

Water in eclogitic garnet and clinopyroxene with oriented quartz and pargasite inclusions, W Norway

Dirk Spengler, AGH University of Krakow, PL

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Introduction

This study analyses 10 eclogites from an ultrahigh-pressure (UHP) metamorphic area in the Western Gneiss Region (WGR) of Norway. Each sample has oriented lamellar to acicular inclusions in clinopyroxene, which are either quartz with pargasite or quartz/albite without pargasite [1]. Oriented quartz in clinopyroxene is typically regarded to have formed by the consumption of vacancies in non-stoichiometric pyroxene, i.e. an isochemical breakdown of a Ca-Eskola component [2, 3]. If applicable then the close spatial association of quartz and pargasite suggests an isochemical origin for the pargasite inclusions too. The H₂O content of nominally anhydrous minerals (NAMs) was quantified and compared with petrological data to evaluate this hypothesis.

UHP rocks in the WGR

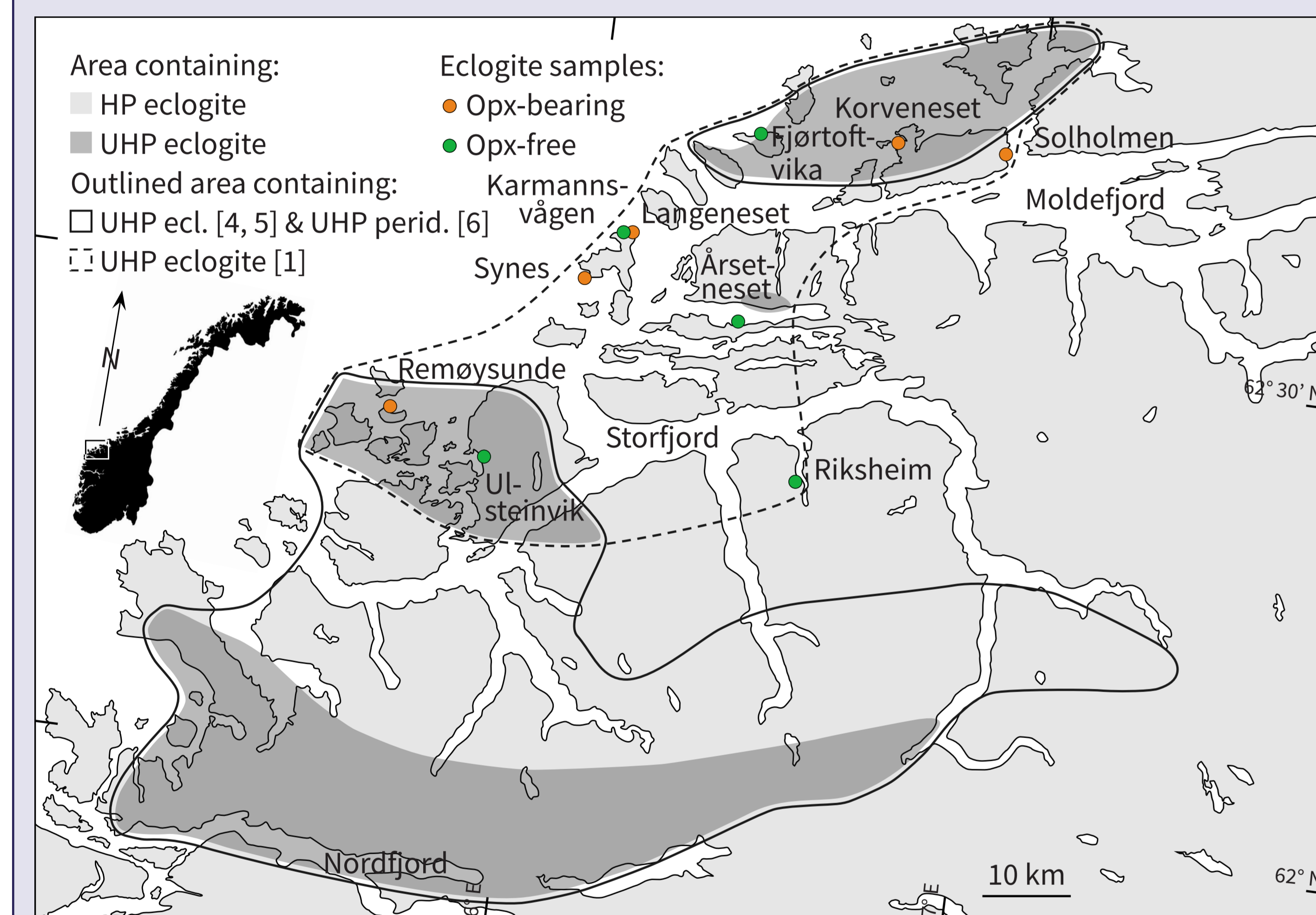


Fig. 1: Simplified map of the WGR that shows sample locations and known areas of UHP metamorphism from eclogite [1, 4, 5] and peridotite [6].

Mafic rocks (eclogite) define three large UHP domains that spread along the coast (shaded in Fig. 1). Ultramafic rocks (garnet pyroxenite enclosed in orogenic garnet peridotite) define UHP exposure that partially overlaps that of eclogite and partially fills the space in between (solid outline). Eclogite of the current sample set (dots) fills the gap between Storffjord and Moldefjord and has recently been shown to bear evidence for UHP metamorphism (dashed outline).

References

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Petrography of oriented inclusions

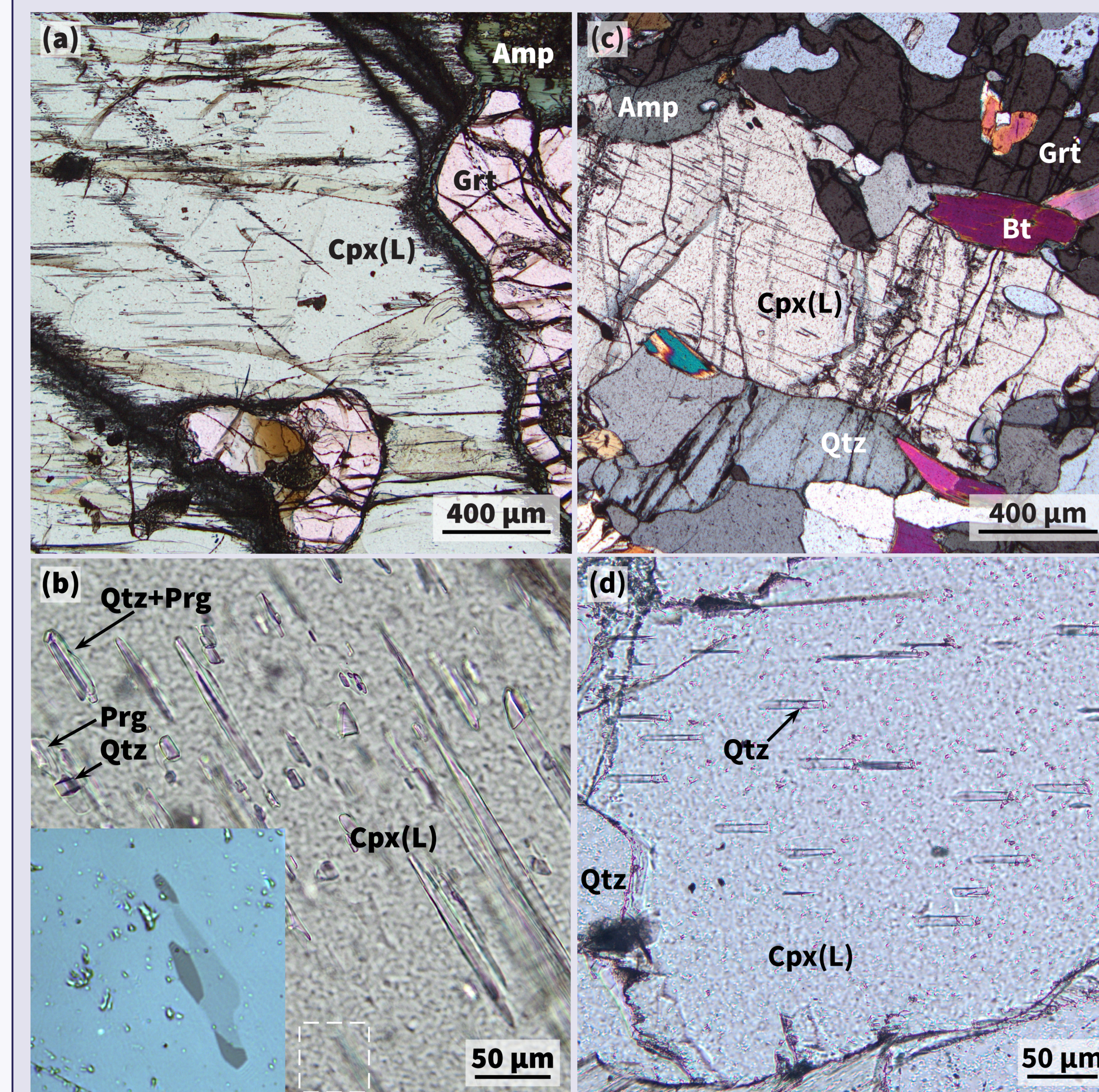


Fig. 2: Inclusions in clinopyroxene. (a, b) Bimineralline needles (PPL, Synes eclogite). Dashed frame shows the position of the inset (reflected light). (c, d) Monomineralic needles (c – PPL, d – nearly XPL, Årsetneset eclogite).

Spectra deconvolution

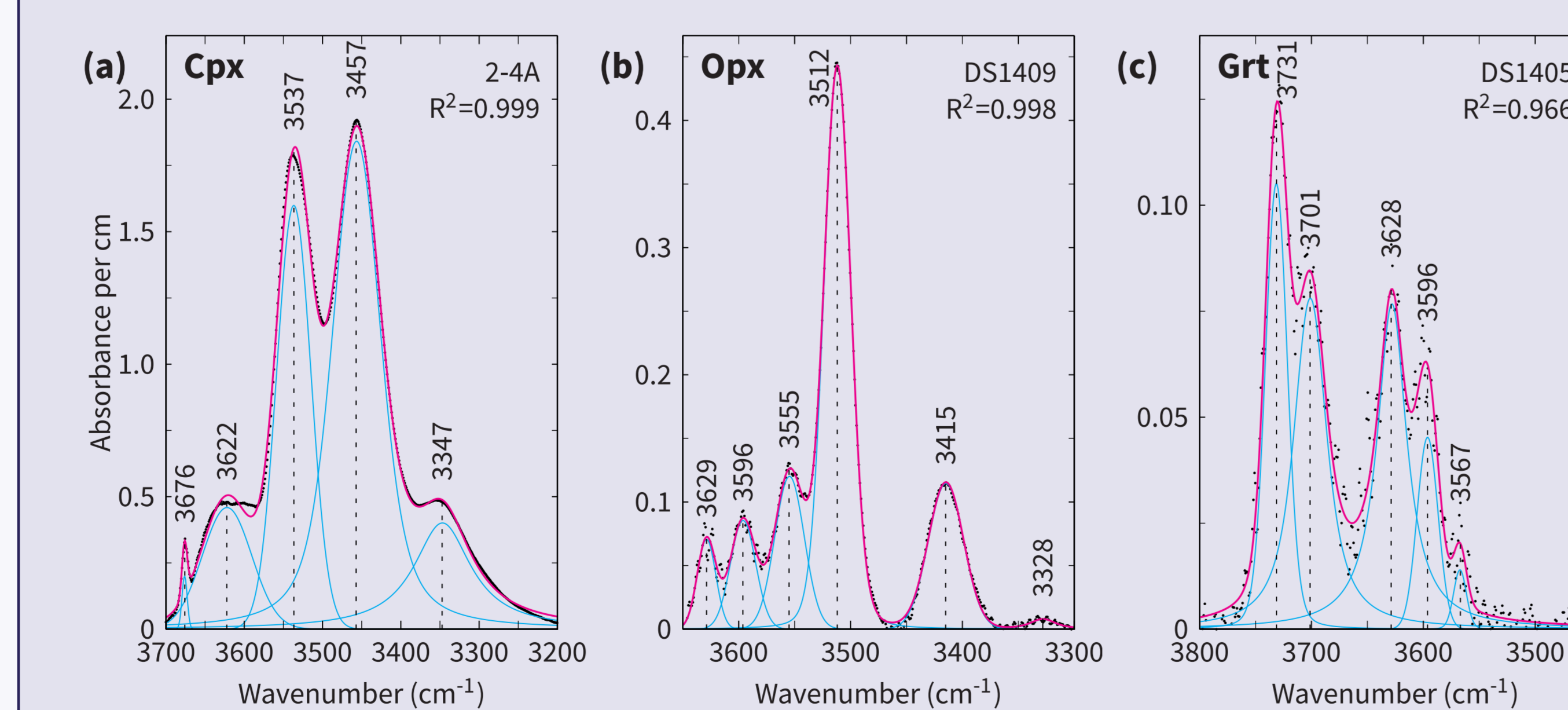


Fig. 4: Baseline corrected and deconvoluted spectra normalised to 1 cm.

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FTIR absorption spectra

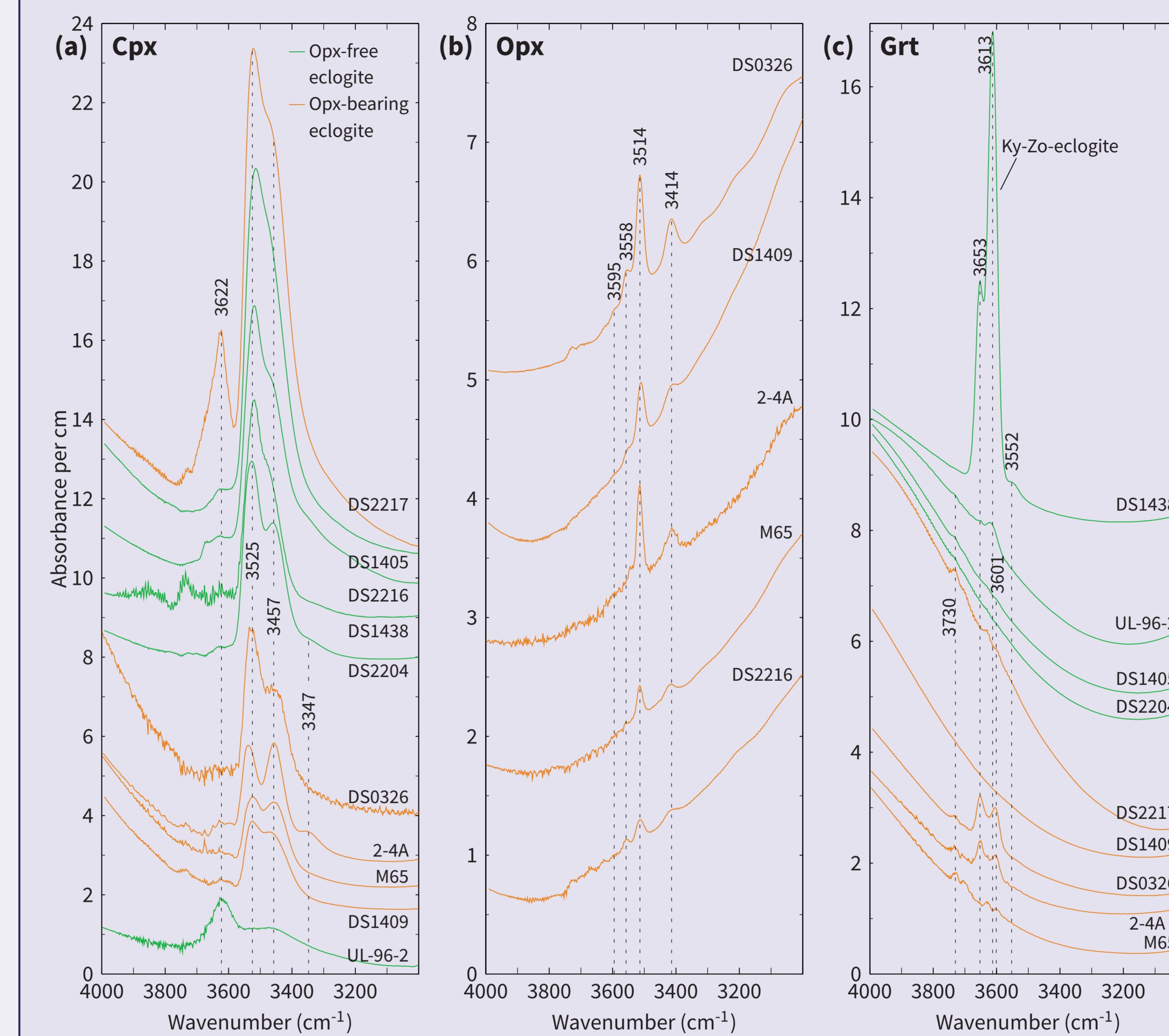


Fig. 3: Representative unpolarised FTIR spectra of (a) clinopyroxene, (b) orthopyroxene and (c) garnet (labels = sample numbers) in the O–H stretching frequency range normalised to 1 cm thickness and offset along the ordinate. Dashed lines indicate the position of selected peaks. Absorption in clinopyroxene with peak positions greater 3550 cm⁻¹ (band III of [7]) may relate to sheet silicate inclusions and were excluded from the quantification.

Quantified H₂O contents

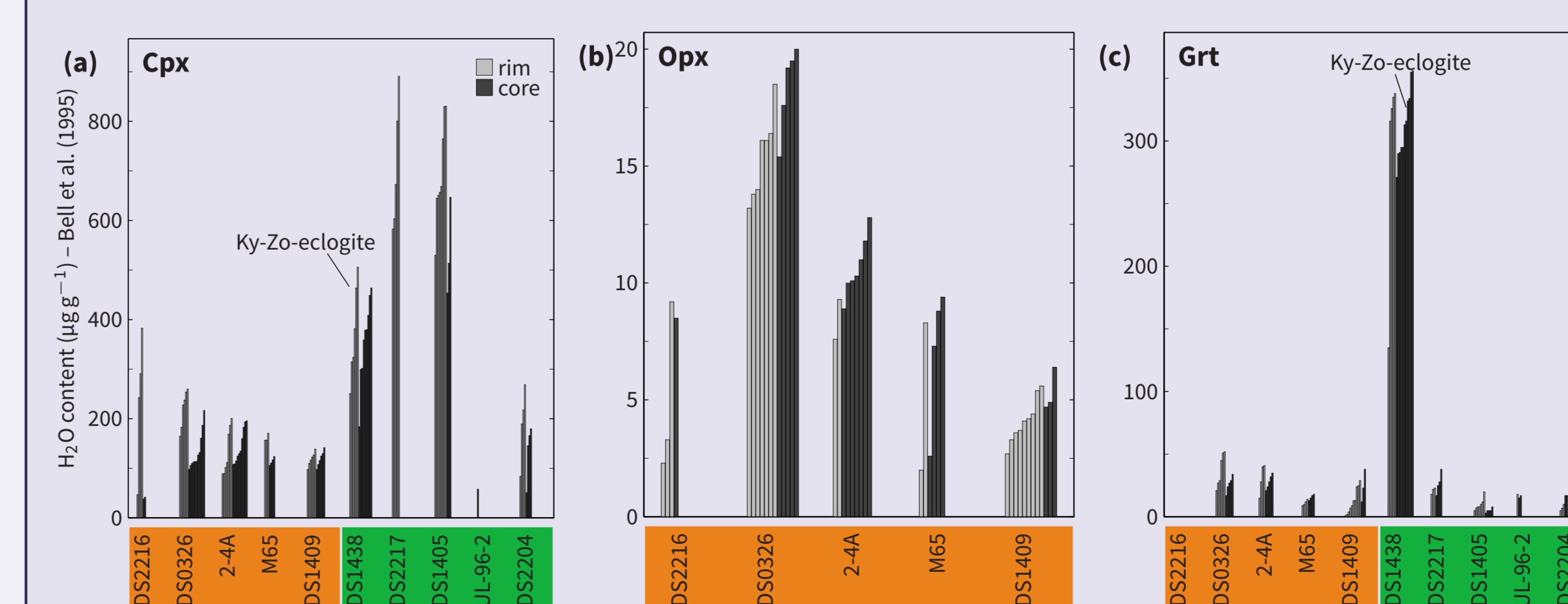


Fig. 5: Water contents in the grain interior (core) and the outermost 200 μm (rim) quantified using individual FTIR spectra and the calibration of [8].

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Discussion

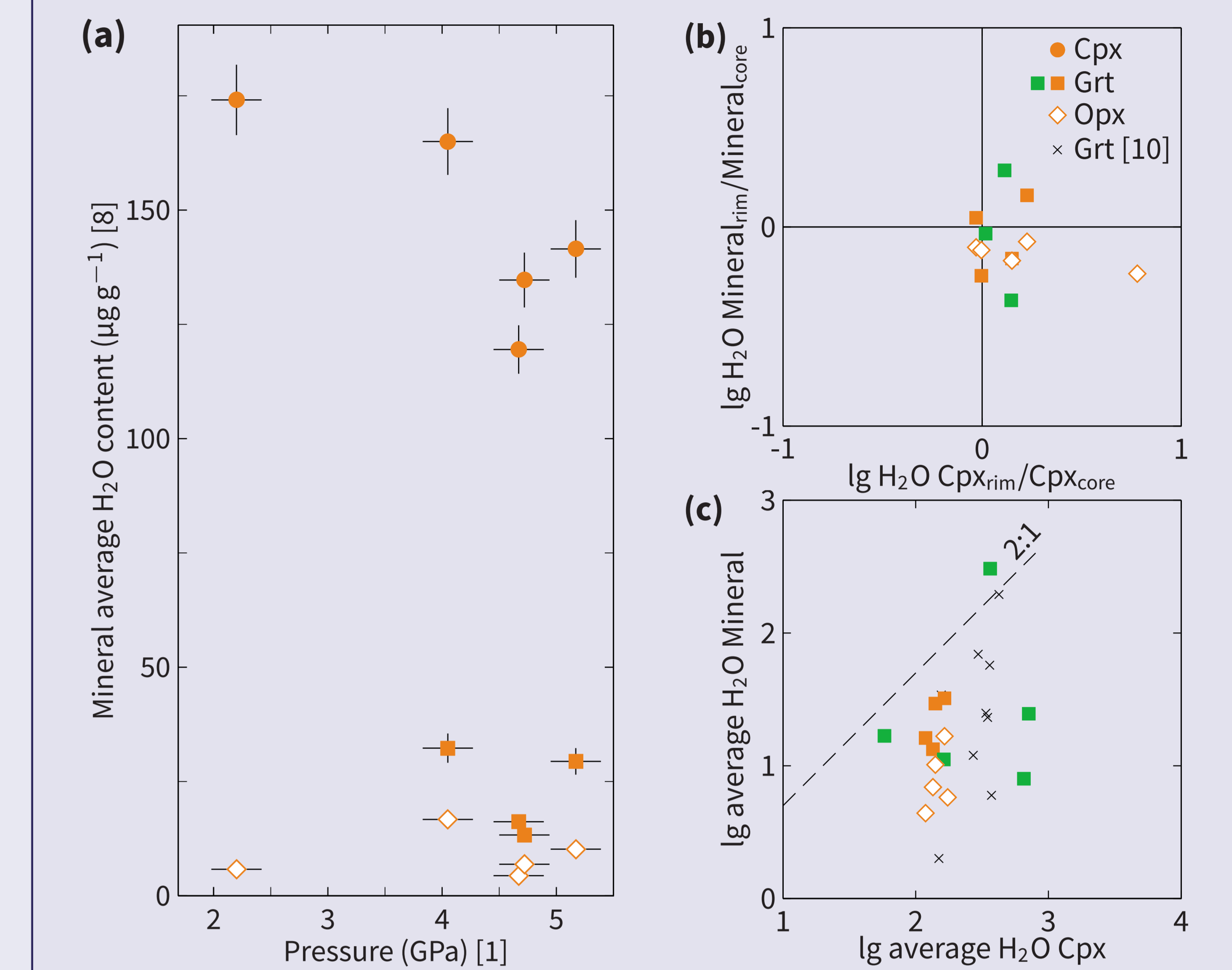


Fig. 6: Bivariate diagrams that show mineral average data for H₂O in NAMs. (a) H₂O content versus metamorphic pressure calculated from the mineral chemistry of orthopyroxene and garnet [1]. Uncertainties refer to 1σ. (b) Ratios calculated from the H₂O content of mineral rims and cores. (c) Mineral H₂O contents. Other information shows typical subsolidus H₂O partitioning between pyroxenes (dashed line, [9]) and with garnet (crosses, [10]).

Conclusion

- (1) The H₂O content of clinopyroxene correlates inversely with retrograde metamorphic pressure.
- (2) Orthopyroxene records H₂O loss at grain rims.
- (3) Precipitation of pargasite lamellae in clinopyroxene from penetrating external H₂O seems unlikely, because cogenetic orthopyroxene lacks chemical equilibrium in the amphibole stability field and shows non-typical H₂O partitioning.

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