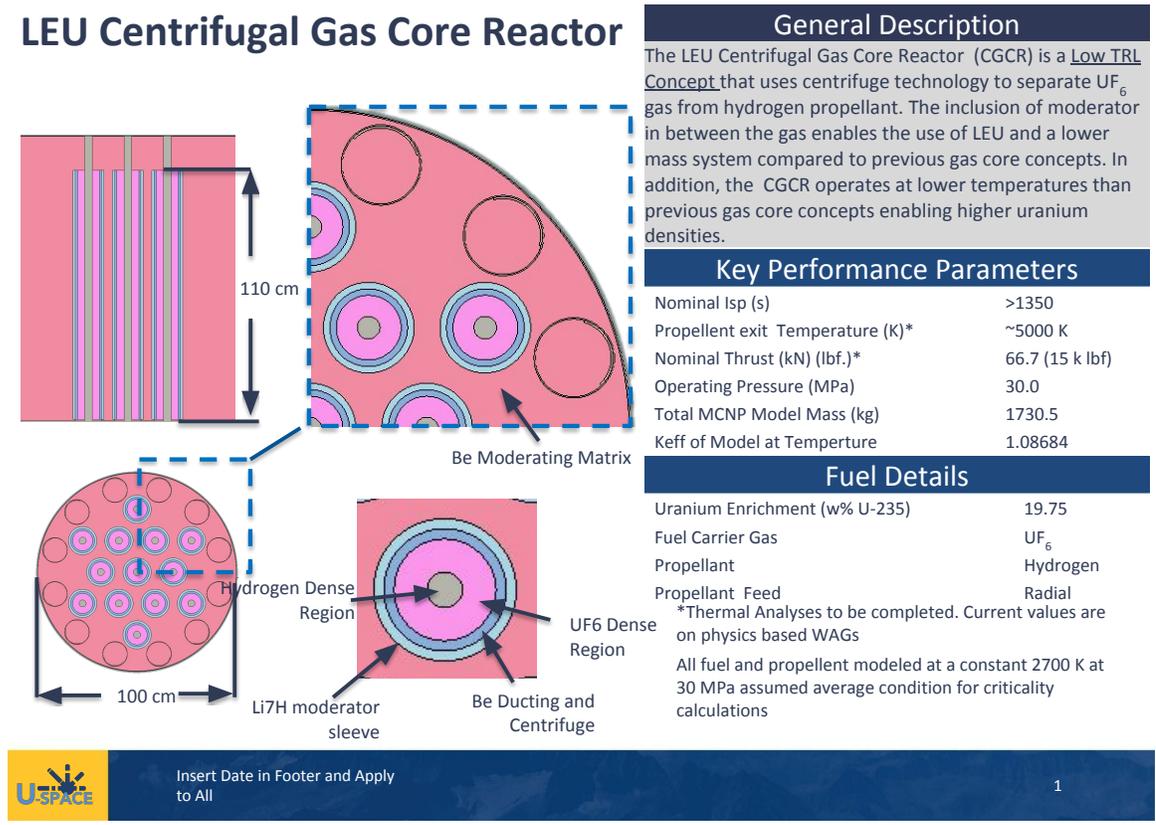
to plasma ionization and potential magnetohydrodynamic instabilities. This research may also extend to other fluid system applications outside of NTP. The successful conclusion of this research has been to demonstrate a first model of CGCR using low enriched uranium metal, centrifugal separation, and fuel/propellant heat transfer.

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APPENDIX A: ADDITIONAL FIGURES

Figure A.1: Preliminary reference design of LEU CGCR with  $UF_6$  as fuel. Analysis done by U-space, provided by NASA



## APPENDIX B: PROGRAM FILES

## B.1 MCNP INPUT

```

1 c Cell Cards
2 1 1 -0.0005061 -1 -26 imp:n,p 1 $ hot h2
3 2 2 -0.059768 1 -2 -25 -26 imp:n,p 1 $ Uranium metal LEU
4 3 2 -0.059817 2 -3 -25 -26 imp:n,p 1 $
5 4 2 -0.059873 3 -4 -25 -26 imp:n,p 1 $
6 5 2 -2.704474 4 -5 -25 -26 imp:n,p 1 $
7 6 2 -13.204924 5 -6 -25 -26 imp:n,p 1 $
8 7 2 -13.585757 6 -7 -25 -26 imp:n,p 1 $
9 8 2 -13.956008 7 -8 -25 -26 imp:n,p 1 $
10 9 2 -14.316250 8 -9 -25 -26 imp:n,p 1 $
11 10 2 -14.667010 9 -10 -25 -26 imp:n,p 1 $
12 11 2 -15.008775 10 -11 -25 -26 imp:n,p 1 $
13 12 2 -15.341993 11 -12 -25 -26 imp:n,p 1 $
14 13 2 -15.667083 12 -13 -25 -26 imp:n,p 1 $
15 14 2 -15.984431 13 -14 -25 -26 imp:n,p 1 $
16 15 2 -16.294397 14 -15 -25 -26 imp:n,p 1 $
17 16 2 -16.597318 15 -16 -25 -26 imp:n,p 1 $
18 17 2 -16.893506 16 -17 -25 -26 imp:n,p 1 $
19 18 2 -17.513585 17 -18 -25 -26 imp:n,p 1 $
20 19 2 -19.100000 18 -19 -25 -26 imp:n,p 1 $
21 20 2 -19.100000 19 -20 -25 -26 imp:n,p 1 $
22 21 2 -19.100000 20 -21 -25 -26 imp:n,p 1 $
23 22 5 -0.85423595 21 -22 -25 -26 imp:n,p 1 $ graphite/H2 ducting
24 23 6 -0.0084719 22 -23 -25 -26 imp:n,p 1 $ cold h2
25 24 3 -0.783 23 -24 -25 -26 imp:n,p 1 $ Li7H reflector
26 c reactor vessel
27 25 4 -1.85 (24 -26):(1 -24 25 -26) imp:n,p 1 $ Beryllium mod
28 26 0 26 imp:n,p 0 $ void outside
29
30 c Surface Cards
31 c solution cylinder
32 1 cz 3.0 $ hot H2/LEU cyl surf
33 2 cz 3.1
34 3 cz 3.2
35 4 cz 3.3
36 5 cz 3.4
37 6 cz 3.5
38 7 cz 3.6
39 8 cz 3.7
40 9 cz 3.8
41 10 cz 3.9
42 11 cz 4.0
43 12 cz 4.1
44 13 cz 4.2
45 14 cz 4.3
46 15 cz 4.4
47 16 cz 4.5
48 17 cz 4.6
49 18 cz 4.7
50 19 cz 4.8
51 20 cz 4.9
52 21 cz 5 $ inner porous centrifuge wall surf
53 22 cz 6 $ outer porous centrifuge wall surf
54 23 cz 6.1 $ inner reflect surf
55 24 cz 7.1 $ outer reflect surf
56 25 pz 50 $ centrifuge top
57 c reactor vessel
58 26 rcc 0 0 -50 0 0 110 50 $ outer Mod surf
59
60 c Data Cards
61 mode n p
62 kcode 5000 1 20 100 $ Calculate keff
63 ksrc 0 4.05 0
64 c Materials
65 M1 1001.84c 1.0 $ Hot H2 @ 2700K
66 M2 92235.84c -0.1975 $ LEU w%U-235 19.75
67 92238.84c -0.8025
68 M3 3007.86c -0.97988 $ Li7H
69 1001.86c -0.02012

```

```
70 M4 4009.80c 1.0 $ Beryllium
71 MT4 be.20t
72 M5 6000.82c -0.995041 $ porous(50%) graphite saturated with H2 @900K
73 1001.82c -0.004959
74 M6 1002.85c 1.0 $ liquid H2
75 M7 6000.86c 1.0 $ graphite
76 M7T grph.20t
77 c Tallies
78 e0:n 0.1 1.0 20.0 $ Default energy bins (MeV)
79 f2:n,p 26.1 26.2 26.3 T $ photon & neutron flux through surface
80 f4:n 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 $ flux in cells
81 16 17 18 19 20 21 22 23 24 25
82 f6:n,p 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16
83 17 18 19 20 21 22 23 24 25 $ energy deposition
84
85
```



54	1 16- 1	15	2	-16.294397	14 -15 -25 -26	imp:n,p
55	17- 1	16	2	-16.597318	15 -16 -25 -26	imp:n,p
56	18- 1	17	2	-16.893506	16 -17 -25 -26	imp:n,p
57	19- 1	18	2	-17.513585	17 -18 -25 -26	imp:n,p
58	20- 1	19	2	-19.100000	18 -19 -25 -26	imp:n,p
59	21- 1	20	2	-19.100000	19 -20 -25 -26	imp:n,p
60	22- 1	21	2	-19.100000	20 -21 -25 -26	imp:n,p
61	23- 1	22	5	-0.85423595	21 -22 -25 -26	imp:n,p
62	24- 1	23	6	-0.0084719	22 -23 -25 -26	imp:n,p
63	25- 1	24	3	-0.783	23 -24 -25 -26	imp:n,p
64	26- vessel	c reactor				
65	27- 1	25	4	-1.85	(24 -26):(1 -24 25 -26)	imp:n,p
66	28- 0	26	0		26	imp:n,p
67	29-					
68	30- Cards	c Surface				
69	31- cylinder	c solution				
70	32- surf	1	cz	3.0	\$ hot H2/LEU cyl	
71	33- 3.1	2	cz			
72	34- 3.2	3	cz			
73	35- 3.3	4	cz			
74	36- 3.4	5	cz			
75	37- 3.5	6	cz			
76	38- 3.6	7	cz			
77	39- 3.7	8	cz			
78	40- 3.8	9	cz			
79	41- 3.9	10	cz			
80	42- 4.0	11	cz			
81	43- 4.1	12	cz			
82	44- 4.2	13	cz			
83	45- 4.3	14	cz			
84	46- 4.4	15	cz			
85	47- 4.5	16	cz			
86	48- 4.6	17	cz			
87	49-	18	cz			

```

      4.7
88    50-    19    cz
      4.8
89    51-    20    cz
      4.9
90    52-    21    cz    5          $ inner porous centrifuge wall
      surf
91    53-    22    cz    6          $ outer porous centrifuge wall
      surf
92    54-    23    cz    6.1        $ inner reflect
      surf
93    55-    24    cz    7.1        $ outer reflect
      surf
94    56-    25    pz    50         $ centrifuge
      top
95    57-    c reactor
      vessel
96    58-    26    rcc    0 0 -50 0 0 110 50          $ outer Mod
      surf
97
      59-

98    60-    c Data
      Cards
99    61-    mode n
      p
100   comment. photonuclear physics may be needed (phys:p).
101    62-    kcode 5000 1 20 100 $ Calculate
      keff
102    63-    ksrc 0 4.05
      0
103    64-    c
      Materials
104    65-    M1    1001.84c    1.0          $ Hot H2 @
      2700K
105    66-    M2    92235.84c    -0.1975    $ LEU w%U-235
      19.75
106    67-    92238.84c
      -0.8025
107    68-    M3    3007.86c    -0.97988    $
      Li7H
108    69-    1001.86c
      -0.02012
109    70-    M4    4009.80c    1.0          $
      Beryllium
110    71-    MT4
      be.20t
111    72-    M5    6000.82c    -0.995041    $ porous(50%) graphite
      saturated with H2
112    73-    1001.82c
      -0.004959
113    74-    M6    1002.85c    1.0          $ liquid
      H2
114    75-    M7    6000.86c    1.0          $
      graphite
115   warning. material    7 is not used in the problem.
116    76-    M7T
      grph.20t
117   warning. material    7 is not used in the problem.
118    77-    c
      Tallies
119    78-    e0:n 0.1 1.0 20.0          $ Default energy bins
      (MeV)
120    79-    f2:n,p 26.1 26.2 26.3 T    $ photon & neutron flux through
      surface
121    80-    f4:n 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15          $ flux in
      cells
122    81-    16 17 18 19 20 21 22 23 24
      25

```

```

123      82-      f6:n,p  1 2 3 4 5 6 7 8 9 10 11 12 13 14 15
124      16
124      83-      17 18 19 20 21 22 23 24 25      $ energy
125      deposition
126
126      84-
127
127      comment. total fission nubar data are being used.
128
129      warning. 1001.82c and 1001.84c are both called for.
130
131      warning. 1001.84c and 1001.86c are both called for.
132
133      warning. 1 of the materials appear at more than one density.
134      1cells
135      print table 60
136
136      atom
136      gram
137      neutron photon photon wt
137      cell mat density density volume mass
137      pieces importance importance generation
138
139      1 1 1 3.02409E-04 5.06100E-04 3.11018E+03 1.57406E+00
139      1 1.0000E+00 1.0000E+00 -1.000E+00
140      2 2 1.51579E-04 5.97680E-02 1.91637E+02 1.14538E+01
140      1 1.0000E+00 1.0000E+00 -1.000E+00
141      3 3 2 1.51703E-04 5.98170E-02 1.97920E+02 1.18390E+01
141      1 1.0000E+00 1.0000E+00 -1.000E+00
142      4 4 2 1.51845E-04 5.98730E-02 2.04204E+02 1.22263E+01
142      1 1.0000E+00 1.0000E+00 -1.000E+00
143      5 5 2 6.85888E-03 2.70447E+00 2.10487E+02 5.69256E+02
143      1 1.0000E+00 1.0000E+00 -1.000E+00
144      6 6 2 3.34893E-02 1.32049E+01 2.16770E+02 2.86243E+03
144      1 1.0000E+00 1.0000E+00 -1.000E+00
145      7 7 2 3.44551E-02 1.35858E+01 2.23053E+02 3.03034E+03
145      1 1.0000E+00 1.0000E+00 -1.000E+00
146      8 8 2 3.53941E-02 1.39560E+01 2.29336E+02 3.20062E+03
146      1 1.0000E+00 1.0000E+00 -1.000E+00
147      9 9 2 3.63078E-02 1.43163E+01 2.35619E+02 3.37319E+03
147      1 1.0000E+00 1.0000E+00 -1.000E+00
148      10 10 2 3.71973E-02 1.46670E+01 2.41903E+02 3.54799E+03
148      1 1.0000E+00 1.0000E+00 -1.000E+00
149      11 11 2 3.80641E-02 1.50088E+01 2.48186E+02 3.72497E+03
149      1 1.0000E+00 1.0000E+00 -1.000E+00
150      12 12 2 3.89092E-02 1.53420E+01 2.54469E+02 3.90406E+03
150      1 1.0000E+00 1.0000E+00 -1.000E+00
151      13 13 2 3.97336E-02 1.56671E+01 2.60752E+02 4.08523E+03
151      1 1.0000E+00 1.0000E+00 -1.000E+00
152      14 14 2 4.05385E-02 1.59844E+01 2.67035E+02 4.26841E+03
152      1 1.0000E+00 1.0000E+00 -1.000E+00
153      15 15 2 4.13246E-02 1.62944E+01 2.73319E+02 4.45356E+03
153      1 1.0000E+00 1.0000E+00 -1.000E+00
154      16 16 2 4.20928E-02 1.65973E+01 2.79602E+02 4.64064E+03
154      1 1.0000E+00 1.0000E+00 -1.000E+00
155      17 17 2 4.28440E-02 1.68935E+01 2.85885E+02 4.82960E+03
155      1 1.0000E+00 1.0000E+00 -1.000E+00
156      18 18 2 4.44166E-02 1.75136E+01 2.92168E+02 5.11691E+03
156      1 1.0000E+00 1.0000E+00 -1.000E+00
157      19 19 2 4.84399E-02 1.91000E+01 2.98451E+02 5.70042E+03
157      1 1.0000E+00 1.0000E+00 -1.000E+00
158      20 20 2 4.84399E-02 1.91000E+01 3.04734E+02 5.82043E+03
158      1 1.0000E+00 1.0000E+00 -1.000E+00
159      21 21 2 4.84399E-02 1.91000E+01 3.11018E+02 5.94044E+03
159      1 1.0000E+00 1.0000E+00 -1.000E+00
160      22 22 5 4.51482E-02 8.54236E-01 3.45575E+03 2.95203E+03
160      1 1.0000E+00 1.0000E+00 -1.000E+00
161      23 23 6 2.53305E-03 8.47190E-03 3.80133E+02 3.22045E+00

```

```

1227 comment.
1228 comment. Source entropy convergence check passed.
1229 comment.
1230 the results of the w test for normality applied to the individual collision,
absorption, and track-length keff cycle values are:
1231
1232         the k( collision) cycle values appear normally distributed at the 95 percent
confidence level
1233         the k(absorption) cycle values appear normally distributed at the 95 percent
confidence level
1234         the k(trk length) cycle values appear normally distributed at the 95 percent
confidence level
1235
1236
1237
-----
1238
|
|
1239 | the final estimated combined collision/absorption/track-length keff = 0.88394 with
an estimated standard deviation of 0.00121 |
1240
|
|
1241 | the estimated 68, 95, & 99 percent keff confidence intervals are 0.88273 to 0.88516,
0.88153 to 0.88636, and 0.88074 to 0.88714 |
1242
|
|
1243 | the final combined (col/abs/tl) prompt removal lifetime = 7.6118E-04 seconds with an
estimated standard deviation of 1.3717E-05 |
1244
|
|
1245 | the average neutron energy causing fission = 4.0417E-01
mev |
1246 | the energy corresponding to the average neutron lethargy causing fission =
1.9465E-05 mev |
1247
|
|
1248 | the percentages of fissions caused by neutrons in the thermal, intermediate, and
fast neutron ranges are: |
1249 |      (<0.625 ev):  50.18%      (0.625 ev - 100 kev):  28.88%      (>100
kev):  20.94% |
1250
|
|
1251 | the average fission neutrons produced per neutron absorbed (capture + fission) in
all cells with fission = 1.7519E+00 |
1252 | the average fission neutrons produced per neutron absorbed (capture + fission) in
all the geometry cells = 1.1759E+00 |
1253
|
|
1254 | the average number of neutrons produced per fission =
2.481 |
1255
|
|
1256
-----
1257
1258
1259 the estimated average keffs, one standard deviations, and 68, 95, and 99 percent
confidence intervals are:
1260

```

```

1553
1554
1555 the first active half of the problem skips 20 cycles and uses 40 active cycles; the
1556 second half skips 60 and uses 40 cycles.
1557 the col/abs/trk-len keff, one standard deviation, and 68, 95, and 99 percent intervals
1558 for each active half of the problem are:
1559
1560          problem      keff      standard deviation      68% confidence
1561          95% confidence      99% confidence
1562
1563          first half      0.88310      0.00172      0.88136 to 0.88484
1564          0.87961 to 0.88659      0.87842 to 0.88778
1565          second half      0.88484      0.00177      0.88306 to 0.88663
1566          0.88126 to 0.88842      0.88005 to 0.88964
1567          final result      0.88394      0.00121      0.88273 to 0.88516
1568          0.88153 to 0.88636      0.88074 to 0.88714
1569
1570 the first and second half values of k(collisions/absorption/track length) appear to be
1571 the same at the 68 percent confidence level.
1572
1573 ltally      4      nps =      499664
1574 tally type 4      track length estimate of particle flux.      units
1575 1/cm**2
1576 particle(s): neutrons
1577 number of histories used for normalizing tallies =      400000.00
1578
1579 volumes
1580 cell:      1      2      3      4
1581 5      6      7
1582          3.11018E+03      1.91637E+02      1.97920E+02      2.04204E+02
1583          2.10487E+02      2.16770E+02      2.23053E+02
1584 cell:      8      9      10      11
1585 12      13      14
1586          2.29336E+02      2.35619E+02      2.41903E+02      2.48186E+02
1587          2.54469E+02      2.60752E+02      2.67035E+02
1588 cell:      15      16      17      18
1589 19      20      21
1590          2.73319E+02      2.79602E+02      2.85885E+02      2.92168E+02
1591          2.98451E+02      3.04734E+02      3.11018E+02
1592 cell:      22      23      24
1593 25
1594          3.45575E+03      3.80133E+02      4.14690E+03      8.47818E+05
1595
1596 cell
1597 1
1598
1599 energy
1600 1.0000E-01      3.91664E-04      0.0042
1601 1.0000E+00      5.52768E-04      0.0036
1602 2.0000E+01      3.66582E-04      0.0041
1603 total      1.31101E-03      0.0023
1604
1605 cell
1606 2
1607
1608 energy
1609 1.0000E-01      3.98594E-04      0.0048
1610 1.0000E+00      6.14332E-04      0.0041
1611 2.0000E+01      4.13450E-04      0.0047
1612 total      1.42638E-03      0.0026
1613
1614 cell
1615 3
1616
1617 energy
1618 1.0000E-01      3.95625E-04      0.0048
1619 1.0000E+00      6.15733E-04      0.0040
1620 2.0000E+01      4.17378E-04      0.0047
1621 total      1.42874E-03      0.0025

```



1969	energy			
1970	1.0000E-01	2.55082E-04	0.0113	
1971	1.0000E+00	7.82706E-05	0.0038	
1972	2.0000E+01	1.17579E-04	0.0044	
1973	total	4.50931E-04	0.0066	
1974				
1975	cell			
	5			
1976	energy			
1977	1.0000E-01	2.44964E-04	0.0095	
1978	1.0000E+00	7.87479E-05	0.0037	
1979	2.0000E+01	1.17989E-04	0.0043	
1980	total	4.41700E-04	0.0055	
1981				
1982	cell			
	6			
1983	energy			
1984	1.0000E-01	2.33354E-04	0.0077	
1985	1.0000E+00	7.91752E-05	0.0034	
1986	2.0000E+01	1.18645E-04	0.0040	
1987	total	4.31174E-04	0.0044	
1988				
1989	cell			
	7			
1990	energy			
1991	1.0000E-01	2.25796E-04	0.0069	
1992	1.0000E+00	7.97257E-05	0.0033	
1993	2.0000E+01	1.20013E-04	0.0039	
1994	total	4.25534E-04	0.0039	
1995				
1996	cell			
	8			
1997	energy			
1998	1.0000E-01	2.23873E-04	0.0064	
1999	1.0000E+00	7.99264E-05	0.0032	
2000	2.0000E+01	1.21318E-04	0.0037	
2001	total	4.25117E-04	0.0036	
2002				
2003	cell			
	9			
2004	energy			
2005	1.0000E-01	2.26951E-04	0.0062	
2006	1.0000E+00	8.02480E-05	0.0031	
2007	2.0000E+01	1.22145E-04	0.0036	
2008	total	4.29344E-04	0.0036	
2009				
2010	cell			
	10			
2011	energy			
2012	1.0000E-01	2.31420E-04	0.0061	
2013	1.0000E+00	8.06878E-05	0.0030	
2014	2.0000E+01	1.22493E-04	0.0035	
2015	total	4.34601E-04	0.0035	
2016				
2017	cell			
	11			
2018	energy			
2019	1.0000E-01	2.40245E-04	0.0060	
2020	1.0000E+00	8.09582E-05	0.0030	
2021	2.0000E+01	1.24229E-04	0.0034	
2022	total	4.45433E-04	0.0034	
2023				

2024	cell			
	12			
2025	energy			
2026	1.0000E-01	2.53460E-04	0.0058	
2027	1.0000E+00	8.14408E-05	0.0029	
2028	2.0000E+01	1.25999E-04	0.0034	
2029	total	4.60900E-04	0.0034	
2030				
2031	cell			
	13			
2032	energy			
2033	1.0000E-01	2.70570E-04	0.0057	
2034	1.0000E+00	8.18886E-05	0.0028	
2035	2.0000E+01	1.27269E-04	0.0033	
2036	total	4.79727E-04	0.0034	
2037				
2038	cell			
	14			
2039	energy			
2040	1.0000E-01	2.95127E-04	0.0057	
2041	1.0000E+00	8.22675E-05	0.0028	
2042	2.0000E+01	1.29374E-04	0.0033	
2043	total	5.06768E-04	0.0035	
2044				
2045	cell			
	15			
2046	energy			
2047	1.0000E-01	3.30298E-04	0.0056	
2048	1.0000E+00	8.28928E-05	0.0027	
2049	2.0000E+01	1.31225E-04	0.0032	
2050	total	5.44416E-04	0.0035	
2051				
2052	cell			
	16			
2053	energy			
2054	1.0000E-01	3.80192E-04	0.0055	
2055	1.0000E+00	8.36352E-05	0.0026	
2056	2.0000E+01	1.33559E-04	0.0031	
2057	total	5.97387E-04	0.0036	
2058				
2059	cell			
	17			
2060	energy			
2061	1.0000E-01	4.59363E-04	0.0053	
2062	1.0000E+00	8.45924E-05	0.0026	
2063	2.0000E+01	1.36398E-04	0.0031	
2064	total	6.80353E-04	0.0037	
2065				
2066	cell			
	18			
2067	energy			
2068	1.0000E-01	5.94594E-04	0.0050	
2069	1.0000E+00	8.57502E-05	0.0025	
2070	2.0000E+01	1.40085E-04	0.0030	
2071	total	8.20429E-04	0.0037	
2072				
2073	cell			
	19			
2074	energy			
2075	1.0000E-01	8.50686E-04	0.0045	
2076	1.0000E+00	8.75293E-05	0.0025	

2077 2.0000E+01 1.44803E-04 0.0030  
 2078 total 1.08302E-03 0.0036

2079  
 2080 cell  
 20

2081 energy  
 2082 1.0000E-01 1.40953E-03 0.0038  
 2083 1.0000E+00 8.98949E-05 0.0024  
 2084 2.0000E+01 1.49875E-04 0.0029  
 2085 total 1.64930E-03 0.0033

2086  
 2087 cell  
 21

2088 energy  
 2089 1.0000E-01 3.08790E-03 0.0030  
 2090 1.0000E+00 9.25687E-05 0.0023  
 2091 2.0000E+01 1.50971E-04 0.0027  
 2092 total 3.33144E-03 0.0028

2093  
 2094 cell  
 22

2095 energy  
 2096 1.0000E-01 7.49227E-07 0.0037  
 2097 1.0000E+00 1.14453E-05 0.0024  
 2098 2.0000E+01 2.36452E-05 0.0026  
 2099 total 3.58397E-05 0.0019

2100  
 2101 cell  
 23

2102 energy  
 2103 1.0000E-01 4.74226E-06 0.0046  
 2104 1.0000E+00 9.68406E-05 0.0029  
 2105 2.0000E+01 2.14180E-04 0.0028  
 2106 total 3.15763E-04 0.0020

2107  
 2108 cell  
 24

2109 energy  
 2110 1.0000E-01 1.24860E-06 0.0032  
 2111 1.0000E+00 1.56405E-05 0.0021  
 2112 2.0000E+01 3.35636E-05 0.0023  
 2113 total 5.04527E-05 0.0016

2114  
 2115 cell  
 25

2116 energy  
 2117 1.0000E-01 8.38639E-08 0.0015  
 2118 1.0000E+00 5.70470E-07 0.0017  
 2119 2.0000E+01 1.23694E-06 0.0026  
 2120 total 1.89127E-06 0.0019

2121  
 2122

2123 \*\*\*\*\* the nps-dependent tfc bin check results are suspect because there are only 1  
 nps tally values to analyze \*\*\*\*\*

2124  
 2125  
 2126

=====

2127  
 2128

results of 10 statistical checks for the estimated answer for the tally  
 fluctuation chart (tfc) bin of tally 6

2129

```

2130  tfc bin    --mean--    -----relative error-----    ----variance of the
      variance----    --figure of merit--    -pdf-
2131  behavior  behavior  value decrease decrease rate  value  decrease
      decrease rate      value      behavior      slope
2132
2133  desired    random    <0.10    yes    1/sqrt(nps)    <0.10    yes
      1/nps      constant  random    >3.00
2134  observed    random    0.00    yes    yes    0.00    yes
      yes      constant  random    9.76
2135  passed?    yes      yes    yes    yes    yes    yes
      yes      yes      yes    yes
2136
2137
=====
2138
2139
2140  this tally meets the statistical criteria used to form confidence intervals: check the
      tally fluctuation chart to verify.
2141  the results in other bins associated with this tally may not meet these statistical
      criteria.
2142
2143  ----- estimated confidence intervals: -----
2144
2145  estimated asymmetric confidence interval(1,2,3 sigma): 1.1756E-03 to 1.1819E-03;
      1.1724E-03 to 1.1851E-03; 1.1692E-03 to 1.1882E-03
2146  estimated symmetric confidence interval(1,2,3 sigma): 1.1756E-03 to 1.1819E-03;
      1.1724E-03 to 1.1850E-03; 1.1692E-03 to 1.1882E-03
2147
2148  lanalysis of the results in the tally fluctuation chart bin (tfc) for tally      6
      with nps =      499664  print table 160
2149
2150
2151  normed average tally per history = 1.17872E-03      unnormed average tally per
      history = 1.85537E-03
2152  estimated tally relative error = 0.0027      estimated variance of the
      variance = 0.0001
2153  relative error from zero tallies = 0.0015      relative error from nonzero
      scores = 0.0022
2154
2155  number of nonzero history tallies =      211767      efficiency for the nonzero
      tallies = 0.5294
2156  history number of largest tally =      108618      largest unnormalized history
      tally = 8.74292E-02
2157  (largest tally)/(average tally) = 4.71222E+01      (largest tally)/(avg nonzero
      tally)= 2.49473E+01
2158
2159  (confidence interval shift)/mean = 0.0000      shifted confidence interval
      center = 1.17873E-03
2160
2161
2162  if the largest history score sampled so far were to occur on the next history, the
      tfc bin quantities would change as follows:
2163  nps =      400513 for this table because 20 keff cycles and      99151 histories
      were skipped before tally accumulation.
2164
2165      estimated quantities      value at nps      value at nps+1
      value(nps+1)/value(nps)-1.
2166
2167      mean      1.17872E-03
      1.17885E-03      0.000115
2168      relative error      2.68439E-03
      2.68714E-03      0.001026
2169      variance of the variance      5.83958E-05
      6.15324E-05      0.053712
2170      shifted center      1.17873E-03
      1.17873E-03      0.000000
2171      figure of merit      5.24856E+04
      5.23781E+04      -0.002049

```

```

2172
2173 the estimated inverse power slope of the 198 largest tallies starting at 3.00801E-02
      is 9.7646
2174 the large score tail of the empirical history score probability density function
      appears to have no unsampled regions.
2175
2176 fom = (histories/minute)*(f(x) signal-to-noise ratio)**2 = (1.513E+05)*( 5.890E-01)**2
      = (1.513E+05)*(3.469E-01) = 5.249E+04
2177
2178 lstatus of the statistical checks used to form confidence intervals for the mean for
      each tally bin
2179
2180
2181 tally result of statistical checks for the tfc bin (the first check not passed is
      listed) and error magnitude check for all bins
2182
2183 4 passed the 10 statistical checks for the tally fluctuation chart bin
      result
2184 passed all bin error check: 100 tally bins all have relative errors less
      than 0.10 with no zero bins
2185
2186 2 missed 1 of 10 tfc bin checks: the slope of decrease of largest tallies is
      less than the minimum acceptable value of 3.0
2187 passed all bin error check: 16 tally bins all have relative errors less
      than 0.10 with no zero bins
2188
2189 6 passed the 10 statistical checks for the tally fluctuation chart bin
      result
2190 passed all bin error check: 100 tally bins all have relative errors less
      than 0.10 with no zero bins
2191
2192
2193 the 10 statistical checks are only for the tally fluctuation chart bin and do not
      apply to other tally bins.
2194
2195 warning. 1 of the 3 tally fluctuation chart bins did not pass all 10
      statistical checks.
2196 ltally fluctuation charts
2197
2198
2199
2200
2201
2202
2203
2204 dump no. 2 on file INP_U_OD_10_ID_6_dens_prof_pure_vls_4r nps = 499664
      coll = 135022750 ctm =
2205 3.09 nrn = 1253109253
2206
2207 11 warning messages so far.
2208
2209
2210 run terminated when 100 kcode cycles were done.
2211
2212 computer time = 3.11 minutes
2213
2214 mcnp version 6 05/08/13 04/27/20
      10:03:46 probid = 04/27/20 10:00:27
2215

```