

# PATENT EXAMINATION QUALITY MANAGEMENT AT NOIP

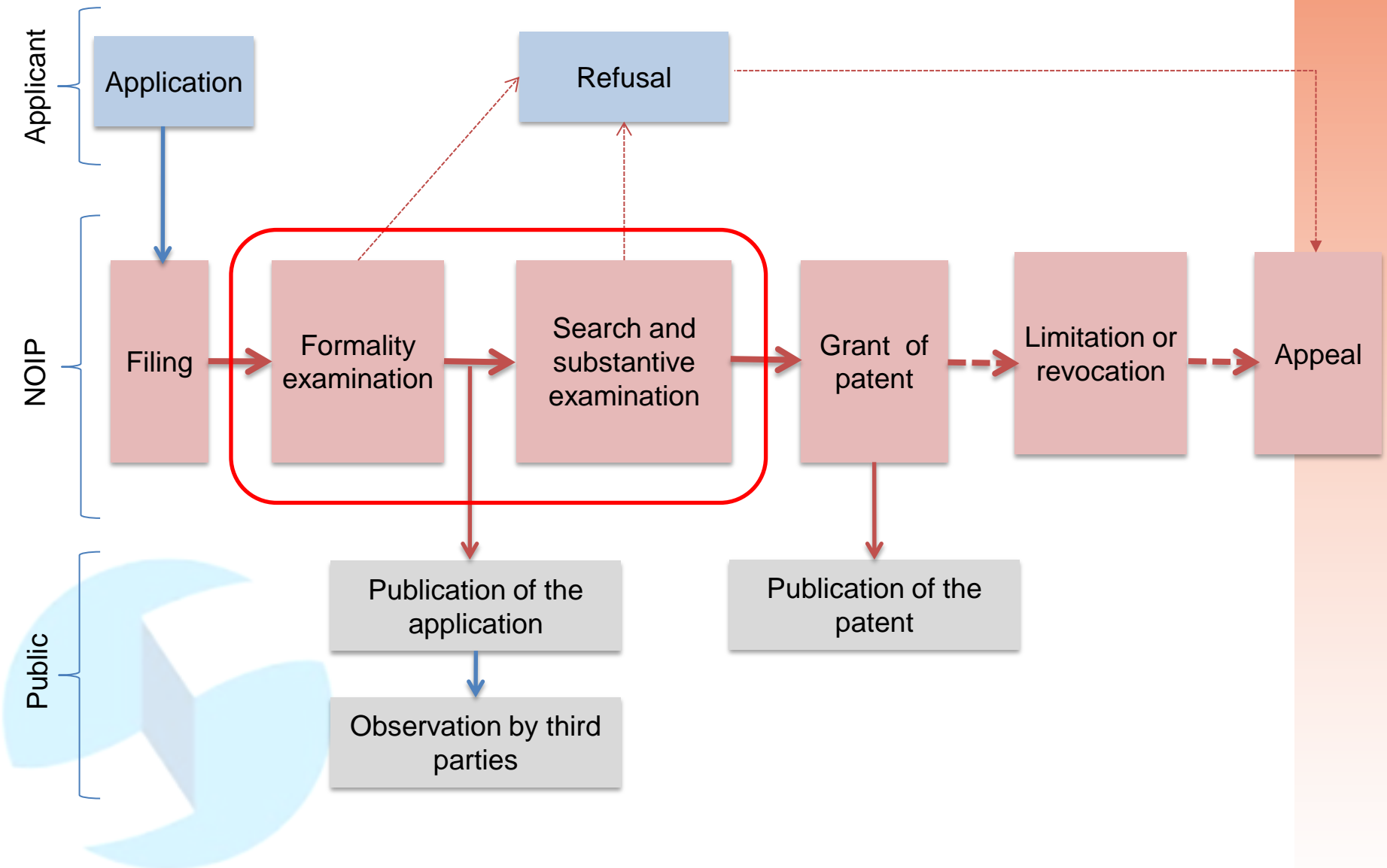


# Outline

- Relation to examination workflow;
- Persons responsible for managing examination quality;
- Documents to be used for managing examination quality.

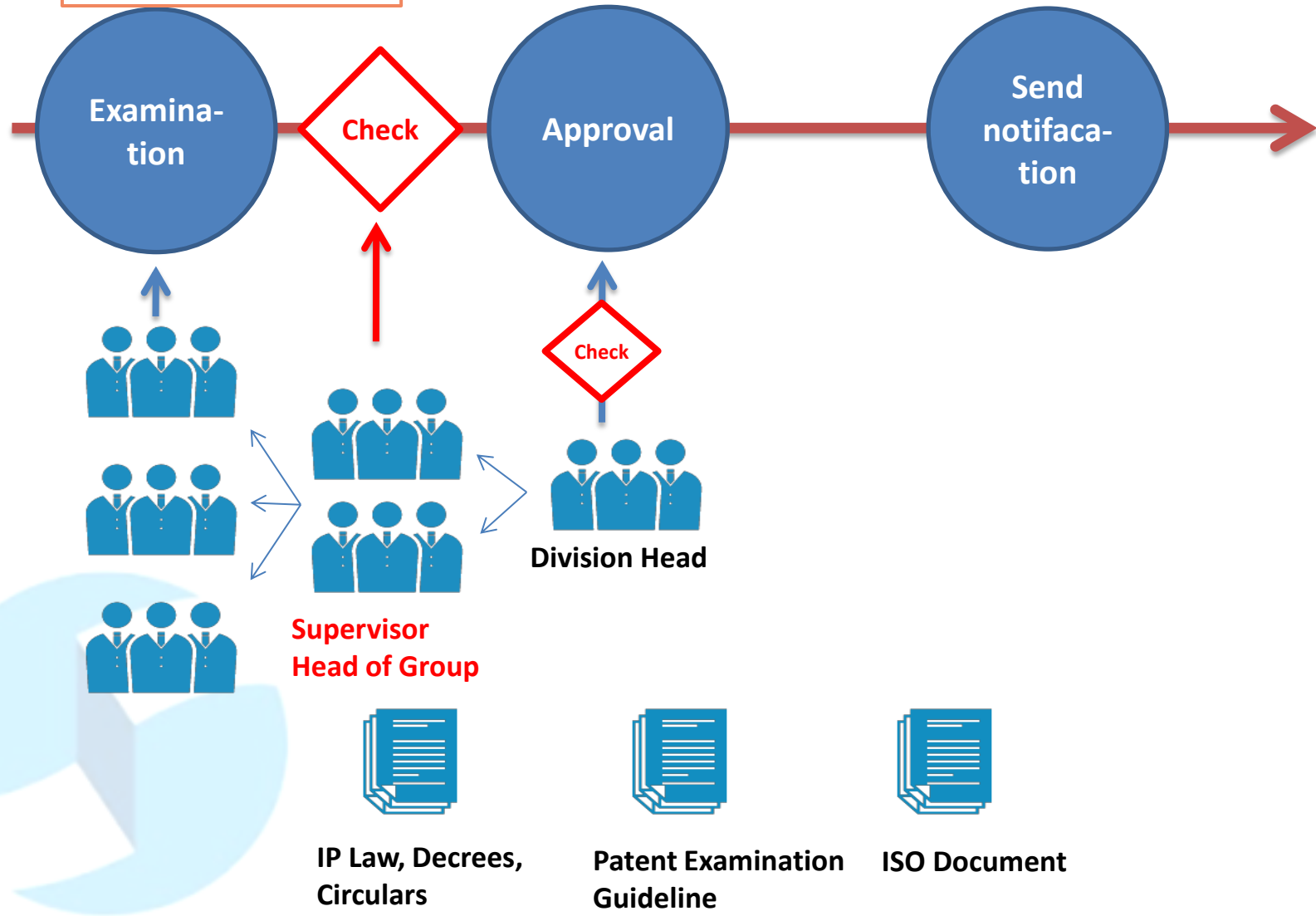


# Grant procedure



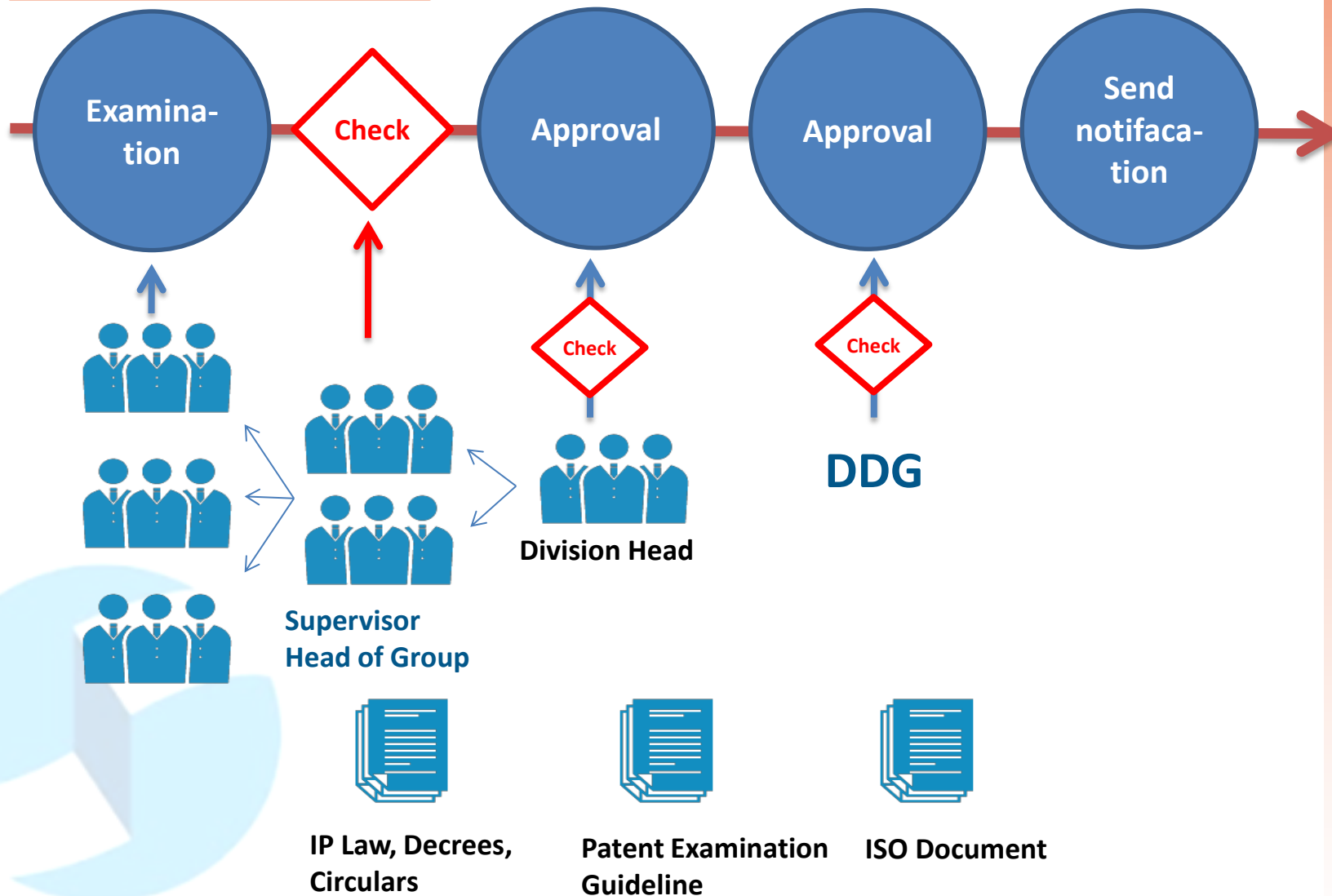
# Workflow of patent examinations and quality management

- ❖ FE notification
- ❖ SE: reason for refusal



# Workflow of patent examinations and quality management

❖ SE final action: grant, reject



# Workflow of patent examinations and quality management

- Head of group/supervisor:
  - ❖ Check every work product
  - ❖ FE: required documents; deadlines to submit documents; national phase entry of PCT application; ...
  - ❖ SE: technical concepts to be searched; formulation of search query; decisions complying with Law, Regulations; notifications easy to understand; form of notification; ...

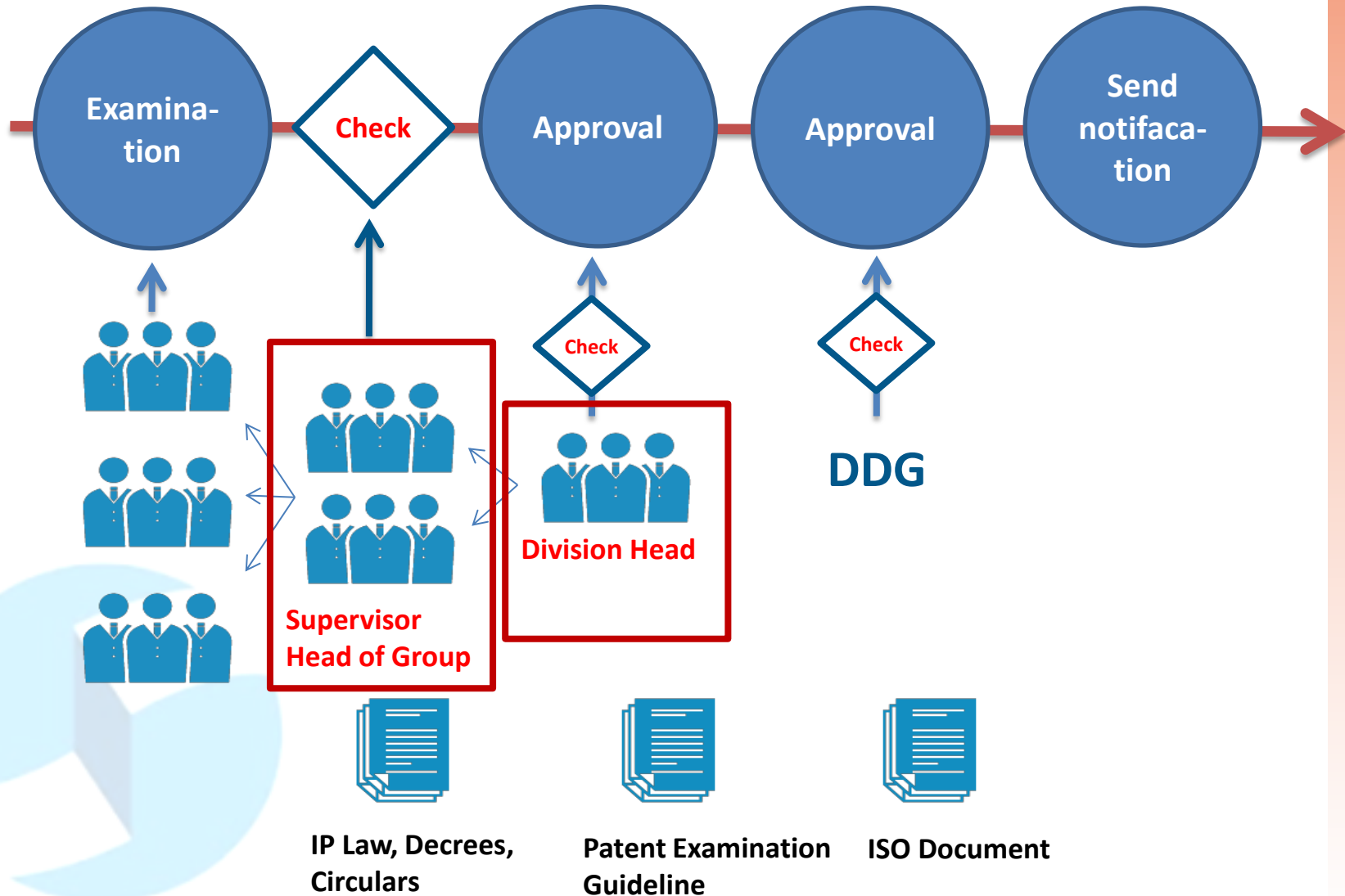
# Workflow of patent examinations and quality management

- Division heads and DDG:
  - ❖ Focus on legal points: the compliance with Law and Regulations;
  - ❖ Sometimes give comment on search results;



# Persons responsible for managing examination quality

❖ SE final action: grant, reject



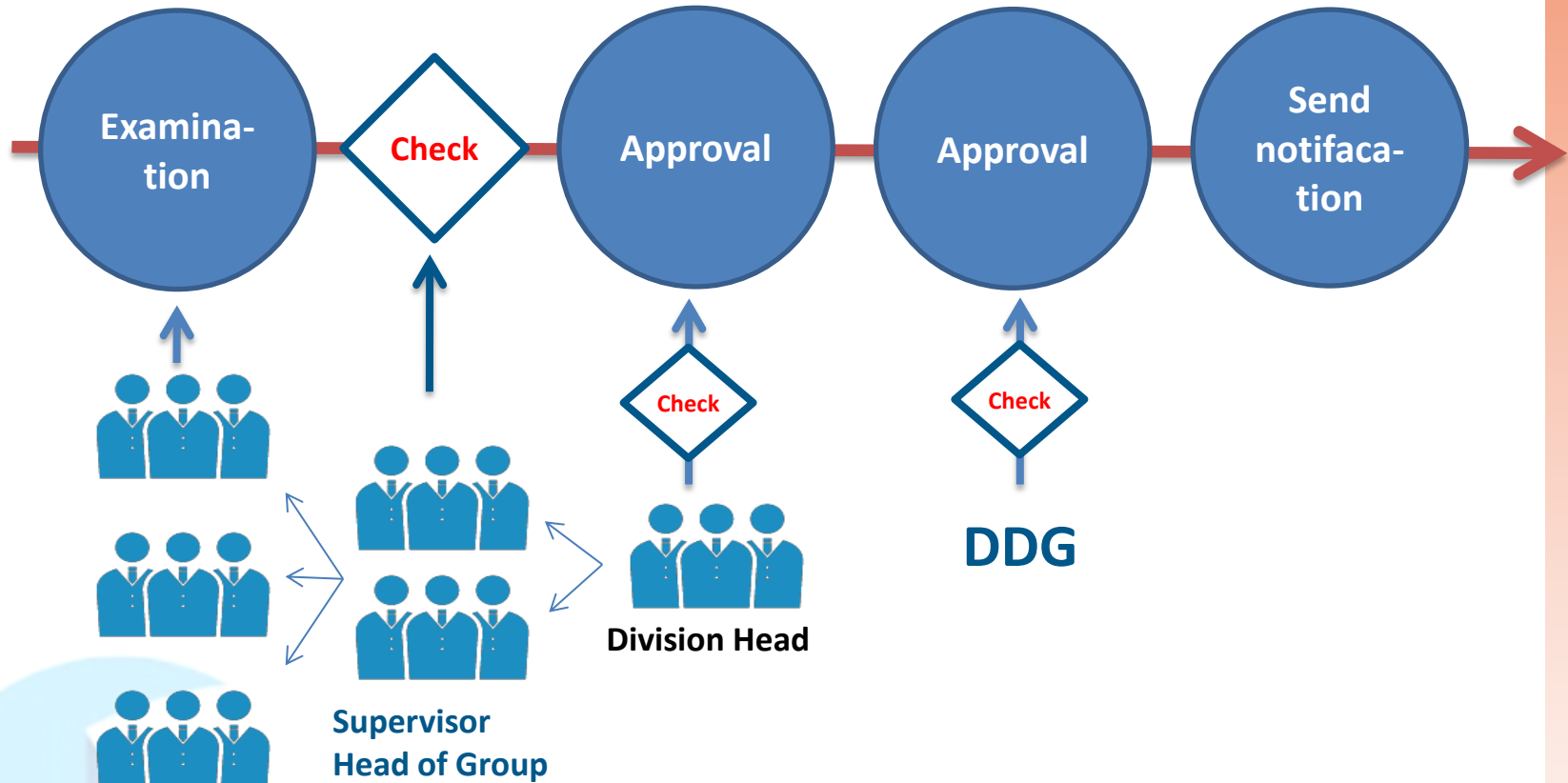


# Persons responsible for managing examination quality

- Supervisor:
  - Five year experience as an examiner
  - Good knowledge and performance in at least 3 recent years
- Head of group:
  - Three year experience as a supervisor
  - Good knowledge and performance
- Division head:
  - Generally more than 10 year experience in examination division
  - Involved in drafting legislative documents
- The Office provides trainings to improve knowledge and skills: administration, English, ... but no specific training on examination quality management.

# Documents to be used for managing examination quality

❖ SE final action: grant, reject



**IP Law, Decrees,  
Circulars**



**Patent Examination  
Guideline**



**ISO Document**

# Documents to be used for managing examination quality

- The requirements and instructions specified in legislative documents and patent examination guideline
- ISO documents only deal with the procedure, timeline matters
- Lack of specific guidance for persons responsible for managing examination quality

# Thank you for attention!

Any question?

