The formation and chronology of the PAT 91501 impact-melt L-chondrite with vesicle-metal-sulfide assemblages

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Abstract – The L chondrite Patuxent Range (PAT) 91501 is an 8.5-kg unshocked, homogeneous, igneous-textured impact melt that cooled slowly compared to other meteoritic impact melts in a crater floor melt sheet or sub-crater dike (Mittlefehldt and Lindstrom, 2001). We conducted mineralogical and tomographic studies of previously unstudied mm- to cm-sized metal-sulfide-vesicle assemblages and chronologic studies of the silicate host. Metal-sulfide clasts constitute about 1 vol.%, comprise zoned taenite, troilite and pentlandite, and exhibit a consistent orientation between metal and sulfide and of metal-sulfide contacts. Vesicles make up ~2 vol.% and exhibit a similar orientation of long axes. $^{39}$Ar-$^{40}$Ar measurements date the time of impact at $4.461 \pm 0.008$ Gyr B.P. Cosmogenic noble gases and $^{10}$Be and $^{26}$Al activities suggest a pre-atmospheric radius of 40-60 cm and a cosmic ray exposure age of 25-29 Myr, similar to ages of a cluster of L chondrites. PAT 91501 dates the oldest known impact on the L chondrite parent body. The dominant vesicle-forming gas was $S_2$ (~15-20 ppm), which formed in equilibrium with impact-melted sulfides. The meteorite formed in an impact melt dike beneath a crater, as did other impact melted L chondrites, such as Chico. Cooling and solidification occurred over ~2 hours. During this time, ~90% of metal and sulfide segregated from the local melt. Remaining metal and sulfide grains oriented themselves in the local gravitational field, a feature nearly unique among meteorites. Many of these metal-sulfide grains adhered to vesicles to form aggregates that may have been close to neutrally buoyant. These aggregates would have been carried upward with the residual melt, inhibiting further buoyancy-driven segregation. Although similar processes operated individually in other chondritic impact melts, their interaction produced the unique assemblage observed in PAT 91501.
1. INTRODUCTION

Impact is one of three fundamental processes, along with accretion and differentiation, which formed and modified asteroidal bodies. From nebular accretion through to modern times, impact has left its traces in the ubiquitous cratered surfaces, distorted shapes, and telltale signs of shocked minerals and melts recorded in meteorites. Despite this abundant evidence, our knowledge of the physical processes and the timing of impact on asteroidal bodies remains incomplete. While we have many well-documented impact craters on Earth to serve as our guide to interpreting impact phenomena on other planets, we have only random sampling of asteroidal impact craters. Further, our knowledge of chronology is limited because the samples we do have may not accurately reflect the impact flux throughout the history of the Solar System. Continuing study of impact-derived meteorites can help fill these gaps in our knowledge.

Melt veined meteorites, impact melt breccias, and impact melts are not uncommon among the known ordinary chondrite population. They have been studied extensively to understand their origin (Rubin, 1985; Stöffler et al., 1991). In particular, considerable effort has focused on relating these meteorites to the physical setting of their formation, starting with the location of their melt. Melt can form on the floor and walls of a crater or in subsurface dikes that may extend beyond its walls. Each of these settings provides a unique physical and thermal environment for the incorporation of clastic material and cooling history.

Impact rates were much higher in the early history of the solar system (Hartmann et al., 2000), but those impacts are probably recorded in the asteroid belt by the population of small bodies produced by the break-up of larger precursor asteroids.
Impact craters observed on asteroids today are more recent events, consistent with the fact that most strong impact events in chondrites occurred within the past 1 Gyr, as determined by Ar-Ar radiometric dating (Bogard, 1995, and references therein). For example, many L-chondrites show Ar-Ar impact heating ages clustering near 0.5 Gyr, perhaps dating the time of disruption of the parent body (Haack et al., 1996). Interestingly, chronological evidence for collisional events very early in asteroid history is sparse.

This paper presents a multidisciplinary study of PAT 91501, a vesicular, impact melted L chondrite (Score and Lindstrom, 1992). Vesicles have been reported in only two other ordinary chondrite impact melts: Shaw (Taylor et al., 1979) and Cat Mountain (Kring et al., 1996). Although these meteorites are chemically and petrologically well-characterized (Harvey and Roeder, 1994, Mittlefehldt and Lindstrom, 2001), no study has addressed the implications of the presence of vesicles in impact melt rocks. Our objectives were to document the meteorite’s impact and cosmic ray exposure history and to understand the genesis of the unusual vesicular nature of this meteorite.

2. SAMPLES AND ANALYTICAL TECHNIQUES

Patuxent Range (PAT) 91501 was recovered in Antarctica during the 1991-1992 collecting season. It was transported more than 800 m prior to cataloging (R.P. Harvey, pers. comm.). Numerous large and small pieces, totaling more than 8.5 kg, were collected. Their relative positions in the meteoroid are unknown. In the same locale, two small metal-sulfide nodules (PAT 91516 and 91528; Clarke, 1994) were recovered. As discussed below, these meteorites are petrologically identical to metal-sulfide nodules
from PAT 95101 and are almost certainly samples from the same original mass (R.P. Harvey, pers. comm.).

PAT 91501 was originally classified as an L7 chondrite (Score and Lindstrom, 1992) based on textural features, mineral chemistry and oxygen isotopic composition, although it was noted that it was similar in many respects to the Shaw L chondrite impact melt. On further investigation (Harvey and Roedder, 1994; Mittlefehldt and Lindstrom, 2001), it was determined to be a near-total impact melt of an L chondrite.

One of the most striking features of PAT 91501 is the mm- to cm-sized vesicles seen on cut surfaces of the sample (Fig. 1), as originally noted by Marlow et al. (1992). We focused on PAT 91501 because it contains large vesicles that are visible in hand sample (Fig. 1), there is abundant material, and it has been described as a total impact melt (Mittlefehldt and Lindstrom, 2001). Visual inspection of PAT 91501 ,50 (2814.3 g) and ,78 (127.6 g) show clastless, light colored surfaces with cm-sized vesicles and metal/troilite aggregates.

2.1 Petrology

All available thin sections of PAT 91501 (,26; ,27; ,28; ,95; and ,111) at the Smithsonian National Museum Natural History, as well as sections of PAT 91516 and PAT 91528, were examined in both reflected and transmitted light with an optical microscope. Metal and troilite compositions were analyzed using a JEOL JXA 8900R electron microprobe at the Smithsonian. Analytical conditions were 20kV and 20nA. Well-known standards were used and analyses were corrected using a manufacturer supplied ZAF correction routine. Sulfur isotopes were analyzed using the 6f ion
microprobe at the Carnegie Institution of Washington utilizing Canyon Diablo troilite as the standard.

The two hand samples described above (91501, 50 and 91501, 78) were imaged at the High Resolution X-ray Computed Tomography facility at the University of Texas at Austin (UTCT), which is described in detail by Ketcham and Carlson (2001). The focus of our work was to determine the distribution of vesicles, metal and sulfide, which are easily distinguished based on their large density contrast from the silicate matrix. Sample PAT 91501, 50 was scanned using the high-energy subsystem, with X-rays set at 420 kV and 4.7 mA, with a focal spot size of 1.8 mm. The samples were scanned in air and, to reduce scan artifacts, the beam was pre-filtered with 1.58 mm of brass. Each slice was reconstructed from 1800 views, with an acquisition time of 128 ms per view. A total of 141 (1024x1024) slices were acquired with a thickness and spacing of 0.5 mm, imaging a 196 mm field of view. The final scan images were post-processed for ring artifact removal. Sample PAT 91501, 78 was imaged using the microfocal subsystem, with X-rays at 180 kV and 0.25 mA, and a focal spot size of approximately 0.05 mm. The sample was sealed and placed in a cylinder and surrounded by water, which was used as a wedge calibration to reduce scan artifacts. Data for 31 slice images were acquired during each rotation of the sample; over each rotation, 1000 views were acquired with an acquisition time of 267 ms per view. A total of 927 (1024x1024) slices at 0.0726 mm intervals, each showing a 67 mm field of view, were acquired. Scans were reconstructed using a software correction to further reduce beam hardening artifacts. Animations, including flipbooks for the 2D computed tomography scans and 3D rotational renderings are available at UTCT. (Movie and flipbook of PAT 91501, 50 can be seen at
http://web.mac.com/metritedoc/
Measurements of vesicles and metal/troilite particles from the CT data volume were made using Blob3D software (Ketcham, 2005), and visualizations were made using Amira® version 3.1.

2.2 Chronology (\(^{39}\)Ar-\(^{40}\)Ar ages and cosmic-ray exposure ages)

A 48-mg whole rock sample of PAT91501, along with multiple samples of the NL-25 hornblende age standard. This irradiation converted a portion of the \(^{39}\)K into \(^{39}\)Ar, and the \(^{40}\)Ar/\(^{39}\)Ar ratio is proportional to the K-\(^{40}\)Ar age. The irradiation constant (J-value) was 0.025210 ±0.000125. Ar was released from PAT 91501 in 34 stepwise temperature extractions and its isotopic composition was measured on a mass spectrometer. Experimental details are given in Bogard et al. (1995).

Two unirradiated whole rock samples of PAT 91501, taken from different locations in the meteorite (see below), were degassed in either two or four stepwise temperature extractions and the He, Ne, and Ar released were analyzed on a mass spectrometer. All noble gas analyses were made at NASA-JSC.

We analyzed chips from four different specimens (subsamples 34, 38, 40, and 42) of PAT 91501. Sample 34 was located adjacent to sample 106, which was analyzed for noble gases. Using facilities at Rutgers, the four specimens were ground and weighed. After addition of Al and Be carriers, the powders were dissolved in strong mineral acids. Beryllium and aluminum were separated by ion exchange, precipitated as the hydroxides, and ignited to the oxides as described by Vogt and Herpers (1988). The activities of \(^{10}\)Be and \(^{26}\)Al were measured by accelerator mass spectrometry at the University of Pennsylvania as described by Middleton and Klein (1986) and Middleton et al. (1983).
As the $^{10}\text{Be}$ activity of sample 40 was unaccountably low and inconsistent with the $^{26}\text{Al}$ activity, we do not report it.

### 3. PREVIOUS WORK

The petrology of the silicate portion of PAT 91501 is reported by Harvey and Roeder (1994) and Mittlefehldt and Lindstrom (2001) and we briefly review this work. PAT 91501 is an unshocked, homogeneous, igneous-textured rock of broadly L chondrite mineralogy and chemistry. Major element mineral chemistries were shown to be consistent with those of L chondrite; the minor element chemistry of olivine and low-Ca pyroxene, on the other hand, is consistent with melting (Mittlefehldt and Lindstrom, 2001); depletions of Zn and Br and sequestration of siderophile and chalcophile elements into the large, heterogeneously-distributed metal-sulfide aggregates were observed (Mittlefehldt and Lindstrom, 2001). Relic material includes rare chondrules, as well as opaque-inclusion-rich olivine and some low-Ca pyroxene grains that comprise ~10 vol.% of the meteorite, but distinct clasts commonly found in impact melt breccias are absent. Mittlefehldt and Lindstrom (2001) concluded that PAT 91501 is an impact melt of an L chondrite that crystallized at a cooling rate slower than that typical for impacts melts and likely formed in a melt sheet on the crater floor or in a sub-crater melt dike.

### 4. RESULTS

We report our analyses of the metal-sulfide-vesicle assemblages, based on both microscopic examination and computed tomography, and the results of the chronological analyses for both $^{39}\text{Ar}-^{40}\text{Ar}$ and cosmogenic noble gases and radionuclides.

#### 4.1 Petrography of metal-sulfide assemblages
PAT 91501 contains both vesicles and rounded metal-sulfide nodules that reach 1 cm in diameter. Previous studies have focused primarily on the silicate portion (Harvey and Roedder, 1994; Mittlefehldt and Lindstrom, 2001), with neither study reporting detailed examination of a metal-sulfide nodule in thin section. As we discuss later, metal-sulfide nodules are rare, with less than 1 per cm$^3$. Apart from a single chemical analysis of troilite reported by Mittlefehldt and Lindstrom (2001), they have never been studied from PAT 91501.

We examined a 5-mm-diameter metal-sulfide nodule adjacent to a 5-mm-diameter vesicle in subsample 111. This nodule consists of a core of Fe,Ni metal (2 by 3.5 mm) rimmed by sulfide, with the two phases exhibiting numerous mutual protrusions into each other. The sulfide is dominantly troilite, although minor (<1 vol.% of the sulfide) pentlandite ($\text{Fe}_{6.15}\text{Ni}_{2.62}\text{Co}_{0.10}\Sigma=8.88\text{S}_8$) is observed at troilite-metal, troilite-silicate and troilite-vesicle boundaries. Schreibersite rims are often found at the border between metal and troilite/pentlandite. The S isotopic composition of pentlandite (3 analyses yield $\delta^{34}\text{S}$ of 0.5-1.9‰) and troilite (7 analyses yield $\delta^{34}\text{S}$ of 0.4-1.2‰) are essentially identical. No polycrystallinity or twinning is observed in the troilite, confirming the observation of Mittlefehldt and Lindstrom (2001) that PAT 91501 experienced minimal secondary shock after its crystallization. The metal is composed of two domains (Fig. 2a). Rimming each domain is a 50 µm thick region of high-Ni (up to 45 wt.%; Fig. 2b) taenite that is relatively inclusion free. Adjacent to this, Ni decreases systematically from ~40 wt.% to ~20 wt.% and this zoned metal often contains 10-30 µm troilite and 1-5 µm schreibersite inclusions. The center of the largest domain, which appears to have been
bisected, is martensitic, with irregular Ni concentrations of 20-25 wt.%, and contains
troilite and schreibersite inclusions that can reach tens of microns.

While metal-sulfide nodules from PAT 91501 have not been previously described,
the published descriptions (Clarke, 1994) for the small iron meteorites PAT 91516 (1.58
g) and PAT 91528 (3.34 g) are essentially identical to that given here for PAT 91501.
The only substantive difference is that Clarke observed a larger number of metal
domains, particularly in PAT 91516, and these were often separated by sinuous troilite.

4.2 Computed Tomography

We used computed tomography (CT) to survey the distribution of vesicles, metal
and sulfide in two samples of PAT 91501 (see flipbooks in supplemental data). Figure 3
is a single frame of a 3 dimensional, rotational visualization made from the CT scan of
PAT 91501,50, in which vesicles and metal-sulfide intergrowths are highlighted. In this
sample, 5085 vesicles were measured, which comprise ~2 volume percent of the sample.
The sizes of the vesicles range in diameter from 0.6 to 14 mm. In contrast, analysis of
PAT 91501,78 yielded 36685 vesicles ranging in size from 0.2 to ~6mm in diameter.
The difference in the numbers and the size range of vesicles is due to the fact that the
smaller sample was scanned at a much higher resolution. In both samples, tiny vesicles
(<1mm diameter) dominate the population. Vesicles in both samples are homogenously
distributed and have a median aspect ratio of 1.4, indicating moderate elongation.

The CT scans revealed the existence and distribution of several large metal-
sulfide intergrowths (Fig. 3). Together, metal and sulfide represent less than 1 volume
percent of the sample. We measured 255 and 142 metal grains in sample ,50 and ,78,
respectively. Metal ranges in size from 0.7 to 8.6 mm in the larger sample and comprises
0.27 vol%, while in the smaller sample metal ranges from 0.1 to 3.8 mm and represents 0.35 vol%; as with the vesicles, the higher-resolution scan of the smaller sample permitted us to measure particles too small to be resolved in the scan of the larger specimen. Sulfide is more abundant than metal in both samples and occupies approximately twice the volume as metal. Sulfide accounts for 0.4 vol% in the larger sample (.50) and for 0.56 vol% in the smaller sample. We measured 404 sulfide grains in sample .50 and 540 grains in sample .78. Sulfide is overall larger than metal and ranges from 0.6 to 12.7mm in .50 and from 0.2 to 4.7mm in .78. As with vesicles, tiny grains (< 1mm) comprise the mode of both the metal and sulfide size distributions (see flipbooks and 3D renderings in supplemental data).

PAT 91501 (.50) contains 169 grains in which metal and sulfide are in contact. These particles were noted by earlier workers (Score and Lindstrom, 1992; Mittlefehldt and Lindstrom, 2001) and attributed to formation as immiscible melts prior to silicate crystallization. Interestingly, these particles exhibit a consistent orientation of the metal and sulfide relative to each other and to the meteorite as a whole. Figure 4a is a stereo plot of the normals to the planes defined by the contact between metal and sulfide with the size of the each circle proportional to the area of the contact. Although some scatter is observed in this plot, particularly for smaller metal-sulfide pairs, the majority of larger particles defines a tight cluster trending 255° and plunging 45°; note that these orientations are with respect to the scan data, and are not geographical.

The CT scans also document the relationship between vesicles, metal, and sulfide. Larger vesicles appear to have metal-sulfide intergrowths associated with them. In the CT scan of the larger sample of PAT 91501, we found 18 instances where vesicles are in
contact with metal only, sulfide only or metal-sulfide intergrowths. In the higher resolution CT scan of PAT 91501, 78, there are nearly 200 vesicles in contact with metal, sulfide or metal-sulfide. The vast majority of the largest vesicles are in contact with metal and/or sulfide.

The elongation of vesicles allows us to examine their orientation as well. Fig. 4b is a stereo plot of the orientations of the vesicle long axes from the main mass of PAT 91501 (,50) with the circle areas proportional to vesicle volume. Again, considerable scatter is observed, particularly among the smaller vesicles. However, the larger vesicles define a distinct cluster trending 300° and plunging 40°, with the main outlier attributable to contact with an irregular metal-sulfide mass. This cluster is offset ~33° from the orientation defined by the normals to the metal-sulfide contacts.

4.3 Ar-Ar Age

The PAT 91501 Ar-Ar age spectrum (Fig. 5) appears complex but can be interpreted to yield a reliable age. The rate of release of 39Ar and changes in the K/Ca ratio and the Ar-Ar age as a function of extraction temperature all suggest that 39Ar is contained in three distinct diffusion domains—0-17%, 17-80%, and 80-100% 39Ar release (Fig 5). The 39Ar release data can be modeled by standard diffusion theory in terms of the parameter $D/a^2$, where $D$ is the diffusion coefficient and $a$ is the average diffusion length for Ar in the degassing grains. On an Arrhenius plot (argon released vs. $1/T$; not shown), data for these three domains give separate linear trends, each one characterized by a different value of $D/a^2$. Above 80% 39Ar release, the observed decreases in age and K/Ca are interpreted to represent release of excess 39Ar recoiled during irradiation into the surfaces of pyroxene grains. Below ~17% 39Ar release, the higher ages are
interpreted to represent loss of recoiled $^{39}$Ar from surfaces of grains possessing a relatively larger K/Ca ratio. Between ~19% and 80% of the $^{39}$Ar release, the K/Ca ratio is relatively constant and the Ar-Ar ages describe a plateau. Ten extractions releasing 19-78% of the $^{39}$Ar define an age of 4.463 ±0.009 Gyr, where the age uncertainty is approximately one-sigma and includes the uncertainty in the irradiation constant, $J$. Seven extractions releasing 30-78% of the $^{39}$Ar give an age of 4.461 ±0.008 Gyr. To examine these data in an isochron plot, we adopted the cosmogenic $^{38}$Ar concentration given below and used the measured $^{37}$Ar/$^{36}$Ar ratios for each extraction to apportion the measured $^{36}$Ar into trapped and cosmogenic components. An isochron plot of $^{40}$Ar/$^{36}$Ar versus $^{39}$Ar/$^{36}$Ar, using trapped $^{36}$Ar, is highly linear ($R^2=0.9995$) and its slope yields an age of 4.466 ±0.0012 Myr, in agreement with the plateau age. The isochron intercept value of $^{40}$Ar/$^{36}$Ar =-79±156 suggests all $^{40}$Ar released in these extractions is radiogenic. The total age summed across all extractions is 4.442 Gyr and suggests that little to no $^{40}$Ar was lost from the sample by diffusion over geologic time. We conclude that impact resetting of the K-Ar age occurred 4.46 ±0.01 Gyr ago.

4.4 Cosmogenic Noble Gases and Radionuclides.

PAT 91501 ,109 (33.4 mg) was heated in two temperature steps and sample ,106 (50.0 mg) was heated in four steps (Table 1). In both samples approximately half of the $^3$He was released at 500°C. In sample ,106 the peak of the Ne release occurred at 900-1200 °C, and the peak of the $^{38}$Ar release occurred at 1200 °C. Measured $^3$He is entirely cosmogenic. The summed $^{20}$Ne/$^{22}$Ne ratios of 0.845-0.847 indicate that measured Ne is also entirely cosmogenic. Consequently we summed concentrations for each Ne isotope across all extractions to obtain total cosmogenic abundances. Measured $^{36}$Ar/$^{38}$Ar ratios
varied over 0.72-1.75 and indicate the release of trapped Ar, which is mostly adsorbed atmospheric Ar, particularly at lower extraction temperatures. We assumed $^{36}$Ar/$^{38}$Ar ratios of 5.32 for trapped Ar and 0.67 for cosmogenic Ar and calculated the abundances of cosmogenic $^{38}$Ar for each extraction. The $^{38}$Ar$_{cos}$ abundances were then summed across each extraction to obtain the total abundance of $^{38}$Ar$_{cos}$. From analyses of He, Ne, and Ar delivered from a standard gas pipette, we estimate the uncertainty in these abundances as $\sim \pm 10\%$.

Cosmogenic abundances and $^{22}$Ne/$^{21}$Ne ratios for the two PAT samples are given in Table 2. Cosmogenic abundances of $^3$He, $^{21}$Ne, and $^{38}$Ar in the two samples agree with each other within their individual uncertainties of $\pm 10\%$. The measured $^{20}$Ne/$^{21}$Ne ratios for ,109 and ,106 are identical at 0.847 $\pm 0.005$ and 0.845 $\pm 0.015$. The measured $^{22}$Ne/$^{21}$Ne ratios of 1.084 $\pm 0.003$ and 1.097 $\pm 0.003$ differ slightly, which probably reflects a shielding difference. A plot of $^3$He/$^{21}$Ne versus $^{22}$Ne/$^{21}$Ne defines a shielding trend for many chondrites (Eberhardt et al., 1966). Sample ,109 plots on this shielding trend, but sample ,106 plots slightly above the trend, as a consequence of its lower $^{21}$Ne concentration. This observation may imply that in our ,106 sample the concentration of Mg, the main target for $^{21}$Ne$_{cos}$ production, was slightly less than the chondritic value. There is no suggestion of diffusive loss of $^3$He in either sample, in spite of the observation that He degassed at relative low temperature in the laboratory (Table 1).

The measured abundances of $^{10}$Be for three PAT samples agree within their uncertainties (Table 2). The measured activities of $^{26}$Al in four PAT samples (Table 2) span a range of $\sim 17\%$.

4.5 PAT Pre-Atmospheric Size
The $^{22}\text{Ne}^{21}\text{Ne}$ ratio of ~1.09 indicates that the pre-atmospheric shielding experienced by PAT 91501 was somewhat greater than that for typical chondrites. The maximum dimension of the recovered meteorite was ~19 cm, which, for the purpose of modeling calculations, sets a minimum radius in space of ~10 cm. Modeling of the $^{22}\text{Ne}^{21}\text{Ne}$ ratio in L-chondrites (Leya et al., 2000) predicts that as the meteoroid radius increases, $^{22}\text{Ne}^{21}\text{Ne}$ ratios as low as ~1.09 are first reached in the center of a body with a pre-atmospheric radius of ~ 30 cm. Thus a somewhat larger body presumably carried the physically separate samples that we analyzed. According to the calculations of both Leya et al. (2000) and Masarik et al. (2001), $^{22}\text{Ne}^{21}\text{Ne}$ ratios plateau at 1.09±0.01 for pre-atmospheric depths from 10 to ≥30 cm in L chondrites with radii of 40 cm.

The $^{22}\text{Ne}^{21}\text{Ne}$ ratio is not useful for setting an upper bound on the pre-atmospheric radius. For this purpose we use the $^{26}\text{Al}$ activity. After a cosmic ray exposure lasting more than 20 My (see below) activities of $^{26}\text{Al}$ (and $^{10}\text{Be}$) would have reached saturation and are therefore equal to average production rates in space – assuming the terrestrial age of PAT was less than 50 kyr or so as suggested by the normal $^{26}\text{Al}/^{10}\text{Be}$ ratios for three samples. According to the calculations of Leya et al. (2000), only meteoroids with radii between 32 cm and 85 cm have the range of $^{26}\text{Al}$ activities observed in PAT. The $^{10}\text{Be}$ activities of PAT 91501 are comparable to those of the L5 chondrite St-Robert (Leya et al., 2001), which is thought to have had a pre-atmospheric radius between 40 and 60 cm. We conclude that the pre-atmosphere radius of PAT 91501 was in this range.

4.6 Cosmic Ray Exposure Age.
Cosmic Ray Exposure (CRE) ages of stony meteorites were initiated by impacts that reduced meteoroids to objects meter-size or smaller and are almost all <100 Myr for stones (Herzog, 2004). To calculate cosmic ray exposure ages for PAT (Table 2), we used the cosmogenic production rates for L-chondrites given by Eugster (1988), except that the $^{38}$Ar production rate was lowered by 11%, as suggested by Graf and Marti (1995). The production rates were corrected for shielding using the measured $^{22}$Ne/$^{21}$Ne ratios. The differences among ages calculated from He, Ne, and Ar for a given sample are greater than the differences in the same age between the two samples. This pattern suggests that most of the apparent variation in CRE age is produced by our choice of production rates. Because cosmogenic Ar is more sensitive to likely compositional variations and because there is some chance that cosmogenic Ar was incompletely extracted, we give greater weight to the $^3$He and $^{21}$Ne ages and obtain a CRE age for PAT 91501 of 25-29 Myr.

We also calculated CRE ages based on the $^{26}$Al-$^{21}$Ne-$^{22}$Ne/$^{21}$Ne and $^{10}$Be-$^{21}$Ne-$^{22}$Ne/$^{21}$Ne equations of Graf et al. (1990a) by using data for the two samples known to have been adjacent to each other .34 and .106. The results, 29.6 Myr and 25.5 Myr, respectively, are in the same range as the CRE ages calculated from the noble gases alone. Finally, we calculated the $^{10}$Be-$^{21}$Ne CRE age for the .34 -.106 pair by using the formula of Leya et al. (2000) after modifying it for a $^{10}$Be half life of 1.5 My. This age, 21.9 My, is about 15-26% lower than the others. Leya et al. (2000) observed that their equation for $^{10}$Be-$^{21}$Ne CRE ages gives a low result for another large L-chondrite, Knyahinya (pretmospheric radius ~45 cm; Graf et al., 1990b). They attribute the
discrepancy to their model’s underestimation of $^{10}\text{Be}$ production rates in meteoroids the size of Knyahinya and larger.

5. DISCUSSION

Among ordinary chondrites, the L chondrites record a particularly severe history of impact bombardment, with almost 5% of this group containing shock melts (La Croix and McCoy, 2007). In this regard, PAT 91501 is not atypical. Indeed, its similarity to the impact-melted Shaw L chondrite was noted during its initial description (Marlow et al., 1992). However, the ancient age, vesicular nature, presence of preserved, cm-sized metal-troilite intergrowths, and orientation of both the vesicles and metal-sulfide particles are unusual. These features promise new insights into the timing of and physical processes occurring during the formation of this impact-melted L chondrite.

5.1 Chronology

Chronological evidence for collisional events on asteroids very early in Solar System history is sparse. Among achondrites, some unbrecciated eucrites may have been excavated from depth on Vesta by a large impact ~4.48 Gyr ago (Yamaguchi et al., 2001; Bogard and Garrison, 2003). Within the chondrites, McCoy et al. (1995) reported ages of enstatite chondrite impact melts dating to before 4.3 Gyr and Dixon et al. (2004) suggested that Ar-Ar ages of ~4.27 Gyr for a few LL-chondrites may date the time of one or more impact events on the parent body. Taken alone, Ar-Ar ages between ~4.38 to ~4.52 Gyr can be ambiguous, as ancient ages may reflect either a late impact or slow cooling after parent body metamorphism (Turner et al., 1978; Pellas & Fiéni, 1988). In contrast, impact melts provide a more direct means for dating the timing of collisional events. Most impact melts give Ar-Ar ages less than 1 Gyr, suggesting that melting and
re-solidification took place recently, either during events confined to the surfaces of modern asteroids or, perhaps, when collisions on asteroids melted partially and launched meteoroids into Earth-crossing orbits. This population of more recently-formed impact melt rocks includes the vesicular meteorites Cat Mountain (Kring et al., 1996) and Chico (Bogard et al., 1995).

In contrast, PAT 91501 dates to the earliest history of the Solar System at 4.461 Gyr. Until this work, Shaw was the only ordinary chondrite known to be a near total impact melt (Taylor et al., 1979) and have an Ar-Ar age consistent with an early (>4.0 Gyr ago) impact (Turner et al., 1978). Indeed, PAT 91501 shares a number of features with Shaw, particularly its light-colored lithology, petrographic texture and clast-free nature (Taylor et al., 1979). Based on its cosmogenic noble gas concentrations, Shaw has a much younger, nominal one-stage CRE age of ~0.6 My, although in all likelihood, Shaw had a complex exposure history with a first stage that probably lasted >10 Myr (Herzog, 1997). In any event, cosmic-ray exposure ages greater than 1 Gyr are unheard of in stones and thus Shaw’s old Ar/Ar age indicates that the meteoroid did not melt (and hence lose an appreciable fraction of its radiogenic ⁴⁰Ar) when it was launched from the asteroid. The 4.46 Gyr impact event that formed PAT 91501 apparently took place considerably earlier than those impacts that reset the Ar-Ar ages of Shaw (4.40 ±0.03 and 4.42 ±0.03 Gyr; Turner et al, 1978). We conclude that PAT 91501 and Shaw formed in different impact events on the L-chondrite parent body, and that the two meteorites were not located in close proximity. There seems little question that PAT 91501 is closely related to the bulk of L chondrites and this relationship is supported by the CRE age for PAT of 25-29 Myr, which lies within a diffuse ~22-30 Myr cluster in the distribution of
L-chondrite CRE ages. The 4.46 Gyr impact for PAT 91501 falls within the range of Ar-
Ar metamorphic ages of relatively unshocked chondrites (Turner et al., 1978; Pellas &
Fiéni, 1988). This observation implies that the L parent body experienced a significant
impact while it was still relatively warm. In all likelihood, these events occurred on the
original L chondrite parent body prior to any subsequent collisions and breakups that
would have formed modern asteroids. These early impacts left PAT 91501 deeply buried
until it was excavated and launched toward Earth ~28 Myr ago.

5.2 Vesicle Formation

PAT 91501 is remarkable for its mm- to cm-sized vesicles. Vesicles of this size
have never before been observed in an impact-melt rock. The few vesicular meteorites
that have been investigated in detail are basaltic eucrites or angrites, where vesicles are
formed by gases liberated or generated during silicate partial melting (McCoy et al., 2006
and references therein). In terrestrial systems, H?O is the typical vesicle-forming gas, as
it is abundant in the Earth’s crust and exsolves from basaltic magmas at relatively
shallow depths (Oppenheimer, 2004). In contrast, chemical analyses (Jarosewich, 1990)
and the presence of abundant Fe,Ni metal suggest that ordinary chondrites likely were
very dry systems and, thus, H?O is unlikely as a major vesicle-forming gas. McCoy et al.
(2006) argued that a mixed CO-CO? gas was responsible for vesicle formation in
asteroidal basalts and the contribution of such a gas cannot be unequivocally eliminated.
The contribution of volatiles from the impactor, such as ice in a cometary body, or
volatilization of silicates at superheated temperatures also seems unlikely, although
impossible to rule out.
A much more likely source of volatiles is sulfur vaporization during impact melting. Numerous previous studies have pointed to the role of sulfur vaporization during metamorphic and impact processes of ordinary chondrites. Lauretta et al. (1997) showed that a small amount of sulfur vaporizes at the metamorphic temperatures of ordinary chondrites. Sulfur vaporization is also a common problem in ordinary chondrite melting experiments (e.g., Jurewicz et al., 1995) and has been invoked to explain the formation of sulfide-rich regions in the Smyer H chondrite impact melt breccia (Rubin, 2002). In PAT 91501, the larger vesicles have metal-sulfide intergrowths associated with them, suggestive of formation by sulfide vaporization during impact melting.

We can calculate the amount of S gas required to create the abundance of vesicles (~2 vol.%) documented with computer tomography. The formula for the bulk density $\beta$ of a vesicular material is

$$1/\beta = n/\rho_g + (1 - n)/\rho_{ng}$$

where $\rho_g$ is the gas density, $\rho_{ng}$ is the density of the non-gas part (i.e. solid or liquid) and $n$ is the mass fraction of gas. If the conditions are such that the gas law holds at least approximately, the density of the gas is given by

$$\rho_g = (mP)/(QT)$$

where $m$ is the molecular weight of the gas, $P$ is its pressure, $Q$ is the universal gas constant (8314 J/kmol) and $T$ is the gas temperature. Substituting for $\rho_g$:

$$1/\beta = (nQT)/(mP) + (1 - n)/\rho_{ng}$$

The two terms on the right are the partial volumes of gas and non-gas, respectively, so the gas volume fraction $v_g$ is given by

$$v_g = [(nQT)/(mP)] / [(nQT)/(mP) + (1 - n)/\rho_{ng}]$$
which is more conveniently re-arranged as

$$n / (1 - n) = (v_g m P) / [(1 - v_g) \rho_{ng} Q T]$$  \hspace{1cm} (5)$$

Using the values we estimated, $v_g = 0.02$, $T = 1670 \text{ K}$, $\rho_{ng} = 3520 \text{ kg m}^{-3}$, $m = 64$ for $S_2$ or $SO_2$, and $P = 5 \times 10^5 \text{ Pa}$ for a depth of a few km in a 50 km radius asteroid, appropriate to the lithostatic load likely to occur in a silicate melt at depth, we find $[n / (1 - n)] = 1.448 \times 10^{-5}$, so that $n = 1.448 \times 10^{-5}$ to the same precision. If we regard this as the abundance of $S_2$, it indicates that $\sim 15$ ppm of gas are necessary for formation of the vesicles, in excellent agreement with earlier calculations of $S_2$ generated in equilibrium with sulfide in PAT 91501.

Alternatively, we can estimate the amount of $S_2$ gas produced by vaporization if all the FeS in a chondrite melted using the following equation from Lauretta et al. (1997):

$$\text{Fe}_{1-x}S \rightarrow 0.99 \text{Fe}_{1-x}S + 0.005S_2$$  \hspace{1cm} (6)$$

Thus, each mole of sulfide liberates 0.005 moles of $S_2$. So for a typical L-chondrite mode of FeS (~4.2 vol%), it would be expected that $\sim 210$ ppm of $S_2$ would form during melting. While this may seem like a small amount, the vesicle volume produced by this amount of gas would be much greater than that observed in PAT 91501. As we discuss in the next section, the amount of sulfide present in the impact melt likely results from gravitational segregation of the dense metal-sulfide particles in the silicate melt. It is likely that the amount of sulfide that actually vaporized is much closer to about one-tenth that of average L chondrites, thus, a $S_2$ gas abundance of $\sim 20$ ppm is probably more reasonable.

The abundance of 15-20 ppm $S_2$ required for vesicle formation and in equilibrium with sulfide is a vanishingly small amount, as also noted for the abundance of CO-CO$_2$ in
vesicular angrites and eucrites by McCoy et al. (2006). Thus, it is no surprise that
evidence for its condensation cannot be found. No sulfide or sulfur linings have been
observed on vesicle walls in PAT 91501, although moderate terrestrial weathering has
occurred in the meteorite and hydrated iron oxides of terrestrial origin commonly occur
as vesicle linings. We considered the possibility that pentlandite found in the metal-
sulfide assemblages might reflect S volatilization. However, no isotopic fractionation
consistent with S volatilization was observed between pentlandite and troilite and, as
discussed later, it appears more likely that pentlandite is an equilibrium phase formed
during cooling in the Fe-Ni-S system.

5.3 Physical Setting and formation of PAT 91501

While PAT 91501 joins a growing list of impact-melted rocks from the L
chondrite parent body, its ancient age and large metal-sulfide nodules and vesicles and
their striking orientation are unique. Whereas it lacks the abundant clasts observed in
many impact melt breccias, similar clast-poor lithologies are observed in Shaw and, most
notably, as a 30 cm wide vein in Chico (Bogard et al., 1995).

Mittlefehldt and Lindstrom (2001) suggested that PAT 91501, because of its
homogenously melted nature and relatively slow cooling compared to other impact melts,
could be part of an impact melt basal layer found on the floor of a crater (Melosh, 1989)
or as a melt dike injected into surrounding country rock (Stöffler et al., 1991). Our work
provides additional constraints to distinguish between these two settings. There are
several reasons to question the formation of PAT 91501 in a crater floor melt sheet. On
Earth, these melt sheets tend to experience rapid cooling and be clast laden. Only in the
very largest terrestrial craters (e.g., Manicouagan, Sudbury) where impact melt sheets
exceed 200 m in thickness are clast-poor, igneous textured rocks observed (Keil et al., 1997). Likewise, fragmentation of a vesicular lava flow or impact melt sheet will occur at the surface of a low-gravity, atmosphereless body. To achieve the equivalent of terrestrial atmospheric pressure, McCoy et al. (2006) calculated that a melt sheet ~130 m thick would be needed on a body ~250 km in radius. These two estimates are in good agreement and suggest the need for a melt sheet in excess of 100 m thickness. An impact event capable of producing such a thick melt sheet on an asteroid would, instead, collisionally disrupt the body (Keil et al., 1997). Thus, we suggest that a melt dike injected into the surrounding country rock below the impact crater is a more viable setting for the formation of PAT 91501.

Injection of molten chondritic material into the surrounding country rock provides both moderate pressure necessary for vesicle retention and a thermal environment conducive to rapid cooling without quenching. We have been unable to constrain the cooling rate. Although zoning within the large taenite particles might normally be taken as indicative of cooling, we argue instead that the assemblage taenite (γ)-troilite-pentlandite is an equilibrium assemblage formed during cooling at temperatures between ~300-500 °C, consistent with phase relations in the Fe-rich portion of the Fe-Ni-S system (Ma et al., 1998). With the surrounding country rock cooler than the melt, solidification could have occurred in a matter of hours. The absence of distinct clasts in PAT 91501 is not inconsistent with such a model. While most impact melt breccias, by definition, contain clastic material from the country rock, subcrater melt dikes on Earth exhibit a range of widths (Keil et al., 1997) and it is reasonable to assume that PAT 91501 sampled one of the wider, clast-free portions of a dike. Indeed, Bogard et al. (1995) argue that a
30 cm wide zone of clast-poor impact melt in Chico samples such an intrusive dike. At a maximum dimension of ~20 cm, PAT 91501 would not be extraordinary in this regard.

Although cooling and crystallization may have occurred relatively rapidly in this dike, we suggest that it was far from a quiescent environment. Despite the preservation of metal and sulfide as mm- to cm-sized nodules, it is clear from the bulk elemental composition that metal and sulfide were lost from the system. Comparison of metal and sulfide abundances in PAT 91501 (~0.3 and ~0.5 vol.%, respectively) with those reported for average L chondrites (3.7 and 4.2 vol.%, respectively; McSween et al., 1991) suggests that the melt from which PAT 91501 crystallized lost ~90% of the metal and sulfide component prior to solidification. This loss is not surprising, given the marked density contrast between molten metal, sulfide and silicate. In fact, a similar density contrast exists between molten silicates and the vesicles, leading to rapid rise within the melt. The velocity, \( u \), of settling or rising is determined using the Stoke’s velocity equation (Turcotte and Schubert, 2002)

\[
\frac{\nu}{u} = \frac{l}{3} \frac{\Delta \rho r^2 g}{\eta}
\]

(7)

where \( \Delta \rho \) is the difference in density between the metal, sulfide or vesicles and the silicate melt; \( r \) is the radius of the grain of metal or sulfide or vesicle; \( g \) is gravity for an assumed 50 km radius parent body (0.012 m/s\(^2\)); and \( \eta \) is the viscosity of the silicate melt through which the metal, sulfide or vesicle is moving. We estimated the liquidus temperature of bulk L-chondrite composition to be between 1400 and 1600 °C, which affects the viscosity of the melt. Using the maximum size of the metal, sulfide and vesicles determined from the CT scan, we calculate that the largest vesicle would rise at ~3 m/hr while the largest metal particle would sink at ~2 m/hour.
One of the most astonishing results from the CT scans is the orientation of the metal/sulfide intergrowths. These orientations, reflected in the relative orientation of metal to sulfide (Fig. 3) and the orientations of metal-sulfide contacts and vesicle elongation – are consistent with formation in a gravitational field. In this respect, PAT 91501 is exceptional in that we know which way was up while on the asteroid. This orientation is illustrated in Fig. 3. To the best of our knowledge, only one other meteorite can claim such a distinction. In the Cape York meteorite (Kracher et al., 1977; Buchwald, 1987), elongate troilite inclusions contain chromites concentrated at one end and phosphates at the other, which may be indicative of formation in a gravitational field (Kracher and Kurat, 1975), but have also been attributed to melt migration in a thermal gradient (Buchwald, 1987).

In practice, calculated velocities probably represent theoretical maximums, as the vesicles likely coalesced during rise while the metal particles typically contain significant amounts of attached, less-dense sulfide. Nonetheless, these calculations suggest that metal-sulfide particles and vesicles should have rapidly segregated from the volume of melt that eventually crystallized to form PAT 91501.

Unless this rock happened to capture a snapshot of metal-sulfide particles sinking and vesicles rising, the retention of any vesicles or metal-sulfide requires another explanation. Far from being dominated by gravitational settling or rising under the influence of buoyancy alone, we suggest that the system was also influenced by the movement of melt within the fracture and the binding of dense metal-sulfide and buoyant vesicles to produce particles of near-neutral buoyancy. When the melt was injected into the cold country rock, it began to rise due to the marked thermal difference between the
Using the method of Wilson and Head (1981), we calculate that the melt rose at a velocity of 0.028 m/s through the dike and solidified due to cooling after migration of ~220 m (McCoy et al., 2006). At this rate, the magma solidified after rising through the dike for ~2 hours. Importantly, the rate of rise of the melt through the dike was roughly an order of magnitude faster than the rate of metal-sulfide settling or vesicle rise. Thus, settling of metal and sulfide to the bottom of the dike was inhibited by the rapid rise of melt through the dike.

It also bears noting that the gravitational vector inferred from the metal-sulfide contacts is inconsistent with the orientation of the vesicle long axes (Fig. 4). This slight offset may result from minor turbulence in the rising magma, or possibly an additional lateral component of melt movement that would be reflected in the vesicle shapes but not the gravitational settling of the metal.

Finally, the preservation of metal-sulfide-vesicle assemblages may result from the offsetting differences in density and buoyancy. It is interesting to note that the upward velocities of the average-sized gas bubbles responsible for the vesicles and the downward velocity of the average-sized metal-sulfide grains are very similar at all temperatures. If surface tension forces bind bubbles and grains of comparable size together, offsetting buoyancy may be created that would cause the linked bubbles and sulfide grains to be suspended, or at least to move only very slowly, in the melt. Neutral buoyancy has been suggested for magnetite and vesicles in the Bishop Tuff, where a vesicle either scavenged magnetite crystals from the melt or served as a nucleation point for magnetite growth, in the pre-eruptive magma (Gualda and Anderson, 2007). The attainment of neutral
buoyancy in the upward moving melt from which PAT 91501 crystallized might explain the retention of even large metal-sulfide particles.

### 6. CONCLUSIONS

Among the abundant impact melt rocks and breccias from the L chondrite parent body, PAT 91501 is unique in exhibiting cm-sized metal-sulfide particles and vesicles, for the remarkable alignment of these particles, and for its ancient age. Sulfur volatilization must have been a ubiquitous process during impact melting of chondritic materials and other meteorites (e.g., Chico) are known that reasonably sample impact melt dikes injected into the crater basement. These other meteorites do not exhibit the large vesicles seen in PAT 91501. This sample must have formed by a combination of a particularly large, early impact on the L chondrite parent body that formed unusually wide, clast free melt veins where the combination of relatively slow cooling and crystallization, coalescence and rise of vesicles, coalescence and sinking of metal-sulfide particles, formation of metal-sulfide-vesicle aggregates creating neutrally buoyant assemblages, and upward flow of magma in the dike. Although similar processes must have occurred in the formation of other chondritic impact melt rocks, they did not combine in the unique combination that formed PAT 91501.

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EAR-0345710 and EAR-0113480.
REFERENCES


Table 1. Noble gas abundances in two samples of PAT 91501

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<th>Sample, °C</th>
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<th>$^4\text{He}$</th>
<th>$^{20}\text{Ne}$</th>
<th>$^{21}\text{Ne}$</th>
<th>$^{22}\text{Ne}$</th>
<th>$^{36}\text{Ar}$</th>
<th>$^{38}\text{Ar}$</th>
<th>$^{40}\text{Ar}$</th>
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<td>8.58</td>
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<td>0.12</td>
<td>1.15</td>
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Table 2. Abundances of cosmogenic species and cosmic-ray exposure ages of PAT 91501.

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<th>Sample, °C</th>
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<th>$^{38}\text{Ar}$</th>
<th>$^{20}\text{Ne}/^{22}\text{Ne}$</th>
<th>$^{22}\text{Ne}/^{21}\text{Ne}$</th>
<th>$^9\text{Be}$</th>
<th>$^{26}\text{Al}$</th>
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</table>

Noble gas concentrations in $10^4$ cm$^3$ STP/g. Cosmic-ray exposure ages, $T$, in Myr. $^9\text{Be}$ and $^{26}\text{Al}$ activities in dpm/kg; uncertainties are estimated to be ±7%. $T_{10-21}$ and $T_{26-21}$ after Graf et al. (1990a).
Figure Captions

Figure 1. Photograph of PAT 91501,50. Numerous vesicles and metal-sulfide grains (up to cm-sized) are visible on the cut surface. Cracks throughout sample are likely due to terrestrial weathering. Scale cube is 1cm on a side.

Figure 2. A) Reflected light optical photomicrograph of an intergrown metal-sulfide particle in contact with a vesicle in PAT 91501, (111). The particle has been etched to show the metallographic texture consisting of mainly taenite (t) and martensite (m). Troilite (tr) with small particles of embedded pentlandite (p) rims the entire particle. B) Nickel composition (following traverse illustrated in A) across the two domains showing high-Ni inclusion-free (or poor) taenite rims grading into intermediate-Ni martensitic cores.

Figure 3. A single frame from the 3 dimensional rotation visualization made from the CT scan of PAT 91501, (50), in which vesicles and metal-sulfide intergrowths are highlighted. Metal (yellow), sulfide (magenta) and vesicles (blue bubbles) are set in a semi-transparent outline of the specimen pictured in Figure 1. Arrow points to prominent, large vesicle seen in Fig. 1. The specimen is oriented as it would have been at the time of crystallization as suggested by the metal-sulfide orientations (sulfide above metal in all instances). Note, however, that long axes of vesicles and metal-sulfide masses are offset somewhat to the left.

Figure 4. Stereo plots from PAT 91501, (50) of a) the normals to the planes defined by the contact between metal and sulfide with the size of the each circle proportional to the area of the contact and b) orientations of the vesicle long axes with the circle areas proportional to vesicle volume. Clustering of orientations are observed for
both metal-sulfide contacts and vesicle elongation. See text for discussion of orientation directions.

Figure 5. Ar-Ar ages (Gyr, rectangles, left scale) and K/Ca ratios (stepped line, right scale) as a function of cumulative release of $^{39}$Ar for temperature extractions of a melt sample of PAT 91501. Seven extractions releasing 30-78% of the $^{39}$Ar give an age of 4.461 ±0.008 Gyr, which we interpret to be the formation time of the PAT 91501.