

EliteGroTM-Adv.

XENOGENEIC-FREE Cell Culture Supplement





Efficient

- Only 5% EliteGro[™] and EliteGro[™]-Adv. replace up to 20% FBS
- Faster cell population doublings (<30 hrs).
- Low cell seeding density.



EPG-050 / EPG-500 50ml / 500ml



EliteGro[™]-Adv.

EPA-050 / EPA-500 50ml / 500ml No heparin required



- U.S. FDA CDER **DMF 032759**.
- U.S. FDA CBER MF 27525.
- GMP-compliant facility in the USA.
- Material from U.S. FDA-registered and AABB accredited supplier and intending use in human transfusion.
- No xenogeneic component;
 No anticoagulant added.





Reliable

- Stable and consistent supply.
- Minimal variation from lot to lot.
- Professional management team possesses 10-year experience support.

EliteGro[™]-Adv. (GMP grade)

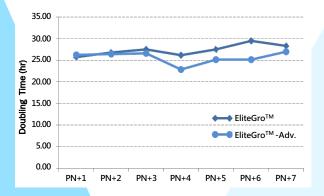
EPAGMP-050 / EPAGMP-500 50ml / 500ml No heparin required



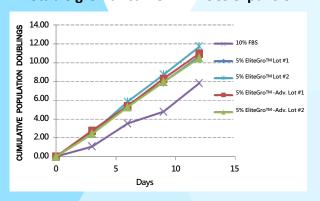
EFFICIENT, SAFE, RELIABLE

EliteGroTM and EliteGroTM-Adv. provide abundant human source growth factors and work as effective replacement for FBS (fetal bovine serum) to support cell expansion. EliteGroTM and EliteGroTM-Adv. are animal serum-free, xenogeneic-free cell culture supplements for research or commercial development of cell therapies. Successfully support cell proliferation of multiple types of mesenchymal stem cells without morphology or characterization change.

Less doubling time and stable growth curve in hMSCs expansion

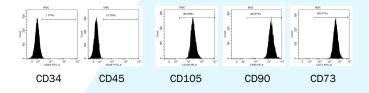


Stable growth curve in hMSCs expansion

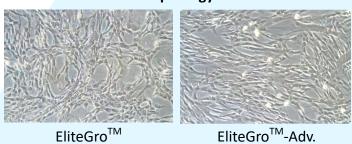


Characterization of hMSCs

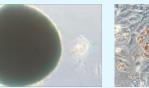
In 5% EliteGro[™] and EliteGro[™]-Adv. medium, hMSCs keep positive for the hMSCs markers (CD90, CD73 and CD105) and negative for CD34 and CD45.



Human hMSCs morphology



Differentiation ability of hMSCs







Chondrocytes

Adipocytes

Osteocytes

Biomedical EliteCell Corp is a reliable team with more than 10 years extensive manufacture and market experiences in supplement products. Collaborating with FDA licensed blood center in the USA for raw material traceability and safety. GMP-compliant facility in Texas has integrated quality management system and follows FDA regulation 21 CFR Part 820.

