

LA-UR -92-46

Los Alamos National Laboratory is operated by the University of California for the United States Department of Energy under contract W-7405 -ENG-36.

TITLE: TARGET/BLANKET CONCEPTUAL DESIGN FOR THE LOS ALAMOS ATW CONCEPT

AUTHOR(S): John R. Ireland and Michael W. Cappiello, N-12

SUBMITTED TO: Specialists' Meeting on Accelerator-Based Transmutation
Paul Scherrer Institute (PSI)
Wuerenlingen/Villigen, Switzerland
March 24-26, 1992

By acceptance of this article, the publisher recognizes that the U.S. Government retains a nonexclusive, royalty-free license to publish or reproduce the published form of this contribution, or to allow others to do so, for U.S. Government purposes.

The Los Alamos National Laboratory requests that the publisher identify this article as work performed under the auspices of the U.S. Department of Energy

Los Alamos Los Alamos National Laboratory
Los Alamos, New Mexico 87545

TARGET/BLANKET CONCEPTUAL DESIGN FOR THE
LOS ALAMOS ATW CONCEPT

January 1992

AUTHORS AND ACKNOWLEDGMENTS

AUTHORS:

M. Cappiello
J. Ireland
J. Sapir
G. Farnum

CONTRIBUTORS:

G. Russel
W. Sommer
W. Rider
B. Krohn
W. Wilson
R. Perry
R. LaBauve
P. Lisowski
E. Arthur
L. Linford
C. Cummings
C. Cappiello
R. Manzanares

EDITORIAL:

K. Ames

WORD PROCESSING:

G. Mirabal

TARGET/BLANKET CONCEPTUAL DESIGN FOR THE LOS ALAMOS ATW CONCEPT

I. INTRODUCTION

The Los Alamos Accelerator Transmutation of Waste (**ATW**) concept has many potential applications that include defense waste transmutation, defense **material** production (i.e., **tritium** and **^{238}Pu**), and the **transmutation** of hazardous nuclear wastes **from** commercial **nuclear reactors** (fission products and **actinides**). A more advanced long-term **Los Alamos** effort is investigating the potential of an accelerator-driven system to produce fission energy with a minimal nuclear waste stream. All applications employ a high-energy (800- to **1600-MeV**), **high-current** (25-250 **mA**) proton linear accelerator as the driver. In this report, we discuss only the target/blanket conceptual design for the **commercial nuclear** waste application.

The basis of all **ATW** applications is the use of high-energy protons **from** a **linear** accelerator to produce neutrons through **spallation** instead of nuclear chain reactions. The protons are directed onto a high-atomic-number target material to produce neutrons, which are then moderated in a **surrounding** heavy-water (**D_2O**) blanket. The neutrons are slowed to thermal energy in the moderator and are absorbed in the fission product waste, or they are used in the fission and transmutation of **actinides**. Because of the intense flux of generated neutrons, fission products that have small neutron absorption cross sections, such as **^{99}Tc** , can be transmuted to stable isotopes in significant quantities. Fissile actinides such as **^{239}Pu** can be fissioned directly.

Because of the high flux levels, very dilute mixtures in the blanket region can be used, which therefore reduce the potential for criticality accidents and power excursions. The transmutation isotopes are continuously transported in double-walled piping systems through the blanket region, through heat exchangers, and into separation process loops. Heavy-water aqueous solutions and slurries **are** used as the transport media.

II. DESIGN SUMMARY

A schematic of the base-case ATW design is shown in Fig. 1. The linear accelerator operates at **1600 MeV** at a continuous-wave current of **250 mA**. The primary proton beam is then split into four beams, each having a **current** of **62.5 mA** and an energy of **1600 MeV**. Each of the four beams is directed into four separate **target/blanket modules**. **A modular design approach** allows **maintenance or replacement of components with relative ease** and adds to the overall

reliability of the system. The modules have separate cooling loops for their respective targets, moderators, and blankets, and operate independently from each other.

The high-energy proton beam strikes a centrally located **spallation** target to produce an intense source of neutrons. The base-case target design is comprised of heavy-water-cooled tungsten rods. The power density is very high, and therefore requires sub-cooled boiling as the heat-transfer mechanism. Because of the intense proton and neutron **environment**, the target assembly will **require** routine replacement

Each target/blanket module is designed to burn the **actinides** and fission products from approximately two light-water reactors and produce approximately **1542-MW_t** power. Figure 2 shows a schematic of the blanket and balance-of-plant components for electric energy production. The blanket region and balance-of-plant design is based on existing heavy-water reactor technology employed in the **CANDU** reactor system. Here, the fuel is contained in double-walled pressure tubes that pass through the blanket-

One main difference between the **CANDU** design and the ATW design is that an aqueous homogeneous continuous flow primary loop is employed in the ATW design, whereas solid heterogeneous fuel is used in **CANDU**. It is necessary to use the liquid fuel approach for actinide burning in ATW to take advantage of the high neutron flux and the rapid burn rates, as well as facilitate the continuous processing. This approach complicates the design somewhat because the fuel will be **circulated** both inside and outside the blanket region. To prevent the radioactive liquid fuel **from** coming into contact with the steam loop that drives the turbine-generator set, an intermediate light-water heat-transfer loop is added. The additional intermediate loop adds another degree of safety margin in case of accidental radioactive release from a rupture **from** the liquid fuel primary loop. The heat from the intermediate loop is **transferred** to a light-water secondary loop for steam generation. In this concept, energy produced in the blanket **from** the fission of the **actinides** is recovered and used to produce electricity with about 30% thermal efficiency. This recovery of useful energy greatly reduces the overall cost of the facility.

A proposed layout for the target/blanket is shown in Fig. 3. The solid **tungsten/lead** target is surrounded by an **annulus** of technetium in **D₂O**, an actinide slurry region, and a **D₂O** reflector. The target/blanket design is evolving in an effort to enhance the safety, reliability, and **efficiency** (with respect to neutron utilization). For example, it may be possible to keep the actinide slurry within the blanket region, and transfer heat to the heavy-water moderator that is pumped outside the blanket. This retention will greatly reduce the overall volume of actinide slurry and reduce the risk of spills outside the blanket.

Another option that is being considered is the use of a flowing liquid lead target. The use of such a target adds complexity to the design but has the potential to increase the neutron utilization efficiency. In addition, if greater accelerator currents are required, the liquid target can

more easily remove the heat. The base-case target/blanket design and some of the design options are presented in more detail below.

111. TARGET/BLANKET DESIGN OBJECTIVES

The design of the target/blanket region for ATW is complex. As the target intercepts the proton beam, the heating rates are exceptionally **high (9 MW/l peak)** and the neutron **flux levels** produced are much higher than those for commercial nuclear reactors. The neutrons are very expensive to produce when an accelerated proton beam is used. Therefore, the design of the **target/blanket** is crucial; it must use the neutrons as **efficiently** as possible while **maximizing** safety and reliability considerations. We have specified several goals in the design of the **target/blanket** (values given are for one light-water reactor waste discharge):

- **transmute** 33 kg of technetium and 7 kg of iodine per **year**;
- transmute 300 kg of **actinide** mixture per year (**neptunium**, americium, plutonium);
- minimize parasitic capture in structures and **target**;
- use **materials** that are compatible with fluids to minimize **corrosion**;
- use materials that avoid **generation** of long-lived radioactive **nuclides**;
- use materials that can withstand intense irradiation,
- provide a safe system that will easily meet nuclear facility licensing and safety **goals**;
- maintain **subcriticality** during normal and off-normal situations;
- use passive modes of heat removal where **possible**;
- ensure easy operation and **maintenance**; and
- minimize long-term research and development activities and costs.

IV. TARGET DESIGN OPTIONS

The protons **from** the split beam are directed onto the neutron production target region. There are several high atomic number materials that can be used for the **spallation** target including lead (**Pb**), lead-bismuth (**Pb-Bi**), tungsten (**W**), tantalum (**Ta**), and uranium (**U**). Through **spallation** in a lead **target**, approximately 50 neutrons are produced for each incident **1600-MeV** proton. Neutron production is reduced to about 22 neutrons per proton for the **800-MeV** application, and if a tungsten target is used, the neutron production decreases to approximately 18 neutrons per proton for the **800-MeV** accelerator. For heat transfer and power density considerations, approximately **75%** of the thermal energy in the beam is deposited directly into the

target; the remainder of the power is removed by neutrons and gammas that leak into the blanket. For the 1600-MeV, 62.5-mA proton beam, approximately 100-MW_t power is **generated**, of which 75 MW_t is deposited in the **target**.

For the **ATW** commercial waste application, two target design options have been considered. The base-case design uses solid tungsten rods cooled with heavy water. As an option, a flowing liquid lead or lead-bismuth system is proposed, either of which act as the **spallation** target material and heat transport medium.

A. Solid Target Design

The solid target is made of separate regions of tungsten and lead (Fig. 4). Because of its high melting point (3300° C), tungsten would be used primarily in the high thermal energy region of the target. Because the interior of the solid target is pressurized, a window is required to separate the accelerator vacuum **from** the target region. For this configuration, heavy-water cooling is required for the window and the tungsten rods to keep the temperature gradients at reasonable values. The neutron absorption in **D₂O** is small and has the added benefit of moderating the neutrons. However, **D₂O** will interact with the proton beam and cause some degradation in neutron production.

A region of solid lead surrounds the tungsten target. The lead provides a boost in the **neutron** production from additional **spallation** and **(n,xn)** reactions. The lead is cooled with **D₂O** to keep it well below the melting point. A disadvantage to using tungsten as the **main** target material is that it has a significant neutron absorption cross section. Enriching the tungsten in the isotope **¹⁸⁴W** would reduce the absorption approximately by a factor of 10 but would be very expensive. To increase the neutron leakage **from** the tungsten and therefore reduce the parasitic absorption, we have introduced flux traps in the spaces between the tungsten rod units. We have successfully demonstrated this concept at the **Los Alamos** Neutron Scattering Center (**LANSCE**).

1. Target Detail. The target is designed as a separate assembly that can easily be removed and replaced during routine maintenance. The target assembly consists of a spherical shaped window, several tungsten rod units, and coolant pipes. The entire assembly is encased in a single structure and is removed and replaced as a single component. The window and tungsten rod units are cooled with **D₂O**. The regions between the tungsten rod units are **filled** with helium gas that is at a slightly higher pressure than the **D₂O** coolant (about 3.4 M_Pa), which maintains a compressive force on the container of each unit and reduces the mechanical stress. The window and the containers for the individual tungsten units are made of **Inconel 718**. The structural container for the entire assembly is made of an aluminum alloy.

2. Window. At high-energy accelerator facilities, a metal **interface** is used to maintain a vacuum in the accelerator and to protect the equipment in the experimental area. Because of the

intense energy of the ATW accelerator, a windowless system is desirable, and it is possible that the flowing liquid target may provide that capability. However, if a solid target is used, a window is required. Even if the liquid target is used, it may be necessary to first test the system with a window to determine if the vacuum pumps can keep up with the pressure demand before actual operation takes place without a window. Therefore it is important to study existing window designs and to determine their applicability to ATW.

Results of research at the Los Alamos Meson Physics Facility (**LAMPF**) have provided considerable data about window survivability. The **LAMPF** accelerator is 800 MeV, 1 mA. The double-walled flat window used on the current beam stop at **LAMPF** has undergone 17,500 h at 20-30 $\mu\text{A}/\text{cm}^2$, for a total proton fluence of $1.3\text{e}22$ p/cm^2 . This window is made of **Inconel 718**. Another **Inconel** window is used for the vacuum-t-air interface. It is replaced yearly and accumulates approximately $2.9\text{e}21$ p/cm^2 . The proposed window for the Swiss Nuclear Institute Neutron Source (**SINQ**) is being tested at **LAMPF**. This material is a steel (**Fe10.5Cr**) and has also accumulated $2.9\text{e}21$ p/cm^2 of fluence proton.

For comparison to ALTU, we assume that the **1600-MeV, 62.5-mA** proton beam is roughly equivalent to 800 MeV, 140 mA with respect to material damage. This provides an average flux of about 224 $\mu\text{A}/\text{cm}^2$ and in one full-power year $3.52\text{e}22$ p/cm^2 of proton fluence. This fluence is a factor of 2.7 times the current fluence on the **LAMPF** beam stop still in service. Based on this experience, it may be reasonable to assume that a window in ATW would require replacement about every 6 months.

3. Tungsten Rod Units. The proton beam that strikes the target is a 25-cm square spatial pattern with a nearly flat intensity profile. The tungsten rod units are also square and are 30 cm on a side. The **D₂O** coolant enters one side of the unit through an inlet plenum and exits the opposite side through a similar plenum. The coolant feed for each unit is separate, which allows for pressure and integrity checks before final assembly. Because the beam degrades as it passes through the units, there is a reduction in the power density from one unit to the next. To flatten the axial power distribution and the neutron source, we vary the amount of tungsten in each unit. An orifice between each unit may be required. In the horizontal direction, because the power density is flat across the tungsten square, there is no need for orifices within each individual unit. Although the rod units are square, they fit inside a cylindrical structure that is about 40 cm in diameter. The rest of the surrounding blanket is also cylindrical.

Within a single tungsten rod unit, the rods are encased in a thin shell of **Inconel 718** (Fig. 5). This contains the **D₂O** coolant and provides a structural support for the rods. The rods are set in several sets of two rows, each with flow passages in between. From a distance, the rods appear to form a plate with a bumpy surface. The rods are held in place at the ends in a manner that allows them to expand and contract, which thereby significantly reduces the thermal and

mechanical stresses. The number of rod sets (or “plates”) in each unit can vary to flatten the axial power distribution. **In** each unit, the tungsten occupies **about** 50% of the **free** volume. The rest is occupied by the **D₂O** coolant.

4. Heat-Transfer Analysis. Detailed heat-transfer calculations were performed for a typical channel in the peak power density region (the tungsten unit that is first hit by the proton beam). **In** this region, the calculated power density in the tungsten is 8.64 **MW/l**, and in the **D₂O** 0.327 **MW/l**. For our geometry, this is equivalent to a heat flux of about 10.8 **MW/m²**. For the heat-transfer analysis, “plate” geometry was assumed with the plate thickness equivalent to a **two**-row set of rods. The calculations were **performed** with a **multiphase** flow code that can predict the onset of boiling with reasonable accuracy. The tungsten “plate” was modeled as a slab with 30 **axial** nodes placed along its length. Each **axial** node *contained* 10 horizontal nodes to **resolve** the temperature distribution between the middle of the “plate” and the surface that is in contact with the coolant. The results show that for a coolant pressure of 5 **MPa**, inlet temperature 300 K, and a velocity of 13 m/s, the flow will remain single phase. The peak tungsten temperature is about 450 K, which is small compared with the melting point (3800 K). If **subcooled** boiling is used as the heat-transfer mechanism, the pressure may be reduced significantly. **Experimental** results for fusion reactor applications (**Refs. 1–3**) show that heat fluxes greater than 20 **MW/m²** may be obtained. Therefore, it appears that adequate cooling can be achieved.

The hydraulic design of the target will be **verified** with full-scale tests to experimentally check the flow distribution through the assembly and each tungsten unit. **Subscale** tests will be used to **verify** the thermal design to ensure that cooling is adequate for both normal and off-normal conditions.

5. Decay Heat. The decay heat has been predicted for the main components of the target assembly using the **LAHET/MCNP/CINDER** code system. The results show that after 1 year of operation at 80% availability, the decay heat in the tungsten amounts to 0.4% of the total heat generated at **full** power. *This* heat decays quickly and after 1 day is down to 0.1 %.

B. Liquid Target Design

The flowing liquid **spallation** target is a pure lead material (melting point 600 K). The lead target material has a high atomic number, good heat-transport properties, and is compatible with **low-chromium**, low-nickel stainless steel alloys such as **Croloy**. A lead-bismuth **eutectic** was originally chosen as the material of choice for the liquid target option because of its low melting point, high boiling **point**, low vapor pressure, and high atomic number. However, because of the production of ²¹⁰Po from the bismuth (a major **radiologic** hazard), this material has fallen out of favor. **Instead**, lead is now the material of choice and is the focus of much of our attention for the flowing target system. Although lead operates at a higher temperature, it appears the difficulties

with respect to cooling, vapor pressure, and operation can be overcome. Evidently, the Soviets have experience with liquid lead coolants for some of their reactors. Still, lead produces some long-lived waste products when irradiated in the proton beam and becomes a mixed waste (toxic and radioactive). When compared with the solid tungsten target, this puts the flowing lead system at a disadvantage. One advantage of the flowing liquid lead design is that a window between the beam and the target may be unnecessary if the vapor pressure of the liquid at the surface is sufficiently low that the vacuum may be maintained.

The orientation of the flowing liquid target is vertical with the liquid lead flowing upward through an **annulus**, turning, and then flowing downward into a cone-shaped open channel. Here the fluid interacts with the proton beam and is in direct contact with a vacuum. As the target material is flowing downward through the channel, the **spallation** reactions cause **significant** heat generation. The fluid coalesces in the central target area that is about 50 cm in diameter and flows downward at about 2 m/s. The flow rate is sufficiently high to convect the heat away at a reasonable temperature. In the target region, the bulk temperature increase is 100 K (**from** 700 K to 800 K). The flow exits the bottom of the target region and is pumped through a heat exchanger. The heat is transferred to a secondary sodium-potassium (**NaK**) **eutectic** heat-transport loop and then dumped to the atmosphere through a forced-air heat exchanger. The loop components such as the pump and heat exchanger will require development and testing to ensure an adequate design. The **NaK** system and dump heat exchanger are similar to the heat dump system used in the Fast Flux Test Facility.

The neutron damage to the container wall may require replacement **every** 1 or 2 years. The intense flux occurs in a relatively small 50-cm section of the wall. Here the static head of fluid is very low (0.09 **MPa** or 13.2 psi) and the calculated hoop stress only 4.68 **MPa** (680 psi). Because of the low stress, material lifetimes may be increased. A double-wall construction is required to provide thermal insulation between the target and the blanket, and to prevent liquid metal-water interaction in the event of a leak.

v. **BLANKET DESIGN**

The ATW blanket design for each of the four modules is based on the existing **CANDU** reactor design. The primary reasons for this choice is that the **CANDU** system uses heavy water as the coolant and moderator, and the power level is approximately the same for each module as in the **CANDU-3**. Therefore, ATW can benefit from this experience base, which will minimize overall research and development costs.

A. Comparison to CANDU Heat Transport System

Table 1 provides a comparison of key design parameters for **CANDU-3** and one target/blanket **module** of ATW.

B. MATERIAL COMPATIBILITY

The current ATW base-case design employs zirconium alloys for containment vessels and piping in heavy-water-cooled environments. Temperatures **from < 100° C** to 300° C are anticipated. Neutron flux of about $2 \times 10^{15} \text{ n/cm}^2\text{-s}$ is anticipated and the neutron spectrum varies between mostly thermal and mostly **fast-reactor-like**, but because of a lead shield, not **spallation-like**. In addition, the presence of technetium salts in the slurry require that the **pH** of the slurry be approximately 11. With respect to the performance of **zircalloys** in the anticipated environment there is a wealth of information.

Zircalloys were developed for cladding in light-water reactors. Therefore their performance in a thermal neutron spectrum and water environment is excellent. In the above application, **zircalloys** will perform adequately provided a number of precautions are addressed.

It is important to note that almost every light-water reactor in the world has been shutdown at least once because of corrosion problems, so no system is likely to be perfect. The **zircalloys** are the best materials at moderate temperatures (< 400° C) in water environments. **Zircalloys** (and all other reactor metals) are reactive with water and obtain their protection **from** a tightly adherent oxide surface layer. Light-water reactors typically add LiOH to the water to obtain a **pH** of 9 to 11 as a corrosion inhibitor. At a higher pH, LiOH enhances **corrosion**. The mechanism of LiOH inhibition is not completely understood, but lithium interaction with the oxide protective layer is thought to play a role. Therefore, it is likely that technetium salts with a **pH** of 11 will not be corrosive to **zircalloys**, but some caution is required.

In addition, corrosion in **zircalloys** is enhanced by dissolved oxygen in the water. **Zircalloys** are sensitive to hydrogen **embrittlement**. Hydrogen enters the alloy by the reaction $\text{Zr} + \text{H}_2\text{O} \rightarrow \text{ZrH} + \text{O}$. **ZrH** forms interstitial platelets that decrease the alloys' strength. At much higher temperatures such as would be experienced in a reactor coolant loss, **zirconium** reacts with water to form hydrogen gas by the reaction $\text{Zr} + \text{H}_2\text{O} \rightarrow \text{ZrO}_2 + 2\text{H}_2 + 140 \text{ kcal/mole}$. This situation obviously causes an explosive hazard. **Zircaloy-4** is similar to **zircaloy-2** except that **zircaloy-4** has no nickel content.

Zircalloys have been irradiated up to 100 displacements per atom (**dpa**) thermal neutron **fluence**. The amount of swelling and the decrease in ductility caused by thermal neutrons is known and can be accommodated by proper design. However, many variables are important in determining the response of any material to irradiation and extrapolation to different conditions is

not straightforward. Performance (expected service life) will be dependent on the irradiation temperature and flux, the neutron **spectrum**, and chemistry of corrosion.

Erosion is an issue with **ATW** systems using metal-oxide (ceramic particulate) slurries. Flowing water enhances corrosion, especially at bends in tubing and in pumps. Because the **materials** depend on an oxide layer to prevent corrosion, an abrasive slurry that abrades or pits the layer will increase the corrosion rate. This effect can be very severe at high flow velocities. Overall, for the current **ATW** application, **zircaloy-4** would be the preferred alloy.

VI. PRELIMINARY PHYSICS ANALYSIS OF THE BLANKET

The aqueous blanket incorporates a flowing **D₂O** “slurry” of pressurized water reactor actinide wastes contained in **CANDU-type** pressure tubes. The double-walled pressure tubes are immersed in a **D₂O** moderator that surrounds the neutron-producing target. Long-lived **fission-product** wastes are circulated through the blanket, either directly in the moderator or in separate tubes.

The composition of the actinide waste was obtained **from an ORIGEN** point depletion calculation using a **CANDU** neutron spectrum. Table 2 shows the relative isotopic composition of the light-water reactor actinide wastes and the equilibrium composition **after irradiation** in a flux of 1.0×10^{15} that was used in our calculations. Plutonium isotopes constitute almost 90% of the feed material mass but only 50% of the equilibrium mass. The primary **fissile** isotopes are ²³⁹Pu and ²⁴¹Pu.

A. LATTICE CELL CALCULATIONS

Figure 6 shows a schematic of the lattice cell employed in our initial calculations. The cell contains the actinide slurry surrounded by double-walled zirconium pressure tubes and the associated **D₂O** moderator. Dimensions and area fractions are shown for the configuration that was eventually chosen for the reference design.

Initial calculations investigated the effects of **actinide** concentration, tube size, and tube spacing on **k_{eff}** (or **k_∞** in the case of the non-leakage unit cell). Figure 7 shows the dependency of **k_∞** on actinide concentration. Initially increasing the actinide concentration from 5 g/l has a pronounced positive effect on **k_∞**, but the curve flattens out at approximately 30 g/l. At low actinide concentrations, the neutron balance is dominated by captures in the **zirconium** tubes, which account for **60%** of all captures at 5 g/l. Initial introduction of **fissile** material effectively competes with the zirconium captures, reducing those captures to 13% at 50 g/l where that effect becomes relatively small. However, increasing the **actinide** concentration also hardens the neutron spectrum in the slurry, which increases the **epithermal** and resonance captures in the **non-fissile** isotopes

(primarily ^{240}Pu and ^{242}Pu) relative to the predominantly thermal fission reactions in the **fissile** isotopes. Once the zirconium captures become less **important**, the spectral-hardening effect tends to mitigate further increases in k_{∞} . A finite system might continue to exhibit an increasing k_{eff} dependency if the addition of **fissile** material competes with neutron leakage. However, the proposed ATW blanket is a relatively large, low leakage system, and should show the same behavior as found in the lattice-cell calculations. Therefore, for the hexagonal lattice configuration used in these calculations (**10-cm-diam.** tubes spaced 17.0 cm apart), it appears that the maximum attainable k_{∞} is approximately 1.0. Considering that we need additional reactivity to compensate for leakage, **target**, and structure absorption, as well as fission product poisoning and burning, it is **doubtful** that this value of k_{∞} will provide the high multiplication needed for **satisfactory** system performance.

The effects of the slurry-tube radius and pitch are shown in Figs. 8a and 8b, which are **parametric** plots of the same data. For the tube-size investigation, the relative areas of slurry, walls, and moderator were maintained as the cell dimensions were varied. Figure 8 shows that at a given pitch, k_{∞} increases only slightly with increasing tube size, but increasing the tube pitch (spacing) can yield significant improvements in k_{∞} . A larger tube pitch increases the moderator/slurry ratio. This allows fission neutrons that leave the slurry to slow down and **thermalize** in the D_2O moderator before returning to the slurry, thereby reducing the probability of resonance capture in the **nonfissile** actinides. This effect can be seen in Fig. 9, which plots the actinide capture-to-fission ratio as a **function** of tube spacing (capture is defined as a non-fission absorption). Increasing the pitch strongly reduces the overall parasitic capture probability. Because a constant **actinide** slurry area is required for heat-transport considerations, increasing the **slurry/moderator ratio** will also increase the blanket size.

The final cell calculations extended the range of **actinide** concentrations up to 100 g/l. These results are shown in Fig. 10. The lower curve in Fig. 10 is an extension of the plot in Fig. 7 and confirms the inability to increase k_{∞} for that configuration (**17.0-cm** pitch). However, at the larger spacings, increasing the actinide **concentration** can achieve further gains in k_{∞} .

Based on these cell calculations, we selected a 5-cm-radius tube, a 23.8-cm pitch, and a 75-g/l **actinide** concentration as the reference lattice configuration to be used in the blanket calculations. The resulting value of k_{∞} , shown in Fig. 5 as a large dot, is 1.158. A neutron balance for the reference **lattice-cell** configuration is shown in Table 3.

B. BLANKET CALCULATIONS

A schematic of the target-blanket configuration is shown in Fig. 3. The target is a homogenized version of the solid tungsten rod design. Primary regions in our one-dimensional model are the homogenized tungsten rods, a **Pb-D₂O** region, an aluminum target structure, and a

zircaloy tank wall followed by a thin 1 molar **Tc/D₂O** buffer region whose thickness was varied. A **D₂O** outer reflector surrounds the “active” **actinide** blanket region.

Because ONEDANT is a one-dimensional code (cylindrical geometry in this case) that cannot accept azimuthal geometry changes, we must homogenize the lattice cell for incorporation into the blanket calculations. This was done by flux-volume-weighting the constituent materials and the cross sections in the unit cell to preserve reaction rates. The accuracy of this approximation must be checked by a more rigorous **MCNP** calculation that can more accurately model the detailed geometry.

Before homogenizing the lattice cell, we incorporated **¹⁰B** into the **actinide** slurry to simulate the buildup of fission products (excluding technetium and iodine). The **¹⁰B** concentration was adjusted to produce the same fission product absorption rate relative to the total **actinide** absorption that were calculated for a 10-day buildup at a flux of 1.0×10^{15} . The addition of the appropriate amount of **¹⁰B** reduced the k_{∞} of the lattice cell by $0.057 \Delta k$.

The blanket calculations incorporated the homogenized (flux-weighted) cell cross sections that contained the **¹⁰B** simulated lumped fission products. The k_{eff} of an infinite-length blanket configuration was $0.057 \Delta k$ less than the corresponding cell value. This reduction is due to radial leakage and neutron capture in blanket materials. An additional 0.035 was lost in the inclusion of an axial-leakage approximation into the one-dimensional calculation. The resulting blanket k_{eff} was 1.008.

The next step in the blanket calculations was to simulate burning of long-lived fission products produced in the pressurized water reactor cycle as well as in the blanket itself. Again, relative neutron captures per fission required to burn technetium and iodine were obtained from the **ORIGEN** calculation. The following procedure was followed for these calculations:

1. An operating value of k_{eff} was selected
2. To simulate fission product burning, an **eigenvalue** calculation was performed in which technetium was added homogeneously throughout the **actinide** region and the outer **reflector** to reduce the k_{eff} from the initial value of 1.008 to the target value selected in step 1.
3. A source problem was run with the appropriate technetium concentration determined in step 2.
4. Results from step 3 (power and the ratio of technetium captures per fission) were normalized to a **800-MeV, 140-mA** beam (or equivalently to a **1600-MeV, 62.5-mA** beam) and compared with the goal values required to support two light-water reactors.

5. The above steps were repeated for varying thicknesses of the inner **Tc/D₂O** buffer region.

These steps were followed for operating **k_{eff}** values of 0.95 and 0.93. The results are shown in Figs. 11 and 12, respectively. Two figures of merit are plotted as a **function** of the **Tc-D₂O** buffer region **thickness**: (1) the blanket power and (2) the technetium burn rate, both relative to the respective goal values required to service two pressurized water reactors. From Fig. 7 we see that at an operating **k_{eff}** of 0.95 and no technetium buffer region, sufficient neutron multiplication exists to produce approximately 1.2 times the 1640 **MW** needed to burn the **actinide** wastes from two pressurized water reactors. However, at this relatively high **k_{eff}** there are not enough excess neutrons available to burn the required technetium atoms (only 72% of the goal technetium burn is achieved). Increasing the buffer region thickness puts technetium atoms in a high flux region and increases the technetium captures relative to the fissions. However, **capturing** the source neutrons before they are multiplied in the actinide region reduces the effective multiplication and the resulting power. These conflicting trends are shown in Figs. 11 and 12. At the point where the two curves **intersect**, the two parameters (**actinide** and fission-product burn) are balanced for the needs of the pressurized water **reactor** cycle. At a **k_{eff}** of 0.95 and a buffer region thickness of 1.9 cm, the blanket will operate at a power level of 1542 **MW** and support 1.88 pressurized water reactors. A **neutron** balance for this **configuration** is shown in Table 4.

Operating at a **k_{eff}** of 0.93 allows the incorporation of more technetium atoms and provides more excess neutrons so that for a given buffer-region thickness more technetium will be burned. However, Fig. 12 shows that even with no buffer region, the blanket will burn only 88% of the technetium goal amount. At a **k_{eff}** of 0.93 there is also insufficient multiplication to meet the power goal, and adding a buffer region to increase the technetium burn will further aggravate the power deficiency. Thus, a blanket operating at a **k_{eff}** of 0.93 would not have any technetium buffer region and would service the equivalent of approximately 1.7 pressurized water **reactors**.

In calculating the achievable technetium burn, we included radial neutron leakage as being captured in technetium. This could be achieved by increasing the technetium concentration in the outer part of the radial reflector and/or by making it thicker.

The reference blanket has an **actinide** inventory of 325 kg in a 250 cm length. The corresponding technetium inventories are 82.0 and 99.2 kg in blankets operating at a **k_{eff}** of 0.95 and 0.93, respectively. The one-dimensional fluxes in the **actinide** blanket were calculated to be 1.38×10^{15} and 1.26×10^{15} **n/cm³-s** for **k_{eff}**s of 0.95 and 0.93, respectively. The **two**-dimensional fluxes will probably be lower throughout a **significant** part of the blanket.

In summary, we have performed one-dimensional physics calculations to investigate the performance of an aqueous ATW blanket to burn **actinides** and long-lived fission products **from** the

pressurized water reactor fuel cycle. These calculations indicate **that** a blanket combined with a 800-MeV, **140-mA** (or equivalently a 1600-MeV, **62.5-mA**) accelerator maybe able to transmute wastes from almost two pressurized water reactors.

These calculations must be considered very preliminary. They are based on equilibrium **actinide** and fission product concentrations calculated **from** a point depletion code using a **CANDU** neutron spectrum. Because the blanket performance is very sensitive to data derived **from** these depletion calculations, a second iteration using flux and spectra determined from the initial blanket study must be **performed**. Also, more rigorous two- and three-dimensional physics calculations must be performed to check the accuracy of the one-dimensional calculations and to account for spatial effects. Eventually, two-dimensional depletion calculations should be performed on the blanket design, not just for equilibrium operation, but for the entire ATW operating cycle. Additional cross-section generation effort is also needed. In particular, we need cross sections for several **actinides** not currently available, such as ^{248}Cm , and for D_2O at the relatively high **slurry** temperature of 600 K.

VII. CONCLUSION

A conceptual design for the target/blanket of the Los **Alamos** ATW concept has been presented. The neutronics, mechanical design, and heat transfer have been investigated in some detail for the base-case design. Much more work needs to be done, but at this point it appears that the design is feasible and will approach the design goal of supporting two commercial power reactors with each **target/blanket** module.

REFERENCES

1. Falter, H. D. et al., "Tests of High-Heat-Flux Target Plates for the JET Divertor," American Nuclear Society Transactions, November 1991, pp. 655-666.
2. **Celata**, G. P. et al., "Burnout in Highly **Subcooled** Water-Flow Boiling in Small-Diameter Tubes," American Nuclear Society Transactions, November 1991, pp. 657-658.
3. 130yd, R. D., "**Subcooled** Flow Boiling Critical Heat Flux (**CHF**) and its Application to Fusion Energy Components," Fusion Technology, Vol. 7, January 1985, pp. 7-30.

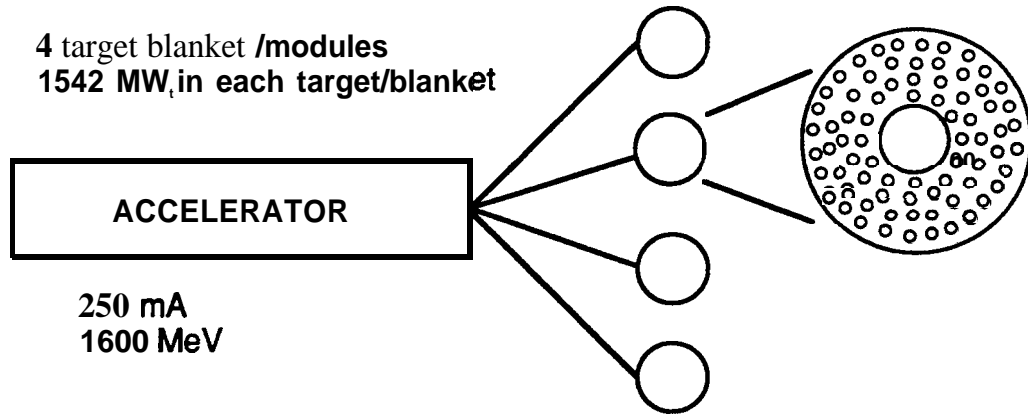


Fig. 1.
A'W schematic.

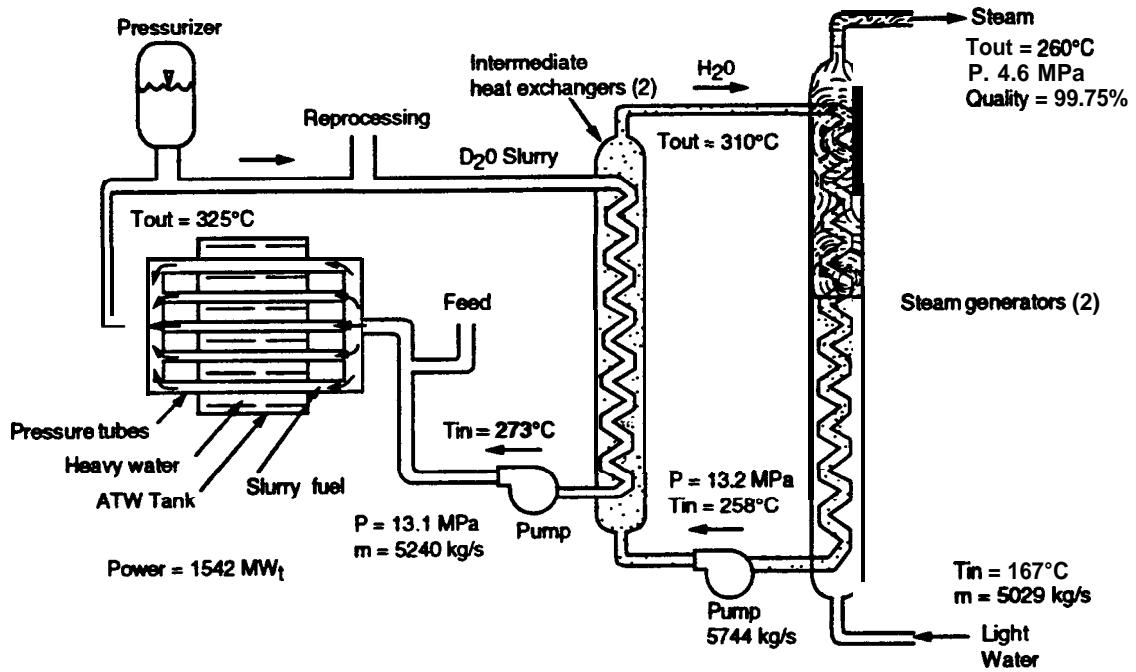


Fig. 2.
ATW balance of plant for the actinide slurry.

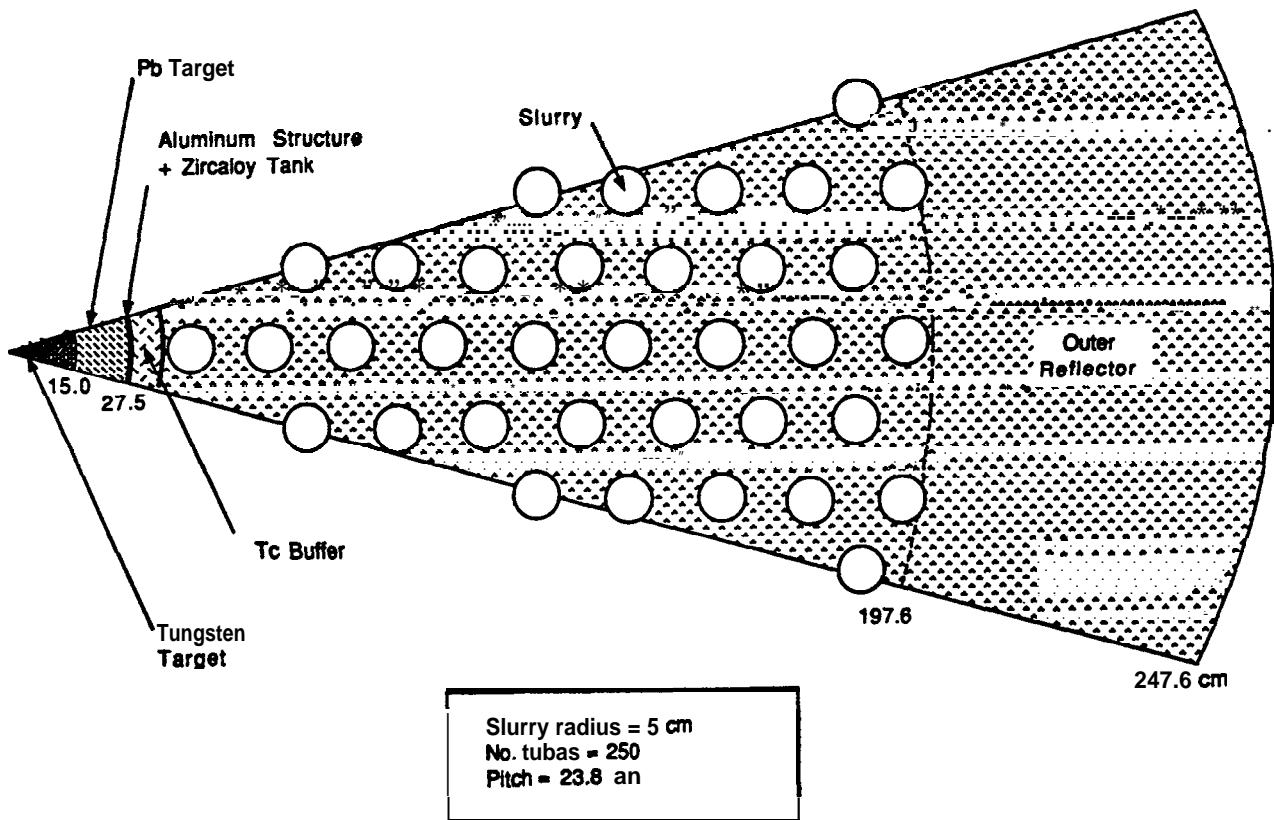


Fig. 3.
ATW target/blanket layout.

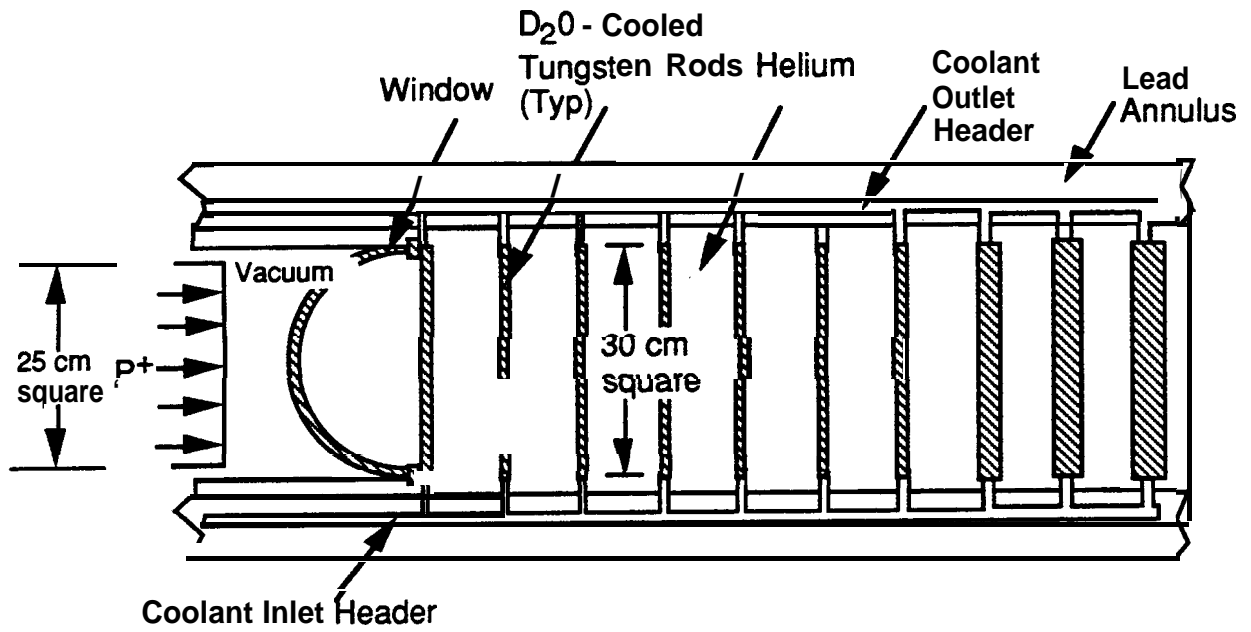


Fig. 4.
Solid target design.

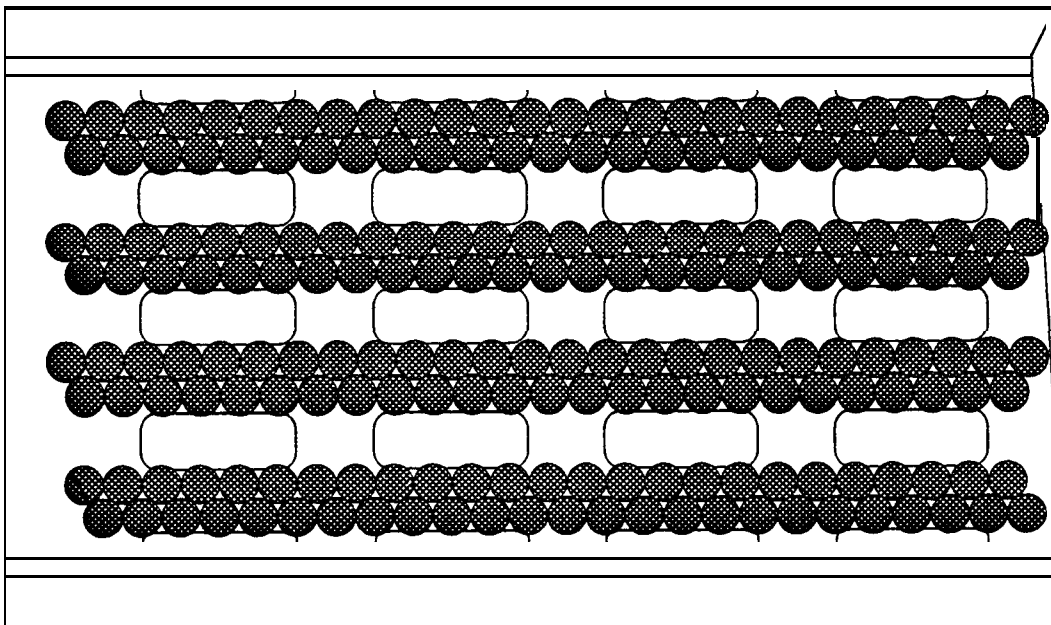
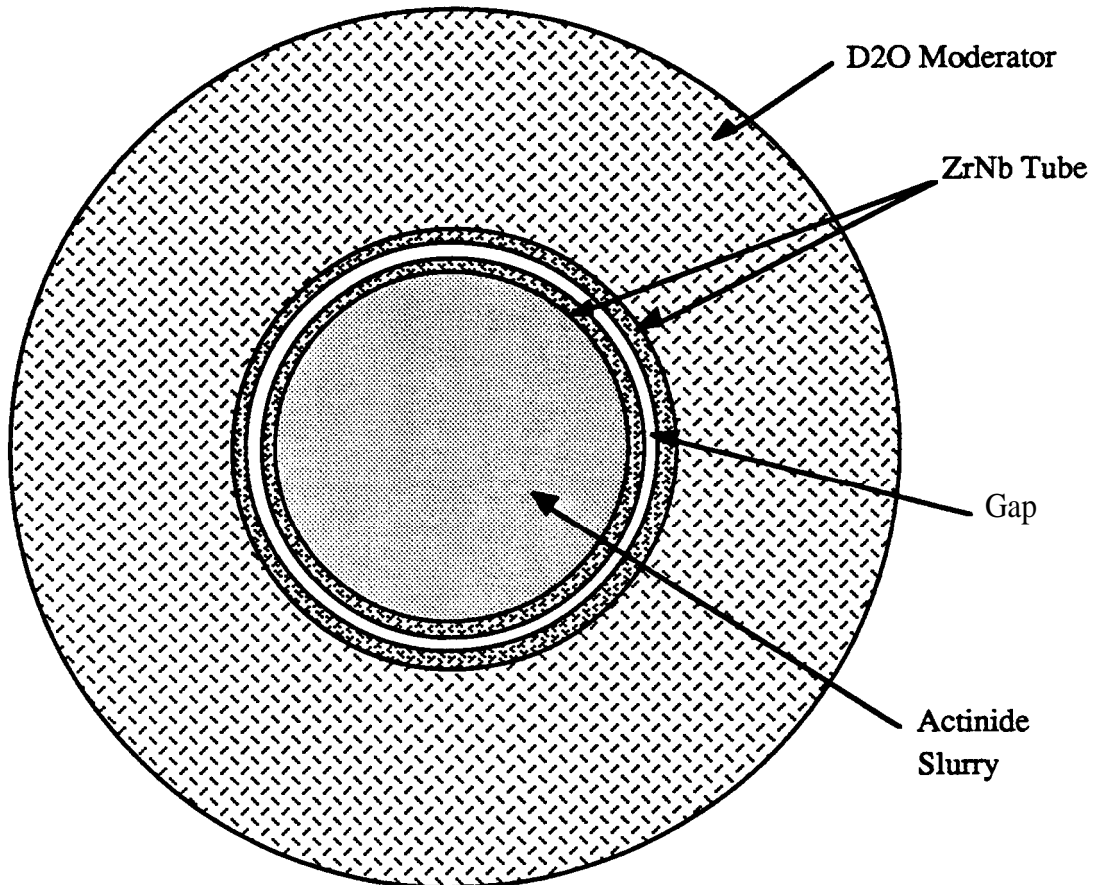


Fig. 5.
End view of the tungsten rod unit.



<u>Region</u>	<u>Radius</u> <u>(cm)</u>	<u>Thickness</u> <u>(cm)</u>	<u>Volume</u> <u>Fraction</u>
slurry	5.0		0.160
Inner Tube	5.42	0.42	0.028
Gap	5.81	0.39	0.028
Outer Tube	6.23	0.42	0.032
Moderator	12.52	6.29	0.752

Pitch = 23.8 cm (9.4 in.)

Fig. 6.
Blanket lattice cell.

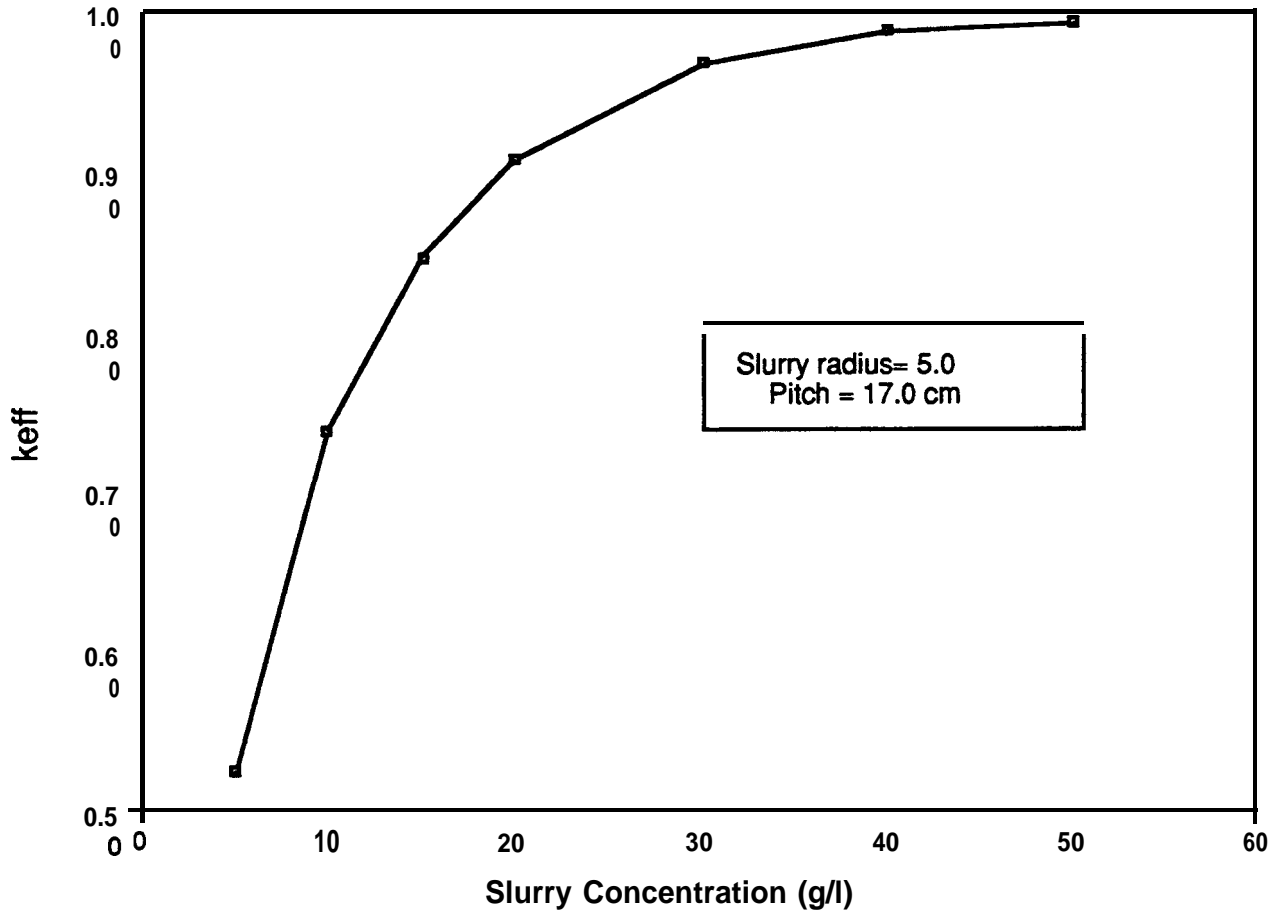


Fig. 7.
 k_{eff} vs slurry concentration for the cell calculation.

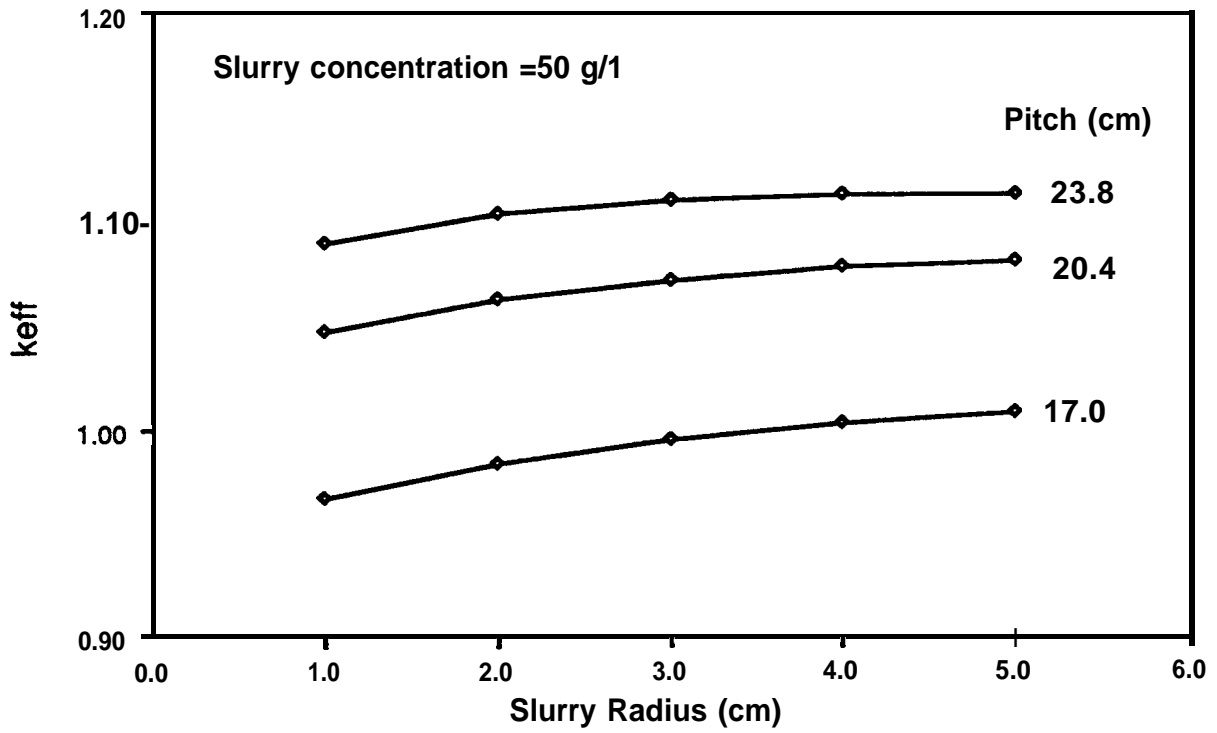


Fig. 8a.
keff vs shiny radius.

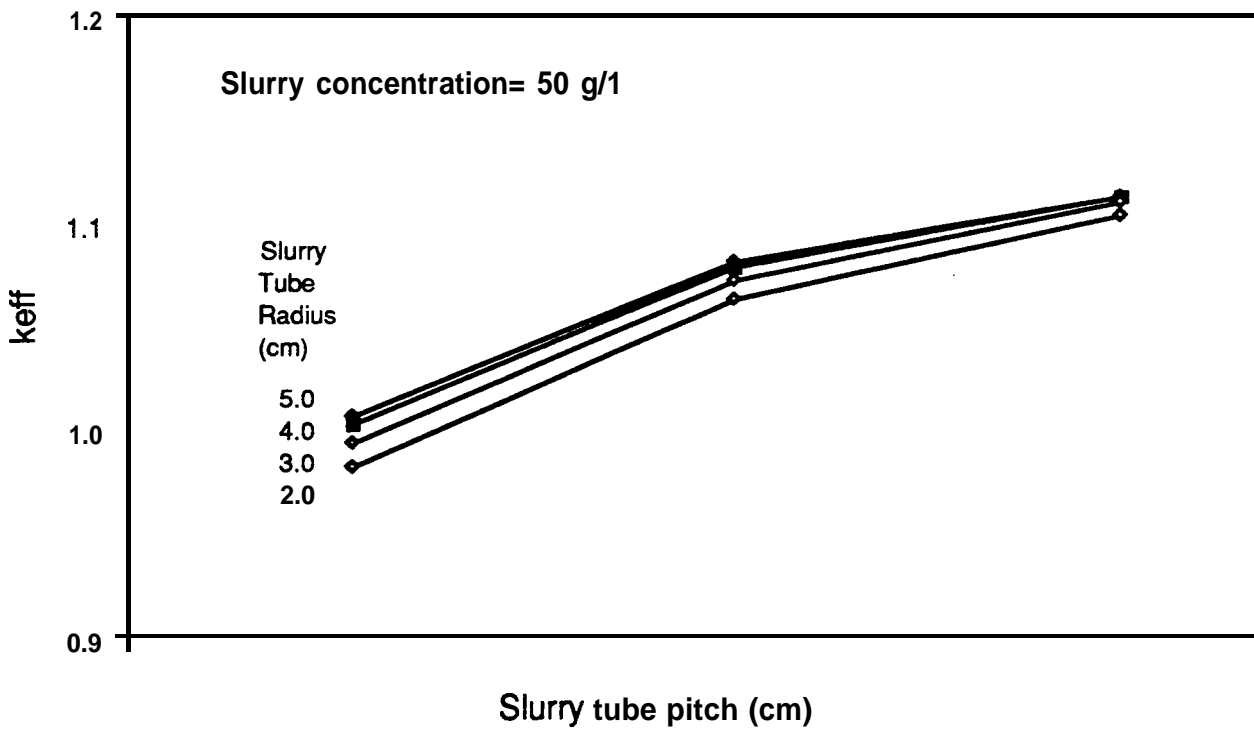


Fig. 8b.
keff vs slurry tube pitch.

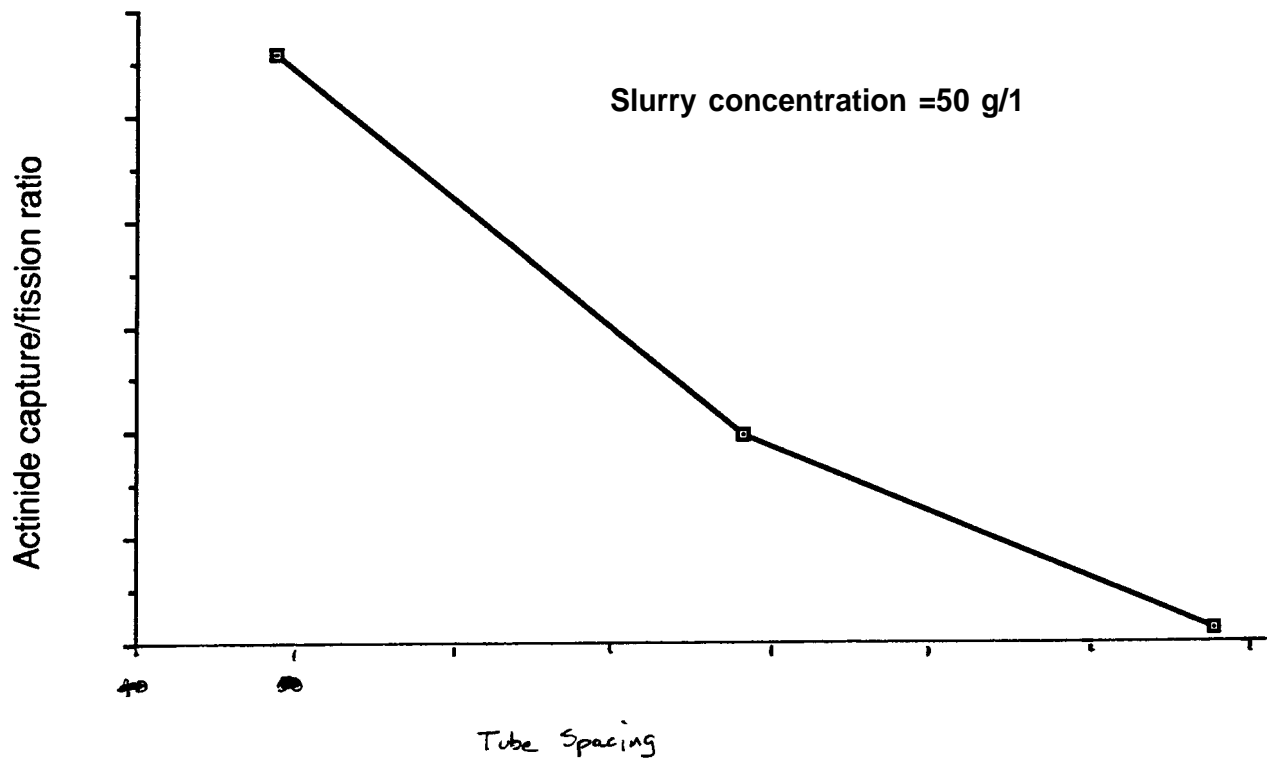


Fig. 9.
Capture/fission ratio vs slurry tube spacing.

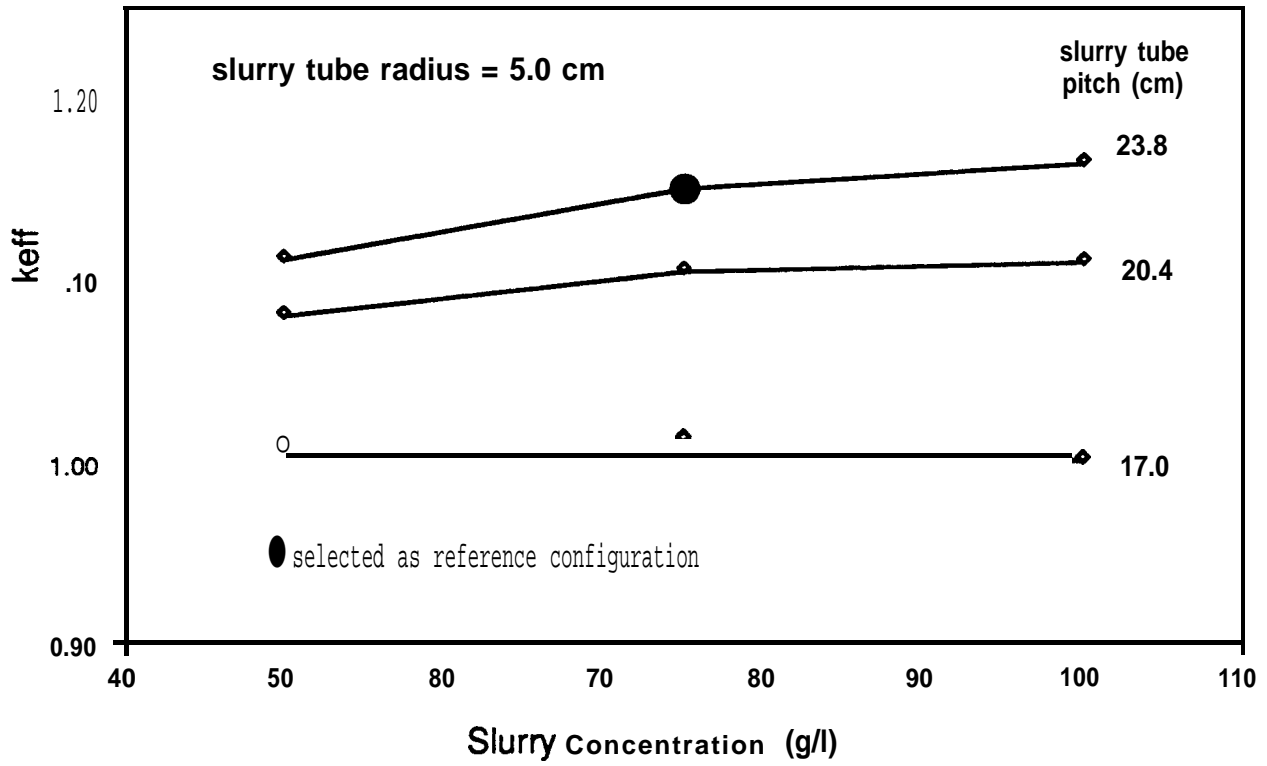


Fig. 10.
 k_{eff} vs slurry concentration.

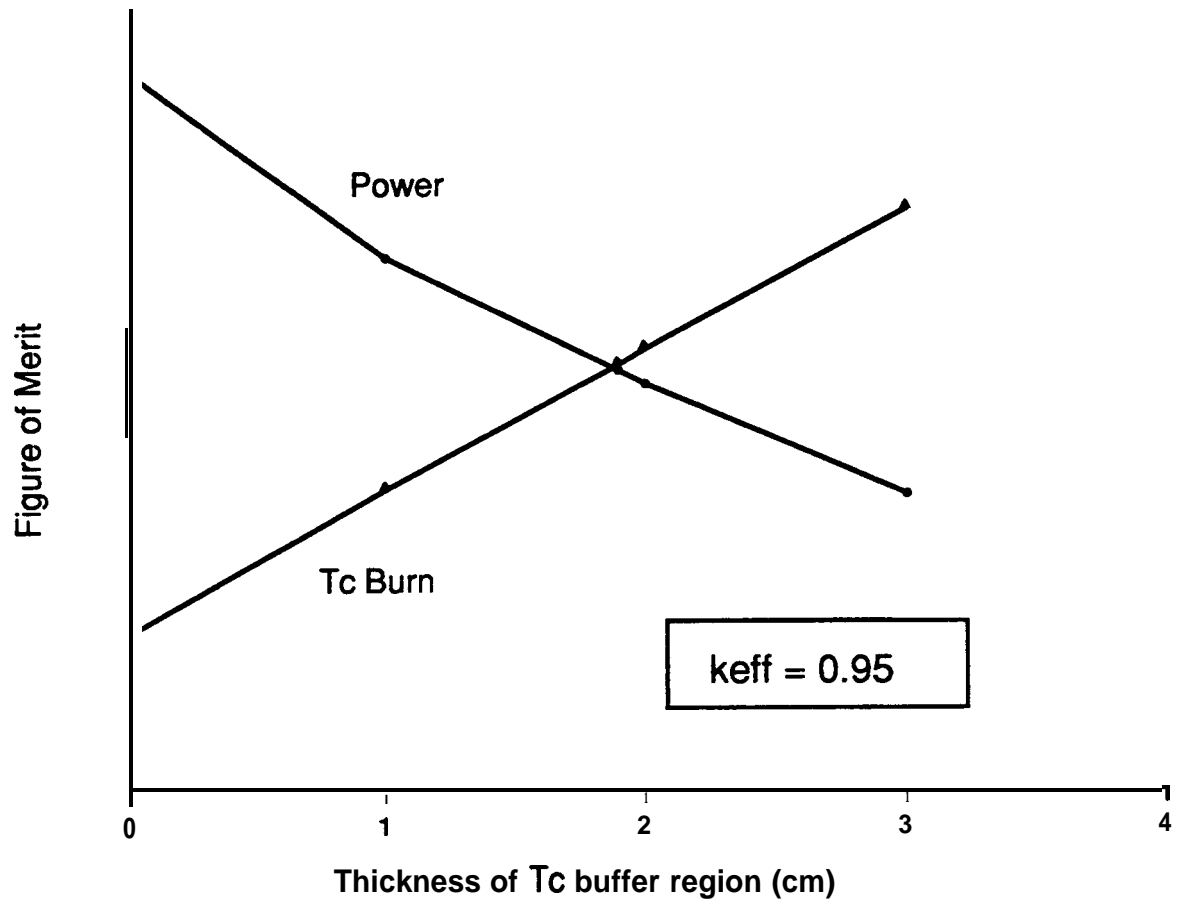


Fig. 11.
Blanket performance at k_{eff} of 0.95.

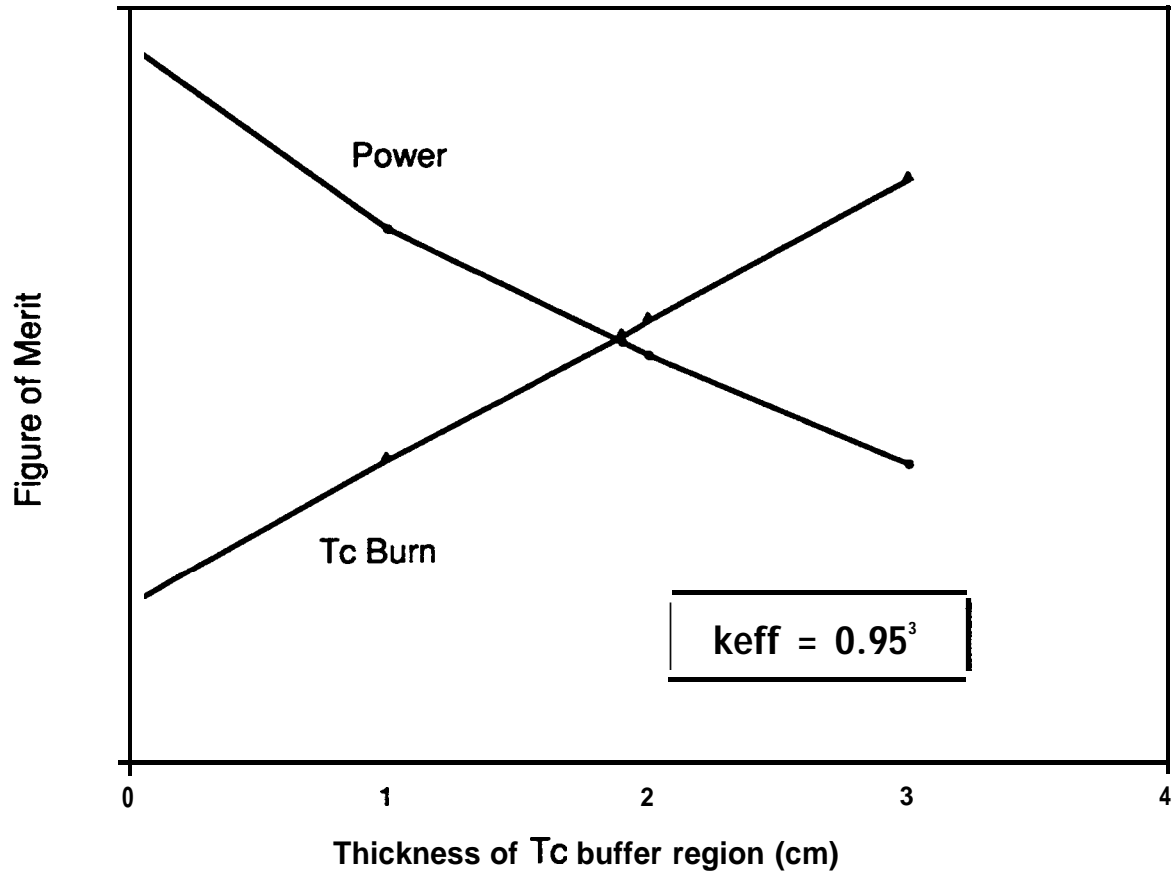


Fig. 12.
Blanket performance at k_{eff} of 0.93.

TABLE 1
KEY DESIGN PARAMETER COMPARISON

PARAMETER	CANDU-3	ATW
1. Blanket Arrangement		
Type	horizontal pressure tube	same
coolant	pressurized heavy water	same
Moderator	heavy water	same
Number of Fuel Assemblies	232	250
Fuel Assembly Material	Zirconium-Niobium	same
Total mass of Fuel	53174 kg	1550 kg
		(total primary loop)
k_{eff}	1.0	0.95
2. Fuel		
Fuel	compacted/sintered	aqueous actinide
	natural UO_2 pellets	solution (75 gin/l);
		(Pu, Np, Am, Cm)
Form	fuel bundle assembly;	flowing fuel solution
	37 elements/assembly	
Bundle Length	0.495 m	
Bundle Outer Diameter	0.1024 m	0.10 m
Bundles/Fuel Assembly	12	
3. Heat Transport System		
Number of Steam Generators	2	same
Steam Generator Type	vertical u-tube	same
Number of Heat-Transport Pumps	2	same
Pump Type	vertical, centrifugal, single suction, double discharge	same
Number of Intermediate Heat Exchangers (II-IX)		2
IHX Type		vertical once-through
Number of IHX Pumps	-	2

PARAMETER	CANDU-3	ATW
IHX Pump Type		vertical, centrifugal
		single suction, double
		discharge
Blanket Outlet Pressure	9.9 MPa	13.1 MPa
Blanket Outlet Temp.	310 c	325 C
Blanket Inlet Temp.	258 C	273 C
Total Flowrate	5300 kg/s	5240 kg/s
Steam Outlet Temp.	260 c	same
Feedwater Inlet Temp.	187 C	same
steam Quality	99.75%	same
Steam Pressure	4.6 MPa	same
IHX outlet Temp.		310 c
IHX Outlet Pressure		13.2 MPa
IHX Inlet Temp.		258 C
IHX Flowrate		5744 kg/s
4. Power		
Total Fission Heat	1440.3 MW _t	1542 MW _t
Net_ElectricalOutput	450 MWe	487 MWe

TABLE 2
RELATIVE ISOTOPIC COMPOSITIONS

Isotope	PWR Discharge	Equilibrium PHI=1.0E15
237Np	0.0449	0.0317
238Np	0.0000	0.0003
239Np	0.0000	0.0000
Total Np	0.0449	0.0321
238Pu	0.0140	0.0230
239Pu	0.5148	0.0610
240Pu	0.2372	0.1003
241 Pu	0.0785	0.0467
242Pu	0.0481	0.2630
243Pu*	0.0000	0.0002
244Pu*	0.0000	0.0026
Total Pu	0.8925	0.4968
241Am	0.0513	0.0048
242Am	0.0000	0.0001
242mAm*	0.0001	0.0001
243Am	0.0092	0.0874
244 Am*	0.0000	0.0000
244mAm*	0.0000	0.0000
245Am*	0.0000	0.0000
Total Am	0.0607	0.0924
242Cm	0.0000	0.0213
243 Cm*	0.0000	0.0006
244Cm	0.0018	0.2794
245 Cm'	0.0001	0.0051
246 Cm*	0.0000	0.0004
247 Cm*	0.0000	0.0094
248 Cm*	0.0000	0.0625
240 Cm*	0.0000	0.0000
Total Cm	0.0019	0.3787
	1.0000	1.0000

•Multigroup cross sections not currently available.

TABLE 3
LATTICE CELL NEUTRON BALANCE

Isotope	Absorption	Fissions
^{237}Np	0.0219	0.0000
^{238}Np	0.0026	0.0024
^{239}Np	0.0000	0.0000
^{238}Pu	0.0435	0.0014
^{239}Pu	0.2777	0.1928
^{240}Pu	0.1513	0.0002
^{241}Pu	0.2594	0.1915
^{242}Pu	0.0380	0.0003
^{241}Am	0.0122	0.0001
^{242}Am	0.0064	0.0053
^{243}Am	0.0439	0.0001
^{242}Cm	0.0020	0.0002
^{244}Cm	0.0243	0.0013
zirconium	0.1033	
D₂O	0.0134	
TOTAL	1.0000	0.3956

TABLE 4
REFERENCE BLANKET NEUTRON BALANCE

keff	0.95
Sources	
Accelerator	1.000
Fissions Neutrons	8.994
Total Sources	9.994
Absorption	
Tungsten Region	0.4146
Lead Region	0.0880
Al + Zirc	0.0330
Tc Buffer	0.1855
Zirc	0.0106
Active Blanket	
Slurry	7.338
Clad	0.829
Moderator	0.097
Tc (Self + LWR)	0.427
Total Active Blanket	8.691
Outer Reflector	
Tc	0.044
D ₂ O	0.020
Total Outer Refl.	0.064
Total Abs.	9.487
Leakage	
Radial	0.167
Axial	0.343
Total Leakage	0.510
Total Losses	9.997
Fissions	3.072
Tc Captures	0.656