1.0g sample of triglyceride was dissolved in 4ml solvent.

The solution then was placed in distillation flask and the residual material was collect by wash the beaker with 1 ml solvent.

25ml of 0.5M KOH was added.

Control was set up with additional 1:1 ethanol solvent for final volume that identical to the experimental solution

Reflux condenser was set up and being placed in boiling water bath for 30 minutes.

The flask was allowed to cool and 3 drop of indicator solution was added to both flask and being titrate with 0.5M HCl solution.

The weight of KOH used was calculated and saponification number was determined.

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R Bogdan

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