



Crossing the Digital Divide in Journal Publishing:

전자학술지와 인쇄본 발행 업무의 일원화

이 춘 실

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Agenda

- Journal Publishing/Production Workflows
- Journal Editing Workflows
- Recommendations
- Production Output Examples



Journal Editing & Publishing Workflows of most journals currently published in Korea

Manuscript
Accepted

Editing &
Preparation

Production

Distribution



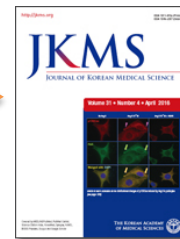
Digital
Divide



Online



Print



For efficient and fast publishing

- 이원화 되어 있는 학술지 발행 과정 개선은 당면 과제
- Problems → To be resolved
 - Print publication, then online (Separate processes)
→ E-journal/Print at the same time (One workflow)
 - Manual processing → Automatic generation
 - Human Errors → Machine/Systematic detection
(error prone → error free)

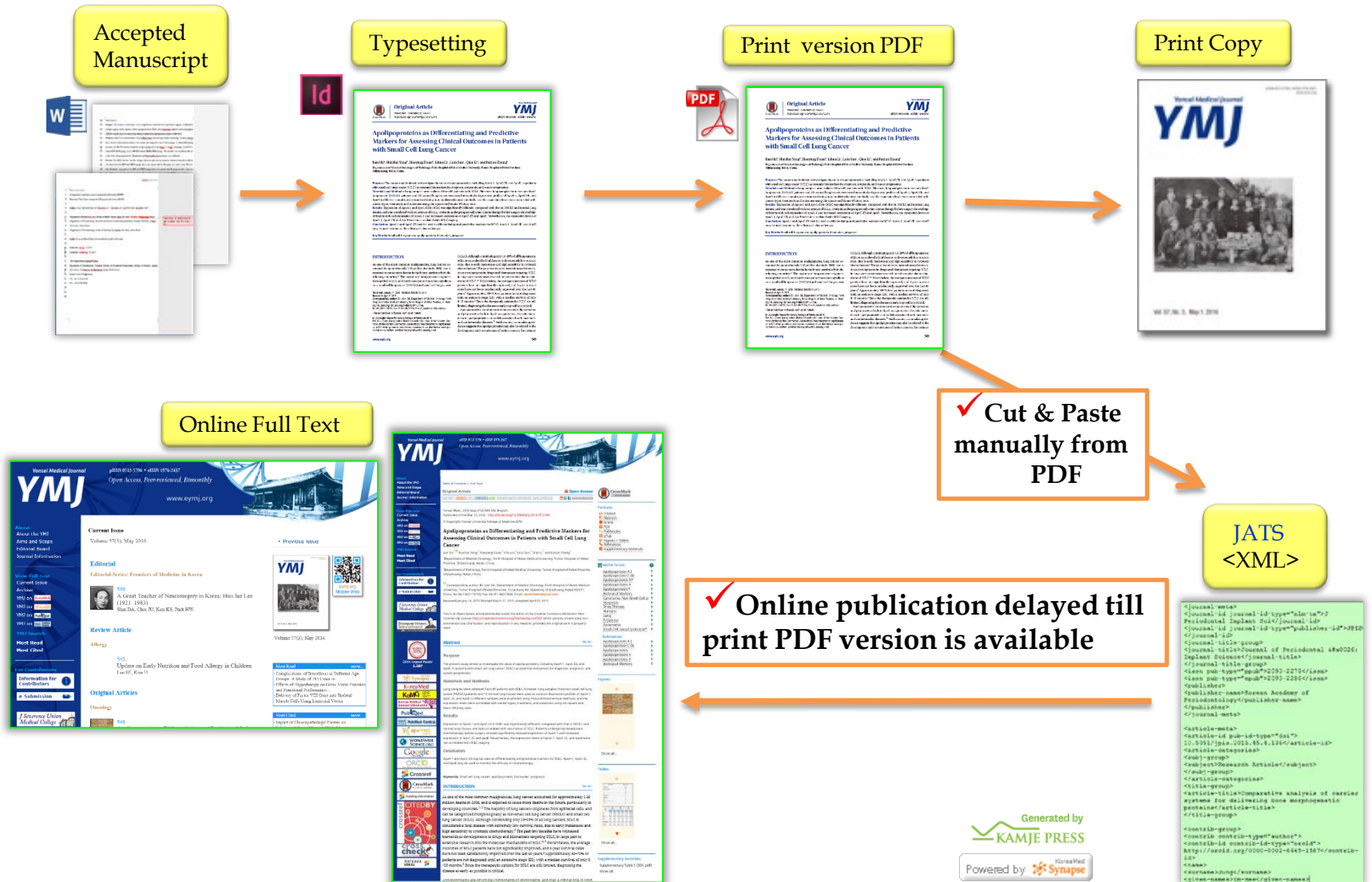
Time, Human resource and Cost savings

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Journal Publishing/Production Workflows

Journal Publishing/Production Workflow (1): From Final Print version PDF to JATS XML



From Final Print version PDF to JATS XML

- XML file production from print version PDFs
- Manual cut & paste
- Slow, Tedious, Error-prone
- Online publication delayed till print PDF version is available
- Current publishing workflows for most Korean (medical) journals





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Korean Medical Journal Information

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KoreaMed Synapse

PubMed Central

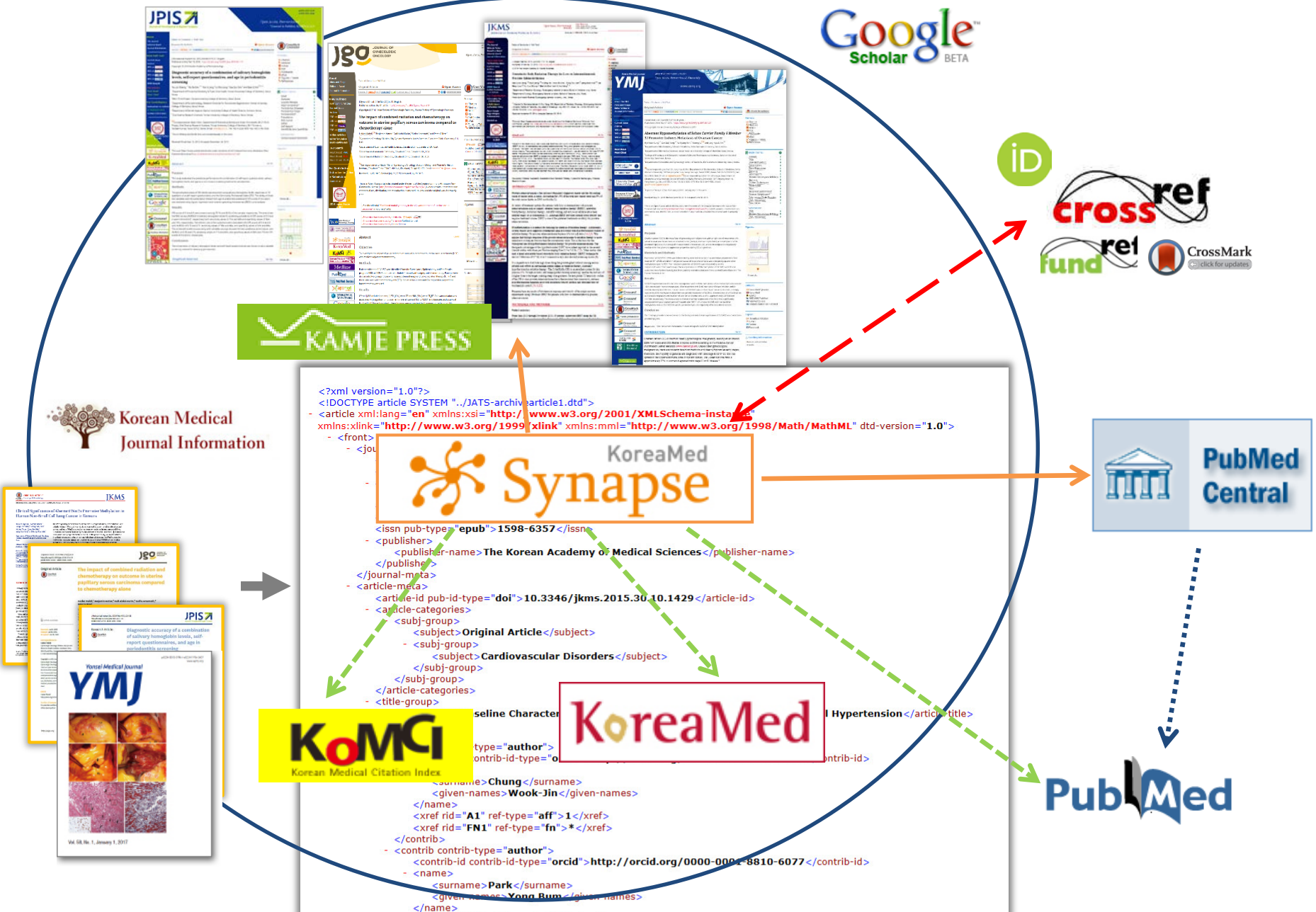
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KoMCI
Korean Medical Citation Index

KoreaMed

PubMed

Synapse & its byproduct/derivative databases



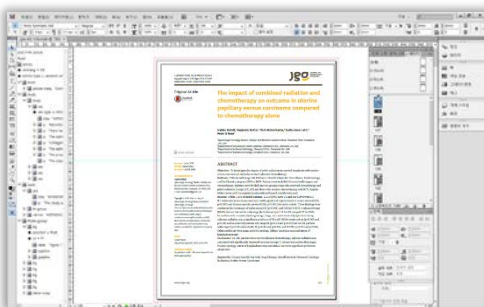
Journal Editing/Publishing/Production Workflow (1)

[Online Full Text](#)

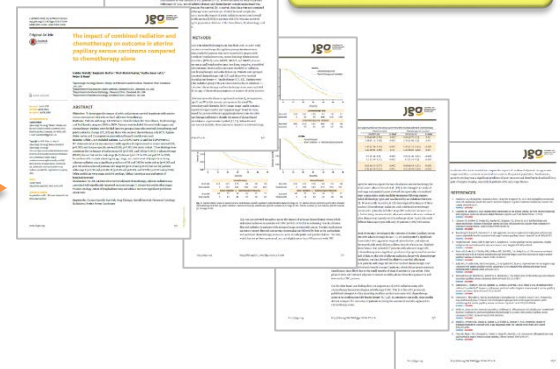
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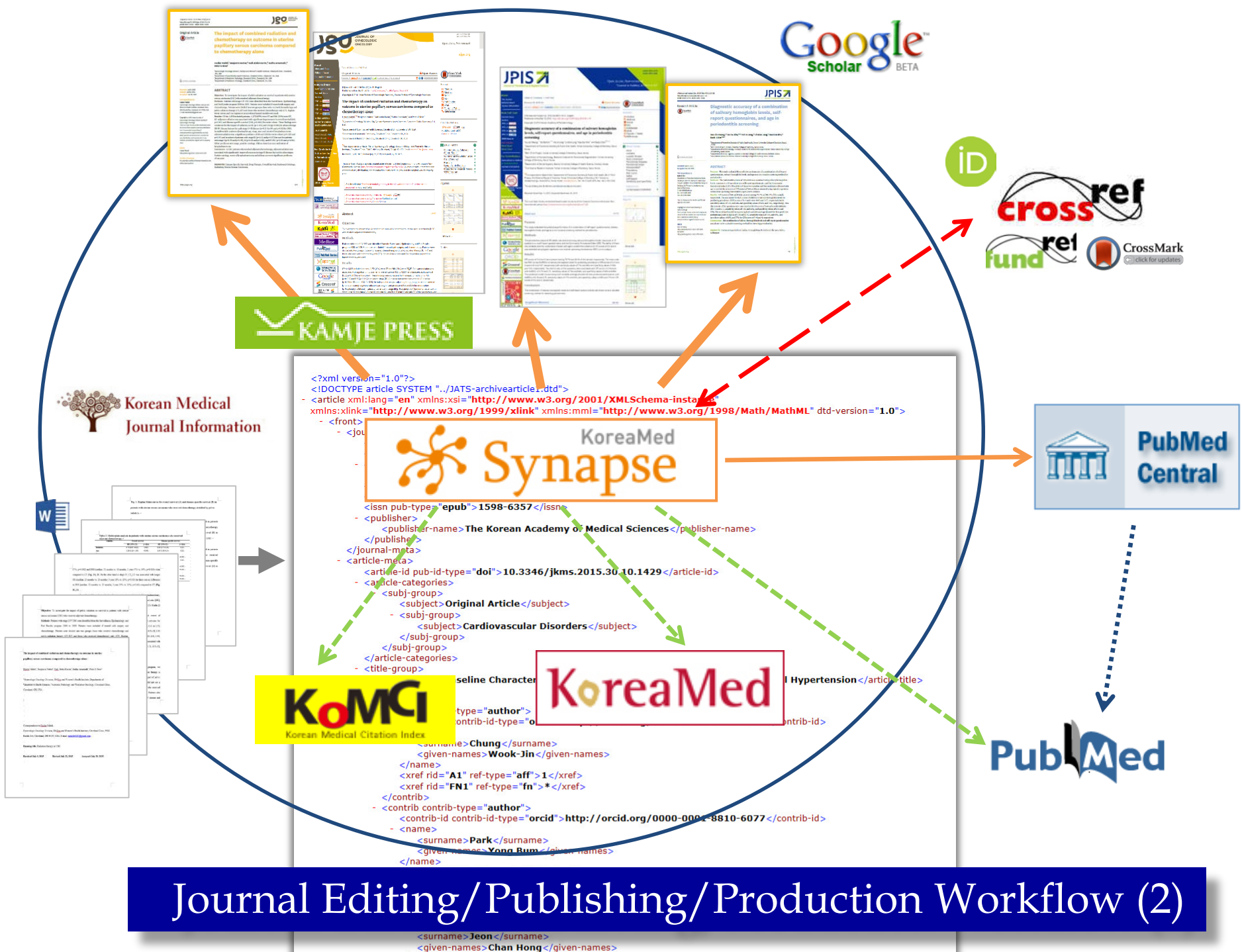


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Interactive PDF





Journal Publishing/Production Workflow (3): One JATS XML for Online, Typesetting and Print

Accepted,
Fully edited
Manuscript

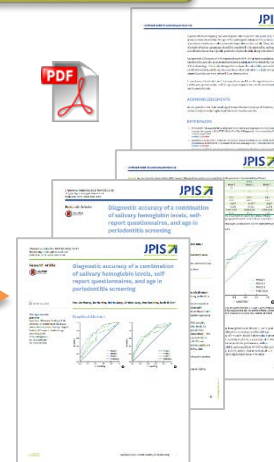
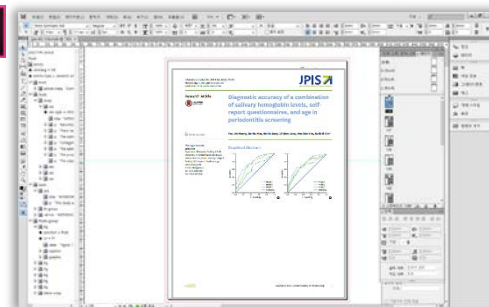
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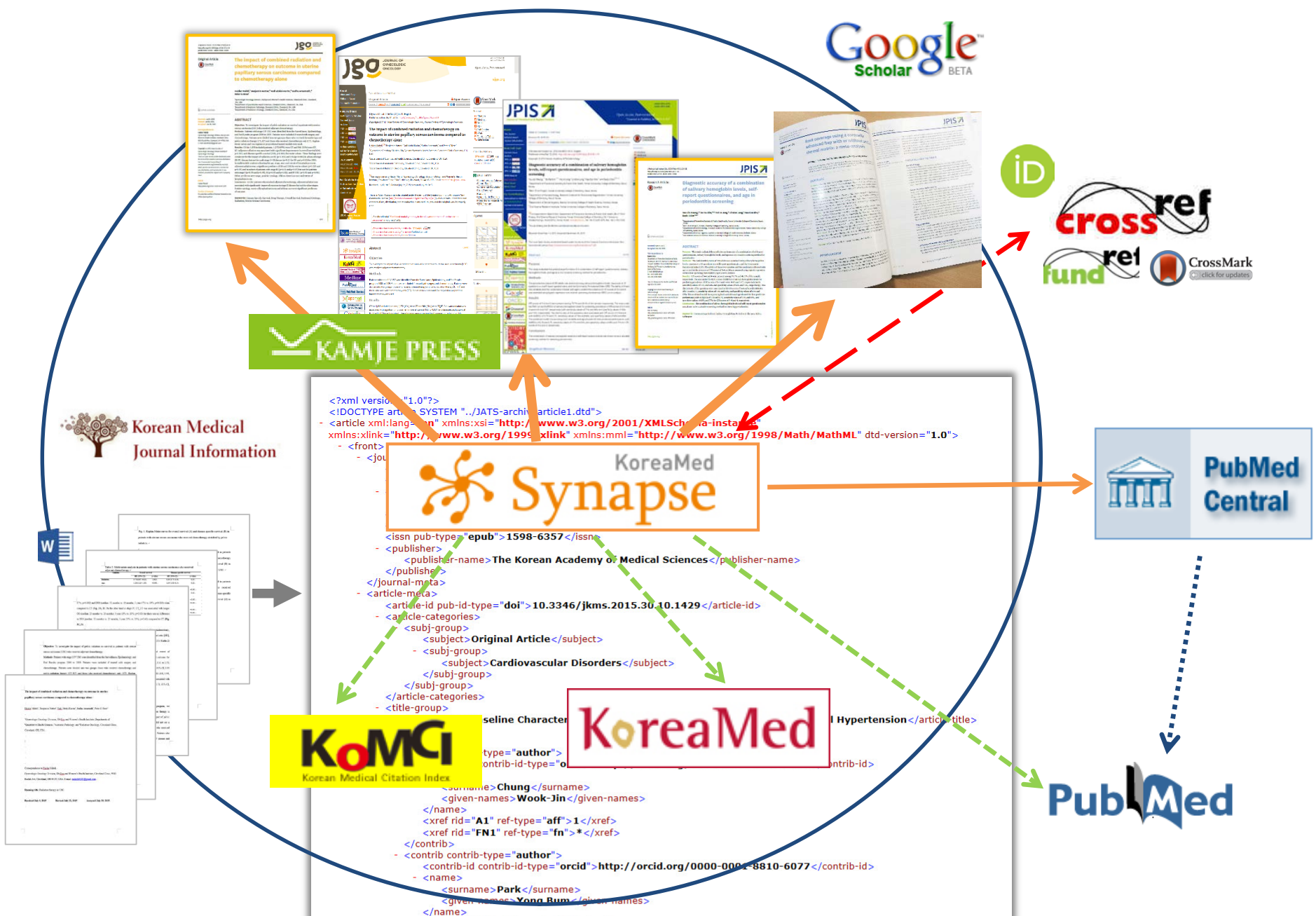
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Journal Editing/Publishing/Production Workflow (3)

One JATS XML for Online and Typesetting

- JATS XML conversion from “Author Manuscripts (Word file)”
 - Online Full text from JATS XML
 - At the same time, automated generation of (InDesign) Typesetting/Layout file from JATS XML
- Online Interactive PDF
- Print



More Efficient Publishing Workflows

- No more cut & paste of texts
Automated conversion of manuscript texts to JATS XML
- Speedy
- One-point (integrated) publishing
- E-journal only
E-journal and Print
- Interactive PDF (vs. static PDF)



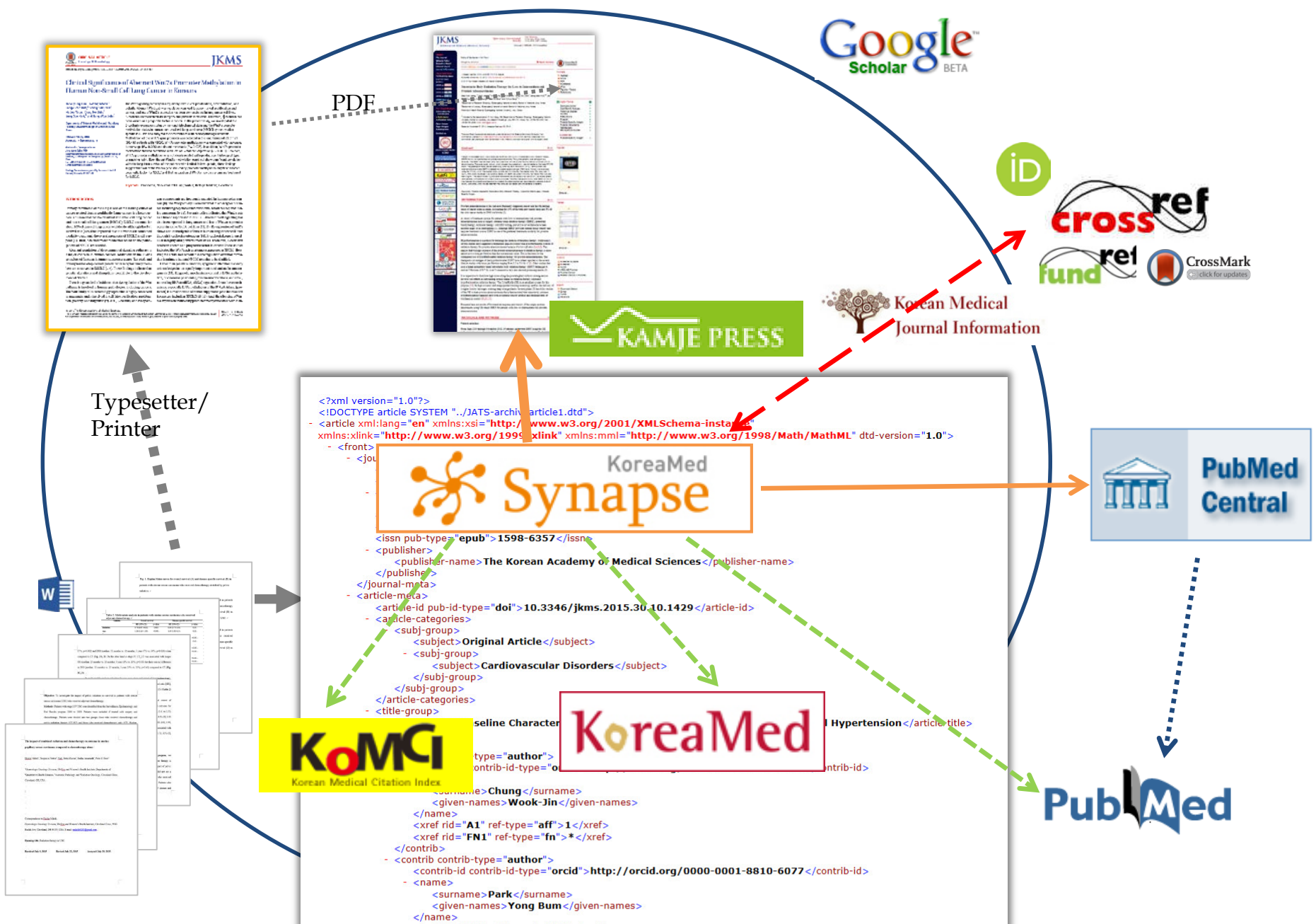
Separate Processing of Online and Print versions



Separate Processing of Online and Print Versions

- Online version and Print version published as two separate processing from the same fully edited manuscript
- Version Control problems
Any changes made on one side should be reflected on the other side





Journal Editing/Publishing/Production Workflow (4)

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Journal Editing Workflows

Any changes made after XML file production/InDesign Typesetting

- Require revisions on XML files to regenerate Online Full text and PDF
- Or, two revisions both on Online Full text (XML) and on PDF
- Better to have all the manuscript editing done before hands (before XML production)

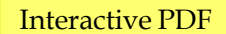


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Online Full Text

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Journal Editing/Production Workflow (3): One JATS XML for Online, Typesetting and Print

Accepted,
Fully edited
Manuscript

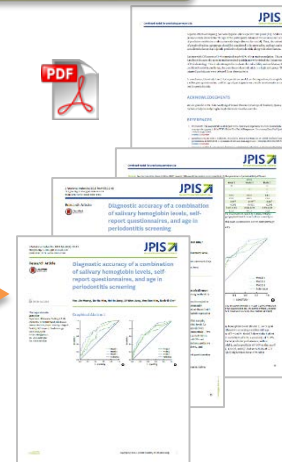
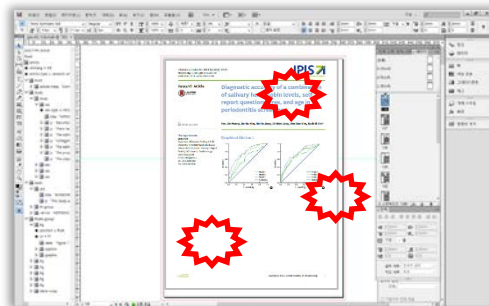
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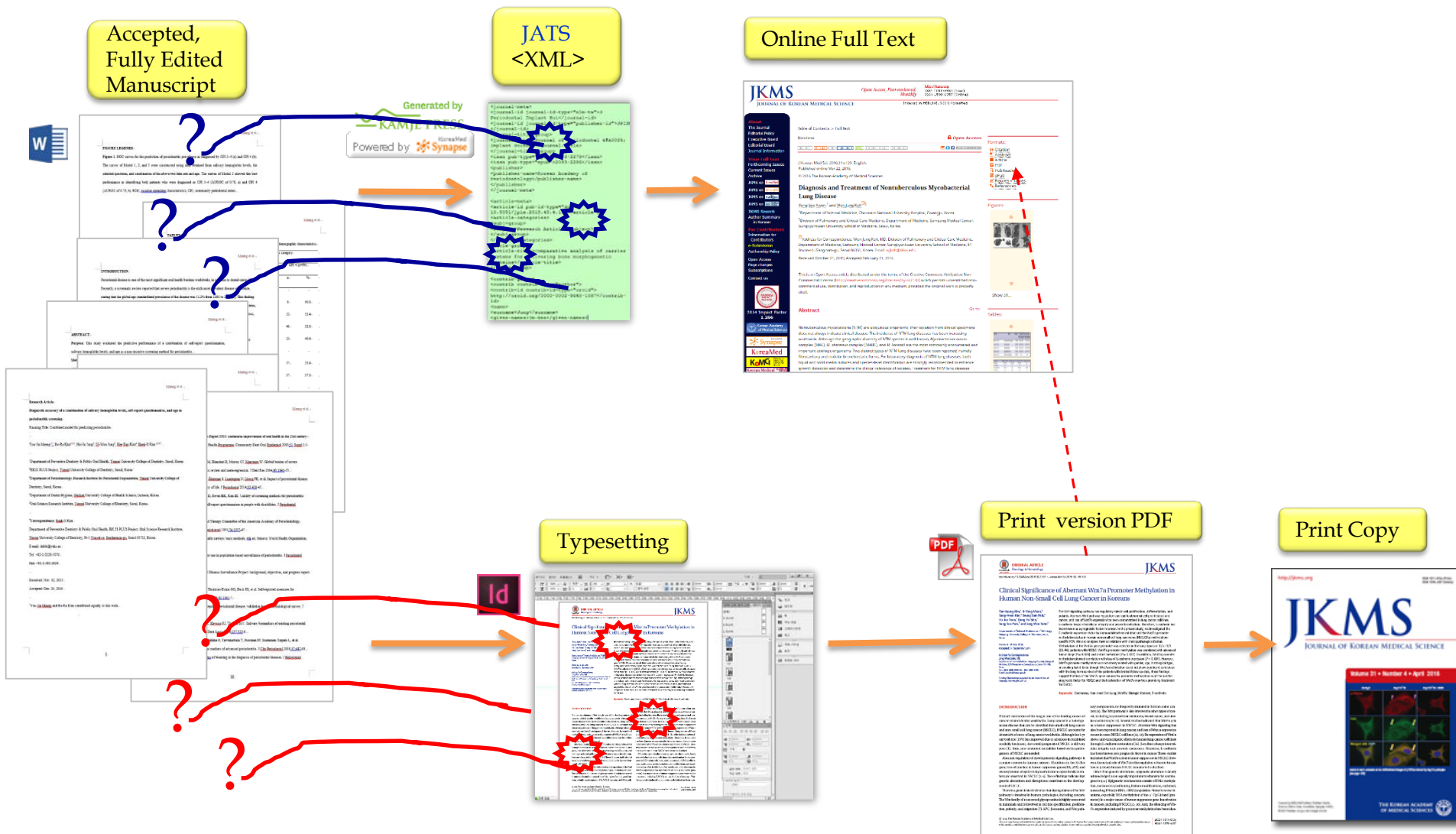
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Interactive PDF

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Journal Editing/Production Workflow (4): Separate Processing of Online and Print version



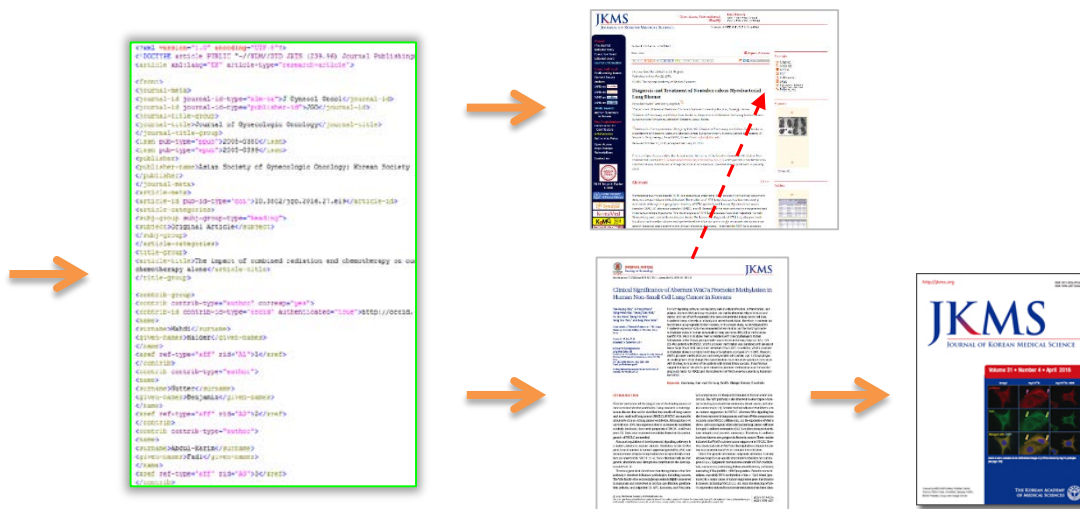
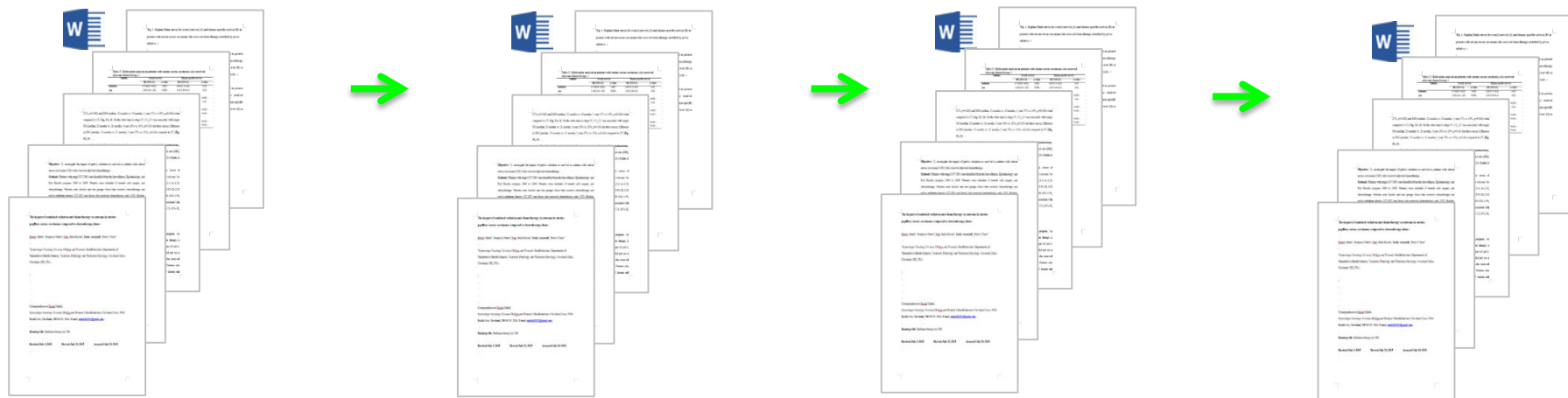
Modified Editing Workflows: Fully Edited “Final” Manuscripts

Accepted
Manuscript

Manuscript
Editing

English
Correction

Author
Proofreading



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Recommendations

Recommendation (1)

- Prepare fully edited “Final” manuscripts
 - Formats
 - Copy editing
 - Manuscript editing
 - English correction
 - Author proofreading



Recommendation (2)

- Produce JATS XML file from fully edited manuscripts
- Automate JATS XML file production and processing
 - to reduce manual cut & paste work
 - to detect any systematic, technical errors



Recommendation (3)

Once a proper JATS XML file is produced

- Generate both online version and PDF (via InDesign) version of full text using the same XML file.

Interactive PDF with internal and external links can be made from fully marked up XML files.

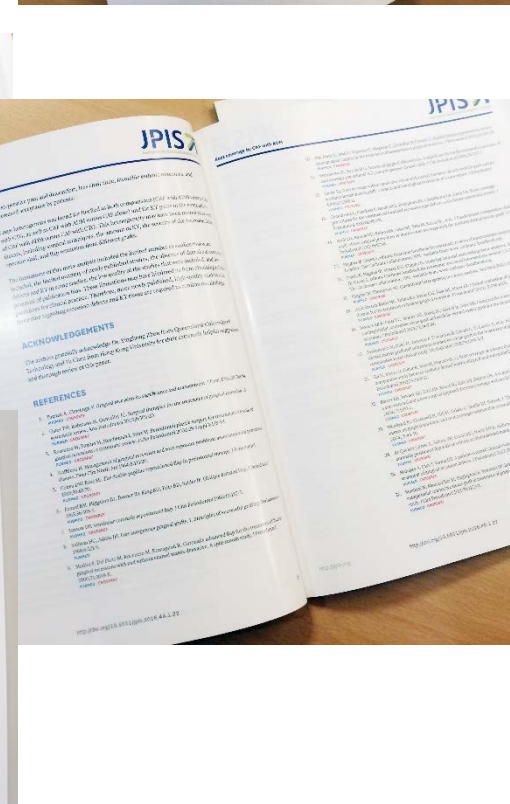
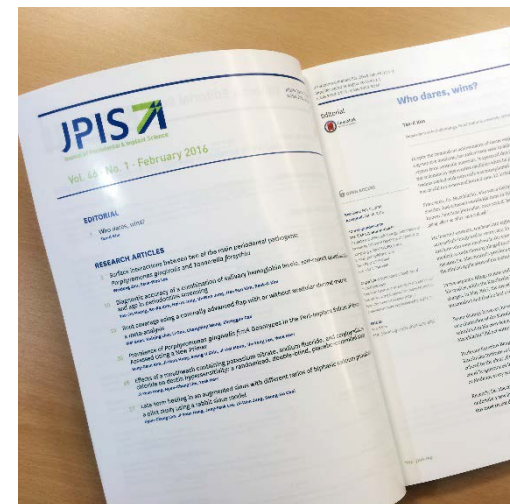
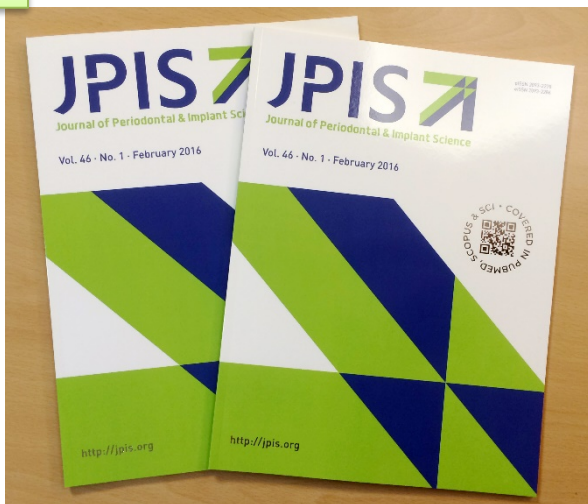




Production Output Examples

Journal Publishing Workflow (2) & (3):
One JATS XML for Online and Typesetting





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CITATION

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Impact on survival with adjuvant radiotherapy for clear cell, mucinous, and endometrioid ovarian cancer: the SEER experience from 2004 to 2011

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Abstract

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Objective

Evaluate the impact of radiotherapy on cause specific survival (CSS) and overall survival (OS) for stage (I–III) clear cell, mucinous, and endometrioid ovarian cancer.



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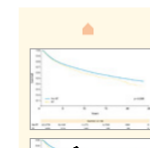
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MeSH Terms:

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Female
Humans
Incidence
Mucins*
Ovarian Neoplasms*
Ovary
Radiotherapy
Radiotherapy, Adjuvant*

Substances:
Mucins

Figures:



Original Article



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This paper was presented at Poster Presentation at American Society for Therapeutic Radiology and Oncology, October 2015.
<http://arjod.org/0000-0000-0675-616X>
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This paper was presented at Poster Presentation at American Society for Therapeutic Radiology and Oncology, October 2015.

Conflict of Interest:
No potential conflict of interest relevant to this article was reported.

<http://ejgo.org>

Impact on survival with adjuvant radiotherapy for clear cell, mucinous, and endometrioid ovarian cancer: the SEER experience from 2004 to 2011

Sagar C. Patel,¹ Jonathan Frandsen,² Sudershan Bhatia,^{1,3} David Gaffney²

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ABSTRACT

Objective: Evaluate the impact of radiotherapy on cause specific survival (CSS) and overall survival (OS) for stage (I–III) clear cell, mucinous, and endometrioid ovarian cancer. **Methods:** We analyzed incidence, survival, and treatments from the Surveillance, Epidemiology, and End Results (SEER) Program from 2004 to 2011 for clear cell, mucinous, and endometrioid histologies of the ovary for stages (I–III). We examined CSS and OS for all three histologies combined and each histology with relation to the use of adjuvant radiation therapy (RT). Survival analysis was calculated by Kaplan-Meier and log-rank analysis. **Results:** CSS was higher in individuals not receiving RT at 5 years (81% vs. 74%) and 10 years (74% vs. 69%, $p=0.003$). OS was higher in individuals not receiving RT at 5 years (76% vs. 73%) and 10 years (64% vs. 59%, $p=0.039$). Stage III patients receiving RT had a higher OS at 5 years (54% vs. 44%) and 10 year intervals (36% vs. 30%, $p=0.037$). Stage III patients with mucinous histology receiving RT had a higher OS at 5 years (50% vs. 36%) and 10 years (45% vs. 26%, $p=0.052$). **Conclusion:** Those receiving RT had a lower CSS and OS at 5 and 10 years. However, subgroup analysis revealed a benefit of RT in terms of OS for all stage III patients and for stage III patients with mucinous histology.

Keywords: Ovary; Radiation; Survival; Uncommon Histology

INTRODUCTION

In the United States, ovarian cancer is the most common cause of gynecologic-related cancer mortality [1]. Specifically, nearly 23,000 women are diagnosed with ovarian cancer yearly, and of these, 14,000 women die every year. A majority of these women are diagnosed with advanced disease, namely, The International Federation of Gynecology and Obstetrics stage III disease [1]. The established standard of care for ovarian cancer has been total abdominal hysterectomy, bilateral salpingo-oophorectomy, omentectomy, washings, and suspicious node removal with adjuvant platinum based chemotherapy, based on extent of primary and nodal involvement, margins, and residual disease [2]. Unfortunately, the 5-year overall survival (OS) for ovarian cancer is about 40% and the median progression free survival for advanced ovarian cancer is about 18 months [1].

References

Go to:

- Jayson GC, Kohn EC, Kitchener HC, Ledermann JA. Ovarian cancer. *Lancet* 2014;384:1376–1388. [PUBMED](#) [CROSSREF](#)
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RT, clear cell, mucinous, endometrioid ovarian cancer

In conclusion, over 2004 to 2011, only 3% of all clear cell, endometrioid, and mucinous ovarian cancer cases were treated with adjuvant RT. Subgroup analysis revealed a benefit of RT in terms of OS for all stage III patients and for stage III patients with mucinous histology. These findings together with previous studies that demonstrated a potential survival benefit of adjuvant RT for stage I and II patients in these histologies suggest a role of RT. Therefore, further investigation should be performed in the indication for RT, dose and volume treated, RT techniques and delivery, treatment compliance, and the patient's functional status for non-metastatic clear cell, mucinous, and endometrioid ovarian cancer.

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- Jayson GC, Kohn EC, Kitchener HC, Ledermann JA. Ovarian cancer. *Lancet* 2014;384:1376–1388. [PUBMED](#) [CROSSREF](#)
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- Rai B, Bansal A, Patel FD, Sharma SC. Radiotherapy for ovarian cancers - redefining the role. *Asian Pac J Cancer Prev* 2014;15:4759–4763. [PUBMED](#) [CROSSREF](#)
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- Sorbe B, Swedish-Norwegian Ovarian Cancer Study Group. Consolidation treatment of advanced (FIGO stage III) ovarian carcinoma in complete surgical remission after induction chemotherapy: a randomized, controlled, clinical trial comparing whole abdominal radiotherapy, chemotherapy, and no further treatment. *Int J Gynecol Cancer* 2003;13:278–286. [PUBMED](#) [CROSSREF](#)

Tumor FDG uptake heterogeneity in cervical cancer

Table 1. Clinicopathological characteristics of patients who underwent positron emission tomography/computed tomography before operation for cervical cancer (n=85)

Characteristic	No. (%)
Age (yr), median (range)	47 (27-69)
Progression-free survival (mo), median (range)	32 (6-83)
FIGO stage	
Ib1	59 (69.4)
Ib2	11 (12.9)
IIA	15 (17.6)
Histology	
Squamous cell carcinoma	59 (69.4)
Adenocarcinoma	16 (18.8)
Adenosquamous carcinoma	5 (5.9)
Others	5 (5.9)
Tumor diameter, median (range)	3.2 (0.5-9.5)
Lymph node metastasis	20 (23.5)
Parametrium invasion	15 (17.7)
Recurrence	14 (16.5)

FIGO, International Federation of Gynecology and Obstetrics.

primary tumor size ($r=0.455$, $p<0.001$), depth of cervical stromal invasion ($r=0.425$, $p=0.001$), and PM invasion ($r=0.227$, $p=0.039$). **Supplementary Table 2** depicts Descriptive statistics for each MTV by SUV threshold.

3. Cut-off value for tumor heterogeneity

The ROC curves used to analyse the IFH in relation to PFS. IFH at an SUV of 2.0 was used for ROC analysis, and the area under the curve was 0.661, and 0.418 was determined to be the cut-off value.

4. Tumor heterogeneity and recurrence

Table 2 summarizes the prognostic values of all of the parameters investigated in the current study. Except FIGO stage which was calculated as categorized variable, other parameters were calculated as continuous variables. Cox proportional hazard analysis revealed that recurrence was significantly associated with TLG_{total} ($p<0.001$), MTV_{total} ($p<0.001$), SUV_{total} ($p=0.015$), FIGO stage ($p=0.015$), SUV_{total} ($p=0.004$), and IFH ($p=0.005$). The Kaplan-Meier survival graphs (**Fig. 2**) showed that PFS differed significantly in groups of subjects categorized based on IFH ($p=0.013$, log-rank test).

Table 2. Analyses of prognostic factors for progression-free survival in patients with cervical cancer

Variable	Test for PFS	HR	95% CI	p-value
Age (yr)		1.000	0.946-1.057	0.999
FIGO stage	II vs. I	3.746	1.290-10.881	0.015
Tumor size		1.177	0.924-1.500	0.188
LN metastasis	Present vs. absent	1.716	0.516-5.497	0.363
PM invasion	Present vs. absent	1.899	0.594-6.046	0.279
SUV _{total}		1.040	1.008-1.072	0.011
MTV _{total}		1.020	1.008-1.032	0.001
TLG _{total}		1.001	1.001-1.002	<0.001
SUV _{total}		1.105	1.033-1.184	0.004
IFH		2.456	2.853-119.310	0.005

FIGO, International Federation of Gynecology and Obstetrics; HR, hazard ratio; IFH, intratumoral [¹⁸F] fluorodeoxyglucose (FDG) uptake heterogeneity; LN, lymph node; MTV, metabolic tumor value; PFS, progression-free survival; PM, parametrium; SUV, standardized uptake value; TLG, total lesion glycolysis.

Tumor FDG uptake heterogeneity in cervical cancer

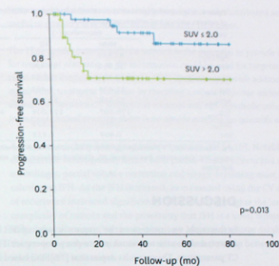


Fig. 2. Kaplan-Meier survival graph shows significantly different progression-free survival between the groups categorized by intratumoral [¹⁸F] fluorodeoxyglucose uptake heterogeneity above (blue line) and below (green line) cut-off value (0.418; $p=0.013$, log-rank test).

5. Prediction of recurrence

Table 3 presents the multivariate regression analysis of the prognostic values of the parameters determined to be significant in univariate analysis. Multivariate analysis identified IFH (HR, 756.997; 95% CI, 2.047 to 279,923.191; $p=0.028$) was the only independent risk factor for recurrence in the current study.

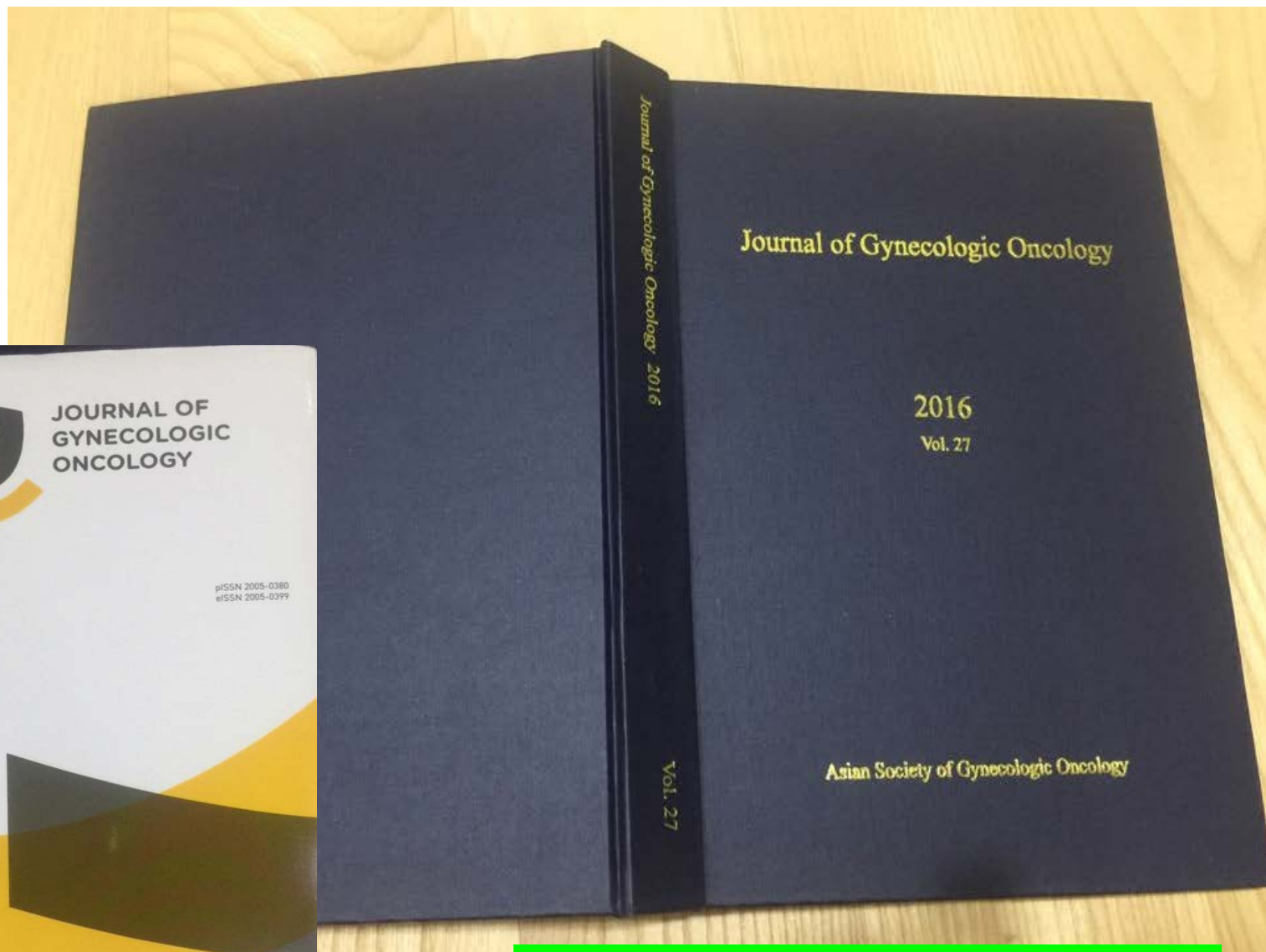
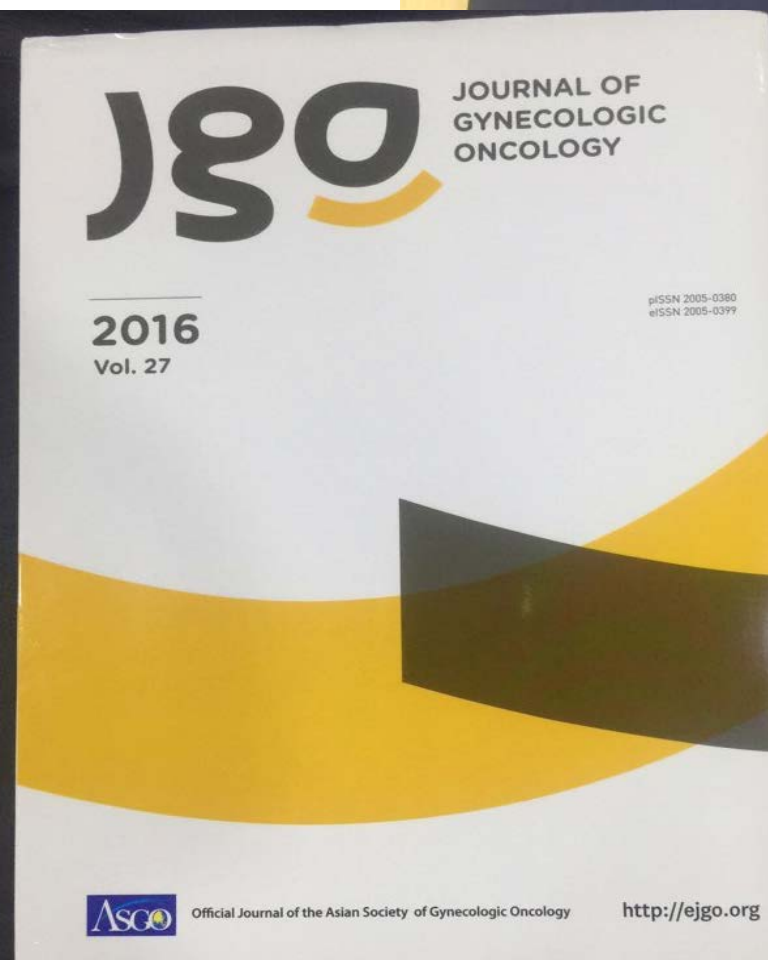
Table 4 summarizes the clinico-pathological and PET/CT derived characteristics of patients without and with recurrence. There were significant differences of PFS, MTV_{total}, TLG_{total}, SUV_{total}, and IFH between patients with and without recurrence. There was significant difference ($p=0.047$) between the mean IFH values of non-recurrent and recurrent groups (**Fig. 3**).

Table 3. Multivariate analyses of prognostic factors for progression-free survival

Variable	HR	95% CI	p-value
IFH	756.997	2.047-279,923.191	0.028
FIGO stage*	1.817	0.418-7.911	0.426
MTV _{total}	0.981	0.946-1.017	0.293
TLG _{total}	1.005	0.999-1.011	0.118
SUV _{total}	0.832	0.637-1.055	0.129
SUV _{total}	1.059	0.958-1.165	0.283

FIGO, International Federation of Gynecology and Obstetrics; HR, hazard ratio; IFH, intratumoral [¹⁸F] fluorodeoxyglucose (FDG) uptake heterogeneity; LN, lymph node; MTV, metabolic tumor value; PFS, progression-free survival; SUV, standardized uptake value; TLG, total lesion glycolysis.

*Test for PFS: I vs. II.



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permanent archiving



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Clin Nutr Res. 2016 Jul;5(3):180-189. English.

Published online Jul 26, 2016. <http://dx.doi.org/10.7762/cnr.2016.5.3.180>

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The Association between Coffee Consumption and Bone Status in Young Adult Males according to Calcium Intake Level

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Abstract

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The purpose of this study was to investigate the association between coffee consumption and bone status (bone mineral density and bone metabolism-related markers) according to calcium intake level in Korean young adult males. Healthy and nonsmoking males (19-26 years, n = 330) participated in this study. Anthropometric measurements, dietary habits, and nutrient intakes were surveyed. Bone status of the calcaneus was measured by using quantitative ultrasound (QUS). Bone metabolism-related markers

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Body Height
Body Weight
Calcaneus
Calcium*
Coffee*
Collagen Type I
Energy Intake
Food Habits
Humans
Male*
Metabolism
Miners
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Conflict of Interest
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ABSTRACT

The purpose of this study was to investigate the association between coffee consumption and bone status (bone mineral density and bone metabolism-related markers) according to calcium intake level in Korean young adult males. Healthy and nonsmoking males (19-26 years, n = 330) participated in this study. Anthropometric measurements, dietary habits, and nutrient intakes were surveyed. Bone status of the calcaneus was measured by using quantitative ultrasound (QUS). Bone metabolism-related markers including serum total alkaline phosphatase activity (TALP), N-mid osteocalcin (OC), and type I collagen C-terminal telopeptide (ICTP) were analyzed. The subjects were divided into two groups based on daily calcium intake level: a calcium-sufficient group (calcium intake $\geq 75\%$ RI, n = 171) and a calcium-deficient group (calcium intake $< 75\%$ RI, n = 159). Each group was then further divided into three subgroups based on daily average coffee consumption: no-coffee, less than one serving of coffee per day, and one or more servings of coffee per day. There were no significant differences in height, body weight, body mass index, energy intake, or calcium intake among the three coffee consumption subgroups. QUS parameters and serum ICTP, TALP, and OC were not significantly different among either the two calcium-intake groups or the three coffee consumption subgroups. Our results may show that current coffee consumption level in Korean young men is not significantly associated with their bone status and metabolism according to the calcium intake level.

Keywords: Coffee consumption; Calcium intake; Bone status; Young adult males

INTRODUCTION

Osteoporosis has been a global public health problem particularly among postmenopausal women, but its prevalence in males is getting increasing [1-3]. Recently, the prevalence of osteoporosis in Korean men aged over 65 years was reported as 15.2%, which was significantly lower than that in Korean women (57.6%), but osteoporosis is also becoming a major disease in aged men [4].

Bone mass increases primarily during the growth period, and peak bone mass is generally completed in between age 18 and 30. It has been reported that peak bone mass is one of important determinants of osteoporosis [5-7]. Genetic factors are the strongest determinants

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High ApoB & the Increased Risk of Newly-onset Diabetes

Table 3. Multiple stepwise regression analysis to find major contributors to the increase of fasting blood glucose level during the short period (3 months)

Variables	Standardized b-coefficients	p value	R	p value
Model 1				
ΔApoB	0.315	0.002	0.315	0.002
Model 2				
ΔApoB	0.281	0.005	0.382	0.001
ΔWaist circumference	0.219	0.029		

Dependent variable: changed levels of fasting blood glucose; independent variables: changed levels of body mass index, waist circumference, systolic blood pressure, diastolic blood pressure, triglyceride, total cholesterol, HDL-cholesterol, LDL-cholesterol, non-HDL-cholesterol, ApoB, ApoA and adiponectin; Δ, changed levels between at baseline and after 3-month.

as major contributors to the net change of glucose levels. In model 1, increased ApoB levels contributed to the increase of fasting glucose levels (standardized b-coefficient: 0.315, $p = 0.002$). In model 2, increased levels of ApoB (standardized b-coefficient: 0.281, $p = 0.005$) and waist circumference (standardized b-coefficient: 0.219, $p = 0.029$) contributed to the increase of fasting glucose levels, respectively ($r = 0.382$, $p = 0.001$).

DISCUSSION

This study suggested that serum ApoB levels may be closely associated with the increased risk of diabetes in Korean men. In our study, subjects with higher ApoB levels (≥ 87.0 mg/dL) showed approximately 2 times higher risk of newly-onset diabetes than those with lower ApoB levels (< 87.0 mg/dL) even after the adjustment for age, BMI, BP, TG, HDL-C, LDL-C, non-HDL-C, ApoA and adiponectin. In addition, multiple stepwise regression analysis revealed that changed ApoB levels were a main contributor to the changed glucose level.

Recent studies have reported that ApoB levels were associated with diabetes [8,12,21,22]; for example, plasma ApoB levels was associated with the incidence of type 2 diabetes, and the risk of diabetes might be better predicted by ApoB levels than by LDL-C or HDL-C. These reports may suggest the potential for the use of ApoB in the assessment for the risk of type 2 diabetes [21,22]. In a large Finnish cohort studies, Salomaa et al. [12] found that ApoB and adiponectin among the screened 31 novel biomarkers, were the best biomarkers for the prediction of the risk of diabetes incidence. On the other hand, Davidson [5] demonstrated that the predictive value of the ApoB/ApoA1 might be stronger and better than the use of either apolipoprotein alone or together. According to Harper and Jacobson [11], the levels of non-HDL-C strongly correlated with the levels of ApoB, and both of non-HDL-C and ApoB rather than LDL-C, may better predict overall cardiovascular risk. Non-HDL-C is calculated by subtracting HDL-cholesterol from total cholesterol, and it reflects circulating levels of the atherogenic ApoB-containing lipoproteins including LDL-C, VLDL, intermediate density lipoprotein cholesterol, chylomicron remnants, and lipoprotein [11,23]. Non-HDL-C has as a major advantage in aspect that it can be calculated by all lipid profiles, and it measures all the ApoB-containing lipoproteins that are considered atherogenic [23].

In our study, serum ApoB levels turned out to be a good predictor for the risk of newly-onset diabetes even after the adjustment. In addition, net change of ApoB levels together with that of waist circumference were found as main influencing factors on the net change of glucose levels. It may be related that an absolute value of ApoB indicates the particle number of LDL as well as represents approximate LDL size thus, the measurement of ApoB levels may better reflect diabetic status, particularly newly-onset diabetes [6,13].

High ApoB & the Increased Risk of Newly-onset Diabetes

The present investigation may have several limitations. First, the study design was rather based on cross-sectional observation, not on a case-control design, because study participants (NFG, IFG and newly-onset diabetes) were classified by screening their fasting glucose levels even though some people were in the borderline of criteria had re-examination. Second, the period of follow-up observation (3 months) was so short, and the followed-up sample size was small. Thirds, study subjects were exclusively Korean men, thus the results may not be applicable to women or other ethnic samples whose clinical and biochemical characteristics may differ from those in our population. Thus, it is necessary to investigate the relationship between ApoB levels and the risk of newly-onset diabetes in other ethnic population as well as much larger population including women. In addition, prospective longitudinal observation is needed in order to investigate if ApoB levels can be used as a better prognostic marker for the risk of newly-onset diabetes and diabetic complication.

CONCLUSION

This study shows that ApoB levels are closely associated with the increased risk of newly-onset diabetes in Korean men, which suggests a possibility of serum ApoB levels as an early predictor for the risk of diabetes.

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The Intra- and Inter-rater Reliability and the Learning Curve for a Simple Neurological Score for Rats

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Abstract

Objective

To measure the intra- and inter-rater reliability of a simple sensorimotor performance test for rats, and to

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Motor Activity
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Rats, Sprague-Dawley
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ABSTRACT

Objective: To measure the intra- and inter-rater reliability of a simple sensorimotor performance test for rats, and to evaluate the learning efficiency of a novice rater for the test.

Method: Middle cerebral arteries were occluded by intraluminal sutures in 25 male Sprague-Dawley rats (10–12 weeks old). The sensorimotor performance test was performed by a novice and an experienced rater, with each rater performing the test twice each week for 3 consecutive weeks. A ten-minute standardized video about the rating method was shown to the novice rater after the second test each week.

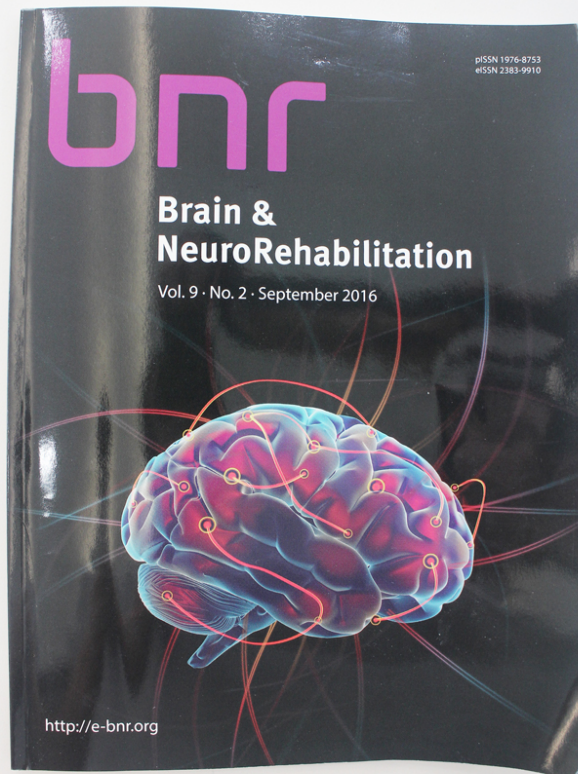
Results: The intra- and inter-rater agreement was determined using Cohen's weighted kappa coefficient. The intra-rater reliability was initially poor for the novice (κ [95% confidence interval], 0.31[-0.02, 0.64]), but it improved significantly after 3-week self-education using the standardized video (0.81 [0.69, 0.93], showing almost perfect agreement. The reliability of the experienced researcher was good at all times (κ [0.64, 0.76, 0.71, for week 1, 2, 3, respectively], indicating substantial agreement. The inter-rater reliability showed clear improvement after self-education (κ [0.44, 0.69, 0.69, for week 1, 2, 3, respectively]. Although the total sum score was highly reliable, some of the individual items showed lower intra- and inter-rater agreement. However, each rater showed greater within-rater variability for different subtests.

Conclusion: The simple sensorimotor performance test showed high degree of intra- and inter-rater agreement when performed by experienced or properly educated raters. The inaccuracy of the novice was rectified by 3-week self-education using a video.

Keywords: Learning curve; Motor activity; Behavioral research; Reproducibility of results

INTRODUCTION

Both practicality and reliability are essential prerequisites of outcome measures in neuroscience research using laboratory animals. Although the volume of infarct tissue or histological changes such as the number of specific cells and optical density of tissue markers have been frequently used, tracking of these changes longitudinally within the same brain is limited. Therefore, neurological functional tests still serve as the valuable adjuncts even in this golden era of molecular biology.





Thank you!

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