

CHAPTER III

CONCLUSION

The acylation cyclization of a β -substituted allene using an acyl organocobalt reagent was not diastereoselective. Alternatively, the palladium catalyzed cyclization of β -hydroxy allene was found to proceed with 2:1 stereoselectivity. Such β -substituted allenes could therefore be precursors for amphogynine alkaloids. Future work should involve the use of alternative N-protecting groups, such as carbamates or novel sulfonamides.



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