

RESEARCH ARTICLE

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Key Points:

- A multiscale spectrum of mountain waves was observed during a mesospheric mountain wave event over New Zealand
- Temperatures of the mountain waves were derived using sodium density mixing ratios
- Average momentum fluxes associated with observed mountain waves were large compared to zonal means

Supporting Information:

- Figure S1
- Data Set S1
- Data Set S2
- Data Set S3
- Data Set S4
- Data Set S5
- Data Set S6
- Data Set S7

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Momentum Flux Spectra of a Mountain Wave Event Over New Zealand

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Abstract During the Deep Propagating Gravity Wave Experiment (DEEPWAVE) 13 July 2014 research flight over the South Island of New Zealand, a multiscale spectrum of mountain waves (MWs) was observed. High-resolution measurements of sodium densities were available from ~70 to 100 km for the duration of this flight. A comprehensive technique is presented for obtaining temperature perturbations, T' , from sodium mixing ratios over a range of altitudes, and these T' were used to calculate the momentum flux (MF) spectra with respect to horizontal wavelengths, λ_H , for each flight segment. Spectral analysis revealed MWs with spectral power centered at λ_H of ~80, 120, and 220 km. The temperature amplitudes of these MWs varied between the four cross-mountain flight legs occurring between 6:10UT and 9:10UT. The average spectral T' amplitudes near 80 km in altitude ranged from 7–13 K for the 220 km λ_H MW and 4–8 K for the smaller λ_H MWs. These amplitudes decayed significantly up to 90 km, where a critical level for MWs was present. The average MF per unit mass near 80 km in altitude ranged from ~13 to 60 m²/s² across the varying spectra over the duration of the research flight and decayed to ~0 by 88 km in altitude. These MFs are large compared to zonal means and highlight the importance of MWs in the momentum budget of the mesosphere and lower thermosphere at times when they reach these altitudes.

1. Introduction

Gravity waves (GWs) have been known for many years to play major roles in the dynamics and structure of the atmosphere from the Earth's surface into the mesosphere and lower thermosphere (MLT). A key role is the transport and deposition of momentum from sources at lower altitudes to regions of dissipation at higher altitudes. Momentum flux (MF) divergence causes flow accelerations in the direction of horizontal GW propagation, and the cumulative effects in the MLT are decelerations and even reversals of the zonal flow in both hemispheres and an induced residual circulation from the summer to winter hemisphere near the mesopause (Fritts & Alexander, 2003; Garcia & Solomon, 1985; Holton, 1982, 1984). Orography is a major source of GW generation, and orographic GWs are a key component in parameterizations used in global circulation and climate models (Alexander et al., 2010; Kim et al., 2003). Multiple satellite observations have demonstrated GW hot spot regions in the stratosphere over major orography, suggesting the significant role that mountain waves (MWs) play at these higher altitudes (Eckermann & Preusse, 1999; Gong et al., 2012; Hoffmann et al., 2013; Jiang et al., 2005; McLandress et al., 2000; Wu et al., 2006).

The Deep Propagating Gravity Wave Experiment (DEEPWAVE) observed GWs in the vicinity of an orographic GW hot spot over the South Island of New Zealand during June–July 2014 (Fritts et al., 2016). The National Center for Atmospheric Research/National Science Foundation (NCAR/NSF) Gulfstream V (GV) aircraft flew 25 research flights (RFs) out of Christchurch, NZ. Onboard GV instruments used during this campaign included both Rayleigh and sodium (Na) lidars allowing for measurements of the stratosphere and MLT, and an advanced mesospheric temperature mapper (AMTM), which provided temperatures derived from OH airglow emissions centered near ~87 km. Additionally, flight-level data were collected by in situ instrumentation. Through these campaign measurements, multiple MWs were observed throughout the atmosphere, and a number of these have been analyzed to date (Bossert et al., 2015, 2017; Bramberger

et al., 2017; Eckermann et al., 2016; Fritts et al., 2016; Kruse et al., 2016; Pautet et al., 2016; Smith & Kruse, 2017). This paper focuses on the multiscale MW event on 13 July 2014 observed from the GV RF, and calculations of its changing MF spectra over a range of altitudes and spatial regions. These calculations are performed using a presented technique for obtaining temperatures from mixing ratios associated with significant sodium density perturbations on the lower side of the sodium layer.

2. Measurements

During the RF on 13 July 2014, the sodium lidar was run in a special high-resolution mode that provided sodium density measurements at 1-s sampling using one frequency; however, this nonstandard single-frequency measurement configuration prohibited the derivation of temperatures provided by the standard two laser frequency measurements. These high-resolution measurements allowed for the observation of small-scale secondary GWs within the sodium layer with horizontal wavelengths $\lambda_H < 40$ km along the flight track (Bossert et al., 2017), as well as a $\lambda_H \sim 200$ –240 km MW, which persisted for the entire flight duration (Bossert et al., 2015). There were four flight passes during the RF across the South Island, and these all traversed the highest peak, Mt. Cook. These flight passes took place between 6:10UT and 9:10UT, with sodium lidar data acquisition beginning at 6:20UT.

The data used in this study were filtered using a boxcar moving average window over 1.8 km (6 altitude bins) and 15 s (15 temporal bins), the latter corresponding to an along-track horizontal resolution of ~ 3.6 km given a mean GV cruise speed of 240 m/s. During the second and third passes, the lidar data were affected by intermittent problems with the laser's locking to the sodium line. In order to use all of the data, a temporal low-pass filter was applied to remove locking-related anomalies from the data series while retaining GW horizontal wavelengths of $\lambda_H \sim 30$ km and larger. The applied filter had a passband of 100 s (~ 24 km along track) and a stopband of 50 s (~ 12 km along track).

3. Temperature and MF Measurements and Validation

Several methods are available to infer GW-induced temperature perturbations diagnostically from observed GW-induced sodium density perturbations. Sodium density perturbations have previously been used to calculate temperature perturbations associated with GWs (Bossert et al., 2014; Shelton et al., 1980; Swenson & Gardner, 1998), and sodium lidars have also been previously used to calculate MFs associated with GWs (Acott et al., 2011; Gardner & Liu, 2007). Assuming hydrostatic GWs and purely vertical and linear gradients in background temperatures and sodium densities, GW-induced perturbations in temperature can be derived from sodium density perturbations using parcel-advection methods (see Eckermann et al., 1998). The diagnostic relation relating sodium density perturbations (assuming conservation of the sodium mixing ratio and potential temperature following air motions) is given in equation (1),

$$\rho'_{\text{Na}} = \left[\frac{\bar{\rho}_{\text{Na}}}{\gamma H} + \frac{\partial \bar{\rho}_{\text{Na}}}{\partial z} \right] \left(\frac{gT'}{N^2 \bar{T}} \right) \quad (1)$$

where ρ'_{Na} is the sodium density perturbation, $\bar{\rho}_{\text{Na}}$ is the background sodium density, T' is the temperature perturbation, \bar{T} is the background temperature, γ is the ratio of specific heats, g is the gravitational acceleration, N is the background buoyancy frequency in rad/s, and H is the pressure scale height.

The GW-induced T' can also be calculated by using the observed vertical displacement of Na mixing-ratio isopleths. This approach was used as an approximate calculation for T' for the large scale ~ 240 km MW present during the 13 July 2014 RF in Bossert et al. (2015), as well as the ~ 40 km MW observed over the Auckland Islands the following day (Eckermann et al., 2016). In the absence of chemistry, the mixing ratio of any minor constituent is conserved under adiabatic isobaric advection. In such cases, the observed vertical displacement of isopleths of tracer mixing ratio from their undisturbed equilibrium altitudes, Δz_{mix} , provides a direct measurement of the GW-induced vertical displacement perturbation. The resulting temperature perturbation T' due to adiabatic expansion and compression of the air parcel during this vertical advection is given by equation (2),

Table 1
Variables Commonly Referred to in This Paper

Parameter	Description
T'	gravity wave temperature amplitude
\bar{T}	mean background temperature
g	gravitational acceleration
λ_H	horizontal wavelength
ρ'_{Na}	sodium density perturbation
N	buoyancy frequency
$\bar{\rho}_{Na}$	mean sodium density
γ	ratio of specific heats
Γ	adiabatic lapse rate
R	ideal gas constant
M	average atmospheric molecular weight
$k_H = 2\pi/\lambda_H$	horizontal wavenumber
$m = 2\pi/\lambda_z$	vertical wavenumber
u_H'	horizontal wind perturbation in GW propagation direction
U_H	horizontal wind in GW propagation direction
$H = R\bar{T}(z)/Mg$	pressure scale height

$$T' = -\Delta z_{mix} \left(\Gamma + \frac{dT}{dz} \right) \quad (2)$$

where Γ is the adiabatic lapse rate 9.5 K/km and $\frac{dT}{dz}$ is the background temperature gradient. Table 1 gives an overview of the variables used in equations (1) and (2) as well as those used later in this paper. Equation (1) is more accurate for cases in which perturbations are observed within the sodium layer with smaller parcel altitude displacement. Equation (2) works better for larger amplitude waves since Δz_{mix} is quantified directly from observed mixing ratios. Both methods also assume approximate hydrostatic equilibrium and adiabatic motion, which may not always be strictly valid for large-amplitude and short-scale GWs. For the observed GWs on 13 July, the amplitudes of the MWs on the bottom side of the layer were large enough to significantly perturb sodium densities from the layer. In this case, it is difficult to assess a relative background sodium density with respect to observed sodium density perturbations; thus, equation (1) would not be applicable. Therefore, equation (2) is used for the subsequent temperature analysis.

3.1. Temperature Calculations From Modeled Sodium Density

A previous 2-D simulation of the observed $\lambda_H \sim 240$ km MW during the 13 July 2014 event by Heale et al. (2017) provided GW-induced perturbations in both temperature and sodium densities. Sodium densities from this simulation were also published and used for the investigation of arising secondary features associated with primary MW breaking (Bossert et al., 2017). The fields from this simulation allow for sodium mixing ratio perturbations to be calculated and converted into temperature perturbations using equation (2), which can then be compared and objectively validated against the directly simulated temperature perturbations from the model. Sodium mixing ratios were derived using background densities from the Navy Global Environmental Model (NAVEM). The calculation for the sodium mixing ratio is given in equation (3),

$$R_{Na} = \frac{\rho_{Na}(z)T(z)}{\rho_{atm}(z_o)T(z_o) \exp \left[-\int_{z_o}^z \frac{dz}{H(z)} \right]} \quad (3)$$

where ρ_{atm} is the atmospheric background density. For the purposes of mixing ratio calculations in this paper, which use the background atmospheric density from NAVEM reanalysis, equation (3) simplifies to $R_{Na} = \rho_{Na}(z)/\rho_{atm}(z)$. From these mixing ratios, T' was calculated using equation (2) and the background temperature profile from NAVEM. Figure 1 shows the modeled T' and modeled sodium densities over a region of the 240 km MW with breaking and associated features occurring and corresponding T' calculated from the mixing ratio displacements for 49 different mixing ratio isopleths. As can be seen from Figures 1a

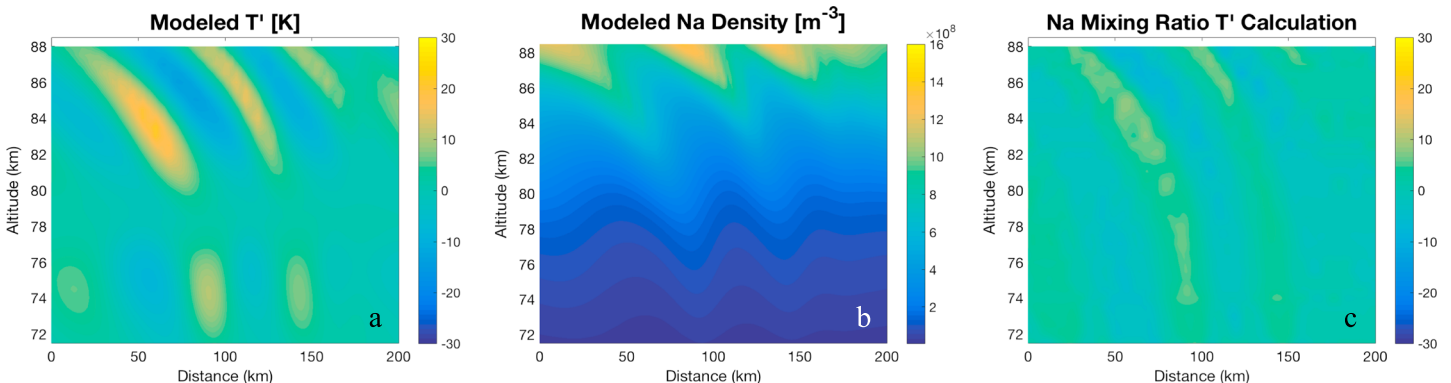


Figure 1. (a) The modeled T' from Heale et al. (2017). (b) The associated sodium densities from this model. (c) The T' calculated from the sodium mixing ratio calculated using equations (2) and (3).

and 1c, there is a reasonable match between the actual T' and calculated T' . The method is only applicable to adiabatic motions and will not provide a reliable estimate under conditions where horizontal advection is significant and diabatic motions are dominant. Additionally, because the method provides an average T' over an entire parcel displacement, when a parcel is displaced through a region of temperature fluctuations that are smaller in vertical scale than the overall parcel displacement, these fluctuations will not be resolved. This is apparent in the modeled data, as there is a small region of evanescence between 77 and 79 km evident in Figure 1a, and this is not resolved in the mixing ratio calculated temperatures shown in Figure 1c. The model comparisons in Figure 1 demonstrate that the calculation of T' from mixing ratios as would be calculated with sodium density measurements provides an estimate on MW T' amplitudes spatially and vertically in the absence of chemistry, which allows for estimates of MF when no other high-resolution direct methods of temperature measurement are available.

3.2. Influences of Sodium Chemistry

In order to test the chemical response of the Na layer to wave-driven oscillations in temperature, pressure, and the mixing ratios of minor constituents, we employed a full time-resolved model of Na chemistry. The 1-D model is essentially that described by Plane (2004), with recent updates to the rate coefficients of some reactions, which were remeasured in the laboratory (Gómez-Martín et al., 2016; Gómez-Martín et al., 2017). The 12 neutral reactions, 10 ion-molecule reactions, and 5 photochemical reactions are listed in Table 2. Na, Na⁺, NaOH, and NaHCO₃ are treated explicitly in the model, and the minor species (NaO, NaO₂, NaO⁺, Na⁺.N₂, and Na⁺.CO₂) are assumed to be in chemical steady state (Plane, 2004).

Table 2
Neutral and Ionic Gas-Phase Reactions in the Sodium Model

Number	Reaction	Rate Coefficient ^a
Neutral chemistry		
R1	Na + O ₃ → NaO + O ₂	$1.1 \times 10^{-9} \exp(-116/T)$
R2	NaO + O → Na + O ₂	$2.2 \times 10^{-10} (T/200)^{1/2}$
R3	NaO + O ₃ → Na + 2O ₂	$3.2 \times 10^{-10} (\exp(-550/T))$
R4	NaO + H ₂ → NaOH + H	4.9×10^{-12}
R5	NaO + CO → Na + CO ₂	9.0×10^{-11}
R6	NaO + H ₂ O → NaOH + OH	$5.1 \times 10^{-10} \exp(-240/T)$
R7	NaOH + H → Na + H ₂ O	3.9×10^{-11}
R8	NaOH + CO ₂ (+M) → NaHCO ₃	$1.2 \times 10^{-27} (T/200)^{-4.12}$
R9	NaHCO ₃ + H → Na + H ₂ CO ₃	$1.84 \times 10^{-13} T^{0.777} \exp(-1014/T)$
R10	Na + O ₂ (+M) → NaO ₂	$5.0 \times 10^{-30} (T/200)^{-1.22}$
R11	NaO ₂ + O → NaO + O ₂	$5.0 \times 10^{-10} \exp(-940/T)$
R12	2NaHCO ₃ (+M) → dimer	$8.8 \times 10^{-10} (T/200 K)^{-0.23}$
Ion-molecule Chemistry		
R20	Na + O ₂ ⁺ → Na ⁺ + O ₂	2.7×10^{-9}
R21	Na + NO ⁺ → Na ⁺ + NO	8.0×10^{-10}
R22	Na ⁺ + N ₂ (+M) → Na.N ₂ ⁺	$4.8 \times 10^{-30} (T/200)^{-2.2}$
R23	Na ⁺ + CO ₂ (+M) → Na.CO ₂ ⁺	$3.7 \times 10^{-29} (T/200)^{-2.9}$
R24	Na.N ₂ ⁺ + X → Na.X ⁺ + N ₂ (X = CO ₂ , H ₂ O)	6×10^{-10}
R25	Na.N ₂ ⁺ + O → NaO ⁺ + N ₂	4×10^{-10}
R26	NaO ⁺ + O → Na ⁺ + O ₂	1×10^{-11}
R27	Na.O ⁺ + N ₂ → Na.N ₂ ⁺ + O	1×10^{-12}
R28	Na.O ⁺ + O ₂ → Na ⁺ + O ₃	5×10^{-12}
R29	Na.Y ⁺ + e ⁻ → Na + Y (Y = N ₂ , CO ₂ , H ₂ O, O)	$1 \times 10^{-6} (T/200)^{-1/2}$
R30	Na ⁺ + e ⁻ → Na + hν	$3.9 \times 10^{-12} (T/200)^{-0.74}$
Photochemical reactions		
R31	NaO + hν → Na + O	5.5×10^{-2}
R32	NaO ₂ + hν → Na + O ₂	1.9×10^{-2}
R33	NaOH + hν → Na + OH	1.8×10^{-2}
R34	NaHCO ₃ + hν → Na + HCO ₃	1.3×10^{-4}
R35	Na + hν → Na ⁺ + e ⁻	2×10^{-5}

^aUnits: unimolecular, s⁻¹; bimolecular, cm³ · molecule⁻¹ · s⁻¹; termolecular, cm⁶ · molecule⁻² · s⁻¹. Rate coefficients are from Plane (2004), apart from R4, R5, R6, and R8 from Gómez-Martín et al. (2016), R7 from Gómez-Martín et al. (2017), and R30, which is calculated by Badnell (2006).

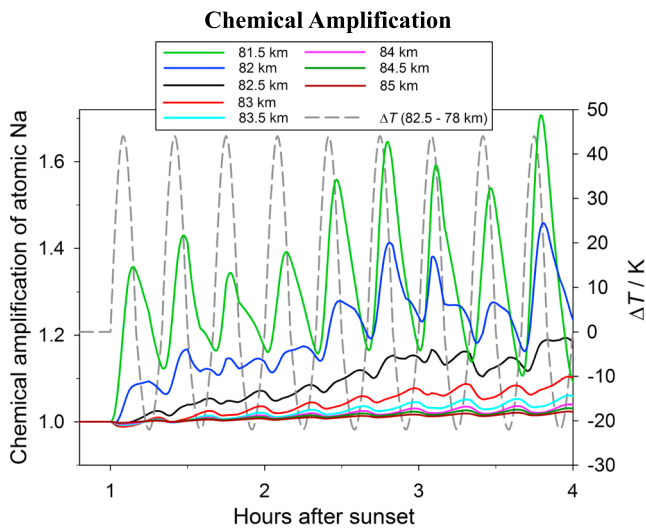


Figure 2. The chemical amplification factor (CAF) of atomic Na over 4 hr, for a wave period of 20 min. The black line shows the CAF when the Na vmr before the wave is 1.1×10^{-12} , which is at 82.5 km. The wave then displaces the air parcel down to 78 km, and up to 84.8 km (i.e., approximately 50% of the downward displacement). The resulting oscillation of the air parcel temperature is shown with the dashed grey line (right-hand ordinate axis). The CAFs for air parcels starting at other altitudes between 81.5 and 85 km, and displaced down to 78 km, are also shown in the figure.

The model was run under midlatitude conditions where all the minor atmospheric constituents, which control the Na chemistry (O_3 , O, H, NO^+ , O_2^+ , and electrons), vary diurnally. A time step of 30 s was used. One hour after sunset the wave was turned on with a period of 20 min. In order to examine the effect of wave-driven chemistry on the underside of the layer consistent with the DEEPWAVE observations, we considered vertical displacements from mean heights between 81.5 and 85 km down to 78 km. The upward excursions were set to be 50% of the downward displacements due to strong constraints on the wave amplitude approaching a critical level at ~ 90 km.

The model was used to determine the chemical amplification factor (CAF) of atomic Na, defined as the Na vmr in an air parcel when it is displaced by the wave and full chemistry operates, divided by the Na vmr when chemistry is turned off and Na is treated as an inert tracer. Figure 2 illustrates how the CAF varies over 4 hr for the wave period of 20 min. The CAF is largest, up to 1.6, starting from a mean height of 81.5 km. This is where the Na vmr is very small (4.8×10^{-14}) at the base of the layer, and there is a very large reservoir of $NaHCO_3$. This reservoir species is converted into Na via reaction R9 (Table 2), which has a significant activation energy and thus produces Na rapidly due to adiabatic heating of the air parcel during the downward excursion. During the subsequent upward excursion, the CAF decreases as the temperature falls, though the CAF does not return to 1. This is because of nonlinear processes in the model; the third-order reactions R8, R10, and R12; and the smaller upward wave excursion.

At higher mean altitudes, the ratio of Na to $NaHCO_3$ increases rapidly, and so the CAF is essentially unity starting from a mean height of 85 km (Figure 2). At a mean height of 82.5 km, where the Na vmr is 1.1×10^{-12} and thus detectable by the airborne lidar operating at high time resolution, the CAF is less than 1.2.

3.3. Temperature Calculations From Measured Sodium Densities

Given the potential for mixing ratios to provide an estimate of GW T' , they are used here for the calculations of T' associated with the multiple scales of MWs observed from the GV on 13 July 2014. Although the mixing ratios have been used previously for rough temperature estimates (Bossert et al., 2015; Eckermann et al., 2016), the analysis presented here corrects for chemical influences on the bottom side of the layer and investigates changes in temperature perturbation estimates with altitude. Additionally, the data presented here have been filtered using a low-pass filter instead of the long temporal boxcar averaging which was previously applied in Bossert et al. (2015). This allows for a more precise assessment of the mixing ratio perturbations and temperature amplitudes due to $\lambda_H < 100$ km. The mixing ratios calculated using equation (3) for the four flight segments across the South Island are shown in Figure 3. For the T' calculation, mixing ratios are calculated along 27 mixing ratio isopleths with average altitudes ranging from 79 to 89 km. The average altitude of each isopleth is used for the altitude of the calculated T' from equation (2). A correction factor accounting for the CAF shown in Figure 2 is then applied to the calculated T' . Correction factors plotted over a range of mixing ratios are shown in Figure 4. T' amplitudes are divided by the correction factors for each mixing ratio isopleth. Figure 5a shows the mixing ratio contours with one highlighted in red, which corresponds to an average altitude of 81.5 km. The corresponding temperature perturbations for this contour are shown in Figure 5b in red, and the corrected temperature perturbations accounting for the CAF are shown in black. The calculated temperature perturbations for each mixing ratio contour during the second pass are shown in Figure 5c. The calculated T' from mixing ratio contours for each pass are shown in Figure 6. The background dT/dz used in equation (2) was obtained from NAVGEM reanalysis (Eckermann et al., 2018), reinterpolated onto a constant geometric height grid for the 13 July 2014 flight. The temperature profile was the average of all reanalysis grid points within a circular region centered near Mt. Cook with a 600 km great circle radius, and for times from 0600–0900 UTC. These temperatures are shown in Figure 7.

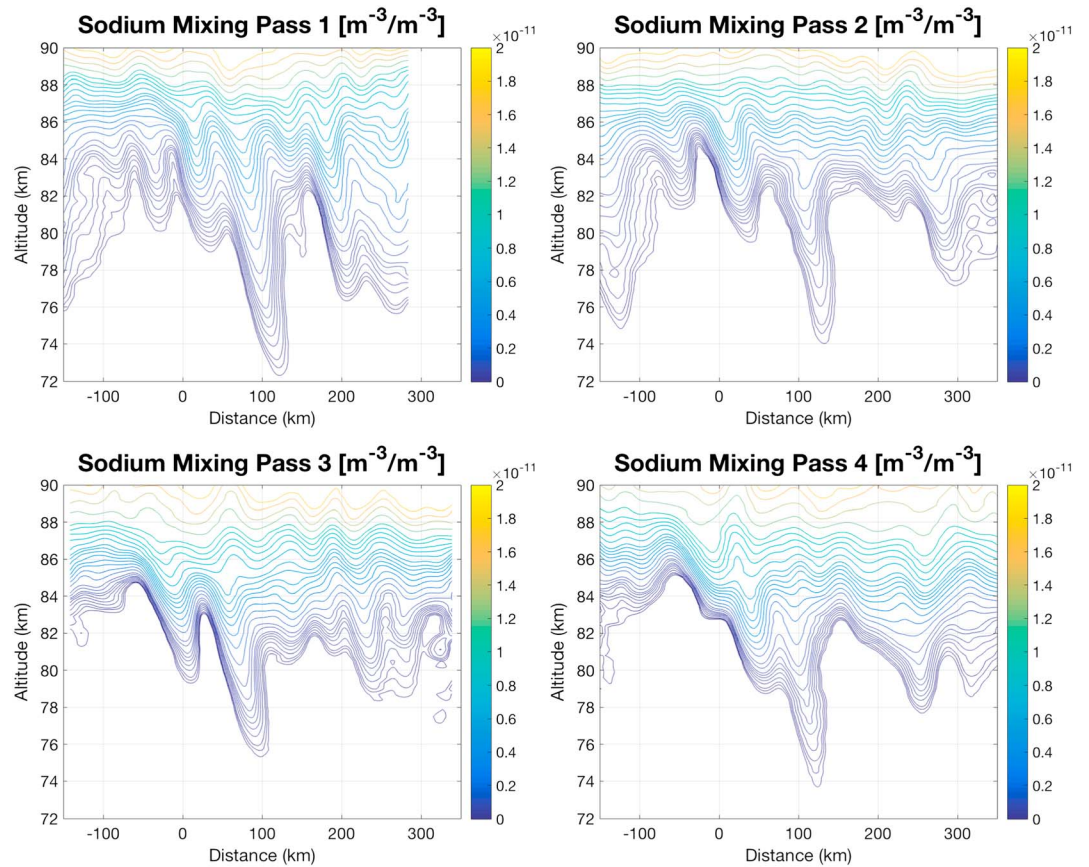


Figure 3. Sodium mixing ratio contours for all four passes calculated from observations using equation (3).

The T' given in Figure 6 reveal a superposition of multiple GWs with a range of different λ_H . In order to estimate the T' for each specific GW present, a fast Fourier transform (FFT) is applied to the along-track data at any given height, which provides a spectral amplitude for each transect. For each pass, data were zero-padded to length of 4,000 km. To reduce FFT sidelobes, data for the first and last 24 km were smoothed to a value of zero. The resulting spectral amplitudes are given in Figure 8. The FFT for each pass reveals spectral power centered around $\lambda_H \sim 220, 120,$ and 80 km, with the spectral power being most significant on the first and second passes. The λ_H are consistent with observations at the flight level of the GV, and stratospheric lidar observations of MWs (Fritts et al., 2018), suggesting that these observed wavelengths are likely associated with MWs. As expected for MWs, all amplitudes for the $\lambda_H \sim 220, 120,$ and 80 km decay upon approach to the MW critical level near 90 km. However, we also note that there may be contributions from GWs, which are not MWs. For example, scales of $\lambda_H < 60$ km are observed during each transect, which persist or arise near the critical level, indicating a non-zero phase speed. Additionally, in these nonMW cases, it is difficult to predict the associated phase speed, orientation, and thus, related MF. As demonstrated by the FFT plots, there is variation in MW amplitudes between each pass, and variation in the amplitudes between different λ_H , which is expected as forcing conditions have variation in time, and varying λ_H will have different responses to atmospheric fluctuations during upward propagation. Additionally, differences in dissipation can occur based on the initial amplitude of the wave and the

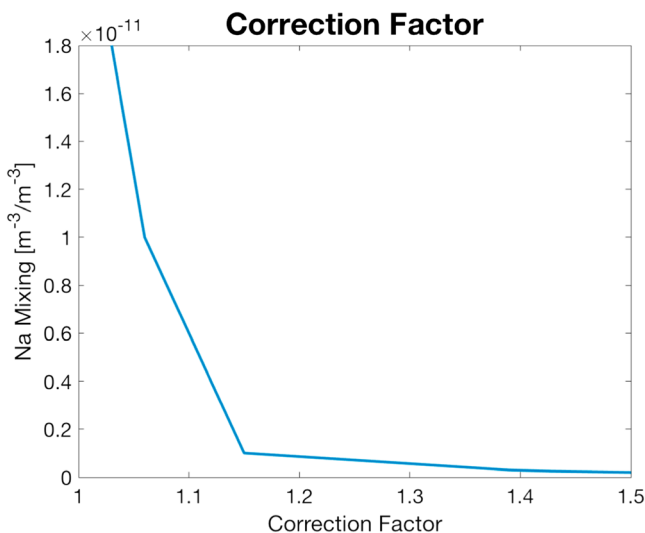


Figure 4. The correction factor obtained from Figure 2 and applied to the T' calculated from sodium mixing ratio displacements.

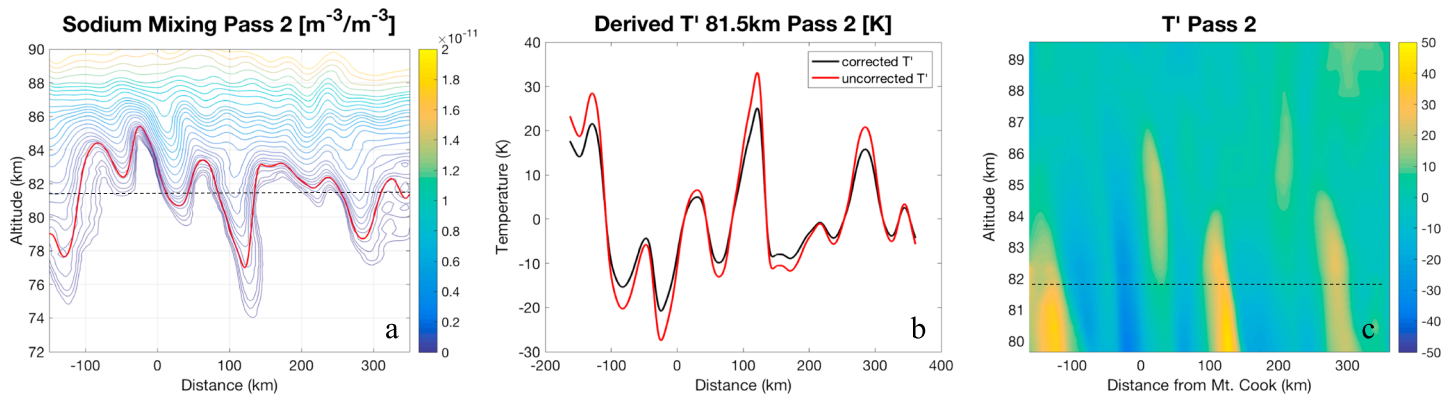


Figure 5. (a) Sodium mixing ratio contours for the second pass with a selected contour highlighted in red and the corresponding mean altitude of the contour shown by the black dotted line. (b) The corresponding T' derived from this contour using equation (2) in red, and the T' corrected for chemical amplification in black. (c) The corrected T' derived for each contour, and the black dotted line corresponds to the altitude for the temperature perturbations given in plot (b).

horizontal wavelength, as breaking is tied to multiple aspects of the GW and occurs upon approach to the critical level (Andreassen et al., 1998). The average spectral T' near 80 km in altitude varied from 7 to 13 K for the 220 km λ_H MW between each pass and ranged from 4 to 8 K for the smaller λ_H MWs for each pass. The λ_H and corresponding spectral average T' shown in Figure 8 are used to calculate MF spectra, which is discussed in the following section.

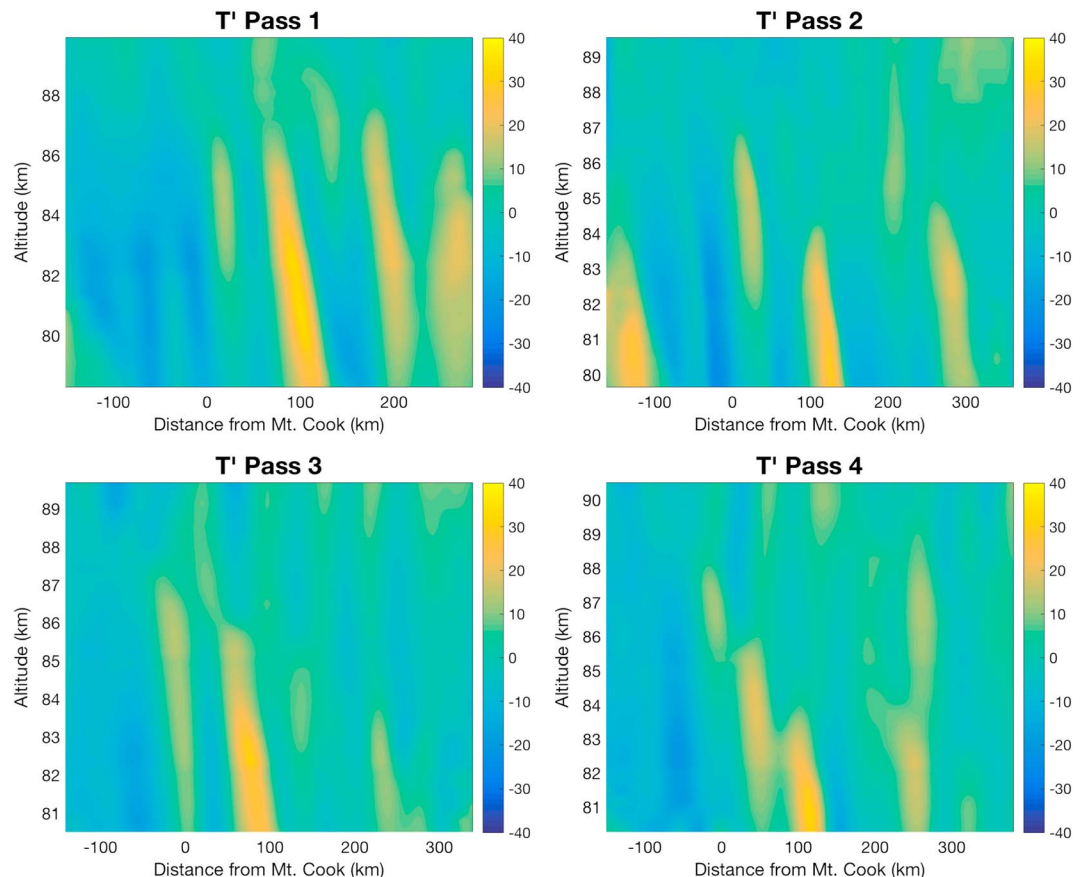


Figure 6. Calculated T' for each contour shown in Figure 2 using equation (2).

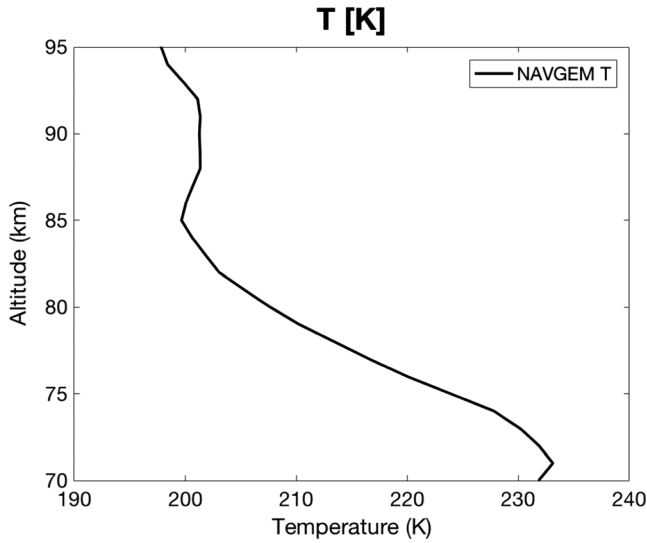


Figure 7. Average NAVGEM temperatures for the duration of RF22 from 6:00 to 9:00 UTC over a 600 km radius circle centered near Mt. Cook.

3.4. MF Spectra Calculations

Assuming hydrostatic motions, the MF per unit mass for a given GW can be calculated as (Bossert et al., 2015; Ern et al., 2004; Fritts et al., 2018),

$$MF = \langle u'_H w' \rangle = \left\langle \frac{1}{2} \left(\frac{T' g}{\bar{T} N} \right)^2 \left(\frac{k_H}{m} \right) \right\rangle \quad (4)$$

where u'_H and w' are the horizontal and vertical wind perturbations of the GW, k_H and m are the horizontal and vertical wavenumbers, and braces denote the mean over the observation segment. While k_H can be obtained from the FFT and corresponding spectral temperature amplitude calculation, m must be calculated using the dispersion relation (Fritts & Alexander, 2003),

$$m^2 = \frac{N^2}{(c - U_H)^2} - \frac{1}{4H^2} - k_H^2 \quad (5)$$

where c is the Earth-relative phase speed assumed to be ~ 0 m/s for the observed presumed MWs and U_H is the background wind in the direction of the MW horizontal wavenumber vector. Winds for this calculation are obtained from the Kingston meteor radar located on Tasmania, and these have previously been used for analysis in Bossert et al. (2015). Given the small meridional wind magnitudes above 80 km, and the near-zonal propagation direction

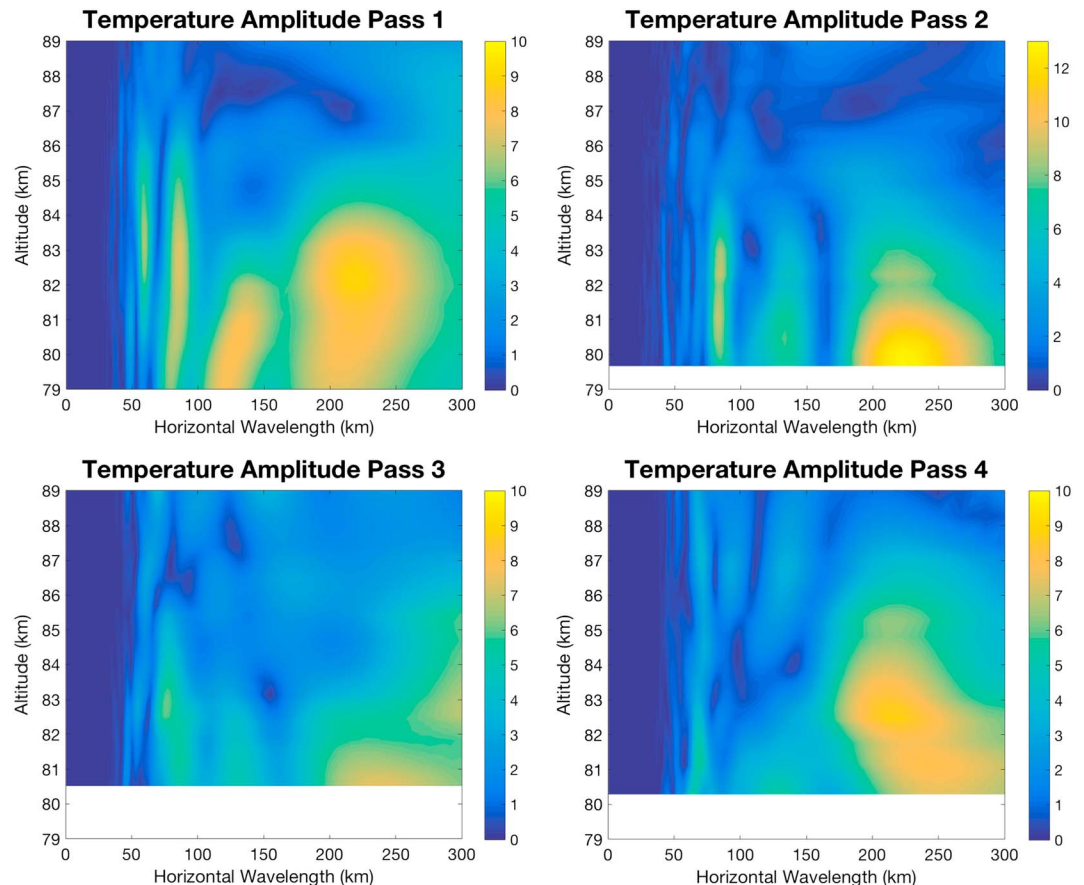


Figure 8. Spectral average T' for each pass calculated from an FFT of the T' shown in Figure 6.

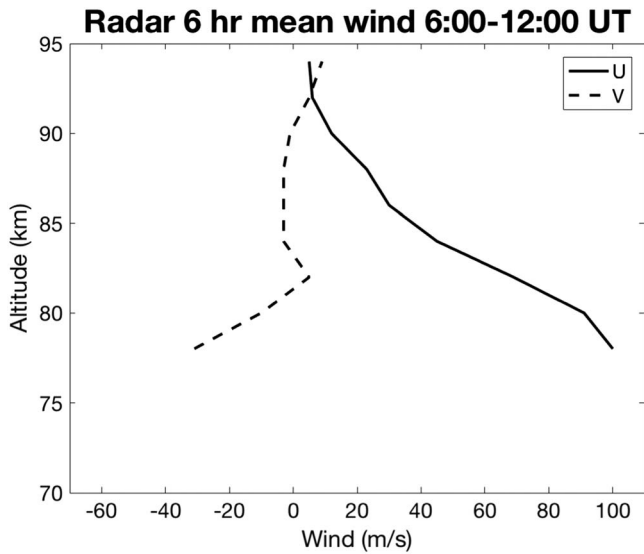


Figure 9. Zonal and Meridional Kingston Meteor radar winds on 13 July 2014 averaged from 6:00 to 12:00UT used in analysis by Bossert et al. (2015).

of the MWs, the zonal winds are used in equation (5) for the calculation of the vertical wavenumber. It is also assumed that the λ_H calculated from the FFT are approximately equal to the λ_H of the actual GWs, although it should be noted that there may be slight offsets in the calculation due to MW orientations slightly off-axis from a zonal propagation. AMTM observations shown in Bossert et al. (2015) demonstrated a nearly zonal orientation of the $\lambda_H \sim 220$ km MW, and other MWs generated under the same forcing conditions would be assumed to have similar orientation. The zonal and meridional winds are shown in Figure 9. The zonal winds imply a critical level near 90 km for stationary MWs. These winds were previously used in Bossert et al. (2015), and an offset time was used to account for tidal influences. It is clear that there is a critical level given the strong decay of MWs leading to 90 km. However, given the distance of the radar observations on Tasmania, the winds used in these calculations provide an estimate and not an exact value and may have associated errors. Given the MW decay in altitude is in agreement with the altitude of the critical level measured by the radar, the radar winds appear to provide a reasonable estimate of background winds with regard to the critical level location. Using equation (4) with the temperature amplitudes given in Figure 6, spectral MF values are obtained over a range of altitudes. The resulting spectral MFs per unit mass are shown in Figure 10. The MF calculations demonstrate the variation of MF between each pass, and between differing λ_H , which is not unexpected given the variability, which can arise in both forcing conditions and atmospheric

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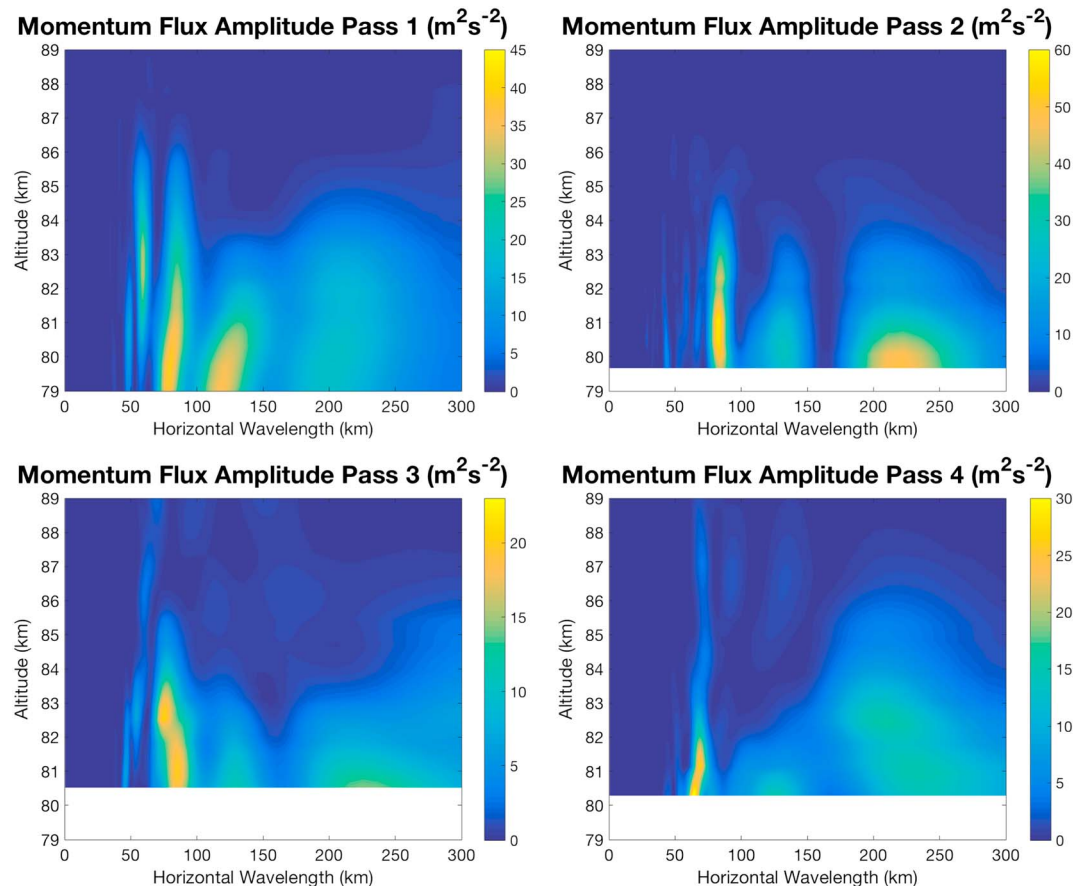


Figure 10. Spectral MF calculated using equation (4), spectral average T' shown in Figure 8, and NAVGEM zonal winds given in Figure 9.

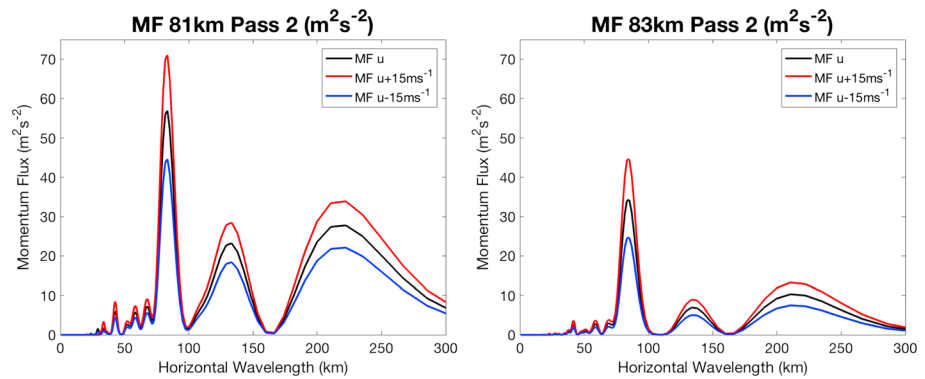


Figure 11. MF calculations for $u_H \pm 15$ m/s calculated at 81 and 83 km show error deviations of $\sim 25\%$ for a corresponding wind error of 15 m/s.

propagation conditions over time. It is most notable that the MF quickly decays in altitude, which is expected given the critical level near ~ 90 km. The first and second GV transects observed the largest MW T' and thus account for the largest MF contributions. The smaller λ_H of 80 and 120 km have MF values of about $35\text{--}60$ m^2/s^2 on the first and second pass near 81 km, with values significantly decreasing up to 85 km. MF values for these λ_H during the third and fourth passes decrease to values of $\sim 10\text{--}25$ m^2/s^2 near 80 km. The 220 km λ_H MW had the largest MF contribution during the second pass with MF values of ~ 50 m^2/s^2 near 80 km. These values are between 10 and 25 m^2/s^2 near 80 km in altitude for the remaining three passes. While horizontal scales of less than 60 km are included in these calculations, these shorter λ_H may have associated ground relative phase speeds, so it is unlikely that the MF calculations associated with these GWs are accurate given the assumed zero ground relative phase speed expected for MWs in the MF calculation. Additionally, as mentioned previously, the winds used in these calculations may have associated errors. To quantify potential errors, the MF was also calculated for winds ± 15 m/s. These calculations are shown for the second pass at 81 and 83 km in Figure 11 and demonstrate errors of $\sim 25\%$ for these potential wind deviations from measurements. It should also be noted that T' calculations have larger associated errors lower in altitude due to the varying chemistry at lower altitudes. Additionally, these calculations are an average estimate over the entire flight path, and localized MFs due to stronger phases over the mountains and directly in the lee of the mountains likely have larger associated MFs. Regardless of this, the spectral MF calculations show MF values that are larger than the average background MF values previously measured by radars and satellites, which have ranged from ~ 1 to 20 m^2/s^2 (Ern et al., 2018; Fritts et al., 2010, 2012; Fritts & Vincent, 1987; Murphy & Vincent, 1993; Nakamura et al., 1993; Reid et al., 1988; Tsuda et al., 1990; Vincent & Reid, 1983; Wang & Fritts, 1990). This finding is qualitatively in good agreement with Hertzog et al. (2012), who showed that MW events are particularly intermittent and strong events can carry very large MFs. Fritts et al. (2018) measured MFs associated with multiscale GW events observed in the OH layer for two separate events, finding maximum MF values for GW packets ranging from ~ 60 to 940 m^2/s^2 . The large MF value in that case was for a short, 10-min period GW that propagated through the OH layer within 45 min. For the 13 July event presented here, MF values were calculated over a sustained period of GW activity over several hours and demonstrate the spectral nature of MWs and varying MF over a range of MWs present in the MLT over New Zealand during this flight.

4. Conclusions and Discussion

The results presented here demonstrate the significant values and variability of MF for a multiscale MW event using a technique for estimating the MF spectra of MWs observed in the sodium density layer during the DEEPWAVE campaign. This technique utilizes observed vertical excursions in sodium mixing ratio isopleths to estimate T' at a given altitude using diagnostic relations derived from parcel relations valid for adiabatic GWs. Na chemistry is shown to have a minor effect on the estimates of T' . The average T' for each wavenumber is then used to assess the mean MF for that GW within the resolved spectrum. Importantly, this necessarily underestimates the total MF when GW amplitudes vary along the flight pass, as it provides an average value.

The results using this T' measurement technique for the 13 July 2014 DEEPWAVE flight demonstrate MFs distributed among a broad range of λ_H along the flight transects. Strongest T' and MF values were observed centered near $\lambda_H \sim 220, 120,$ and 80 km, which may be due to multiscale MW generation at the ground level or due to harmonic MW generation from the primary 220 km MW. The observations demonstrate the variability of MWs, which contribute to the vertical transport of horizontal momentum over a GW hot spot. The calculated T' and corresponding MF for MWs within each flight pass spectrum both decrease strongly approaching the critical level near 90 km. However, below this region, MF values are significant over a range of MW λ_H , which are sustained over the period of the several-hour RF.

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