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Review

# A Review of Zirconolite Solid Solution Regimes for Plutonium and Candidate Neutron Absorbing Additives

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**Abstract:** Should the decision be made to immobilise the UK Pu inventory through a campaign of Hot Isostatic Pressing (HIP) in a zirconolite matrix, prior to placement in a geological disposal facility (GDF), a suite of disposability criteria must be satisfied. A GDF safety case should be able to demonstrate that post-closure criticality is not a significant concern by demonstrating that such an event would have a low likelihood of occurring and low consequence if it were to occur. In the case of ceramic wasteforms, an effective means of criticality control may be the co-incorporation of a requisite quantity of a suitable neutron absorbing additive, either through co-immobilisation within the host structure or the encapsulation of discrete particles within the grain structure. Following an initial screening of a range of potential neutron absorbing additives, a literature-based assessment of the solid solution limits of a number of potential additives (Gd, Hf, Sm, In, Cd, B) in the candidate zirconolite (CaZrTi<sub>2</sub>O<sub>7</sub>) wasteform is presented. Key areas of research that are in need of development to further support the safety case for nuclearised HIP for Pu inventories are discussed.

**Keywords:** zirconolite; immobilisation; plutonium; wasteform; hot isostatic pressing; criticality



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## 1. Status of the United Kingdom's Plutonium Inventory

Upon completion of domestic reprocessing of spent nuclear fuel (SNF) derived from the civil reactor fleet, the United Kingdom will have accumulated a substantial inventory of separated PuO<sub>2</sub> in excess of 140 teHM (tonnes heavy metal equivalent) [1,2]. This inventory is the result of the PUREX reprocessing (Plutonium Uranium Reduction Extraction) of SNF. In the PUREX process, irradiated fuel assemblies are discharged from a reactor, cooled, sheared and dissolved in HNO<sub>3</sub>, with Pu/U subsequently recovered in a series of solvent extraction steps. Pu(IV) and Pu(III) nitrates are coextracted from the aqueous raffinate by complexation with tributyl phosphate (TBP), followed by precipitation as Pu insoluble oxalates, Pu<sup>4+</sup>(C<sub>2</sub>O<sub>4</sub>)<sub>3</sub>·6H<sub>2</sub>O and Pu<sup>3+</sup><sub>2</sub>(C<sub>2</sub>O<sub>4</sub>)<sub>3</sub>·10H<sub>2</sub>O, via the addition of oxalic acid. The precipitated oxalates are subsequently converted to PuO<sub>2</sub> via calcination in the air or an inert atmosphere, with calcination temperature and time generally inversely proportional to specific surface area (m<sup>2</sup>/g), prior to interim storage awaiting a decision on long term management [3]. The Nuclear Decommissioning Authority (NDA) is responsible for identifying a preferred long term strategy for placing this material beyond reach, satisfying proliferation and safety concerns, that the site license companies will implement. Potential routes towards Pu recycling and/or disposal were defined in the Credible Options Analysis, utilising a life cycle approach to outline and evaluate: (1) pathways to disposition within the constraints of financial and socioeconomic benefit; (2) safety and compatibility with the disposal environment; and (3) readiness and maturity of necessary technology [4]. Several iterations of the Credible Options framework have identified two high level options that are feasible: reuse of the bulk inventory as mixed oxide fuels (MOX) or chemical

immobilisation of the inventory in a suitable matrix. In either case, it is accepted that the timescales for the implementation of any strategy will be several decades at a minimum, and consideration must be given to the quality and source of the Pu feedstock. A summary of UK Pu stocks reported to IAEA under INFCIRC/549 is listed in Table 1. As of 2022, and outlined in the NDA 2022–2025 Business Plan, all separated Pu has been produced and consolidation at Sellafield has been completed. At this stage, a campaign of repackaging for long term storage is underway, with all cans not suitable for extended storage undergoing repackaging. The merits of a MOX fuel-based strategy will not be discussed in this article, and the reader can be referred to the latest NDA Strategy and Position Paper (2019) [1]. Legacy storage of PuO<sub>2</sub> inventories over the past ~60 y has resulted in a fraction of the inventory becoming contaminated with Cl; whilst it is envisaged that this contaminated material will still be suitable for reuse, there is a sizeable inventory of MOX scraps and unirradiated fuel assemblies that will not be suitable for reuse. Any Pu materials designated for disposal (both of high purity and contaminated status) will necessitate specifically tailored ceramic or glass–ceramic wasteforms for immobilisation prior to disposition within a geological disposal facility (GDF).

**Table 1.** 2021 annual figures for holdings of civil unirradiated plutonium in the United Kingdom (reported to IAEA under INFCIRC/549 [5]).

Source	Amount (teHM)
Unirradiated separated plutonium in product stores at reprocessing plants	137.0
Unirradiated separated plutonium in the course of manufacture or fabrication and plutonium contained in unirradiated semi-fabricated or unfinished products at fuel or other fabricating plants elsewhere	0.4
Plutonium contained in unirradiated MOX fuel or other fabricated products at reactor sites or elsewhere	2.0
Unirradiated separated plutonium held elsewhere	1.2
Unirradiated separated plutonium owned by foreign bodies under UK safeguards	24.1
<b>Total</b>	<b>164.7</b>

## 2. Criticality Safety and the Role of Neutron Absorbing Additives in Waste Packages

To construct a site specific safety case for the geological disposal of higher activity radioactive wastes in the UK, Nuclear Waste Services (NWS), a subsidiary of the NDA and the body responsible for the design, construction, and continued operation of the as yet un-sited GDF, maintains a generic disposal safety case outlining the claims, arguments, and evidence that underpin GDF deliverability. A key aspect of the safety case is an assessment of waste package criticality during transport, GDF operation, and post closure [6]. Generic Criticality Safety Assessments (CSAs), as described by Hicks (2007), are considered for waste packages containing one or more of the following four waste components: irradiated natural and slightly enriched uranium; low enriched uranium; high enriched uranium and; separated plutonium [7]. In the interest of brevity, this report will focus only on separated Pu. The reader is referred to Solano et al., (2012) for a comprehensive discussion of criticality control in intermediate level waste and SNF packages [8].

Criticality control measures guard against an inadvertent self-sustaining chain reaction within immobilised fissile material. During the construction and operational phase of the GDF, waste packages will be transported to the facility and placed in a suitable geometrical arrangement in pre-designated areas. The safety of workers and the general public must therefore be ensured by two general approaches: (1) producing waste packages that remain sub-critical by placing an upper limit on the quantity of fissile material contained within and; (2) by optimising the physical form of the waste package such that a critical chain reaction is not possible, either through an arrangement in specific dimensions and geometry that are capable of mitigating the accumulation and critical mass of fissile material, or

through the addition of a neutron absorbing additive. The condition of criticality is met when radionuclides capable of undergoing fission are arranged such that a self-sustaining chain reaction may occur. This is described by the notation  $K_{eff} = 1$ , i.e., the rate of neutrons produced by per fission and the rate of neutrons lost by absorption, or leakage, is equal to unity. Under this condition, a self-sustaining reaction will occur.

Following the closure of the GDF, criticality assurance may be provided through a series of engineered barriers, providing a layered defence known as the multi-barrier concept [9]; however, it is accepted that on the timescale of  $>10^3$  years, water will slowly ingress, resulting in the deterioration of the physical containment of waste packages. The release of radioactive material into the near field is considered to be inevitable, with the possibility of new configurations that could result in criticality. An arrangement of material that is critical would likely produce an increase in temperature and result in the generation of fission products. A suitable additive for mitigating criticality has a high thermal neutron cross-section and may easily be integrated into a wasteform package to reduce the overall neutron flux, thus reducing  $K_{eff} < 1$  and mitigating the possibility of criticality. Inhibiting the neutron flux by the inclusion of additives may also afford the opportunity to increase the upper wasteloading limit, improving the volume footprint of the repository environment and reducing the number of waste packages, in turn reducing the number of handling operations and associated dose uptake to workers. There are a number of candidates that have been proposed as suitable additives for Pu-containing ceramics. For example, Nadykto and Timofeeva (2001) [10] estimated that in the case of thermal neutrons, the additive:Pu nuclei ratio to avoid a nuclear chain reaction for B, Gd, Cd, Sm, and Eu was 1.6, 0.023, 0.45, 0.2, and 0.25, respectively. The incorporation of the additive in the wasteform could be accomplished in a number of ways.

There is not currently a consensus regarding the most effective means or efficacy of adding the neutron absorbing additive to a wasteform on an industrial scale; however, the addition would likely have to be constrained within the batch formulation and powder preparation stage. The engineering unit operations for a HIP-based strategy are illustrated in Figure 1. Within this process, there are two options that may be suitable, as they do not require additional handling or unit operations associated with them and make use of existing pre-treatment parameters that are already essential to the HIP program. These are as follows:

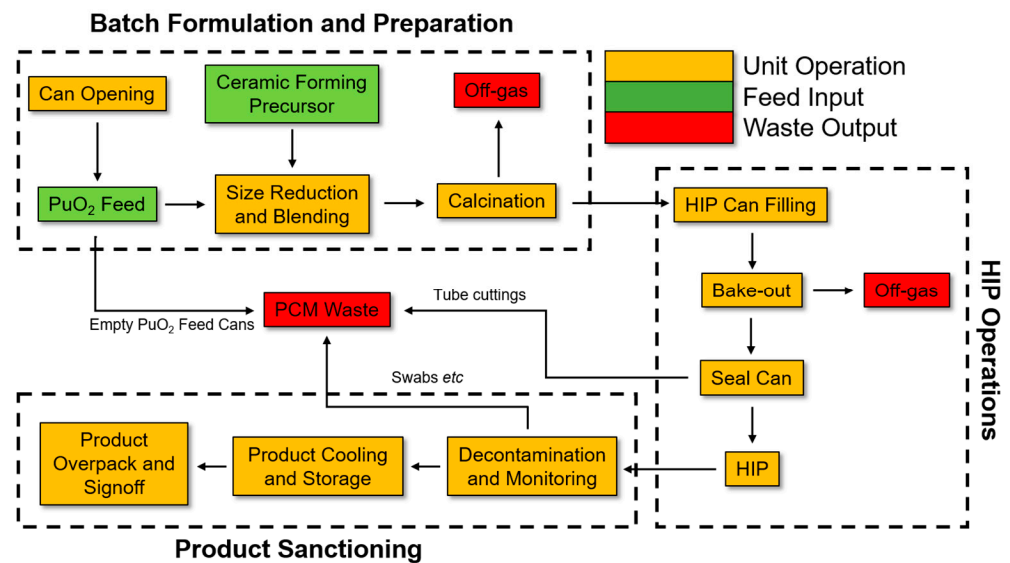


Figure 1. Simplified overview of primary unit operations that may be associated with HIP immobilization strategy for Pu.

(1) **Wasteform tailoring**—In this option, the ceramic composition is tailored such that a pre-determined quantity of additive can be incorporated within the structure without

disrupting the phase assemblage or resulting in the formation of deleterious secondary phases. The addition of trivalent species, e.g.,  $Gd^{3+}$ , would necessitate tailoring of the nominal composition to account for excess/deficiency in charge units across the structure. An appropriate amount of additive is incorporated in the oxide form directly to the milling stage as a function of the total wt. % of the waste package (e.g., direct addition of 2 wt. %  $Gd_2O_3$ ). The inactive component of the ceramic precursor would require a pre-treatment stage consisting of homogenisation and calcination prior to the addition of the  $PuO_2$ . This would be followed by a secondary milling stage to ensure complete oxide dispersion and size reduction to assist the sintering process. In this case, the additive source may be introduced alongside the  $PuO_2$  at the secondary milling stage. Compositions developed for this route are discussed in detail in the following section.

**(2) Additive encapsulation**—Alternatively, the additive may be incorporated into the waste microstructure as discrete particles via encapsulation. This may be more applicable for additives such as boron which, due to its small ionic radius, may not be effectively accommodated in a solid solution with the host ceramic phase. In this instance, a compound such as  $B_4C$  may be encapsulated at intergranular spaces and triple points, forming discrete particulates within the microstructure. An additive such as  $B_4C$  has an extremely high Vickers hardness; therefore, to attain homogeneous distribution throughout the microstructure, the feed would have to be comprised of a fine powder that can easily be dispersed, or the process would need to take advantage of an aggressive milling stage.

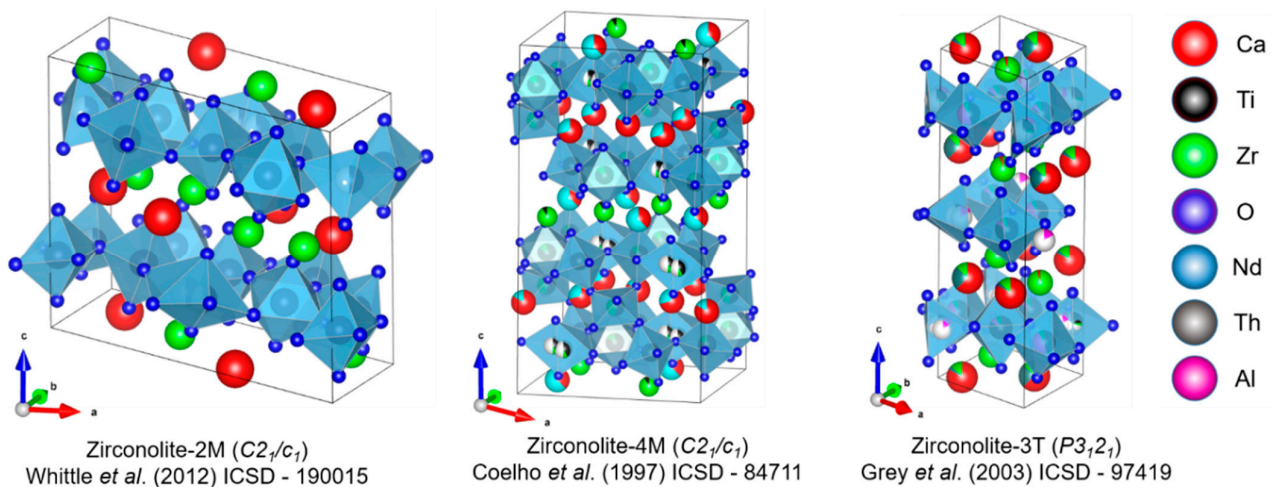
### 3. Review of Zirconolite Chemistry and Evaluation of Potential Solid Solution Regimes for Pu Immobilisation

Technical efforts in the UK to identify a suitable wasteform for separated Pu have considered single and multiphase ceramics based on derivatives of the titanate pyrochlore-rich wasteform developed for the US DoE Plutonium Immobilisation Project (PIP) and vitrified wasteforms based on variations of borosilicate glass (e.g., lanthanide borosilicate (LaBS), alkali tin silicates (ATS)). Encapsulation and dilution of  $PuO_2$  in cementitious wasteforms is still also considered a technically feasible option on the basis of GDF compliance and simplicity [11,12]. Following several years of inactive trials and deselection, occurring primarily in collaboration between the National Nuclear Laboratory (NNL) and the Immobilisation Science Laboratory (ISL), wasteforms favouring zirconolite as the host Pu phase have been selected as the preferred candidate and, as such, a program of systematic active and inactive trials aiming to maximise Pu wasteloading through tailoring a selection of baseline compositions and further probe the underlying crystallography of the zirconolite structure is underway [13–16].

Zirconolite (nominally  $CaZrTi_2O_7$ ) is a rare accessory mineral that is present in a wide variety of terrestrial localities, demonstrating an affinity for an extensive solid solution with, but not limited to,  $Mg^{2+}$ ,  $Al^{3+}$ ,  $Fe^{2+/3+}$ ,  $Nb^{5+}$ ,  $Ta^{5+}$ ,  $Pb^{2+}$ ,  $U^{4+}$ ,  $Th^{4+}$ ,  $Ce^{3+/4+}$ ,  $Pr^{3+}$ ,  $Gd^{3+}$ ,  $Sm^{3+}$ , and  $Nd^{3+}$  [17]. The ability of zirconolite to incorporate such a wide variety of species, with five distinct cation receptor sites ( $Ca^{VIII}$ ,  $Zr^{VII}$ ,  $Ti(I)^{VI}$ ,  $Ti(II)^V$ , and  $Ti(III)^{VI}$ ), makes it an ideal candidate for co-immobilisation of separated Pu, lower valence charge compensator cations such as  $Mg^{2+}$  and  $Al^{3+}$ , and an appropriate amount of a suitable neutron absorbing additive (e.g.,  $Gd^{3+}$ ) to suppress criticality. Zirconolite (the archetype of which is the 2M structure) has been recognised by NDA as a candidate ceramic phase for Pu and, as such, has commissioned a body of work aiming to systematically validate potential solid solutions using a combination of surrogates (Ce/U/Th). It should also be noted that glass–ceramic wasteforms based on the  $CaZrTi_2O_7$ – $Na_2Al_2Si_6O_{16}$  system have also been proposed for Cl-contaminated Pu residues [18–22].

The idealised zirconolite parent structure crystallises in monoclinic symmetry (space group  $C2/c$ ) and comprises two distinct cation layers, where  $Ca^{2+}$  and  $Zr^{4+}$  polyhedra are 8 and 7-fold coordinated to oxygen (i.e.,  $CaO_8$  and  $ZrO_7$ ), respectively, and are interspaced by layers of Ti–O polyhedra arranged in a topology closely related to that of hexagonal tungsten bronze (HTB) (Figure 2). In this theoretical description, Ti occupies three distinct

sites within the HTB layer, two of which are 6-fold coordinated to oxygen forming six-membered rings. The remaining Ti is statistically distributed in the centre of these rings, in a 50% occupied 5-fold coordinated trigonal bi-pyramidal site. In the zirconolite unit cell, planes of Ca/Zr polyhedra and HTB layers are stacked 1:1 along parallel to [001], with HTB positioned at  $z = 0.25$  and  $z = 0.75$  related by a  $180^\circ$  rotation, offset along [130]; subsequent modules at  $z = 0.75$  and  $z = 1.25$  are offset in the [1–30] direction [23]. This structure is referred to as zirconolite-2M, with reference to the two-layer repeating lamellar structure comprising the monoclinic unit cell. A more comprehensive description is provided by Gatehouse et al. in which the zirconolite-2M structure remains stable over the solid solution  $\text{CaZr}_x\text{Ti}_{3-x}\text{O}_7$  constrained by  $0.80 < x < 1.35$  [24], with mixed occupancy of Zr-Ti-O polyhedra also determined to be a function of temperature, between 1350 and 1500 °C [25].



**Figure 2.** Crystal structures of zirconolite-2M, 4M, and 3T polytypes (images reproduced from the Vesta software using Crystallography Information Files (CIF) from Whittle et al. [26], Coelho et al. [25], and Grey et al. [27]).

Zirconolite structured materials are typically produced via high temperature sintering from a mixture of oxide precursors, and although sintering regimes are typically  $>1300^\circ\text{C}$  and  $>10$  h, there does not seem to be any clear correlation between processing conditions and lattice dimensions. A selection of reported lattice parameters of  $\text{CaZrTi}_2\text{O}_7$  and corresponding processing conditions are listed in Table 2. Due to the flexibility of the zirconolite structure, a variety of potential solid solution regimes for Pu immobilisation have been identified, both from a survey of available literature and hypothesised based on feasible (charge balanced) substitutions (Table 3). Depending on the substitution mechanism and prevailing oxygen fugacity during the sintering process, zirconolite may crystallise in several distinct polytypes, characterised in principle by variation in the stacking sequence of Ca/Zr and HTB layers. For example, the incorporation of  $>0.15$  f.u. of  $\text{M}^{4+}$  (Ce/U/Pu) promotes the formation of the zirconolite-4M polytype, described by Coelho et al. as a four layer intergrowth of zirconolite-2M and pyrochlore-type modules, resulting in a doubling of the zirconolite unit cell along the  $c^*$  axis [23]. Zirconolite-4M may also form as a result of near equimolar accommodation of trivalent species within the  $\text{Ca}^{2+}$  and  $\text{Zr}^{4+}$  sites, i.e., zirconolite-4M is essentially an intermediate phase between  $\text{CaZrTi}_2\text{O}_7$  (zirconolite) and  $\text{REE}^{3+}_2\text{Ti}_2\text{O}_7$  (pyrochlore) end-members [28]. Heavily disordered substitution regimes, such as targeting accommodation of  $\text{M}^{3+}$  and  $\text{M}^{4+}$  species between  $\text{Ca}^{2+}$  and  $\text{Ti}^{4+}$  sites, may also result in the formation of other polytypes, such as zirconolite-3T (trigonal unit cell, space group: Pbnm) or zirconolite-3O (orthorhombic unit cell, space group: Acam). For a comprehensive discussion of zirconolite polytypism, the reader is referred to White (1984) [29]. Generally, substitution schemes for the accommodation of  $\text{Pu}^{4+}$  within the  $\text{Ca}^{2+}$

site are facilitated by a charge-coupled substitution, whereby a lower valence cation such as  $M = \text{Fe}^{3+}/\text{Al}^{3+}$  is accommodated within the  $\text{Ti}^{4+}$  site [30]. As stated, a simple isovalent substitution for  $\text{Zr}^{4+}$  has also been demonstrated and may allow greater wasteloading; however, this results in a phase assemblage dominated by zirconolite-4M, using both Ce/U surrogates and Pu [16,31,32].

**Table 2.** Reported unit cell parameters for zirconolite polytypes.

Structure	Symmetry	a (Å)	b (Å)	c (Å)	$\beta$ (°)	Space Group	Origin	Ref.
2M	Monoclinic	12.4435	7.2735	11.3748	10.563	C2/c	Synthetic	[14]
4M	Monoclinic	12.553	7.248	23.081	84.799	C2/c	Synthetic	[23]
3T	Trigonal	7.228	-	16.805	-	P3 <sub>1</sub> 21	Synthetic	[27]
3O	Orthorhombic	10.31	14.48	7.41	-	Acam	Natural	[33]

Several of these solid solution regimes would necessitate Pu to be present as  $\text{Pu}^{3+}$ . This may not prove feasible, as it may involve the conversion of the  $\text{PuO}_2$  into  $\text{Pu}_2\text{O}_3$  prior to ceramic processing, incurring additional feed preparation steps in the process outlined in Figure 1. The conversion of  $\text{Pu}^{3+}$  within the wastefrom, e.g., through the use of strongly reducing conditions during sintering, will also likely drive the formation of deleterious accessory phases such as  $\text{CaTiO}_3$ . Furthermore,  $\text{CaTiO}_3$  has demonstrated an affinity for trivalent  $\text{M}^{3+}$  incorporation and has demonstrated markedly lower chemical durability than the zirconolite phase, and thus may significantly reduce overall wastefrom performance through uptake and subsequent release of  $\text{Pu}^{3+}$  from  $\text{CaTiO}_3$  [34–36].

**Table 3.** List of feasible solid solution regimes for Pu in zirconolite and summary of expected polytype fields.

Solid Solution Regime	Comments	References
$\text{CaZr}_{1-x}\text{Pu}^{4+}_x\text{Ti}_2\text{O}_7$	This system exhibits a polymorphic transition to 4M in the interval $0.10 \leq x \leq 0.20$ . This has been demonstrated in several key investigations using Ce/U/Pu. A mixture of zirconolite-2M and zirconolite-4M could be expected in the compositional interval $0.20 \leq x \leq 0.40$ . Zirconolite-4M is rarely stabilised as a single phase and exists over a narrow compositional range of $0.40 \leq x \leq 0.50$ , accompanied by pyrochlore. Transformation to the pyrochlore structure is common around $x = 0.60$ , with solid solution limits of $\text{CeO}_2$ defined at $x \sim 0.80$ (this could also be expected for $\text{UO}_2$ and $\text{PuO}_2$ ).	[13,16,31,32, 37,38]
$\text{Ca}_{1-x}\text{Pu}^{4+}_x\text{ZrTi}_{2-2x}\text{M}^{3+}_{2x}\text{O}_7$	This solid solution appears to be the most common method of substitution, by which Ce/U/Pu are substituted within the $\text{Ca}^{2+}$ site, and charge balance provided by a small lower valence cation exchanged for $\text{Ti}^{4+}$ (typically $\text{Al}^{3+}$ , $\text{Fe}^{3+}$ , $\text{Cr}^{3+}$ , or $\text{Mg}^{2+}$ . Note: in the case of $\text{Mg}^{2+}$ , the nominal composition becomes $\text{Ca}_{1-x}\text{Pu}_x\text{ZrTi}_{2-x}\text{Mg}_x\text{O}_7$ ). Zirconolite-2M is typically formed near single phase in the compositional range of $0.00 \leq x \leq 0.20$ , with secondary perovskite commonly observed for Ce surrogate compositions due to partial $\text{Ce}^{3+}$ speciation. At elevated wasteloading (i.e., $x = 0.35$ ), free oxides may be observed. Furthermore, the dominant zirconolite polytype at enhanced wasteloading appears to be controlled by choice of $\text{M}^{3+}$ species. For example, $\text{Ca}_{0.65}\text{Ce}_{0.35}\text{ZrTi}_{1.30}\text{Cr}_{0.70}\text{O}_7$ was reported as zirconolite-2M by Blackburn et al., yet the corresponding $\text{Ca}_{0.65}\text{Pu}_{0.35}\text{ZrTi}_{1.30}\text{Fe}_{0.70}\text{O}_7$ composition was reported as zirconolite-3T by Gilbert et al. This may be attributed to the different redox and/or site occupancy behaviour between Ce/Pu and/or ionic radii of Cr/Fe.	[14,31,39–44]
$\text{Ca}_{1-x}\text{Pu}^{3+}_x\text{ZrTi}_{2-x}\text{M}^{3+}_x\text{O}_7$	Systematic studies in the $\text{Ca}_{1-x}\text{Ln}_x\text{ZrTi}_{2-x}(\text{Al,Fe})_x\text{O}_7$ system have been reported for Ln = La, Nd, Gd, Ho, and Yb, which may act as $\text{Pu}^{3+}$ surrogates. The phase evolution of these ceramics is seemingly dictated by ionic radii, with increased solubility for smaller cations such as $\text{Gd}^{3+}$ . Polytype transformation to zirconolite-3O could be expected at elevated substitution ( $x = 0.60$ ) for some $\text{Ln}^{3+}$ surrogates (e.g., Nd) but 2M phase stable up to $x = 0.80$ for others, e.g., $\text{Gd}^{3+}$ .	[45–48]
$\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Pu}^{3+}_{2x}\text{Ti}_2\text{O}_7$	Equimolar substitution of $\text{Pu}^{3+}$ between $\text{Ca}^{2+}$ and $\text{Zr}^{4+}$ could be expected to produce a phase transformation to the pyrochlore structure with elevated Pu content $x \geq 0.60$ via formation of intermediate zirconolite-4M phase. This has been demonstrated for the corresponding $\text{Nd}^{3+}$ , $\text{Y}^{3+}$ , $\text{Sm}^{3+}$ , and $\text{Gd}^{3+}$ solid solutions.	[49–52]
$\text{Ca}_{1-x}\text{Pu}^{4+}_{2x}\text{Zr}_{1-x}\text{Pu}^{4+}_x\text{Ti}_{2-4x}\text{M}^{3+}_{4x}\text{O}_7$	This solid solution mechanism would see $\text{Pu}^{4+}$ cations co-partitioned between $\text{Ca}^{2+}$ and $\text{Zr}^{4+}$ sites. Such targeted solid solutions have been observed to predominantly yield zirconolite-3T.	[53]

## 4. Solid Solution Limits of Potential Neutron Absorbing Additives in Zirconolite

### 4.1. Candidate Neutron Absorbing Additives

Initial screening identified a selection of suitable neutron absorbing additives species (Table 4). Gd, Hf, Sm, Cd, In, and B were identified as suitable candidates on the basis of accessible oxidation states, commercial availability, cost per gram of component oxides, and thermal neutron absorption cross section. The cost index, as outlined in Table 4, is taken from the current Merck reagent catalogue (accurate as of November 2022) and is, therefore, not expected to be representative of the price for large quantities of constituent reagent materials. Nevertheless, these listings provide an order of magnitude comparison. Although the wastefrom design philosophy stipulates that the down-selection of neutron absorbing additives should not be dictated entirely by cost, it is clear from Table 4 that the unit price of specific oxides can vary by three orders of magnitude. Although the elements listed are not an exhaustive list of all potential additives, an initial filter was applied such that any species with a cost index approximately 100 times greater than Gd were excluded.

**Table 4.** Data for evaluating candidate neutron absorbing additives.

Element	Cost of Component Oxide (£/g)	Merck Product Code	Purity	Thermal Neutron Cross Section ( $\sigma_a$ /barns)
Gd	2.76	278513	99.9% trace metals basis	$4.90 \times 10^4$
Hf	151.00	203394	$\geq 99.95\%$	104
Sm	4.20	228672	99.9% trace metals basis	$5.92 \times 10^3$
Cd	7.00	202894	$\geq 99.99\%$ trace metals basis	$2.45 \times 10^4$
In	7.50	289418	99.99% trace metals basis	194
B *	1.76	378119	98%	767

\* All elements are listed as oxides except for B, which is anticipated to be  $B_4C$ ; prices are accurate as of November 2022.

### 4.2. Gadolinium Doped Zirconolite

Gadolinium incorporation within zirconolite has been extensively studied since  $Gd^{3+}$  possesses physical characteristics also applicable to that of an effective minor actinide simulant. Nevertheless,  $Gd^{3+}$  may be accommodated in zirconolite to act as a neutron absorbing additive. Gadolinium has been included in a series of polycrystalline phase assemblages, such as the SYNROC wastefrom, for criticality control. However, these studies did not intend to incorporate  $Gd^{3+}$  within specific lattice sites within the zirconolite parent structure; hence, a meaningful discussion of the solid solution limits is not always possible. For example, a zirconolite-rich SYNROC composition was synthesised by HIP at 1280 °C in the work of Li et al., with 8.96 wt. %  $Gd_2O_3$  added for criticality control; however, the partitioning of  $Gd^{3+}$  between the obtained zirconolite, hollandite, pyrochlore, and rutile phases was not discussed [54]. The partitioning of  $Ln^{3+}$  species within a series of SYNROC compositions was investigated by Lumpkin et al., with the conclusion that the partitioning of these elements within the various crystalline phases was largely controlled by ionic radii [55]. Consequently, it was inferred that  $Ln^{3+}$  cations such as  $Gd^{3+}$  demonstrated an affinity for substitution within the large  $Ca^{2+}$  site in the zirconolite structure.

Two zirconolite-rich SYNROC wastefroms were HIPed by Zhang et al. using both U/Th surrogates (targeting 90% zirconolite and 10% hollandite) and Pu (targeting 80% zirconolite, 10% hollandite, and 10% rutile), with Gd and Gd + Hf added to each sample for criticality control, respectively [56]. In both instances, the zirconolite phase was observed to overwhelmingly accommodate both neutron absorbing additives alongside U/Th and Pu. Leaching trials in deionised water at 90 °C demonstrated near congruent release of Gd/Hf with Pu from the wastefrom, with normalised release rates  $< 10^{-5} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$  after 35 d for Hf and Pu. It was concluded that the release of Pu from the wastefrom would likely be accompanied by the release of Hf/Gd, mitigating any potential localised criticality.

The solid solution limits of Gd are more easily discussed when a series of specific solid solutions are targeted. Zhang et al. fabricated the  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Gd}_{2x}\text{Ti}_2\text{O}_7$  ( $0.00 \leq x \leq 1.00$ ) solid solution by conventional sintering at 1400 °C for 48 h using oxide precursors [52]. Zirconolite-2M was produced as the dominant phase in the range of  $0.00 \leq x \leq 0.20$ , with a mixture of zirconolite-2M and zirconolite-4M obtained for  $0.20 \leq x \leq 0.50$ . A cubic pyrochlore phase was detected via powder XRD for  $x \geq 0.40$ , gradually progressing towards a single phase at  $x = 0.60$ . A 7-day ASTM PCT leaching test [57] was performed for compositions corresponding to  $x = 0.20$  (mixture of zirconolite-2M and 4M) and  $x = 0.40$  (mixture of zirconolite-2M, 4M, and pyrochlore) using deionised  $\text{H}_2\text{O}$  as the leachate. The normalised leach rate of Gd from each sample was exceptionally low, measured to be  $4.96 \times 10^{-7} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$  and  $1.78 \times 10^{-7} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$ , respectively.

A systematic investigation was performed by Ma et al. characterising the incorporation of a variety of  $\text{Ln}^{3+}$  species in the zirconolite system  $\text{Ca}_{1-x}\text{Ln}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$  ( $\text{Ln} = \text{La}, \text{Nd}, \text{Gd}, \text{Ho}, \text{and Yb}, 0.10 \leq x \leq 1.00$ ) by sintering oxide precursors at 1400 °C for 100 h [48]. The prevailing phase assemblage and dominant zirconolite polytype were controlled by the choice of  $\text{Ln}^{3+}$  species, with  $\text{La}^{3+}$  failing to form a complete solid solution past the limit of  $x = 0.20$ . When targeting the  $\text{Ca}_{1-x}\text{Gd}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$  system, zirconolite-2M was maintained extensively over the range of  $0.00 \leq x \leq 0.80$ , after which minor  $\text{ZrO}_2$  became prominent as a secondary phase. Interestingly, the replication of this series with  $\text{Nd}^{3+}$  produced the zirconolite-3O phase at  $x \geq 0.70$ , implying that  $\text{Gd}^{3+}$  does not distort the zirconolite structure as much as closely related  $\text{Ln}^{3+}$  species, likely due to size constraints. Interestingly, the  $\text{Ca}_{1-x}\text{Gd}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$  ( $0.00 \leq x \leq 1.00$ ) solid solution synthesised at 1400 °C (5 h) by Stefanovsky et al. produced different phase fields than those reported by Ma et al. [46]. Whilst the microstructure was comprised of a single phase zirconolite over the range of  $0.00 \leq x \leq 0.90$ , at which point minor Gd-rich  $\text{ZrO}_2$  was observed by SEM, electron diffraction data were consistent with the zirconolite phase transforming to 3T or 6T symmetry in the range of  $0.50 \leq x \leq 0.90$ . When targeting  $x = 1.00$ , i.e., nominal composition  $\text{GdZrTiAlO}_7$ , electron diffraction analysis was consistent with zirconolite-3O. The solid solution limits of Gd within the following formulations were probed by Vance et al.:  $\text{Ca}_{1-x}\text{Gd}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$ ,  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Gd}_{2x}\text{Ti}_2\text{O}_7$ , and  $\text{CaZr}_{1-x}\text{Gd}_x\text{Ti}_{2-x}\text{Nb}_x\text{O}_7$  via by conventional sintering at 1450 °C for 20 h [58]. When targeting  $\text{Ca}_{1-x}\text{Gd}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$ , the phase evolution was largely similar to related solid solutions, e.g., Ma et al. [48], with the formation of zirconolite-3O observed at  $x = 0.70$ . A transformation to the pyrochlore structure was identified at  $x = 0.50$  for  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Gd}_{2x}\text{Ti}_2\text{O}_7$ , similar to the phase fields observed by Zhang et al. for Gd [52]. Identical phase fields were also formed for  $\text{CaZr}_{1-x}\text{Gd}_x\text{Ti}_{2-x}\text{Nb}_x\text{O}_7$ , by which the pyrochlore phase was produced above  $x = 0.50$ .

The  $\text{Ca}_{1-x}\text{Ln}_x\text{ZrTi}_{2-x}\text{Fe}_x\text{O}_7$  ( $\text{Ln} = \text{La}, \text{Nd}, \text{Gd}, \text{Ho}, \text{and Yb}, 0.10 \leq x \leq 1.00$ ) solid solution fabricated by Ji et al. [47] complements the data reported by Ma et al. [48], allowing a reasonable comparison to be made between  $\text{Ca}_{1-x}\text{Gd}_x\text{ZrTi}_{2-x}(\text{Al},\text{Fe})_x\text{O}_7$ . The solid solution limits for the zirconolite-2M phase were largely similar for compositions charge balanced with Al, with the 2M phase stabilised over the range  $0.10 \leq x \leq 0.80$  for Fe-substituted compositions. Ji et al. concluded that zirconolite-3O was the dominant polytype over the compositional range  $0.90 \leq x \leq 1.00$ , implying that the end member  $\text{GdZrTiFeO}_7$  may be a near single phase zirconolite-3O; however, no electron diffraction data were provided.

Phase relations in the pseudo-binary system  $(1-x)\text{CaZrTi}_2\text{O}_7 - (x)\text{GdAlO}_3$  ( $x = 0.25, 0.50$  and  $0.75$ ) were investigated by Mikhailenko et al. via a cold press and sinter method at 1400 °C and 1500 °C [59]. Zirconolite was present as the major crystalline phase when targeting  $x = 0.25$ , with ~70 wt. % of the phase assemblage composed of zirconolite with calculated stoichiometry  $\text{Ca}_{0.36}\text{Gd}_{0.60}\text{Zr}_{1.12}\text{Ti}_{1.40}\text{Al}_{0.47}\text{O}_7$ ; however, Gd was still concentrated in the secondary perovskite phase. Increasing the ( $x$ ) parameter to 0.75 produced a phase assemblage comprised of 45 wt. % zirconolite, 45 wt. % perovskite, and 10% hibonite ( $\text{CaAl}_2\text{O}_9$ ). Interestingly, this pseudo-binary system produced a greater relative fraction of zirconolite than the corresponding Sm system, also fabricated by Mikhailenko

et al. when targeting  $x = 0.75$  [60]. Leaching of these ceramics was performed according to the Australian Powder No. 1 (P1) test, with distilled water used as the leachate (0.2 g of material; 20 mL distilled water; static test at 95 °C for 6 h before solution replenished; continued for 10 d). The normalised leach rate of Gd was  $\sim 10^{-4} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$  over the 10 d period when the nominal ( $x$ ) parameter was 0.25; the release rate of Gd increased by an order of magnitude of  $x = 0.75$ , attributed to excess perovskite content.

In summary, the evidence that  $\text{Gd}^{3+}$  readily substitutes within the  $\text{Ca}^{2+}$  and  $\text{Zr}^{4+}$  sites in the zirconolite structure is extensive, with a number of corroborating studies reporting high wasteloading within the  $\text{Ca}^{2+}$  site. The possibility of equimolar substitution of  $\text{Gd}^{3+}$  between  $\text{Ca}^{2+}$  and  $\text{Zr}^{4+}$  sites is also demonstrated, although this was reported to produce the zirconolite-4M phase at moderate levels of substitution. Considering the relatively low ratio of additive:Pu atoms necessary to reduce  $K_{\text{eff}} < 1$  (refer to back to Table 4), the incorporation of Gd within zirconolite should not be expected to destabilise the zirconolite phase, or indeed to produce a polytypical transformation, given that roughly 1 atom of Gd is necessary to offset 100 Pu atoms. Considering the dilution level of Gd needed to satisfy these neutronic requirements, the most effective means of Gd incorporation would likely be achieved by co-incorporation of  $\text{Gd}^{3+}$  across  $\text{Ca}^{2+}$  and  $\text{Zr}^{4+}$  sites, negating the need to alter the preferred formulation to accommodate for charge imbalance. Moreover, the cost index of Gd is attractive relative to Hf and In.

#### 4.3. Hafnium Doped Zirconolite

From a solid solution perspective, Hf may be the most suitable additive, as  $\text{Hf}^{4+}$  readily forms a complete solid solution with  $\text{Zr}^{4+}$  in the zirconolite structure on the basis of near-identical ionic radii (0.78 Å vs. 0.76 Å for  $\text{Zr}^{4+}$  and  $\text{Hf}^{4+}$  in 7-fold coordination, respectively) [61]. Furthermore, the interchange of these species results in a negligible impact on the overall structural properties. Three compositions corresponding to  $\text{CaZrTi}_2\text{O}_7$ ,  $\text{CaHf}_{0.5}\text{Zr}_{0.5}\text{Ti}_2\text{O}_7$ , and  $\text{CaHfTi}_2\text{O}_7$  were fabricated by Vance et al. with a sintering temperature of 1450 °C maintained for 20 h [58]. There was negligible variation in the unit cell parameters of the zirconolite phase (Table 5), suggesting that in any optimised ceramic formulation for Pu,  $\text{Zr}^{4+}$  may be interchangeable with a requisite mol. % of  $\text{Hf}^{4+}$  to satisfy criticality requirements.

**Table 5.** Unit cell parameters for Hf-doped zirconolite (reproduced from Vance et al. [58]).

Composition	Unit Cell Parameters			
	a (Å)	b (Å)	c (Å)	$\beta$ (°)
$\text{CaZrTi}_2\text{O}_7$	12.447	7.272	11.384	100.54
$\text{CaHf}_{0.5}\text{Zr}_{0.5}\text{Ti}_2\text{O}_7$	12.434	7.265	11.350	100.58
$\text{CaHfTi}_2\text{O}_7$	12.420	7.262	11.343	100.57

Complete replacement of Zr with Hf in the zirconolite structure was demonstrated by Putnam et al., with single phase  $\text{CaHfTi}_2\text{O}_7$  synthesised from component oxides at 1300 °C, with intermittent regrinding to ensure phase homogeneity [62]. Furthermore,  $\text{CaHfTi}_2\text{O}_7$  was determined to be more thermodynamically stable than stoichiometric  $\text{CaZrTi}_2\text{O}_7$ ; hence, partial replacement of Zr with Hf may be favourable from a wasteform perspective whilst satisfying criticality requirements. Hyatt et al. formulated a series of zirconolites with nominal stoichiometry  $(\text{Ca}_{1-x/2}\text{Gd}_{x/2})(\text{Zr}_{1-5x/2}\text{Ce}_x\text{Hf}_x\text{Gd}_{x/2})\text{Ti}_2\text{O}_7$ , with Ce acting as a structural simulant for Pu and  $\text{Hf}^{4+}$  included as a neutron absorber [63]. The following solid solution limits were systematically defined: zirconolite-2M ( $0.00 \leq x \leq 0.15$ ); mixture of zirconolite-2M and zirconolite-4M ( $0.15 \leq x \leq 0.25$ ); mixture of zirconolite-4M and pyrochlore ( $0.25 \leq x \leq 0.30$ ). It is assumed that the equimolar substitution of  $\text{Gd}^{3+}$  across  $\text{Ca}^{2+}$  and  $\text{Zr}^{4+}$  was responsible for the formation of zirconolite-4M and pyrochlore, as very similar phase fields were observed for  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Gd}_{2x}\text{Ti}_2\text{O}_7$ . This implies that the partial occupation of  $\text{Hf}^{4+}$  within the  $\text{Zr}^{4+}$  site does not disrupt the zirconolite-2M to zirconolite-4M phase transformation to any meaningful extent.

The  $\text{CaZr}_{1-x}\text{Hf}_x\text{Ti}_2\text{O}_7$  ( $0.00 \leq x \leq 1.00$ ) solid solution was fabricated by Zhang et al. using combustion synthesis combined with hot pressing for densification [64]. It was determined that the zirconolite-2M polytype was maintained over the phase evolution, with complete digestion of  $\text{HfO}_2$  within a highly densified microstructure at each compositional interval. The chemical durability of  $\text{CaZr}_{0.4}\text{Hf}_{0.6}\text{Ti}_2\text{O}_7$  was investigated by the ASTM MCC-1 standard leaching test [65] over a 42 d period. The normalised leach rate after the 42 d period was significantly lower for Hf than Ca, corresponding to  $1.11 \times 10^{-8} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$  and  $0.25 \times 10^{-2} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$ , respectively. As there was no Pu surrogate included in the targeted phase composition, no comment can be made as to whether there may be a congruent release of Pu/Hf from the wasteform. Complementary data were published by Zhang et al. utilising the same combustion synthesis route to produce the  $\text{Ca}_{1-x}\text{Hf}_{1-x}\text{Gd}_x\text{Ti}_2\text{O}_7$  solid solution, with  $\text{Gd}^{3+}$  included as a surrogate for minor actinide species such as  $\text{Pu}^{3+}$  (although, as discussed in the previous segment, Gd may also be included as an effective neutron absorbing additive) [66]. A phase transition from  $\text{CaHfTi}_2\text{O}_7$  (zirconolite-2M) to  $\text{Gd}_2\text{Ti}_2\text{O}_7$  (cubic pyrochlore), with a solid solution limit of  $\text{Gd}_2\text{O}_3$  defined at  $x = 0.80$ , was observed. Interestingly, the authors did not report the formation of an intermediate zirconolite-4M phase, as was observed in closely related solid solution regimes (e.g.,  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Gd}_x\text{Ti}_2\text{O}_7$  synthesised by Zhang et al. [52] or  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Nd}_x\text{Ti}_2\text{O}_7$  synthesised by Coelho et al. [23]). After a 42 d leaching period (under MCC-1 conditions), the release of Gd and Hf was measured from the sample corresponding to nominal composition  $x = 0.60$  (i.e.,  $\text{Ca}_{0.4}\text{Hf}_{0.4}\text{Gd}_{1.2}\text{Ti}_2\text{O}_7$ ). The release rates of Gd and Hf were measured to be  $4.72 \times 10^{-7} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$  and  $1.59 \times 10^{-8} \text{ g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$ , respectively, demonstrating that a similar cumulative dissolution trend may be preserved. However, an inspection of the powder XRD data revealed that whilst exceptionally low leach rates were observed, the composition  $x = 0.6$  was consistent with the sample adopting the pyrochlore structure; hence, any meaningful discussion of the durability of a zirconolite-structured material with a corresponding Gd/Hf concentration is not possible (since both materials were not single phase).

The incorporation of  $\text{Hf}^{4+}$  in  $\text{Nd}^{3+}$ -zirconolite was investigated by Caurant et al., targeting the solid solution  $\text{Ca}_{1-x}\text{Nd}_x\text{HfTi}_{2-x}\text{Al}_x\text{O}_7$ , with  $\text{Nd}^{3+}$  acting as a structural simulant for  $\text{Pu}^{3+}$  [67]. Compositions targeting  $x = 0.01$  and  $x = 0.20$  were fabricated by solid-state reaction at  $1460^\circ\text{C}$  between component oxides. It was confirmed that zirconolite remained in the 2M polytype for each targeted composition, alongside minor  $\text{CaTiO}_3$  and Hf-Ti-O rich phases, accounting for a negligible portion of the phase assemblage. Mixed occupancy of  $\text{Hf}^{4+}$  and  $\text{Ti}^{4+}$  ions were confirmed, consistent with data reported for  $\text{CaZr}_x\text{Ti}_{3-x}\text{O}_7$ , with  $\text{Nd}^{3+}$  exclusively entering the  $\text{Ca}^{2+}$  site, in accordance with the target composition. Loiseau et al. prepared the corresponding Zr solid solution ( $\text{Ca}_{1-x}\text{Nd}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$ ,  $x \leq 0.8$ ), allowing useful comparison between the Zr/Hf series [68]. In agreement with Caurant et al., the zirconolite-2M structure was stable in the corresponding compositional range ( $x \leq 0.20$ ). Furthermore, zirconolite occupied the 2M structure until  $x = 0.7$ . When targeting a nominal Nd concentration of  $x = 0.8$ , the zirconolite-3O polytype was formed with orthorhombic symmetry (space group  $\text{Acam}$ ). Based on these observations, it would be expected that the Hf equivalent composition would also occupy 3O symmetry at elevated  $\text{Nd}^{3+}$  content. The solid solution limit for which zirconolite formed the 3O polytype was also consistent with the work of Ma et al. [48] discussed previously.

Tetravalent  $\text{Hf}^{4+}$  was utilised in place of  $\text{Zr}^{4+}$  in the work of Begg et al., with the aim to investigate the substitution of Pu for  $\text{Ca}^{2+}$  without the addition of a charge compensation cation, targeting  $\text{Ca}_{0.9}\text{Pu}_{0.1}\text{HfTi}_2\text{O}_7$ . A slight excess of Pu was reported such that the actual concentration was  $<0.1$  f.u. [44]. Sintering in the air at  $1400^\circ\text{C}$  produced a near-single phase zirconolite alongside  $\sim 5$  wt. % pyrochlore with the composition  $\text{Ca}_{0.92}\text{Pu}_{0.56}\text{Hf}_{0.75}\text{Ti}_{1.74}\text{O}_7$ , and minor  $\text{TiO}_2/\text{HfTiO}_4$ . Annealing the sample in 3.5%  $\text{H}_2/\text{N}_2$  was sufficient to reduce 80%  $\text{Pu}^{3+}$ ; however, no disruption to the phase assemblage was observed, demonstrating the ability of the zirconolite phase to accommodate for changes in Pu valence, most likely compensated by Hf/Ti vacancies.

In summary, there appears to be an inherent advantage in incorporating Hf in zirconolite insofar as the ionic radius of Hf is nearly identical to that of Zr ( $\text{Hf}^{\text{VII}} = 0.76 \text{ \AA}$ ;  $\text{Zr}^{\text{VII}} = 0.78 \text{ \AA}$ ). In practice, the replacement of  $\text{Zr}^{4+}$  with  $\text{Hf}^{4+}$  should not impact zirconolite phase formation; moreover, the composition should not need tailoring to account for excess charge unit considerations. The primary drawback to the use of  $\text{HfO}_2$  is the high cost relative to Gd/Sm/B; hence, the complete replacement of Zr with Hf, i.e.,  $(\text{Ca}_{0.80}\text{Pu}_{0.20})(\text{Hf}_{0.90}\text{Pu}_{0.10})\text{Ti}_{1.60}\text{Al}_{0.40}\text{O}_7$ , would likely incur a significant cost to head-end operations. Nevertheless, the partial replacement of Zr by Hf, with the view to only satisfy the additive:Pu ratio, remains an attractive prospect, and it is reasonable to assume such substitution would not influence Pu partitioning or zirconolite phase formation.

#### 4.4. Samarium Doped Zirconolite

Several formal studies have utilised  $\text{Sm}^{3+}$  as a surrogate for either minor actinide species or as a neutron absorbing additive. It was demonstrated in early studies, such as that of Kesson et al., that zirconolite may incorporate  $\text{Sm}^{3+}$  with concentrations as high as 0.67 f.u. [30]. A zirconolite-structured material with the composition  $(\text{Ca}_{0.55}\text{Sm}_{0.45})(\text{Sm}_{0.22}\text{Zr}_{0.71}\text{Mg}_{0.06})\text{Ti}_{1.99}\text{O}_7$  was synthesised by hot pressing a 1:1 molar mixture of  $\text{CaZrTi}_2\text{O}_7$ — $\text{Sm}_2\text{Ti}_2\text{O}_7$  at 1400 °C and 1459 °C, producing a two-phase mixture consisting of zirconolite (with the above composition) and pyrochlore. This study provided many initial insights into  $\text{Ln}^{3+}$  accommodation within the zirconolite structure, demonstrating that larger  $\text{Ln}^{3+}$  species were preferentially partitioned within the  $\text{Ca}^{2+}$  site on account of size constraints, whilst smaller  $\text{Ln}^{3+}$  may be accommodated in the  $\text{Zr}^{4+}$  site. For the formation of an  $\text{Mg}^{2+}/\text{Nb}^{5+}$  substituted zirconolite sample, with composition  $(\text{Ca}_{0.57}\text{Sm}_{0.32}\text{Zr}_{0.06})\text{Zr}_{1.00}(\text{Ti}_{1.14}\text{Mg}_{0.40}\text{Nb}_{0.40}\text{Zr}_{0.06})\text{O}_7$ , Kesson et al. demonstrated that  $\text{Sm}^{3+}$  might be incorporated into the zirconolite structure with considerable charge balance species retained on the  $\text{Ti}^{4+}$  site.

Mikhailenko et al. fabricated zirconolite ceramics in the pseudo-binary system  $(1-x)\text{CaZrTi}_2\text{O}_7 - (x)\text{SmAlO}_3$ , for  $x = 0.25, 0.50$ , and  $0.75$ , using a conventional cold press and sinter method between 1400 and 1500 °C [60]. When targeting  $x = 0.25$ , a two-phase mixture of zirconolite and perovskite was obtained, accounting for 80% and 20% of the overall phase fraction, respectively, with  $\text{Sm}^{3+}$  concentrated in the zirconolite phase. Increasing the  $(x)$  parameter served to greatly reduce the quantity of the zirconolite phase, with a mixture of zirconolite (25%), perovskite (60%), and free oxide (15%, Sm-stabilised  $\text{ZrO}_2$ ) produced when targeting  $x = 0.75$ . Coincidentally,  $\text{Sm}^{3+}$  was overwhelmingly concentrated in the perovskite phase, demonstrating the tendency of  $\text{ABO}_3$  phases to accommodate  $\text{Ln}^{3+}$  species. At each interval of  $(x)$ , powder diffraction data were consistent with zirconolite-3T of trigonal symmetry.

Jafar et al. synthesised a series of zirconolites targeting the  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Sm}_{2x}\text{Ti}_2\text{O}_7$  solid solution, with  $0.00 \leq x \leq 1.00$ , by mixed oxide synthesis at 1300 °C [50]. Three distinct phase fields were identified, comprising zirconolite-2M, zirconolite-4M, and cubic pyrochlore. Specifically, a single phase zirconolite was produced when targeting  $x = 0.00$ , with a mixture of zirconolite-2M and a secondary perovskite phase obtained in the interval  $0.05 \leq x \leq 0.10$ . Zirconolite-2M and zirconolite-4M were observed to coexist (alongside the perovskite phase) in the range of  $0.15 \leq x \leq 0.25$ . Zirconolite-4M was the only polytype present for  $0.30 \leq x \leq 0.35$ , with the appearance of a pyrochlore structured phase at  $x = 0.40$ , while Sm-rich pyrochlore was produced as a single phase for  $x \geq 0.60$ . The presence of the  $\text{CaTiO}_3$  secondary phase was attributed to the preferential occupation of  $\text{Sm}^{3+}$  within the 8-fold coordinated  $\text{Ca}^{2+}$  site, presumably due to ionic size compatibility ( $v_{\text{VIII}}\text{Sm}^{3+} = 1.07 \text{ \AA}$ ). The behaviour of  $\text{Sm}^{3+}$  in zirconolite in the  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Sm}_{2x}\text{Ti}_2\text{O}_7$  system is similar to that of analogous solid solutions containing  $\text{Nd}^{3+}$ ,  $\text{Gd}^{3+}$ ,  $\text{Y}^{3+}$ , etc.; therefore, it may be expected that alternative substitution regimes would not significantly deviate from the expected phase evolution.

Zhang et al. targeted two  $\text{Sm}^{3+}$  incorporation schemes:  $\text{Ca}_{1-x}\text{Sm}_x\text{ZrTi}_{2-x}\text{Al}_{2x}\text{O}_7$  ( $0.00 \leq x \leq 0.50$ ) and  $\text{Ca}_{1-x}\text{Zr}_{1-x}\text{Sm}_{2x}\text{Ti}_2\text{O}_7$  ( $0.00 \leq x \leq 1.00$ ), using a combustion synthesis

method, as previously described [69]. The substitution scheme targeting  $\text{Sm}^{3+}$  incorporation within  $\text{Ca}^{2+}$  and  $\text{Zr}^{4+}$  was identical to that of Jafar et al. [50]; however, the phase fields were inconsistent, inasmuch as zirconolite-4M was not identified as an intermediate phase in transition between  $\text{CaZrTi}_2\text{O}_7$  and  $\text{Sm}_2\text{Ti}_2\text{O}_7$ . It is unclear why, in this instance, a Sm-rich zirconolite-4M phase was not produced, but it is possibly a result of the discrepancy between fabrication parameters (Jafar et al. utilised a conventional solid state sintering method from component oxides, whereas Zhang et al. used the combustion synthesis method, with  $\text{CuO}$  as the oxidant). Synthesis of  $\text{Ca}_{1-x}\text{Sm}_x\text{ZrTi}_{2-x}\text{Al}_{2x}\text{O}_7$  produced a heterogeneous phase assemblage in which undigested component oxides were present in the microstructure at all levels of the targeted Sm concentration. Furthermore, as the phase assemblage evolved towards  $x = 0.50$ , powder XRD data were consistent with zirconolite occupying the 3T polytype. These data were not consistent with those produced by Ma et al. in the  $\text{Ca}_{1-x}\text{Ln}_x\text{ZrTi}_{2-x}\text{Al}_x\text{O}_7$  system for  $\text{Ln} = \text{La}, \text{Nd}, \text{Gd}, \text{Ho},$  and  $\text{Yb}$ , insofar as zirconolite-3T was not detected [48]. As the ionic radii of  $\text{Sm}^{3+}$  (1.07 Å) fall between that of  $\text{Nd}^{3+}$  (1.11 Å) and  $\text{Ho}^{3+}$  (1.01 Å) in eight-fold coordination, this may imply that the zirconolite-3T phase in this system is stabilised over a very narrow compositional range.

In summary, whilst there are numerous studies that have successfully incorporated  $\text{Sm}^{3+}$  within zirconolite, this has typically been with the view to simulate trivalent minor actinide species rather than as a neutron absorber. Nevertheless, despite some discrepancies as to polytype formation at extensive Sm substitution, the phase fields produced when incorporating  $\text{Sm}^{3+}$  in zirconolite are largely analogous to  $\text{Gd}^{3+}$ . The similarities between Sm and Gd (i.e., the cost index and the absorption cross section) signify that whilst both could be expected to be an effective neutron absorbing additive, there is no apparent benefit that Sm would confer over Gd.

#### 4.5. Cadmium Doped Zirconolite

The authors are not aware of any studies that have systematically investigated the direct incorporation of  $\text{Cd}^{2+}$  within the zirconolite structure. Nevertheless, the ionic radii of  $\text{Ca}^{2+}$  and  $\text{Cd}^{2+}$  are nearly identical (1.12 Å and 1.10 Å in eight-fold coordination, respectively); hence, a continuous solid solution between  $\text{CaZrTi}_2\text{O}_7$  and  $\text{CdZrTi}_2\text{O}_7$  is feasible. Similar to  $\text{Hf}^{4+}$  zirconolite, this may allow the substitution of a requisite quantity of  $\text{Cd}^{2+}$  within an optimised composition with negligible impact on the obtained structure. A Cd-zirconolite was reported by Leturcq et al.; zirconolite with nominal composition  $\text{Ca}_{0.863}\text{Nd}_{0.137}\text{ZrTi}_{1.863}\text{Al}_{0.137}\text{O}_7$  was homogenised before the addition of 1 wt. %  $\text{CdO}$  and sintering as a pellet at 1450 °C in the air for 4 h [70]. SEM analysis confirmed that whilst Cd was partially incorporated within the zirconolite phase, Cd was also partitioned within a minor  $\text{CaTiO}_3$  phase (presumably accommodated within the  $\text{Ca}^{2+}$  site in both instances). A series of  $\text{A}^{2+}\text{A}^{4+}\text{Ti}_2\text{O}_7$ -type charge-coupled pyrochlore materials with  $\text{A}^{2+} = \text{Cd}^{2+}$  were reported by McCauley and Hummel.  $\text{CdCeTi}_2\text{O}_7$  was successfully synthesised as a single phase cubic pyrochlore phase under a heat treatment of 1150 °C for 24 h, whereas the targeted  $\text{CdThTi}_2\text{O}_7$  and  $\text{CdHfTi}_2\text{O}_7$  phases, prepared at 1250 °C for 24 h and 96 h, respectively, produced a majority pyrochlore phase alongside residual impurities. Interestingly, it was reported that the  $\text{CdZrTi}_2\text{O}_7$  phase, which would be expected to yield the zirconolite structure, instead formed a single phase cubic pyrochlore with the lattice dimension  $a = 10.02 \pm 1$  Å, indicating that  $\text{Cd}^{2+}$  may best be accommodated as a neutron absorbing additive at relatively low concentrations to avoid a partial phase transformation to pyrochlore. Nevertheless, confidence that  $\text{Cd}^{2+}$  may be accommodated in zirconolite in dilute proportions without significant disruption to the phase assemblage may be taken from the fact that several other pertinent Ca/Cd analogues have been reported in the literature, e.g.,  $\text{CaTiO}_3$  [71] and  $\text{CdTiO}_3$  [72],  $\text{Ca}_2\text{Nb}_2\text{O}_7$  [73] and  $\text{Cd}_2\text{Nb}_2\text{O}_7$  [74], and  $\text{CaMoO}_4$  [75] and  $\text{CdMoO}_4$  [76].

In summary, the advantage of  $\text{Cd}^{2+}$  within the formulation may be similar to that of  $\text{Hf}^{4+}$  insofar as  $\text{Cd}^{2+}$  may be able to effectively replace  $\text{Ca}^{2+}$  as the A-site cation in zirconolite on the basis of ionic radii, though the phase  $\text{CdZrTi}_2\text{O}_7$  has not yet been

reported. Furthermore, Cd is an attractive additive due to its relatively low cost index in relation to Gd/Hf/In. The lack of literature regarding Cd-doped zirconolite may contribute significantly towards its down selection. Moreover, the handling of Cd is accompanied by toxicity concerns, which would be a notable consideration during precursor production and the downstream handling of powders. For example, in the work of McCauley and Hummel, it was necessary to produce Cd<sup>2+</sup>-containing compounds within platinum tubes to avoid volatilisation. This may prove unattractive for the synthesis of Pu-containing compounds. The reduction of CdO to Cd metal may also be a concern, particularly for compositions produced by HIP, for which the Fe-rich environment of the HIP container produces a reducing environment. On the basis of these stipulations, Cd may not be selected as a practically useful additive species.

#### 4.6. Indium Doped Zirconolite

Similarly to Cd, there are few studies that have incorporated In<sup>3+</sup> in zirconolite. A suite of zirconolites targeting nominal compositions CaZrTi<sub>2-x</sub>In<sub>x</sub>O<sub>7-x/2</sub> ( $x = 0.25, 0.50$  and  $1.00$ ) was prepared by Begg et al., reacted at 1400 °C for 40 h [77]. Here, In<sup>3+</sup> was introduced as a surrogate for Ti<sup>3+</sup> in order to simulate highly reducing conditions. Zirconolite was not produced as a single phase at any targeted level of In<sup>3+</sup> concentration (with the exception of  $x = 0.00$ ). Targeting a nominal concentration of  $x = 0.25$  produced a two phase mixture of Ca<sub>0.85</sub>Zr<sub>1.16</sub>Ti<sub>1.67</sub>In<sub>0.33</sub>O<sub>7</sub> (zirconolite) and Ca<sub>0.98</sub>Zr<sub>0.01</sub>Ti<sub>0.99</sub>In<sub>0.01</sub>O<sub>3</sub> (perovskite), comprising 90 wt. % and 10 wt. % of the phase assemblage, respectively. Increasing the nominal In<sup>3+</sup> concentration to  $x = 0.50$  significantly disrupted the phase assemblage, such that 20 wt. % of the overall yield was composed of a Ca<sub>0.17</sub>Zr<sub>0.35</sub>Ti<sub>0.33</sub>In<sub>0.25</sub>O<sub>1.9</sub> fluorite-structured phase. Targeting the nominal composition CaZrTiInO<sub>6.5</sub> (i.e.,  $x = 1.00$ ) completely destabilised the zirconolite phase, with the phase assemblage comprised of 40 wt. % perovskite and 60 wt. % fluorite. Nevertheless, considering the estimated ionic radius of In<sup>3+</sup> in 7-fold coordination (0.86 Å), In<sup>3+</sup> may be more easily accommodated within the Zr<sup>4+</sup> site, replacing half of the necessary Al<sup>3+</sup> cations necessary in a substitution regime such as Ca<sub>1-x</sub>Pu<sub>x</sub>Zr<sub>1-x</sub>In<sub>x</sub>Ti<sub>2-x</sub>Al<sub>x</sub>O<sub>7</sub>. A recent study by Blackburn et al. produced a detailed characterisation of the Ca<sub>1-x</sub>Zr<sub>1-x</sub>In<sub>2x</sub>Ti<sub>2</sub>O<sub>7</sub> solid solution [78]. It was confirmed that a single phase zirconolite-2M material could be stabilised for  $x \leq 0.20$ ; however, contrary to formulation, it was determined that In<sup>3+</sup> was dominant in the Ti<sup>4+</sup> site, resulting in structural rearrangement and significant disruption of the phase assemblage beyond  $x \geq 0.30$ . Indicating that In<sup>3+</sup> has a preference for the Ti<sup>4+</sup> site, a series of Ca<sub>1-x</sub>U<sub>x</sub>ZrTi<sub>2-2x</sub>In<sub>2x</sub>O<sub>7</sub> compositions were fabricated targeting dominant U<sup>4+</sup> and U<sup>5+</sup>, with U acting as a structural surrogate for Pu. It was confirmed that at relatively low concentrations ( $x = 0.05$  and  $0.10$ ), single phase zirconolite-2M when processed under both argon and air. These data suggest that an additive of In<sup>3+</sup> could potentially be included as a charge balancing species in candidate zirconolite solid solution regimes.

In summary, the deployment of In<sup>3+</sup> as a potential neutron absorbing additive in zirconolite is feasible when targeting substitution for Ti<sup>4+</sup>. However, deployment at an industrial scale may be inhibited by the large associated cost that would be necessary to ensure sufficient neutron absorption. Furthermore, the additive:Pu ratio required for In<sup>3+</sup> to provide the necessary criticality control is an order of magnitude greater than other trivalent species, such as Gd<sup>3+</sup> or Sm<sup>3+</sup>. Although there is some literature on this subject, it may be stated that at the present time, there is not sufficient evidence to provide confidence in the ability of In<sup>3+</sup> to successfully act as a neutron absorber in zirconolite.

#### 4.7. Boron Doped Zirconolite

Boron cannot be accommodated in a solid solution with zirconolite on the basis of ionic radii (B<sup>VI</sup> = 0.27 Å); therefore, it must be distributed as discrete intergranular particulates throughout the microstructure, which would most likely concentrate at grain boundaries and triple points. Furthermore, unlike the other additives considered, boron would be present as the carbide B<sub>4</sub>C rather than oxide. Gong et al. introduced B<sub>4</sub>C powders into

mixtures of Y-stabilised ZrO<sub>2</sub>, targeting 0.71, 1.40, and 2.12 wt. %, corresponding to a Pu content of 9.3, 17.6, and 25.3 wt. %, respectively [79]. B<sub>4</sub>C was found to be spatially distributed in the YSZ matrix, with no void space or reaction between the YSZ-B<sub>4</sub>C observed. A thorough survey of the literature did not identify any reports detailing the inclusion of B<sub>4</sub>C in zirconolite waste packages.

## 5. Gap Analysis

The objective of this review was to provide an overview of established solid solution regimes for Pu incorporation in zirconolite and insight into potential pathways towards the formulation of a zirconolite-rich wastefrom capable of accommodating a requisite quantity of a suitable neutron absorbing additive. Within the available literature, there remain several high level issues that have not been adequately addressed (at least in a manner that is systematic enough to provide confidence in the approach) that are required to inform a decision regarding the most effective route towards disposal in a zirconolite matrix. These are detailed herein.

**(1) Choice of substitution scheme and favourability of dominant polytype:** The choice of solid solution regime appears to be favoured towards a coupled A-site substitution, by which Pu<sup>4+</sup> is accommodated in solid solution with Ca<sup>2+</sup> sites, with a smaller lower valence cation, such as Al<sup>3+</sup> distributed across the available Ti<sup>4+</sup> sites to provide charge balance. The choice of substitution scheme and, to a lesser extent, the partial oxygen pressure of the atmosphere introduced during the sintering process have been shown to control polytype formation. Whilst it has not been explicitly defined as a criterion for the design of zirconolite wastefroms, it is generally accepted that maintaining zirconolite in its 2M form (or indeed, a single phase polytype of the 3O/3T species) may be preferable, rather than a mixture of polytypes. Despite this, we are not aware of any studies that conclusively determine that the chemical durability of zirconolite wastefroms is compromised, to any meaningful and quantifiable extent, by the presence of multiple zirconolite polytypes, and indeed whether any of the isolated polytypes exhibit any variation in leaching properties.

**(2) Processing conditions for additive incorporation:** In order to support an immobilisation program for Pu, the construction of a pilot scale facility is necessary, ideally allowing for the physical optimisation of all chemical engineering aspects and processing parameters. As outlined in Figure 1, there are several head-end powder handling operations that may need to be (1) optimised for handling  $\alpha$ -emitters such as Pu and (2) configured correctly to ensure efficient and homogeneous feed into downstream operations such as can filling. One or more of the head-end operations will make use of high intensity homogenisation/size reduction techniques in order to sufficiently mix the feed material and ensure PuO<sub>2</sub> is completely dispersed in the wastefrom. Ball milling and attrition milling are both attractive options that take advantage of pre-existing ceramic processing technologies. The milling stage would ideally utilise dry processing, in which the homogenised inactive component of the feedstock is blended with the PuO<sub>2</sub> and mixed with a lubricant to aid the milling process before discharge into HIP cans. A variety of organic lubricants were utilised by Squire et al. in the attrition milling of zirconolite (target composition Ca<sub>0.75</sub>Ce<sub>0.25</sub>ZrTi<sub>1.625</sub>Fe<sub>0.375</sub>O<sub>7</sub>), consolidated by HIP [80]. Examination of the product materials post-HIP evidenced poor densification due to incomplete lubricant burnout, resulting in the incomplete formation of a single phase product. It was discussed that the use of fumed metal oxides might provide an inherent advantage insofar as they may act as both a milling aid and form part of the wastefrom itself. Examples include the use of hydrophobic Al<sub>2</sub>O<sub>3</sub> as a fumed powder, or indeed Ti<sub>2</sub>O<sub>3</sub>, which may, in turn, allow the use of Ti<sup>3+</sup> as a charge compensation species, negating the need for Al<sup>3+</sup>.

**(3) Validation of surrogate studies using plutonium:** The use of chemical surrogates is prevalent in most laboratory-scale wastefrom trials, as a precursor to active validation trials, on the basis of cost and expediency. Ce<sup>4+</sup> and U<sup>4+</sup> are typically employed as surrogates for Pu<sup>4+</sup> in the immobilised form, both in ceramic and vitrified wastefroms. However, it is widely accepted that surrogates do not produce wastefrom characteristics that can be

directly translated to Pu-substituted compositions. Discrepancies between the surrogate and 'real' compositions are typically a result of redox activity, with the tendency of Ce to reduce to the  $Ce^{3+}$  form considerably greater than Pu. For example, Blackburn et al. recently demonstrated that the partial oxygen pressure of the sintering environment applied for  $CaZr_{1-x}Ce_xTi_2O_7$  ceramics significantly altered the obtained phase assemblage [16]. Vance et al. demonstrated that the sintering of  $CaZr_{1-x}U_xTi_2O_7$  ceramics in the air significantly disrupted the phase assemblage due to the formation of  $U^{5+}$  [31]. As these behaviours were not replicated in the corresponding  $CaZr_{1-x}Pu_xTi_2O_7$  solid solution synthesised by Begg et al. [32], the use of chemical surrogates must be done so with caution. The physical characteristics (such as morphology and density) of constituent oxide powders must also be taken into account.

(4) Relative leach rates of neutron absorbing additive and plutonium/uranium: There is, at present, an absence of systematic studies aiming to evaluate the relative leach rates of neutron absorbing elements alongside Pu and U (as  $^{235}U$  is a key decay product from  $^{239}Pu$ ). In order to ensure that the likelihood of post closure criticality remains low, and if low consequence should it occur, it is necessary to understand whether the neutron absorbing species and fissile inventory are likely to leach congruently into the near field environment. In an event whereby the absorbing additive is preferentially leached with respect to Pu, the wastefrom may be exposed to an in-package criticality event, whereas an out of package event would see Pu leached at higher rates than the chosen additive. Systematic studies aiming to quantify the leach rates of surrogates (U and Ce) and neutron absorbing species (Gd and Hf) from representative wastefrom compositions, e.g.,  $(Ca,U,Gd)(Zr,Hf,Gd)(Ti,Al)_2O_7$  are therefore necessary, as these data could feasibly be used to model criticality scenarios and, in turn, allow selection of the most appropriate additive.

## 6. Conclusions

The aim of this review was to provide an overview of the current status of the UK immobilisation program for Pu inventories, with a specific focus on efforts to optimise the baseline zirconolite formulation to accommodate a requisite quantity of neutron absorbing additive. A detailed summary of Pu/additive solubility in a range of established zirconolite solid solution regimes was also provided, alongside a review of zirconolite crystallography. A selection of potential neutron absorbing additives was evaluated through an initial screening, with the aim to down-select any candidates that did not appear attractive on the basis of cost and neutronics. Accordingly, a literature review of successful candidates, Gd, Hf, Sm, Cd, In, and B, was performed, aiming to understand their respective solubility within the candidate Pu-zirconolite phase. Where possible, a review of the solid solution limits with respect to polytype transformations and oxide solubility was provided. For each additive, a high-level summary was presented, identifying both Gd and Hf as the most promising oxide compounds.

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