Magnesium and chromium isotope evidence for initial melting by radioactive decay of $^{26}$Al and late stage impact-melting of the ureilite parent body

Elishevah M.M.E. van Kooten *, Martin Schiller, Martin Bizzarro

Centre for Star and Planet Formation and Natural History Museum of Denmark, University of Copenhagen, DK-1350 Copenhagen, Denmark

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Abstract

Polymict ureilites are meteoritic breccias that provide insights into the differentiation history of the ureilite parent body. We have sampled a total of 24 clasts from the polymict ureilite Dar al Gani 319, representing a variety of lithologies such as mantle residues, cumulates and crustal fragments that are genetically related to monomict ureilites. In addition, we sampled four non-indigenous dark clasts and two chondrule-containing clasts from the same meteorite. We report on the petrology and the bulk mass-dependent and mass-independent magnesium and chromium isotope systematics of these clasts. The DaG 319 polymict ureilite consists predominantly of clasts related to Main Group ureilite residues (MG clasts) with varying Mg#s (0.74–0.91), as well as a significant fraction of olivine-orthopyroxene clasts related to Hughes Type ureilites (HT clasts) with consistently high Mg#s (~0.89). In addition, DaG 319 contains less abundant feldspathic clasts that are thought to represent melts derived from the ureilite mantle. A significant mass-dependent Mg-isotope fractionation totaling $\Delta_{25}^{25}\text{Mg} = -450$ ppm was found between isotopically light feldspathic clasts ($\mu^{25}\text{Mg} = -305 \pm 25$ to $15 \pm 12$ ppm), MG clasts ($\mu^{25}\text{Mg} = -23 \pm 51$ ppm) and HT clasts ($\mu^{25}\text{Mg} = 157 \pm 21$ ppm). We suggest that this isotopic offset is the result of equilibrium isotope fractionation during melting in the presence of an isotopically light magnesite component. We propose Mg-carbonates to be stable in the upper ureilite mantle, and pure carbon phases such as graphite to be stable at higher pressures. This is consistent with HT clasts lacking carbon-related phases, whereas MG clasts contain abundant carbon. The timing of differentiation events for the ureilitic clasts are constrained by high precision $^{53}\text{Mn}-^{53}\text{Cr}$ systematics and $^{26}\text{Al}-^{26}\text{Mg}$ model ages. We show that a dichotomy of ages exist between the differentiation of main group ureilite residues and HT cumulates rapidly after CAI formation and later remelting of cumulates with corresponding feldspathic melts, at 3.8 ± 1.3 Myr after CAI formation. Assuming an initial $^{26}\text{Al}^{/}/^{27}\text{Al}$ abundance $[(^{26}\text{Al}^{/}/^{27}\text{Al})_0 = 1.33(0.04) \times 10^{-5}]$ similar to the angrite parent body, the early melting event is best explained by heat production from $^{26}\text{Al}$ whereas the late event is more likely caused by a major impact. Variations in $^{54}\text{Cr}$ between MG clasts and HT clasts agree with a carbonaceous chondrite impactor onto the ureilite parent body. This impactor may be represented by abundant dark clasts found in polymict ureilites, which have $\mu^{26}\text{Mg}$ and $\mu^{4}\text{Cr}$ signatures similar to CI chondrites. Similar volatile-rich dark clasts found in other meteorite breccias provide insights into the timing of volatile influx to the accretion region of the terrestrial planets.

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1. INTRODUCTION

Achondrites are basaltic and plutonic fragments of differentiated asteroidal and planetary bodies. Having
undergone most of their differentiation such as melting and crystallization in the first few million years of Solar System formation (Lugmair and Shukolyukov, 1998; Bizzarro et al., 2005; Amelin, 2008; Kleine et al., 2012; Schiller et al., 2010, 2011, 2015a), these meteorites provide insights into the earliest stages of the magmatic evolution of planets. Ureilites are the second most abundant group of achondrites and contain fragments of mostly mantle-derived materials, which can be divided into dominantly unbrecciated main group (or monomict) ureilites and rarer polymict (multiple clast types) ureilites. Most main group ureilites consist of olivine-pyroxene (dominantly pigeonite) mantle restites with varying magnesium numbers (Mg# = Mg/(Fe + Mg)), constant Mn/Mg ratios (Goodrich et al., 1987; Goodrich and Delaney, 2000) and high carbon abundances (~3 wt.%). These characteristic features suggest that ureilites are the product of melting reactions that reduce FeO-containing silicates to Fe-metal and Mg-rich silicates during partial melting in the presence of carbon (Goodrich et al., 1987, 2007; Warren and Kallemeyn, 1992; Walker and Grove, 1993; Singletary and Grove, 2003). Alternatively, the range of Mg#s can be explained by heterogeneous accretion of the ureilite parent body (UPB) (Warren and Kallemeyn, 1992; Warren and Huber, 2006; Warren, 2012). Monomict ureilites Mg#s correlate with oxygen isotope signatures of ureilites that fall on the Carbonaceous Chondrite Anhydrous Mineral (CCAM) line (Clayton and Mayeda, 1988, 1996), suggesting that these isotope signatures are of primitive origin (Sanders et al., 2017). However, rapid removal of partial melts formed during potential melting can preserve initial heterogeneity of oxygen isotopes as well. Fast migration of melts is ensured by the melting process, where the formation of CO and CO2 gases contribute to explosive volcanism of partial melts, thereby potentially explaining the absence of these igneous rocks in our meteorite collection (Takeda, 1987; Warren and Kallemeyn, 1992; Scott et al., 1993). However, feldspathic clasts are present in polymict ureilites and are suggested to represent fragments of these SiO2-rich melts (Ikeda and Prinz, 2001; Ikeda et al., 2003; Cohen et al., 2004; Bischoff et al., 2014). These plagioclase-rich clasts are small (<mm-sized) fragments with variable textures and compositions (Cohen et al., 2004), ranging from mafic (olivine-augite and anorthite-rich) to felsic mineralogies (albite-rich) indicating different degrees of fractional crystallization and cooling histories. Polymict ureilites also contain clasts of monomict ureilites (type I ureilite clasts; Ikeda et al., 2000), olivine-orthopyroxene-augite clasts similar to a group of unusual ureilites typified by Hughes 009 (type II ureilite clasts; Ikeda et al., 2000; Ikeda and Prinz, 2001; Goodrich et al., 2001) and olivine-augite clasts that are thought to represent para-cumulates (Goodrich et al., 2004).

The paradoxical blend of high degree partial melting (~25-30%) required to produce the composition of the olivine-pigeonite restites (Scott et al., 1993; Goodrich, 1999; Goodrich et al., 2007), together with noble gas signatures (Göbel et al., 1978), siderophile trace element patterns in metal (Rankenburg et al., 2008), anomalous nitrogen (Grady and Pillinger, 1988) and oxygen isotope signatures that resemble that of primitive chondrites (Clayton and Mayeda, 1988, 1996), complicates the development of a ureilite differentiation model that satisfies all constraints. For example, the timeline and stages of magmatic differentiation based on the groupings of feldspathic clasts (one-stage or multi-stage) and the effect of the disruptive event that formed the polymict ureilites remain unresolved. Most of these issues have been addressed by melting models that have used major elements, trace elements and rare earth element (REE) patterns to constrain the physical conditions of differentiation and the chemical nature of the UPB and its precursor material (Singletary and Grove, 2003; Goodrich et al., 2007; Barrat et al., 2016b). In addition, 26Al-26Mg, 53Mn-53Cr and 182Hf-182W short-lived isotope, as well as U-Pb and 176Lu-182Hf long-lived isotope systematics have been used to constrain the formation history of feldspathic clasts and monomict ureilites (Kita et al., 2003, 2007; Goodrich et al., 2010; Qin et al., 2010b; Yamakawa et al., 2010; Bischoff et al., 2014; Amelin et al., 2015; Buddé et al., 2015; Bast et al., 2016). However, the lack of an established genetic relationship between these feldspathic clasts and other ureilite components limits their use to develop a unified model for the magmatic evolution of the UPB. More precise isochron data may reveal previously unseen relationships between different lithologies. The focus of this work lies therefore in defining genetic relationships between various igneous reservoirs of the UPB and constraining its differentiation history by using the 26Al-26Mg and 53Mn-53Cr short-lived radioisotope systems along with mass-dependent magnesium and chromium isotope compositions. The combination of chronology of magmatic processes, mass-dependent isotope fractionation patterns and 53Cr nucleosynthetic anomalies can be utilized to outline the physical processes that resulted in the differentiation of the UPB. The various ureilite components (e.g., feldspathic, type I and type II ureilite clasts) were sampled from the polymict ureilite Dar al Gani 319, which is well known for the diversity of its ureilite clasts (Ikeda et al., 2000, 2003; Ikeda and Prinz, 2001; Cohen et al., 2004). In addition, we have sampled a variety of chondritic clasts present in this meteorite to identify the nature of potential precursor materials to the UPB. We describe relationships between various ureilite clasts and provide a timeline for a two-stage differentiation history of the UPB. Furthermore, we provide constraints on the chemical and physical mechanisms involved in these differentiation events.

2. SAMPLES

Samples from this study were taken from the polymict ureilite DaG 319 and general descriptions of these clast types are given in overview Table 1, whereas detailed descriptions, images and compositional data of individual samples and their phases can be found in the Electronix Annex. DaG 319 is a micro-breccia and multiple clasts have been studied extensively for their mineralogy, petrology, trace element and oxygen isotope composition (Ikeda et al., 2000, 2003; Ikeda and Prinz, 2001; Cohen et al., 2004; Goodrich et al., 2004; Kita et al., 2004, 2006).
Although monomict ureilite clasts dominate in DaG 319, this meteorite also contains a variety of feldspathic igneous clasts that may be indigenous to the UPB (<2 vol.%) as well as chondritic clasts (<2 vol.%).

For the purpose of Mg- and Cr-isotope analyses, we sampled the following clasts:

1. We sampled nine type I ureilitic clasts as defined by Ikeda et al. (2000) that are identical to monomict ureilites (hereafter MG clasts, main group clasts). These clasts were mainly present as separate entities of mm-sized olivine and pyroxene with a wide range of Mg#s (Fo74–92). We sampled four olivine and four pyroxene clasts in addition to one clast containing both minerals.

2. We sampled five clasts classified as type II ureilitic clasts or Hughes Type ureilites (hereafter HT clasts, Hughes Type clasts; Ikeda et al., 2000; Goodrich et al., 2001). These clasts are cm-sized objects containing olivine and low-Ca pyroxene (Wo < 5) with a narrow range of high Mg#s (Fo ~74–92).

3. Ten feldspathic clasts were sampled, which were generally smaller (<1 mm) than MG clasts and include albitic, olivine-augite and labradoritic lithologies (Cohen et al., 2004).

4. Lastly, we sampled six chondritic clasts. Four of these are primitive carbonaceous clasts or ‘dark clasts’, which make up the bulk of the chondritic clasts in DaG 319 and were previously considered as potential precursor material to the UPB (Ikeda et al., 2003). Dark clasts are fragments of carbonaceous chondrites consisting of hydrothermally altered chondritic matrix. They have so far been classified as CI (Ivuna-type) and CM (Mighei-type) like clasts based on the composition of their phyllosilicates (serpentinite-smectite solid solution) and the presence of sulfides and magnetite. Oxygen isotope compositions of dark clasts typically follow the CCAM line, but also plot above the terrestrial fractionation line (Clayton and Mayeda, 1988). In addition to the dark clasts, two chondritic clasts that contain chondrules were sampled.

3. ANALYTICAL METHODS

3.1. Mineral chemistry

Initial investigations of the DaG 319 meteorite were conducted on polished thick sections using the Phillips XL40 scanning electron microscope located at the Geological Survey of Denmark and Greenland. Back scattered electron images (BSE) were taken of all samples selected for Mg- and Cr-isotope analyses, along with semi-quantitative elemental maps by energy dispersive spectroscopy (EDS). Quantitative compositional information of individual minerals was retrieved by electron microprobe with the JEOL JXA-8200 Superprobe at the University of Copenhagen. The microprobe operated at 15 keV accelerating voltage, with a 15 nA beam current and a fully focused
4 μm beam using five wavelength dispersive spectrometers. Sample analyses were carried out along standard measurements of in house silicates and oxides, described in Sandrin et al. (2009).

3.2. Magnesium and chromium isotope analyses

DaG 319 clasts were sampled from the thick sections with the computer-assisted NewWave micro-drill, using tungsten carbide drill bits at the Centre for Star and Planet formation (Copenhagen). Powders from the drilled samples were transferred with double distilled MQ water to clean Savillex beakers. Olivine and pyroxene-rich samples (MG and HT clasts) were typically between 0.5 and 1.0 mg in size, whereas feldspathic clasts samples were smaller, yielding <100 μg of sample material. This resulted in 100–400 μg Mg and 2–10 μg Cr for MG and HT clasts and 2–10 μg Mg and 60–500 ng of Cr for feldspathic clasts. To avoid contaminations from surrounding material, which can potentially have very contrasting Mg- and Cr-isotope signatures, care was taken not to drill close to the clast boundaries. After drilling, the individual drill spots were examined visually under a microscope for possible contamination. Any samples where contamination was suspected were discarded. Samples were digested first on a hotplate at 150 °C in a mixture of concentrated HNO3/HF acid for 2 days and subsequently for 1 day in Parr bombs at 210 °C. Afterwards, sample solutions were dried down and treated with aqua regia for one more day. Olivine and pyroxene MG clasts were not completely dissolved at this point, due to remaining carbon residues. These samples were further treated for 3 additional days in concentrated HNO3/HF acid, using Parr-bombs at 210 °C. After this treatment, all Cr and Mg was in solution (including potential ilmenite and spinel grains that have previously been observed in ureilic materials) and ready for elemental purification by ion exchange chromatography. A 5% aliquot was taken from the sample solutions before commencing with the purification process and used to determine the 27Al/26Mg and 55Mn/52Cr ratios on the ThermoFisher X-series quadrupole Ionization Coupled Plasma Mass Spectrometer (ICPMS) at the Centre for Star and Planet Formation (Copenhagen), which are accurate to 2%. Purification of Mg and Cr was done by methods described in Bizzarro et al. (2011) and Van Kooten et al. (2016).

The isotopic composition of the purified Mg was determined by the standard-sample bracketing technique using the Neptune Plus Multi Collector (MC)-ICPMS at the Natural History Museum of Denmark based on protocols described in Bizzarro et al. (2011) and Van Kooten et al. (2016). Samples were typically analyzed with a signal intensity of 20–45 V on mass 24Mg (all collectors are connected to amplifiers with 1011 Ω feedback resistors) for smaller feldspathic samples and 80–100 V on 24Mg was measured on a detector connected to a 1010 Ω feedback resistor, whereas 25Mg and 26Mg were measured on detectors connected to 1011 Ω feedback resistors) for larger samples. Apart for two feldspathic clasts that did not contain sufficient amounts of Mg, all samples we systematically analyzed ten times (Table 2). Mg-isotope data are reported in the μ-notation as deviations from the DTS-2b standard [μ25Mg DTS-2b = −122 ± 17 ppm (2SD, Bizzarro et al., 2011)] according to the following formula:

$$\mu^{25}_\text{Mg} = \left( \frac{\mu^{25}_\text{Mg}_{\text{sample}}}{\mu^{25}_\text{Mg}_{\text{DTS-2b}}} - 1 \right) \times 10^6$$  \hspace{1cm} (1)$$

where x represents mass 25 or 26. The mass-independent component of 26Mg (μ26Mg) is reported in the same fashion, but represents deviations from the internally normalized 26Mg/24Mg of the sample from the reference standard, normalized to 25Mg/24Mg = 0.126896 (Bizzarro et al., 2011) using the exponential mass fractionation law. All Mg data reduction was conducted off-line using the Iolite data reduction package which runs within Igor Pro (Paton et al., 2011) and changes in mass bias with time were interpolated using a smoothed cubic spline. For each analysis, the mean and standard error of the measured ratios were calculated using a 2SD threshold to reject outliers. Individual analyses of a sample were combined to produce an average weighted by the propagated uncertainties of individual analyses and reported final uncertainties are the 2SE of the mean. The external reproducibility of the technique utilized here to acquire our Mg isotope data has been estimated by Bizzarro et al. (2011) to be 20 and 2.5 ppm for the 25Mg and 26Mg, respectively. This estimate is based on the repeated analysis of various terrestrial rock standards as well as extraterrestrial samples. In the current study, we have verified that our external reproducibility is in line with earlier estimates, through repeated analysis of individual column-processed rock digestion aliquots of various USGS terrestrial rock standards acquired over a one-year period that coincide with the time the data reported in this paper were acquired. In detail, we analyzed four aliquots of the BHVO-2 standard and seven aliquots of the DTS-2b standard. Note that nine out of the 11 standard analyses were reported in Olsen et al. (2016) and Van Kooten et al. (2016). Based on these data, we derive an external reproducibility of 22 and 2.5 for the 25Mg and 26Mg, respectively, which is in line with the estimate reported by Bizzarro et al. (2011) using the same technique. The Cr-isotope composition of all samples was measured by thermal ionization mass spectrometry (Triton TIMS) at the Centre for Star and Planet Formation. We used the small chemistry column purification techniques described by Schiller et al. (2014), which we adapted to include a last purification step to remove sodium and to minimize organics and elemental impurities introduced during the separation. This step utilizes an Eichrom AG50 X8 200–400 mesh cation resin and sample pretreatments to ensure full recovery of Cr after removing residual unwanted impurities (i.e., organics, Na, Fe, Ti, Al) that can cause matrix effects or interference on the Cr signal. First, the samples are pretreated in 50 μl 1 M HNO3 + 10 μl concentrated H2O2 for >5 days at room temperature to promote the speciation of Cr3+ (Larsen et al., 2016a). Subsequent dilution to a 0.5 M HNO3 loading solution and elution of 1.3 ml of 0.5 M HNO3 on a 100 μl cation resin, resulted in the removal of Na, K and organics. Adding an additional three column volumes of 1 M HF removed any remaining
trace of Fe, Al and Ti. Cr was then collected by elution with 6 M HCl. Cr yields are typically >95% and washes of samples with lower yields were subjected to an additional cycle of the last cation purification step and recombined. Chromium isotope analyses were conducted using a hybrid method of total evaporation and standard-sample bracketing fully described in Van Kooten et al. (2016). Samples were typically measured 16 times distributed over multiple sessions. Smaller samples were measured using fewer filaments according to their sample size (1 filament = 15 ng). The SRM-3112a standard was measured concurrently over all sessions. Cr-isotope data is presented in the $\varepsilon$-notation relative to this standard:

$$\varepsilon^{54}\text{Cr} = \left( \frac{^{54}\text{Cr}}{^{52}\text{Cr}} \right)_{\text{sample}} \left/ \left( \frac{^{54}\text{Cr}}{^{52}\text{Cr}} \right)_{\text{SRM-3112a}} - 1 \right. \times 10^4$$

(2)

where $x$ represents mass 53 and 54 and the $^{54}\text{Cr}/^{52}\text{Cr}$ values reflect the internally normalized values, normalized to $^{54}\text{Cr}/^{52}\text{Cr} = 19.2832$ (Trinquier et al., 2008a) using the exponential mass fractionation law. Individual analyses of a sample were combined to produce an average weighted over propagated uncertainties of individual analyses and reported final uncertainties are the 2SE of the mean using a 2SD outlier rejection scheme. From terrestrial rock standards (e.g., DTS-2b, BHVO-2) processed in this work and Van Kooten et al. (2016) that were measured over the same period as our samples, we estimate an external reproducibility of 6 ppm (2SD) and 12 ppm on mass-bias corrected $^{54}\text{Cr}$ and $^{52}\text{Cr}$ values, respectively.

Additional mass-dependent Cr-isotope measurements by MC-ICPMS were carried out on six MG and four HT clasts. Of these, one MG clast and all HT clasts were also
analyzed by TIMS. Mass-dependent Cr-isotope analyses were carried out by standard-sample bracketing following procedures described in Schiller et al. (2014) at the Centre for Star and Planet Formation (Copenhagen) with the Neptune Plus MC-ICPMS. Interferences from argon oxides and nitrides on the $^{52}$Cr, $^{53}$Cr and $^{54}$Cr signal were minimized by using a trifluoro-methane (CHF$_3$) supplementary gas and running in a ‘cold plasma’ mode with an RF power of 700 W. At high resolution this reduced the $^{40}$Ar$^{16}$O and $^{40}$Ar$^{14}$N signals to below 0.5 V and 0.2 V, respectively. Analyses were carried out on the low mass-side of the Cr peak, typically at mass 51.908. This peak position was confirmed before each standard analysis by peak centering on the $^{50}$Cr peak, which is the isotope least affected by gas based interferences. All samples were analyzed as ~1 ppm solutions and a 25 μl/min uptake rate, which resulted in a total Cr beam of ~20 V. Each individual sample run comprised 834 s (16.77 s x 50 blocks), consuming ~400 ng. Each sample was run five times, whereas the DTS-2b standard was run 20 times. All data was reduced using the Iolite software package in Igor Pro. Changes in mass bias with time were interpolated using a smoothed cubic spline and data are presented as the weighted mean and 2SE error. Repeated analyses of the DTS-2b rock standard using this setup resulted in an external reproducibility of 12 ppm (2SD) and 28 ppm on internally normalized $\varepsilon^{50}$Cr and $\varepsilon^{52}$Cr values, respectively. Mass-dependent Cr-isotopes are presented in the $\delta^{50}$Cr notation:

$$\delta^{50}\text{Cr} = \left[ \frac{\left(^{50}\text{Cr} / ^{52}\text{Cr}\right)_{\text{sample}}}{\left(^{50}\text{Cr} / ^{52}\text{Cr}\right)_{\text{SRM-312a}}} - 1 \right] \times 10^3$$  (3)

where $x$ represent 50, 53 and 54. During measurements, interferences of $^{50}$Ti, $^{50}$V and $^{54}$Fe on $^{50}$Cr and $^{54}$Cr were monitored by measuring $^{49}$Ti, $^{51}$V and $^{56}$Fe signals. The interference of $^{50}$V on $^{50}$Cr was typically <1 ppm of signal strength, whereas those of $^{50}$Ti on $^{50}$Cr and $^{54}$Fe on $^{54}$Cr were one or two orders of magnitude larger, respectively. Corrections of <10$^4$ ppm on $^{52}$Fe yielded accurate and reproducible $\varepsilon^{54}$Cr values (Schiller et al., 2014). This resulted in an external reproducibility of 28 ppm on $\delta^{50}$Cr.

4. RESULTS

4.1. Compositions of DaG 319 clasts

4.1.1. Main Group (MG) and Hughes Type (HT) clasts

Thick sections of the DaG 319 meteorite are dominated (~90 vol.%) by mm-sized olivine and pyroxene grains similar to monomict ureilites (Fig. 1a). Pyroxenes are present in lower abundances and are generally smaller (<1 mm). These MG clasts are typically mono-mineralic crystals interpersed in a fine-grained groundmass consisting of smaller ureilite clasts, sulfides, FeNi-metal, dark clasts and carbon-rich matter. Some MG clasts also occur as aggregates of olivine and pyroxene. Olivine clasts typically have relatively thin rims (<100 μm) that are Mg-poor with respect to their cores and most clasts contain veins of carbon-rich matter (see Electronic Annex). Main Group clasts have variable magnesium numbers (Mg# = Mg/(Fe + Mg)) ranging from 0.74–0.91, but have constant Mn/Mg ratios (Fig. 2; Electronic Annex, Tables S2 and S3). Main Group pyroxenes and olivines fall on different slopes in Fe/Mg versus Fe/ Mn space, as shown previously by Goodrich and Delaney (2000). In contrast, the HT clasts in these sections (~10 vol.%) are generally more Mg-rich and have a very narrow range of Mg#s (~0.89; Electronic Annex, Table S1). Hughes Type clasts are named for their resemblance to unusual ureilites similar to Hughes 009, since the Mn/Mg ratios of HT olivines are generally offset from the linear correlation determined for MG clasts and correspond to Hughes Type compositions (Fig. 2; Downes et al., 2008). However, HT clasts are distinct from Hughes Type ureilites in that their pyroxene Mn/Mg ratios fall on the trend for MG pyroxenes and are, thus, offset from the field characterizing pyroxenes from Hughes Type ureilites. In addition, Hughes Type clasts contain 7–52 vol.% augite and 23–47 vol.% olivine (Goodrich et al., 2001), whereas HT clasts from DaG 319 contain relatively low-Ca pyroxenes and olivine contents of ~40 vol.% (Electronic Annex, Table S1 and Fig. S4). Hughes Type clasts are typically mm- to cm-sized and contain fairly euhedral, submm olivine (Fig. 1b) surrounding mostly anhedral pyroxenes that are enriched in Al compared to MG clasts pyroxenes (typically 1.2–1.3 wt % Al$_2$O$_3$; Electronic Annex, Tables S2 and S3). Olivine and pyroxene grains show triple junctions with 120° angles and have no preferred orientation.

4.1.2. Feldspathic clasts

A detailed classification and description of the feldspathic (plagioclase-bearing) clasts in DaG 319 has been previously reported by Ikeda and Prinz (2001) and Ikeda et al. (2003). Their classification is based on texture and further divides feldspathic clasts into porphyritic, pilotaxitic, glassy and gabbroic types. Furthermore, Cohen et al. (2004) defined multiple populations of feldspathic clasts (albitic, labradoritic, olivine-augite) based on their mineralogy and texture. Most of the feldspathic clasts analyzed here are albitic (n = 8), one sample is labradoritic and three samples have an olivine-augite composition (Fig. 1, see also Electronic Annex). Albitic clasts have varying porphyritic textures: (1) Three of the clasts contain a groundmass of albitic feldspar (An < 5) with embedded needle-shaped enstatite crystals (Fig. 1c), (2) four clasts are dominated by albite with typically minor anhedral pigeonite-augite phenocrysts (Fig. 1d) and (3) one clast contains relatively large (~100 μm) euhedral phenocrysts of enstatite and albite in a pilotaxitic matrix of pyroxene with plagioclase needles (Fig. 1e). The labradoritic clast is dominated by enstatite phenocrystals in a labradoritic groundmass with An ~70 (Fig. 1f). Two out of three olivine-augite feldspathic clasts have a porphyritic texture with forsterite and augite phenocrysts embedded in an albitic groundmass (Fig. 1g), whereas one clast is dominated by augite and contains minor forsterite and albite (Fig. 1h). Similar to MG and HT clasts, olivine-augite and labradoritic clasts are characterized by pyroxene with high Mg#s between 0.8 and 1.0 (Electronic Annex, Table S4), whereas albitic clasts
have pyroxenes with intermediate Mg#s (0.5–0.7). All pyroxenes in feldspathic clasts have variable Mn/Mg ratios and fall on a trend determined for fractional crystallization from the main ureilite body (Fig. 2). Of these clasts, two clasts show shock-related textures: clasts BSa3 and TR3 contain melt pockets between albite grains with augite compositions. Clast RT3 contains a granoblastic texture of porphyritic augite and olivine, which is indicative of recrystallization during thermal overprinting (Electronic Annex, Fig. S8).

Fig. 1. Compilation of back scattered electron images from various clast types in the DaG 319 polymict ureilite, with (a) typical distribution of clast types, with mostly main group/monomict (MG) clasts of olivine (ol) and more rare pyroxene (px), where the in white lineated clasts (WB2, 3, 4) are sampled for Mg- and Cr-isotope analyses; (b) WB1: olivine-orthopyroxene (HT) clast; (c) RT1a (like RT1b, BSb1): albitic clast with enstatite needles; (d) BSa2: albitic clast with augite phenocrysts; (e) DUO2-3: Albitic clast with albite (Ab) and enstatite (En) phenocrysts; (f) WB2: labradoritic clast with En and anorthite (An); (g) RT3 (like BSa3, TR3): olivine-augite feldspathic clast in albitic groundmass; (h) DUO1-2: olivine-augite clast with minor albite; (i) LC6: Dark clast with sulfides, magnetite and carbonates in a phyllosilicate matrix and (j) TR7: chondrule-containing clast.
Compositions than Earth’s mantle, with lower Mg-isotope compositions for all measured DaG 319 samples are presented in Table 2 and Fig. 3. The difference in 26Mg values for MG clasts and HT clasts, we calculate the effects of kinetic and equilibrium isotope fractionation on the 26Mg values of these clasts (Table 2). We show that correcting the Mg-isotope data by equilibrium (µ26Mg_eq) instead of kinetic (µ26Mg_kin) fractionation law has little effect on the µ26Mg values for MG clasts (µ26Mg_eq = -10.6 ± 5.4 ppm), but increases the 26Mg for HT clasts (µ26Mg_eq = 8.0 ± 5.0 ppm) and significantly lowers the spread in data as indicated by a reduction in MSWD values from 15 for µ26Mg_kin to 4.7 for µ26Mg_eq. This suggests that mass-dependent Mg-isotope fractionation exhibited by HT clasts is best explained by equilibrium rather than kinetic law, since we expect no variation in µ26Mg values for clasts that appear to be co-genetic based on very constant Mg#s and Mn/Mg ratios and a narrow range of µ26Mg values.

Labradoritic and olivine-augite feldspathic clasts (n = 4) have 27Al/26Mg ratios between 0.077 and 0.146 and have varying µ26Mg_eq values ranging between -8.9 ± 2.4 ppm and 8.5 ± 1.3 ppm (Fig. 4). Albitic clasts (n = 6) have higher 27Al/26Mg ratios ranging from 0.6 to 5.3 with µ26Mg eq excesses between -8.1 ± 2.2 ppm and 24.9 ± 3.5 ppm. Unlike HT clasts, we cannot point to a superior isotope fractionation law for feldspathic clasts (e.g., equilibrium or kinetic mass-bias corrected µ26Mg), since the spread in µ26Mg values is at least in part related to variations in 27Al/26Mg.

The four analyzed dark clasts have µ26Mg values ranging between 4.3 ± 2.0 ppm and 8.6 ± 1.2 ppm, with an average of µ26Mg = 6.2 ± 3.6 ppm (2SD). These values are comparable with CI-type carbonaceous chondrites (Larsen et al., 2011). The chondrule-containing clasts have µ26Mg values of −3.8 ± 3.0 ppm and −5.2 ± 3.3 ppm for TR7 and BSa4, respectively. Their Mg-isotope compositions suggest a link to ordinary chondrites (Larsen et al., 2011).

4.3. Chromium isotopes

4.3.1. Mass-dependent Cr-isotope compositions

The 50Cr/52Cr ratios of MG and HT clasts were obtained by MC-ICPMS and are presented in Table 3.
and Fig. 5. The accuracy and reproducibility of this method has been tested by repeat analyses of the dunite DTS-2b rock standard, yielding a $\delta^{50}\text{Cr}$ value of 0.314 ± 0.006. This is in good agreement with the value of 0.320 ± 0.096 (2SD) for DTS-2b obtained by Schiller et al. (2014). Analyses of MG and HT clasts show that there is a resolvable Cr-isotope difference of ~190 ppm between the two group averages (Fig. 5), where $\delta^{50}\text{Cr}$ is 0.140 ± 0.090 (2SD) for five MG clasts and 0.331 ± 0.080 (2SD) for HT clasts. Thus, in contrast to the $\delta^{25}\text{Mg}$ values, the $\delta^{50}\text{Cr}$ values indicate a lighter stable isotope composition for HT clasts relative to MG clasts.

4.3.2. $\varepsilon^{54}\text{Cr}$ values

Cr-isotope data obtained by TIMS are presented in Table 4 and Fig. 6. We analyzed four MG clasts that have $\varepsilon^{54}\text{Cr}$ values ranging between $-1.10 \pm 0.12$ and $-0.74 \pm 0.07$ (Fig. 6b). This data range is confirmed by MC-ICPMS data from BSa5 and five additional MG clasts with $\varepsilon^{54}\text{Cr}$ values between $-1.28 \pm 0.29$ and $-0.74 \pm 0.05$ (Table 3). The average $\varepsilon^{54}\text{Cr}$ value of $-0.94 \pm 0.11$ (2SE) for MG clasts is in agreement with literature data for monomict ureilites [$\varepsilon^{54}\text{Cr} = -1.08 \pm 0.25$ (2SE), Larsen et al., 2011; $\varepsilon^{54}\text{Cr} = -0.92 \pm 0.02$ (2SE), Yamakawa et al., 2010], but is $^{54}\text{Cr}$-depleted relative to multiple ureilite fragments from the polymict ureilite Almahata Sitta [$\varepsilon^{54}\text{Cr} = -0.77 \pm 0.03$ (2SE), Qin et al., 2010b]. The HT clasts have overall more positive $\varepsilon^{54}\text{Cr}$ values than the MG clasts and, with an average of $-0.79 \pm 0.10$ (2SE), are thus closer in composition to Almahata Sitta. The $\varepsilon^{54}\text{Cr}$ data for HT individual clasts as well as their average [$\varepsilon^{54}\text{Cr} = -0.79 \pm 0.14$ (2SE)] obtained by TIMS are reproduced by MC-ICPMS, highlighting the accuracy between both approaches and suggesting an offset in $\varepsilon^{54}\text{Cr}$ between MG and HT clasts of up to 15 ppm. In Fig. 7a, we plot $\varepsilon^{54}\text{Cr}$ values of MG and HT analyses against their $\delta^{50}\text{Cr}$ values, in which we show that no significant correlation exists. If anything, $\delta^{50}\text{Cr}$ seems to increase with $\varepsilon^{54}\text{Cr}$, an effect that is opposite to what is expected from equilibrium isotope fractionation-related $\varepsilon^{54}\text{Cr}$ excesses (Fig. 7b). However, recalculating HT clast $\varepsilon^{54}\text{Cr}$ values with an equilibrium law (in a similar approach to Mg-isotopes) reduces
the 2SD value to 0.22 as opposed to kinetic law yielding a 2SD of 0.35. Similar to the mass-dependent Mg-isotope data, this reduction in scatter suggests that the HT-related reservoir is affected by equilibrium isotope fractionation of Cr.

The analyzed feldspathic clasts show relatively large variations in $\varepsilon^{54}$Cr (Table 4). Their range spans from $-1.18 \pm 0.11$, similar to MG clasts, up to $1.16 \pm 0.5$ for clast TR3. The bulk of feldspathic clasts have $\varepsilon^{54}$Cr values between $-1.18 \pm 0.11$ and $-0.54 \pm 0.15$. We note that the anomalously high $\varepsilon^{54}$Cr value of clast TR3 may be related to its shock-melting history (see Electronic Annex). The small size of the feldspathic clasts (<500 ng Cr) did not allow for repeat analyses by MC-ICPMS. The dark clasts from DaG 319 range from $\varepsilon^{54}$Cr = 1.30 ± 0.12 to 1.86 ± 0.11, averaging around the CI carbonaceous chondrite value of 1.56 ± 0.13 (2SD, Shukolyukov and Lugmair, 2006; Trinquier et al., 2007; Qin et al., 2010a; Van Kooten et al., 2016). Combined with their $\mu^{56}$Mg composition, this suggests that the dark clasts are closely related to CI chondrites. The chondrule-containing clasts have $\varepsilon^{54}$Cr values of $-0.09 \pm 0.15$ and 0.40 ± 0.40 (Table 4) corresponding to ordinary and enstatite chondrites with average bulk values of $-0.34 \pm 0.14$ and +0.08 ± 0.10, respectively (Trinquier et al., 2007; Qin et al., 2010b).

4.3.3. $^{53}$Mn-$^{53}$Cr systematics

In Fig. 6a, we present $\varepsilon^{53}$Cr data from MG and HT clasts together with literature data from whole rock monomict ureilites (Yamakawa et al., 2010) and Almahata Sitta fragments (Qin et al., 2010b). Main Group clasts have an average $\varepsilon^{53}$Cr composition of 0.13 ± 0.14 (2SD), which is similar to monomict ureilites. This average value is confirmed by MC-ICPMS analyses on BSa5 and five additional MG clasts (Table 3; $\varepsilon^{53}$Cr = 0.12 ± 0.21 (2SD)). Corresponding $^{55}$Mn/$^{52}$Cr ratios of MG clasts range between 0.5 and 0.8, in agreement with data from Yamakawa et al. (2010) for monomict ureilites. In contrast, HT clasts have higher $^{55}$Mn/$^{52}$Cr ratios of ~1.2 and $\varepsilon^{53}$Cr values that average 0.43 ± 0.06 (2SD). Almahata Sitta fragments consist of lithologies that are similar to both MG clasts and HT clasts (Zolensky et al., 2010) and have $\varepsilon^{53}$Cr values and corresponding $^{55}$Mn/$^{52}$Cr ratios that fall between monomict ureilites reported by Yamakawa et al. (2010) and HT clasts in this work (Fig. 8). The $\varepsilon^{53}$Cr data for individual feldspathic clasts varies from 0.07 ± 0.06 to 1.31 ± 0.27 (2SE). In Fig. 8, we present $\varepsilon^{53}$Cr versus $^{55}$Mn/$^{52}$Cr data of MG clasts and HT clasts, as well as data from Yamakawa et al. (2010) and Qin et al. (2010b). Our results from sampled main group olivine and pyroxene cores (MG clasts) plot on a slope equivalent to $(^{55}$Mn/$^{55}$Mn)$_0 = (6.8$

<table>
<thead>
<tr>
<th>Mass-dependent Cr-isotope data and mass-bias corrected $\varepsilon^{53}$Cr and $\varepsilon^{54}$Cr data for ureilite clasts measured by MC-ICPMS. Internal errors are presented as 2SE.</th>
<th>$^{55}$Mn/$^{52}$Cr</th>
<th>$\varepsilon^{54}$Cr</th>
<th>$\varepsilon^{53}$Cr</th>
<th>$\delta^{50}$Cr</th>
</tr>
</thead>
<tbody>
<tr>
<td>Terrestrial standards</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DTS-2b (3)</td>
<td>0.04 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.314 ± 0.006</td>
<td></td>
</tr>
<tr>
<td>Main group clasts</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BSa5 (olpx)</td>
<td>0.5570</td>
<td>-0.74 ± 0.06</td>
<td>0.09 ± 0.07</td>
<td>0.369 ± 0.027</td>
</tr>
<tr>
<td>BSbpx (px)</td>
<td>0.4978</td>
<td>-1.28 ± 0.29</td>
<td>-0.05 ± 0.10</td>
<td>0.130 ± 0.015</td>
</tr>
<tr>
<td>DUO1px (px)</td>
<td>0.7200</td>
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<td>0.14 ± 0.10</td>
<td>0.155 ± 0.027</td>
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<td>Bsapx (px)</td>
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<td>0.19 ± 0.10</td>
<td>0.072 ± 0.024</td>
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<tr>
<td>DUO1ol (ol)</td>
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<td>0.22 ± 0.05</td>
<td>0.148 ± 0.005</td>
</tr>
<tr>
<td>WBO7 (ol)</td>
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<td>-0.79 ± 0.14</td>
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<td>0.196 ± 0.017</td>
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<tr>
<td>Average</td>
<td>0.6311</td>
<td>-0.95 ± 0.39</td>
<td>0.12 ± 0.21</td>
<td>0.140 ± 0.090</td>
</tr>
<tr>
<td>Hughes type clasts</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DUO1-1 (olpx)</td>
<td>1.1578</td>
<td>-0.83 ± 0.16</td>
<td>0.35 ± 0.55</td>
<td>0.353 ± 0.032</td>
</tr>
<tr>
<td>DUO2-1 (olpx)</td>
<td>1.1732</td>
<td></td>
<td>0.30 ± 0.15</td>
<td>0.292 ± 0.016</td>
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<tr>
<td>RT2 (olpx)</td>
<td>1.1116</td>
<td>-0.66 ± 0.23</td>
<td>0.54 ± 0.12</td>
<td>0.303 ± 0.029</td>
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<tr>
<td>TR4 (olpx)</td>
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<td>-0.90 ± 0.15</td>
<td>0.42 ± 0.08</td>
<td>0.376 ± 0.010</td>
</tr>
<tr>
<td>Average</td>
<td>1.1561</td>
<td>-0.79 ± 0.25</td>
<td>0.40 ± 0.21</td>
<td>0.331 ± 0.080</td>
</tr>
</tbody>
</table>
Table 4  
$\epsilon^{53}$Cr and $\epsilon^{54}$Cr values of ureilite clasts measured by TIMS. $N$ is the amount of analyses done for each sample. Internal errors for individual clasts are given in 2SE. Average values are presented with a 2SD error.

<table>
<thead>
<tr>
<th>$^{53}$Mn/$^{52}$Cr</th>
<th>$\epsilon^{54}$Cr</th>
<th>$\epsilon^{53}$Cr</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>DTS-2b (2)</td>
<td>-0.03 ± 0.13</td>
<td>0.06 ± 0.05</td>
<td>28</td>
</tr>
<tr>
<td>Dark clasts</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>LC2b</td>
<td>0.7877</td>
<td>1.44 ± 0.18</td>
<td>16</td>
</tr>
<tr>
<td>LC3</td>
<td>0.6979</td>
<td>1.30 ± 0.12</td>
<td>14</td>
</tr>
<tr>
<td>LC4</td>
<td>0.6922</td>
<td>1.58 ± 0.18</td>
<td>10</td>
</tr>
<tr>
<td>LC6</td>
<td>0.5790</td>
<td>1.86 ± 0.11</td>
<td>14</td>
</tr>
<tr>
<td>Average</td>
<td>1.55 ± 0.48</td>
<td>0.00 ± 0.20</td>
<td></td>
</tr>
<tr>
<td>Chondritic clasts</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>TR7</td>
<td>0.6103</td>
<td>-0.09 ± 0.15</td>
<td>10</td>
</tr>
<tr>
<td>BSa4</td>
<td>0.7135</td>
<td>0.40 ± 0.40</td>
<td>5</td>
</tr>
<tr>
<td>Main group clasts</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>WB3 (px)</td>
<td>0.5069</td>
<td>-1.10 ± 0.12</td>
<td>14</td>
</tr>
<tr>
<td>WB4 (ol)</td>
<td>0.8000</td>
<td>-0.90 ± 0.18</td>
<td>15</td>
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<tr>
<td>WB5 (ol)</td>
<td>0.7044</td>
<td>-0.98 ± 0.13</td>
<td>16</td>
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<tr>
<td>BSa5</td>
<td>0.5570</td>
<td>-0.74 ± 0.07</td>
<td>16</td>
</tr>
<tr>
<td>Average</td>
<td>-0.93 ± 0.30</td>
<td>0.13 ± 0.14</td>
<td></td>
</tr>
<tr>
<td>Hughes type clasts</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>WB1</td>
<td>1.1797</td>
<td>-0.86 ± 0.08</td>
<td>14</td>
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<td>DUO1-1</td>
<td>1.1578</td>
<td>-0.89 ± 0.11</td>
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</tr>
<tr>
<td>DUO2-1</td>
<td>1.1732</td>
<td>-0.83 ± 0.09</td>
<td>16</td>
</tr>
<tr>
<td>RT2</td>
<td>1.1116</td>
<td>-0.61 ± 0.09</td>
<td>11</td>
</tr>
<tr>
<td>TR4</td>
<td>1.1818</td>
<td>-0.75 ± 0.11</td>
<td>16</td>
</tr>
<tr>
<td>Average</td>
<td>-0.79 ± 0.22</td>
<td>0.43 ± 0.06</td>
<td></td>
</tr>
<tr>
<td>Feldspathic clasts</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DUO2-3 (Ab)</td>
<td>3.9447</td>
<td>-0.72 ± 0.08</td>
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<tr>
<td>BSa2 (Ab)</td>
<td>1.0117</td>
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<td>BSb1 (Ab)</td>
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<td>RT1b (Ab)</td>
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<tr>
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<td>8</td>
</tr>
<tr>
<td>DUO1-2 (OA)</td>
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<td>-0.91 ± 0.15</td>
<td>8</td>
</tr>
<tr>
<td>RT3 (OA)</td>
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<td>0.02 ± 0.14</td>
<td>6</td>
</tr>
<tr>
<td>BSa3 (OA)</td>
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<td>-0.56 ± 0.25</td>
<td>8</td>
</tr>
<tr>
<td>TR3 (Ab)</td>
<td>1.4400</td>
<td>1.16 ± 0.50</td>
<td>3</td>
</tr>
<tr>
<td>Average</td>
<td>-0.76 ± 1.52</td>
<td>0.70 ± 1.08</td>
<td></td>
</tr>
</tbody>
</table>

5. DISCUSSION

5.1. Origin of the Mg mass-dependent isotope fractionation

Our $\mu^{25}$Mg data indicate a significant amount of mass-dependent Mg-isotope fractionation between various ureilite clasts that translates into a range of $\sim 450$ ppm. Feldspathic clasts display a wide range of isotopically light Mg compositions ($\mu^{25}$Mg = $-305 \pm 25$ ppm to +15 ± 12 ppm), whereas HT clasts consistently record heavy Mg-isotope signatures ($\mu^{25}$Mg = 157 ± 27 ppm). Main Group clasts are intermediate between these groups with $\mu^{25}$Mg values between $-50 \pm 14$ ppm and $-11 \pm 7$ ppm. We discuss below potential mechanisms that can account for the observed mass-dependent Mg-isotope fractionation.

5.1.1. Initial Mg-isotope heterogeneity

An important mechanism to discuss is the possibility that all variation is caused by initial heterogeneity of mass-dependent Mg-isotopes during accretion of polymict ureilites or of the initial UPB. This would either suggest that the clasts are not genetically related to each other or that, similarly to oxygen isotopes, this initial heterogeneity was preserved during parent body processes (Clayton and Mayeda, 1988; Clayton and Mayeda, 1996). However, the total amount of variation in $\mu^{25}$Mg values found for Solar System planetary reservoirs is approximately 100 ppm (Larsen et al., 2016b), suggesting that the relatively large variation found in ureilite clasts cannot be accounted for by heterogeneity alone.

5.1.2. Equilibrium isotope fractionation

Partial melting and/or fractional crystallization can cause equilibrium isotope fractionation for which the partition coefficient is dependent on the relative difference in coordination numbers of the element of interest in both minerals and melts (Schauble, 2011). In the case of magnesium isotopes, tetrahedral sites in minerals are preferentially occupied by the heavier isotopes, whereas light isotopes prefer the octahedral sites. For example, spinel with Mg occupying tetrahedral sites has a higher $\mu^{25}$Mg value than co-genetic silicates like olivine or pyroxene with Mg occupying octahedral sites. Sclerolite melts are thought to contain a mixture of tetrahedral, pentahedral and octahedral sites that is dependent on their composition (Wilding et al., 2004; Henderson et al., 2006) and should, thus, be isotopically heavier than co-genetic silicates (Schauble, 2011). Although for terrestrial reservoirs (e.g., mid-ocean ridge basalts, ocean island basalts and Earth’s mantle), the amount of Mg equilibrium fractionation during differentiation appears to be small (Schiller et al., 2010; Teng et al., 2007, 2010; Huang et al., 2011), mass-dependent Mg-isotope fractionation has been observed during magma differentiation on asteroids (Schiller et al., 2017). Thus, if feldspathic clasts are partial melts with relatively high Mg coordination numbers, these clasts should be isotopically heavier than their residues. In contrast to this
prediction, the clasts are lighter than main ureilite body clasts by up to 250 ppm and lighter than HT clasts by up to 450 ppm.

A possible solution to this apparent contradiction is the presence of a magnesite component in the partial melt with a low Mg coordination number, which should be isotopically light relative to co-genetic silicates (Schauble, 2011; Yang et al., 2012). As a result, the isotope effect is opposite to that of ‘normal’ equilibrium isotope fractionation such that the magnesite-containing melt will be depleted in the heavy Mg-isotopes relative to the residue. The stability of magnesite in the ureilite mantle is dependent on pressure, temperature and oxygen fugacity. At lower pressures and temperatures, oxidized carbon becomes stable over pure carbon phases such as graphite over a larger range of oxygen fugacities (Goodrich et al., 2007; Stagno and Frost, 2010; Stagno et al., 2011). We show in Fig. 9 the stability of graphite in a pressure versus fO2 diagram, where with increasing temperature, graphite is stable over CO + CO2 at lower pressures. The modeled peak temperature for equilibrated ureilite residues (1276 °C, Goodrich et al., 2007) results in the stability of CO and CO2 over carbon in the upper ~5 km of the ureilite mantle. In Earth’s mantle, the C—CO + CO2 equilibrium line defines the transition between graphite and magnesite, or between diamond and magnesite in the lower mantle (Stagno and Frost, 2010; Stagno et al., 2011). Hence, we show in Fig. 9 that the upper ureilite mantle may be characterized by the presence of oxidized carbon as a stable mantle phase, whereas in the lower mantle carbon is reduced to graphite. Pure carbon phases are so far observed in ureilite mantle residues defined as ‘type I ureilite clasts’ (Ikeda et al., 2000; Kita et al., 2004) or MG clasts in this work, whereas carbon veins are absent in crystals from ‘type II ureilite clasts’ or HT clasts in this work. Hence, MG clasts with their range of Mg#s, may represent smelting-related residues derived from varying depths in the stability regime of pure carbon, whereas HT clasts may be derived from shallower mantle regions compatible with magnesite. We emphasize that the smelting process is not necessary to invoke separate stability regimes for carbon and magnesite.

To discuss the potential presence of magnesite in HT clasts it is relevant to first consider the magmatic origin of HT clasts, which represent either cumulates or residues from the ureilite mantle. They can be linked through Fe/Mg versus Mn/Mg ratios of their olivines and pyroxenes to the unusual Hughes Type ureilites (Fig. 2), which are suggested to represent cumulates (Downes et al., 2008). However, unlike Hughes Type ureilites, HT clasts have a higher abundance of olivine (>40 vol.%) and do not contain high-Ca pyroxenes such as augite. On Earth, such peridotite compositions (e.g., harzburgites) are rarely associated with cumulate origins. Furthermore, HT clasts do not show a characteristic crystal alignment expected for cumulates (Electronic Annex, Fig. S4). Collectively, the petrological features of HT clasts indicate that they are residues, although their mixed characteristics suggest that they may have experienced multiple differentiation events and may have primarily formed as cumulates. We interpret that the low carbon abundances (Goodrich et al., 2001; Ikeda and Prinz, 2001; Kita et al., 2004) and high $^{25}$Mg values of HT clasts relative to MG clasts suggest that HT clasts represent cumulates from which a isotopically light magnesite melt component was extracted. These low $^{25}$Mg values are recorded by the feldspathic clasts. It is possible to roughly estimate the amount of magnesite necessary to produce the isotopic differences observed between HT clasts and feldspathic clasts, since similar effects are observed on Earth. Terrestrial basalts from the North China craton are found to be on average 150 ppm lighter for $^{25}$Mg than their source mantle peridotites (Yang et al., 2012). This is attributed to a 20% dolomitic melt derived from carbonated...
mantle material, where dolomite has on average 20 wt.% MgO and $\delta^{26}\text{Mg} = 2.2$ ‰ (range between 1.09 ‰ and 5.5 ‰). Magnesite is more Mg-rich than dolomite with MgO = 48 wt.% and is predicted to be isotopically lighter than dolomite by $\delta^{24}\text{Mg} = 5\%$ at 1000 °C (Schauble, 2011). Hence, a lower percentage of magnesite-melt is required to achieve the same offset: ~8% magnesite for an offset of 150 ppm. Since the offset between HT clasts and feldspathic clasts is on average 300 ppm, we estimate this would require on average ~16% of the total melt to be magnesite. This is, however, dependent on the equilibrium temperature and Mg-isotope composition of the magnesite.

The $^{53}\text{Mn}/^{52}\text{Cr}$ systematics of HT clasts agree with our suggestion that these clasts represent mantle rocks that...
5.1.3. Disequilibrium isotope fractionation

Mass-dependent Mg-isotope variations as seen in feldspathic clasts can also be the result of disequilibrium isotope fractionation during magmatic processes. For example, experienced multiple differentiation events, since HT clasts intersect with MG clasts as well as with feldspathic clast isochrons (Fig. 8, see also Section 5.4). Their petrological characteristics suggest that they may have initially formed as cumulates as implied from their link to Hughes type ureilites (Fig. 2), but experienced remelting at a later stage, depleting their composition in high-Ca pyroxene. Partial melts from depleted sources are in agreement with trace element compositions of feldspathic clasts (Kita et al., 2004).

For a partial melt extracted from a magnesite-rich source, it is unclear whether this magnesite would have been enriched in trace elements such as Ba, U and Sr, that are normally compatible with carbonates (Ionov, 1996). Nevertheless, the Almahata Sitta trachyandesite shows enrichments in Ba, U and Sr relative to CI chondrites (Bischoff et al., 2014), which do not exclude the presence of magnesite in the source. In addition, silica-rich melts as well as alkali basalts similar to the Almahata Sitta and feldspathic clast composition are frequently associated with carbonate melts (Woolley and Kjarsgaard, 2008; Zhang et al., 2017). If indeed feldspathic clasts are melts from depleted sources represented by HT clasts during a secondary differentiation event, the presence of magnesite in cumulates required for the Mg-isotope signatures of feldspathic clasts needs to be explained. Since magnesite would be incompatible in the melt, we expect no magnesite present in cumulates formed during melting of the ureilite mantle. However, if we assume a smelting model for the ureilite mantle (Goodrich et al., 1987, 2007; Warren and Kelleineym, 1992; Walker and Grove, 1993; Singleton and Grove, 2003), CO and CO₂ gases released during smelting will react with forsterite and water to form magnesite (Kelemen et al., 2011) in the upper 5 km of the ureilite mantle (Fig. 9). The presence of water in this reaction increases the carbonation, but is not required to form magnesite (Dabirian et al., 2012). Nevertheless, some authors argue for the presence of water on the UPB (Goodrich et al., 2015).

5.1.3. Disequilibrium isotope fractionation

Mass-dependent Mg-isotope variations as seen in feldspathic clasts can also be the result of disequilibrium isotope fractionation during magmatic processes. For example,
be enriched in the light isotopes (Richter et al., 2008; Richter et al., 2009) and result in significantly larger Mg-isotope fractionation than observed from equilibrium fractionation. For example, Xiao et al. (2013) find $\mu^{25}$Mg differences of $\sim$200 ppm between olivine and orthopyroxene in lherzolites, caused by consumption of orthopyroxene that is replaced by clinopyroxene during melting. Disequilibrium fractional melting was previously suggested to explain REE patterns in ureilites (Goodrich et al., 2007). However, Barrat et al. (2016b) propose from a lithophile trace element study of ureilites that disequilibrium melting is unlikely. Furthermore, both our Mg and Cr mass-dependent isotope data support equilibrium isotope effects for HT clasts (see Section 4.2.2 and 4.3.2).

5.2. Implications for mass-dependent Cr-isotope fractionation

In addition to a mass-dependent Mg-isotope offset, a significant difference in $\delta^{56}$Cr of 190 ppm exists between lighter HT and heavier MG clasts. We considered equilibrium isotope fractionation by melting in the presence of magnesite as a viable mechanism to create HT clasts heavy in Mg-isotopes. Due to the lack of Cr incorporation in magnesite, this process does not affect Cr-isotope fractionation. However, in agreement with the isotopically light HT clasts, equilibrium isotope fractionation will enrich the melt in heavy Cr-isotopes during partial melting, whereas the residue/cumulate is isotopically light (Mouyier et al., 2011). The magnitude of mass-dependent Cr-isotope fractionation is most likely controlled by the oxidation states of Cr (Schönberg et al., 2008; Schönberg et al., 2016). For example, the amount of fractionation observed in terrestrial mantle and crust derived rocks is $<0.2\%$/amu (Schönberg et al., 2016), which is larger than the mass-dependent isotopic variation observed between ureilite mantle residues even though the variability in oxidation states of Earth’s mantle is thought to be significantly lower than that of the ureilite mantle (Papike et al., 2005; Goodrich et al., 2013; Bonnand et al., 2016). Hence, the amount of mass-dependent isotope fractionation between MG and HT clasts can be explained by differences in oxidation states within the ureilite mantle.

Collectively, the combined Mg and Cr mass-dependent isotope data suggest that equilibrium fractionation is the most likely mechanism involved in the differentiation of HT clasts, which will be summarized below. As we discuss in Section 5.1, two processes can result in a light $\mu^{25}$Mg composition for feldspathic clasts (1) equilibrium fractionation by melting a magnesite component that is isotopically light and (2) disequilibrium isotope fractionation between residue/cumulate and partial melt. If HT clasts represent cumulates of a partial melt from main group ureilites, which has been inferred based on the Mn/Mg relationship between HT clasts and Hughes type ureilites (Downes et al., 2008), equilibrium fractionation will result in a melt with isotopically heavy Cr- and Mg-isotope compositions relative to the cumulates (Schauble, 2011). Kinetic isotope fractionation, however, will result in heavy Cr- and Mg-isotope compositions for the HT clasts relative to the melt. We find that the HT clasts have high $\delta^{49}$Cr as well as $\mu^{25}$Mg values, which we interpret as reflecting extraction of the feldspathic melt in the presence of oxidized carbon (i.e., magnesite) during equilibrium isotope fractionation.

5.3. Source of the $^{54}$Cr heterogeneity in the ureilite parent body

If the variability in $\varepsilon^{54}$Cr-isotope compositions is the result of equilibrium isotope fractionation, we expect an increase of $\varepsilon^{54}$Cr correlated with a decrease of $\delta^{50}$Cr (Fig. 7b). Hence, MG clasts with lower $\delta^{50}$Cr values should be depleted in $^{54}$Cr relative to HT clasts. In contrast, we find an opposite trend, suggesting that the $\varepsilon^{54}$Cr variation in ureilites is unrelated to mass-dependent isotope fractionation processes. The presence of variable mass-independent $^{54}$Cr abundances in meteorites and their components have been linked to different accretion regions in the protoplanetary disk (Trinquier et al., 2007, 2009; Larsen et al., 2011; Olsen et al., 2016). In particular, carbonaceous chondrites have variable, positive $\varepsilon^{54}$Cr values with respect to terrestrial planets. On a planetesimal scale, $\varepsilon^{54}$Cr values of distinct bulk Solar System objects are fairly constant, whereas individual components of primitive meteorites (e.g., chondrules, the potential building blocks of planets, Johansen et al., 2015) can exhibit large variations in $\varepsilon^{54}$Cr (Van Kooten et al., 2016; Olsen et al., 2016). Differentiation on planetary bodies such as the UPB can homogenize initial isotopic heterogeneity present in its precursor material. Oxygen isotope signatures from ureilite mantle material still exhibit large mass-independent isotope variability with a total range of $\sim$2‰ on the $\Delta$17O (Clayton and Mayeda, 1988), suggesting that initial heterogeneity of oxygen isotopes was not erased during parent body processes on the UPB. Similarly, the range in $\varepsilon^{54}$Cr values measured in HT and MG clasts ($\Delta\varepsilon^{54}$Cr$_{max} = \sim$65 ppm), as well as the more variable feldspathic clasts, may also represent primordial $^{54}$Cr heterogeneity. However, Cr in olivine diffuses faster than oxygen by at least two orders of magnitude at similar temperatures and oxygen fugacities (Ryerson et al., 1989; Dohmen et al., 2002; Ito and Ganguly, 2006; Chakraborty, 2010) and is, thus, much more susceptible to isotopic homogenization than oxygen. Hence, preserving isotope heterogeneity of $^{54}$Cr in the ureilite mantle during its differentiation may prove more difficult than maintaining oxygen isotope heterogeneity. A potentially more viable mechanism to create $^{54}$Cr variability between different ureilite mantle clasts is late admixing of $^{54}$Cr-rich material to the upper ureilite mantle, perhaps via impacts of material similar to carbonaceous chondrites. This would likely affect impact-derived melts to a larger degree, thereby explaining the $^{54}$Cr enrichments in HT and feldspathic clasts relative to MG clasts. The addition of carbonaceous chondrite material to the UPB can potentially explain several other features of monomict and polymict ureilites outlined in Section 5.6.

5.4. Chronological significance of $^{53}$Mn-$^{53}$Cr data

5.4.1. $^{53}$Mn-$^{53}$Cr data

To investigate the potential chronological relationships between individual ureilite components, we plot $\varepsilon^{53}$Cr
versus $^{55}\text{Mn}/^{52}\text{Cr}$ values for ureilites (Fig. 8), where we compare HT, MG and feldspathic clasts to Almahata Sitta ureilite fragments (Qin et al., 2010b) and bulk monomict ureilites (Yamakawa et al., 2010). Isochron data referred to in this section can be found in Table 5. Here, all calculated absolute ages are anchored to the D’Orbigny angrite with an absolute age of 4563.59 ± 0.20 Ma (Amelin, 2008) and a $(^{53}\text{Mn}/^{52}\text{Mn})_0$ ratio of (3.24 ± 0.04) $\times 10^{-6}$ (Glavin et al., 2004). MG clasts represent different olivine and pyroxene cores similar to main group ureilite compositions. Seven MG clasts form an isochron with $(^{53}\text{Mn}/^{52}\text{Mn})_0 = (6.79 ± 2.00) \times 10^{-6}$, corresponding to an absolute age of 4567.5 ± 1.9 Ma and within error of the timing of CV CAI formation at 4567.3 ± 0.16 Ma (Connelly et al., 2012). Extrapolation of this slope shows that HT clasts fall on the MG clast isochron. A combined isochron of HT and MG clasts results in a $(^{53}\text{Mn}/^{52}\text{Mn})_0$ of (7.50 ± 0.83) $\times 10^{-6}$, corresponding to an absolute age of 4568.1 ± 56 Ma. The $^{53}\text{Mn}/^{53}\text{Cr}$ ages obtained for MG clasts are in agreement with $^{26}\text{Al}^{26}\text{Mg}$ ages for bulk monomict ureilites (Baker et al., 2012), with an absolute age of 4566.7 ± 0.7 Ma when tied to the D’Orbigny angrite (Schiller et al., 2015a, see also Section 5.5 for further discussion). These ages are significantly older than $^{53}\text{Mn}/^{53}\text{Cr}$ ages obtained for bulk ureilites (Qin et al., 2010b; Yamakawa et al., 2010; Fig. 8, Table 5). However, ages from Qin et al. (2010b) and Yamakawa et al. (2010) are in good agreement for our feldspathic clasts (excluding BSa3 and TR3, see also Section 4.3.3), that corresponds to $(^{53}\text{Mn}/^{55}\text{Mn})_0 = (3.65 ± 1.5) \times 10^{-6}$ and an absolute age of 4564.1 ± 1.9 Ma. HT clasts also fall on the feldspathic clast isochron and when included yield $(^{53}\text{Mn}/^{52}\text{Mn})_0 = (3.62 ± 0.82) \times 10^{-6}$. The feldspathic isochron is in agreement with $^{53}\text{Mn}/^{53}\text{Cr}$ ages from in situ isotope analyses by secondary ionization mass spectroscopy (SIMS) on feldspathic clasts from polymict ureilites (Kita et al., 2004; Goodrich et al., 2010), the $^{26}\text{Al}/^{26}\text{Mg}$ age of the Almahata Sitta trachyandesite (Bischoff et al., 2014) and $^{182}\text{Hf}/^{182}\text{W}$ ages of bulk monomict ureilites (Budde et al., 2015). However, the $^{182}\text{Hf}/^{182}\text{W}$ age is not directly comparable to other ages, since it represents (1) a model age and (2) it discards six datapoints that when included would result in an age comparable to MG clasts and monomict ureilites (Baker et al., 2012). Budde et al. (2015) argue to discard these points based on contamination from terrestrial weathering, increasing $^{182}\text{Hf}/^{182}\text{W}$ ratios and $^{182}\text{W}$ abundances. However, three out of six ‘contaminated’ ureilites have similar $^{182}\text{Hf}$ concentrations as ‘uncontaminated’ ureilites and have $^{182}\text{W}$ ratios within the typical range of ureilites. Including these datapoints yields an age within error of CAI formation, which is in agreement with Baker et al. (2012) and our ages for monomict ureilites. Thus, we disregard the conclusions of Budde et al. (2015) in our discussion. Finally, the scatter of feldspathic clasts in $^{53}\text{Cr}$ versus $^{55}\text{Mn}/^{52}\text{Cr}$ space suggests that (1) not all feldspathic clasts were derived during the same event, (2) feldspathic clasts were not derived from the same source, or (3) $^{53}\text{Mn}/^{52}\text{Cr}$ systematics of the clasts were disturbed. Although we cannot confidently rule out any of these possibilities at this time apart from clasts BSa3, TR3 and possibly RT3 (see Electronic Annex), feldspathic clast textures show no signs of secondary processes such as shock melting or terrestrial weathering. Furthermore, the distribution feldspathic clasts in $^{53}\text{Mn}/^{53}\text{Cr}$ and $^{26}\text{Al}/^{26}\text{Mg}$ space excludes the formation of feldspathic clasts from a single source (MG clasts or HT clasts) at different times, but may involve these clasts originating from multiple sources at the same time or over a period of time (Figs. 4 and 8).

5.4.2. The age dichotomy explained

The picture emerging from our new chronological data and other work (Yamakawa et al., 2010; Qin et al., 2010b; Goodrich et al., 2010; Bischoff et al., 2014) shows a dichotomy in age populations with separate events dated at 4567.1 ± 1.1 Ma and 4563.5 ± 1.5 Ma (Fig. 10). This raises the question if this age difference is real or an artefact related to using different dating techniques in different laboratories. Feldspathic clasts have been dated with at least four different techniques (e.g., $^{53}\text{Mn}/^{53}\text{Cr}$ and $^{26}\text{Al}/^{26}\text{Mg}$ systematics by SIMS, MC-ICPMS and TIMS) and yield consistent ages averaging to 4563.1 ± 1.6 (2SD) and, therefore, must represent a real geological event. To better understand the significance of the age dichotomy, we investigate below the type of materials sampled as well as the petrological history of these materials with respect to diffusion of elements relevant for dating techniques. Olivine grains from monomict ureilites contain highly reduced magnesian metal-bearing rims that were suggested to be formed by reactions between hot olivine and graphite, likely during rapid decompression from an impact-related event (Takeda, 1987). Hence, these reduction rims represent thermal overprinting of the differentiation event that formed the ureilite residues, resulting in a potential resetting of short-lived chronometers during the lifetime of the parent radionuclide. Thus, by sampling the cores of olivines and pyroxenes rather than bulk monomict ureilites, it is more likely that the age obtained represents the actual differentiation event that formed these residues and possibly also the ureilite core (Barrat et al., 2015) rather than later resetting. The $^{53}\text{Mn}/^{53}\text{Cr}$ ages of bulk monomict ureilites, however, include these reduction rims (Yamakawa et al., 2010; Qin et al., 2010b) that were formed while $^{53}\text{Mn}$ with a half-life of 3.7 Myr was still extant. Ito and Ganguly (2006) show that diffusion of Cr in olivine, using pallasites as an example, can reset the $^{53}\text{Mn}/^{53}\text{Cr}$ ages of meteorites. If the ages obtained by Yamakawa et al. (2010) and Qin et al. (2010b) indeed record the resetting ages of monomict ureilites, then this event would not likely have had an impact on the $^{26}\text{Al}/^{26}\text{Mg}$ systematics of the bulk rocks, since (1) virtually all Al was extracted from ureilite residues during partial melting and (2) at that time no $^{26}\text{Al}$ was present when assuming an initial $(^{26}\text{Al}/^{27}\text{Al})_0$ similar to the angrite parent body (Schiller et al., 2015a). Hence, we suggest that the $^{26}\text{Al}/^{26}\text{Mg}$ age of 0.6 Myr after CAI formation by Baker et al. (2012) for bulk ureilites is more robust and likely records the timing of differentiation for monomict ureilite residues.
5.5. Modeling of ureilite differentiation events using initial $\mu^{26}\text{Mg}^+$ compositions

Modeling the $\mu^{26}\text{Mg}^+$ evolution of asteroidal bodies requires knowledge of the initial abundance and distribution of $^{26}\text{Al}$ in the solar protoplanetary disk. Although the canonical $^{26}\text{Al}/^{27}\text{Al}$ of $5.25 \times 10^{-5}$ observed in CV CAIs is typically assumed to represent the initial $^{26}\text{Al}$ abundance of the Solar System as a whole, a number of recent studies have challenged this assumption (Larsen et al., 2011, 2016b; Schiller et al., 2015a,b; Van Kooten et al., 2016). In particular, Schiller et al. (2015a) conducted a careful comparison for the $^{26}\text{Al}$-$^{26}\text{Mg}$ and U-corrected Pb-Pb dates of rapidly-cooled angrite meteorites to determine the initial $^{26}\text{Al}/^{27}\text{Al}$ ratio of inner Solar System solids. Their results establish that the initial $^{26}\text{Al}/^{27}\text{Al}$ ratio of inner Solar System reservoirs may have been reduced relative to the canonical value, a framework that we adopt in the current study (see Connelly et al. (2017) for further discussion). Correlated variability of $^{26}\text{Mg}^+$ and $^{54}\text{Cr}$ between bulk Solar System materials is thought to be the result of preferential destruction by thermal processing in the protoplanetary disk of labile $^{54}\text{Cr}$- and $^{26}\text{Al}$-rich carriers (Trinquier et al., 2007; Qin et al., 2010a,b; Yamakawa et al., 2010) and are therefore thought to have a correspondingly low initial bulk $^{26}\text{Al}/^{27}\text{Al}$ ratio. Due to the lack of representative bulk ureilite materials it is not possible to derive a robust estimate of the initial $^{26}\text{Al}/^{27}\text{Al}$ ratio of the UPB. Rather, we use the angrite parent body with $^{54}\text{Cr} = -0.36 \pm 0.07$ (Trinquier et al., 2007) for which the initial bulk $^{26}\text{Al}/^{27}\text{Al}$ ratio is calculated to be $1.33^{+0.21}_{-0.18} \times 10^{-5}$ (Schiller et al., 2015a) to provide an upper limit on the initial $^{26}\text{Al}$ abundance of the UPB. We model the evolution of the bulk UPB (Fig. 11) assuming an upper limit initial $^{26}\text{Al}/^{27}\text{Al}$ ratio of $1.33^{+0.21}_{-0.18} \times 10^{-5}$ (Schiller et al., 2015a), a solar $^{26}\text{Al}/^{26}\text{Mg}$ ratio of $0.00781 \pm 0.00029$ (Paton et al., 2012) and an initial $\mu^{26}\text{Mg}^+$ of $-15.9$ ppm (Larsen et al., 2011). This results in a maximum $\mu^{26}\text{Mg}^+$ ingrowth of up to $-6$ ppm for the bulk UPB (Fig. 11).

Main Group clasts, similarly to monomict ureilites, have an average $\mu^{26}\text{Mg}^+$ of $-10.6 \pm 5.3$ ppm, and our model ages suggest that these melt formed approximately 0.7 Myr after $t = 0$ (CV CAI formation; Connelly et al., 2012). We note that using an initial $^{26}\text{Al}/^{27}\text{Al}$ ratio of $(1.08 \pm 0.29) \times 10^{-5}$ for the bulk UPB as suggested by Larsen et al. (2011) yields a slightly younger age for ureilite residues of 0.9 Myr (Fig. 11, dotted line), but still within the age range of MG clasts and monomict ureilites (Baker et al., 2012). Alternatively, using a canonical $^{26}\text{Al}/^{27}\text{Al}$ ratio requires extremely rapid accretion and differentiation of the UPB at $t = 0$.

Hughes Type clasts either represent mantle residues or cumulates related to partial melting in the ureilite mantle.
models, carbonaceous chondrites (CI, CM, CV) with Ca/Al = 2.5 CI are typically used as the precursor material to the UPB. This is based on the high carbon content in monomict ureilites and the oxygen isotope data corresponding to a slope $/C_{24}$ line (Kita et al., 2004; Goodrich et al., 2007). However, Cr, Ca, O and Tm isotope data essentially precludes that ureilites were derived from carbonaceous chondrites (Trinquier et al., 2007; Qin et al., 2010b; Yamakawa et al., 2010; Warren, 2011; Schiller et al., 2015b; Barrat et al., 2016a). Additionally, the effect of basalt removal on MgO-SiO$_2$ and FeO-SiO$_2$ relationships suggests a non-carbonaceous origin for ureilites (Warren, 2011). Hence, in our model we use ordinary chondrites as precursor material to the UPB. A modeled 25% partial melt from an ordinary chondrite bulk composition yields Al/Mg ratios between 0.7 and 1.4, for fractional and equilibrium crystallization, respectively. In Fig. 11, we show the modeled $\mu^{26}\text{Mg}^+$ evolution of the range of partial melts and derivative cumulates with $\mu^{26}\text{Mg}^+$ values of 2.3 ± 6.3 ppm or 8.0 ± 5.0 ppm, assuming either kinetic or equilibrium processes during the mass-dependent isotope fractionation, respectively. The resulting differentiation age of HT clasts cumulates is then 0.9–1.5 Myr after CAI formation. It is apparent from Fig. 11 that HT clasts cannot have formed as primary residues from the bulk UPB, since this would have resulted in $\mu^{26}\text{Mg}^+ < -6$ ppm. However, a second melting event involving HT clasts cumulates around 3.8 Myr after CAI formation can result in the olivine-rich, low-Ca compositions observed for HT clasts and produce the $\mu^{26}\text{Mg}^+$ values measured in feldspathic clasts. This model agrees with the notion that feldspathic clasts are derived from depleted sources, based on their trace element and REE patterns (Kita et al., 2004). A late melting event, required to produce feldspathic clasts, cannot be initiated by heating from the decay of $^{26}\text{Al}$ as shown by the decay curve with $(^{26}\text{Al}/^{27}\text{Al})_0 = 1.33 \pm 0.21 \times 10^{-5}$ in Fig. 10: at 3.5 Myr after $t = 0$, virtually no $^{26}\text{Al}$ remains and the parent body cools. Hence, this event requires an alternative heating mechanism, possibly by major impact(s). We note that a disruptive impact to the UPB is thought to be prerequisite to the formation of reduction rims in monomict ureilite olivines (Takeda, 1987).

5.6. Disruption of the ureilite parent body

Polymict ureilites are breccias containing clasts that are derived from various depths in the ureilite mantle. This suggests that these meteorites accreted after a major disruptive
event able to exhume materials from tens of kilometers deep. Such a large disruptive impact event is expected to leave its mark on the geological record of the UPB, for example in the form of shock or decompression melting. Late stage melting, 3.8 ± 1.3 Myr after CAI formation, is best explained by impact heating, since essentially no $^{26}$Al is left for heating by radioactive decay (Fig. 10). Furthermore, the high amount of carbon in ureilites, the $\delta^{15}$N isotope anomalies, enrichment in noble gases and grain lineation in ureilites have been attributed to an impact event (Warren and Kallemeyn, 1988; Rubin, 1988; Warren and Rubin, 2010; Day et al., 2017), although the carbon abundance in ureilites is more likely to be the result of a magmatic process (Rubin, 1988). The relative enrichment of $^{15}$N is only found in polymict ureilites, suggesting that the impact projectile was carbonaceous in nature (Grady and Pillinger, 1988) in contrast to the non-carbonaceous UPB. While the impactor that is invoked to have exhumed the monomict ureilites and the impactor that would have caused impact melting do not necessarily have to represent the same event, the data at hand does not exclude the possibility that the exhumation and melting are related to the same impact event. In this scenario, an impactor can exhume material from kilometers deep, while at the same time causing decompression melting in shallow source cumulates. Ureilite material ejected during the impact may then have accreted back to the surface and subsequently mixed with newly formed feldspathic clasts forming a polymict breccia.

The $\varepsilon^{54}$Cr and $\mu^{26}$Mg' values of various chondritic clasts in DaG319 exclude these materials as precursors to the UPB. However, based on the high abundance of dark clasts in the breccias relative to other chondritic clasts (i.e., R, CB, EC and OC), we speculate that the dark clasts may be derived from the proposed large impactor. As mentioned before, these dark clasts are very similar in $\mu^{26}$Mg' and $\varepsilon^{54}$Cr isotope composition to CI carbonaceous chondrites. However, their position in $\varepsilon^{53}$Cr versus $\varepsilon^{54}$Cr space (Fig. 12) suggests that these clasts have experienced less thermal processing than CI chondrites and, thus, are more pristine. The primitive nature of these dark clasts suggests that their parent body may have accreted at a greater heliocentric distance compared to other carbonaceous chondrites. Hence, the inward migration of a larger asteroidal body with a primitive carbonaceous chondrite composition may reflect an epoch of disruption in the outer Solar System (Johnson et al., 2016). Indeed, the volatile-rich dark clasts in our study proposed to be fragments of a carbonaceous impactor are found in abundance in other meteorite breccias as well (Bischoff et al., 2006). A study of these objects might give insights into the flux of volatile-rich asteroidal bodies from the outer Solar System to the accretion region of the terrestrial planets. Dark clasts as well as impact related lithologies from planetary bodies may have the potential to place constraints on astrophysical models representing the dynamical evolution of our Solar System.

6. CONCLUSIONS

We studied the petrology and chemistry as well as the Mg- and Cr-isotope systematics of ureilite clasts from the polymict ureilite DaG 319. Our data provides the following

![Fig. 12. Plots of $\varepsilon^{53}$Cr versus $\varepsilon^{54}$Cr. Data for carbonaceous chondrites (orange) (Trinquier et al., 2007, 2009; Shukolyukov and Lugmair, 2006; Qin et al., 2010a; Van Kooten et al., 2016; Göpel et al., 2015), ordinary chondrite data (light blue) (Trinquier et al., 2007, 2009; Qin et al., 2010a), dark clasts in DaG 319 from this study (green), chondrule-containing clasts from this study (dark blue) and ureilite clasts from this study, where black represents data from TIMS and gray from MC-ICPMS. Variations in $\varepsilon^{54}$Cr between bulk Solar System materials can be attributed to thermal processing of $^{54}$Cr-rich dust (Trinquier et al., 2009), whereas unidirectional changes in $\varepsilon^{53}$Cr correspond to decay from $^{53}$Mn to $^{53}$Cr as seen for feldspathic and HT clasts.](image-url)
constraints on the melting and differentiation history of the UPB:

(1) The DaG 319 polymict ureilite consists foremost of MG clasts related to monomict ureilitic residues with varying Mg#s (0.74–0.91), as well as a large fraction of olivine-orthopyroxene (HT) clasts with consistently high Mg#s (0.89). Based on petrological and mineralogical characteristics, MG clasts are residues, whereas HT clasts were primarily formed as cumulates and further partial melting. A third type of clasts is recognized in DaG 319, consisting of feldspathic clasts that represent partial melts from the ureilite mantle.

(2) Significant mass-dependent Mg-isotope fractionation is present between feldspathic clasts and clasts related to ureilitic mantle residues and cumulates ($\Delta_{26}^{28}$Mg = 450 ppm), which can be best explained by equilibrium isotope fractionation during melting including a magnesite component enriching the melt in light Mg-isotopes. We suggest that Mg-carbonates were stable in the upper ureilite mantle, whereas pure carbon phases such as graphite were stable at higher pressures and depths. This is in agreement with a lack of carbon-related phases in HT clasts, whereas MG clasts contain affulent carbon.

(3) The timing of the differentiation events leading to the formation of MG and HT clasts are constrained by $^{54}$Mn-$^{53}$Cr systematics and $^{26}$Al-$^{26}$Mg model ages, which show that a primary differentiation event resulting in the formation of main group ureilite residues and related to decay of $^{26}$Al occurred early, $\sim$0.7 Myr after CAI formation. We suggest that a later ($\sim$3.8 Myr after $t = 0$) major disruptive impact was responsible for remelting of HT clasts, formation of feldspathic melts and resetting of $^{54}$Mn-$^{53}$Cr ages for monomict ureilitic residues. These feldspathic melts may have been extracted from multiple sources throughout the UPB.

(4) The MG clasts and HT clasts show variable $\varepsilon^{54}$Cr values that are unrelated to mass-dependent Cr-isotope fractionation. We propose that the variations in $\varepsilon^{54}$Cr are the result of admixing of carbonaceous chondritic matter during an impact-related disruptive event that caused late-stage melting in the upper ureilite mantle $\sim$3.8 Myr after CAI formation. The impactor may be represented by pristine dark clasts found in DaG 319.

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**APPENDIX A. SUPPLEMENTARY MATERIAL**

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.gca.2017.03.033.

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