mRNA was confirmed by RT-PCR and protein by immunocytochemistry in the drug resistant clones. In order to study the effect of Nurr1 protein on the differentiation pattern of ES cells, one of the positive clones (MBNr24) was allowed to form embryoid body (EB) for 2 days and were induced to differentiate for another 4 days using RA (1 μ M) and AA (50 mM) (2-/4+ protocol) followed by selection in N2 medium for 10 or 20 days.

Results: After 10 days in N2 medium, cells immunoreactive to anti-GFAP, anti-TH, or anti-NF200 anti-bodies were 38.8%, 11%, and 20.5%, respectively. After 20 days in N2 medium, cells expressing GFAP, TH, or NF200 were 28%, 15% and 44.8%, respectively but approximately 9% of MB03 expressed TH protein when the cells were induced to differentiate using a similar prorocol.

Conclusions: These results suggest that ectopic expression of Nurr1 enhances generation of TH+ cells as well as neuronal cells when hES cells were differentiated by 2-/4+ protocol.

P-49 Improvement of Motor Behavior of Parkinson's Disease Animal Model by Nurr1-transfected Human Embryonic Stem Cells

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Background & Objectives: The purpose of this study is to evaluate an efficacy of in vitro differentiated human embryonic stem (hES, MB03) cells expressing Nurr1 in relief of symptomatic motor behavior of Parkinson's disease (PD) animal models.

Method: MB03 was genetically modified to express Nurr1 protein and was induced to differentiate according to 2-/4+ protocol using retinoic acid and ascorbic acid. The differentiation-induced cells were selected for 10 to 20 days thereafter in N2 medium. Upon selection, cells expressing GFAP, TH, or NF200 were 38.8%, 11%, and 20.5%, respectively. In order to examine therapeutic effects of the differentiated cells in PD animal model, rats were unilaterally lesioned by administration of 6-hydroxydopamine HCl (6-OHDA) into medial forebrain region (MFB, AP -4.4 mm, ML 1.2 mm, DV 7.8 mm with incision bar set at -2.4 mm), as a reference to bregma and the surface of the skull. Confirmation of successful lesion by apomorphine-induced rotational behavior, differentiated cells were transplanted into the striatum (AP 1.0, ML 3.5, DV -5.0; AP 0.6, ML 2.5, DV -4.5).

Results: Improvements of asymmetric motor behavior by the transplanta- tion were examined every two weeks after the surgery. In two weeks, numbers of rotation by the experimental rats were -14.8+33.9% (p<0.05) of the number before transplantation, however, the ratio increased slightly to 13.6+56.3% in six weeks. In contrast, the ratio of sham-grafted animals ranged from 112.3+8.5% to 139.2+28.9% during the examination.

Conclusions: This result suggests that MBNr24 cells differentiated in vitro survived at least for 6 weeks when grafted into brains of PD animal model, and that symptomatic motor behavior was improved.