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K–Ca ages of authigenic sediments: examples from Paleozoic glauconite and applications to low-temperature thermochronometry

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Abstract K–Ca ages of Cambrian glauconites from the Llano uplift, central Texas, were determined in order to reevaluate the ability of the K-Ca system to constrain the timing of deposition of sedimentary packages. All of the K-Ca ages presented here were found to be younger than their stratigraphic ages. In addition to being too young, the K-Ca ages are also highly variable, ranging in age from Silurian to Permian. The oldest subset of glauconite ages are in agreement with previously published Rb-Sr ages from the same outcrop and provide further evidence for there having been a post-depositional thermal or recrystallization event that reset both the Rb-Sr and K-Ca systems. The range of younger glauconite K-Ca ages is similar to the distribution of available apatite fission track ages for the Llano basement. K-Ca ages are interpreted as thermochronologic data reflecting partial retention of Ca in thermally fluctuating basin conditions. Estimates of the closure temperature of Ca in glauconite are found to be 60–90°C for cooling rates of ~0.5–1°C/My. The K–Ca system is potentially useful as a low-temperature thermochronometer with closure temperatures <100°C for glauconite.

Keywords K–Ca isotopes · Thermochronology · Sedimentary rocks

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Introduction

Reliable dating of sedimentary rocks is important to the calibration of the geologic time scale and to the timing and cyclicity of depositional events. Determining the ages of sedimentary units is difficult, however, unless they contain useful biostratigraphic markers or interbedded volcanics (e.g. Odin 1982). It is also made difficult by the fact that sedimentation can occur over relatively lengthy amounts of geologic time. For that reason, direct dating of deposition/ sedimentation is not always possible, but can be closely constrained through radiometric dating of authigenic and diagenetic sedimentary minerals. Depending upon the minerals targeted, however, the diagenetic age can be highly variable and considerably younger than the depositional age (e.g. Brookins et al. 1980; Obradovich 1988). It is not clear how much of that variability is a result of depositional processes and how much is a function resetting during the subsequent thermal history of the sedimentary basin (Evernden et al. 1961).

Both the Rb–Sr and the (K)Ar–Ar systems have been shown to commonly yield erroneously young ages, most likely due to the loss of radiogenic daughter products through diffusion or ion exchange processes (e.g. Hurley et al. 1960, 1961; Thompson and Hower 1973; Morton and Long 1980; Brookins et al. 1980). Given the stable, nonvolatile nature of calcium, the relatively short half-life of ⁴⁰K, and the fact that potassium and calcium are both major constituents in rock-forming minerals, the K–Ca system has great potential for geochronometric applications (Marhsall and DePaolo 1982). Comparative studies examining the K–Ca system have suggested that Ca is a more immobile and retentive daughter and K–Ca ages are more resistant to resetting (Marshall et al. 1986; Shih et al. 1994; Nagler and Villa 2000). K–Ar ages of authigenic feldspar

(Marshall et al. 1986) and Ar-Ar ages of pegmatitic micas (Nagler and Villa 2000) are consistently younger than the corresponding K-Ca ages, due to diffusive loss of ⁴⁰Ar, even at low temperatures. Likewise, Rb-Sr mineral isochron ages of lunar granites are younger than K-Ca isochron ages of the same granites (Shih et al. 1994). Although there are studies indicating that K-Ca geochronology is particularly useful for sedimentary dating (e.g. Polevaya et al. 1958; Marshall et al. 1986; Baadsgaard 1987; Gopalan 2008), published K-Ca sedimentary dates are relatively few. Because ⁴⁰Ca is the most abundant stable isotope of calcium, enrichments in radiogenic ⁴⁰Ca are relatively small and only measurable in materials that are old (Paleozoic-Precambrian) and/or have high K/Ca values. A new analytical method for measuring radiogenic Ca ratios using multi-collector ICP-MS has contributed to improvements in rapid and precise Ca isotopic analysis (Cecil 2009).

Here, we present new K–Ca ages from authigenic Cambrian glauconites from the Llano Uplift as a means for evaluating the utility and robustness of the K–Ca system in dating sedimentary minerals. The principal findings are that the K–Ca ages are younger than the sedimentary deposition of dated units, but they may record heating during diagenesis and cooling below closure temperatures less than 100°C, thus making the K–Ca system a potentially useful low-temperature thermochronometer.

Approaches to sedimentary geochronology

Sedimentary minerals having chemistries appropriate for geochronometric dating are commonly targeted to constrain depositional timing. Recent innovations in this field include U-Pb dating of diagenetic xenotime (McNaughton et al. 1999), monazite (Evans et al. 2002) and organic-rich calcite (Becker et al. 2002). More classical approaches have focused on sedimentary minerals rich in K (and Rb) and are therefore suitable for dating using K-Ar, Ar-Ar and Rb-Sr techniques. Such minerals have included authigenic K-spar (Marshall et al. 1986), illite (Clauer et al. 1997; Srodon et al. 2002) and K-rich evaporites such as sylvite, carnallite, langbeinite or polyhalite (e.g. Baadsgaard 1987; Brookins et al. 1980; Renne et al. 2001). Most of these techniques have proven problematic and have achieved only moderate success. In the case of illite, it can be difficult to distinguish detrital from authigenic populations and in the case of evaporite samples, dating is difficult given the readiness of radiogenic ⁴⁰Ar to diffuse out of certain K-salt species (Brookins et al. 1980).

Perhaps the most commonly used mineral for dating of siliciclastic sediments is glauconite, a K-rich phyllosilicate that forms from clay precursors in a marine environment (Rongbing 1989; Clauer et al. 1992; Stille and Clauer

1994: Meunier and El Albani 2007). In fact, almost half of the absolute ages used in constructing the Mesozoic and Cenozoic time scale are derived from Ar-Ar ages of glaucony series minerals (Smith et al. 1998). Glauconites are particularly attractive targets because they are common in sediments of all ages from the Precambrian throughout the Phanerozoic, are easily recognized in the field, and can be determined to be authigenic on the basis of their chemical, textural and X-ray characteristics (Odin 1982). Glauconite age data, however, are often viewed with skepticism (e.g. Obradovich 1988), as ages are frequently discrepant between systems and are overall too young (Hurley et al. 1960; Hurley 1961; Thompson and Hower 1973; Morton and Long 1980) and sometimes too old (Owens and Sohl 1973; Montag and Seidemann 1981). Leaching experiments for Rb-Sr analysis (Morton and Long 1980) and single grain Ar-Ar laser probe analysis (Smith et al. 1998) have not significantly improved the reliability of glauconite ages.

Several early pioneering studies were conducted, in which K–Ca ages of lepidolites (Ahrens 1951), glauconite (Herzog 1956), and sylvite (Polevaya et al. 1958) were determined. K–Ca geochronology is seldom used, however, due to difficulties measuring Ca isotopes with conventional TIMS methods, and the general difficulty of measuring sedimentary deposits, which form slowly and are prone to alteration and non-closed system behavior. A handful of more recent studies using K–Ca to date K-rich evaporites (Baadsgaard 1987) and glauconites (Gopalan 2008) have emerged, but K–Ca sedimentary geochronological data are relatively few.

Sample preparation and analytical methods

Isotopic and geochronologic data from seven glauconite samples and six K-rich evaporite samples are presented here. Glauconitic sandstone samples were hand crushed, sieved and divided into the following size fractions: 63-105, 105-150, 150-247, and 247-420 µm, which correspond to samples named GL 105, GL 150, GL 247, and GL 420, respectively. Glauconites were separated from the size fractions via magnetic separation using standard techniques described in Clauer et al. (2005). Two glauconite aliquots from each size group were powdered and approximately 15-25 mg of sample powder were weighed and transferred to 15 mL screw top Teflon vials. We performed leaching experiments on one set of aliquots by first rinsing the samples with acetone and water and then putting them in 1 mL of 1 M HCl. Samples were leached for \sim 15 min, including 5 min of ultrasonication, and then centrifuged. The leachates were saved for analysis, and the residues, along with non-leached samples, were dissolved

following the silicate dissolution procedure outlined in Cecil (2009).

Once fully dissolved, all samples were spiked with ⁴⁴Ca and ⁴¹K tracers and allowed to homogenize for 24 h on a hot plate at 120°C. K and Ca were separated using 1-cmdiameter quartz columns filled with Dowex AG50 W-X8 resin (200–400 mesh) to a height of 17 cm. Elemental separates were evaporated to dryness and later redissolved and diluted in 2% HNO₃.

Isotopic measurements were performed on a GV IsoP-robe MC-ICP-MS, using gas-phase reactions in a hexapole collision cell to minimize isobaric Ar interference (Cecil 2009). All samples were introduced in 2% HNO₃ through an Aridus desolvating nebulizer. Sample ${}^{40}Ca/{}^{42}Ca$ ratio measurements were corrected using a sample-bracketing technique, made possible by the good reproducibility (0.02% RSD) of the standard ratios. ${}^{39}K/{}^{41}K$ ratios were also normalized to the standard ratio (Cecil 2009).

K-Ca dating of Cambrian glauconite

Glauconite samples were collected from the Upper Cambrian Lion Mountain sandstone (Riley Fm) in the Llano uplift, central Texas. The Llano uplift is a Proterozoic basement high, which exposes polydeformed metamorphic and granitic rocks associated with a Grenvillian orogenic belt bordering the southern margin of Laurentia (Mosher 1998). It is rimmed by Cambrian–Pennsylvanian terrigenous and shallow marine sedimentary units that dip gently away from the basement uplift, forming a broad anticlinorium with the older (basement) rocks in the core. Apatite fission track age distribution in the basement rocks forming the core of the Llano uplift have Paleozoic ages, showing that the core of the Llano uplift was also buried during the deposition and diagenesis of the Paleozoic section (Corrigan et al. 1998) and that the burial depth was around 2 km. The Paleozoic section is in turn flanked by a thin on-lapping section of Cretaceous carbonates (Fig. 1). The 15-m-thick Lion Mountain sandstone is a dark green, medium-grained unit, composed primarily of glauconite pellets and larger carbonaceous nodules (McBride 1988) that lies conformably above the lower Riley Formation Hickory sandstone and Cap Mountain limestone members (see Morton and Long 1980, for a more detailed description of the stratigraphy).

Glauconite samples were separated into four size fractions and divided into two sets of aliquots for leaching experiments. The size separation was performed in order to test whether or not more mature or pure glauconite existed in a specific size range and to determine whether the abundance of radiogenic Ca and/or age was dependent on the size of the glauconite grains analyzed. Leaching was performed in order to remove loosely bound Ca and to determine the effect of common Ca on the K–Ca age. Seven K–Ca ages range broadly from 223 to 434 Ma (Table 1). There is no apparent correlation neither between age and grain size, nor between age and leached/unleached sample aliquots. It should be noted, however, that the three Permian-aged samples reported here were not leached. Leachates extracted during chemical sample preparation were analyzed and ${}^{40}Ca/{}^{42}Ca$ ratios were found to be indistinguishable from standard ${}^{40}Ca/{}^{42}Ca$ values, indicating that all of the Ca removed during leaching was likely common, loosely bound Ca from associated carbonate.

Comparison of K-Ca, Rb-Sr, and AFT ages

The oldest K-Ca ages (434 and 429 Ma) are identical within error with Rb-Sr ages from glauconites sampled at the same outcrop, as determined by Morton and Long (1980) (Fig. 2a). These Silurian ages are younger than the assumed depositional age of 515 Ma by approximately 16%. Morton and Long (1980) interpret the 430 m.y. age to represent the timing of maximum burial and recrystallization that occurred during deposition of at least 700 m of strata from the late Cambrian to the early Ordovician and before an inferred period of unroofing between the middle Ordovician and the early Devonian. Because all Rb-Sr ages cluster between 410 and 430 Ma, recrystallization is thought to chemically stabilize the glauconite, making it impervious to alteration and resetting of the Rb-Sr system. The fact that the oldest K-Ca ages are coincident with the Rb-Sr ages provides further evidence for there having been a Silurian diagenetic/recrystallization event. Unlike the clustered Rb-Sr ages, however, the K-Ca ages presented here are highly variable and are as young as 223 Ma.

Although the widely distributed K–Ca ages are not similar to Rb–Sr ages, the range of K–Ca ages (434– 223 Ma) is the same as the range of apatite fission track ages from the Llano basement (425–240 Ma) (Corrigan et al. (1998). Age information from the Precambrian basement and the overlying strata are not directly comparable; however, their close stratigraphic relationship (less than 200 m) implies a shared post-Cambrian burial history. Modeling of the wide range of Paleozoic apatite fission track ages and the negatively skewed nature of the track length distributions suggest a protracted burial history, with maximum burial (depths between 2 and 2.5 km, assuming a geothermal gradient of 25°C/km) occurring in the Permian (after Corrigan et al. 1998).

The distribution of the K–Ca ages corresponds to a period of time in which the Llano basement and overlying Cambrian units were at depths of 1–2 km and burial-related heating resulted in temperatures of at least 50–60°C, based on the integrated thermal history of the Llano uplift



Fig. 1 Generalized geologic map of the Llano uplift, Central Texas, showing the relationship between the Precambrian basement high and the surrounding Paleozoic and Cretaceous strata. K–Ca (this study) and Rb–Sr (Morton and Long 1980) ages presented in Fig. 2 are from

Cambrian sandstones sampled at the location marked with a *star*. Apatite fission track (AFT) ages (Corrigan et al. 1998) are marked with *solid circles*. *B* Lake Buchanan, *LBJ* Lake LBJ. Modified from Morton and Long (1980), Corrigan et al. (1998), and Mosher (1998)

Table 1 K-Ca analytical ages of Cambrian Lion Mountain glauconite

Sample name*	⁴⁰ Ca/ ⁴² Ca norm**	K (ppm)	Ca (ppm)	K/Ca	⁴⁰ K/ ⁴² Ca	€ _{Ca}	Age [†]
GL 105	151.184	39815.90	1077.66	36.95	0.682	11.14	434 ± 9
GL 150	151.315	54122.28	442.46	122.32	2.259	19.8	246 ± 11
GL 247	151.288	36949.89	309.28	119.54	2.206	18.02	230 ± 10
GL 420	151.273	36949.89	316.64	116.69	2.151	17.04	223 ± 6
GL 105L	151.223	34141.50	502.89	67.89	1.239	13.7	429 ± 10
GL 247L	151.232	27581.43	360.38	76.53	1.393	14.29	285 ± 7
GL 420L	151.125	15183.97	515.81	29.44	0.536	7.19	365 ± 9

* The numbers in the sample names correspond to the size fractions described in the text. The "L" indicates that the sample was leached for 15 min with 1 M HCl

** Normalized to the six standards bracketing the sample analysis, which in turn are normalized to a 40 Ca/ 42 Ca ratio of 151.016 (Marshall and DePaolo 1982)

[†] Errors are a function of the 1σ deviation of the mean standard isotopic ratios bracketing the sample analysis

from Corrigan et al. 1998 (Fig. 2b). We interpret this as indicating that the K–Ca system in glauconite is sensitive to partial retention at temperatures greater than 50°C, particularly if those elevated temperatures are sustained for periods of geologic time on the order of 10^7-10^8 years. This suggests that K–Ca is behaving as a low-temperature thermochronometer, and the glauconites studied here are being held in a partial Ca retention zone for much of the

mid-late Paleozoic. The lack of K–Ca ages younger than 223 Ma indicates that an unroofing event had occurred by mid-Triassic time, and glauconites experienced no further reheating to greater than 50°C. This is consistent with the timing of post-Ouachita orogenic cooling and exhumation described in Corrigan et al. 1998.

The K–Ca isochron age of all glauconite fractions measured in this study is 375 \pm 50 Ma. This isochron can





be reproduced by about 30% partial retention of Ca for first of 300 My of glauconite residence in the upper crust and 223 My of continuous decay and complete retention of Ca in the glauconite mineral structure (Fig. 3).

Closure temperature of Ca in glauconite

Preliminary estimates of the closure temperature of Ca in glauconite were made using the classic relationships described by Dodson (1973):



Fig. 3 K–Ca average isochron age for all fractions of glauconite analyzed in this study. The *dashed line* represents a modeled isochron assuming 223 My of decay and 100% retention of Ca and 300 My of decay and 30% of Ca retention, as discussed in text

$$E/RT_c = \ln \left(A\tau D_o/r^2\right)$$

and

 $\tau = R T_c^2 / E \left(\mathrm{d}T / \mathrm{d}t \right)$

where *E* is activation energy, *R* is the universal gas constant, T_c is closure temperature, *A* is a geometric constant (A = 27, using a cylindrical model), $D_0 = \text{frequency factor}$, *r* is the effective diffusive radius, and dT/dt is cooling rate. T_c of Ca in glauconite can only be loosely estimated, as very little is known about the parameters governing glauconite. An activation energy (*E*) of 4.182 kJ/mol is chosen based upon the determined activation energy of Sr in biotite (5.02 kJ/mol–Dodson 1973), and the estimated ratio of E_{Sr}/E_{Ca} in micas (Fletcher et al. 1997). Frequency factor (D_0) is ill constrained as there is no diffusion information presently available for Ca in glauconite. We assign a D_0 value of 1, intermediate between estimated D_0 for Sr in biotite (Fletcher et al. 1997) and Ar in muscovite (Harrison et al. 2009).

Unlike in most other systems where the mineral grain itself functions as the effective diffusive domain, the complexity of the glauconite structure makes it difficult to measure the diffusive radius. Glauconite forms by the uptake of K and Fe in degraded micaceous silicates. It does so most readily in micro-reducing environments created by the empty shells of marine microorganisms (Clauer et al. 1992; Stille and Clauer 1994). Morphologically, therefore, glauconite tends to occur as rounded, sand-sized grains, or pellets that are typically 0.1-0.5 mm in size. (For reference, the glauconite grains studied here were sieved into size fractions ranging from 63 to 420 µm). The individual glauconite crystallites that are forming inside of the pellets,



Fig. 4 Closure temperature of Ca in glauconite as a function of cooling rate. *Curves* were modeled using classic relationships described in Dodson, 1973. The *gray rectangle* represents the range of temperatures (inferred from apatite fission track data–Corrigan et al. 1998) thought to be partially resetting Cambrian glauconites

however, are much smaller and are capable of adsorbing and expelling ions during their slow maturation (Stille and Clauer 1994). There are numerous internal glauconite morphologies, but the most commonly described are randomly oriented mica flakes. Although the dimensions of the glauconite microcrystals are almost certainly variable, they could be micron to submicron in size. It is not known whether or not the exterior pellet acts as a diffusive boundary or if diffusion is occurring on a smaller scale within the pellet.

Closure temperatures for Ca in glauconite were modeled as a function of cooling rates using the parameters described earlier (Fig. 4). The different curves represent the relationship between T_c and dT/dt for four different effective diffusive radii, ranging from 1 to 1,000 microns. Typical pellet sizes are in the 100 to 1,000 micron range, whereas microcrystalline glauconitic clays within the pellets might be 1 to 10 microns in size (or conceivably smaller). Closure temperatures at a given cooling rate are higher for larger effective diffusive radii and lower for smaller diffusive domains. Estimates of closure temperature of Ca in glauconite are highly dependent upon the cooling rate and diffusive radius chosen. Apatite fission track data from the Llano basement are indicative of a slow, protracted cooling history (Corrigan et al. 1998). Erosion of the Llano region was likely slow for much of the Paleozoic, as basement rocks appear not have been exhumed to levels above the partial annealing zone for AFT. For that reason, we estimate cooling rates to be in the range of 0.5-1°C/My. Given the sizes of the glauconite grains analyzed (63–420 μ m), and the probability that Ca loss is occurring at scales smaller than the pellet size, 10 μ m is chosen as the preferred *r* value. For those cooling rate and diffusive radius parameters, the closure temperature of Ca in glauconite is around 85°C. These temperatures are slightly higher than the temperatures estimated to partially reset the K–Ca system in glauconite based on AFT data. Although they cannot be considered definitive values, because so little is known about Ca diffusion in glauconite, the broad range of closure temperatures presented here is generally very low and supports the concept of K–Ca behaving as a low-temperature thermochronometer in basin systems.

Discussion

Assuming that the ages reported here are meaningful, there are three plausible interpretations of the younger than depositional ages measured in this study are as follows: (a) protracted authigenic growth of glauconite over large geologic periods, (b) late K addition, via diagenetic processes, and (c) diffusive Ca loss partially above a closure temperature.

It is widely accepted that glauconite grows at the sweater–sediment interface over extended periods of time (Smith et al. 1998). However, protracted authigenic growth of glauconite takes place over time periods of hundred of thousand of years to as much as 5 My (Odin and Dodson 1982; Smith et al. 1998; Meunier and El Albani 2007), and not tens to hundreds of million years. We therefore rule out the hypothesis that the distribution of ages is related to extended authigenic growth.

Late addition of potassium during diagenesis is more difficult to quantify. Qualitative indicators such as freshness of hand samples argue against significant chemical disturbance during diagenesis in the studied rocks. In addition, the concentration of K (%) in our glauconites is well within the range (1–8%) of authigenic glauconites (Thompson and Hower 1973), suggesting it is unlikely that these samples have been subjected by significant uptake of potassium after deposition. We therefore suggest that the studied glauconites have not been subjected to intake of potassium subsequent to their initial growth during the Cambrian.

The third hypothesis, diffusive Ca loss is the most likely explanation for the distribution of ages, and is consistent with independent geologic and thermochronologic constraints from the region. The average isochron age for the studied samples is 375 Ma, which requires that about 30% of the radiogenically produced Ca was retained in the glauconite structure prior to unroofing of these rocks at the end of Permian (Fig. 4).

Conclusions

K-Ca ages from Cambrian glauconites are highly variable and demonstrate that the K-Ca system cannot be used to reliably reproduce depositional ages of this sedimentary package. In the examples presented here, K-Ca ages are younger than presumed depositional ages by 16-56%. The distribution of K-Ca ages in glauconite, however, can yield useful information about the thermal and hydrologic histories of basins. Data from the Llano uplift suggest that the mobility of Ca in glauconite is sensitive to burial to shallow crustal depths and thermal fluctuation, such that Ca ions exchange out of the mineral structure at temperatures above $\sim 50^{\circ}$ C. This is slightly lower than closure temperature estimates of Ca in glauconite, which range from ~60 to 100°C for slow cooling rates of 1°C/My. More rigorous work investigating the diffusive behavior of Ca in micas needs to be done, but on the basis of preliminary results presented here, K-Ca in glauconite has the potential to be used as a low-temperature basin thermochronometer.

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