



énergie atomique • énergies alternatives

A new model for spray interactions.

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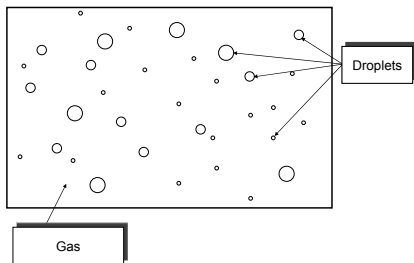
1 Context

- Diesel engines,
- Ariane boosters,
- Aerosols.

2 Goal :

- Having a test code able to deal with various applications,
- Treating cases with numerous droplets of different sizes,
- Establishing test cases.

- 1 Droplets are injected into a gas.



- 2 Euler equations are used for the motion of the gas.
- 3 A kinetic model is set for droplets.

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- u_p : droplet velocity,
- e_p, T_p : internal energy and temperature of a droplet,
- r : droplet radius,
- ρ_p : droplet density,
- m_p : droplet mass
- $f(t, x, u_p, e_p, r)$: droplet density function ,
- Q : collision kernel,
- Q_b : break-up kernel,
- ρ : gas density,
- u : gas velocity,
- e : gas internal energy,
- p : gas pressure,
- T : gas temperature,
- α : gas volume fraction,



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- λ : gas thermal conductivity,
- η : gas dynamic viscosity,
- C : gas specific heat,
- $Pr = \frac{\eta C}{\lambda}$: Prandtl number ,
- Ma : Mach number of the gas,
- $Re = \frac{2r\rho|u_p - u|}{\eta}$: droplet Reynolds number,
- $D_p = \frac{1}{2}\rho\pi r^2 C_d |u_p - u|$: drag coefficient
 $C_d(\alpha, Re, Ma)$,
- $Nu(Re, Pr)$: Nusselt number, ($Nu = 1 + 0.3Re^{1/2}Pr^{1/3}$)
- $\Gamma_p = \frac{\partial u_p}{\partial t}$: droplet acceleration ,
- $\phi_p = \frac{\partial e_p}{\partial t}$: droplet thermal flux.

Set of equations



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$$\partial_t(\alpha\rho) + \nabla_x \cdot (\alpha\rho u) = 0,$$

$$\partial_t(\alpha\rho u) + \nabla_x \cdot (\alpha\rho u \otimes u) + \nabla_x p = - \int_{u_p, e_p, r} m_p \Gamma_p f du_p de_p dr,$$

$$\partial_t(\alpha\rho e) + \nabla_x \cdot (\alpha\rho e u) + p[\partial_t \alpha + \nabla_x \cdot (\alpha u)]$$

$$= \int_{u_p, e_p, r} \left[(m_p \Gamma_p + \frac{m_p}{\rho_p} \nabla_x p) \cdot (u - u_p) - 4\pi r \lambda Nu (T - T_p) \right] f du_p de_p dr,$$

$$\alpha = 1 - \int_{u_p, e_p, r} \frac{4}{3} \pi r^3 f du_p de_p dr = 1 - \int_{u_p, e_p, r} \frac{m_p}{\rho_p} f du_p de_p dr,$$

$$\partial_t f + u_p \cdot \nabla_x f + \nabla_{u_p} \cdot (f \Gamma_p) + \partial_{e_p} (f \phi_p) = Q(f, f) + Q_b(f),$$

$$m_p \Gamma_p = -\frac{4}{3} \pi r^3 \nabla_x p - D_p(u_p - u),$$

$$m_p \phi_p = 4\pi r \lambda Nu (T - T_p),$$

$$p = P(\rho, e),$$

$$e_p = E(T_p).$$

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Why a new model ?

- Difficulties arise with small gas volume fractions.
- Lagrangian/Eulerian simulations are costly and badly scalable on a parallelism level : we need to make computations on droplets as less as possible and choose properly the ones we keep.
- Methods of moments for the droplet density functions have proven to be efficient for some applications (work of M.MASSOT AND AL. : 'Eulerian quadrature based moment models for dilute polydisperse evaporating sprays').

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- For small radii, droplets are rapidly at equilibrium with the gas (velocity/pressure/temperature) due to exchange terms : characteristic times for droplets are proportionnal to r^2 .
- Droplets can break up so obtaining small radii is frequent : there is always a characteristic radius (depending on the flow) for which equilibrium can be taken into account.



- We separate the phase space of the droplet density function in two parts :
 - the first part is made of droplets at equilibrium with the gas (velocity, pressure and temperature),
 - the second one contains the remaining droplets.
- Droplets at equilibrium are solved through a mix model.
- Others are still solved through a kinetic equation.

$$\partial_t(\rho) + \nabla_x \cdot (\rho u) = 0,$$

$$\partial_t(\rho u) + \nabla_x \cdot (\rho u \otimes u) + \nabla_x p = - \int_{u_p, e_p, r} m_p \Gamma_p f du_p de_p dr,$$

$$\partial_t f + u_p \cdot \nabla_x f + \nabla_{u_p} \cdot (f \Gamma_p) = Q_b(f),$$

$$m_p \Gamma_p = -D_p(u_p - u),$$

$$\rho = P(\rho).$$

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We split f in two parts :

$$f = f_1(t, x, v) + f_2(t, x, v). \quad (1)$$

f_1 (resp. f_2) is a droplet density function for droplets of radius r_1 (resp. r_2).

Toy model (2)



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$$\partial_t f_1 + u_p \cdot \nabla_x f_1 + \nabla_{u_p} \cdot (f_1 \Gamma_1) = -F f_1,$$

$$\partial_t f_2 + u_p \cdot \nabla_x f_2 + \nabla_{u_p} \cdot (f_2 \Gamma_2) = F f_1 \left(\frac{r_1}{r_2} \right)^3,$$

$$\Gamma_1 = \frac{D}{r_1^2} (u - u_p),$$

$$\Gamma_2 = \frac{D}{r_2^2} (u - u_p).$$

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Toy model (3)



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Let's define $\varepsilon = r_2/r_1$. We get

$$\begin{aligned} \partial_t(\rho^\varepsilon) + \nabla_x \cdot (\rho^\varepsilon u^\varepsilon) &= 0, \\ \partial_t(\rho^\varepsilon u^\varepsilon) + \nabla_x \cdot (\rho^\varepsilon u^\varepsilon \otimes u^\varepsilon) + \nabla_x p^\varepsilon \\ &= \rho_p \int_{u_p} \left(\frac{D \frac{4}{3} \pi r_1^3}{r_1^2} f_1 + \frac{D \frac{4}{3} \pi \varepsilon^3 r_1^3}{\varepsilon^2 r_1^2} f_2 \right) (u_p - u^\varepsilon) du_p, \\ \partial_t f_1 + u_p \cdot \nabla_x f_1 + \nabla_{u_p} \cdot (f_1 D \frac{u^\varepsilon - u_p}{r_1^2}) &= -F f_1, \\ \partial_t f_2 + u_p \cdot \nabla_x f_2 + \nabla_{u_p} \cdot (f_2 D \frac{u^\varepsilon - u_p}{\varepsilon^2 r_1^2}) &= F f_2 \left(\frac{1}{\varepsilon} \right)^3, \\ p &= P(\rho^\varepsilon). \end{aligned}$$

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Kinetic energy : limit when $\varepsilon \rightarrow 0$ (1)



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$$\partial_t \left(\left(\frac{1}{2} \rho^\varepsilon (u^\varepsilon)^2 \right) + \frac{c}{\gamma - 1} (\rho^\varepsilon)^\gamma \right) + \nabla_x \cdot \left(\frac{1}{2} \rho^\varepsilon (u^\varepsilon)^2 u^\varepsilon \right) = \rho_p u^\varepsilon \left[\int_{u_p} f_1 \frac{D^4 \pi r_1^3}{r_1^2} (u_p - u^\varepsilon) du_p + \int_{u_p} f_2 \frac{D^4 \pi \varepsilon^3 r_1^3}{\varepsilon^2 r_1^2} (u_p - u^\varepsilon) du_p \right],$$

$$\partial_t \int_{u_p} f_1 \rho_p \frac{u_p^2}{2} du_p + \nabla_x \cdot \left(\int_{u_p} f_1 \rho_p \frac{u_p^2}{2} u_p du_p \right) = \int_{u_p} f_1 \frac{D}{r_1^2} (u_p - u^\varepsilon) \cdot du_p - \int_{u_p} F f_1 \rho_p \frac{u_p^2}{2},$$

Kinetic energy : limit when $\varepsilon \rightarrow 0$ (2)



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$$\partial_t \int_{u_p} f_2 \rho_p \frac{u_p^2}{2} du_p + \nabla_x \cdot \left(\int_{u_p} f_2 \rho_p \frac{u_p^2}{2} u_p du_p \right) = \int_{u_p} f_2 \frac{D}{\varepsilon^2 r_2^2} (u_p - u^\varepsilon) \cdot du_p + \frac{1}{\varepsilon^3} \int_{u_p} F f_1 \rho_p \frac{u_p^2}{2}.$$

Kinetic energy : limit when $\varepsilon \rightarrow 0$ (3)



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$$\begin{aligned} & \partial_t \left(\left(\frac{1}{2} \rho^\varepsilon (u^\varepsilon)^2 \right) + \frac{c}{\gamma - 1} (\rho^\varepsilon)^\gamma \right) + \nabla_x \cdot \left(\frac{1}{2} \rho^\varepsilon (u^\varepsilon)^2 u^\varepsilon \right) + \\ & \partial_t \int_{u_p} f_1 \frac{4}{3} \pi \rho_p r_1^3 \frac{u_p^2}{2} du_p + \nabla_x \cdot \int_{u_p} f_1 \frac{4}{3} \pi \rho_p r_1^3 \frac{u_p^2}{2} u_p du_p + \\ & \partial_t \int_{u_p} f_2 \frac{4}{3} \pi \rho_p \varepsilon^3 r_1^3 \frac{u_p^2}{2} du_p + \nabla_x \cdot \int_{u_p} f_2 \frac{4}{3} \pi \rho_p \varepsilon^3 r_1^3 \frac{u_p^2}{2} u_p du_p = \\ & -\rho_p \left[\int_{u_p} f_1 \frac{D^4 \pi r_1^3}{r_1^2} (u_p - u^\varepsilon)^2 du_p + \int_{u_p} f_2 \frac{D^4 \pi \varepsilon^3 r_1^3}{\varepsilon^2 r_1^2} (u_p - u^\varepsilon)^2 du_p \right]. \end{aligned}$$

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By letting tend ε to zero (at a constant mass for small droplets *i.e.*

$\int_{u_p} f_2 \frac{4}{3} \pi \rho_p \varepsilon^3 r_1^3 du_p = cte$), we must have $\int_{u_p} f_2 \frac{D^4 \pi r_1^3}{r_1^2} (u_p - u^\varepsilon)^2 du_p = O(\varepsilon^2)$, in

order to keep kinetic energy positive, so that f_2 is a Dirac mass in velocity (the gas velocity).

New set of equations



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Let's $\tilde{\rho}(t, x)$ be defined by $\int_{u_p} \frac{4}{3} \rho_p \varepsilon^3 r_1^3 du_p$ a partial droplet density, then we get

the following system :

$$\partial_t(\rho) + \nabla_x \cdot (\rho u) = 0,$$

$$\partial_t(\tilde{\rho}) + \nabla_x \cdot (\tilde{\rho} u) = \rho_p \frac{4}{3} \pi r_1^3 \int_{u_p} F f_1 du_p,$$

$$\partial_t((\rho + \tilde{\rho})u) + \nabla_x \cdot ((\rho + \tilde{\rho})u \otimes u) + \nabla_x p = -\rho_p \int_{u_p} \left(\frac{D \frac{4}{3} \pi r_1^3}{r_1^2} f_1 \right) (u - u_p) du_p,$$

$$+ \int_{u_p} \left(F \frac{4}{3} \rho_p \pi r_1^3 f_1 \right) u_p du_p$$

$$\partial_t f_1 + u_p \cdot \nabla_x f_1 + \nabla_{u_p} \cdot \left(f_1 D \frac{u - u_p}{r_1^2} \right) = -F f_1,$$

$$p = P(\rho).$$

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Let's define $\rho_m = \rho + \tilde{\rho}$, the gas mass concentration $c^+ = \frac{\rho}{\rho_m}$ and the small droplet concentration $c^- = \frac{\tilde{\rho}}{\rho_m}$. We get :

$$\partial_t(\rho_m c^+) + \nabla_x \cdot (\rho_m c^+ u) = 0,$$

$$\partial_t(\rho_m c^-) + \nabla_x \cdot (\rho_m c^- u) = \rho_p \frac{4}{3} \pi r_1^3 \int_{u_p} F f_1 du_p,$$

$$\partial_t(\rho_m u) + \nabla_x \cdot (\rho_m u \otimes u) + \nabla_x p = -\rho_p \int_{u_p} \left(\frac{D \frac{4}{3} \pi r_1^3}{r_1^2} f_1 (u - u_p) \right) du_p de_p dr$$

$$+ \int_{u_p} \left(F \frac{4}{3} \rho_p \pi r_1^3 f_1 \right) u_p du_p$$

$$\partial_t f_1 + u_p \cdot \nabla_x f_1 + \nabla_{u_p} \cdot (f_1 D \frac{u - u_p}{r_1^2}) = -F f_1,$$

$$c^+ + c^- = 1,$$

$$p = P(\rho).$$

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Let's define :

$$\beta = \int_{u_p, e_p, r} \frac{4}{3} \pi r^3 f_2 du_p de_p dr,$$

as the volume fraction occupied by small droplets at equilibrium with the gas.

$$\partial_t(\alpha\rho) + \nabla_x \cdot (\alpha\rho u) = 0,$$

$$\partial_t(\beta\rho_p) + \nabla_x \cdot (\beta\rho_p u) = \int_{u_p, e_p, r} F f_1 \rho_p \frac{4}{3} \pi r_1^3 du_p de_p dr,$$

$$\partial_t((\alpha\rho + \beta\rho_p)u) + \nabla_x \cdot ((\alpha\rho + \beta\rho_p)u \otimes u) + \alpha_m \nabla_x p =$$

$$\int_{u_p, e_p, r} m_1 \frac{D}{r_1^2} (u_p - u) f_1 du_p de_p dr + \int_{u_p, e_p, r} F f_1 \rho_p \frac{4}{3} \pi r_1^3 u_p du_p de_p dr,$$

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$$\begin{aligned} & \partial_t(\alpha\rho e + \beta\rho_p E_p) + \nabla_x \cdot ((\alpha\rho e + \beta\rho_p E_p)u) + p[\partial_t\alpha_m + \nabla_x \cdot (\alpha_m u)] \\ &= \int_{u_p, e_p, r} \left[(m_p \Gamma_p + \frac{m_p}{\rho_p} \nabla_x p) \cdot (u - u_p) - 4\pi r \lambda Nu (T - T_p) \right] f_1 du_p de_p dr \\ &+ \rho_p \frac{4}{3} \pi r_1^3 \int_{u_p, e_p, r} F f_1 e_p du_p de_p dr + p \frac{4}{3} \pi r_1^3 \int_{u_p, e_p, r} F f_1 du_p de_p dr, \\ \alpha_m &= 1 - \int_{u_p, e_p, r} \frac{4}{3} \pi r^3 f_1 du_p de_p dr = 1 - \int_{u_p, e_p, r} \frac{m_p}{\rho_p} f_1 du_p de_p dr, \\ \partial_t f_1 + u_p \cdot \nabla_x f_1 + \nabla_{u_p} \cdot (f_1 \Gamma_1) + \partial_{e_p} (f_1 \phi_1) &= -F f_1 + Q_1(f_1, f_1), \\ \Gamma_1 &= -\frac{D}{r_1^2} (u_p - u) - \frac{m_1}{\rho_p} \nabla_x p, \\ m_1 \phi_1 &= 4\pi r_1 \lambda Nu(r_1) (T - T_p), \\ p &= P(\rho, T), \\ E_p &= E(T), \\ e_p &= E(T_p). \end{aligned}$$

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Mix model for thick sprays



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$$\partial_t(\alpha_m \rho_m c^+) + \nabla_x \cdot (\alpha_m (\rho_m c^+ u)) = 0,$$

$$\partial_t(\alpha_m \rho_m c^-) + \nabla_x \cdot (\alpha_m \rho_m c^- u) = \rho_p \frac{4}{3} \pi r_1^3 \int_{u_p} F f_1 (u_p - u) du_p de_p dr,$$

$$\partial_t(\alpha_m \rho_m u) + \nabla_x \cdot (\alpha_m \rho_m u \otimes u) + \alpha_m \nabla_x p =$$

$$\int_{u_p, e_p, r} m_1 \frac{D}{r_1^2} (u_p - u) f du_p de_p dr + \int_{u_p, e_p, r} F f_1 \rho_p \frac{4}{3} \pi r_1^3 u_p du_p de_p dr,$$

$$\partial_t \alpha_m \rho_m e_m + \nabla_x \cdot (\alpha_m \rho_m e_m u) + p [\partial_t \alpha_m + \nabla_x \cdot (\alpha_m u)]$$

$$= \int_{u_p, e_p, r} \left[(m_p \Gamma_p + \frac{m_p}{\rho_p} \nabla_x p) \cdot (u - u_p) - 4\pi r \lambda Nu (T - T_p) \right] f_1 du_p de_p dr$$

$$+ \int_{u_p, e_p, r} F \rho_p \frac{4}{3} \pi r_1^3 f_1 e_p du_p de_p dr + p \int_{u_p, e_p, r} F \frac{4}{3} \pi r_1^3 f_1 du_p de_p dr,$$

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$$\alpha_m = 1 - \int_{u_p, e_p, r} \frac{4}{3} \pi r^3 f_1 du_p de_p dr = 1 - \int_{u_p, e_p, r} \frac{m_p}{\rho_p} f_1 du_p de_p dr,$$

$$\partial_t f_1 + u_p \cdot \nabla_x f_1 + \nabla_{u_p} \cdot (f_1 \Gamma_1) + \partial_{e_p} (f_1 \phi_1) = -F f_1 + Q_1(f_1, f_1),$$

$$\Gamma_1 = -\frac{D}{r_1^2} (u_p - u) - \frac{m_1}{\rho_p} \nabla_x p,$$

$$m_1 \phi_1 = 4\pi r_1 \lambda Nu(r_1) (T - T_p),$$

$$p = P(\rho, T),$$

$$E_p = E(T),$$

$$e_p = E(T_p).$$

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- The model is obtained for two sizes of droplet,
- Generally a transition radius r_t is defined because the distribution in droplet sizes is continuous : droplets of radii larger than r_t are dealt with the Boltzmann equation whereas others are put in the mix,
- Other criteria can be added : for instance in combustion at a given temperature droplets vaporize and can be put in the mix.

We present a case of interactions between two jets of metallic droplets moving in the air with the following properties :

- the initial droplet radius is $6\mu m$,
- the jet coming from the right is two times heavier than the one from the left,
- the jet coming from the right comes faster ($-2500m.s^{-1}$) than the one from the left ($1800m.s^{-1}$),
- the gas is at rest,
- the gas volume fraction is set between 0.95 et 1 in each jet.

We test different transition radii for droplets and compare results

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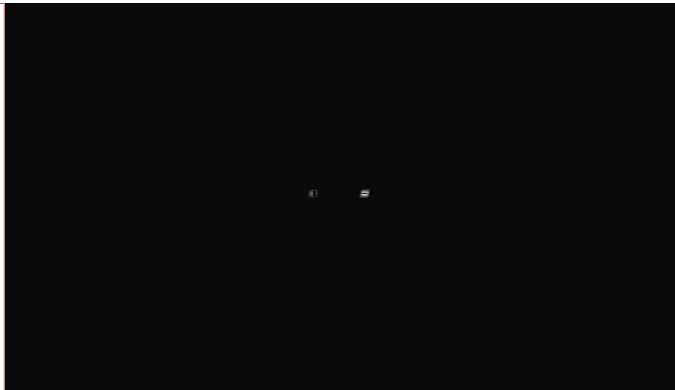
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PRESSION

1.165e+05
1.160e+05
1.155e+05
1.150e+05
1.145e+05
1.141e+05
1.136e+05
1.131e+05
1.126e+05
1.121e+05
1.116e+05
1.111e+05
1.106e+05
1.101e+05
1.096e+05
1.091e+05
1.086e+05
1.082e+05
1.077e+05
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1.067e+05
1.062e+05
1.057e+05
1.052e+05
1.047e+05
1.042e+05
1.037e+05
1.032e+05
1.027e+05
1.023e+05
1.018e+05
1.013e+05
1.008e+05
1.003e+05
9.979e+04
9.930e+04



First Test : results (2)



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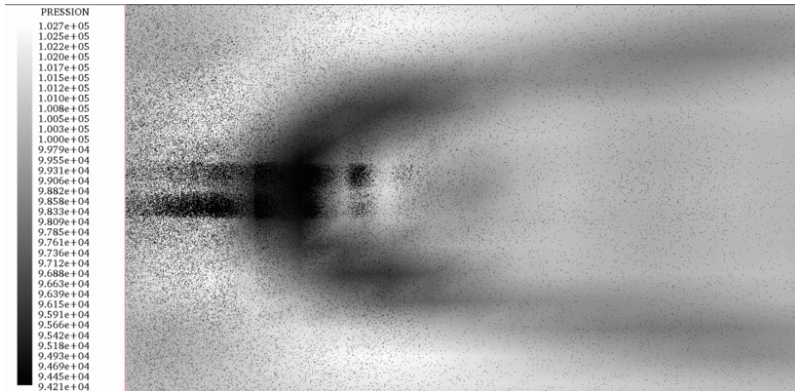
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Pressure for different radii of transition r_t
($r_t = 0$ (without mix), 20nm, 80nm)

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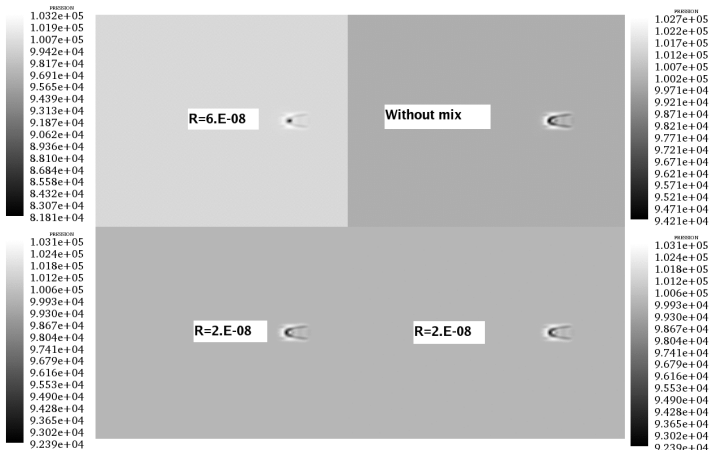
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First Test : results (4)



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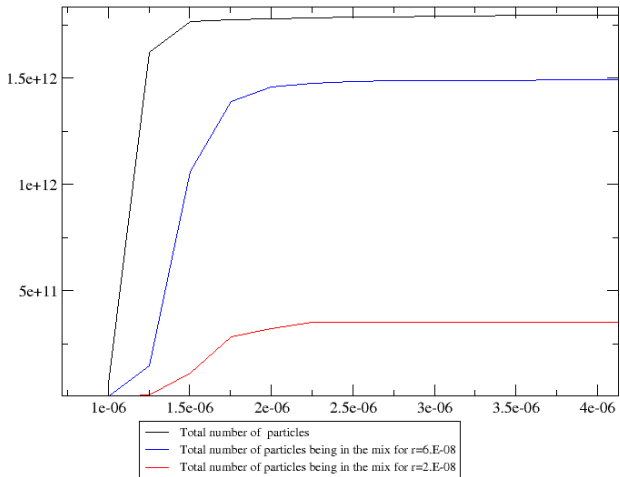
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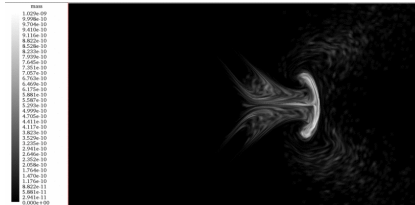
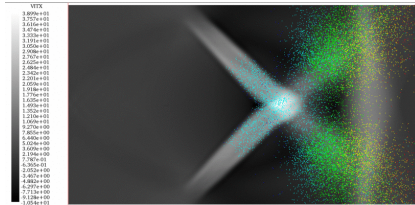
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Location of droplets mass for the mix model on the top (strict criterium for radius of transition) and droplet model on the bottom

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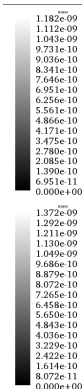
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Location of droplets mass for the mix model on the top (large criterium for radius of transition) and droplet model on the bottom

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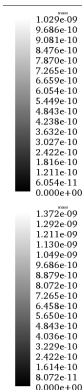
Hydrodynamic limit

Test case

Colliding jets

Crossing jets

Conclusions





Introduction

A new model

Context
Hydrodynamic
limit
Hydrodynamic
limit

Test case

Colliding jets
Crossing jets

Conclusions

- The model can deal with various applications.
- More computations are needed to understand the transition towards mix.
- More realistic data should be used for tests.