# 293T Cell Culture

### Materials

#### 293T medium

name	dose	ratio
DMEM	$500 \mathrm{mL}$	-
FBS	$50~\mathrm{mL}$	1:10
p/s	5  mL	1:100

### Methods

#### Culture (from stock)

- defrost 293T cell stock in 37°C water bath
- PBS wash x1
  - -centrifuge 1200 rpm x5min
  - aspiration
- $\bullet\,$  ressuspend with 10 mL 293T medium and apply into T25
- after confirmation with a microscope, transfer to the incubator
- culture for 5 days (approx.)

#### Immunofluorescense

#### day1

- prepare adherent 24-well dish for observation
- aspirate fibro. in the wells
- apply 293T medium 500uL each
- after pipetting for several times, transfer culturing suspension 200uL each from T25 into wells
- 37°C incubation o/n

### day2

- confirm with microscope that 293T firmly adhere to the floor
- aspiration
- process with PFA x45min (approx.) in RT
- PBS wash x4
- process with 0.3% TX x15min (approx.) in RT
- PBS wash x2
- apply Primary Ab Mix
  - e.g.) Primary Ab Mix (for 1 well)

name	dose	ratio
Blocking One	$500~\mathrm{uL}$	-
anti-xxx Mouse mAb	$0.5~\mathrm{uL}$	1:1000
anti-yyy Rabbit mAb	$0.5~\mathrm{uL}$	1:1000

• 4°C o/n with shading

## day3

- PBS wash x3
- apply Secondary Ab Mix
  - e.g.) Secondary Ab Mix (for 1 well)

dose	ratio
$500~\mathrm{uL}$	-
$0.5~\mathrm{uL}$	1:1000
$0.5~\mathrm{uL}$	1:1000
	500 uL 0.5 uL

- shade in RT x1hr (approx.)
- PBS wash x2
- $\bullet$  apply 1:1000 hochest

name	dose	ratio
PBS	500 uL	_
hochest	$0.5~\mathrm{uL}$	1:1000

- shade in RT x5min (approx.)
- PBS wash x2
- wait until wells get dry
- observe with KEYENCE