

Lattice thermal conductivity of MgO periclase at high pressure

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The heat flow at the core-mantle boundary affects the core evolution, the thermal convection of the mantle and core, and geomagnetism from driving the geodynamo. The core-mantle boundary region is thermal boundary layer, therefore heat is transported dominantly by conduction from the core to the mantle. Previous studies have estimated thermal conductivity at the lowermost mantle in a wide range as 5-30 W/m/K (e.g. Lay et al. 2008). However, thermal conductivity of the lower mantle minerals is not well constrained yet due to a difficulty in measurement at high pressure condition. In this study, the lattice thermal conductivity of MgO periclase was obtained up to 100 GPa by measuring thermal diffusivity at room temperature in a diamond-anvil cell. We measured thermal diffusivity with the light pulse thermo-reflectance technique, which was recently developed (Yagi et al. 2011). This is an only method to measure the thermal diffusivity at high-pressure condition corresponding to the core-mantle boundary. Based on the pressure effect revealed in this study and the temperature dependence proposed previously (e.g. Hofmeister 1999), the lattice thermal conductivity of MgO periclase at the core-mantle boundary region was calculated. Combined with the values for MgSiO₃ perovskite and post-perovskite recently determined (Ohta et al. 2012), the lattice thermal conductivity at the base of the mantle was estimated. Here we will discuss the core-mantle heat flow.

Keywords: thermal conductivity, periclase, core-mantle boundary heat flow, thermal diffusivity

Single crystal elasticity of Pbnm-CaIrO₃, Cmcm-CaIrO₃, and Antigorite determined by inelastic X-ray Scattering

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Single crystal elasticity of Pbnm-, Cmcm- CaIrO₃ and Antigorite was measured at Spring8 BL35 by using inelastic X-ray scattering method. The former two phases are well known analog of perovskite (Pv) and post perovskite (pPv) MgSiO₃, respectively. The last one is important hydrous mineral in subduction slab.

Keywords: Perovskite, Post-Perovskite, Antigorite, crystal elasticity, inelastic X ray scattering

Tomographic properties from mantle convection in a 3-D spherical shell with the self-consistently calculated mineralogy

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We use numerical thermo-chemical mantle convection simulations in a 3-D spherical shell with the self-consistently calculated mineral physics to investigate various statistical properties using global tomography models (root-mean-square, cross-correlations and spectral amplitude of seismic anomalies). Four compositional models are assumed here, which has been already used in a paper published recently [Nakagawa et al., 2012]. In order to be the similar resolution to current global tomographic images, we use spherical harmonic filter technique instead of tomographic filtering technique [e.g. Davies, et al., 2012]. The truncation of spherical harmonic filter is used as degree of 16, 24, 32 and 40 to compare with the original resolution (96x288x96x2, which is equivalent to 384x192x96 or degree of 192 of spherical harmonic expansion). The statistical properties of tomographic image could be explained by thermo-chemical cases (e.g. RMS amplitude and anti-correlation in the deep mantle). However, it would be difficult to explain the ratio of V_s and V_p or V_b for thermo-chemical origin of mantle heterogeneities, which is similar profiles to isochemical cases. On the other hand, the ratio of V_s and density could explain the mineral physics constraint [e.g. Karato and Karki, 2001] with thermo-chemical origin of mantle heterogeneity. In conclusion, in order to assess global tomographic image as a consequence of mantle convection, RMS amplitude, cross-correlation and the ratio of V_s and V_b ratio might have useful information to understand what happens in a convecting mantle inferred from tomographic images in the deep mantle.

Keywords: Mantle convection, Self-consistently calculated mineralogy, Seismic tomography, Spherical harmonic filtering, statistical analysis

Numerical investigations of effects of spatial variations in physical properties on the mantle convection patterns

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1. Introduction

There exist large differences in temperature and pressure within the mantle of terrestrial planets. In the Earth's mantle, for example, a pressure increases by about 135GPa with increasing depth, while a temperature does by about 3500K. These large differences in temperature and pressure are expected to yield substantial variations in the physical properties of mantle materials and, hence, significantly affect the dynamic behaviors in the mantle. Indeed, as had been inferred by several earlier studies, the spatial variations in mineral properties play an important role in the internal structures observed in the mantle of the Earth and terrestrial planets. We will study how the spatial variation of physical properties of mantle materials affects the flow patterns of thermal convection.

2. Model and Procedure of numerical experiments

We carried out numerical experiments of thermal convection in highly viscous and incompressible fluids, in order to study the influences of the spatial variations in physical properties of fluids (viscosity, thermal conductivity and thermal expansivity) on the convecting flow patterns in the mantle of terrestrial planets. We present the results of a series of numerical calculations using (1) a linear stability analysis on the onset of thermal convection in fluids confined in planar layers, and (2) a nonlinear (finite-amplitude) time-dependent thermal convection in a two-dimensional Cartesian box of aspect ratio (width/height) of 6, with systematically varying the magnitude of (i) decrease in viscosity with temperature, (ii) increase in thermal conductivity with pressure (or depth), and (iii) decrease in thermal expansivity with pressure. By comparing the results with those in the presence of their spatial variations, we will discuss the changes on the critical conditions, dominant vertical flow structures, and the convection regimes caused by their spatial variations.

3. Results

From the changes in flow patterns with increasing the amplitudes of temperature dependence of viscosity, we successfully identified the transition into the 'stagnant lid' (ST) regime, where the convection occurs only beneath a thick and stagnant lid of cold fluid at the top surface. We also found by both linear and nonlinear numerical calculations, that the transition takes place regardless of the spatial variations in thermal conductivity and/or expansivity. However, detailed analysis of the numerical results showed a quantitative difference in the critical condition for the onset of ST convection due to the presence of spatial variations in thermal conductivity and expansivity. Especially we focused on the horizontal wave number of perturbation which is largely decreased by the introduction of spatial variations in these properties. We further developed an analytical model of convection cells which consider the thickness of stagnant lid and convective vigor beneath it. The model successfully reproduced the mechanism of increasing horizontal length scale of ST regime convection cells for each condition of spatial variations in physical properties.

4. Discussion and concluding remarks

The results of present studies indicate that, under certain conditions, the convection of fluids with strongly temperature-dependent viscosity takes place which is characterized simultaneously by (i) large horizontal length scales of convective cells and (ii) thick stiff lid of highly viscous fluid above it. This is in a stark contrast with earlier numerical studies using constant thermal conductivity and expansivity where the convection beneath stagnant lids is always associated with cells with small horizontal length scales. Our findings therefore highlight the essential roles of the spatial variation of the thermal conductivity and thermal expansivity on the convection patterns in the mantle of terrestrial planets.

Keywords: mantle convection, viscosity, thermal conductivity, thermal expansivity, stagnant-lid convection

High-pressure transitions in FeCr₂O₄ and crystal structures of new post-spinel phases

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Introduction

FeCr₂O₄ chromite has the spinel (Sp) structure, and forms the solid solution with MgAl₂O₄ and MgCr₂O₄ which occurs in the Earth's mantle. The chromitite composed of mostly FeCr₂O₄-rich spinel includes high-pressure minerals such as diamond and coesite, which suggest possible indicators of material cycles in the deep mantle (Arai, 2010; Yamamoto et al., 2009). Chen et al. (2003) reported occurrence of the high-pressure polymorphs of FeCr₂O₄-rich composition in Suizhou meteorite and synthesized them with diamond anvil cell. They demonstrated that the natural chromite spinel transforms to calcium ferrite (CF)-type at 12.5 GPa and to calcium titanate (CT)-type above 20 GPa. High-pressure transition study of pure FeCr₂O₄ end-member demonstrated that cubic spinel of FeCr₂O₄ transforms to tetragonal at high-pressure and room temperature (Kyono et al., 2012). In this study, we report experimental results on the phase relations in FeCr₂O₄ at high pressure and high temperature and the crystal structure analyses of new post-spinel phases.

Experimental methods

FeCr₂O₄ spinel was synthesized from a mixture of Fe₂O₃ and Cr₂O₃ with a 1:2 molar ratio by heating at 1200C for 24 h in a controlled oxygen fugacity using a mixture of H₂, CO₂ and Ar with volume ratios of 1:1:2. High-pressure experiments were made by quenching method at 12-28 GPa and 800-1600C with a Kawai-type 6-8 multianvil high-pressure apparatus at Gakushuin University. Phase identification of each recovered sample was made with powder and microfocus X-ray diffractometers, and compositional analysis was made with a SEM-EDS. The recovered FeCr₂O₄ sample was observed by a TEM at Geodynamics Research Center of Ehime University. Angle-dispersive synchrotron X-ray diffraction measurements of some recovered samples were made at ambient conditions using the beam line BL02-B2 at SPring-8 for Rietveld analysis with the RIETAN-FP software (Izumi and Momma, 2007).

Results and discussion

Sp-type FeCr₂O₄ first dissociates into a mixture of Fe₂Cr₂O₅ + Cr₂O₃ at about 15 GPa, and further transforms to a CF-like phase at lower temperature than 1300C and CT-type FeCr₂O₄ at higher temperature at around 20 GPa. Although the CF-like phase had the same space group as the CF-type structure from the result of TEM observation, the X-ray diffraction pattern was somewhat different from that of the CF-type structure. Rietveld refinement confirmed that the FeCr₂O₄ synthesized above about 20 GPa at higher temperature than 1300C has the CT-type structure and that Fe₂Cr₂O₅ phase is isostructural to the high-pressure Mg₂Al₂O₅ phase (Enomoto et al., 2009). The CT-type phase with pure FeCr₂O₄ composition and the Mg₂Al₂O₅-type Fe₂Cr₂O₅ were synthesized for the first time, and the structure analyses of the phases were carried out.

Keywords: post-spinel, high-pressure, Rietveld analysis, chromite, FeCr₂O₄, TEM observation

Effects of Thermodynamic Properties on the Geometrical Evolution of Subducting Slabs

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In this study, we perform a series of numerical experiments to investigate the effects of thermodynamic properties on the geometrical evolution of subducting slabs. We calculate density, thermal expansivity, and heat capacity of mantle mineral assemblages of a lherzolite composition over a range of pressure and temperature (PT) conditions applicable to the Earth's mantle, using the thermodynamic database of Stixrude and Lithgow-Bertelloni (2011) and the thermodynamic calculation code *Perple_X* (Connolly, 2009). Following Nakagawa et al. (2009), we assume that thermal diffusivity follows a theoretical power-law relationship with density and derive thermal conductivity from the calculated density, expansivity, and diffusivity. The calculations show that density, expansivity, and conductivity varies significantly with depth; for example, the ranges of their values for a typical mantle geotherm are 3300-5100 kg/m³, 1.5-3.5 10⁻⁵/K, and 3-18 W/m K, respectively. The change in heat capacity is relatively small (< 5%). We incorporate the effects of these thermodynamic properties into a 2-D finite element code with compressible convection formulations under the anelastic liquid approximation (Lee and King, 2009) and develop a thermodynamically consistent dynamic subduction model with kinematic boundary conditions. In the model, we use a composite mantle rheology that accounts for both diffusion and dislocation creep for the upper mantle with rheological parameterization for wet olivine (Hirth and Kohlstedt, 2003). For the lower mantle, following Billen and Hirth (2007) and Lee and King (2011), we adjust the rheological parameter values for wet olivine diffusion creep to test the effects of viscosity contrast between the upper and lower mantle on slab evolution. In models with PT-dependent density, lithostatic pressure in the lower mantle at a given depth is higher than a case with a constant density (by ~800 kg/m³ at the core-mantle boundary). The higher pressure leads to stronger mantle due to the pressure dependence of the mantle viscosity, leading to a different viscosity structure from the case with a constant density. This change in the viscosity structure due to PT-dependent density alone can have a significant effect on the simulation of slab evolution; for example, for a given set of rheological parameters, a model with PT-dependent density predicts buckling of the slab in the lower mantle while a model with constant density shows no buckling. To focus on the effects of thermodynamic properties, we remove this rheological effect of density variation by adjusting the rheological parameters for the lower mantle to maintain a similar viscosity structure for each set of experiments. When no viscosity contrast is imposed between the upper and the lower mantle, the model predicts that the slab sinks vertically into the lower mantle without experiencing much resistance regardless of the effects of thermodynamic properties. When viscosity contrast of ~10-100 is imposed, the model with constant thermodynamic properties predicts the buckling of the slab immediately below the transition zone. In contrast, the thermodynamically consistent model with the same viscosity structure predicts that the slab sinks sub-vertically into the lower mantle, and slab buckling tends to occur in the bottom half of the lower mantle. When large viscosity contrast (>100) is imposed, however, slab buckling occurs immediately below the transition zone even in a thermodynamically consistent model. These modeling results indicate that in numerical simulations, particularly those with viscosity contrast of <~100, noticeably different slab geometry can evolve, depending on the treatment of thermodynamic properties.

Keywords: Dynamic slab model, Thermodynamic properties, Mantle viscosity, Slab geometry, Slab buckling

Sound velocities of CaSiO₃ perovskite

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Calcium silicate perovskite (CPv) is the most significant Ca-bearing mineral of the mantle transition zone and the third most abundant phase of the lower mantle within the pyrolite assumption (O'Neill and Jeanloz, 1990; Ringwood, 1976). In addition, it is believed that subducted materials such as Mid-Ocean Ridge Basalts (MORB) may involve substantial amount of CPv. Thus the high-pressure and high-temperature structural and elastic behaviours of CPv have been extensively studied for their importance in understanding the internal structure of the Earth mantle. However, and besides reports by theoretical calculations, high-pressure experimental measurements of sound velocities of CPv are few (Li et al., 2004; Kudo et al., 2012).

Here, we examined the sound velocities of CaSiO₃ perovskite in situ at high-pressure and high-temperature up to 23 GPa and 1700 K using a combination of ultrasonic interferometry and synchrotron X-ray diffraction techniques within a DIA-type multi-anvil press apparatus at BL04B1 in SPring-8. The velocities of P- and S-wave appeared to behave quasi-linearly within the P and T range studied. Linear fitting for the tetragonal structure of CPv at 300 K yielded $V_P(0) = 10.171$ km/s and $V_S(0) = 5.285$ km/s. Generally our results agreed with Kudo et al. (2012), with V_S being slower than previous estimates by theoretical and experimental methods. In addition we present new high P and T data for the cubic structure of CPv, which displayed velocities about ~5% faster than the tetragonal structure at $T > 500$ K.

This new results suggested that V_P and V_S of CPv are high compared to surrounding mantle in the uppermost part of the mantle transition region (MTR). In the lowermost part of the MTR, CPv would have velocities comparable to PREM velocities, which make this phase barely detectable in this region. On the other hand, velocities of CPv would be significantly slower than lower mantle.

Keywords: CaSiO₃ perovskite, high-pressure, high-temperature, ultrasonic interferometry, X-ray diffraction

Electrical conductivity of stishovite: effects of Al₂O₃ and water

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Stishovite is one of the major constituent minerals in subducted oceanic crust. At pressures above 25 GPa, the post-garnet assembly transformed from garnetite originated from eclogite contains up to 25 vol.% stishovite. On the other hand, if Archean continental crust mainly consisting of tonalite-trondhjemite-granodiorite (TTG) was destructed and subducted into the deep mantle, its assembly contains more than 80 vol.% stishovite. In this case, stishovite should be considered as a main controlling phase for the electrical conductivity structure of the mantle. Electrical conductivity of stishovite is expected to be very low because migration of Si and O vacancies is quite slow even high temperature. However, electrical conductivity of stishovite may drastically increase because of their fast mobility as a charge carrier when stishovite crystals contain small amounts of Al and H.

In the present study, effects of Al and H components on electrical conductivity of stishovite was investigated to consider as a cause of the high conductivity anomaly observed in the mantle transition zone. The starting materials were prepared from SiO₂ powder with various amounts of Al(OH)₃. Stishovite aggregates were synthesized at 12 GPa and 1673 K. Chemical composition of the run products was SiO₂ with various amounts of Al₂O₃ (0, 1, 5 wt.%) and water. The electrical conductivity measurements were performed at 12 GPa and various temperature conditions to detect effect of water. Relatively dry conditions were accomplished by annealing at 1900 K, proton conduction can be detected under relatively lower temperature conditions less than 1100 K. The conductivity of stishovite increases with increasing H₂O and Al₂O₃ contents. This trend implies that the charge carrier in stishovite is probably interstitial alkali ion in association with substitutional Al in Si site. At low temperatures, the Al-related defect occupied by proton in Si site could be a main charge carrier. Because activation enthalpy for proton conduction is lower than that for Al³⁺-M⁺ pair and its dissociation products, proton conduction could be dominant conduction mechanism below 1100 K.

Electrical conductivity of Al-bearing stishovite (more than 1 wt.% Al₂O₃) is more than one order of magnitude higher than those of wadsleyite and ringwoodite. The high conductivity values observed in the mantle transition zone can be explained by a presence of aluminous stishovite derived from subduction of Archean continental crust.

Keywords: stishovite, electrical conductivity, mantle transition zone

First principles prediction of a new dense hydrous magnesium silicate

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The global circulation of water in the earth is important to investigate the evolution history and dynamics of the earth, since the physical properties (e.g. atomic diffusivity, melting temperature, electrical conductivity and seismic velocities) of the constituent minerals are considerably changed by the presence of water. It has been reported that water is carried into the deep Earth's interior by hydrous minerals such as the dense hydrous magnesium silicates (DHMSs) in the descending cold plate. However, high pressure behavior of DHMSs, especially the stability of phase D which is the highest pressure phase of DHMSs has not been clarified so far. In this study, I explored the possibility of further phase transition and dissociation of phase D into the hydrous or anhydrous minerals. As a result, the new phase which has lower enthalpy than phase D has been found above about 40 GPa. Therefore, there is a possibility that this new phase in subducting slab takes over water and carries into the deeper part of earth's lower mantle. The detail of the structure and the high pressure-temperature phase boundary determined by quasi-harmonic approximation will be shown and the possible geophysical implications will also be discussed at the presentation.

Keywords: DHMS, first principles calculation, hydrous mineral, high pressure

Melting temperature measurements of water using a laser-heated diamond anvil cell technique with CO₂ laser

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The phase relations and physical properties of water at high pressures and temperatures are important to physical, geophysical, and planetary problems. For example, recent molecular dynamics calculations suggest that water is superionic at high densities relevant to planets such as Uranus and Neptune, and this predicted property plays a key role in dynamo models to explain the unusual non-dipolar magnetic field structure of these planets. The advances in combining techniques such as synchrotron x-ray diffraction or in situ optical Raman spectroscopy with diamond anvil cell (DAC) technologies allow us to measure the melting temperature of water to within a few percent for pressures up to 35 GPa. However, at pressures above 35 GPa, recently reported values of melting temperature exhibit significant differences with each other. This discrepancy should be caused by a chemical reaction occurred between the metal absorber contained in the DAC and the dissociated water. By using CO₂ laser for sample heating, the metal is not necessary because water has significant absorption in the wavelength range of CO₂ laser (10.6 micron). We report the melting temperature of water in a diamond anvil cell more than 35 GPa using CO₂ laser heating system.

We performed the experiments using a DAC with diamonds having flats of 300 micron diameter. Third distilled H₂O was loaded into a ~100 micron diameter and ~50 micron thick sample chamber in a DAC. A rhenium gasket was used to contain the sample. The samples were first compressed to a required pressure at room temperature and then heated by two CO₂ lasers with a both-sided heating technique reducing the axial temperature gradient in the sample. The incident angle of radiation of the CO₂ lasers (Synrad 100 W) is about 20 degree. The laser beams were focused by ZnSe lens onto the sample in a DAC. The heated area, which corresponds to about ~30 micron, is imaged from both sides of sample on the slit of the entrance of the spectrometer and the two charge coupled device (CCD) detectors, respectively. The temperature was measured by the spectroradiometric method. The uncertainty in temperature within the 30 micron area was less than approximately 7 % stemming from radial temperature gradients.

Melting was determined by plotting the laser power / sample temperature function and looking for the thermal anomaly associated with melting. We found a temperature plateau arising from the melting of water at each pressure. This temperature was determined as the melting point. Another heating experiment of water including a small amount of Ir powder (<10 micron) was performed for cross-check of the melting temperature determined from the relation between the power and the temperature. The temperature when the powder moves was in good agreement with that of the plateau at each pressure. This fact supports that the temperature of the plateau corresponds to the melting point.

The melting temperatures determined in this experiment are much lower than those of the planetary isentropes of Neptune and Uranus. Therefore, at least, solid water might not exist in the interiors of Uranus and Neptune at 100 GPa.

Keywords: water, laser-heated DAC technique, high pressure

Seismic attenuation in the upper mantle beneath the Ontong Java plateau

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We investigate attenuation structure in the upper mantle beneath the Ontong Java plateau (OJP) to identify the origin of this largest LIPs (Large Igneous Provinces) in the globe. Previous studies have shown that S-wave velocities in the upper mantle beneath the OJP are slower than those in the surrounding regions (Richardson et al., 2000). On the other hand, only a few studies have been performed on the attenuation beneath the OJP. Gomer and Okal (2003) showed that ScS-waves traveling beneath the OJP exhibit weaker attenuation than those traveling in the surrounding regions, from which they suggested that the slow velocities in the upper mantle beneath the OJP are not of thermal origin. However, the attenuation of the ScS waves represents average attenuation in the entire mantle beneath the OJP, not in the upper mantle. We need to determine the attenuation in the upper mantle to compare with the slow velocity anomalies in the upper mantle beneath the OJP.

We analyzed teleseismic broadband waveforms of transverse component from deep earthquakes beneath the Solomon Islands subduction zone. Spectral ratios of the sS and S waves were computed to extract information of the attenuation in the upper mantle beneath the OJP. We made correction for crustal structure in and around the OJP, since the sS waveforms are sensitive to the crustal structure at the surface bounce points. Qs values were estimated to be 30-50 for the sS waves of which bounce points are located in the OJP and 40-100 for the sS waves traveling out of the OJP. The Qs values beneath the OJP are well lower than those estimated in other back-arc regions (70-90; Flanagan and Wiens, 1994). The Qs values outside the OJP are close to those in back-arc region. Differential travel time residuals of sS-S are also measured. The differential residuals are well correlated with attenuation pattern: sS waves bouncing at the OJP travel slower by 3 sec than those outside the OJP, which is consistent with the previous studies (e.g., Richardson et al., 2000). These results suggest that the slow velocity anomalies in the upper mantle beneath the OJP are caused mainly by thermal effects.

Keywords: Ontong Java plateau, Seismic attenuation

The thermal history of the Earth's mantle that evolves in two stages

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The thermal history of the mantle calculated from numerical models of a coupled magmatism-mantle convection system is compared with that of the Earth. Both magmatism and mantle convection with tectonic plates are self-consistently reproduced in a two-dimensional rectangular box. The mantle evolves in two stages in these models. On the earlier stage that continues for 1-2 Gyr, heat producing elements (HPEs) and heat flux from the core (HFC) strongly heat the deep mantle, and frequently let hot materials there ascend to the surface as bursts. The mantle-bursts cause vigorous magmatism, stir the mantle efficiently, and make plates move chaotically. As HPEs and HFC decay, however, mantle-bursts stop. On the later stage, subducted basaltic crusts accumulate on the core mantle boundary to form compositionally dense basaltic piles, and plate motion becomes more stable. The average temperature in the entire mantle T_w steadily decreases with time owing to heat extraction by magmatism and mantle convection. The cooling rate is 80-130 K/Gyr on average depending on the internal heating rate. The thermal history of the upper mantle is, however, quite different from this: The average temperature in the upper mantle T_u drops to about 1800 K within the first 100 Myr, and remains almost constant at 1700-1800 K for the subsequent 3 Gyr or even longer regardless of the internal and basal heating rate as well as the initial temperature; T_u gradually decreases to around 1600 K only after that. The thermostat effect of magmatism keeps T_u below 1800 K on the earlier stage no matter how strongly the mantle is heated or how high the initial temperature is. T_u does not decrease on the later stage till 3 Gyr because subducted slabs stagnate on the CMB and do not return back to cool the upper mantle till that time; the steady decrease in T_w during this period is due to the cooling of the lower mantle. The delayed cooling of the upper mantle makes the heat flux at the surface remain almost constant throughout the 4.5 Gyr history of the mantle. At 4.5 Gyr, the Urey ratio is as low as 0.4-0.5 depending on the internal heating rate, and the lower mantle is significantly colder than expected from adiabatic extrapolation from the upper mantle.

Keywords: mantle evolution, plate tectonics, magmatism, numerical simulation, thermal history

Origin of life: Mechanism of leaking Earth; Fate of cooling Earth

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Supply of nutrients is essential to bear life. To enable a nutrient supply, a landmass must appear above sea level, because continental crust (landmass) is nutrient-enriched. Through weathering, erosion, and transportation of nutrient-enriched rocks, particles are eventually fined to a sufficient size for life to ingest. The emergence of a landmass is caused by a drop in sea level. Here, the cause of sea level change through time is discussed.

Mechanism of change of ocean thickness through time

In the early 2 Ga, after the birth of the ocean, the surface ocean thickness increased through the degassing of a water-rich magma generated from a primordial mantle. This is a theoretical speculation based on the geologic constraints. From the Archean to the Proterozoic Earth, mantle potential temperature was 200-150K higher than that of today. The subduction-zone geotherm, as documented by P-T conditions of regional metamorphic belts, prohibits water transportation into the mantle through the subduction of hydrated slabs.

Since the Archean, OIBs, such as komatiites and picrites, are enriched in H₂O and CO₂, as well as MORBs with minor water and CO₂, all of which tend to increase the ocean volume through time. On the other hand, the volume of the ocean never reaches sufficient size to bury all of the continents in the Precambrian. Presumably, the maximum thickness of the ocean would have been less than 1km.

Subduction of ocean water and hydrated oceanic slabs began at the onset of the Phanerozoic, as discussed below. About 700-600 Ma ago, the ocean thickness started to decrease, with the reduction of about 600 m until now through the fluctuations in the balance between output versus input of water into the mantle.

Phase diagrams of MORB + H₂O and peridotite + H₂O indicate that the ocean level would decrease though subduction of hydrated oceanic slabs if the top of the descending slab changes to temperatures lower than 600°C at Moho depth of 30km through time from hot Archean mantle to the present. The subduction zone geotherm along the surface of the descending slab turns to generate blueschist-facies rocks if it crosses the high-temperature corner of the blueschist facies in a P-T space defining the subduction zone geotherm and passing to the point at 10kb, 600°C. A plot of the P-T conditions of the regional metamorphic belts over the world since the Archean shows that the first appearance of blueschist was ca. 700Ma, and the subduction zone rapidly cooled at the onset of Phanerozoic. The temperature of Moho depth was higher than 600°C before 700Ma, but rapidly cooled below 600°C, thereafter, and down to 200°C at present. This suggests that the initiation of return-flow of seawater into the mantle began in the Latest Proterozoic, as estimated by the phase diagrams. The observed drop in sea-level clearly supports the idea, and the proposed sea-level-change curve shows that ca. 600m thick ocean has been removed from the surface into the mantle, at the 410-660km depth transition zone, which has a capability to store about 5 times of the total mass water of surface oceans.

The sea-level fluctuation of plus minus 300m in the Phanerozoic could be explained by the glacial/non-glacial periods, as well as the partial mantle overturn when high-temperature and fertile lower mantle materials catastrophically replaced the upper mantle, such as during the Cretaceous (120-85Ma), a major period of magmatic-driven activity. Another pulse was during the mid-Paleozoic, when huge batholith belts were formed similar to the Cretaceous pulse. If the rate of decreasing ocean volume continues over the next 1.0-1.5 b.y., the Earth will finally dry up, which will mark the end of life.