For detection of intranuclear inclusion bodies, primary CEL cells were grown on sterile coverslips [Adair, 1978]. Briefly, the growth medium was aspirated out from confluent monolayer and washed twice with sterile medium free serum prior to inoculation of 0.1mL viral supernatant. Culture was added with maintenance medium, DMEM and 2% FBS and kept for 48 hours incubation, harvested and fixed in 10% buffered formalin for 5 minutes and stained with haematoxylin and eosin (HE).
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Protocol status: Working
We use this protocol and it's working

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