

Management of internal derangement of Temporomandibular joint with platelet rich plasma: Pilot study

Dr. Labeed Sami Hasan ASSTAH. (B.D.S, F.I.B.M.S)*

Dr. Emad. Hammody. ABDULLA.**

Dr.Shatha Nasih TAWFEEQ M.Sc.***

Dr. Sinan Bahjat ISSA (M.B.Ch.B., M.Sc., F.I.B.M.S.)

Dr. Abdullah Ibrahim HAMAD.****

Abstract

Introduction: Temporomandibular joint internal derangements are progressive painful conditions that associated with limitation of mouth opening that caused by disruption within the internal aspects of the TMJ.

The aim of this study was to evaluate the management of internal derangement of TMJ regarding pain intensity in temporomandibular joints and muscles of mastication and mouth opening.

Materials and methods: This study was conducted in the Department of oral and maxillofacial surgery-college of dentistry Tikrit University. Twenty-one patients, 14 women and 7men with age range between 18-33 years old, were diagnosed with internal derangement of TMJ that were treated by intraarticular PRP.

Results: show significant reduction of pain experienced by the patient after 12 weeks(P- < 0.0001). And show significant improvement of maximal interincisal mouth opening (P-<0.000). These finding demonstrating clearly that a significant improvement in both pain , mouth opening and clicking in the TMJ was achieved after the management.

Conclusion: PRP injection to TMJ space for treatment of internal derangement are safe, effective, non-invasive method that help to reduce pain, joint noise and increase mouth opening, thus improving the jaw function.

Key words: internal derangement, platelet rich plasma, disc displacement

Introduction

TMJ internal derangements are progressive conditions that caused by disruption within the internal aspects of the TMJ. With the most common form of disruption is anterior displacement of the articular disk from

its normal functional relationship between the mandibular condyle and the glenoid fossa that lead to joint dysfunction, that manifested as painful joint, joint noise, with or without locking of the mouth.(1,2)Pain

Vol.:15 No.:1 2018

^{*}Assistant professor, College of Dentistry-Tikrit University.

^{**}Assisstant Professor. College of Dentistry-Ibn Sina University .

^{***}Department of biology, College of Science. Tikrit University

^{****}Assistant professor. M.Sc. medical physiology. Ph.D. Oral Medicine.

measurement is difficult as it is unpleasant subjective emotional experience that triggered by noxious stimuli which may or may not be associated with actual tissue damages, (11, 12) Pain measurement scales like four-point scale and visual analogue scale (VAS)have been used by author for measurement of this subjective sensation(4, 5, 6). Normal maximal mouth opening, measured as the distance between the maxillary and mandibular incisors edgesin healthy adults and its values range from (35-50mm) .The maximal interincisal opening (MIO) of at least 35mm is used as a cutoff point to determine trismus, (4,7, 8, 9, 10) Regarding the treatment of temporomandibular joint dysfunction pain syndrome, a number of conservative methods are used, including physical therapy procedures, occlusal splints, rehabilitation and specialist psychological support(11, 12, 13, 14) When the conservative methods fail, surgical methods are followed such as disc repositioning meniscectomy, etc.(15, 16)Platelet-rich (PRP) plasma has been medicinally since the 1970s (1)and is obtained by centrifugation of whole autogenous blood which yield platelet concentration in PRP is at least 5-fold greater than that in physiological blood.(17,18), Platelets are a nucleated fragments of megakaryocytes that formed in the marrow with the life span ranges from 8-12 days, (19, 20)Platelet contain more than 30 bioactive proteins, that play considerable role in hemostasis, and tissue regeneration or healing,(21)

After secretion of the active proteins, they bind to target cells, which include fibroblasts, osteoblasts mesenchymal stem cells and endothelial cells. These bindings have different consequences. Can cause cellular proliferation, collagen synthesis, matrix formation, osteoid

production, etc. thus provoking tissue healing and regeneration of tissue.(22, 23, 24, 25, 26) Growth factors secretion by platelets start within 10 min after activation.(25)PRP is used particularly in orthopedic for regeneration of Osteoarthritis-induced changes, in the management of injured ligaments and tendons.(27, 28)In oral surgery, PRP has extensive uses. PRP is used to induce bone generation in sinus lift procedures and induce bone formation in clef surgery, and jaw reconstruction. It is used to fill the bony gap in Oroantral fistula and after bone resection or defect after cyst or tumor removal (18, 29)PRP used also in Alveolar socket preservation procedures after tooth extraction, in implant surgery and in sinus lifting.(30, 31, 32, 33)The efficacy of PRP has been highlighted in many researches as a method of management of TMJ disorders and the safety of this methods has been mentioned due to the use of autologous material.(34, 35, 36)TMJ internal derangements are painful progressive conditions that caused by many factors including different displacement of the articular disk from its normal position, alterations in TMJ internal pressure and/ or alteration of various biochemical constituents of the synovial fluid that may lead to failure of lubrication.(37)

The aim of this study was to evaluate the management of internal derangement of TMJ regarding pain intensity in temporomandibular joints and muscles of mastication and mouth opening.

Material and methods

This study was conducted in the Department of Oral and Maxillofacial surgery-college of Dentistry Tikrit University. Twenty-one patients, 14 women and 7men with age range between 18-33 years old, were

diagnosed with internal derangement of TMJ that were treated by intraarticular PRP

Inclusion criteria:

- 1- Patients with painful joint.
- 2- Patients without systemic disease were selected.
- 3- Patients were free from any type of drug for the last month.
- 4- Have not treated with any form of joint surgery.

The assessment include:

Pain intensity, which was recorded for each patient using a Visual Analog Scale (VAS). A scale starts from (0 to 10) grades in which (0) represents no pain, while (10) represents the worst possible pain..(38) (39). Mouth opening in millimeter was measured as distant between incisal edge of upper and lower incisor teeth Maximum Interincisal Opening (MIO) was also clinically recorded .(38)(39). Clicking was assisted clinically and classified into absent, the same, decrease, and increased

The assessments were performed at the prior to treatment and then at a follow –up post 1 week, 2 weeks, 4 weeks, 12 weeks examination.

Steps of the procedure

- 1- OPG was taken for all patients
- 2- Platelet, RBC and WBC count were done before the procedures to ensure that all blood elements were within normal limits. Patients with any blood disorder, were excluded from the study.
- 3- Preparation of PRP, double PRP extraction technique been with drawl of 10ml of patient venous blood using wide bore needle. Median cubital vein was selected for venipuncture.(38)The blood is transferred directly to tube containing sodium citrate to prevent coagulation process. Then centrifuged 6 minutes at 1500 RPM order to separate RBC

- from plasma and platelets.(38) The supernatant plasma which contain platelets were Transferred into another sterile tube. Centrifuge tube at a higher speed 3000 rpm for 10 minutes to obtain a platelet concentrate. The lower 1/3rd is PRP and upper 2/3rd is platelet-poor plasma (PPP).Remove PPP and we get PRP that is ready for use about (2 mL).
- 4- The injection sites were determined drawing by patient's skin between the middle of the tragus and the outer eye corner. The posterior entrance point is located along this canthotragal line, about 10mm from tragus and 2 mm below the line.(38)(39).In the Department of oral surgery, while sitting on the dental chair, The patients were requested to open the mouth forcing the condyle forward with the surgeon hand palpating the zygomatic arch and the resultant preauricular concavity.(38).The skin injection site was decontaminated with disinfectant. Then whole joint anesthetized area was lidocaine with epinephrine. One mL of plasma was injected into temporomandibular joint.(38),Patients were informed before the procedures about the possibility of experiencing an unpleasant and transient sensation in the joint regions.

Result

Twenty-one patients, 14 women and 7men with age range between 18-33 years old.

The overall mean for pain experience by the patient before the treatment 5.33 ± 2.06 (range 2-9). Table (1) shows the pain index after I

weeks of PRP injection ,After 12 weeks follow up the pain values have been decrease with mean 1.14 ± 2.03.(Range 0-5). The P-value for pain experience was < 0.0001 as shown in Table (2). Theoverallmean for maximal interincisal mouth opening before treatment was 27.8 ± 7.11 mm (range 14-35 mm). After 1 week be $31.1\pm$ 9.01mm, table (3). After 12 weeks follow the up mean maximalinterincisal mouth opening was47.8± 7.7mm (range 35-55). The Pvalue for maximal interincisal distance was <0.0001 as shown in Table (4).the symptoms outcome of clicking shows that it was be the same in 33% of the patient, decrease in 43 %, absent in 14% ,and increase in 10% of patient clicking fig(1).

Discussion

Facial pain is a common and leading cause of impairment in the jaw function. TMJ pain in is a special problem as the pain increase with mandibular function during chewing and talking.(11, 12,) Several articles and studies have been published concerning appropriate the management of internal derangement of TMJ, with conflicting reports of its efficacy. safety and associated complications.(40, 41)Treatment with education, reassurance rest. instructions to avoid contributing factors, and mild analgesic as first-line treatment.(42)An interocclusal appliance can be helpful for patient complains with pain. Physical therapy have been also attempted for reduction of dysfunction.(43)A variety of agents, such as steroidal anti-inflammatory, hyaluronic acid and chondroitin sulphate have been used as noninvasive solution for controlling sign and symptom related to TMJ problem.(44)Injection of PRP TMJ is a minimally invasive modality with

effectiveness in management of TMJ dysfunctions involving disc derangement without blocking have been highlighted in many researches.(45)

This study demonstrates significant pain reduction after 12 weeks Pvalue< 0.0001 (table 2). The maximal interincisal distance was increased significantly after 12 weeks value<0.0001 (table 4). The patients reported reduction in clicking as shown in the figure (1) which demonstrate reduction in the clicking. These finding was in agreement with study "Hancı M et al who describe in his study a significant reduction in pain intensity, joint sound and increase in mouth opening.(3)The result of this study supported by study that performed by Al-Delayme R.M.A who demonstrate that PRP significantly effective in improvement of the extent of mouth opening with reduction of VAS value. (46)Platelets have golden role in tissue healing. It act as reservoir for growth factors including vascular endothelial growth factor (VEGF), transforming factor-beta 1 (TGF-β1), growth factor fibroblast growth (FGF), epidermal growth factor (EGF), and platelet-derived growth factor (PDGF). Once these growth factors activated it stimulate the healing cascade in cartilage, muscle ligament, tendon, and in bone, through the effect of PRP on cell proliferation, cellular metabolism anti-inflammatory effect injection site.(47, 48, 18, 49, 50) Lippross et al. reported that PRP reduced inflammatory mediator synthesis in the synovial membrane. (47)

The findings in our study demonstrating a significant improvement in both pain and mouth opening was achieved after the management. The symptom of clicking in the TMJ was not significant need amore studies. These improvement can

be attributed to PRP properties such as growth modifier and stimulator with its anti-inflammatory effect.

Conclusion

PRP injection to the upper TMJ space for treatment of internal derangement is safe, effective, non-invasive method that help to reduce pain, joint noise or clicking and increase mouth opening, thus improving the jaw function. More studies using the lateral movements of the mandible as a parameter are needed, Also need to depend on other criteria for selection of patient for internal derangement such as RDC criteria.

References

- 1- Kutuk N, Bas B, Gone ZB, Yilmaz C, Balcioglu E, Ozdamer S, Alkan A. Effect of platelet –rich plasma on fibrocartilage, cartilage, and bone repair in Tempotomandibular joint. J .oral maxillofac surg. 2014,feb:72(2):277-84.doi:10:1016/j.joms 2013,09.011.Epub 20 B Nov.13
- 2- Dolwick MF, Katzberg RW, Helms CA. Internal derangements of the temporomandibular joint: fact or fiction? J Prosthet Dent 1983; 49:415-8.
- 3- Mustaf Hanci, Mehtap Karamese, Zekeriga Tosun, Tahsin Murad Akian, Selcuk Duman ,Nedium Savaci. Intraarticular platelet-rich plasma injection for the treatment of tempro mandibular disorders and a comparison with arthrocentosis. Journal of craniomaxillofacial surgery volume 43.issue 1 jaunary 2015, pages162-166.
- 4- Hasan L. Evaluation of Postoperative Complications after Surgical Removal of Impacted Lower Wisdom Teeth: a Prospective Study. Iraqi Dent. J. 2015; 37(2):62-68.
 - htt://www.iraqidentaljournal.com
- 5- Jose Rodrigues Laureano Filho, Emanuel Dias de Oliveirae Silva, Igor Batist Camargo, and Fabiana MV Gouveia. Theinflence of cryotherapy on reduction of swelling, painandtrismus after third molar extraction. J Am Dent Assoc, Vol 136, No 6,774-778.

- 6- AB Barroso, V Lima, GC Guzzo, RA Moraes, MCVasconcellos, MM Bezerra, FAL Viana, RCR Bezerra, GSM Santana, FA Frota-Bezerra, MO Moraes and MEA Moraes. Efficacy and safety of combined piroxicam, dexamethasone, orphenadrine, and cyanocobalamintreatment in mandibular molar surgery. Braz J Med BiolRes, September 2006, Volume 39)9(1241-1247.
- 7- NurhanGüler ,PerranFuldenYumuk et al . Limited Painful Mouth Opening. J Oral MaxillofacSurg 63:1201-1205, 2005.
- 8- Mezitis M, Rallis G, Zacharides N. The normal range of mouth opening. J Oral Maxillofac Surg1984; 47: 1028–1029.
- 9- Rieder CE. Maximum mandibular opening in patients with and without a history of TMJ dysfunction. J Prosthet Dent 1978; **39:** 441–446.
- 10- Tveteras K, Kristensen S. The aetiology and pathogenesis of trismus. Clin Otolaryngol Allied Sci1986;11:383-7
- 11- J. Okeson, Management of Temporomandibular Disorders and Occlusion, Elsevier, San Diego, Calif, USA, 7th edition, 2013.
- 12- M. Pihut, TheEffectiveness of Prosthetic and Pharmacological Masseter Muscle Relaxation as Alternative Treatment for Temporomandibular Joint Dysfunction, Monograph, Krakow, Poland, 2012
- 13- R. Grey, S. Davies, and A. Quayle, "Th clinical guide to temporomandibular disorders. The clinical guide series," British Dental Journal, pp. 23–60, 2003.
- 14- F. Liu and A. Steinkeler, "Epidemiology, diagnosis, and treatment of temporomandibular disorders," Dental Clinics of North America, vol. 57, no. 3, pp. 465–2004, 2013.
- 15- Katzberg RW, Westesson PL, Tallents RH, Drake CM. Anatomic disorders of the temporomandibular joint disc in asymptomatic subjects. J Oral MaxillofacSurg 1996; 54:147-53.
- 16- Ribeiro RF, Tallents RH, Katzberg RW, Murphy WC, Moss ME, Magalhaes AC and other. The prevalence of disc displacement in symptomatic and asymptomatic volunteers aged 6 to 25 years. J Orofac Pain 1997; 11:37-47.
- 17- Sánchez-González DJ, Méndez-Bolaina E, Trejo-Bahena NI (2012) Platelet- rich plasma peptides: key for regeneration. Int J Pept 2012: 532519.
- 18- Prakash S, Thakur A (2011) Platelet concentrates: past, present and future. J Maxillofac Oral Surg 10: 45-4913.

- 19- Conley CL. Hemostasis. In: Mountcastle VB, editor. Medical Physiology. St. Louis: The C.V. Mosby Company; 2004. pp. 1137–46.
- 20- Harrison P, Cramer EM. Platelet alphagranules. Blood Rev. 1993;7:52–62. [PubMed].
- 21- Schliephake H. Bone growth factors in maxillofacial skeletal reconstruction. Int J Oral Maxillofac Surg. 2002;31:469–84. [PubMed].
- 22- Sunitha Raja V, Munirathnam Naidu E. Platelet-rich fibrin: Evolution of a second-generation platelet concentrate. Indian J Dent Res. 2008;19:42–6. [PubMed].
- 23- Cole BJ, Seroyer ST, Filardo G, Bajaj S, Fortier LA. Platelet-rich plasma: Where are we now and where are we going? Sports Health. 2010;2:203–10. [PMC free article] [PubMed].
- 24- Kevy SV, Jacobson MS. Comparison of methods for point of care preparation of autologous platelet gel. J Extra Corpor Technol. 2004;36:28–35. [PubMed].
- 25- Marx RE. Platelet-rich plasma: Evidence to support its use. J Oral Maxillofac Surg. 2004;62:489–96. [PubMed].
- 26- Antoniades HN, Williams LT. Human platelet-derived growth factor: Structure and functions. Fed Proc. 1983;42:2630–4. [PubMed].
- 27- Li M, Zhang C, Ai Z, Yuan T, Feng Y, et al. (2011) [Therapeutic effectiveness of intra-knee-articular injection of plateletrich plasma on knee articular cartilage degeneration]. Zhongguo Xiu Fu Chong Jian Wai KeZaZhi 25: 1192-1196.
- 28- Lyras D, Kazakos K, Verettas D, Polychronidis A, Simopoulos C, et al. (2010) Immunohistochemical study of angiogenesis after local administration of platelet-rich plasma in a patellar tendon defect. IntOrthop 34: 143-148.
- 29- Carlson NE, Roach RB Jr (2002) Plateletrich plasma: clinical applications in dentistry. J Am Dent Assoc 133: 1383-1386.
- 30- Ogundipe OK, Ugboko VI, Owotade FJ. Can autologous Palatelet-rich plasma gel enhance healing after surgical extraction of mandibular third third molars? JoRALMaxillofac Surg.2011;10:2305-2310.doi:
- 31- Cabbar F, Guler N, Kurkcu M, Iseri U, Sencift K. The effect of bovine bone graft with or without platelet-rich plasma on maxillary sinus floor augmentation. JOralMaxillofac Surg.2011;10:2537-2547.doi:

- 32- Anand U, Mehta DS .Evaluation of immediately loaded dental implants bioactivated with platelet –rich plasma placed in the mandibular posterior region: a clinico radiographic study. J Indian Soc Periodontol.2012;89-95.doi:10.4103/0972-124X.94612.[PMC free article][PubMed][Cross Ref].
- 33- Poesc IPW, Ziya-Ghazvini F ,Schicho K, Buchta C, Moser D, Seemann R,E wers R, Schopper C, Application of platelet rich plasma for enhanced bone regeneration in grafted sinus. J Oral Maxillofac Surg. 2012.04.027[PubMed][Cross Ref].
- 34- R. Civinini, A. Macera, L. Nistri, B. Redl, and M. Innocenti, "The use of autologous blood-derived growth factors in bone regeneration," Clinical Cases in Mineral and Bone Metabolism, vol. 8, no. 1, pp. 25–31, 2011.
- 35- Z. Stopa, H. Wanyura, L. Pączek, and D. Samolczyk-Wanyura, "Th level of cytokines and proteolitic enzymes in the articular synovial flude and blood serum of patients with temporomandibular joint disease," Polish Journal of Stomatology, vol. 67, no. 7, pp. 429–443, 2010.
- 36- B. J. Cole, S. T. Seroyer, G. Filardo, S. Bajaj, and L. A. Fortier, "Platelet-rich plasma: where are we now and where are we going?" Sports Health, vol. 2, no. 3, pp. 203–210, 2010.
- 37- Alpaslan C, Bilgihan A, Alpaslan GH, Gu"ner B, Ozgu"rYis M, Erbas, D (2000) Effect of arthrocentesis and sodium hyaluronate injection on nitrite, nitrate, and thiobarbiturate acid-reactive substance levels in the synovial fluid. Oral Surg Oral Med Oral Pathol Oral RadiolEndod 89:686–69.
- 38- Vladimir Machoň, Monika Řehořová, Jiří Šedý, and René Foltán; APlatelet-Rich Plasma in Temporomandibular Joint Osteoarthritis Therapy: A 3-Month Follow-Up Pilot Study: J Arthritis, 2013, 2:2
- 39- M. Pihut, M. Szuta, E. Ferendiuk, and D. Zeńczak-Więckiewicz. Evaluation of Pain Regression in Patients with Temporomandibular Dysfunction Treated by Intra-Articular Platelet-Rich Plasma Injections, A Preliminary Report;BioMed Research International Volume 2014 (2014), Article ID 132369, 7 pages http://dx.doi.org/10.1155/2014/132369
- 40- Kai S, Kai H, Tabata O, et al. Long-term outcomes of nonsurgical treatment in nonreducing anteriorly displaced disk of the temporomandibular joint. Oral Surg



- Oral Med Oral Pathol Oral RadiolEndod1998;85(3):258–67.
- 41- Minakuchi H, Kuboki T, Matsuka Y, et al. Randomized controlled evaluation of non-surgical treatments for temporomandibular joint anterior disk displacement without reduction. J Dent Res 2001;80(3):924–8.
- 42- de Bont LG, Dijkgraaf LC, Stegenga B. Epidemiology and natural progression of articular temporomandibular disorders. Oral Surg Oral Med Oral Pathol Oral RadiolEndod 1997;83(1):72–6.
- 43- Suvinen TI, Hanes KR, Reade PC.
 Outcome of therapy in the conservative management of temporomandibular pain dysfunction disorder. J Oral.Rehabil 1997;24(10):718–24.
- 44- Hayami T (2008) Osteoarthritis of knee joint as a cause of musculoskeletal ambulation disability symptom complex (MADS). Clin Calcium 18:1574-1580.
- 45- E. T. Daif, "Autologous blood injection as a new treatment modality for chronic recurrent temporomandibular joint dislocation," Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology, vol. 109, no. 1, pp. 31–36, 2010.
- 46- Al-Delayme, R.M.A., Alnuamy, S.H., Hamid, F.T. et al. The Efficacy of

- Platelets Rich Plasma Injection in the Superior Joint Space of the Tempromandibular Joint Guided by Ultra Sound in Patients with Non-reducing Disk Displacement. J. Maxillofac. Oral Surg. (2016). doi:10.1007/s12663-016-0911-9.
- 47- Lippross S, Moeller B, Haas H, Tohidnezhad M, Steubesand N, et al. (2011) Intraarticular injection of plateletrich plasma reduces influmnation in a pig model of rheumatoid arthritis of the knee joint. Arthritis Rheum 63: 3344-3353.
- 48- Napolitano M, Matera S, Bossio M, Crescibene A, Costabile E, et al. (2012)Autologous platelet gel for tissue regeneration in degenerative disorders of the knee. Blood Transfus 10: 72-77.
- 49- Middleton KK¹, Barro V, Muller B, Terada S, Fu FH, Evaluation of the effects of platelet-rich plasma (PRP) therapy involved in the healing of sports-related soft tissue injuries. (Iowa Orthop J. 2012;32:150-63.
- 50- Civinini R¹, Macera A, Nistri L, Redl B, InnocentiM.The use of autologous blood-derived growth factors in bone regeneration. Clin Cases Miner Bone Metab. 2011 Jan;8(1):25-31.

Table (1): Statistical comparison regarding the pain index before and after 1 weeks of the PRP injection (df=20).

Study group	Mean pain index ± SD	T- test	S.E.	Confidence Interval	P- value
Before (n= 21)	5.33 ± 2.06	3.2	0.16	0.18 to 0.87	0.0045*
After 1 weeks (n= 21)	4.81 ± 2.14				

^{*} Highly significant

Table (2): Statistical comparison regarding the pain index before and after 12 weeks of the PRP injection(df= 20).

Study group	Mean pain index ± SD	T- test	S.E.	Confidence Interval	P- value
Before op (n= 21)	5.33 ± 2.06	11.55	0.36	3.43 to 4.95	< 0.0001*
After 12 weeks (n= 21)	1.14 ± 2.03				

^{*} Highly significant

Table (3): Statistical comparison regarding the MIO before and after 1 weeks of the operation (df= 20).

Study group	Mean MIO ± SD	T- test	S.E.	Confidence Interval	P- value
Before (n= 21)	27.8 ± 7.11	3.05	1.09	1.05 to 5.62	0.0064*
After 1 weeks (n= 21)	31.1 ± 9.01				

^{*} Highly significant

Table (4): Statistical comparison regarding the MIO before and after 12 weeks of the operation (df= 20).

Study group	Mean MIO ± SD	T- test	S.E.	Confidence Interval	P- value
Before op (n= 21)	27.8 ± 7.11	11.8	1.697	16.5 to 23.6	<0.0001*
After 12 weeks (n= 21)	47.8 ± 7.7				

^{*} Highly significant

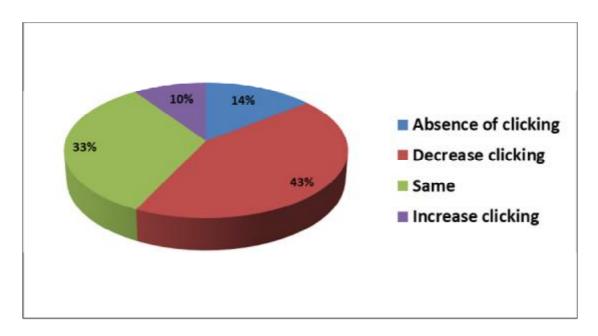


Figure 1: Distribution of cases according to the symptomatic outcome of patient clicking.