

An eClinical Trial System for Cancer that Integrates with Clinical Pathways and Electronic Medical Records

Running head: A New Integrated eClinical Trial System

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Abstract

Background Various information technologies currently are used to improve the efficiency of clinical trials. However, electronic medical records (EMRs) are not yet linked to the electronic data capture (EDC) system. Therefore, the data must be extracted from medical records and transcribed to the EDC system. Clinical pathways are planned process patterns that are used in routine clinical practice and are easily applicable to the medical care and evaluation defined in a trial protocol. However, few clinical pathways are intended to increase the efficiency of clinical trials.

Purpose Our purpose is to describe the design and development of a new clinical trial process model that enables the primary use of EMRs in clinical trials by integrating clinical pathways and EMRs.

Methods We designed a new clinical trial model that uses EMR data directly in clinical trials and developed a system to follow this model. We applied the system to an investigator-initiated clinical trial and examined whether all data were extracted correctly. At the protocol development stage, our model measures endpoints based on

clinical pathways with the same diagnosis. Next, medical record descriptions and the format of the statistical data are defined. According to these observations, screens to entry data which is used for both in clinical practice and for study are prepared into EMRs with an EMR template, and screens are prepared for data checks on our EMR retrieval system (ERS). In an actual trial, patients are registered and randomly assigned to a protocol treatment. The protocol treatment is executed according to clinical pathways, and the data are recorded to EMRs using EMR templates. The data are checked by a local data manager using reports created by the ERS. After edit checks and corrections, the data are extracted by the ERS, archived in portable document format (PDF) with an electronic signature, and transferred in comma separated values format (CSV) to a coordinating centre. At the coordinating centre, the data are checked, integrated, and made available for statistical analysis.

Results We verified that the data could be extracted correctly and found no unexpected problems.

Limitation To execute clinical trials in our system, the EMR template and efficient EMR retrieval systems are required. Additionally, to execute multi-institutional clinical

trials, it is necessary to create templates appropriate for EMRs at all participating sites and for the coordinating centre to validate local templates and procedures.

Conclusion We proposed and pilot tested a new eClinical trial model. Because our model is integrated with routine documentation of clinical practice and clinical trials, redundant data entries were avoided and the burden on the investigator was minimised. The reengineering of the clinical trial process would facilitate the establishment of evidence in the future.

Keywords

clinical research informatics, electronic data capture, CDISC, EMR retrieval, template, data warehouse

BACKGROUND

To improve the efficiency of clinical trials, evaluating and restructuring the information-capture process and the flow of data are critical [1-3]. In many clinical trials, paper-based case report forms (CRFs) are developed for each clinical trial protocol and information is transcribed from source documents, including medical records, to CRFs and from CRFs to the clinical data management system (CDMS). Thus, mistranscriptions and inconsistencies are inevitable in this dual transcription process [2, 4]. Various information technologies currently are used to improve the efficiency of clinical trials, including computerised registration systems, CDMS, and electronic data capture (EDC) systems, among others [2, 4-5]. Elsewhere, medical information technology recently has been promoted. The electronic medical record (EMR) system has been launched in approximately 20% of all institutions in Japan. However, EMRs are not yet linked to the EDC system for clinical research; therefore, the clinical trial staff must manually select the necessary information from EMRs and transcribe it into the EDC system [2]. Thus, the workload at each participating site has not changed with the transition from paper-based CRFs to EDC systems, and there has been no breakthrough in efficiency [2].

The president of the Clinical Data Interchange Standards Consortium (CDISC), Rebecca Daniels Kush, named the next generation of clinical trials ‘eClinical trials’ [3]. She emphasized the importance of optimising clinical trials by redesigning the trial process to utilise primarily electronic processes and to coordinate hospital information systems with the various clinical trials systems [3]. In addition, the CDISC’s Electronic Source Data Interchange (eSDI) initiative, encouraged by the US Food and Drug Administration, was started in 2004 [4]. The eSDI detailed the regulatory requirements and outlined promising scenarios concerning clinical trials using electronic source data (eSource), such as electronic subject diaries, electronic laboratory results, and EMRs. To enhance clinical trials utilising eSource, the eSDI analysed the present regulatory requirements, compiled 12 indispensable requirements, outlined five scenarios that would fit the regulatory requirements, and formed a proposal.

Currently, most clinical trials including standards-based approaches that use data from EMRs are planned according to the concept that the primary use of EMRs is in clinical practice and a secondary use is clinical research [6-11]. To use EMR data secondarily in a clinical trial, it is necessary to obtain various types of information that are stored in EMRs by category, such as the diagnosis and medications taken. However, extraction of

this information is not facilitated by the current EMR system [11-13]. Moreover, not all information that is necessary to execute a clinical trial typically is stored in the EMR [6, 10-12]. If EMRs are used only for the purpose of collecting certain receipts of claims and the treatments for each patient, enough information may be stored in the current EMRs. However, if there is a secondary intention to use the accumulated information from EMRs in clinical research, then the data that are not gathered specifically for research purposes may be incomplete and unreliable [6, 11-12]. To improve the efficiency of clinical trials, it is necessary to establish a method that utilises EMRs directly in clinical trials and enables higher levels of information-sharing and exchange between clinical practice and clinical trials [1-3].

Clinical pathways have been introduced in clinical practice [14, 15]. Clinical pathways are planned process patterns aimed at improving both process quality and resource usage [14, 15]. Clinical pathways include components such as a timeline, the categories of care or activities, and the interventions for specified groups of patients with a particular diagnosis. They are widely used as a treatment plan or as a substitute for physicians' orders [14, 15]. Clinical pathways appear to be easily applicable to the protocol treatment of a clinical trial because they deliver a standard of care to patients with a specific disease

[14, 15]. However, there are few clinical pathways that purport to improve the efficiency of clinical trials.

PURPOSE

Our purpose is to develop a new clinical trial process model that enables the primary use of EMRs in a clinical trial by integrating clinical pathways and EMRs. We report herein on a pilot test of our model in a single-centre clinical trial conducted at our institution.

METHODS

Methods and outline of our model of the clinical trial process

We modelled a new clinical trial process in which practical medical records are used directly in clinical trials and developed a clinical trial system to follow our model. We applied the new system to an investigator-initiated clinical trial and examined whether all of the data needed to conduct the clinical trial were extracted correctly throughout the system. Moreover, by comparing the new model with the conventional method of

conducting clinical trials, we made a qualitative observation of how the new model reduces the workload in clinical trials.

In our model, a clinical trial is performed as follows (see also Figure 1):

1. In the protocol development stage, the methods of measuring endpoints (i.e., protocol treatment and evaluation items) are decided based on the clinical pathways that usually are used for patients with the same diagnosis. Next, medical record descriptions are defined, including eligibility criteria, tests, protocol treatment, efficacy of treatment, adverse events and follow-up. At the same time, the data format and coding conventions for the statistical data are defined to be compatible with statistical software. For example, the description of adverse events is synchronised and standardised with clinical practice and clinical trials according to the Common Terminology Criteria for Adverse Events (CTCAE) [16].

2. According to the defined descriptions, screens are prepared for entry of the additional data required for the clinical trial into EMRs using EMR templates, and screens are prepared for data checks and data extraction by our EMR retrieval system (ERS). The templates are named to enable clinicians to identify them as pertaining only to the trial.

3. Patients who provide informed consent and agree to participate are registered and randomly assigned to one of the protocol treatments.
4. The protocol treatment assigned to the patient is executed according to clinical pathways, and the treatment data are recorded to EMRs using EMR templates.
5. Data are extracted using the ERS and are checked automatically for completeness and consistency and to identify anomalies to be resolved.
6. After edit checks have been satisfied following corrections, the data are extracted by the ERS, archived in portable document format (PDF) with an electronic signature, and transferred in comma separated values format (CSV) to a coordinating centre.
7. At the coordinating centre, the clinical trial data from participating sites are checked, integrated, and made available for statistical analysis.

When it is necessary to update trial data that already have been transferred to the coordinating centre, the process repeats from when the investigator updates the medical records using the clinical trials template. In addition, central pathology reviews or central laboratory findings are integrated with clinical data in the coordinating centre.

Because our new model is integrated, with the routine documentation of clinical practice and the procedures of clinical trials, to avoid redundant data entries, the efficiency of clinical trials can be improved.

Applied information technology

EMR templates

To record clinical trial data to EMRs that could be available for clinical practice, we used EMR template technology [14, 17-21].

In the current EMR system in Japan, various types of information are integrated, including coded data. Coded data include laboratory results that are in the computerised physician order entry system, narrative information, such as observations during physical examinations or progress notes that are written in free-text form, radiological images, and waveform information from electrocardiograms. In particular, it was necessary to convert free-text information to structured information that is expressed by data elements consisting of items and values with codes for use in a clinical trial [17]. The most practical method is to use a template for entering data into the EMR system

[17]. Using templates, the entered narrative information can be made available for data analysis.

In Japan, the EMR template technology already has been implemented for most large hospitals' EMR systems, including ours, and is widely used to standardise medical records. Although multiple vendors supply various templates, when the new information is recorded, our hospital stores key data, such as the template ID, item ID, and entry date and time, in the EMR database. Moreover, our EMR system is managed according to the Japanese Ministry of Health, Labour and Welfare's guidelines on hospital information system management. To guarantee data integrity, all modifications to EMR data are recorded when and by whom each operation is executed, and an audit trail is made automatically. Audit trails record whose, when, and by whom medical records have been accessed. No user can update audit trails or prevent recording them. In addition, after medical records are submitted by medical staff, no user can delete the records. When medical records are updated, the new data are created as updated records, and the old data are retained as deleted records with a 'deletion flag'. This traceability management function is applied to the clinical trial template; all clinical trial data and histories are stored on the EMR system.

EMR retrieval system

We use the ERS that was developed in our translational research centre to extract data from EMRs, verify the completeness and consistency of trial data, and create output data for statistical analysis. The ERS can retrieve data from patients participating in the clinical trial comprehensively and efficiently. To retrieve and report the records in clinical practice directly as data for clinical research, we identified entities from EMRs that are useful for clinical research (i.e., data on patient demographics, diagnosis, physical examination, progress notes recorded in the EMR template, operative notes, laboratory tests, radiological or pathological studies, medications and injections, and other treatments). Next, we designed a multidimensional data model for EMR retrieval for clinical research [11, 22-25]. To ensure that the data retrieval process is practical and independent from the structure of the EMR system, a data warehouse was created by extracting, transforming, and loading information from the EMR system. To retrieve and report information from multiple patients efficiently, an online analytical processing (OLAP) tool was installed [26]. The OLAP tool runs from an Internet browser and has the ability, in hypertext markup language (HTML), to report on information retrieved

from the browser. The reports are created in various formats, such as PDF, CSV, and extensible markup language (XML). The ERS also was applied to the data recorded using the new template. We created reports for checking trial data and extracting data for statistical analysis.

Application to a clinical trial

We applied the system to a single-institution phase II trial for hepatocellular carcinoma.

(‘A randomised controlled study of effectiveness between transcatheter arterial chemoembolisation (TACE) with cisplatin and TACE with epirubicin for multiple hepatocellular carcinomas’). This study was approved by the ethics committee of Kyoto University Hospital and was registered in the UMIN Clinical Trials Registry [27] (registration number UMIN000003162).

Based on the findings from an observational study in our hospital [28, 29], this phase II trial was designed to confirm the effectiveness of TACE in combination with one of the two anticancer agents. A total of 160 subjects will be enrolled, treated by a single TACE therapy, and evaluated for 12 months. Including laboratory tests, the number of data

items per subject is approximately 80, and the number of total records per subject is approximately 15 (Table 1).

Procedures of the clinical trial

Preparation

In the pilot test of our clinical trial process model, we prepared seven templates:

Eligibility Criteria, Target Lesion, Protocol Treatment, Efficacy of Treatment based on the European Association for the Study of the Liver (EASL) standard [30], Efficacy of Treatment based on the Response Evaluation Criteria in Cancer of the Liver (RECICL) standard [31], Adverse Events based on CTCAE 4.0, and Follow-Up (Table 1). Codes and grades (i.e., EASL, RECIEL, CTCAE) were assigned by the investigators when they entered these data into the EMR using the clinical trial templates. Three types of ERS reports were used to check the clinical trial data: a List of Cases, each patient's Case Registration Form, and the Case Report Form (Table 1). We show an example of the EMR template and progress notes from the EMR in Figure 2.

For the local data manager to verify the completeness and consistency of trial data, the Case Report Form can be identified. The Case Report Form summarises all the templates on one screen such that the integrity of all information for each patient can be assessed at a glance. The latest data for Eligibility Criteria, Target Lesion, Protocol Treatment, Efficacy of Treatment, and Follow-Up are displayed by item; Adverse Events are shown in sequence according to the name of the adverse event so that changes in the adverse event grade can be observed (Figure 3).

Patient registration and random assignment of treatment arm

Patients were registered and assigned randomly to a treatment arm in an independent registration centre outside of the EMR system. The investigator received the registration and assignment information and recorded it on the Eligibility Criteria template in the EMRs.

Recording clinical trial data in EMRs

The investigator conducted the protocol treatment and evaluations according to the clinical pathways and recorded the results in templates. The results of required laboratory tests were extracted directly from the EMRs.

Correction of data

Clinical trials often require local-level administrative help with the increased workload [5, 32], but our model provides the local data manager with new responsibilities to ensure trial data quality. In the pilot study, a local data manager—a clinician at the participating site—manually checked the trial data for completeness and consistency using the Case Report Form on the ERS. When the local data manager found incomplete or inconsistent trial data, he asked the investigator to determine whether data should be corrected. The investigator added necessary data using the original EMR templates, thus storing a revision history in the EMRs.

In the pilot study, the data in the EMR system create the edit checks and format the data for statistical analysis; thus a separate EDC system or the CDMS is unnecessary. We executed data checks manually to ensure that the latest medical records were correct.

Extracting data for statistical analysis

We used the ERS to extract data for the statistical analyses according to predefined data formatting and coding conventions. After the data were presented for statistical analysis in the CSV by the ERS, reports of all the data were prepared and archived in PDFs with an electronic signature added to prevent manipulation.

RESULTS

We applied our new system to an actual clinical trial and evaluated its feasibility.

Using the list of registered patients in the registration centre, we determined whether we could extract both the data by templates and the laboratory results of the registered patients in pre-designed forms. We verified that the data could be extracted correctly and found no unexpected problems.

Regarding the clinical staff workload, the investigators did not need to complete CRFs or enter data into an EDC system in addition to the EMRs. Among the prepared templates for the clinical trial, the only one that was added to the pilot study was Eligibility Criteria for patient registration. The investigators had to record information in the medical records for TACE, including the number of tumours, their diameter and extent of vascular invasion, and the therapeutic regimen, such as anticancer drug doses and embolised arteries in the liver. To record the necessary clinical information on a patient's EMR, investigators used the templates based on the clinical pathways for TACE, and they checked the results of laboratory tests shown in the EMRs.

Regarding the data quality control, as the data in the EMR templates were identical to those in the ERS, the use of the EDC system or the CDMS was also unnecessary. When investigators added information to EMRs, local data managers viewed reports in the Case Report Form automatically on the ERS. Local data managers check the data until all of the errors are corrected. In this way, the progress of the clinical trial and the situation for each patient could be assessed at a glance.

Figure 4 shows the procedural differences among a paper-based clinical trial, a clinical trial using the EDC system, and our new model. In contrast with conventional clinical trials, our model made it unnecessary to transcribe data from medical records to CRFs, to execute source data verification (SDV) at the participating site, to transport paper CRFs from the participating site to the coordinating centre, or to make double entries in the CDMS from paper CRFs. In addition, data checking was performed by the local data manager at the participating site, and supplemental data checking and data integration were performed at the central coordinating centre.

LIMITATIONS

To execute clinical trials according to our model, specialised data must be accumulated from EMRs according to each clinical trial protocol. From a technical viewpoint, equipment for the flexible input functions, such as the EMR template, and efficient EMR retrieval systems are required. Consequently, trials must be planned from the stage of protocol development. The pilot test trial was planned to answer the practical questions that clinicians face in clinical practice. If a clinical trial is not practice-based [33-35], then it will not be easy to add medical records using a template. In our model,

the coordinating centre workload for data management and quality monitoring would be reduced. Instead, the workload of the local data manager at the participating site would be increased. In addition, in the test trial, every time errors were found in trial data using the Case Report Form on the ERS, the local data manager had to communicate with the investigator. In the future, more efficient local data-management methods will have to be established [36].

Because the test trial used for the pilot study was executed in a single institution, it was not necessary to integrate the clinical trial data from multiple participating sites. We used the CSV format to present data for statistical analysis and the PDF format to archive the trial data at the participating site. To execute multi-institutional clinical trials, it will be necessary to standardise the descriptions in EMRs, use evidence-based clinical pathways, and create and validate templates across EMRs at all participating sites [37]. Currently, the accuracy of the information in EMRs may vary across sites [12]. For example, there are standard medical terminologies, such as ICD9, ICD10 [38], MedDRA [39], and SNOMED [40], but diagnosis or medications are associated with the payment of medical insurance, and use of classification schemas and terminology are at the discretion of clinicians at each site. Thus, the central coordinating centre will

need to monitor and standardise the use of EMRs and integrate data provided across sites. In addition, a vendor-neutral and platform-independent standard format, such as the CDISC Operational Data Model (ODM) [41] and the CDISC Study Data Tabulation Model (SDTM) [42], will be needed to archive, transfer, and integrate trial data. The advantage is that the EMR system would not need to be retained in the future to access the data or the audit trail [4].

CONCLUSION

We proposed a new clinical trial process model by integrating it with clinical pathways and EMRs, which is the primary use of EMRs in a clinical trial. In a single-centre trial selected for the pilot study, our new model was integrated with the routine documentation of clinical practice and the procedures of clinical trials. Therefore, redundant data entries for the trial were avoided, and the burden on the investigators posed by the trial was minimised.

The basis of our model is that the standardisation of clinical practice from clinical pathways can be applied as a treatment plan for a clinical trial; one primary use of the

EMR system could be to provide data for clinical trials. The most prominent characteristic of our model is that many data management tasks that typically are performed by the central coordinating centre are moved to the participating sites. In effect, the EMRs replace the EDC system and the CDMS. To maintain efficient data quality control, it is important to execute data management near the site of the data occurrence. The earlier an error is detected in a clinical trial, the sooner and more easily and cheaply it can be rectified [3]. Conventionally, the period from when an investigator executes protocol treatment to when the trial data are transcribed in the CRFs and checked by the data manager of the central coordinating centre spans several days (e.g., mean 5 days ranging 1 to 25 in our centre). In contrast, when investigators added information to the EMRs in the test trial, the data were immediately available to the local data manager for checking, which allowed rapid resolution of any anomalies introduced. Such real-time availability of trial data should reduce the time and cost of conducting clinical trials.

In addition, in the eSDI proposal, the third scenario was called ‘Single Source Concept [4],: data need to be entered only once for multiple purposes (research, patient care, safety surveillance) within the context of existing regulations. The fourth scenario was

called ‘EMR Extraction and Investigator Verification’: a clinician would check EMRs for the necessary data for the clinical trial and extract and transfer the data to a sponsor. We believe that our new model fits these scenarios and that we have demonstrated an implementation model for eClinical trials based on the eSDI proposal. According to the eSDI scenario, Title 21 Part 11 of the Code of Federal Regulations [43] starts at the point of the creation of a clinical research record. In other words, it is not necessary to apply these regulations to the medical records at the site. The clinical research record is created at the point of electronic signature signing. When data are extracted from EMRs as clinical trial data, the migration of the data from EMRs to the clinical trial database must be validated at the participating site. In addition, the sponsor has to produce documents on how the procedures of the clinical trial follow all appropriate regulations. Because of the direct extraction of the clinical trial data from EMRs, our model may eliminate or minimize the need for source data verification. The audit plan should focus on the accuracy of the migration of data from EMRs to ensure that data are not changed in the extraction process and that patient confidentiality requirements are met [4]. We believe that by appropriating the efforts of source data verification in favour of validating the clinical trial process, the costs can be reduced and the efficiency of clinical trials can be improved.

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Table 1. The Screen to Record Data for EMRs and the Data Check Screen.

Screen to Record Data for EMRs		Edit Check Screen
Template Name	Item Name	
Eligibility Criteria	Informed consent	List of Cases Case Registration Form
	Registration Number	
	Date of Consent	
	Result of Allocation	
	Selection Criteria 1–8	
	Exclusion Criteria 1–10	
Target Lesion	Number	Case Report Form
	Tumour Location	
	Size (max)	
	Incipient TACE/non-Incipient TACE	
	Vascular Invasion	
	Frequency of TACE	
Protocol Treatment	Date of Protocol Treatment	
	Size (max)	
	Total Dose	
	Number	
	Total Dose for TACE	
	Tumour Location	
	Addition of RFA/PEIT	
	Anticancer Drug	
	Infusional Therapy	
Efficacy of Treatment (EASL) Efficacy of Treatment (RECICL)	Efficacy of Treatment	
	Recurrence Form	
	Plan of Treatment	
	Other	
Adverse Event	Abdominal Pain	
	Cause of Abdominal Pain	
	Diarrhoea	
	Cause of Diarrhoea	

	Nausea	
	Cause of Nausea	
	Vomiting	
	Cause of Vomiting	
	Fever	
	Cause of Fever	
	Performance Status	
	Other	
Follow-Up	Date of Progressive Disease	
	Recurrence Form	
	Treatment after Recurrence	
	Conform to Protocol Treatment	
	Reason of Change	
	Date of Death	

EASL: European Association for the Study of the Liver; EMR: electronic medical record; PEIT: percutaneous ethanol injection therapy; RECICL: Response Evaluation Criteria in Cancer of the Liver; RFA: radiofrequency ablation; TACE: transcatheter arterial chemoembolisation.

Figure captions

Figure 1.

Our new model of the clinical trial process.

Figure 2.

Electronic medical record (EMR) and the EMR template. The template is opened from within our EMR system. The template converts narrative information from progress notes, thereby making the information available as analytical data.

Figure 3.

The List of Cases and the Case Report Form. The Case Report Form can be consulted from the List of Case Registration Forms. The Case Report Form summarises all the templates on one screen so that all information can be assessed at a glance. EASL: European Association for the Study of the Liver; EMR: electronic medical record; RECICL: Response Evaluation Criteria in Cancer of the Liver.

Figure 4.

The pilot study revealed many advantages over a conventional clinical trial process, i.e., unnecessary to transfer information from medical records to the case report form, no need for source data verification at the participating site, unnecessary to transmit the case report form from the participating site to the coordinating centre, and unnecessary

to re-enter data into the clinical data management system (CDMS) from the paper case report form. In addition, data management in the coordinating centre was replaced with edit checks initiated by a local data manager at the participating site. CRF: case report form; CDMS: clinical data management system; DWH: data warehouse.

Figure 1.

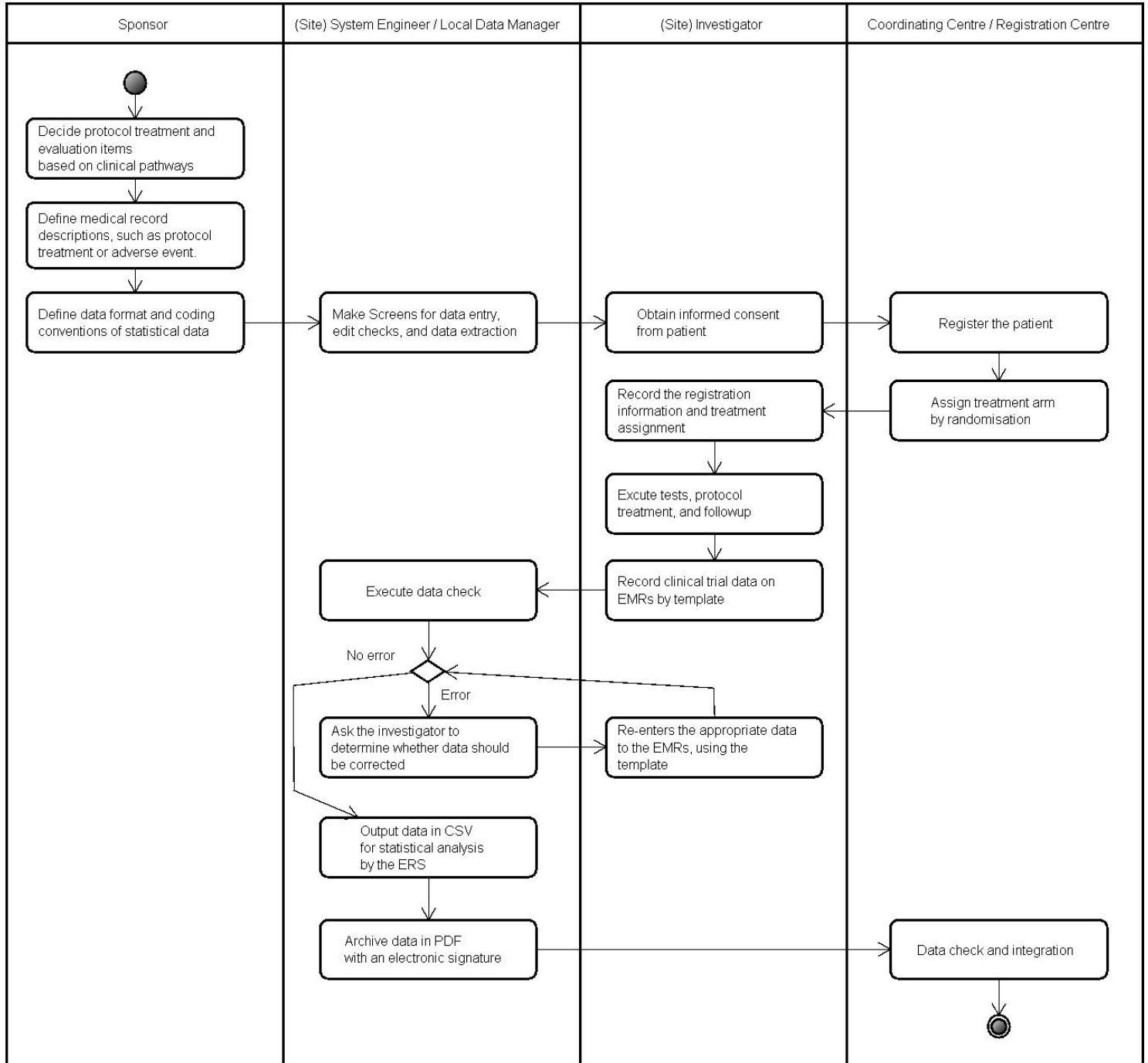


Figure 2.

EMR

京都大学医学部附属病院 [メイン画面]

加テ 看護 ケア 指示 記録 画像 検査 複合 診療 (1) 診療 (2) 部門 個人 診療記録

1933年05月28日生 77歳11ヶ月 女 病棟: 他 担当 血液型:

ナビ | 履歴 | 画像 | 加テ | 病名歴 | スキヤ履歴 | 他病院履歴 | 文書 | 入院歴 | 手

絞り込み条件
プロブレム
全て 全科 全て
カルテ
全て 全科 入外共通 保険指定 全て

詳細を表示 履歴で表示 表示条件設定 付箋一覧 印刷

▲保険情報: 【主】政府【公】精32

D#0 2010-10-12 14:11 肝臓移 外来 医師

F 【治療前情報】
再発治療・直近の治療: 手術,TACE
病巣1 部位 S: 2,4

Template of Target-Lesion

- Number
- Tumor Location
- Size(max)
- Incipient TACE/ non Incipient TACE
- Vascular Invasion
- Frequency of TACE

テンプレート入力

テンプレート名 【TACE臨床試験】標的病変

個数 ☒ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5以上

占拠部位 ☐ S1 ☒ S2 ☐ S3 ☐ S4 ☐ S5 ☐ S6 ☐ S7 ☐ S8

大きさ(最大) ☒ 2cm以下 ☐ 2cm~5cm ☐ 5cm以上

初発・非初発TACE ☒ 初発 ☐ 非初発

脈管浸潤 ☒ なし ☐ 門脈 ☐ 静脈 ☐ 胆管

TACE回数 1 回

OK キャンセル 計算

検査所見

A

P

F

【【TACE臨床試験】標的病変】

個数: 1
占拠部位: S2
大きさ(最大): 2cm以下
初発・非初発TACE: 初発
脈管浸潤: なし
TACE回数: 1 回

Progress Notes of EMR

<< 先頭へ < 前へ 次へ > 直近へ >> 検索 拡張表示

05/02 (月) 15:27

Figure 3.

The screenshot displays a web application interface for managing clinical trial data. The main window is titled "1.TACE試験_登録票_作成" (1.TACE Trial Registration Form Creation). It features a "List of Cases" table and a "Case Report Form for Individual Subject" for a selected subject.

List of Cases Table:

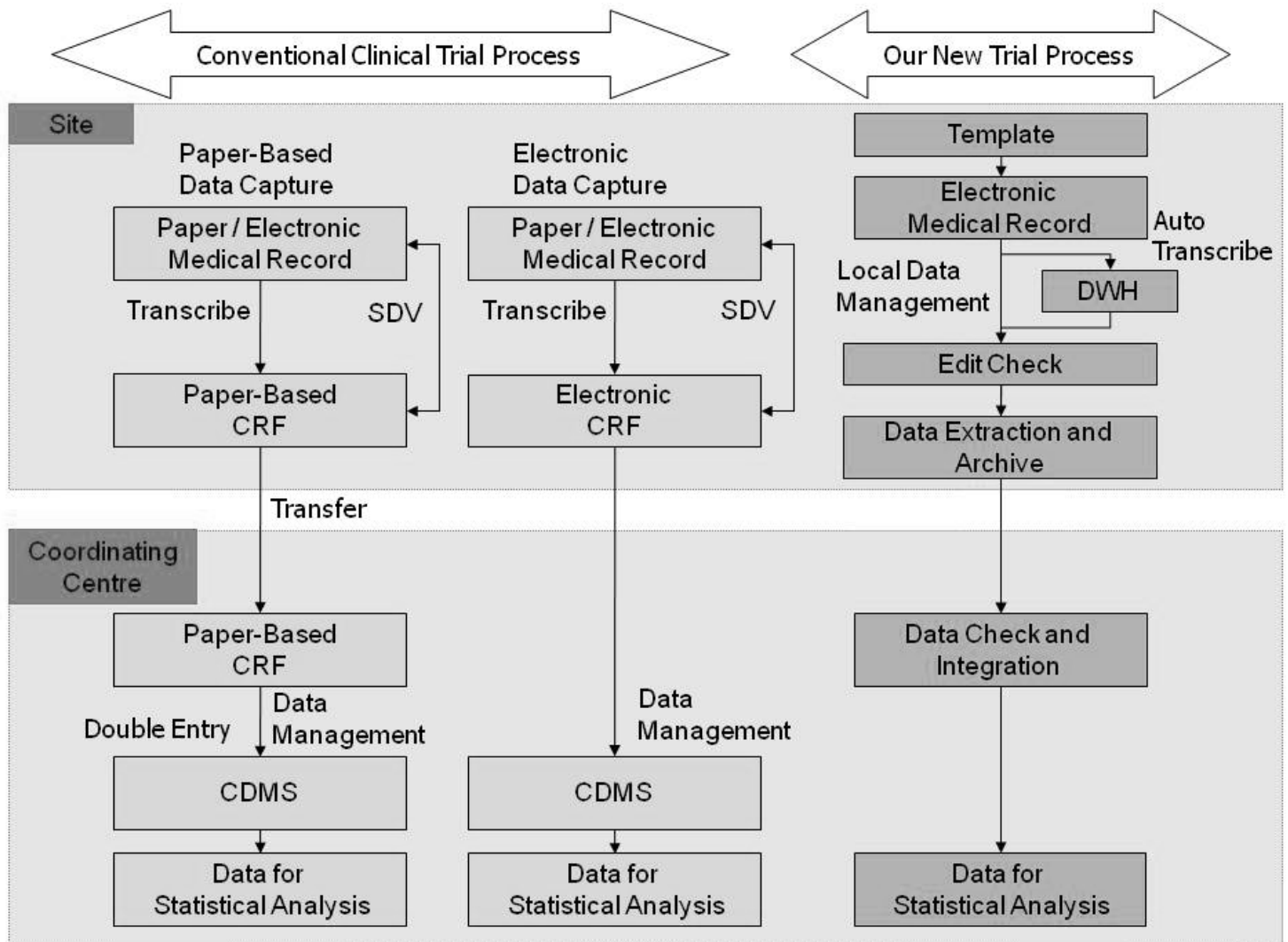
登録番号	患者ID	同意取得日	表示テキスト
TT-001		2010-09-14	臨床試験参加: 不参加
TT-002		2010-10-14	臨床試験参加: 参加
TT-004		2010-11-08	臨床試験参加: 参加
TT-005		2010-11-09	臨床試験参加: 参加
TT-006		2010-11-10	臨床試験参加: 参加
TT-007		2010-11-24	臨床試験参加: 参加
TT-008		2010-12-06	臨床試験参加: 参加
TT-009		2010-12-14	臨床試験参加: 参加
TT-010		2010-12-20	臨床試験参加: 参加
TT-011		2011-01-05	臨床試験参加: 参加
TT-012		2011-01-17	臨床試験参加: 参加
TT-013		2011-01-19	臨床試験参加: 参加
TT-014		2011-01-24	臨床試験参加: 参加
TT-015		2011-02-01	臨床試験参加: 参加
TT-016		2011-02-02	臨床試験参加: 参加
TT-017		2011-02-07	臨床試験参加: 参加
TT-018		2011-02-08	臨床試験参加: 参加
TT-019		2011-03-07	臨床試験参加: 参加

The "Case Report Form for Individual Subject" is displayed for subject ID "TT-017". It includes a "症例報告書" (Case Report Form) section with a table of vital signs and clinical data.

Case Report Form Table:

項目	値	単位	発生日時	検査項目
1	2011/05/10		検査項目: 0	
2	2011/05/10		検査項目: 1	
3	2011/05/10		検査項目: 2	
4	2011/05/10		検査項目: 3	
5	2011/05/10		検査項目: 4	
6	2011/05/10		検査項目: 5	
7	2011/05/10		検査項目: 6	
8	2011/05/10		検査項目: 7	
9	2011/05/10		検査項目: 8	
10	2011/05/10		検査項目: 9	
11	2011/05/10		検査項目: 10	
12	2011/05/10		検査項目: 11	
13	2011/05/10		検査項目: 12	
14	2011/05/10		検査項目: 13	
15	2011/05/10		検査項目: 14	
16	2011/05/10		検査項目: 15	
17	2011/05/10		検査項目: 16	
18	2011/05/10		検査項目: 17	
19	2011/05/10		検査項目: 18	
20	2011/05/10		検査項目: 19	

Figure 4.



List of abbreviations

CDISC: clinical data interchange standards consortium;

CDMS: clinical data management system;

CRF: case report form;

CSV: comma separated values;

CTCAE: common terminology criteria for adverse events;

DWH: data warehouse;

EASL: european association for the study of the liver;

EDC: electronic data capture;

EMR: electronic medical records;

ERS: electronic medical records retrieval system;

eSDI: electronic source data interchange;

eSource: electronic source data;

HTML: hypertext markup language;

ICD: international classification of diseases;

MedDRA: medical dictionary for regulatory activities;

ODM: operational data model;

OLAP: online analytical processing;

PDF: portable document format;

PEIT: percutaneous ethanol injection therapy;

RECICL: response evaluation criteria in cancer of the liver;

RFA: radiofrequency ablation;

SDTM: study data tabulation model;

SDV: source data verification;

SNOMED: systematized nomenclature of medicine;

TACE: transcatheter arterial chemoembolization;

XML: extensible markup language;