Jan 29, 2019 Version 2

Lymphocyte proliferation in poultry species V.2

PLOS One

DOI
dx.doi.org/10.17504/protocols.io.xjpfkmn

Ahmed O.Abass[1], Gamal Mehaisen[1]

[1]Cairo University

Gamal Mehaisen
Cairo University

OPEN ACCESS

DOI: https://dx.doi.org/10.17504/protocols.io.xjpfkmn

External link: https://doi.org/10.1371/journal.pone.0214839

Protocol Citation: Ahmed O.Abass, Gamal Mehaisen 2019. Lymphocyte proliferation in poultry species. **protocols.io https://dx.doi.org/10.17504/protocols.io.xjpfkmn**

Manuscript citation:
Mehaisen GMK, Eshak MG, Elkaiaty AM, Atta A-RMM, Mashaly MM, Abass AO (2017) Comprehensive growth performance, immune function, plasma biochemistry, gene expressions and cell death morphology responses to a daily corticosterone injection course in broiler chickens. PLoS ONE 12(2): e0172684. doi:10.1371/journal.pone.0172684

Protocol status: Working
We use this protocol and it's working

Created: January 29, 2019

Last Modified: January 29, 2019

Protocol Integer ID: 19791

Keywords: lymphocyte proliferation in poultry species, lymphocyte proliferation, poultry species

Troubleshooting

1 The heparinized blood samples were added to separation medium Histopaque®-1077 (cat# 10771, Sigma, USA).

2 Samples were centrifuged at 1030 xg for 20 min at 4°C.

3 Peripheral blood mononuclear cells (PBMCs) were isolated and washed twice with RPMI-1640 (Invitrogen Corp., Grand Island, NY, USA) and then re-suspended in 2 ml of RPMI-1640 complete culture medium.

4 The viable lymphocytes were detected using Trypan Blue dye and plated in triplicate wells (96-well plate) at 1×10^6 cells per well.

5 A 50 µl of either Concanavalin-A (Con-A, 45 µg/ml, cat# C5275, Sigma, USA) or Lipopolysaccharide (LPS, 10 µg/ml, cat# L4391, Sigma, USA) was added to selected wells to induce the proliferation of T lymphocyte and B lymphocyte, respectively.

6 Control wells received 50 µl of RPMI-1640 medium.

7 Cells were then incubated for 48 h at 42 °C with 5 % CO_2.

8 After incubation, 15 µl of 3-[4,5-dimethylthiazol]-2,5-diphenyltetrazolium bromide (MTT, 5 mg/ml, cat# M2128 , Sigma, USA) was added to each well and the cells were incubated for another 4 h.

9 Subsequently, 100 µl of 10% sodium dodecyl sulfate dissolved in 0.04 M HCl solution was added to each well to lyse the cells and solubilize the MTT crystals.

10 Finally, the absorbance at 570 nm was recorded using an automated ELISA microplate reader (ChroMate® Microplate Reader-4300, Awareness Technology Inc., Palm City, FL, USA).

11 Stimulating index (SI) for either T or B cells was calculated as follows: SI = $OD570_{(stimulated\ cells)}$ / $OD570_{(unstimulated\ cells)}$.