

Data Sheet

DESCRIPTION:	CHO-pEE14/tpa/UG37-08gp140REKR
REPOSITORY REFERENCE:	ARP5003
DETAILS OF PRODUCTION:	Stable CHO cell line secreting A-clade oligomeric gp140 (gp120 + external domain of gp41) into ambient medium. Cleavage site between gp120/41 retained. N-terminal signal sequence of gp replaced by that of tissue plasminogen activator (tpa) – cleaved from gp as it is secreted from cell. Gp140 N-termini starts E ₂₈ N ₂₉ L ₃₀ W ₃₁ V ₃₂ ; C-termini ends with 2F5 Mab epitope ELDKWAS. Full details of plasmid available on request
SPECIAL INFORMATION:	Derived from functional A clade gp160 clone p92UG037-08 (Gao et al., AIDS Res.Hum.Retro. 11, 1359-1367 (1994));viral isolate is 92/UG/037 (Molecular clone Accession no.U51190)
GROWTH MEDIA:	Glutamine-free DME supplemented with GS salts (both JRH Biosciences), 5% dialysed foetal calf serum and 200µM L-methionine sulphoximine (MSX). MSX selection must be applied throughout culture. Full culture details on request
PURIFICATION:	Secreted gp140 can be immunoaffinity purified using D7324 (Aalto Bioreagents, Rathfarnham, Dublin, Ireland). Full details available on request
STORAGE:	Liquid Nitrogen

SOURCE:

Dr S.A. Jeffs, NIBSC

REFERENCES:

Jeffs et al (2002) – in press

ACKNOWLEDGEMENTS:

Publications should acknowledge the donor of the reagent and the Programme EVA Centre for AIDS Reagents. Suggested wording can be found on our website at <http://www.nibsc.ac.uk/spotlight/aidsreagent/index.html> in the “Acknowledgements” section.

Please also ensure that you send us a copy of any papers resulting from work using reagents acquired through CFAR (this can be electronically or as a paper copy)