MONAZITE, CRICHTONITE, AND AESCHYNITE IN THE CHICXULUB IMPACT CRATER PEAK RING.

Axel Wittmann¹, ¹Eyring Materials Center, Arizona State University, Tempe, AZ, USA, <u>axel.wittmann@asu.edu</u>.

Introduction: A recent drilling into the peak ring of the K-Pg Chicxulub impact structure recovered granitic basement lithologies with zircon U-Pb crystallization ages between 315 and 330 Ma [1,2] and an average shock metamorphic overprint of 17±1 GPa [3]. This study reports the first descriptions of monazite-(Ce), crichtonite-group, and aeschynite-group minerals from the Chicxulub impact crater with implications for their petrogeneses.

Samples and Methods: I used the JEOI JXA-8530F electron microprobe at ASU to generate X-ray intensity maps for petrographic thin sections from 27 samples of granitic basement rocks of the Lower Peak Ring section [747–1335 meters below sea-floor (mbsf)] in the IODP-ICDP Expedition 364 drillcore M0077A [4]. Quantitative wavelength dispersive spectrometry analyses were performed after confirming the presence of all major and minor cations in the targeted minerals. Monazite analyses followed a modified analytical protocol of [5] using the Probe for EPMA software, including the calculation of (U,Th)-Pb ages.

Results: Monazite occurs as a strongly fractured, 20 μm grain intergrown with allanite and epidote in a granite sample at 1201.3 mbsf, and as a strongly fractured, 60 μm grain in the granite portion of a sample at 952.75 mbsf, ca. 2 cm distance to an impact melt vein. A cluster of euhedral, 5 μm to 30 μm monazite grains in cataclased aplite dike sample at 1196.61 mbsf is associated with allanite, epidote, and thorite. Well preserved, 50 to 170 μm monazite-(Ce) crystals were only found in aplite dike sample at 875.68 mbsf; they are associated with epidote and exhibit two to three sets of planar fractures. EMPA spot analyses indicate compositions of $(Ce_{0.26-0.38}Th_{0.16-0.33}U_{0.01}Pb_{0.00-0.01}Y_{0.03-0.05}La_{0.11-0.20}Ca_{0.04-0.08}Pr_{0.03-0.04}Nd_{0.13-0.17}Sm_{0.03-0.05}Gd_{0.02-0.04}Dy_{0.01})_{\Sigma 1.02-1.14}(P_{0.69-0.87}Si_{0.11-0.26})_{\Sigma 0.91-0.98}$ and crystallization ages of 315 Ma ± 13 Ma $(2\sigma, n=11)$ for monazite in an aplite dike at 875.68 mbsf.

 $\frac{Crichtonite_group\ minerals}{(La_{0.07-0.08}Ce_{0.08-0.10}Dy_{0.10-0.11}Pb_{0.09-0.10})_{\Sigma_{0.35-0.37}Ti_{13.28-13.52}Fe_{5.79-6.03}Mn_{0.77-0.84}(V_{0.07-0.09}Si_{0.03-0.21}Ca_{0.04-0.08}Sc_{0.34-0.41}Na_{0.04-0.07}Al_{0.05-0.19}Mg_{0.00-0.15}Zn_{0.21-0.22})_{\Sigma_{0.85-1.28}O_{38}}]_{n=3}\ occurs\ in\ two\ samples,\ where\ the\ fractured\ 15\ \mu m\ crystals\ are\ associated\ with\ chlorite\ and\ titanite;\ crushed,\ 10\ to\ 30\ \mu m\ grammacioliite-(Y)\ [(Pb_{0.26-0.52}Sr_{0.23-0.38}U_{0.00-0.02})_{\Sigma_{0.62-0.89}}(Y_{0.14-0.32}Mn_{0.22-46}La_{0.00-0.06}Ce_{0.04-0.19}Pr_{0.00-0.03}Nd_{0.02-0.09}Sm_{0.01-0.04}Ca_{0.26-0.61}K_{0.01-0.09}Zr_{0.00-0.06})_{\Sigma_{1.04-1.40}(Ti_{13.25-13.75}}Zn_{0.13-0.24}V_{0.04-0.25}Al_{0.07-0.18}Nb_{0.03-0.06}Na_{0.04-0.09}Si_{0.07-0.42}Fe^{2+}_{1.59-2.05}Fe^{3+}_{3.59-4.39})_{\Sigma_{0.70}}\ _{Total_{\Sigma_{21.93-22.06}}}$

 $\frac{Aeschynite-group\ minerals}{Eu_{0.00-0.01}Gd_{0.01-0.02}Dy_{0.00-0.02}Er_{0.00-0.01})\sum_{1.03-1.17}(Nb_{0.60-0.29}U_{0.04-0.09}Y_{0.02-0.08}La_{0.02-0.03}Ce_{0.13-0.19}Pr_{0.02-0.04}Nd_{0.11-0.20}Sm_{0.02-0.04}Eu_{0.00-0.01}Gd_{0.01-0.02}Dy_{0.00-0.02}Er_{0.00-0.01})\sum_{1.03-1.17}(Nb_{0.60-0.79}Ta_{0.01-0.04}Ti_{0.75-0.93}Si_{0.30-0.41}Fe^{3+}_{0.13-0.29}Mn_{0.01-0.01}Na_{0.00-0.06}Cl_{0.01-0.10})\sum_{2.13-2.26} \frac{1}{Total}\sum_{3.19-3.31}]_{n=34}$ occur in aplite dike sample 875.68 mbsf as euhedral, fragmented, altered and likely hydrated 10 to 150 μ m crystals with allanite and rutile that contains up to 2.4 wt% Nb and 0.4 wt% Ta. Their Ce# and Nb# [6] indicate compositions of aeschynite-(Nd), aeschynite-(Ce), and nioboaeschynite-(Nd).

Discussion: The accessory minerals described here occur well preserved in a few samples of pegmatite and aplite dikes that were emplaced in close temporal association with their host granite [1,2] ~315 Ma ago. Crichtonite has only been described from late-magmatic stage mineralizations tied to tensile fissures and metasomatism [7,8]; grammacioliite-(Y) can crystallize at ~400 °C under metamorphic conditions [9]; this may indicate formation in shrinkage cracks that were exploited as pathways for granitic melt, pneumatolytic and hydrothermal fluids. Brittle fracturing of monazite, crichtonite, grammacioliite-(Y), and aeschynite suggests these phases experienced shock metamorphism during the Chicxulub impact. Because crichtonite and aeschynite-group minerals typically form as secondary minerals, they are evidence for the localized metasomatic and hydrothermal alteration of the granitic Chicxulub peak ring rocks prior to the impact 66 Ma ago. Alteration of aeschynite-group minerals may be tied to the hydrothermal system that was triggered by the Chicxulub impact [10]. Metasomatic and hydrothermal mineral assemblages that predate the impact complicate the interpretation of impact-related hydrothermal mineral assemblages and add complexity to the interpretation of radioisotopic disturbances in mineral chronometers in Chicxulub Lower Peak Ring granitic rocks.

Acknowledgements: I thank captain and crew, drilling team, scientists, and technical staff who participated in shipboard and/or shore-based operations for IODP-ICDP Expedition 364. This study used facilities at Eyring Materials Center at ASU supported by NSF grant NNCI-ECCS-1542160; A.W. received support from NSF grant OCE 1737087.

References: [1] Zhao J. et al. (2020) Gondwana Research 82:128–150. [2] Ross C. H. et al. (2022) GSA Bulletin 334:241–260. [3] Feignon J.-G. et al. (2020) Meteoritics & Planetary Science 55:2206–2223. [4] Morgan J. et al. (2017) Proceedings of the International Ocean Discovery Program 364:176 p. [5] Allaz J. et al. (2019) Microscopy and Microanalysis 25:30–46. [6] Yang Z. et al. (2001) European Journal of Mineralogy 13:1207–1214. [7] Stalder H. A. and Bühler Ch. (1987) Schweizerische mineralogische und petrographische Mitteilungen 67:93–102. [8] Potter E. G. and Mitchell R. H. (2005) Contributions to Mineralogy and Petrology 150:212–229. [9] Theye T. et al. (2010) European Journal of Mineralogy 22:443–452. [10] Kring D. A. et al. (2020) Science Advances 6:eaaz3053.