

UCLH Software Development Community

./sdc

Initial Meeting: 2019-02-15

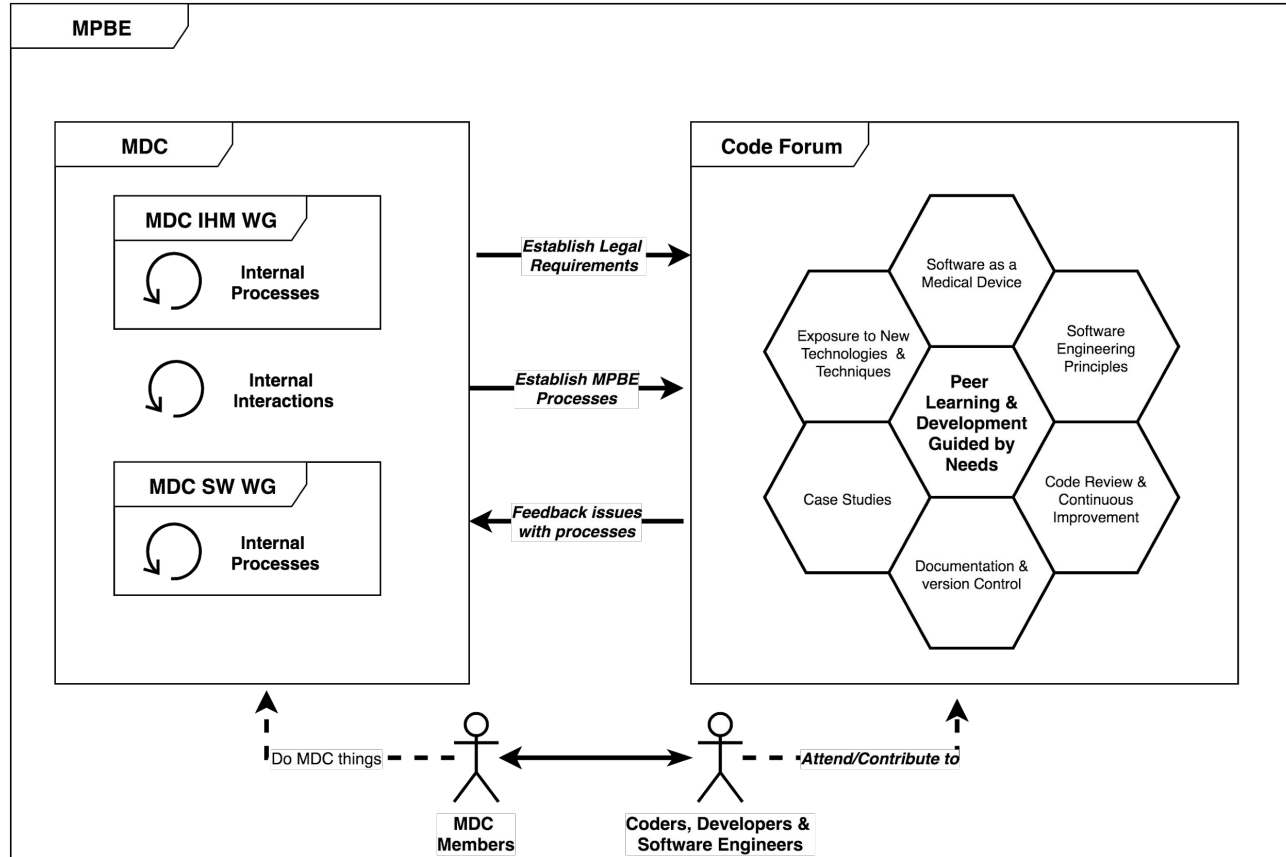
Purpose

- Share best practice for software development
- Peer to peer learning
- Code review
- Rubber ducking
- Develop pragmatic/repeatable approach to Medical Device Software development

How will sdc work?

- Regular informal meetings (fortnightly)
- Aim to establish core group with pool of knowledge to cover aspects of:
 - Software engineering
 - Medical device legislation
 - System/device life-cycle management and support
 - But invite expert guest speakers as appropriate

./sdc and MDC



How will we do this?

- Initial goal of group:
 - Let's build a Medical Software Device
 - Work through development stages of hypothetical device drawing from examples of real 'in-house' medical device applications
 - Covering
 - Requirements gathering
 - Tool selection
 - Version control
 - Documentation
 - **Legislative considerations**
 - Use of SOUP
 - **Code design**
 - Deployment
 - Management
- + whatever road blocks, questions or points requiring further investigation are encountered along the way

Who Are We All?

What's Your Name?

Where are you Based?

What's Your Role?

How Is Software Development Involved?

What Do You Want/Expect From sdc?

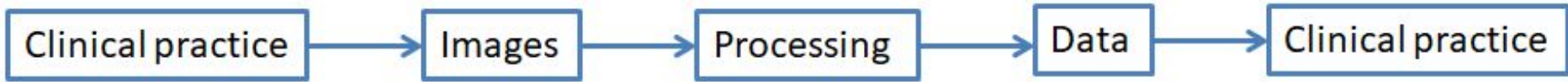


Software Development for Image Processing Pipeline

Drawing from:
Software Development for Neonatal MR
Spectroscopy Processing Pipeline

Overview of our project:

- Image processing pipeline



- Not specific to MR – frequently encountered Image analysis problem

Today's Meeting

- Step 1 – setting requirements for software
 - User requirements/stories
 - Inputs Outputs
 - Capturing initial requirements

Questions for today

What do we need to establish before proceeding?

How do we work out our requirements?

What information are
we missing?

Future Steps

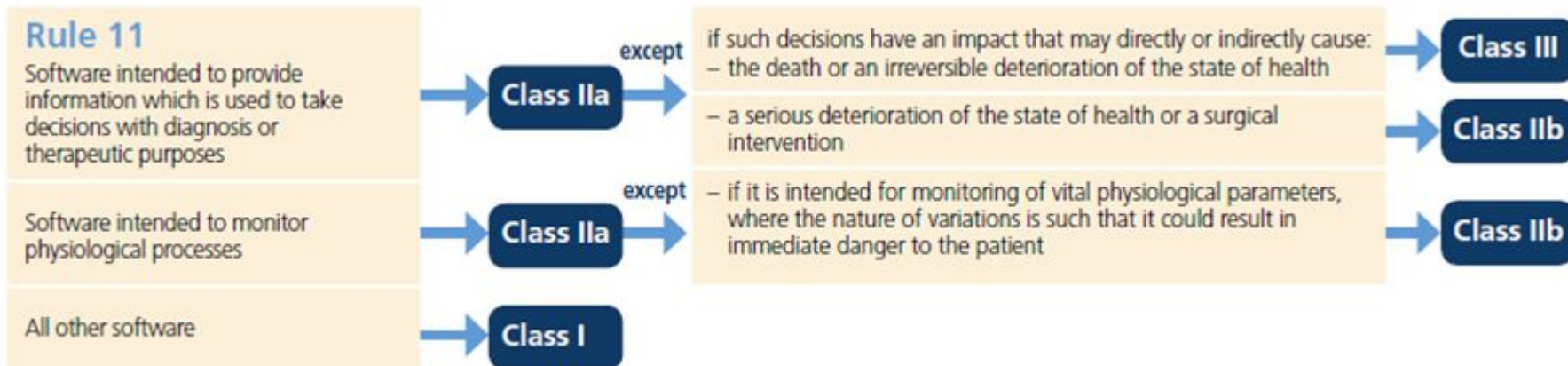
- Early March - Step 2 – setting requirements for development – choosing tools.
 - Choosing a language
 - Using an IDE
 - Version control
 - Shared development
- ~ mid to late March – invited talk on ISO 13485 and Development of Medical Devices
- Early April - Step 3 – Assessing and managing risk.
 - Classifying our pipeline
 - Architectural design choices
 - Target running environment
 - Component choices – 3rd party libraries and SOUP

Future Steps

- **Mid April onwards**

- Step 4 – What does “good” code look like?
- Step 5 – Develop – Commit – Test – Publish
- Step 6 – Infosec
- Step 7 – Software release
- Step 8 – Life-cycle management

Medical Device Classification



Next Steps

What Now?

- Feedback
 - What did you think?
 - What else can we do?
- Knowledge Storage & Comms
 - Redmine?
 - GitHub Repo
- Next Meeting
 - Friday 1st March?
 - Where?