Lower Mantle Melting: Experiments and Thermodynamic Modeling in the System MgO-SiO$_2$

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Abstract

Characterizing and modeling melting relations in the system MgO-SiO$_2$ at lower mantle pressures rely on the location of the eutectic points for MgO-MgSiO$_3$ and MgSiO$_3$-SiO$_2$. While at an uppermost lower mantle pressure there is general consensus on the eutectic composition in the former, large discrepancies exist for MgSiO$_3$-SiO$_2$ from experiments in the diamond anvil cell, ab-initio simulations and models built on them. In order to address this discrepancy, we have performed multi-anvil press experiments at 24 GPa for Mg$_8$Si$_6$O$_{16}$ and Mg$_6$Si$_{16}$O$_{32}$ at temperatures of 2650 ± 100 K and 2750 ± 100 K. In the experiments at 2750 ± 100 K, we observe the presence of partial melt. The recovered Mg$_8$Si$_6$O$_{16}$ sample shows SiO$_2$ stishovite as the liquidus phase, and electron microprobe analysis of the quenched melt determines $X_{\text{SiO}_2} = 0.53 ± 0.03$ as the eutectic composition. We fit a thermodynamic model to describe the melting relations in the MgO-SiO$_2$ system, and extrapolate to core-mantle boundary pressure. At 136 GPa, we predict that the eutectic points have moved further away from enstatite composition, and solidus temperatures remain similar for MgO-MgSiO$_3$ and MgSiO$_3$-SiO$_2$.

Plain Language Summary

Melting of rock in the Earth's deep interior is an important process that has shaped its geological history, and continues to do so to date. However, melting processes at high pressure have not been fully characterized to date, and here we perform experiments that help to better describe models for it. We do so for compositions in the MgSiO$_3$-SiO$_2$ system that complement prior work for MgO-MgSiO$_3$; combined, these experiments put important constraints on melting for the most important chemical components in the Earth's mantle, MgO-SiO$_2$. We find that when silicate melts crystallize, they do form the dominant lowermost mantle mineral, MgSiO$_3$ bridgmanite, only for a limited composition range in terms of the SiO$_2$/MgO ratio; otherwise, MgO or SiO$_2$ form first, and generally at much higher temperatures. Melts in both the MgO-MgSiO$_3$ and MgSiO$_3$-SiO$_2$ systems become fully crystallized at very similar temperatures which suggests that a prior idea that the occurrence of partial melts in the Earth's deepest mantle is associated with different chemical composition does not hold if only the MgO-SiO$_2$ system is considered.

1. Introduction

Silicate melting has shaped the chemical and physical evolution of Earth's interior, from magma ocean formation during accretion (Elkins-Tanton et al., 2003; Elardo et al., 2011; Monteux et al., 2016; Wood & Halliday, 2005) to the continuous extraction of melt through plate tectonic processes. The latter is dominated by the ongoing formation of mid-ocean ridge basalt (MORB) from mantle peridotite (Kelemen et al., 1995). Melts may also form at the boundaries of the mantle transition zone (e.g., Revenaugh & Lipkin, 1994; Schmandt et al., 2014) and near the core-mantle boundary (CMB), where they serve as an explanation (e.g., Andraut et al., 2014; Labrosse et al., 2007) for the occurrence of seismic ultra-low velocity zones (ULVZ; Herrnleun & Jellinek, 2010; Rost et al., 2005). Most deep melting processes are mitigated by the influence of volatile components (Dasgupta, 2018; Johnson et al., 2008), but despite volatile loss of slab material transported into Earth's deep interior (e.g., Chemia et al., 2015), dry melting may occur within the thermal boundary layer at the base of the mantle (Hirose et al., 1999).

Melting experiments on natural rocks (e.g., Andraut et al., 2014; Fiquet et al., 2010; Pradhan et al., 2015) at lower mantle (LM) pressure ($P \geq 24$ GPa) yield valuable insight on solidus and liquidus temperatures ($T$), but the complexity of composition, the small size of samples, especially in diamond anvil cell (DAC) experiments, and large thermal gradients make it difficult to identify the liquidus and solidus in a systematic way. Therefore, the consideration of simplified systems is required in high-$P$ experiments in order to develop a thermodynamic description that can successively be built up by adding components, ideally combining well-controlled multi-anvil
Given a molar ratio MgO/SiO₂ ~ 1.15 or X_{SiO_2} = 0.47, with X_{SiO_2} = (SiO_2/SiO_2 + MgO), for the bulk Earth (Dauphas et al., 2015) and X_{SiO_2} = 0.45 for the bulk silicate Earth (Palme & ONeill, 2003), many experiments on melt relations in MgO-SiO₂ have focused on the MgO-MgSiO₃ subsystem (Liebske & Frost, 2012; Ohnishi et al., 2017). However, SiO₂-enriched reservoirs in the Earth, primarily associated with oceanic crust, have been considered as potential sources for partial melting near the CMB (Andrault et al., 2014; Nomura et al., 2014), and related to the observation of the ULVZ there (Thorne et al., 2019). The location of the eutectic in the Mg-SiO₂-SiO₃ system may further be directly relevant for the aubrite parent body in our solar system with X_{SiO_2} > 0.5 (Dauphas et al., 2015), and for rocky exoplanets as some of the stars in the Hypathia Catalog, taken as a proxy for potential planetary compositions, show X_{SiO_2} ~ 0.6 (Hinkel & Unterborn, 2018).

Experiments on melt relations in the MgO-SiO₂ system at lower mantle P were performed in the MA press by Liebske and Frost (2012), the DAC by Baron et al. (2017), Ohnishi et al. (2017), and Ozawa et al. (2018), modeled based on ab-initio simulations by de Koker et al. (2013), and thermodynamically described by Boukaré et al. (2015), Miyazaki and Korenaga (2019) and Belmonte et al. (2017). Bridgmanite is the phase at the liquidus in the vicinity of the enstatite composition, with two eutectics points bracketing its stability; this stability field has been found to expand with P.

At P = 24 – 25 GPa, the eutectic composition on the MgO-MgSiO₃ join is consistently determined as X_{SiO_2} = 0.43 ± 0.01 (Belmonte et al., 2017; Boukaré et al., 2015; de Koker et al., 2013; Liebske & Frost, 2012; Ohnishi et al., 2017; Ozawa et al., 2018) with some uncertainty in T, ranging from ~2700 K (Liebske & Frost, 2012) to ~2900 K (Boukaré et al., 2015). For MgSiO₃-SiO₂, de Koker et al. (2013) find X_{SiO_2} = 0.62 ± 0.01 at a eutectic temperature Tₑ, lower by 100–150 K than for MgO-MgSiO₃ at 24 GPa, while Belmonte et al. (2017) compute X_{SiO_2} = 0.51 at Tₑ ~ 2800 K, ~50 K higher than for the peridotite eutectic. Ozawa et al. (2018) determine X_{SiO_2} = 0.60 at P = 40 GPa in DAC experiments and extrapolate to X_{SiO_2} = 0.58 and Tₑ ~ 2500 K at P = 25 GPa, by contrast. However, to date melting relations for MgSiO₃-SiO₂ have not been studied in the MA press, although it provides better control of thermal gradients and allows more rigorous textural analysis of the larger sample compared to DAC experiments. Here we perform MA melting experiments at P = 24 GPa with Mg₆Si₇O₂₆ and Mg₃SiO₇ starting compositions, complementing the work by Liebske and Frost (2012).

### 2. Experimental and Analytical Methods

The synthesis of compositions along the MgSiO₃-SiO₂ join (Mg₆Si₇O₂₆ and Mg₃SiO₇) requires MgSiO₃ glass, for which analytical-grade MgO and SiO₂ powders (1:1 molar ratio) are mixed in ethanol, dried under infrared light, and ground to a fine powder. The mixture is melted in a Pt crucible for 30 minutes at 1650 °C and quenched by dropping the crucible into water. The MgSiO₃ glass is repeatedly ground to a fine powder in ethanol and then dried. Mg₆Si₇O₂₆ and Mg₃SiO₇ powders are prepared by mixing the ground enstatite glass with SiO₂ powder; they are stored under vacuum to prevent hydration.

We perform melting experiments using the 1,200 t Sumitomo MA press at Bayerisches Geoinstitut (Keppler & Frost, 2005). The sample powder is put in a Rhenium (Re) capsule (~2 mm in length and ~1 mm in diameter), closed with a Re foil disk on top and bottom, and inserted in a Cr₂O₃-doped MgO octahedron (7 mm edge length). A LaCrO₃ heater inside the octahedron is insulated from the Re capsule by a MgO sleeve (Figure S1 in Supporting Information S1). Cell assembly parts are stored in an oven at 1000°C for 12 hr to dehydrate prior to use. The octahedron is compressed by tungsten carbide anvils with 3 mm corner truncations (7/3 assembly). The assembly is compressed to P = 24 GPa, heated to the target T and kept there for 1 – 2 minutes. The run product is quenched by shutting off the power and slowly decompressed to ambient conditions over 16 h. We perform experiments without thermocouples and extrapolate the heating power-T relation, determined in an independent experiment at P = 24 GPa with the same setup to a power of 433 W (T = 2375 K) using W-Reuters wires as thermocouples, to 450 – 470 W applied here. To account for the P dependence of thermocouple electromotive force, we follow Nishihara et al. (2020) and add a correction to estimate the T of the experiments (Table 1), with an uncertainty of ±100 K. Thermocouples tend to fail at the conditions explored (e.g., Liebske & Frost, 2012), and using power instead makes T conditions in the experiments far more reproducible.
Recovered sample chambers are mounted within epoxy parallel to the cylinder axis and then polished to the central section to observe phase relations along the thermal gradient, which is parallel to the furnace axis. The texture is analyzed with a LEO1530 scanning electron microscope (SEM) at acceleration voltage 15 – 20 kV and beam current 10 nA. Chemical compositions of the run products are determined using a JEOL JXA-8200 electron microprobe (EPMA) at acceleration voltage 15 – 20 kV with focused beam for mineral phases and a beam diameter of 30 μm. Counting times are 20 s on the peak and 10 s on the background for each element. Enstatite and orthoclase standards are used for Mg and Si calibration.

### 3. Sample Analysis

In the experiments for Mg$_6$Si$_4$O$_{16}$ and Mg$_6$Si$_4$O$_{17}$ heated with 450 W ($T = 2650 ± 100$ K) (Table 1), we find a crystalline matrix with well-developed triple junctions (similar to the right side of the sample in Figure 1a), showing that they were performed at sub-solidus conditions, consistent with the prediction by de Koker et al. (2013). In the higher-$T$ experiments ($T = 2750 ± 100$ K) for Mg$_6$Si$_4$O$_{17}$, we observe a quench texture in the recovered sample coexisting with the solid (Figure 1a). Within the quench texture we observe small bridgmanite and stishovite grains (both a few μm in size, also confirmed by X-ray diffraction), with a small amount of liquid between them, suggesting that we have just reached super-solidus $T$, tightly constraining $T_e$, with our value in excellent agreement with the prediction by de Koker et al. (2013). In addition, some of the stishovite grains appear to have grown at the expense of the liquid during quench, as indicated by the lighter-colored rims. The amount of melt preserved in the sample is too small for it to be analyzed by EPMA.

For the Mg$_6$Si$_4$O$_{16}$ sample heated with 470 W ($T = 2750 ± 100$ K), by contrast, we observe a liquid quench texture, coexisting with acicular stishovite crystals, some of which reach lengths of 50 μm and widths of 30 μm, indicating a liquidus assemblage (Figure 1b). The identification of stishovite as the liquidus phase for $X_{SiO_2} = 0.6$ starting composition suggests that the liquid evolves toward a MgSiO$_3$-rich composition. Microprobe analysis of 32 spots in the quenched melt shows values in the range of $X_{SiO_2} = 0.49 – 0.60$, reflecting small-scale heterogeneity (Figure 1b), with an average of $X_{SiO_2} = 0.53 ± 0.03$ (Figure 2) that accurately represents the bulk melt. As the melting experiment for Mg$_6$Si$_4$O$_{16}$ with virtually the same heating power suggests that we are very close to $T_e$, the liquid composition in the Mg$_6$Si$_4$O$_{16}$ experiment constrains the eutectic composition in the MgSiO$_3$-SiO$_2$ system.

This is a significantly lower eutectic $X_{SiO_2}$-value at $P = 24 – 25$ GPa than calculated by de Koker et al. (2013) based on ab-initio simulations with...
$X_{\text{SiO}_2} = 0.63$ (Figure 2) or extrapolated from DAC experiments by Ozawa et al. (2018) with $X_{\text{SiO}_2} = 0.58$, but consistent with $X_{\text{SiO}_2} = 0.51$ modeled by Belmonte et al. (2017).

4. Thermodynamic Description

4.1. Model Formulation and Parameters

We extend the previously established thermodynamic model for melting phase relations in MgO-MgSiO$_3$ from Liebske and Frost (2012) to the MgO-SiO$_2$ system by using the location of both eutectic points as constraints. For a closed formulation with a minimum number of parameters, we represent all liquids through the oxide components MgO and SiO$_2$, and use MgO periclase, MgSiO$_3$, bridgmanite and SiO$_2$ stishovite as solid phases. At both eutectics we equate the chemical potentials ($\mu$),

$$\mu^\text{solid} = \mu^\text{liquid} + \mu^\text{SiO}_2$$

respectively. The chemical potentials of the liquids can be expanded into a standard state potential $\mu^\text{liquid}$ for MgO and $\mu^\text{liquid}$ for SiO$_2$, and an activity term $a_{\text{MgO}} = X_{\text{MgO}}\gamma_{\text{MgO}}$ ($a_{\text{SiO}_2} = X_{\text{SiO}_2}\gamma_{\text{SiO}_2}$), with $\gamma_{\text{MgO}}$ ($\gamma_{\text{SiO}_2}$) the activity coefficient:

$$\mu^\text{solid} = \mu^\text{liquid} + R \cdot T \cdot \ln(X_{\text{MgO}}\gamma_{\text{MgO}}),$$

$$\mu^\text{solid} = \mu^\text{liquid} + R \cdot T \cdot \ln(X_{\text{SiO}_2}\gamma_{\text{SiO}_2}).$$

In a binary symmetric solution model, the activity coefficients are described as

$$R \cdot T \cdot \ln(\gamma_{\text{MgO}}) = W_{\text{MgO-SiO}_2}(1 - X_{\text{MgO}})^2$$

and

$$R \cdot T \cdot \ln(\gamma_{\text{SiO}_2}) = W_{\text{MgO-SiO}_2}(1 - X_{\text{SiO}_2})^2,$$

where $W_{\text{MgO-SiO}_2}$ is a Margules interaction parameter. We formulate a $T$-dependence of $W_{\text{MgO-SiO}_2} = W_0 - W_1 T$, but with experiments performed only at 24 GPa, a $P$ dependent model cannot be fit. Using a more complex solution model, for example, asymmetric as in Baron et al. (2017) and de Koker et al. (2013), does not change the width of the stability field for MgSiO$_3$ plus liquid as pointed out by de Koker et al. (2013), and with the number of data collected here, more complex models are not sufficiently constrained.

The standard state chemical potentials $\mu^\text{liquid}_\text{MgO}$ and $\mu^\text{liquid}_\text{SiO}_2$ are determined from melting of MgO and SiO$_2$, respectively. For MgO, for example,

$$\mu^\text{solid}_\text{MgO} - \mu^\text{solid}_\text{MgO} = \overline{\mu}_{\text{f,MgO}} = 0$$

holds, with $\overline{\mu}_{\text{f,MgO}}$ the Gibbs energy of fusion. Standard state chemical potentials for the liquid components ($\mu^\text{liquid}_\text{MgO}$ and $\mu^\text{liquid}_\text{SiO}_2$) are evaluated based on the thermodynamic model by de Koker et al. (2013) (cf. Supporting Information S1 for an extended presentation); $\mu^\text{solid}_\text{MgO}$, $\mu^\text{solid}_\text{MgSiO}_3$ and $\mu^\text{solid}_\text{SiO}_2$ are based on the Birch-Murnaghan Mie-Grüneisen model (cf. Supporting Information S1 for an extended presentation; e.g., Chust et al., 2017) for

Figure 2. MgO-SiO$_2$ melting relations computed from our thermodynamic model at 24 GPa (bottom black graph), based on the experiments performed in the MgSiO$_3$-SiO$_2$ system, combined with the result for the MgO-MgSiO$_3$ system from Liebske and Frost (2012). The eutectic composition (red circle with error) is determined based on an experiment for MgSiO$_3$ at $P = 24$ GPa and $T = 2750 \pm 100$ K (open circle); starting compositions and temperature of the other three experiments (Table 1) are also shown, with the filled black circles for the experiments at $T = 2650 \pm 100$ K showing experiments that remained below the solidus. Melting relations at $P = 136$ GPa (top black graph) computed from our model. For comparison we include melting relations from the model of de Koker et al. (2013) (dK13, blue) both at $P = 24$ GPa and $P = 136$ GPa. Abbreviations: pc for periclase, brg for bridgmanite, st for stishovite and liq for liquid.
MgO periclase, SiO$_2$ stishovite and MgSiO$_3$ bridgmanite used in de Koker et al. (2013). Both models and corresponding databases are implemented in the BurnMan thermoelastic toolbox (Cottaar et al., 2014).

Using the SiO$_2$ liquid and stishovite parameters in the thermodynamic model of de Koker et al. (2013), we find that an effective interaction parameter of $\pm 200$ kJ/mol $\leq W_{\text{MgO-SiO}_2} \leq \pm 250$ kJ/mol is required to reproduce the solidus temperatures of $T \sim 2750$ K for MgSiO$_3$-SiO$_2$ determined here. Nevertheless, we cannot find a satisfactory solution for $W_{\text{MgO-SiO}_2}$ that describes $X_{\text{SiO}_2} = 0.53 \pm 0.03$ for the eutectic adequately. This is not surprising considering that in the model of de Koker et al. (2013) liquidus $T$ decreases rapidly from the high melting temperature ($T_m$) of stishovite with increasing MgO content. We therefore consider modifications in the coefficients for the thermodynamic description of liquid SiO$_2$ in the model of de Koker et al. (2013) (Tables S1 and S3 in Supporting Information S1). In order to reasonably match $T_m$ of SiO$_2$, we also consider changes in the thermodynamic description for stishovite which we restrict to reference energy and entropy (Table S4 in Supporting Information S1).

Rather than optimizing the thermodynamic parameters for SiO$_2$ liquid and stishovite separately from the Margules parameter $W_{\text{MgO-SiO}_2}$, we jointly determine them by using a least square Monte Carlo minimization. With only two constraints (the eutectics), the problem is underdetermined, and initial guesses and bounds for the parameters are important. We make choices that are guided by physical insight into (a) the thermodynamic parameters of SiO$_2$ liquid, (b) the melting curve of stishovite, (c) the range of values for the effective interaction parameter $W_{\text{MgO-SiO}_2}$ at 24 GPa discussed above, and (d) mixing properties for the liquids along the MgO-SiO$_2$ join estimated by de Koker et al. (2013). The melting point of MgSiO$_3$ at 24 GPa is not used as a constraint; rather, it serves as a consistency test for the results of the mixing model and the thermodynamic parameters for SiO$_2$ liquid. Using 1,000 Monte Carlo passes, we find an optimized thermodynamic description for SiO$_2$ (Tables S1 and S3 in Supporting Information S1) and model a $T$-dependent Margules parameter of

$$W_{\text{MgO-SiO}_2} = -174 \pm 4 \frac{\text{kJ}}{\text{mol}} - 17 \pm 2 \frac{\text{J}}{\text{mol} \cdot \text{K}} T.$$ (9)

The value for $W_2$ is consistent with the entropy of mixing reported by de Koker et al. (2013) from ab-initio simulations. At 25 GPa, minimum values for excess Gibbs energy of mixing in the various thermodynamic models developed for high $P$ applications fall in the range of $-50 - 60$ kJ/mol, with the exception of that of Miyazaki and Korenaga (2019) (Figure S2 in Supporting Information S1). At this (and higher) $P$, the mixing model by de Koker et al. (2013) shows very little asymmetry.

Any $P$ dependence discussed below hinges on the thermodynamic description of the liquid and solid components, and an implicit $P$ dependence of the interaction parameter via $T_m$. In this context, not only the evolution of the eutectic points with $P$ are important, but also the melting curve $T_m - P$ for MgSiO$_3$ as a consistency check as it involves both thermodynamic parameters for SiO$_2$ and the parametrization of $W_{\text{MgO-SiO}_2}$ (Equation 9), similar to its role at 24 GPa.

### 4.2. MgO-SiO$_2$ Melting Relations at Experimental Pressure (24 GPa)

Combining $W_{\text{MgO-SiO}_2}$ and the modified thermodynamic parameters as described above, we compute the MgO-SiO$_2$ phase diagram at $P = 24$ GPa (Figure 2). It hinges on five points: $T_m$ of MgO, MgSiO$_3$, and SiO$_2$ and the two eutectics.

$T_m = 4760$ K for MgO computed by BurnMan at 24 GPa is very close to that reported in de Koker et al. (2013) (4800 K) (Figures 2 and 3a), and agrees with DAC experiments by Du and Lee (2014) and Kimura et al. (2017) in the same $P$ range. Ab-initio results by Alfè (2005) and Soubiran and Militzer (2020) report $T_m$ higher and smaller by $\sim 250$ K, respectively.

For stishovite, we compute $T_m = 4210 \pm 120$ K at 24 GPa, with the uncertainty (two standard deviations) determined by 1,000 Monte Carlo passes, varying the thermodynamic parameters (Tables S1–S4 in Supporting Information S1), slightly higher than $T_m = 3920$ K from de Koker et al. (2013), but consistent with DAC experiments that bracket melting with $T = 3800 - 4300$ K at $P = 22 - 29$ GPa (Shen & Lazor, 1995) and $T = 4000 - 4400$ K at $P = 27 - 29$ GPa (Andrault et al., 2020; Figures 2 and 3b).
In our modified model, bridgmanite melts at 2750 ± 120 K at 24 GPa (Figures 2 and 4a), slightly below \( T_m = 2800 - 2900 \) K bracketed in MA experiments by Ito and Katsura (1992), DAC experiments by Zerr and Boehler (1993), the thermodynamic assessment of Liebske and Frost (2012) (2820 K), and molecular dynamics results by Di Paola and Brodholt (2016) in the same \( P \) range. \( T_m \) from the thermodynamic assessments of de Koker et al. (2013) falls slightly above our error bar estimated in the Monte Carlo simulations, the experiments by Shen and Lazor (1995) report a significantly higher value. The relatively low \( T_m \) of bridgmanite at 24 GPa in our model is directly linked to its narrow stability range on the liquidus in terms of \( x_{SiO_2} \), especially for the SiO\(_2\)-rich side of the phase diagram. Enhancing bridgmanite stability, for example, by a constant shift of the reference energy in the underlying thermodynamic model of the solid, would inevitably move the eutectic point(s) further away from MgSiO\(_3\), making it impossible to reproduce the experimental results.

The eutectic points at 24 GPa themselves are reproduced well in the Monte Carlo simulations with \( T_e = 2690 ± 100 \) K at \( x_{SiO_2} = 0.43 ± 0.01 \) and \( T_e = 2740 ± 120 \) K at \( x_{SiO_2} = 0.53 ± 0.01 \). The eutectic \( T \) of the latter is smaller than \( T_m \) for MgSiO\(_3\) by 10 K only, reflecting the flat liquidus curve in the vicinity of MgSiO\(_3\) (Figure 2). Similar to \( x_{SiO_2} \), a slightly smaller value of \( T_e \) for the MgO-MgSiO\(_3\) eutetic is consistent with the thermodynamic assessment of Belmonte et al. (2017) at uppermost LM pressure.

### 4.3. MgO-SiO\(_2\) Melting Relations at Lower Mantle Pressure

As mentioned above, the melting curves for MgO and SiO\(_2\) provide important anchor points for the phase relations (Equation 8) at higher \( P \), and that of MgSiO\(_3\) serves as a test for the validity and consistency of the liquid model.

For MgO, our melting curve matches DAC data (Du & Lee, 2014; Fu et al., 2018; Kimura et al., 2017) for \( P \) throughout the lower mantle within their margins of error (Figure 3a), reaching \( T_m = 7510 \) K at \( P = 136 \) GPa, CMB pressure, similar to de Koker et al. (2013) and ab-initio simulations by Soubran and Militzer (2020) using thermodynamic integration. Previous assessments by de Koker and Stixrude (2009) and direct two-phase ab-initio simulations by Alfè (2005) show higher \( T_m \) but still within experimental uncertainty.

With the changes in the thermodynamic parameters for SiO\(_2\) liquid and stishovite, \( T_m \) of SiO\(_2\) at \( P = 24 \) GPa is larger by ~ 300 K compared to de Koker et al. (2013), a difference that increases to ~ 550 K at CMB pressure (Figures 2 and 3b). However, our computed melting curve closely follows the experimental data on \( T_m \) by Andraut et al. (2020) up to 80 GPa; for \( P > 30 \) GPa our melting curve is within the margin of error of that computed by Usui and Tsuchiya (2010), and they approach the same value at CMB pressures. As has been pointed out by de Koker et al. (2013), the potential presence of CaCl\(_2\)-structured SiO\(_2\) (Tsuchida & Yagi, 1989) is expected to have a small effect on \( T_m \). However, the reconstructive phase transition of SiO\(_2\) to seifertite in the lowermost mantle (Das et al., 2020; Dubrovinsky et al., 2001) will expand the stability range of solid SiO\(_2\) to higher \( T \), an effect that can inherently not be captured by our model, and may account for the divergence of our model with the data by Andraut et al. (2020) for \( P > 100 \) GPa.

For MgSiO\(_3\), the melting curve predicted by our model shows a steep initial increase with \( P \) (Figure 4), closely following the trend of the DAC experiments (Shen & Lazor, 1995; Zerr & Boehler, 1993) to 40 GPa, and, with larger scatter in the data, to 62 GPa, the highest \( P \) achieved in Zerr and Boehler (1993), and that of the shock wave experiments by Fei et al. (2021) for much of the lower mantle \( P \)-range (50–80 GPa). However, shock wave experiments (Fei et al., 2021; Fratanduono et al., 2018; Mosenfelder et al., 2009) vary largely in their estimates of the...
melting curve. At higher $P$, our predicted $T_m - P$ curve flattens and reaches $5560 \pm 220$ K at 136 GPa. The melting temperature predicted by Stixrude and Karki (2005) at CMB $P$ is within the margin of error of our model, but their melting curve shows a more uniform increase for LM pressure.

The consistency of predicted melting curve evolution with prior data for all three congruently melting compositions along the MgO-SiO$_2$ join allows us to calculate the phase diagram at $P = 136$ GPa with some confidence (Figure 2) and to explore the evolution of both eutectics in terms of $X_{SiO_2}$ (Figure 5) and $T_e$ (Figure 6).

In agreement with previous work, we find that both eutectics move away from MgSiO$_3$ with increasing $P$. For the MgO-MgSiO$_3$ eutectic, $X_{SiO_2}$ shifts to lower values with $P$ significantly faster than in the models of de Koker et al. (2013) and Liebske and Frost (2012) or the experimental brackets determined in the DAC by Ohnishi et al. (2017) between 45 and 70 GPa. The $P$ dependence of the eutectic composition for MgO-MgSiO$_3$ agrees well with a second set of DAC experiments by Ozawa et al. (2018) over the $P$ range of the entire LM. The MgO-MgSiO$_3$ eutectic in our model has a minimum value of $X_{SiO_2} = 0.32 \pm 0.02$ at $P \sim 60$ GPa, beyond which it increases quasi-linearly with $P$ to reach $X_{SiO_2} = 0.35 \pm 0.02$ at CMB pressure. This behavior stems from the different $P$ evolution of the melting curves for MgO (Figure 3a) and MgSiO$_3$ (Figure 4b): At $P$ of the uppermost LM, the melting curve of Mg-SiO$_3$ is steeper than the one of MgO, leading to a rapid decrease in eutectic $X_{SiO_2}$. At $P > 60$ GPa, the $I_m - P$ slope for MgSiO$_3$ is flatter than the one for MgO, shifting eutectic $X_{SiO_2}$ to larger values.

As discussed above, the MgSiO$_3$-SiO$_2$ eutectic composition is close to Mg-SiO$_2$ with $X_{SiO_2} = 0.53 \pm 0.01$ at $P = 24$ GPa in our model, consistent with Belmonte et al. (2017), but significantly lower than $X_{SiO_2}$ calculated by de Koker et al. (2013). As $I_m$ for SiO$_2$ does not increase with $P$ at the same rate as that for MgSiO$_3$, eutectic $X_{SiO_2}$ moves to larger values with $P$ uniformly, reaching a value of $X_{SiO_2} = 0.66 \pm 0.01$ at 136 GPa. Experiments by Ozawa et al. (2018) at $P = 128$ GPa and $P = 139$ GPa agree well with our prediction, as for the MgO-MgSiO$_3$ eutectic, with values that are smaller than the model of de Koker et al. (2013). While Baron et al. (2017) assumed the chemical evolution of the eutectic following de Koker et al. (2013) in their data analysis, the actual composition reported for one sample at $P = 48$ GPa reveals a significantly lower value with $X_{SiO_2} = 0.59$, within the margin of error of our model prediction, and in general agreement with data by Ozawa et al. (2018) at $P = 41$ GPa. With a higher $I_m$ for lowermost mantle $P$ due to the presence of serpentine, $X_{SiO_2}$ would move to slightly smaller values.

Eutectic temperatures in the MgO-MgSiO$_3$ and MgSiO$_3$-SiO$_2$ systems are very close to one another, rising from $T_e = 2690 \pm 100$ K and $T_e = 2740 \pm 120$ K at $P = 24$ GPa to $T_e = 5420 \pm 180$ K and $T_e = 5390 \pm 230$ K at $P = 136$ GPa, respectively (Figures 2 and 6). The MgO-MgSiO$_3$ and MgSiO$_3$-SiO$_2$ eutectic temperatures cross at $P \sim 125$ GPa, reflecting the opposing trends of eutectic $X_{SiO_2}$ (Figure 5). The difference between $T_e$ and the melting curve of MgSiO$_3$ does not exceed 200 K in the entire LM. At CMB pressure, our model shows higher $T_e$ by 500–700 K than the models of de Koker et al. (2013) and Liebske and Frost (2012) for MgO-MgSiO$_3$, and higher $T_e$ by 800 K compared to de Koker et al. (2013) for MgSiO$_3$-SiO$_2$. The eutectic $T$ estimated by Baron et al. (2017) appear rather low in this context.
5. Melting Near the Core-Mantle Boundary

This similarity in $T_\mu$ and therefore the solidus $T_s$ for MgO-MgSiO$_3$ and MgSiO$_3$-SiO$_2$ predicted here suggests that melting of basalt at lower $T$ than peridotite (Andrault et al., 2014; Kuwahara et al., 2018; Pradhan et al., 2015; Tateno et al., 2014, 2018), invoked as an explanation for the occurrence of the ULVZ in the lowermost mantle (e.g., Thorne et al., 2019), is not simply tied to its higher $X_{\text{SiO}_2}$ value. Rather, further differences in chemical composition will influence $T_\mu$ and melt relations either directly within a solid solution (e.g., a higher FeO or Al$_2$O$_3$ content in bridgmanite) or through changes in the resulting phase assemblage. Compared to mantle peridotite (Workman & Hart, 2005), oceanic crust is strongly enriched in Al$_2$O$_3$, CaO, and Na$_2$O, while the FeO content is similar in both lithologies (e.g., Chemia et al., 2015). For oceanic crust this leads to a phase assemblage that differs in various aspects from peridotite at LM pressure: (a) it contains significantly more CaSiO$_3$ perovskite (e.g., Chust et al., 2017) and (b) SiO$_2$ occurs as a free phase rather than the ferropericlase solid solution. Both CaSiO$_3$ perovskite and SiO$_2$ stishovite dissolve a few % of Al$_2$O$_3$ and very little FeO (e.g., Kuwahara et al., 2018; Tateno et al., 2018), while ferropericlase in a peridotitic composition contains up to 20% FeO (e.g., Chust et al., 2017; Kuwahara et al., 2018). Further, (c) for basalt an Al$_2$O$_3$ dominated mineral occurs in the LM, either as Ca-ferrite (Irifune & Ringwood, 1993; Tateno et al., 2018) or the NAL phase (Imada et al., 2012; Kato et al., 2013). As a consequence of these phase relations, bridgmanite in a basaltic composition can be expected to be significantly enriched in FeO compared to a mantle peridotite. A high FeO content in partial melts from basalt compositions at LM pressure (Kuwahara et al., 2018; Pradhan et al., 2015; Tateno et al., 2018) suggests that the solidus $T$ in the resulting bridgmanite is significantly lower than for the eutectic in the MgSiO$_3$-SiO$_2$ system established here. This is further supported by the observation that while CaSiO$_3$ perovskite is the phase on the solidus in peridotite at LM pressure, it becomes the liquidus phase for MORB compositions, and bridgmanite simultaneously becomes the solidus phase (Kuwahara et al., 2018; Pradhan et al., 2015; Tateno et al., 2014, 2018).

However, the chemical complexity of natural or model systems investigated in these studies makes it challenging to disentangle the causes for these discrepancies. In order to analyze the influence of other chemical components in a systematic way by building up a thermodynamic model for mantle compositions, controlled melting experiments in the MA press like the ones performed here are required, ideally for MgO/SiO$_2$ ratios of the eutectic compositions. Not only dictated by chemical abundances, but also the crystal-chemical and phase relation arguments outlined in the previous paragraph, such a model should be built from the two-component system explored here to the FeO-MgO-SiO$_2$ and subsequently the FeO-MgO-Al$_2$O$_3$-SiO$_2$ systems. These will provide successive insights into the melting relations, for example, from the influence of FeO (and Al$_2$O$_3$) in the bridgmanite solid solution and from an additional Al$_2$O$_3$-rich phase in the assemblage.

With bridgmanite the liquidus phase in a peridotite composition (Tateno et al., 2014), the shallow slope of its melting curve through much of the LM opens the possibility that crystallization of the mantle may have indeed initiated from the middle of the mantle, as proposed in a number of studies (Andrault, 2019; Stixrude et al., 2009). Most prominently, this would lead to the formation of a basal magma ocean (Herzberg et al., 2013; Labrosse et al., 2007), and a separation of two magma ocean reservoirs would strongly be aided if the crystallizing layer is neutrally buoyant. While bridgmanite is denser than the coexisting liquid with $X_{\text{SiO}_2}$ < 0.5, the strong partitioning of FeO into the melt (Nomura et al., 2011; Tateno et al., 2014) may lead to a situation in which the density contrast between magma ocean liquid and crystallizing bridgmanite is minimal.

With the eutectic in the MgSiO$_3$-SiO$_2$ system exceeding $X_{\text{SiO}_2} = 0.60$ at $P \sim 70$ GPa, magma oceans in rocky exoplanets would likely crystallize bridgmanite rather than stishovite for most plausible scenarios, and therefore...
not create a very dense $\text{SiO}_2$-dominated layer deep in their mantle that would impede the onset of convection and therefore efficient heat loss from the planet (e.g., Tosi et al., 2013).

6. Conclusions

In multi-anvil experiments at 24 GPa we have measured the compositions of partial melts formed from a Mg$_3$SiO$_5$ starting composition at a temperature of $2750 \pm 100$ K. Electron microscope and microprobe analysis of the recovered experimental charges reveals that stishovite is the crystalline phase at the liquidus, and the coexisting liquid has a composition of $X_{\text{SiO}_2} = 0.53 \pm 0.03$. This implies an Mg$_3$SiO$_5$-$\text{SiO}_2$ eutectic that is very close to Mg$_3$SiO$_5$ bridgmanite in composition which is therefore the liquidus phase in the MgO-$\text{SiO}_2$ system over a compositional range from $X_{\text{SiO}_2} = 0.43$ (Liebske & Frost, 2012) to $X_{\text{SiO}_2} = 0.53$, with very similar eutectic (solidus) temperatures. Combining our eutectic with that of Liebske and Frost (2012) for MgO-Mg$_3$SiO$_5$ and thermodynamic models for the liquid components $\text{SiO}_2$ and MgO as well as the solid phases $\text{SiO}_2$ stishovite, MgO periclase and Mg$_3$SiO$_5$ bridgmanite, we have modeled the melting phase relations in the MgO-$\text{SiO}_2$ system using a symmetric solution model for the liquid components.

The extrapolation of our thermodynamic description for melting relations to higher pressures hinges on its temperature dependence and the melting curves of MgO and $\text{SiO}_2$. Our model describes the melting curve of Mg$_3$SiO$_5$, bridgmanite consistent with diamond anvil cell experiments up to 60 GPa, and recent shock wave experiment to 80 GPa. In our model, the eutectic compositions move to $X_{\text{SiO}_2} = 0.35 \pm 0.02$ and $X_{\text{SiO}_2} = 0.66 \pm 0.02$ at core-mantle boundary pressure, respectively. Eutectic temperatures of these two compositions remain similar, suggesting that lower solidus temperatures of basalt over peridotite in the deep mantle, which has been observed in a series of prior experiments on natural or model compositions, must be related to chemical components other than $\text{SiO}_2$ and MgO.

Data Availability Statement

In compliance with AGU's data availability requirements, the microprobe analysis of the melt in the experiment S7238 (Table 1) at the basis of the current work is available on Figshare (https://doi.org/10.6084/m9.figshare.16837090.v1).

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