

PROCESS MAP OF THE JAIU QUALITY MANAGEMENT SYSTEM

1. Purpose

The QMS Process Map of JAIU establishes the structure of the university's processes, their interrelationships, process owners, key inputs/outputs, performance indicators, and QMS documents that ensure quality management and continuous improvement.

2. QMS Inputs and Outputs (Top Level)

Inputs: regulatory requirements of the Republic of Kazakhstan, accreditation requirements, expectations of interested parties (students, employers, clinical sites), results of monitoring and audits, risks and nonconformities, eBilim data.

Outputs: prepared graduates (competencies), learning and assessment results, quality indicators (KPIs), management reports and decisions, improvement plans (CAPA), supporting QMS records, increased stakeholder satisfaction.

3. Structure of the process model (as shown in the diagram)

The JAIU process model includes 4 levels:

1. **Strategy and Quality** (Management and Leadership)
2. **Core university processes** (education, research and innovation, social/educational)
3. **Support and resources** (human resources, material and technical support, IT, clinical facilities, library, simulation center)
4. **Quality loop and documentation** (monitoring, audit, risks/CAPA, document flow, and records)

4. QMS Process Register (Top Level)

Legend:

M — management processes; **O** — core processes; **P** — support processes; **Q** — quality loop (closed-loop improvement cycle).

4.1 Management processes (M)

| Code | Process | Process Owner | Key Outputs | Key KPIs (examples) | QMS Documents (Reference) |
|-----------|---------------------------------------|-----------------------|--|--|------------------------------------|
| M1 | Strategic Management and Development | Rector's Office | Strategic plan, development plans | Implementation of measures, achievement of strategy KPIs | QMS-01-02 |
| M2 | Quality Management (QMS Coordination) | Quality Council + OQM | Quality policy, decisions, improvement plans | Percentage of CAPAs closed on time, audit plan fulfillment | QMS-01-01, QMS-40-05, QMS-60-03/04 |
| M3 | Risk and Opportunity Management | OVMC | Risk register, response measures | Register validity, percentage of risks with measures | QMS-60-01/02 |

| | | | | | |
|-----------|-----------------------|-------------------|--|--|-----------|
| M4 | QMS Management Review | Quality Committee | MR minutes, decisions, KPI adjustments | MR Frequency, % of Decisions Implemented | QMS-40-05 |
|-----------|-----------------------|-------------------|--|--|-----------|

4.2 Core Processes (O)

| Code | Process | Process Owner | Key Outputs | Key KPIs (examples) | QMS Documents (Reference) |
|-------------|--|---|--|--|----------------------------------|
| 01 | Design, Revision, and Approval of OP/OOP | Academic Department + Methodology Council | Curricula/Educational Programs, Educational Program Descriptions | % of OP revised according to schedule | SMK-20-01/02/03, SMK-20-04/05 |
| 02 | Implementation of the credit-based educational process | UIO + Academic Department + Dean's Office | Schedule, enrollment, academic process | Timeliness of schedules/registers, fulfillment of plans | SMK-20-06, SMK-20-12 |
| 03 | Assessment and Evaluation (ongoing/midterm/final) | Academic Affairs Office + UIO + Departments | Grades, grade sheets, module results | Percentage of grades submitted on time, percentage of outstanding grades | SMK-20-09, SMK-40-06 |
| 04 | Internships, clinical sites, and mentoring | Dean's Office/Internship Department + Departments | Practicum reports, facility agreements | % of sites with agreements/mentors, % of reports submitted on time | SMK-20-11, SMK-30-04 |
| 05 | Simulation training and OSCE/OSPE skills assessment | Simulation Center + Departments | OSCE protocols, checklists, results | % of students who passed OSCE, average score, % of stations without violations | SMK-30-03, SMK-20-14 |
| 06 | Science and Innovation | Science Department | Research, publications, projects | Number of publications/projects, Science KPIs | SMK-80-01...05 |
| 07 | Social and Educational Work | Educational Services | Programs, events, social work KPIs | Participation, satisfaction, KPIs | SMK-90-01...04 |

4.3 Support and Resources (P)

| Code | Process | Process Owner | Key Outputs | KPIs (examples) | QMS Documents (Reference) |
|-------------|---|----------------------|-------------------------------|------------------------|----------------------------------|
| P1 | Staffing and Development of Teaching Staff | HR | Staff, PC, DI | % of positions filled | SMK-10-09, SMK-10-12, SMK-30-08 |
| P2 | IT Infrastructure and Digital Services (eBilim/EDO) | IT | Service availability, reports | Uptime, SLA requests | QMS-30-02, QMS-30-07, QMS-70-04 |

| | | | | | |
|-----------|---|------------|---------------------------------|---------------------------------------|----------------------|
| P3 | Management of training data and documents (records/reports/archive) | UIO | Reports, certificates, archive | Issuance deadlines, data completeness | SMK-10-07, SMK-70-05 |
| P4 | Library and information resources | Library | Collection, access to resources | Attendance, Availability | SMK-30-01 |
| P5 | Logistical Support | AHC | Logistics, logistics records | % of supply, update | SMK-30-05/06 |
| P6 | Occupational Health and Safety | OTiTB/AHC | Training sessions, reports | % of briefings, incidents | QMS-60-07/08/09 |

4.4 Quality System and Documents (Q)

| Code | Process | Process Owner | Key Outputs | KPIs (examples) | QMS Documents (Reference) |
|-------------|--|----------------------|------------------------------------|--|----------------------------------|
| Q1 | Quality Monitoring and Evaluation | OQ | Monitoring reports, questionnaires | % of questionnaires returned, satisfaction index | SMK-40-01/02 |
| Q2 | Internal audit and self-assessment | OVMC | Audit reports, improvement plan | Number of nonconformities, % resolved | QMS-40-04 |
| Q3 | Nonconformity management and CAPA | QMS + process owners | CAPA, corrective actions | % of CAPAs completed on time | QMS-60-03/04 |
| Q4 | Document flow and QMS records (log, version control) | OVMC (+ IT for EDM) | Registry, current versions | % of documents up to date | QMS-70-01/02/03/05 |

5. Linking process maps to QMS documents (summary matrix)

- Strategy/Quality:** QMS-01-01...05, QMS-40-05, QMS-60-01...04
- Training:** QMS-20-01...14 (+ OCCE via QMS-30-03 and QMS-20-14)
- Resources:** QMS-30-01...08, QMS-60-07...09
- Quality loop:** SMK-40-01...08, SMK-70-01...06
- Science:** SMK-80-01...05
- Social/Educational:** SMK-90-01...04
- Ethics:** SMK-50-01...07

6. Note on Updates

The process map is reviewed **at least once a year** and whenever there are changes to the structure of JAIU, accreditation requirements, digital processes (eBilim/EDO), or based on the results of QMS management analysis.

Appendix. Process Card (TEMPLATE)

PROCESS CARD

Process Code: ____

Process Name: _____

Group: M / O / P / Q

Process Owner: _____

Co-executors: _____

Process objective: _____

Process boundaries (start-end): _____

Inputs: _____

Outputs: _____

Key steps (5-10 steps):

1. _____
2. _____
3. _____
4. _____
5. _____

Key Performance Indicators (KPIs) and Target Values:

1. KPI1: _____ (target: __)
2. KPI2: _____ (target: __)
3. KPI3: _____ (target: __)

Process risks (link to risk register):

1. R1: _____ / Mitigation: _____
2. R2: _____ / Mitigation: _____

Records/evidence (what is retained):

1. _____
2. _____

Related QMS documents: _____

Related forms/templates: _____

Monitoring frequency: _____

Review frequency: _____



Appendix. Process Card 02

Process code: 02

Name: Implementation of the educational process using the credit system

Group: 0 (main)

Owner: UIO + Academic Affairs Office

Co-executors: Dean's Office, Departments, IT Service (eBilim), OVMK (Quality Data)

Process objective: To ensure the planning and implementation of the educational process in accordance with the educational programs/curricula, the credit system, and QMS requirements, with full traceability in eBilim.

Boundaries (start-end): from the approval of the curriculum/schedule to the end of the semester and the generation of final grade sheets/reports.

Inputs:

1. Approved educational programs/curricula, syllabi, academic calendar
2. Student body, orders (enrollment/transfer/withdrawal)
3. Faculty workload, resources (classrooms/computer center/facilities)
4. QMS regulations (credit system, eBilim)

Outputs:

1. Schedule, distribution by groups/subgroups
2. Classes held, attendance/activity logs (eBilim)
3. Interim reports (modules), final semester report
4. Performance/risk analytics (for the Academic Council and administration)

Key steps:

1. Entering/updating student enrollment and curricula in eBilim.
2. Creating course loads and schedules.
3. Organizing classes, monitoring attendance/participation.
4. Support for independent study assignments and tasks in the LMS, consultations.
5. Preparation for modular assessments/exams (eligibility checks).
6. Generating grade sheets, recording results, archiving.
7. Transferring data to the OVMK for quality analysis.

KPI:

1. KPI1: Timeliness of schedule approval/publication — target: ___% on time.
2. KPI2: Completeness of eBilim logs — target: ___% of courses/groups.
3. KPI3: Timeliness of generating reports — target: ___% on time.

Risks:

1. R1: Errors in enrollment data/curricula → action: double-check by academic departments + dean's office.
2. R2: eBilim failure/service unavailability → measure: backup procedure + IT SLA.

Records/evidence:

1. Curricula/calendar, schedules, enrollment orders
2. eBilim logs (attendance/activity), ledgers, reports

Related QMS documents: QMS-20-06; QMS-20-12; QMS-10-07; QMS-70-04/05

Forms/templates: schedule template, ledgers, module/semester reports

Monitoring: weekly/monthly; Review: annually.

Appendix. Process Card 03

Process code: 03

Name: Assessment and Evaluation (ongoing/midterm/final)

Group: 0 (main)

Owner: Academic Affairs Office + UIO

Co-executors: Departments/instructors, OVMK, IT (eBilim), Dean's Office

Process objective: To ensure objective, transparent, and consistent assessment of students' knowledge and skills, with recording of results and quality analysis.

Scope: From establishing criteria/FOS to publishing results and analyzing outcomes.

Inputs:

1. FOS, assessment criteria, BRS/modular system
2. Curriculum/syllabi, independent study assignments, tests
3. Group lists/student body, eligibility for assessment

Outputs:

1. Grades by assessment type, test results
2. Module/final grade sheets, protocols (if applicable)
3. Academic performance and risk analytics (arrears, trends)

Key stages:

1. Approval of criteria/FOS and communication to students.
2. Conducting ongoing assessments (assignments, tests, practical exercises).
3. Conducting midterm assessments (Module 1, Module 2).
4. Final assessment (exam/test) based on the approved model.
5. Entering results into eBilim and generating grade sheets.
6. Verification of accuracy, appeals (if any).
7. Analyzing results and initiating CAPA as needed.

KPI:

1. KPI1: Percentage of courses with approved learning outcomes/rubrics — target: ___%.
2. KPI2: Timeliness of entering grades into eBilim — target: ___% on time.
3. KPI3: Percentage of courses with semester-end results analysis — target: ___%.

Risks:

1. R1: Inconsistent grading → measure: standardized rubrics + internal oversight.
2. R2: Violations of academic integrity → measure: proctoring/anti-plagiarism/committee.

Records/evidence:

1. FOS, rubrics, test banks, monitoring reports
2. Registers, minutes, academic performance reports, CAPA decisions

Related QMS documents: QMS-20-08; QMS-20-09; QMS-40-06; QMS-50-01...04

Forms/templates: reports, rubrics, academic performance analysis reports

Monitoring: every module/semester; Review: annually.

Appendix. Process Card 05 (OSCE/OSPE)

Process Code: 05

Title: Simulation-based training and assessment of practical skills (OSCE/OSPE)

Group: 0 (core)

Owner: Simulation Center

Co-implementers: relevant departments, the Dean's Office, clinical mentors (if necessary), OVMK

Process objective: To ensure the development and objective assessment of students' practical competencies at simulation stations using standardized checklists and a transparent procedure.

Scope: From OSCE/OSPE planning to the approval of results and quality analysis.

Inputs:

1. List of skills/competencies for the educational program
2. OSCE/OSPE stations, checklists, criteria, scenarios
3. Schedule, student lists, simulation center resources

Outputs:

1. OSCE/OSPE protocols, individual results
2. Report on station/examiner quality
3. Recommendations for improvement (CAPA) and action plan

Key stages:

1. OSCE/OSPE planning (stations, skills, resources, schedule).
2. Preparation of stations, materials, and standardized checklists.
3. Training of examiners/assistants, assessment calibration.
4. Conducting the OSCE/OSPE according to procedure (route, timing, monitoring).
5. Collection and verification of checklists/scores, preparation of reports.
6. Announcement of results, handling of appeals (if applicable).
7. Analysis of reliability/quality and implementation of improvements.

KPI:

1. KPI1: Percentage of students who passed the OSCE/OSPE as planned — target: ___%.
2. KPI2: Number of stations without critical procedural violations — target: ___/total.
3. KPI3: Examiner calibration results (discrepancy in scores) — target: ≤ ___%.

Risks:

1. R1: Uncalibrated assessment → action: examiner training + monitoring of discrepancies.
2. R2: Lack of resources/stations → action: planning, reserve days, flow distribution.

Records/evidence:

1. OSCE/OSPE plan, checklists, protocols, analysis reports
2. Examiner registry, station readiness reports, CAPA plans

Related QMS documents: QMS-30-03; QMS-20-14; QMS-40-06; QMS-60-03/04

Forms/templates: station checklist, OSCE protocol, analysis report

Monitoring: for each OSCE/semester; Review: annually.

Below are **ready-made templates** for insertion into documents:

1. **Internal Audit Checklist (Q2)**
2. **Approval sheet** (for internal audit regulations)
3. **List of Developers and Responsible Parties** (for internal audit regulations)
4. **Approval sheet** (for CAPA procedure Q3)
5. **List of Developers and Responsible Persons** (for CAPA Procedure Q3)

INTERNAL AUDIT CHECKLIST

Audit Object (Process/Department): _____

Process code: ____ (e.g., 02/03/05/Q1/Q2/Q3)

Audit period: _____

Audit date: “_” _____ 20

Auditor(s): _____

Department representative: _____

A. General Requirements for the QMS

| No. | Verification criterion | Evidence (what to look for) | Compliance (Yes/No/Partially) | Comment/ Non-compliance |
|-----|---|---|-------------------------------|-------------------------|
| A1 | Process defined (purpose, scope, owner) | Process card/procedure, assignment of responsible parties | | |
| A2 | Documents are up-to-date and version-controlled | QMS register, title/version/date, access to the current version | | |
| A3 | Process records/evidence are maintained | Logs, records, protocols, reports, eBilim exports | | |
| A4 | Process KPIs are defined and monitored | KPI report, analytics, improvement plans | | |
| A5 | Process risks are identified and mitigation measures are in place | Risk register, response measures, responsible parties | | |
| A6 | Staff are informed of the requirements | Briefings, memos, meeting minutes | | |

B. Process implementation (step-by-step)

| No. | Verification criterion | Evidence | Yes/No/Partially | Comment/Non-compliance |
|-----|--|--|------------------|------------------------|
| B1 | Process planning is carried out in accordance with regulations | Plan/schedule, timetable/calendar, orders | | |
| B2 | Execution corresponds to the process description | Examples of completed operations, sample documents | | |
| B3 | Control points are met | Control reports, deadline monitoring | | |
| B4 | Data is recorded in the IS (if applicable) | eBilim/EDO records, logs, exports | | |
| B5 | Deviations are identified and recorded | Nonconformity/NS register, memos | | |

C. Nonconformities and Corrective Actions (CAPA)

| No. | Criterion | Evidence | Yes/No/Partially | Comment |
|-----|---|---|------------------|---------|
| C1 | Nonconformities are documented using the established form | NS Register, CAPA cards | | |
| C2 | Causes are analyzed (5 Whys/Ishikawa) | Completed CAPA cards | | |
| C3 | Actions are completed on time | CAPA status, supporting records | | |
| C4 | Effectiveness of actions is verified | "Effective/Not effective" mark, post-verification | | |

D. Specifics by area (select as appropriate)

D-EDUCATION (02/03): schedule, attendance/grade logs, reports, FOS, BRS, performance analysis.

D-OSCE/OSPE (05): stations, checklists, examiner calibration, protocols, appeals, reliability analysis.

D-IT/eBilim: service availability, access rights, backup procedures, SLA, incident log.

Audit Summary:

1. Non-conformities (number): __ critical __ non-critical

2. Observations/opportunities for improvement: __

3. Recommendations: _____

Signatures: Auditor _____ / Department Representative _____

CAPA CARD (Corrective/Preventive Actions)

CAPA Code: CAPA-__-20__

Source: internal audit KPI monitoring student complaint OSCE/exam risk other: _____

Date of Registration: ..20__

Process/Department: _____

Process Code: ____ (02/03/05/Q1/Q2/Q3...)

Classification: critical non-critical observation/improvement

A. Description of non-conformity

| Field | Fill in |
|--|---------|
| Description of nonconformity (what happened) | |

| | |
|--|---|
| Requirement/criterion violated (reference) | QMS document: __ section __ / standard __ |
| Evidence (facts, references to records) | eBilim/statement/protocol/report/screenshot/report No. __ |
| Scope (groups/period/quantity) | |
| Immediate actions (if necessary) | |

B. Root Cause Analysis

| Method | Fill in |
|--------------------------|---|
| Analysis method (select) | <input type="checkbox"/> 5 Whys <input type="checkbox"/> Ishikawa <input type="checkbox"/> Data Analysis <input type="checkbox"/> Expert Assessment |
| Root Cause #1 | |
| Root Cause #2 (if any) | |
| Systemic cause (if any) | |

C. Action Plan (CAPA Plan)

| No. | Action | Type | Responsible | Deadline | Resources | Performance indicator | Status |
|-----|--------|--|-------------|----------|-----------|-----------------------|---|
| 1 | | <input type="checkbox"/> CA <input type="checkbox"/> PA | | | | | <input type="checkbox"/> Open <input type="checkbox"/> In progress <input type="checkbox"/> Completed |
| 2 | | <input type="checkbox"/> CA <input type="checkbox"/> PA | | | | | |
| 3 | | <input type="checkbox"/> CA <input type="checkbox"/> PA | | | | | |

CA — Corrective Action (addresses the root cause). **PA** — Preventive Action (prevents recurrence).

D. Effectiveness Check (after completion)

| Field | Filling |
|--------------------------------------|---|
| How to verify effectiveness (method) | <input type="checkbox"/> Follow-up audit <input type="checkbox"/> KPI monitoring <input type="checkbox"/> Record sampling <input type="checkbox"/> Survey/questionnaire |
| Verification date | ..20__ |
| Result | <input type="checkbox"/> Effective <input type="checkbox"/> Partially effective <input type="checkbox"/> Ineffective |
| Comments and next steps | |

E. CAPA Closure

| Field | Fill in |
|------------------------------|------------------------------|
| Closure date | ..20__ |
| Supporting documents/records | No. ___ / reference / folder |
| Closed by (OVMC) | Full Name, Signature, Date |
| Approved by process owner | Full name, signature, date |

REGISTER OF NON-CONFORMITIES AND CAPA (maintained by OVMK / Quality Department)

| No. | Date | Source | Process/Department | Process Code | Description of NS | Criticality | Process Owner | CAPA Code | D |
|-----|------|--------|--------------------|--------------|-------------------|--|---------------|-----------|---|
| 1 | | | | | | <input type="checkbox"/> closed <input type="checkbox"/> not closed | | | |
| 2 | | | | | | <input type="checkbox"/> full-time <input type="checkbox"/> part-time | | | |
| 3 | | | | | | <input type="checkbox"/> cr <input type="checkbox"/> non-cr | | | |

Source (example): Q2 audit, Q1 monitoring, performance analysis, OSCE/OSPE, student complaint, risk register.

All nonconformities are recorded in the QMS register on the day of detection (or no later than 3 business days). For critical nonconformities, a CAPA plan is developed within 5 business days. CAPA closure is permitted only after verifying effectiveness and the availability of supporting records.

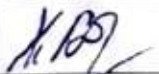
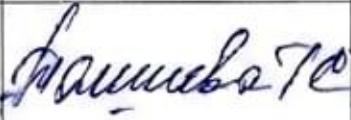







CHANGE LOG

| Change No. | Basis for Amendment | Pages | Summary of the amendment | Revision | Signature | Date |
|------------|---------------------|-------|--------------------------|----------|-----------|------|
| 1 | | | | | | |
| 2 | | | | | | |
| 3 | | | | | | |

Edition: 1000

Effective date: “ ” 20

APPROVAL SHEET

| No | Position / Role | Full Name | Signature | Date |
|----|---|--|---|-----------|
| 1 | Developed by | Kanetova D.E. |  | 29.12.25 |
| 2 | Approved: head of the responsible department |  |  | 29.12.25 |
| 3 | Approved: Head of the Educational and Informational Department | Kanetova D.E. |  | 29.12.25 |
| 4 | Approved: leading specialist for quality | Kalmuratova A. |  | 29.12.25 |
| 4 | Approved: head of the legal affairs and human resources department / lawyer | Sydykova B.J. |  | 29.12.25 |
| 5 | Approved: vice-rector for academic affairs | Sadyrova N.A. |  | 29.12.25 |
| 6 | Approved: vice-rector for science, SR and GE | Asilova Z.A. |  | 29.12.25 |
| 7 | Endorsed / considered in the established manner | JASU Scientific Council |  | 29.12.25. |

