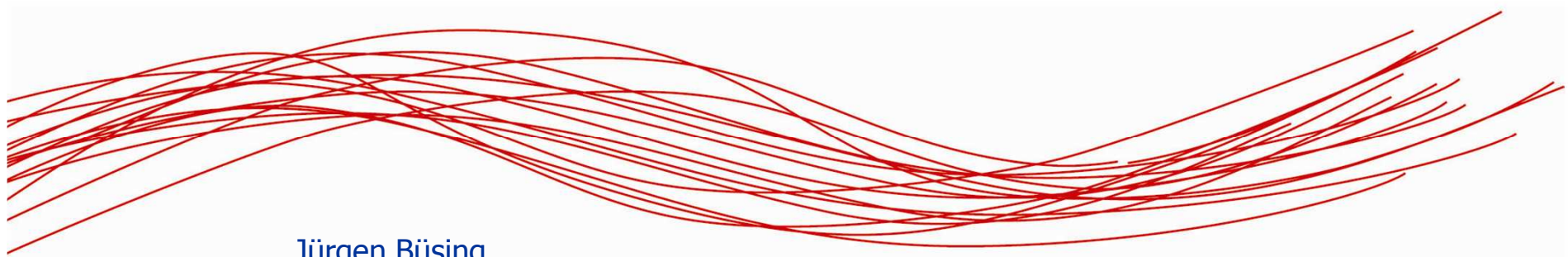


COORDINATION ACTIONS and RESEARCH POLICY



Jürgen Büsing
F.4: Advanced Therapies and
Systems Medicine

Directorate F
DG Research and Innovation

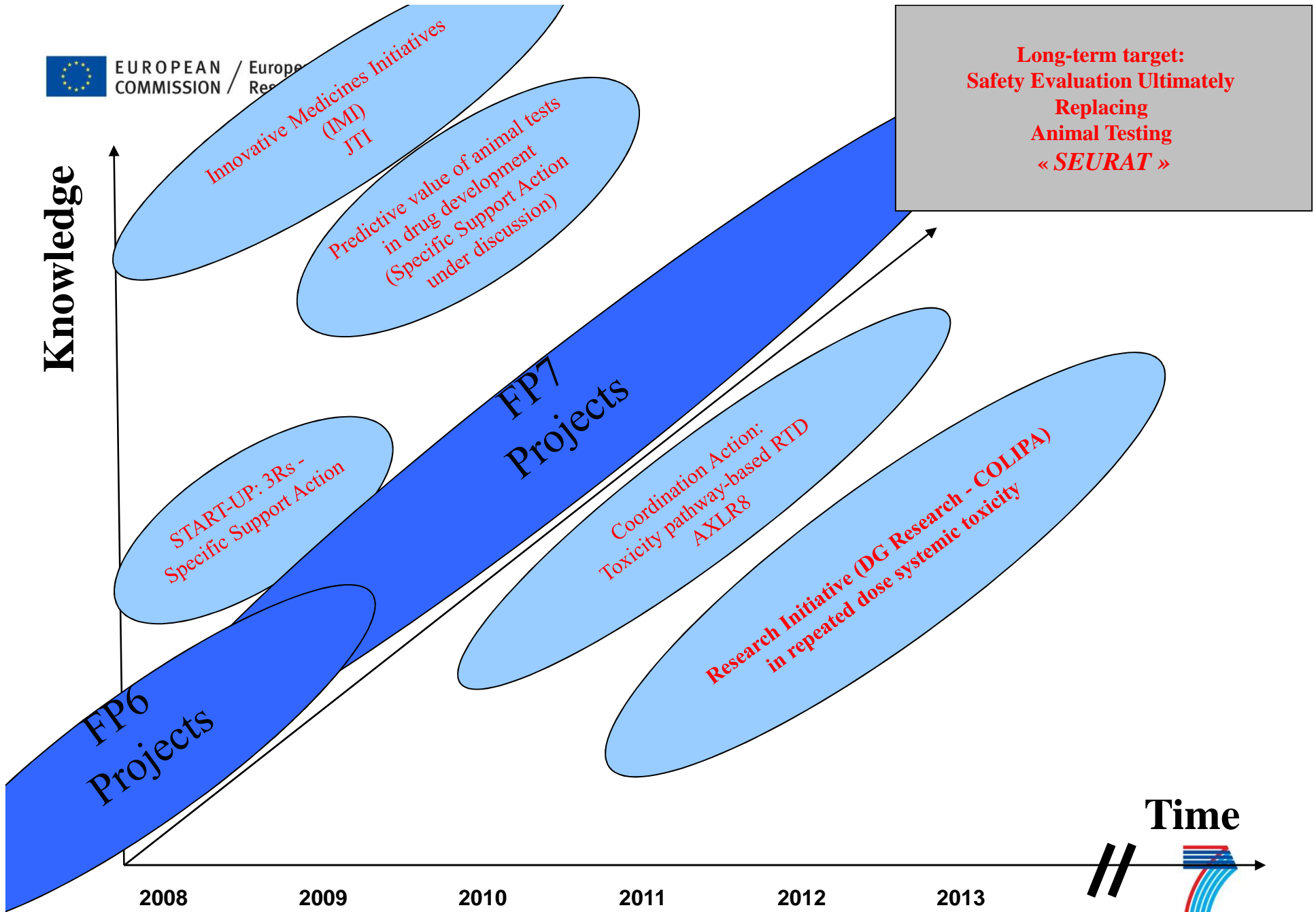
AXLR8-2 WORKSHOP 2011
23 – 25 MAY 2011, BERLIN, GERMANY





EUROPEAN COMMISSION / Europe Research

Knowledge



Long-term target:
Safety Evaluation Ultimately
Replacing
Animal Testing
« SEURAT »

Time

2008 2009 2010 2011 2012 2013



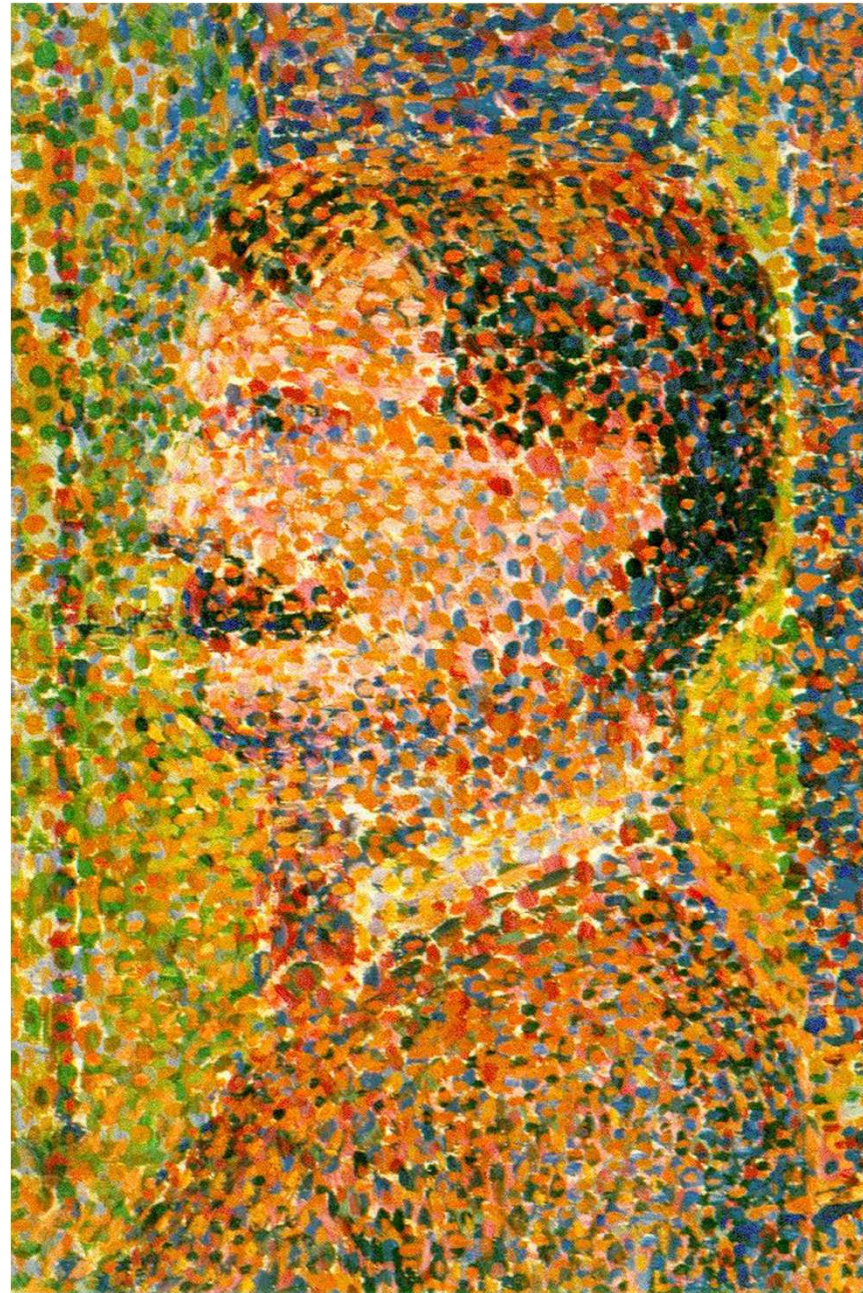
Acronym 'SEURAT':

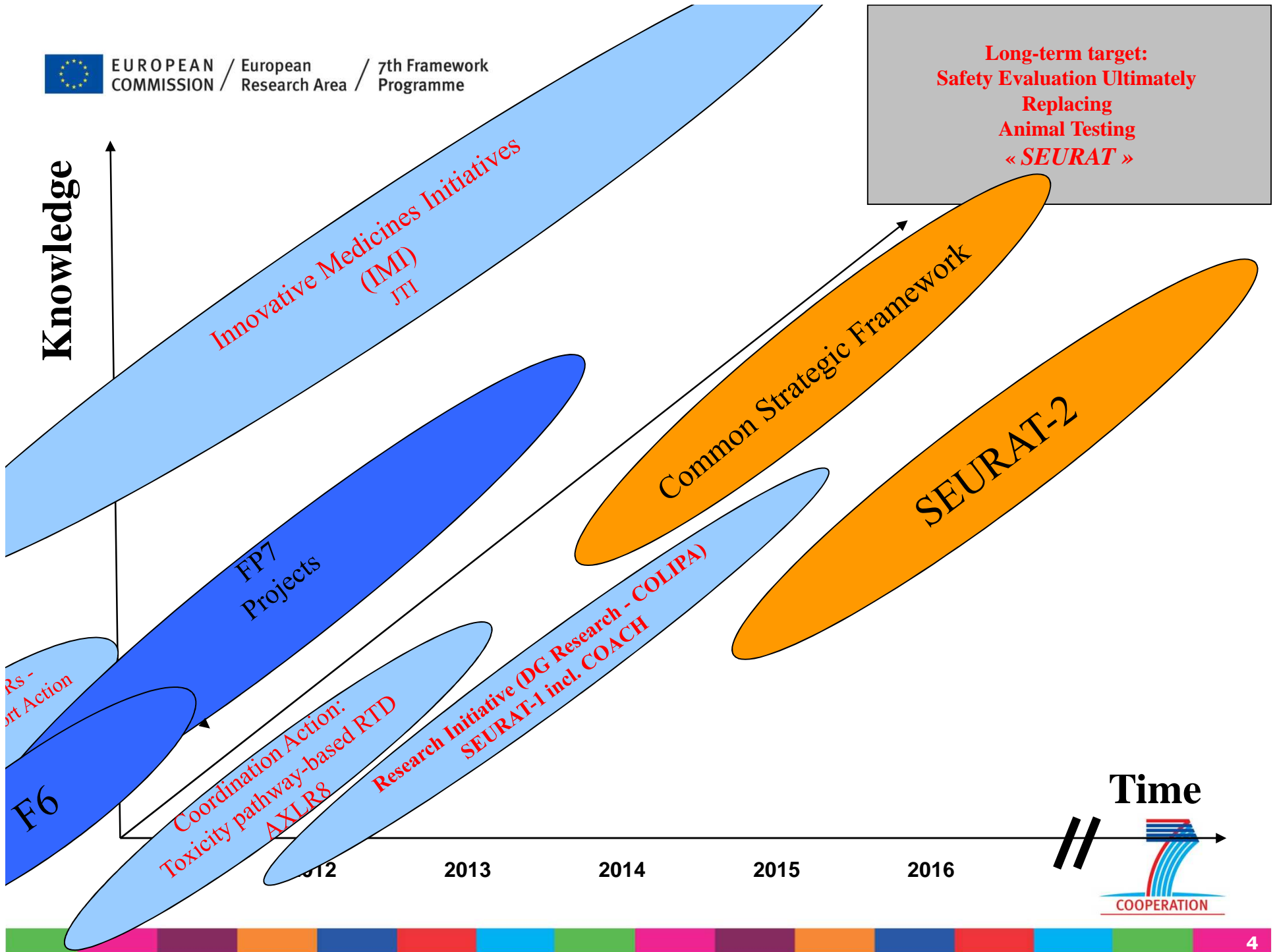
R.A. Pedersen et al., 2008

Introduction of 'The SEURAT Strategy' into the Health Programme:

M. Hallen, 2008

Georges-Pierre Seurat : detail from *La Parade* (1889) showing pointillism







Timetable

2005: Vision: R&D Programme on 'replacement of repeated dose systemic toxicity'

**2007: Call for proposals for a Coordination Action on Repeated dose systemic toxicity
(no proposals)**

2007/ 2008: first contacts Colipa-Commission (Research Initiative)

2008/ 2009: preparation of work programme by group of experts

Mid 2009: launch of call for proposals; deadline: beginning of 2010

Spring 2010: evaluation of proposals

Summer-autumn 2010: contract negotiations

January 2011: start of Research Initiative

Research Initiative

December 2015: end of Research Initiative SEURAT-1

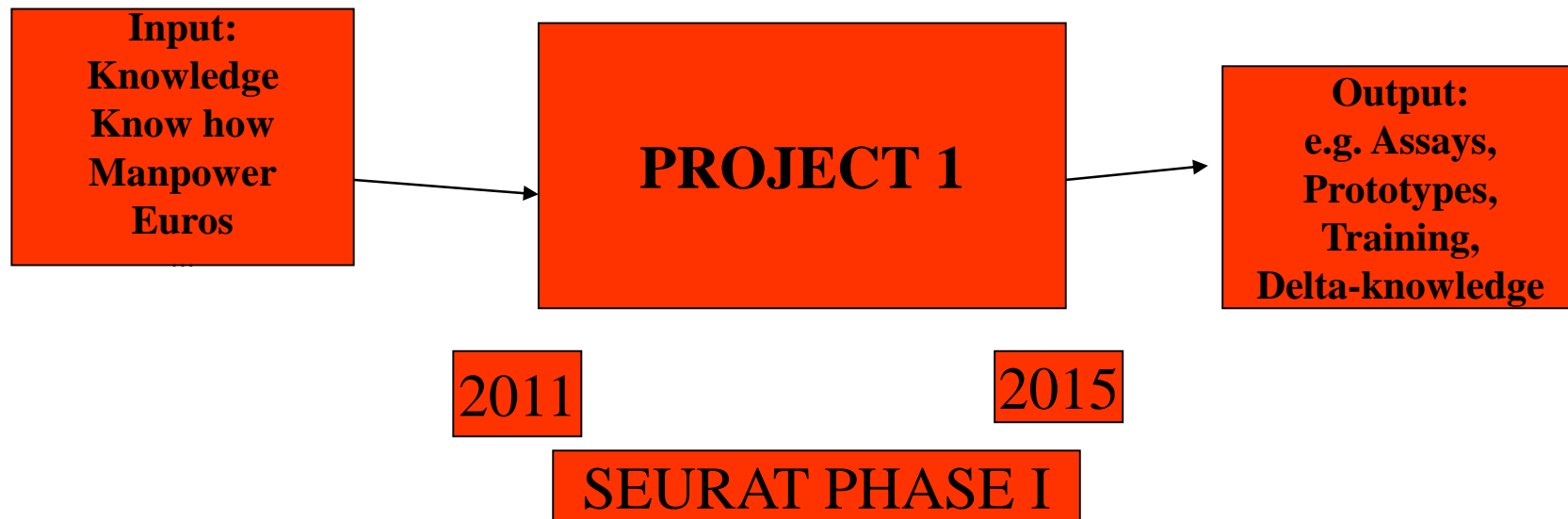
December 2016: end of Coordination Action COACH.



Urgent Tasks:

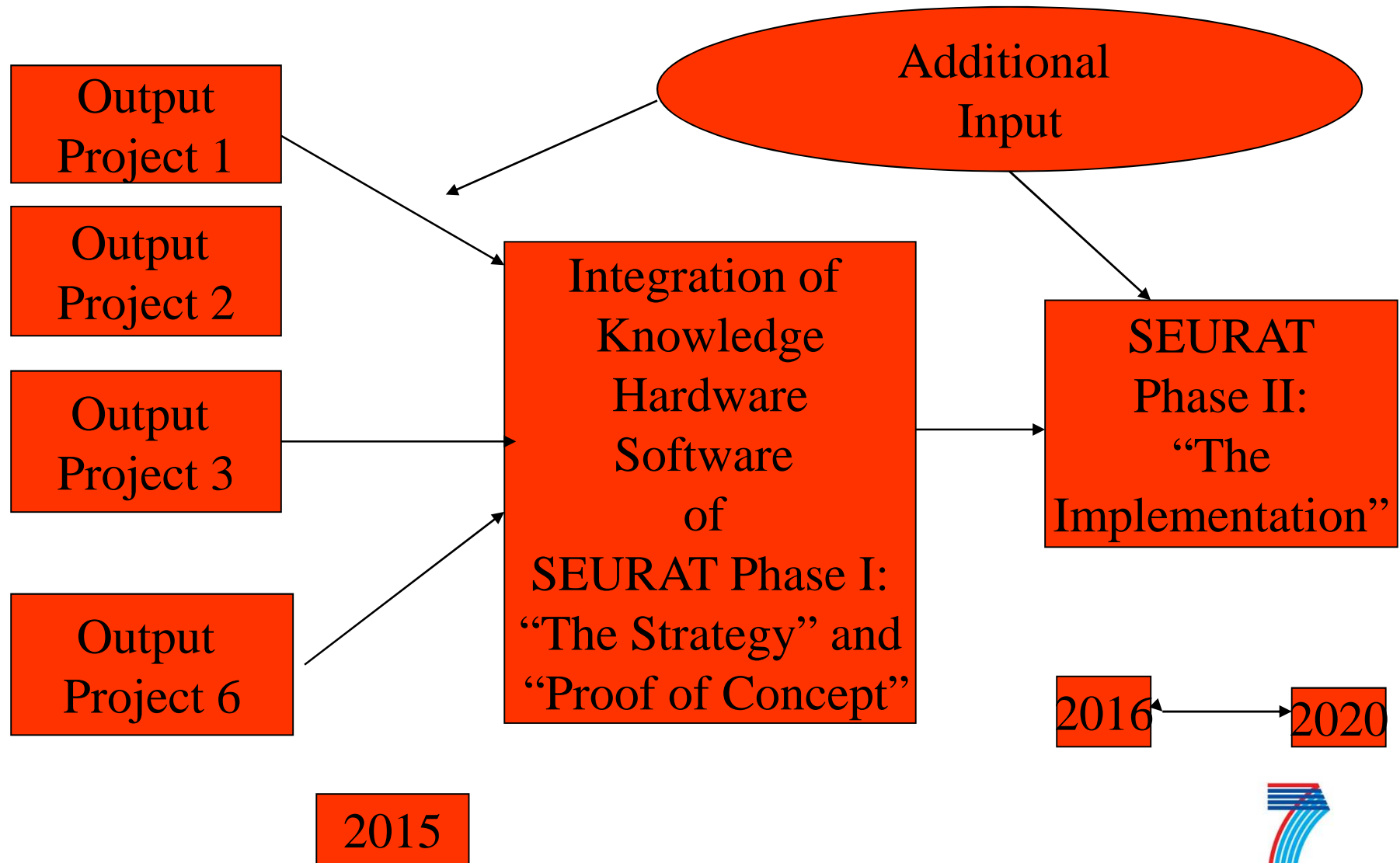
- Analyse and describe "SEURAT" (long-term target!) in detail and identify relevant 'actors'
- Monitor Projects and progress in "state-of-the art"
- Assess impact of PPP between Commission and Cosmetics Industry
- Clearly define transition from SEURAT-1 to SEURAT-2
- Develop science and business plan and roadmap for period after 2013
- Take into consideration other relevant activities (avoid duplication; create synergies)
- Develop a research plan for "Innovative Toxicity Testing in Advanced Therapies and Systems Medicine"
- Prepare Executive Summary of present workshop on 25 May 2011







Transition towards SEURAT Phase II



Cooperation within the cluster and communication with the Commission

In order to deliver the envisaged benefits of the Research Cluster on new testing methods, the individual projects and the coordinating action must cooperate to achieve common goals and sustainable benefits.

It is of the utmost importance that all beneficiaries of all Projects cooperate and that all activities are geared to one another. This demands coordination and supervision beyond the six independent Projects.

It is therefore required that the Coordination Action provides for a governing body in the form of a Scientific Panel which will monitor and coordinate all the Projects. This Scientific Panel will include the coordinators of all six projects and further external experts. The Scientific Panel will stimulate communication between the projects, will monitor progress, define future tasks of the cluster, and will be organised by the Co-ordination Action.

The chairperson of the Scientific Panel acts through the COACH secretariat as the contact point between the Commission on the Research Cluster.



Recommendation from the 2010 AXLR8 workshop:

...“Taking a ‘systems’ approach to human biology that relies on mechanisms and pathways has great value for society, contributing both to improved toxicity testing and to the fundamental molecular elucidation of human disease.” ...

The goal is to do the best science ... with a purpose; and the purpose is improved human health and safety.

