

Effectiveness of Ginger Jelly Candy to Reduce Intensity and Duration of Primary Dysmenorrhea in Adolescents

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Abstract. Dysmenorrhea is one of the most common gynecologic complaints experienced by teenagers and more than half of menstruating women. The adverse effects of dysmenorrhea affect the psychological and economy aspects. The purpose of this study was to analyze the differences decrease of intensity and duration of primary dysmenorrhea between adolescents who were given ginger jelly candies and given ibuprofen.

This research is an experimental Quasy research with nonequivalent pre testpost test control group design. Population in this research is student of Health Polytechnic of Health Ministry of Tasikmalaya which experienced primary dysmenorrhea with medium degree of dysmenorrhea which amount 94 people that divided into 2 groups. 47 respondents were given ginger jelly candy and 47 respondents were given ibuprofen. Samples were taken in consecutive sampling. The results showed that ginger jelly candy can decrease the intensity and duration of dysmenorrhea (p < 0.001). After comparison, there was a difference of decrease in the intensity of dysmenorrhea after the first 6 h of jelly candy and ibuprofen treatment (p < 0.05) but no difference in duration of dysmenorrhea between jelly candy and ibuprofen treatment (p = 0.991) Conclusions in this study are jelly ginger and ibuprofen can both reduce the intensity and duration of dismenorrhea in adolescents. Therefore, the decrease of intensity and duration of dismenorrhea in the group of adolescents treated with ibuprofen were higher than adolescents treated with ginger jelly candies.

Keywords: Ginger jelly · primary dysmenorrhea · adolescents

1 Introduction

Dysmenorrhea is one of the common gynecological complaints experienced by adolescents and affects more than half of menstruating women. The detrimental impact of dysmenorrhea affects psychological and economic aspects so that it affects the quality of life of adolescents themselves. One of the main causes of primary dysmenorrhea is the increased production of uterine prostaglandins derived from cyclooxygenase which is induced by pro-inflammatory cytokines and produces prostaglandins that mediate the inflammatory response, resulting in increased myometrial contractility, uterine ischemia, sensitivity of pain fibers, and finally causing pain during menstruation. Management of dysmenorrhea consists of pharmacological and non-pharmacological approaches. The main pharmacological treatment is the administration of NSAIDs, one of which is ibuprofen, while the popular non-pharmacological treatment is the administration of herbs or phytopharmaceuticals. Ginger as an herbal product contains pharmacological properties as well as NSAIDs. Research and clinical trials conducted by Ozgoli et al., from Iran, showed that ginger has the same analgesic effect as ibuprofen or mefenamic acid, so it can be used to reduce the intensity and duration of dysmenorrhea. By processing ginger into functional food ingredients in the form of jelly candy, it is hoped that it will be more acceptable among adolescents to overcome the problem of primary dysmenorrhea.

2 Research Method

The design of this study is a quasi-experimental research design with a nonequivalent pre-test-post-test control group design The research was conducted at Tasikmalaya, West Java to 94 adolescents midwifery students at Health Polytechnic of the Ministry of Health with primary dysmenorrhea as the study samples.

3 Results

Table 1 describes the pain intensity of adolescent dysmenorrhea before and after being treated in both groups. The table shows that before being given treatment, the average pain intensity of the two groups was at a score of 5 and 6 while after being given treatment the average pain intensity score in the ibuprofen group decreased to 1 and 2. This decrease in intensity was higher than the ginger jelly candy group with a pain intensity score. 3 and 4. The comparison of pain intensity before and after being given treatment showed a p value < 0.001, it showed that there was a significant decrease in pain intensity before and after being given given treatment showed a p value score.

Table 2 shows that the decrease in pain scores from 2 h to 4 h and from 2 h to 6 h has a p value of < 0.05, which means that there is a significant difference in the intensity decrease of dysmenorrhea in the decrease in pain scores in the first 6 h after giving ginger jelly candy and ibuprofen.. The decrease in pain scores from 2 h to 8 h, 2 h to 10 h and 2 h to 12 h had a p value of > 0.05 which means there was no difference in the decrease in the intensity of dysmenorrhea between adolescents who were given ginger jelly candy and adolescents who were given ibuprofen.

From Table 3 known that the median duration of dysmenorrhea before being given ginger jelly candy was 17 h and before being given ibuprofen was 18 h with a p value of 0.020, whereas after being given treatment, either consumption of ginger jelly candy or ibuprofen, the median duration of pain was reduced to 16 h with a value of p 0.991. The decrease in the duration of dysmenorrhea between adolescents who were given ginger jelly candy and adolescents who were given ibuprofen was p = 0.006, which means that there was no difference in the decrease in duration between adolescents who were given given ginger jelly candy and adolescents who were given ibuprofen. Comparison of the duration of dysmenorrhea before and after administration of ginger jelly candy or ibuprofen p < 0.001. This shows that both ginger jelly candy and ibuprofen can reduce

the duration of dysmenorrhea in other words that there is a significant decrease in the duration of dysmenorrhea before and after treatment.

4 Discussion

Table 1 and Table 3 show that before being given treatment (cycle I/pretest), there was no statistically significant difference in terms of intensity and duration of dysmenorrhea in

| | Treatment | | _ |
|-----------------|------------------------|------------------------|-----------------------------|
| | Ginger | Ibuprofen | |
| Pain Intensity | Jelly | (n = 47) | p value [*] |
| 1 and intensity | Candy | | |
| | (n = 47) | | |
| Pre data : | | | |
| Due 2 hours | $((1 7)^{**})$ | ϵ $(A = 7)$ | 0 272 |
| Pre 2 nours | 0(4 - 7) | 0(4-7) | 0,572 |
| Pre 4 nours | 6(4-6) | 5(4-6) | 0,084 |
| Pre o nours | 5(4-6) | 5(4-6) | 0,387 |
| Pre 8 nours | 5(4-6) | 5(4-6) | 0,285 |
| Pre 10 nours | 5(4-6) | 5(4-6) | 0,221 |
| Pre 12 hours | 5(3-6) | 5 (3 – 6) | 0,133 |
| Post Data : | | | |
| Post 2 hours | 4(2-6) | 2(2-5) | < 0,001 |
| Post 4 hours | 4(1-6) | 2(0-3) | < 0,001 |
| Post 6 hours | 4(1-6) | 2(0-3) | < 0,001 |
| Post 8 hours | 3(1-5) | 1(0-3) | < 0,001 |
| Post 10 hours | 3(1-5) | 1(0-3) | < 0,001 |
| Post 12 hours | 3(1-5) | 1(0-3) | < 0,001 |
| | | | ŗ |
| Pre vs Post | | | |
| Comparison : | | n<0.001 | |
| 2 hours | n<0.001 | p < 0.001 p < 0.001 | |
| 4 hours | p < 0.001 n < 0.001 | p < 0.001 p < 0.001 | |
| 6 hours | p < 0.001 | p < 0.001 | |
| 8 hours | p < 0,001 | p < 0.001 | |
| 10 hours | p < 0.001 | p < 0.001 | |
| 12 hours | p < 0.001 | p<0,001 | |
| | p<0,001 | | |

| Tuble 1. Comparison of the intensity of dysinenormed (pre and post) in the two ireatment group | Table 1. | Comparison of t | he intensity of dysm | nenorrhea (pre and po | st) in the two treatment | groups |
|---|----------|-----------------|----------------------|-----------------------|--------------------------|--------|
|---|----------|-----------------|----------------------|-----------------------|--------------------------|--------|

note : *) Mann-Whitney test, **) median value and range; ***) Wilcoxon test

| | Trea | _ | |
|--------------|----------|----------------------|----------------------|
| Dain score | Ginger | Ibuprofen $(n = 47)$ | Nilai |
| decrease | candy | (11 - 47) | \boldsymbol{p}^{*} |
| decrease | (n = 47) | | |
| From 2 hours | | | |
| to 4 hours | 0 (-1 – | 1 (0-4) | 0,002 |
| From 2 hours | 2) ** | 1(0-5) | 0,020 |
| to 6 hours | 0 (-1 – | 1(0-5) | 0,161 |
| From 2 hours | 3) | 1 (0-5) | 0,207 |
| to 8 hours | 1 (0 – | 1(0-5) | 0,142 |
| From 2 hours | 3) | | |
| to 10 hours | 1 (0 – | | |
| From 2 hours | 3) | | |
| to 12 hours | 1(0-3) | | |

Table 2. Comparison of Dysmenorrhea Intensity Score Decrease in the two Treatment groups

note : *) Mann-Whitney test, **) median value and range

Table 3. Comparison of the duration of dysmenorrhea (pre and post) in the two treatment groups

| | Treati | | |
|--|-----------------------------------|----------------------|-------------------------|
| Pain Duration (hour) | Ginger jelly candy (n = 47) | Ibuprofen $(n = 47)$ | p value [*] |
| Pre Data : | 17 (12-25) | 18 (15-22) | 0,020 |
| Post Data : | 16 (10 – 19) | 16 (11 – 19) | 0,991 |
| Decrease in the duration of dysmenorrhea | 1 (0-8) | 2 (0-6) | 0,006 |
| Pre vs post comparison | p<0,001*** | p<0,001 | |

the treatment group (ginger jelly candy) and control group (ibuprofen). This condition was caused by the fact that the respondents in this study both had moderate degrees of dysmenorrhea and homogeneous characteristics in terms of age, menarche, length of menstruation and BMI. After being given treatment (cycle II/posttest) giving ginger ielly candy or ibuprofen to both groups on the first day of menstruation, there was a decrease in the intensity and duration of dysmenorrhea in both groups where the mean value of decreased intensity and duration of dysmenorrhea in the treatment group was lower than the control group. These average values indicate the success of giving ginger jelly candy in reducing the intensity and duration of dysmenorrhea although the decrease in the intensity of dysmenorrhea is lower than ibuprofen. These findings explain ginger as a herbal product that has pharmacological properties as well as NSAIDs. This is in line with research and clinical trials conducted by Ozgoli et al., from Iran, which showed that ginger has the same analgesic effect as ibuprofen or mefenamic acid, so it can be used as a treatment to reduce the intensity and duration of dysmenorrhea. The results of Rahnama et.al's research showed that consuming 1500 mg powdered ginger capsules 2 days before the onset of the menstrual cycle until the third day of menstruation was significantly better in reducing the intensity and duration of dysmenorrhea compared to placebo. In this study, ginger extract was given as much as 2×50 mg. With a span of 6 h when converted is equivalent to 2×1000 mg of ginger powder. This dose increase is safe because based on the results of previous studies giving 50 mg of ginger extract in a single dose did not cause any side effects. In this study, none of the respondents complained of any side effects.

Table 1 shows that there is a decrease in the intensity of dysmenorrhea starting in the first 2 h after giving ginger jelly candy or ibuprofen, while Table 3 shows that there is a decrease in the duration of dysmenorrhea in the first 24 h after giving ginger jelly candy or ibuprofen. This is in accordance with research on the absorption of ibuprofen by Rambe et al. that the absorption of ibuprofen is fast through the stomach and maximum plasma levels are reached after 1.2 h. The plasma half.life is about 2 h. Ninety percent of ibuprofen is bound to plasma proteins. Onset is approximately 30 min. Studies on the pharmacological effects of ginger extract in human clinical trials, analyzed human plasma samples and detected low concentrations of free 10.gingerol and shogaol 6., while mostly 6., 8., 10. And gingerol and shogaol 6.present in plasma as glucuronide and sulfate metabolites. The pharmacokinetics of 6., 8., 10. And gingerols and shogaol 6.and their metabolites were analyzed. The half.life of all compounds and their metabolites is between 1 and 3 h. A pharmacokinetic study conducted by Yanke Yu et al. said that after oral administration of 2 g of ginger extract, the active compound gingerol content began to be detected in plasma 1 h after and began to be undetectable 2 h later.. Meanwhile, the active compound content of shogaol began to be detected in plasma 2 h after the oral dose and began to be undetectable 4 h after. The average half.life of the active compounds metabolized gingerol and shogaol is 1.3 h.

The mechanism of decreasing the intensity and duration of dysmenorrhea is in accordance with the pathophysiology that primary dysmenorrhea occurs due to the mechanism of endometrial prostaglandins and leukotrienes. After ovulation occurs in response to increased production of progesterone, fatty acids will increase in the phospholipids of cell membranes. Then arachidonic acid and other omega-7 fatty acids are released and initiate

a flow of prostaglandins and leukotrienes in the uterus which results in the mediation of the inflammatory response, menstrual cramps, and other menstrual molimina. The product of arachidonic acid metabolism is prostaglandin (PG) F2-alpha, which is a cyclooxygenase (COX) which causes hypertonus and vasoconstriction in the myometrium, resulting in ischemia and menstrual pain. However, the role of prostaglandins and leukotrienes has not been explained in detail and requires further research. To reduce the primary dysmenorrhea, one of the ways is by inhibiting the COX-2 pathway. The active substances found in ginger such as 10 gingerols, 8-shogaol and 10-shogaol can inhibit the COX-2 pathway, so that the formation of prostaglandins from arachidonic acid is disrupted. Other factors that can have an effect on reducing the intensity and duration of dysmenorrhea. In this study, the results of the GCMS test of the main compound identified in the thick ginger extract was 6-shogaol, so it was suspected that 6-shogaol also played a role as a COX-2 inhibitor. Ginger extract contains various components, hence it is important to identify the compounds responsible for its pharmacological effects. In a study mentioned that 6-, 8-, and 10-gingerol and 6-shogaol act as anti-inflammatory, antibacterial, antipyretic, anti-lipidemic, antitumorigenic, and antiangiogenic. In most clinical trials, the content of the active ingredients in ginger extract was not measured. The use of non-standardized ginger extracts in different clinical studies partly explains the mixed results of clinical studies in addition to the various study designs and dosing regimens. A study by Schwertner et al. showed that variable amounts of 6-gingerol, 6shogaol, 8-gingerol, and 10-gingerol in different brands of ginger dietary supplements. In different products, 6-gingerol ranged from 0.00-9.43 mg/g, 6-shogaol ranged from 0.16-2.18 mg/g, 8-gingerol ranged from 0.00-1.10 mg/g, and 10 gingerols ranging from 0.00-1.40 mg/g.42 Active compounds in ginger such as gingerdiol and gingerdione, shogaol, beta-carotene, capsaicin, caffeic acid and curcumin act as inhibitors of cyclooxygenase (COX) and lipooxygenase resulting in inhibition of cyclooxygenase (COX) and lipooxygenase. Leukotrienes and prostaglandin synthesis and acts as an anti-inflammatory and analgesic drug in primary dysmenorrhea.

5 Conclusion

Ginger jelly candy can reduce the intensity and duration of primary dysmenorrhea, although the decrease in the intensity and duration of primary dysmenorrhea in adolescents who consume ibuprofen is higher than adolescents who consume ginger jelly candy. However, giving ginger jelly candy can be an alternative treatment for dysmenorrhea for adolescents who do not like to take drugs.

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Bibliography

- 1. Tria Wahyuningrum S. Penanganan Dismenore Pada Remaja Putri Di Madrasah Aliyah Negeri Mojosari Kabupaten Mojokerto. Jurnal Keperawatan Bina Sehat. 2015;12(2).
- 2. Rakhma A. Gambaran derajat Dismenore dan upaya penanganannya pada siswi Sekolah Menengah Kejuruan Arjuna Depok Jawa Barat. 2012.
- 3. Risnomarta SD. Hubungan OAINS pada Pengobatan Dismenorea dengan Kejadian Dispepsia pada Mahasiswi Fakultas Kedokteran Universitas Andalas. Jurnal Kesehatan Andalas. 2015;4(2).
- Gupta R, Kaur S, Singh A. Comparison to assess the effectiveness of active exercises and dietary ginger vs. active exercises on primary dysmenorrhea among adolescent girls. Nursing and Midwifery Research. 2013;9(4):153.
- 5. Ju H, Jones M, Mishra G. The prevalence and risk factors of dysmenorrhea. Epidemiologic reviews. 2013:mxt009.
- 6. Kashani L, Mohammadi M, Heidari M, Akhondzadeh S. Herbal Medicine in the Treatment of Primary Dysmenorrhea. Journal of Medicinal Plants. 2015;1(53):1-5.
- Mirabi P, Alamolhoda SH, Esmaeilzadeh S, Mojab F. Effect of medicinal herbs on primary dysmenorrhoea-a systematic review. Iranian journal of pharmaceutical research: IJPR. 2014;13(3):757.
- 8. Ozgoli G, Goli M, Moattar F. Comparison of effects of ginger, mefenamic acid, and ibuprofen on pain in women with primary dysmenorrhea. The Journal of alternative and complementary medicine. 2009;15(2):129-32.
- Rahnama P, Montazeri A, Huseini HF, Kianbakht S, Naseri M. Effect of Zingiber officinale R. rhizomes (ginger) on pain relief in primary dysmenorrhea: a placebo randomized trial. BMC complementary and alternative medicine. 2012;12(1):92.
- Gupta SA, Jain M, Gupta A, Soni G. Prevalence of Dysmenorrhea among Adolescent Girls: A Cross-sectional Study from Urban Central India. International Journal of Preventive, Curative & Community Medicine. 2015;1(1):35-7.
- 11. Salmalian H, Saghebi R, Moghadamnia AA, Bijani A, Faramarzi M, Amiri FN, et al. Comparative effect of Thymus vulgaris and ibuprofen on primary dysmenorrhea: A triple-blind clinical study. Caspian journal of internal medicine. 2014;5(2):82.
- 12. Grandi G, Ferrari S, Xholli A, Cannoletta M, Palma F, Romani C, et al. Prevalence of menstrual pain in young women: what is dysmenorrhea. J Pain Res. 2012;5:169-74.
- 13. Jaafarpour M, Hatefi M, Khani A, Khajavikhan J. Comparative Effect of Cinnamon and Ibuprofen for Treatment of Primary Dysmenorrhea: A Randomized Double-Blind Clinical Trial. 2015.
- 14. Gumilar RA. Pengaruh Pendidikan Kesehatan Terhadap Perubahan Tingkat Pengetahuan Dan Sikap Remaja Putri Tentang Penanganan Dismenore Di Smpn 2 Kartasura: Universitas Muhammadiyah Surakarta; 2014.
- 15. Jenabi E. The effect of ginger for relieving of primary dysmenorrhoea. J Pak Med Assoc. 2013;63(1):8-10.
- Unsal A, Ayranci U, Tozun M, Arslan G, Calik E. Prevalence of dysmenorrhea and its effect on quality of life among a group of female university students. Upsala journal of medical sciences. 2010;115(2):138-45.
- 17. Mahvash N, Eidy A, Mehdi K, Zahra MT, Mani M, Shahla H. The effect of physical activity on primary dysmenorrhea of female university students. World Applied Sciences Journal. 2012;17(10):1246-52.
- Proctor M, Farquhar C. Diagnosis and management of dysmenorrhoea. British Medical Journal. 2006;7550:1134.

- 19. Lefebvre G, Pinsonneault O, Antao V, Black A, Burnett M, Feldman K, et al. Primary dysmenorrhea consensus guideline. J Obstet Gynaecol Can. 2005;27(12):1117-46.
- Bachtiar A. Pengaruh Ekstrak Jahe (Zingiber Officinale) Terhadap Tanda Dan Gejala Osteoartritis Pada Pasien Rawat Jalan Di Puskesmas Pandan Wangi Kota Malang. Program Magister Ilmu Keperawatan Kekhususan KMB: Fakultas Ilmu Keperawatan, Depok; 2010.
- 21. van Breemen RB, Tao Y, Li W. Cyclooxygenase-2 inhibitors in ginger (Zingiber officinale). Fitoterapia. 2011;82(1):38-43.
- Agarwal AK, Agarwal A. A study of dysmenorrhea during menstruation in adolescent girls. Indian journal of community medicine: official publication of Indian Association of Preventive & Social Medicine. 2010;35(1):159.
- 23. Sinaga C. Perbedaan Cara Mengatasi Stres dalam Aktivitas Belajar antara Remaja Laki-laki dan Perempuan di SMA Plus Pematang Raya kabupaten Simalungun. 2011.
- Anindita AY. Pengaruh Kebiasaan Mengkonsumsi Minuman Kunyit Asam Terhadap Keluhan Dismenorea Primer Pada Remaja Putri Di Kotamadya Surakarta: Sebelas Maret University; 2010.
- 25. Novia I, Puspitasari N. Faktor Risiko yang Mempengaruhi Kejadian Dismenore Primer. The Indonesian Journal of Public Health. 2008;4(2):96-104.
- 26. Dawood MY. Nonsteroidal anti-inflammatory drugs and changing attitudes toward dysmenorrhea. The American journal of medicine. 1988;84(5):23-9.
- 27. Osayande AS, Mehulic S. Diagnosis and initial management of dysmenorrhea. Am Fam Physician. 2014;89(5):341-6.
- 28. Marjoribanks J, Proctor M, Farquhar C, Derks RS. Nonsteroidal anti-inflammatory drugs for dysmenorrhoea. The Cochrane Library. 2010.
- 29. Wong CL, Farquhar C, Roberts H, Proctor M. Oral contraceptive pill as treatment for primary dysmenorrhoea. The Cochrane Library. 2009.
- Proctor M, Murphy PA. Herbal and dietary therapies for primary and secondary dysmenorrhoea. The Cochrane Library. 2001.
- 31. Proctor M, Farquhar C, Stones W, He L, Zhu X, Brown J. Transcutaneous electrical nerve stimulation for primary dysmenorrhoea. The Cochrane Library. 2002.
- 32. Smith CA, Zhu X, He L, Song J. Acupuncture for dysmenorrhoea. The Cochrane Library. 2011.
- Smith CA, Crowther CA, Petrucco O, Beilby J, Dent H. Acupuncture to treat primary dysmenorrhea in women: a randomized controlled trial. Evidence-Based Complementary and Alternative Medicine. 2011;2011.
- 34. Proctor M, Murphy PA, Pattison HM, Suckling JA, Farquhar C. Behavioural interventions for dysmenorrhoea. The Cochrane Library. 2007.
- MY. D. Primary dysmenorrhea: Advances in pathogenesis and management. Obstet Gynecol. 2006;108:428–41.
- 36. McDowell I. Measuring health: a guide to rating scales and questionnaires: Oxford University Press; 2006.
- 37. Dixon J, Bird H. Reproducibility along a 10 cm vertical visual analogue scale. Annals of the Rheumatic Diseases. 1981;40(1):87-9.
- 38. Hawker GA, Mian S, Kendzerska T, French M. Measures of adult pain: Visual analog scale for pain (vas pain), numeric rating scale for pain (nrs pain), mcgill pain questionnaire (mpq), short-form mcgill pain questionnaire (sf-mpq), chronic pain grade scale (cpgs), short form-36 bodily pain scale (sf-36 bps), and measure of intermittent and constant osteoarthritis pain (icoap). Arthritis care & research. 2011;63(S11):S240-S52.
- 39. Yang J, Chen J, Lao L, Yang M, Chen J, Bo L, et al. Effectiveness study of moxibustion on pain relief in primary dysmenorrhea: study protocol of a randomized controlled trial. Evidence-Based Complementary and Alternative Medicine. 2014;2014.

- 40. Hernani WC. Kandungan Bahan Aktif Jahe dan Pemanfaatannya dalam Bidang Kesehatan. Balai Besar Penelitian dan Pengembangan Pasca Panen Pertanian Bogor. 2011.
- 41. Winarti C, Nurdjanah N. Peluang tanaman rempah dan obat sebagai sumber pangan fungsional. Jurnal Litbang Pertanian. 2005;24(2):47-55.
- 42. Yu Y, Zick S, Li X, Zou P, Wright B, Sun D. Examination of the pharmacokinetics of active ingredients of ginger in humans. The AAPS journal. 2011;13(3):417-26.
- 43. Chrubasik S, Pittler M, Roufogalis B. Zingiberis rhizoma: a comprehensive review on the ginger effect and efficacy profiles. Phytomedicine. 2005;12(9):684-701.
- 44. Aisyah F, Sari Ip. Uji Toksisitas Akut Ekstrak Etanolik Rimpang Jahe Putih Besar (Zingiber Majus Rumph) Pada Tikus Jantan Galur Wistar: Universitas Gadjah Mada; 2013.
- 45. Sundari LPR. Pemberian Kapsul Zink Per Oral Selama Empat Hari Sebelum Haid Menurunkan Kadar Prostaglandin Dan Nyeri Haid Pada Penderita Nyeri Haid Primer Program Studi Biomedik. 2011;Universitas Udayana.
- 46. Zukhrullah M, Aswad M. Kajian Beberapa Senyawa Antiinflamasi: Docking Terhadap Siklooksigenase-2 Secara In Silico. Majalah Farmasi dan Farmakologi. 2012;16(1).
- 47. Connolly TP. Cyclooxygenase-2 inhibitors in gynecologic practice. Clinical medicine & research. 2003;1(2):105-10.
- 48. Octaviana P. Kualitas Permen Jelly dari Albedo Kulit Jeruk Bali (Citrus grandis L. Osbeck) dan Rosela (Hibiscus sabdariffa L.) dengan Penambahan Sorbitol: UAJY; 2013.
- 49. Hasniarti. Studi Pembuatan Permen Buah Dengen. Program studi Ilmu dan Teknologi Pangan. 2012;Universitas Hasanuddin(Makasar).
- Bushra R, Aslam N. An overview of clinical pharmacology of Ibuprofen. Oman Med J. 2010;25(3):155-1661.
- Rambe AZ. Uji Rasio Laju Absorpsi Ibuprofen pada Usus Halus Kelinci (Oryctolagus cuniculus) yang Dikeringkan dengan Freeze Dryer Dibandingkan dengan Usus Halus Kelinci (Oryctolagus cuniculus) Segar. 2014.
- Supriadi MY, Dono Wahyuno. Jahe (Zingiber Officinale Rosc) Status Teknologi Hasil Penelitian Jahe. Balai Penelitian Tanaman Obat Dan Aromatik. 2011;Pusat Penelitian Dan Pengembangan Perkebunan Badan Penelitian Dan Pengembangan Pertanian Kementrian Pertanian. (ISBN 978-979-548-031-0).
- RI K. Suplemen III. Farmakope Herbal Indonesia. Edisi I ed. RI K, editor. Jakarta: Kemenkes RI; 2013.
- 54. DeMAN JM. Kimia Makanan. Edisi kedua ed. Bandung: ITB; 1997.

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