

# Treatment of microphysics in the MPI-Hamburg LES model

Frank Müller, Andreas Chlond

MPI for Meteorology, Hamburg



## Outline

- ♦ The microphysical modules in the MPI-LES model
- ♦ Cloud droplet formation – activation of aerosols
- ♦ Comments on submitted results
- ♦ Outlook



# The microphysical modules in the MPI-LES model

- 3 (+ 1) microphysical schemes are implemented in the current LES version:

- 1.no-rain microphysics: sat. adjustment to form cloud water (including sub-scale condensation), no droplets (Sommeria and Deardorff, 1977)

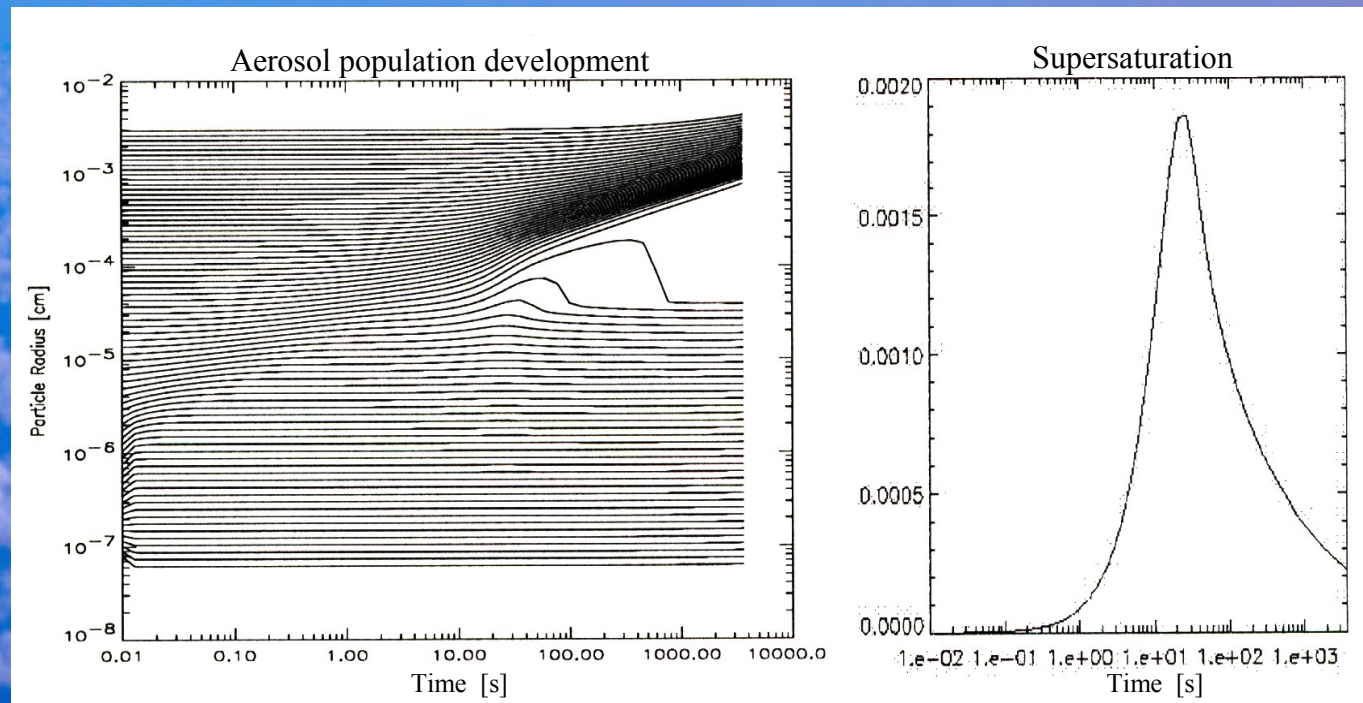
- 2.classical 'Kessler' scheme:  $q_c$ ,  $q_r$ , sat. adjustment, auto-conversion, accretion, sedimentation and evaporation of  $q_r$  (Kessler, 1969)

- 3.'Lüpkes' scheme:  $q_c$ ,  $q_r$ ,  $N_r$ , 2.+ rain drop self-collection, sedimentation of  $N_r$  (Lüpkes, 1989; Lüpkes, 1991)

- 4.two-moment scheme for cloud and rain water, resp.:  $q_c$ ,  $q_r$ ,  $N_c$ ,  $N_r$ , 3.+ self-collection of  $N_c$ , sedimentation of  $q_c$  and  $N_c$ , activation (Seifert and Beheng, 2000; Seifert, 2002)



# Cloud droplet formation – activation of aerosols



- A detailed treatment of aerosol population development is very computer resource demanding.
- Since non-activated aerosols are in thermodyn. equilibrium aerosol activation is in general a good approximation to describe the corresponding cloud droplet sources



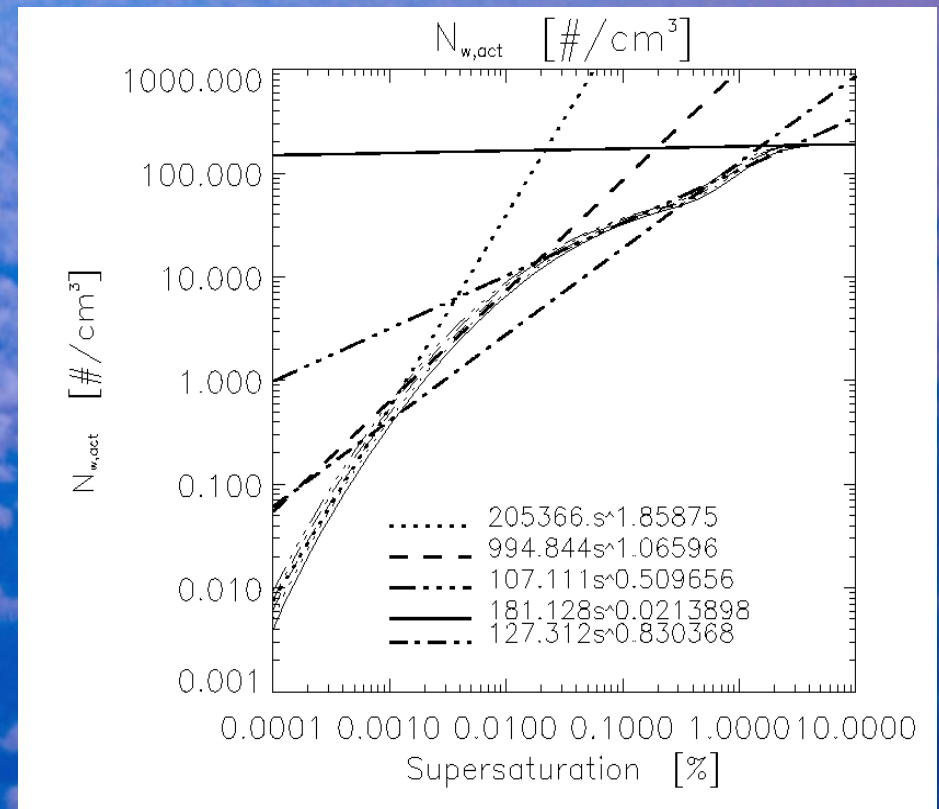
# Cloud droplet formation – activation of aerosols (cont.)

- Activation is based on  $N_{CCN}$  spectra (Twomey, 1959; Twomey and Wojciechowski, 1969; Hudson, 1980)

$$N_{CCN}(s) = C s^k$$

- The power law parameters  $C$  and  $k$ , given in the legend, were fitted to the activation spectrum based on the given aerosol size distribution.
- The activation spectrum is gained from:

$$N_{CCN}(s) = - \int_{r_{ap}(s_c=0)}^{r_{ap}(s_c=s)} f_{ap}(r_{ap}) dr_{ap} = \int_{r_c}^{\infty} f_{ap}(r_{ap}) dr_{ap}$$





## Cloud droplet formation – activation of aerosols (cont.)

- To calculate the no. of newly activated cloud drops the maximum supersaturation has to be determined:

$$\frac{ds_{vw}}{dt} = A_1 w + A_2 \frac{dq_c}{dt} = f(s_{vw}) = 0$$

$$A_1 = \frac{\epsilon l_{21} g}{R_a T^2 c_{pa}} - \frac{g}{R_a T} \quad A_2 = \frac{p}{\epsilon e_{sat,w}} + \frac{\epsilon l_{21}^2}{R_a T^2 c_{pa}}$$

- The determination of the cloud water mixing ratio tendency is crucial!

$$\frac{dq_c}{dt} = 2\pi \frac{\rho_w}{\rho_a} (2 A_3)^{2/3} s(t) \int_0^s n(s') \left[ \int_{t_o}^t s(t') dt' \right]^{\frac{1}{2}} ds'$$

- Using the droplet growth equation yields:

$$r \frac{dr}{dt} \approx A_3 s \quad \text{with} \quad A_3 = \left[ \frac{\rho_w R_v T}{e_{sat,w} D_v} + \frac{l_{21} \rho_w}{k_a T} \left( \frac{l_{21}}{R_v T} - 1 \right) \right]^{-1}$$

$$r(t) = \left[ r(t_o)^2 + 2 A_3 \int_{t_o}^t s(t') dt' \right]^{\frac{1}{2}}$$



## Cloud droplet formation – activation of aerosols (cont.)

- According to the original work of Twomey (1959) the activation spectrum is written as:

$$n(s) = k C s^{k-1}$$

- *Taking into account only the loss of water vapor due to condensation of newly activated droplets* results in a change of cloud water mixing ratio as follows:

$$\frac{dq_c}{dt} = 2\pi \frac{\rho_w}{\rho_a} A_3 k C \left[ \frac{A_3}{A_1 w} \right]^{\frac{1}{2}} s^{k+2} B \left( \frac{k}{2}, \frac{3}{2} \right)$$

- Maximum supersaturation  $s_{\max}$  and  $N_{\text{CCN}}$  are given by:

$$s_{\max} = C^{-\frac{1}{k+2}} \left[ \frac{(A_1 w)^{\frac{3}{2}}}{2\pi \frac{\rho_w}{\rho_a} A_2 A_3^{\frac{3}{2}} k B} \right]^{\frac{1}{k+2}} \quad N_{\text{CCN}, \max} = C^{\frac{2}{k+2}} \left[ \frac{(A_1 w)^{\frac{3}{2}}}{2\pi \frac{\rho_w}{\rho_a} A_2 A_3^{\frac{3}{2}} k B} \right]^{\frac{k}{k+2}}$$



# Cloud droplet formation – activation of aerosols (cont.)

- Extending the previous derivation by considering also *condensation on already existing cloud droplets* yields:

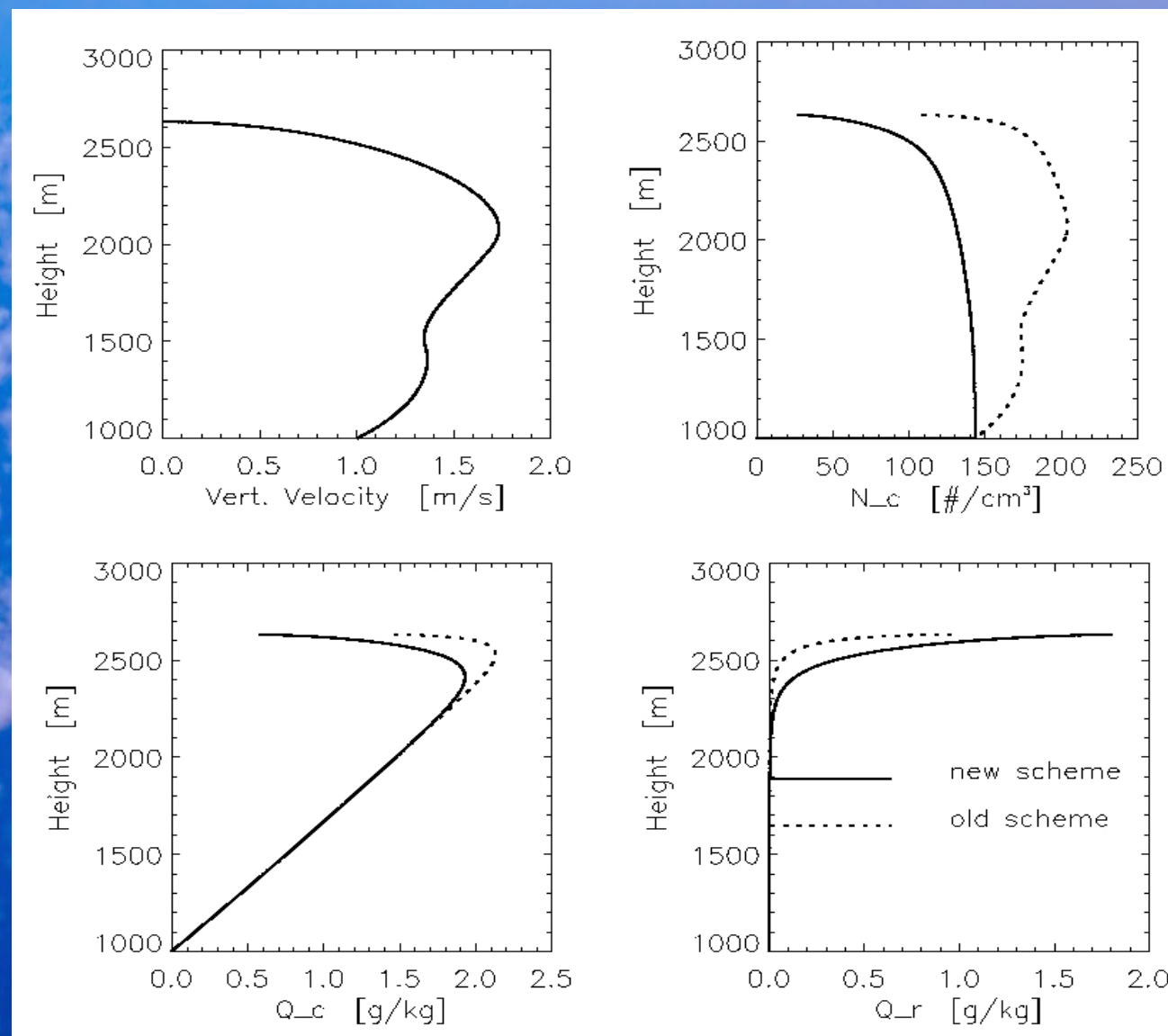
$$\begin{aligned} \frac{dq_l}{dt} &= \frac{dq_l}{dt}_{act} + \frac{dq_l}{dt}_{cond} \\ &= 2\pi \frac{\rho_w}{\rho_a} A_3 k c B\left(\frac{k}{2}, \frac{3}{2}\right) \left[\frac{A_3}{A_1 w}\right]^{1/2} + 4\pi \frac{\rho_w}{\rho_a} A_3 \bar{N}_c \left[\bar{r}_c^2 + \frac{A_3}{A_1 w} s^2\right]^{1/2} \end{aligned}$$

- The maximum supersaturation  $s_{max}$  is given by:

$$\begin{aligned} s_{max}^{n+1} &= s_{max}^n - \frac{f(s)}{f'(s)} \\ f'(s_{vw}) &= 2\pi \frac{\rho_w}{\rho_a} A_2 A_3 \left[ k(k+2) \left[\frac{A_3}{A_1 w}\right]^{1/2} s_{vw}^{k+2} + 2\bar{N}_c \left[\bar{r}_c^2 + \frac{A_3}{A_1 w}\right]^{1/2} + \right. \\ &\quad \left. + 2\bar{N}_c s_{vw} \frac{A_3}{A_1 w} \left[\bar{r}_c^2 + \frac{A_3}{A_1 w} s_{vw}^2\right]^{1/2} \right] \end{aligned}$$



## Cloud droplet formation – activation of aerosols (cont.)



Simulation results from an entraining air parcel model using the two-moment scheme.

..... std. Activation (Twomey, 1959)  
—— modified activation scheme



## Comments on submitted results

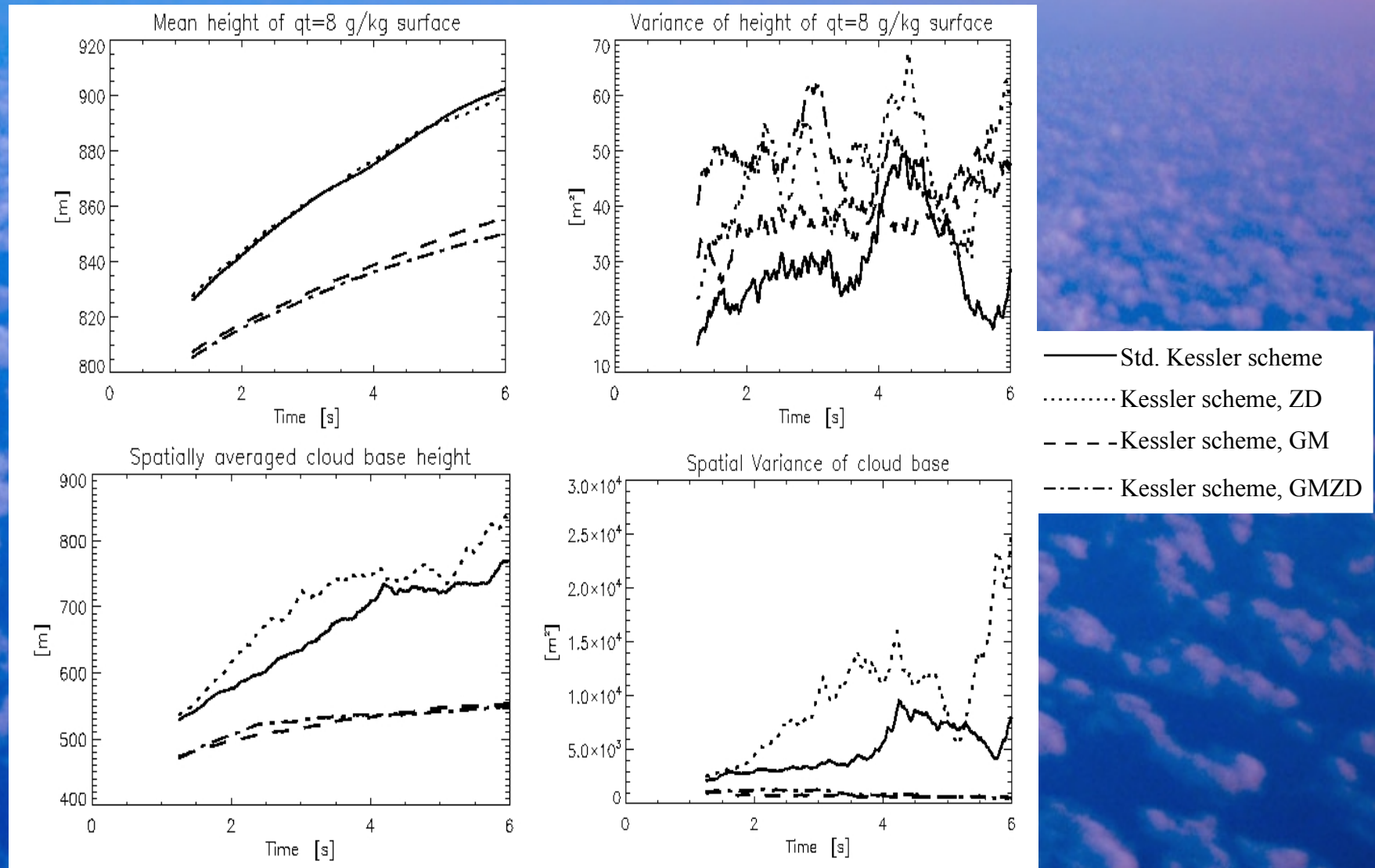
Our submitted results had the following features:

- Too fast increasing cloud top and cloud base
- Cloud breakup (Lüpkes scheme) with  $1 < cc < 0.4$
- Gradually diminishing LWP:  $t=6h: 70d/m^2 < LWP < 20g/m^2$
- Precipitation only with Kessler scheme

What is the reason for this behaviour? Wrong physics? Bad numerics?

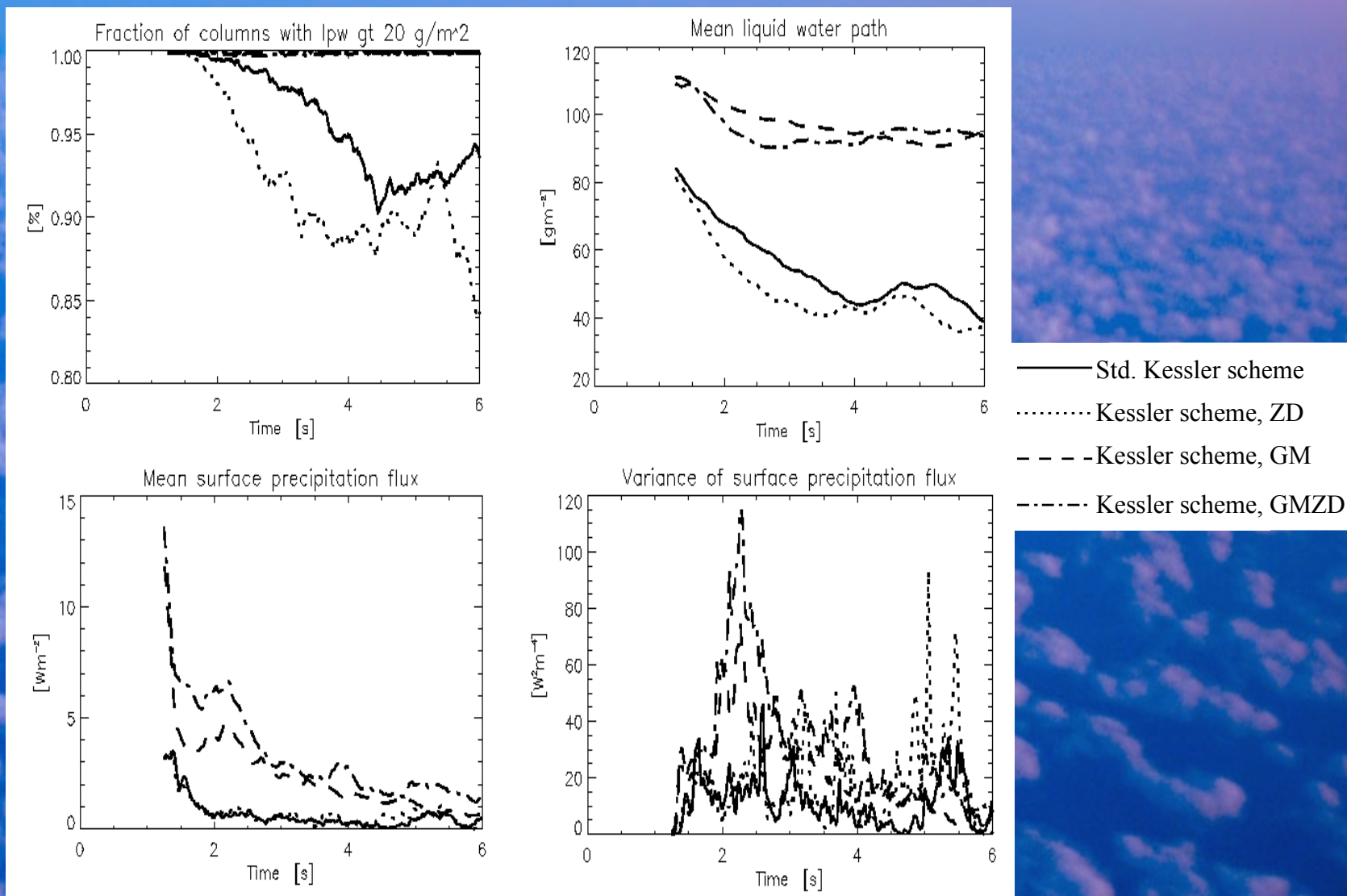


## Comments on submitted results (cont.)





## Comments on submitted results (cont.)





# Outlook

- Perform simulations with the new two-moment scheme
- Implementation of a detailed (size resolved) cloud microphysical scheme
- Look at aerosol-cloud interactions

