CONTENTS

	Page
INTRODUCTION	1
REVIEW OF LITERATURE	3
1- Nomenclature and Synonyms	3
2- History and geographic distribution of CPD	3
2.1 CPD in the World	3
2.2 CPD in Egypt	5
3- Classification of C PD virus	6
4- Physical properties of CPD virus	7
5 Chemical composition of CPD virus	8
5.1 envelope	8
5.2 genome organization	8
5.3 proteins and function	9
6- Strains and isolates of <i>CPD</i> virus	10
7- Effect of Physico-chemical agents	12
8- Antigenic properties	14
9- CPD virus and replication	15
9.1 Virus adsorption and entry into the cell	15
9.2 Virus Uncoating	16
9.3 Virus Replication	16
10- Haemagglutination properties	18

11- Laboratory diagnosis	19
11.1 Samples	19
11.2 CPD virus isolation	19
11.2.1 In fertile eggs	19
11.2.2 On tissue culture	20
11.3 CPD virus Serological identification	22
11.3.1 Complement fixation test (CFT)	22
11.3.2 Fluorescent antibody technique (FAT)	23
11.3.3 Agar gel precipitation test (AGPT)	23
11.3.4 Haemagglutination inhibition test (HI)	23
11.3.5 Enzyme linked immunosorbant assay (ELISA).	24
11.4 CPD virus Non serological identification	24
11.4.1 Electron microscope (E.M)	24
11.4.2 Polymerase chain reaction (PCR)	26
12- Immunity and resistance to CPD viral infection	27
13- Interferon production	30
14- immunization or vaccination	31
MATERIAL AND METHODS	35
1-materials	35
1.1 Virus strain	35
1.2 Samples	35
1.3 Glycerol buffer – saline	36
1.4 antibiotic solution	36

1.5 Fertile eggs
1.6 Hyper immune sera
1.7 Anti species (Anti sheep IgG conjugate)36
1.8 Nobel Agar36
1.9 Phosphate buffer saline (PBS)37
1.10 reagents used for electron microscopic examination37
1.11 reagents used for polymerase chain reaction39
1.12. Red blood cells for Haemagglutination test40
2- methods
2.1 preparation of suspected CPD virus samples41
2.2. isolation of suspected CPD virus in SPF – ECE41
2.3 preparation of viral antigen from inoculated fertile eggs41
2.4 Titration of the virus isolates in SPF- ECE42
2.5 serological identification of suspected CPD virus isolate42
2.5.1 Agar gel precipitation test42
2.5.2 Indirect fluorescent antibody technique43
2.6 Non serological identification of suspected CPD virus isolate.44
2.6.1 Electron microscope44
2.6.2 Polymerase chain reaction45
2.7 Haemagglutination properties47
2.7.1 Haemagglutination test
2.7.2 Haemagglutination inhibition test47
RESULTS48

DISCUSSION	70
SUMMARY	74
REFERENCES	76
ARABIC SUMMARY	-

List of Abbreviation

CPD : Contagious pustular dermatitis.

ECEs : Emberyonated chicken eggs.

AGPT : Agar Gel Precipitation Test.

IFAT : *Indirect fluorescent antibody technique*.

PCR : Polymerase chain reaction.

NZ2 strain : New Zealand type 2 strain.

DNA : Deoxyribonucleic Acid.

FMD : Foot and mouth disease.

VEGF : *Vascular endothelial growth factor.*

ITR : Inverted terminal repeats.

dsDNA : Double stranded deoxyribonucleic Acid.

RNA : Ribonucleic Acid.

UV light : Ultra violet light.

CAM : Coriallantoic membrane.

pfu : Pock forming units.

CPE : Cytopathic Effect.

MDOK : Marbin-Darby ovine kidney.

FAT : Fluorescent antibody technique.

AGID : Agar gel immunodifusion.

E.M. : Electron microscope.

ELIZA : Enzyme-linked immunosorbant assay

 $TCID_{50}$: Tissue culture infective dose 50

OVIL-10 : Ovine virus interleukin-10

CFT : Complement fixation test.

MHC : Major Histocompatibility Complex.

GM-CSF: Granulocyte-macrophage colony- stimulating

factor.

CTL : Cytotoxic T cells.

IL-10 : Interleukin- 10.

INF-Y : Interferon-Y.

SPF : Specific pathogen free.

PBS : Phosphate buffer saline.

EID₅₀ : Embryo lethal dose 50.

HA : Haemagglutination.

HI : Haemagglutination inhibition.

RBCs : Red blood corpuscles.

nm : nanometer.

G+C: Guanine + cytosine.

Kbp : kilo base pair.

NZ7 : New Zealand type 7 strain.

RNA : Ribonucleic Acid.

mRNA : messenger ribonucleic Acid.

INF : interferon.

SPF : specific pathogen free.

Mg : milligram.

Iu : international unite.UK : united kingdom.TE : tris EDTA buffer.

EDTA : ethylene diamine tetra acetic acid.

Mg cl2 : magnesium chloride.

SDS : sodium dedocyle sulphate.

DMAE : Dimethyl amino ethanol.

NSA : Nonenyl Succinic anhydride Madified.

DER-736 : epoxy resin.

ERL-420 : vinyl cyclohexene dioxide.

7. SUMMARY.

This research aimed to isolate and recognize CPD virus in sheep and goat herds and the research investigate the following:

- 1- The affected sheep and goats show clinical signs like nodules, pustules, scars and crusts which are obviously appeared on lips, gums, mouth commisures and eyelids.
- 2- Samples from these affected animals were collected in order to isolate and recognize the CPD virus with normal serological and non serological means.
- 3- Preliminary identification of the CPD virus antigen using electron microscopic examination for skin samples which give characteristic appearance of the virus (ovoid in shape, with rounded ends and characteristic ball of wool appearance).
- 4- Trials for isolation of CPD virus:

The prepared samples from skin lesions were inoculated in emberyonated chicken eggs by the CAM route which showed thickening, oedema and hemorrhage of CAMs tissue.

- 5- Titration of CPD virus isolates revealed that the titer of collected sheep isolates was 5 \log_{10} EID₅₀ / ml while the titer of collected goat's isolates was $4\log_{10}$ EID₅₀ / ml.
- 6- The inoculated CAMs were harvested and AGPT and IFAT tests were done which gave positive results for viral antigen.
- 7- PCR test was done using known primer on harvested samples from CAM and tissue samples. Amplification and running of characteristic fragments of CPD viral DNA on Ethidium Bromide stained Agarose Gel then staining of DNA gave

typical results in many segments on DNA fragment which indicate that the given sample contain CPD viral antigen.

- 8- Characterization and identification of CPD virus revealed that:
 - The isolated virus has no haemagglutinating property to avian RBCs of (chicken, duck, goose, turkey and pigeon RBCs).
 - The isolated virus has no haemagglutinating property to mammalian RBCs of (sheep, goat, donkey, horse, cattle and buffalo RBCs).
 - The isolated virus has haemagglutinating property to rabbit RBCs
- The haemagglutination inhibition was applied and gave positive results in form of button shape pattern.

From the previous we can conclude that the isolated virus from affected animals (sheep and goats) assumed to be CPD virus as confirmed by AGPT, IFAT and PCR.