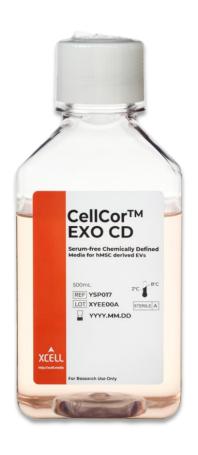


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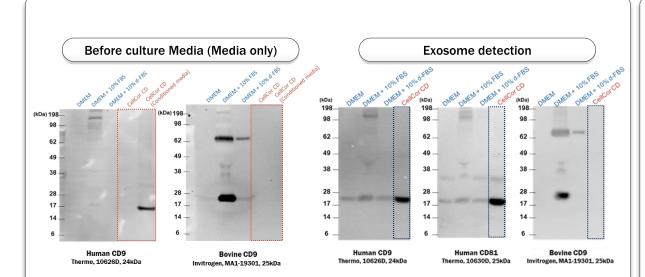
Scope	Mass production, research and drug development of exosomes
Superiority	Eliminate more than 95% of impuritiesConvenience for culture
Application	 Adipose derived MSC Bone marrow derived MSC Umbilical cord derived MSC Various origin MSC
Competitiveness	Stability of cellSafety, homogeneity



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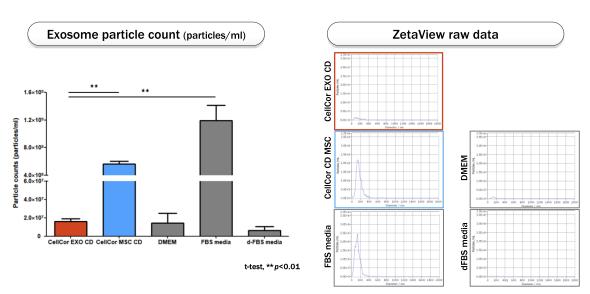
Comparison of culture test with FBS contained medium

Exosome specific marker



- Experimental Method: MSC were incubated in each media for 3 days, and then exosome was isolated and Western blot
 was performed.
- Measuring and analysis instrument: Tangential Flow Filtration (TFF, PALL) / iBrightTM CL1000 Imaging System (Invitrogen)
- * CellCor EXO CD did not contain bovine-derived exosomes, and showed the highest exosome production

Purity



- Experimental Method: Measure the number of particles in before culture medium using ZetaView.
- Measuring and analysis instrument: Zetaview (Particle metrix)
- CellCor EXO CD was confirmed to have the highest purity as it contained the lowest number of particles.

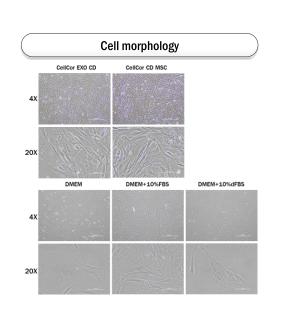


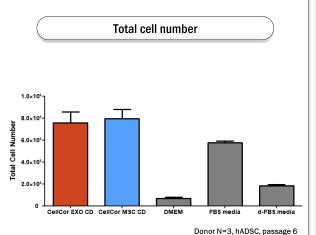
Comparison of culture test with FBS contained medium

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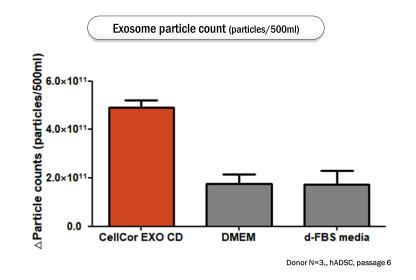
Cell Proliferation





- Experimental Method: MSC were seeded 1x10⁵cells on T25 flask, and cultured during 5 days. The number of MSCs were counted.
- Measuring and analyzing instrument: NucleoCounter® NC-250 (ChemoMetec)
- ❖ More cells were proliferated with CellCor EXO CD in the same culture environment.
- ❖ Proliferation was about 3.8 times better than d-FBS media.

Exosome production



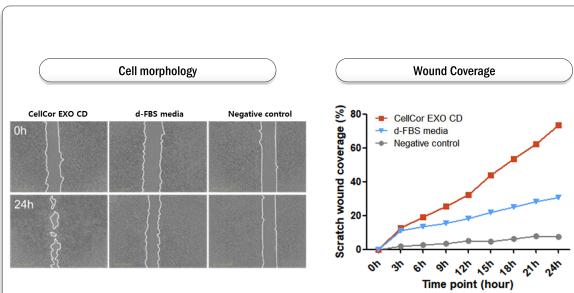
- Experimental Method: MSC were seeded 1x10⁵cells on T25 flask, and cultured during 5 days. The number of MSC-derived exosomes was counted.
- Measuring and analyzing instrument: NucleoCounter® NC-250 (ChemoMetec), Zetaview (Particle metrix)
- It was confirmed that the production of exosomes from CellCor EXO CD is superior to that of each medium.
- It was verified that the production volume is about 2.5 times (or more) and is a very efficient medium for mass production.



Comparison of culture test with FBS contained medium

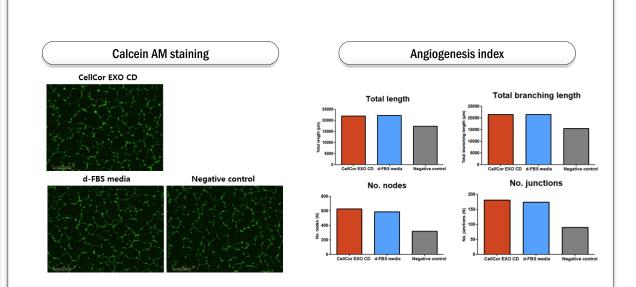
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Wound healing assay



- Experimental Method: HaCaT cell were cultured on 48well plate, after wounding the cells, the cells were treat with
 exosomes to evaluate the wound recovery ability.
- Measuring and analyzing instrument: Incucyte ZOOM software (Sartorius)
- ❖ As a result of comparing cell wound healing capabilities by treating MSC-derived exosome on CellCor EXO CD or each media, it was confirmed that exosomes secreted from CellCor EXO CD had excellent proliferation and regeneration effects of damaged cells.

Angiogenesis asssay



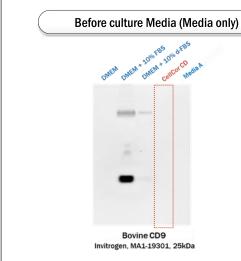
- Experimental Method: HUVEC cell culture on 24well plate with 1x10⁹ particles/ml of exosomes, and evaluate angiogenic index after 17h.
- Measuring and analyzing instrument: Incucyte ZOOM software (Sartorius), ImageJ
- **❖** The exosome produced using CellCor EXO CD showed a high angiogenic index.

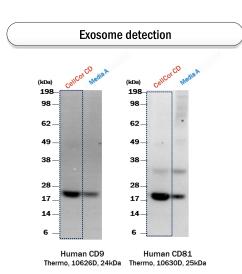


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Comparison of culture test with Commercial medium

Exosome Specific Marker

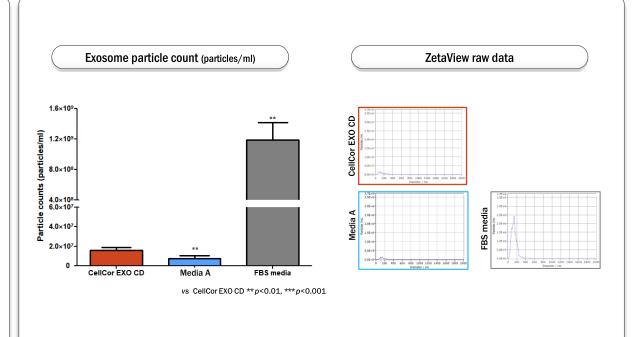




- Experimental Method: MSC were incubated in each media for 3 days, and then exosome was isolated and performed Western blot.
- Measuring and analyzing instrument: Tangential Flow Filtration (TFF, PALL) / iBright™ CL1000 Imaging System (Invitrogen)

CellCor EXO CD did not contain bovine-derived exosomes, and showed the highest exosome production

Purity



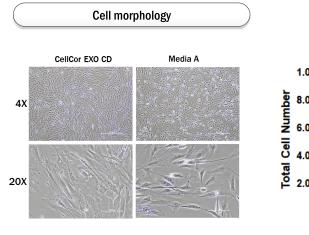
- Experimental Method: Measure the number of particles in before culture medium using ZetaView.
- Measuring and analyzing instrument: Zetaview (Particle metrix)
- CellCor EXO CD was confirmed to have the highest purity as it contained the lowest number of particles.

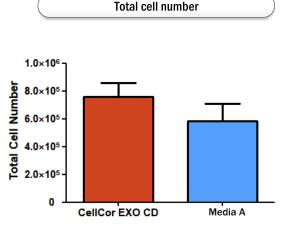


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Comparison of culture test with Commercial medium

Cell Proliferation

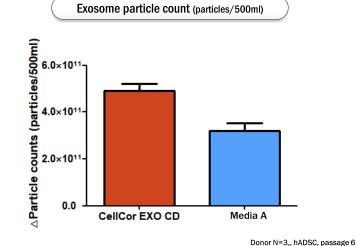




Donor N=3,, hADSC, passage 6

- Experimental Method: MSC were seeded 1x10⁵cells on T25 flask, and cultured during 5 days. The number of MSCs were counted.
- Measuring and analyzing instrument: NucleoCounter® NC-250 (ChemoMetec)
- ❖ More cells were proliferated with CellCor EXO CD in the same culture environment.
- ❖ Proliferation was about 1.3 times better than Commercial A media.

Exosome Production



- Experimental Method: MSC were seeded 1x10⁵cells on T25 flask, and cultured during 5 days. The number of MSC-derived exosomes was counted.
- Measuring and analyzing instrument: NucleoCounter® NC-250 (ChemoMetec), Zetaview (Particle metrix)
- It was verified that the production volume is about 1.6 times (or more) and is a very efficient medium for mass production.

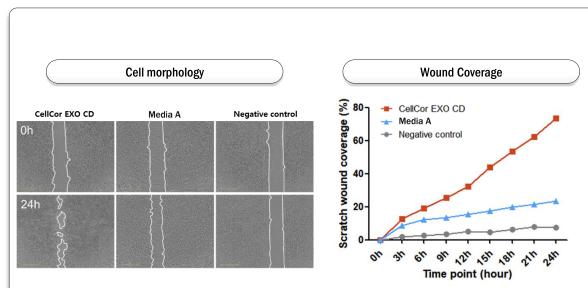


Comparison of culture test with Commercial medium

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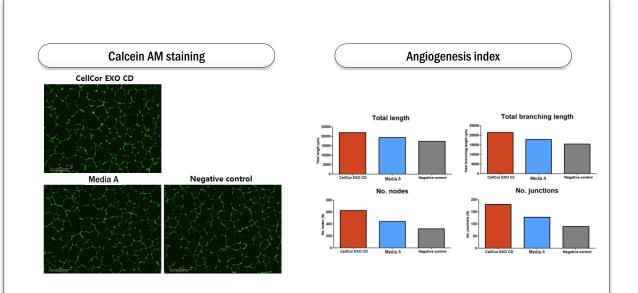
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Wound healing assay



- Experimental Method: HaCaT cell were cultured on 48well plate, after wounding the cells, the cells were treat with
 exosomes to evaluate the wound recovery ability.
- Measuring and analyzing instrument: Incucyte ZOOM software (Sartorius)
- As a result of comparing cell wound healing capabilities by treating MSC-derived exosome on CellCor EXO CD or each media, it was confirmed that exosomes secreted from CellCor EXO CD had excellent proliferation and regeneration effects of damaged cells.

Angiogenesis assay

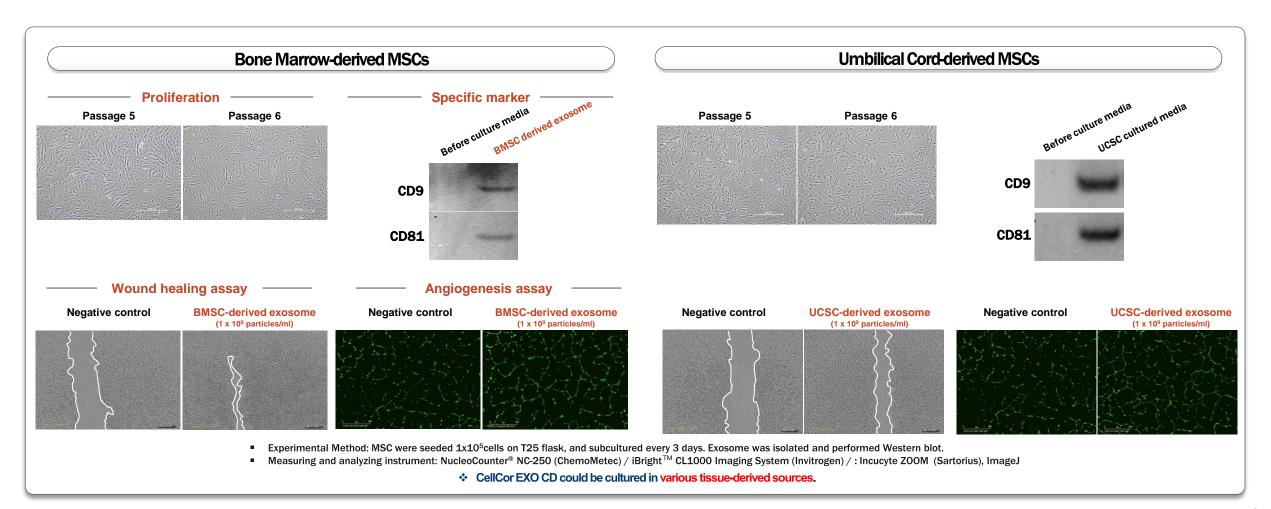


- Experimental Method: HUVEC cell culture on 24well plate with 1x10⁹ particles/ml of exosomes, and evaluate angiogenic index after 17h.
- Measuring and analyzing instrument: Incucyte ZOOM software (Sartorius), ImageJ
- The exosome produced using CellCor EXO CD showed a high angiogenic index.



Comparison of culture test with various tissue origin

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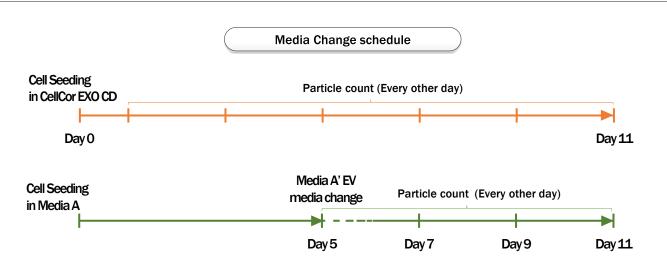


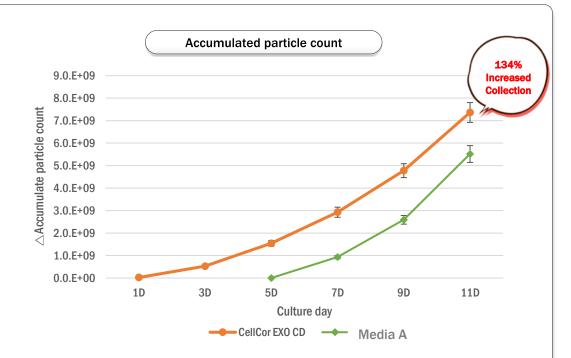


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Analytical Comparison Compared to Competing Media

Exosome particle count





- Method
 - 1) CellCor EXO CD : Seed ADSCs 1x10⁵ cells /25 T flask and culture with CellCor EXO CD without media change
 - 2) Media A: Seed ADSCs 1x10⁵ cells /25 T flask and culture with Media A'-MSC media for 5 days and change to Media A' and culture for 6 more days
- Analysis Method: Collect the conditioned medium in a 2-days period, isolate the exosomes, and conduct particle count using Zetaview
 Calculate the secreted amount of particles by subtracting the number of particles that existed in the media before culture
- Instrument Used : Zetaview (Particlematrix)
- CellCor EXO CD can be used for both expansion and collection of EVs during the period of 3 ~ 11 days without any media change

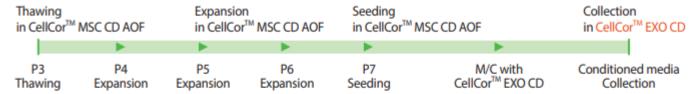


Enabling Flexible Research Design

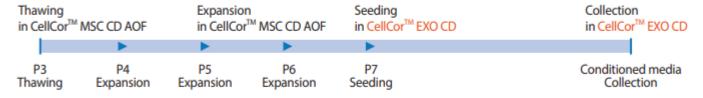
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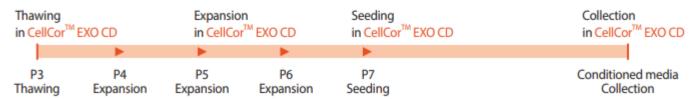
■ Media Change Process

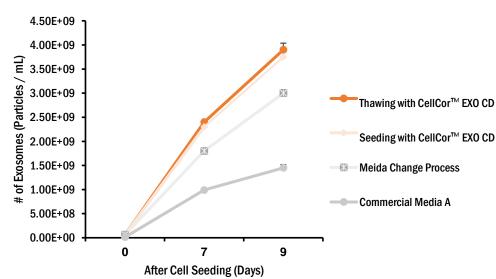


■ Seeding with CellCorTM EXO CD Process



■ Thawing with CellCorTM EXO CD Process





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