## **CHAPTER V**

## **CONCLUSION**

Proteins from Sesbania grandiflora flowers, leguminous plant, were characterized by using electrophoresis, chromatography and mass spectrometry. Crude proteins from Sesbania grandiflora flowers were extracted and fractionation precipitated with ammonium sulfate. The amount of protein is higher when using the higher amount of ammonium sulfate, the most concentration was found in 60% precipitation and the lowest concentration was found in 20% precipitation. Biological activity tests of these proteins found that 60% and 90% crude protein show hemagglutination and α-glucosidase inhibitory activity. For 60% crude protein, 13.5 mg of SGF60 was purified in only one step by using DEAE-cellulose anion exchange chromatography. This protein show 31 HU/mg of specific hemagglutination activity with rabbit red blood cell and 82% inhibition of α-glucosidase inhibitory activity. Approximate molecular weight of SGF60 from SDS-PAGE electrophoresis is 40 kDa. To identify this protein, tandem mass spectrometry using ESI-Q-TOF has been use. Peptide sequencing shows that partial amino acid sequence of SGF60 matched with p27SJ, a novel protein inhibited HIV-1, from Hypericum perforatum and DING protein from Solanum tuberosum. For 90% crude protein characterization, 2.16 mg SGF90 can be purified in two steps. Gel filtration chromatography using Superdex-200 has been used in first step and DEAE-cellulose anion exchange chromatography was used in second step. Molecular weight of this protein is 63 kDa. To identify amino acid sequences of this protein, ESI-Q-TOF has been use. The database searching of peptide sequences show that partial amino acid sequence of SGF90 protein matched with the partial amino acid sequence of beta-glucosidase (At5g36890) and beta-glucosidase F8K4.3 protein from Arabidopsis thaliana. SGF90 can inhibit α-glucosidase enzyme (74%inhibition) and agglutinate rabbit erythrocyte at 3.8 HU/mg. In addition, the protein profile of Sesbania grandiflora from 2D-gel electrophoresis presents at least six proteins. These proteins were analyzed by peptide mass mapping using MALDI-TOF MS. From the search results, there are no signification match between the results and proteins in database.