Global MedTech Compliance Conference 2017

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Interactive session on Code Convergence

Voting results

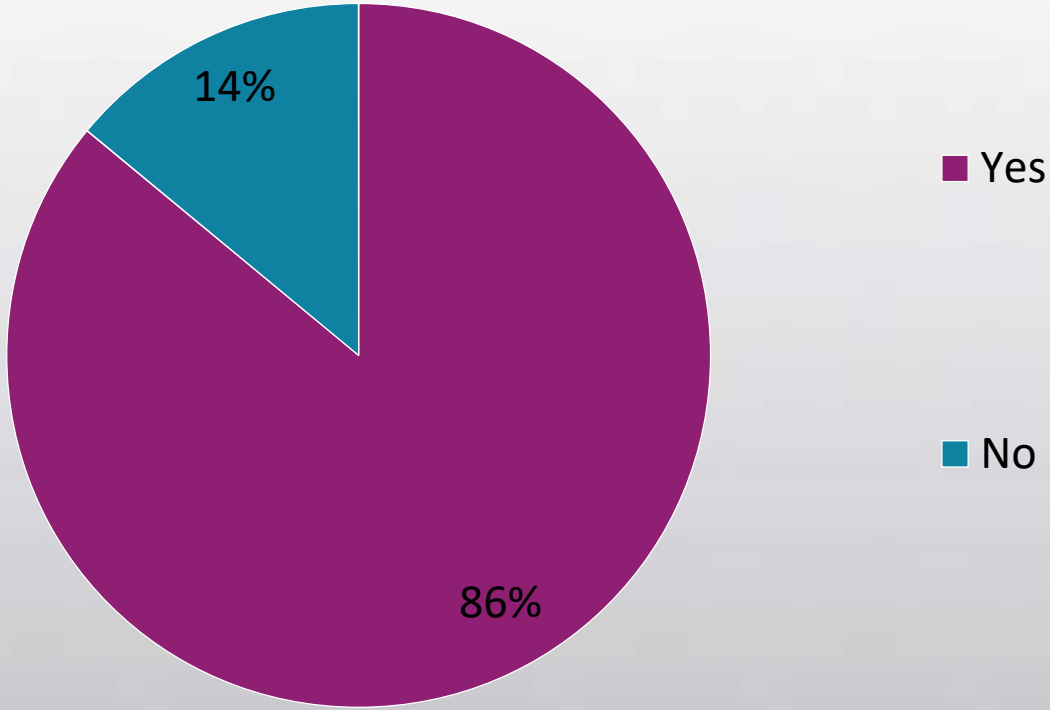
4 May 2017, Amsterdam

1) A global MedTech Code

Should we start the process to come to a global MedTech Code?

1. Yes
2. No

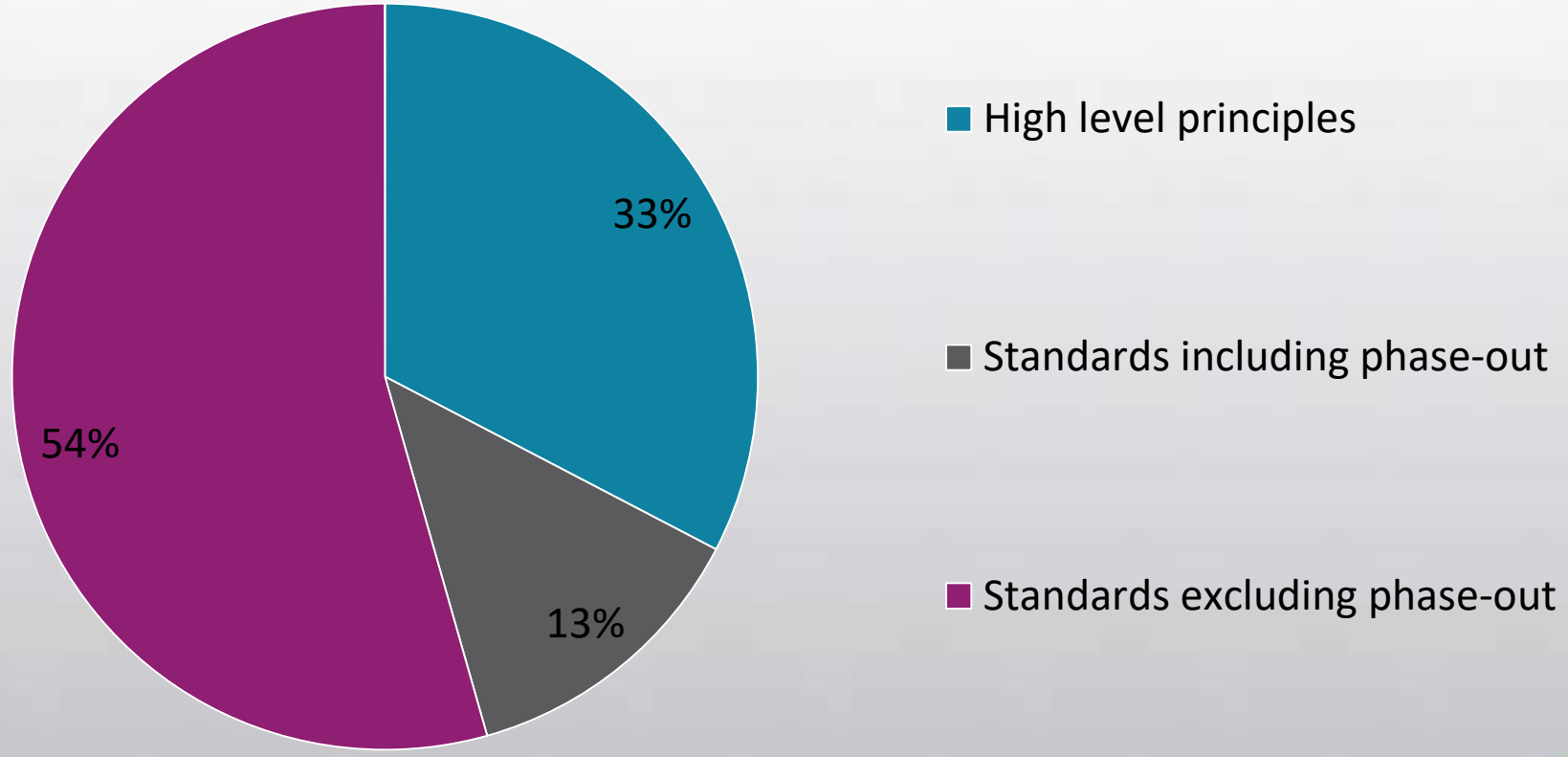
Results



The Code should have the level of detail in line with:

1. High level principles
2. Standards for each type of interactions between industry (e.g. gifts) excluding phase-out of direct sponsorship of Healthcare Professionals and Transparency
3. Standards for each type of interactions between industry (e.g. gifts) including phase-out of direct sponsorship of HCPs and Transparency

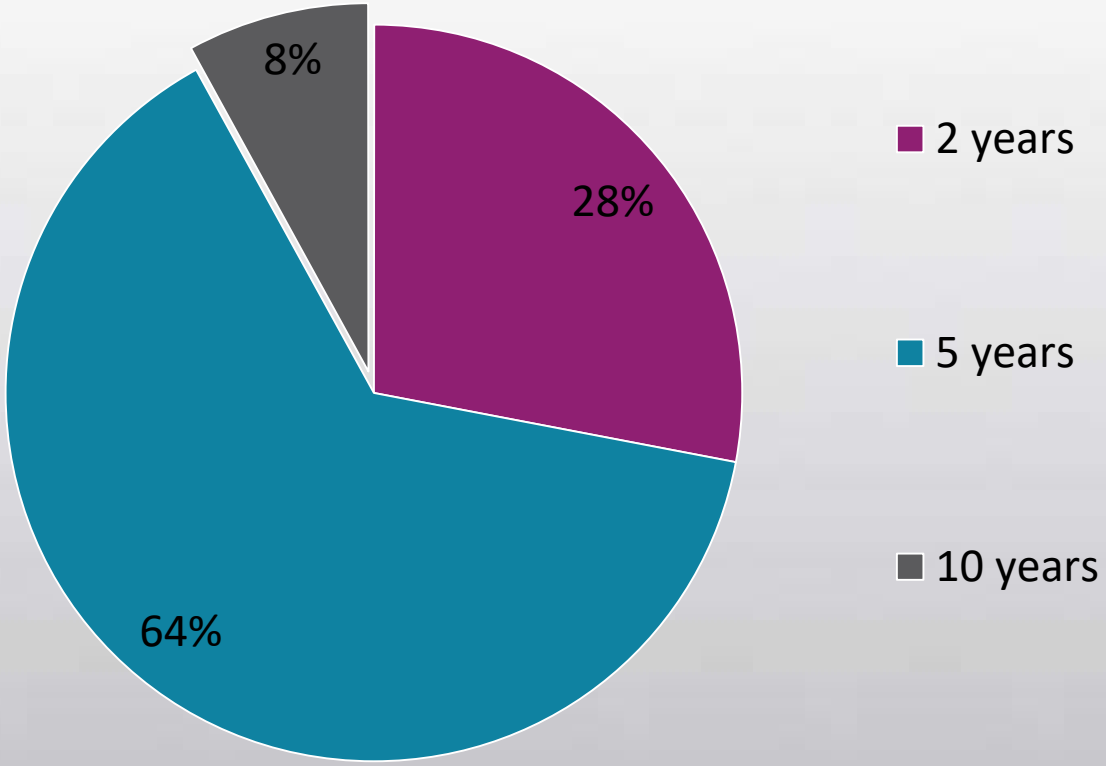
Results


33%
13%
54%
High level principles
Standards including phase-out
Standards excluding phase-out

We should allocate industry resources to make the Code a reality within (?) years.

1. 2 years
2. 5 years
3. 10 years

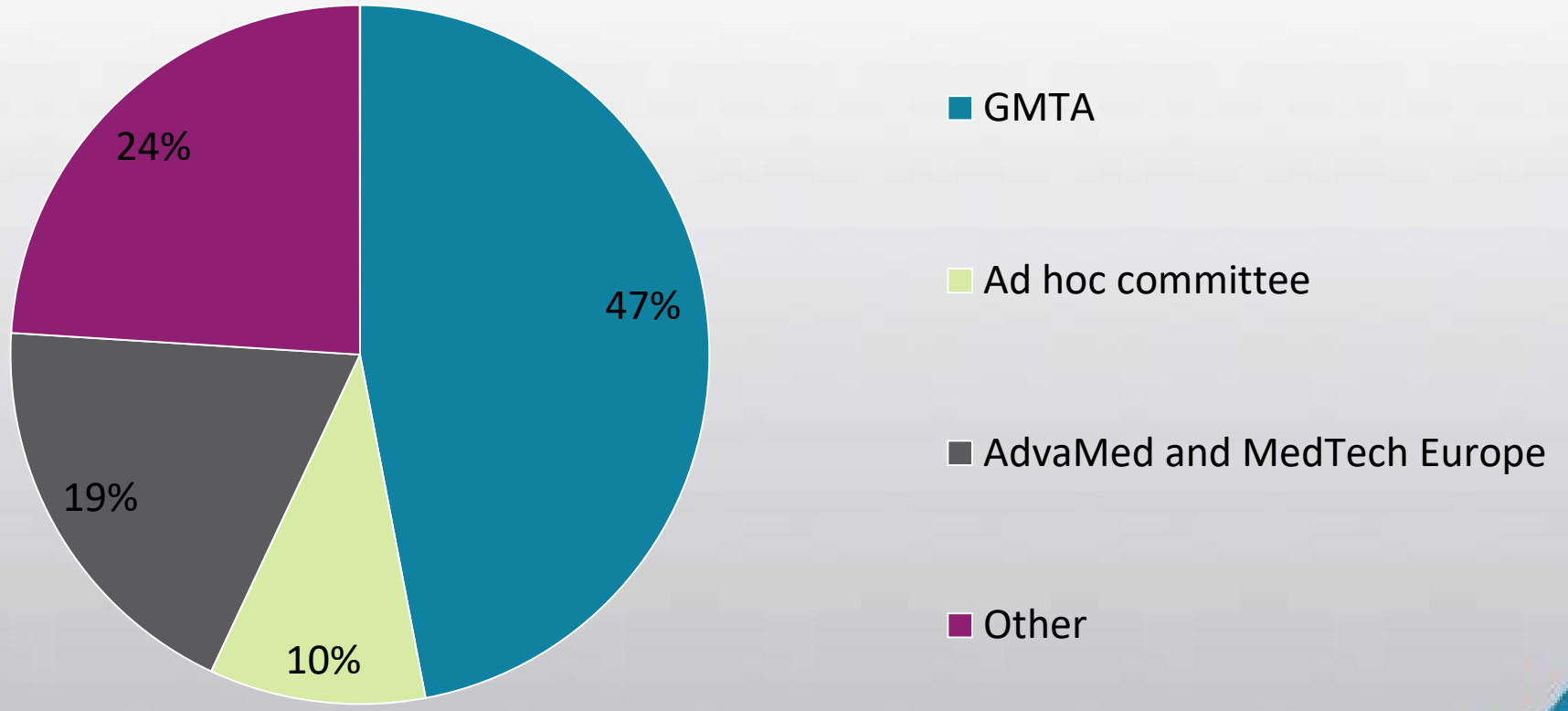
Results



Who should drive the development of the global MedTech Code?

1. GMTA (Global MedTech Alliance)
2. Ad hoc global group of compliance officers from interested companies
3. Advamed together with MedTech Europe, without other associations
4. Other

Results

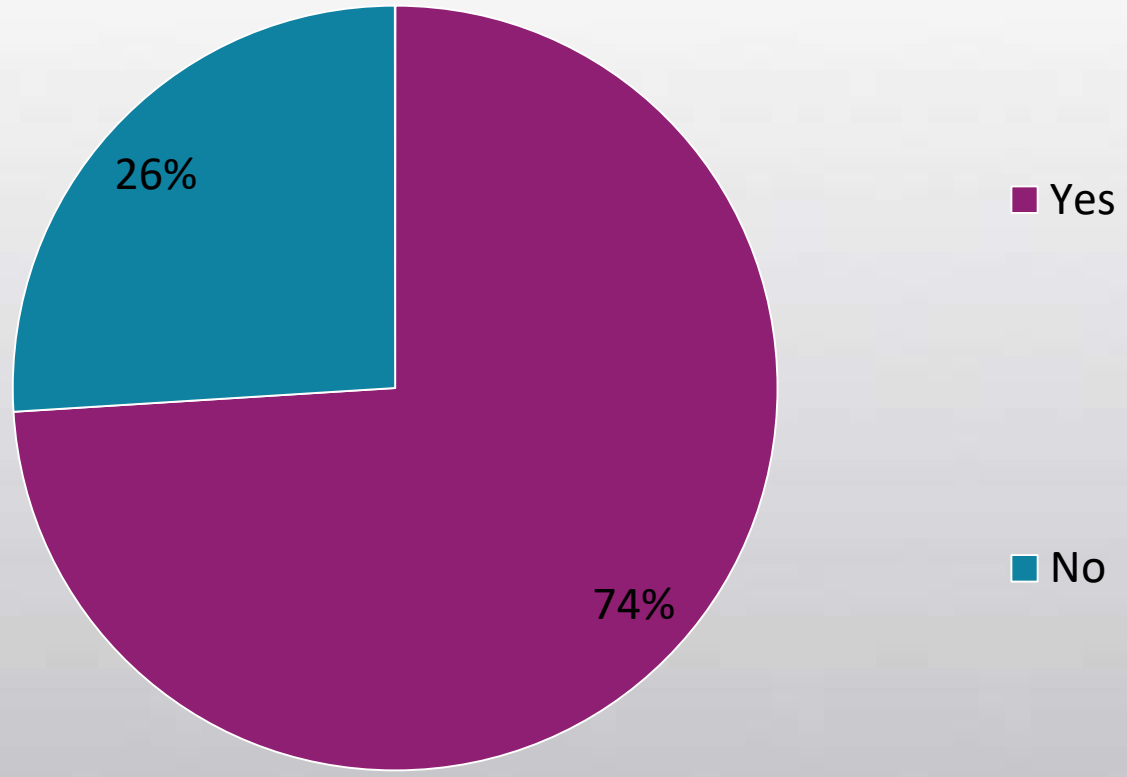

24%
47%
19%
10%
GMTA
Ad hoc committee
AdvaMed and MedTech Europe
Other

2) A Lifesciences Industry Code

Should we start the process to come to a Lifesciences Industry Code, as a model?

1. Yes
2. No

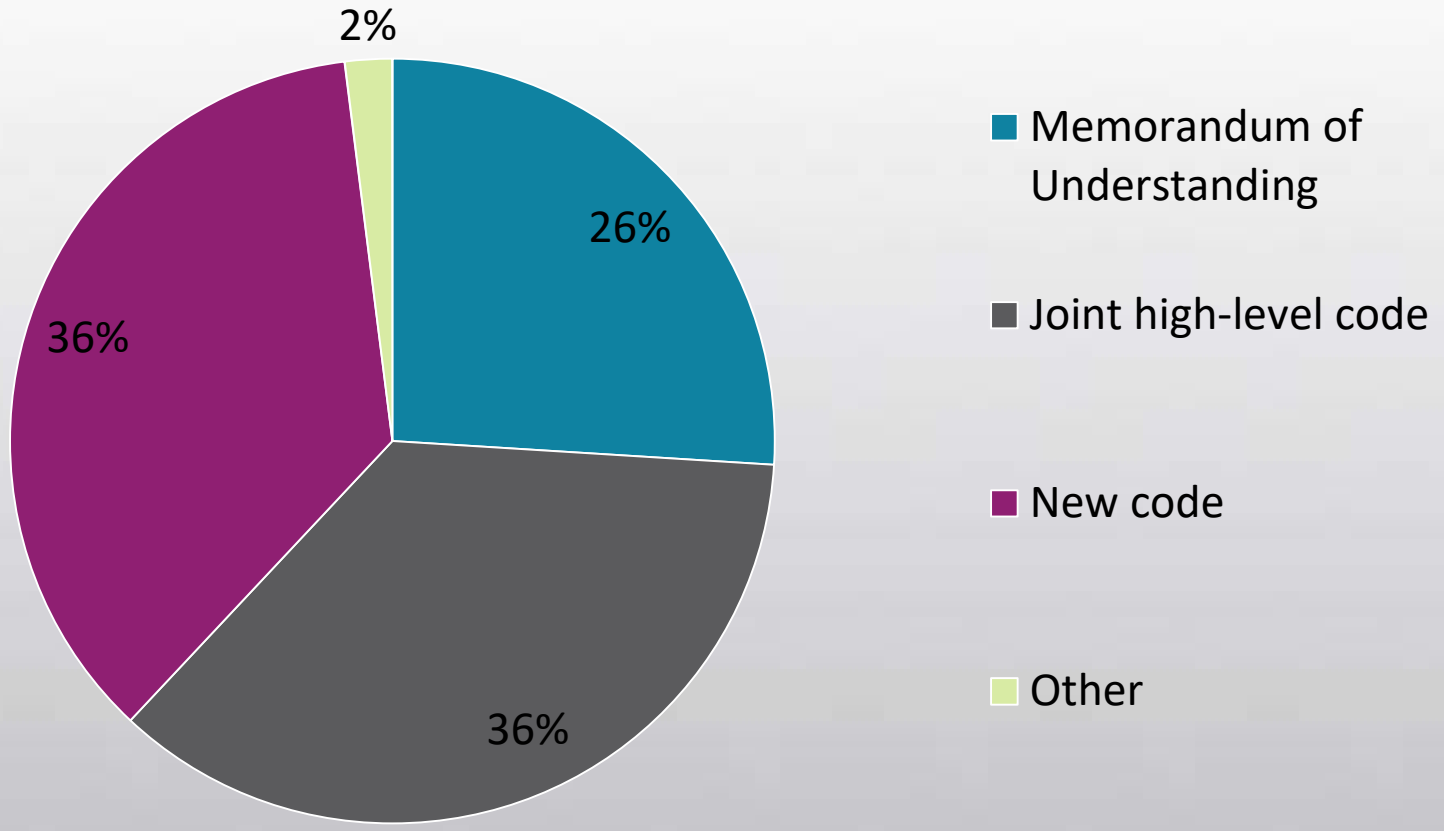
Results



What level of detail should such a convergence entail?

1. Memorandum of Understanding, including high level principles
2. Joint high-level code, which would include closer alignment but existing codes would still continue to co-exist
3. Actual new code, replacing all the existing Codes
4. Other (e.g. mutual recognition process of some sort)

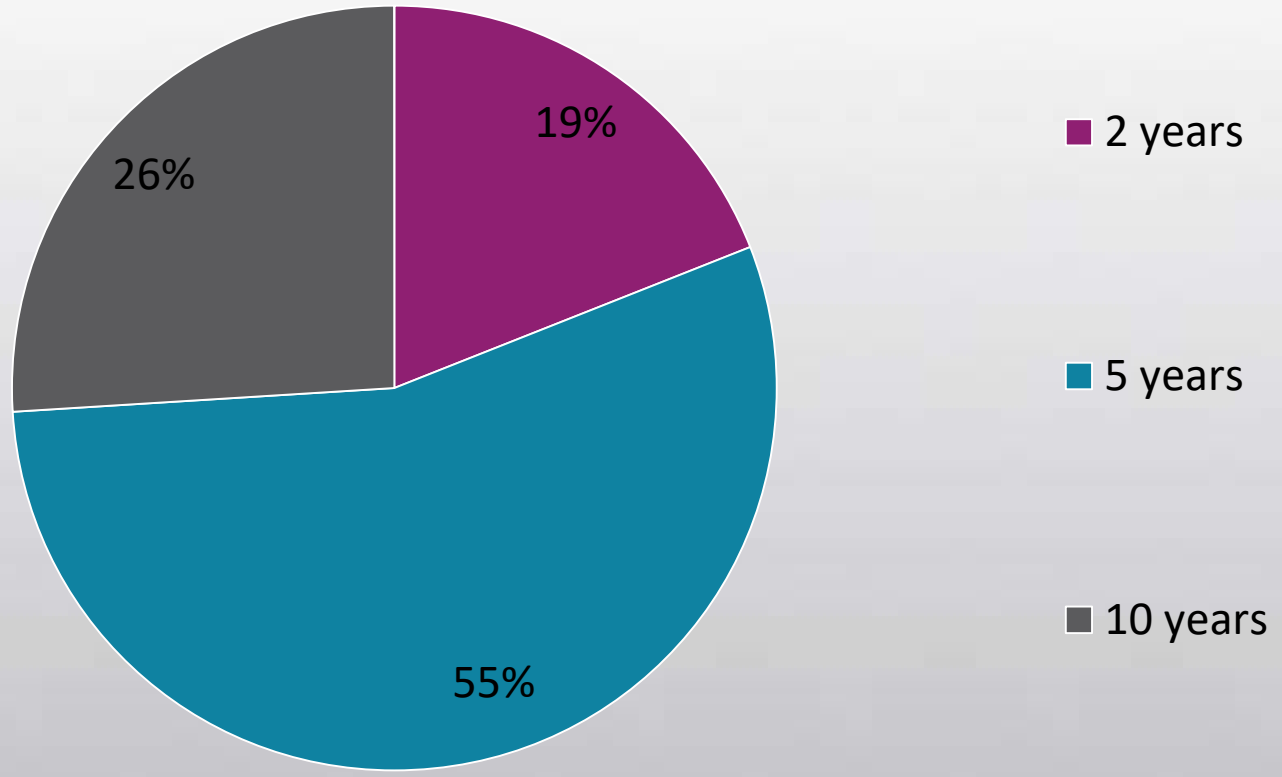
Results


2%
26%
36%
36%
Memorandum of Understanding
Joint high-level code
New code
Other

We should allocate industry resources to make the Code a reality within (?) years

1. 2 years
2. 5 years
3. 10 years

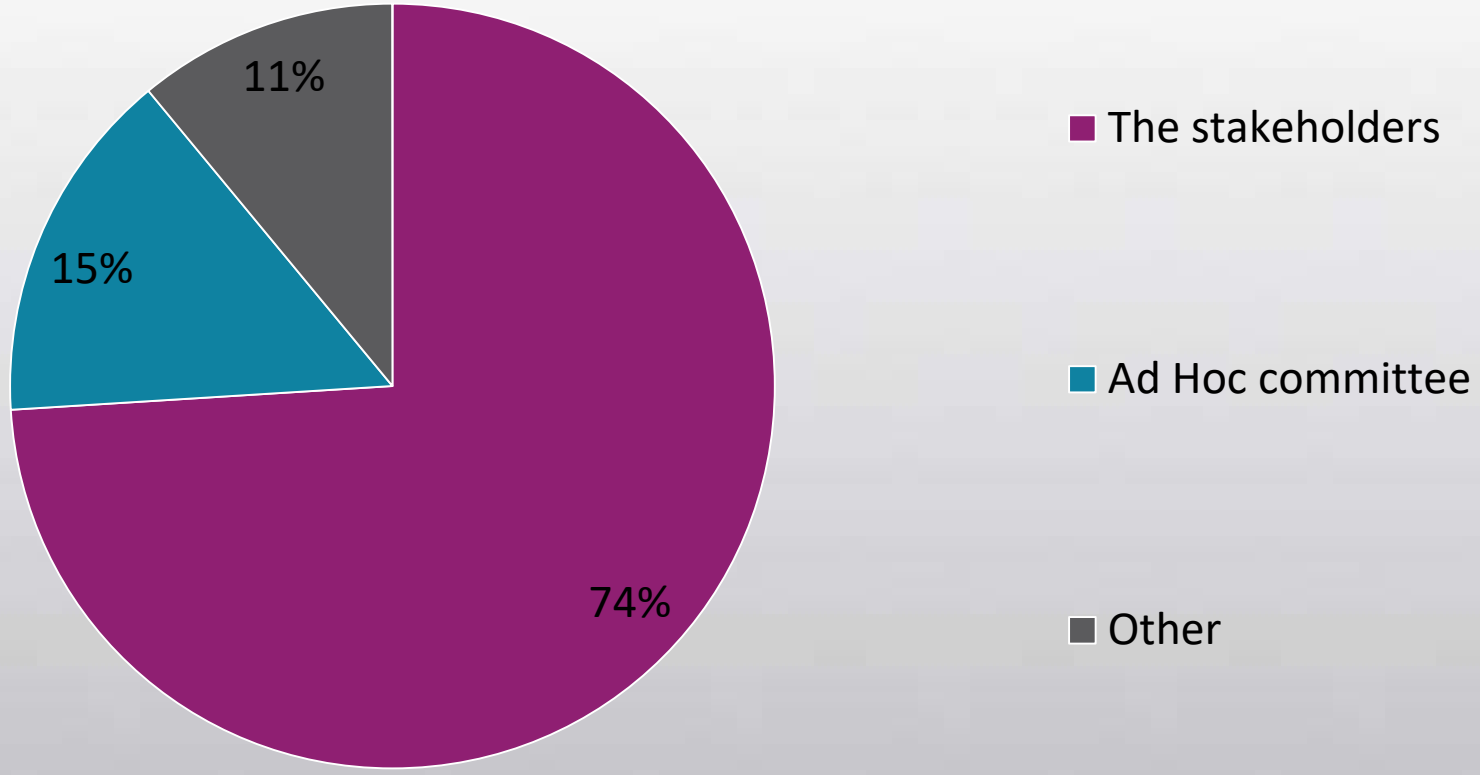
Results



Who should drive the development of the Lifesciences Industry Code?

1. EFPIA, Medicines for Europe, MedTech Europe, AdvaMed and COCIR
2. Ad hoc global group of compliance officers from all industries sitting together
3. Other

Results

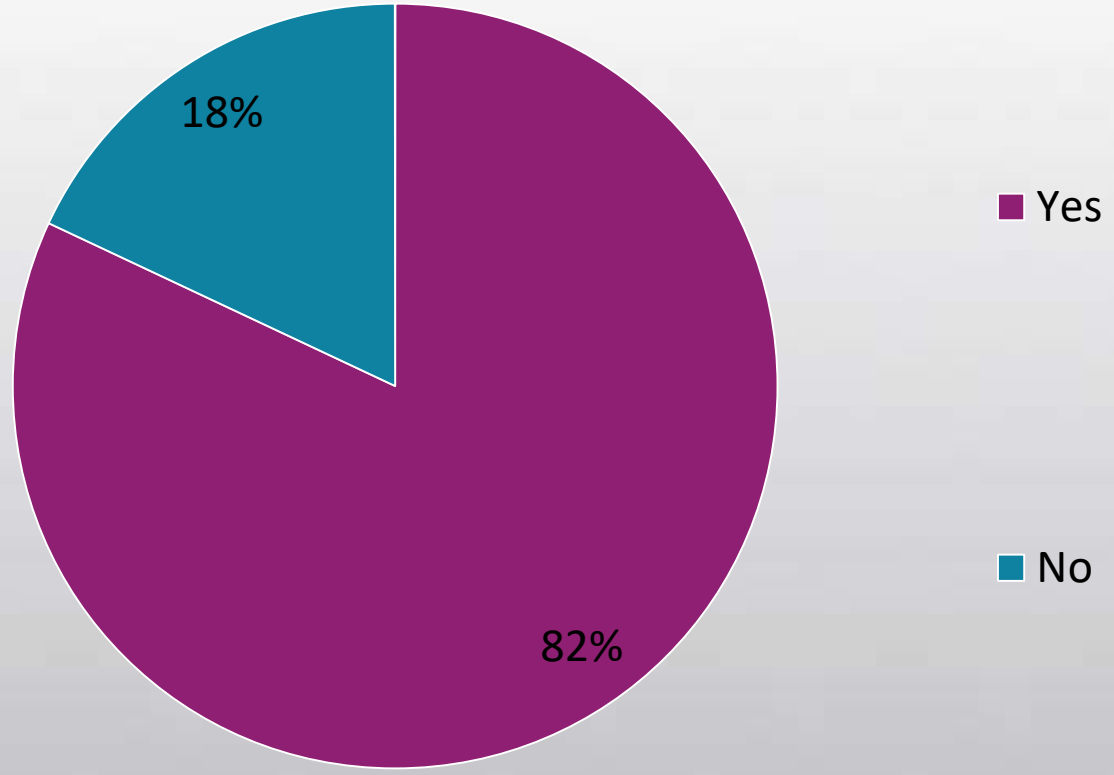


3) Stakeholders Code

Should we start the process to come to a Stakeholders Code?

1. Yes
2. No

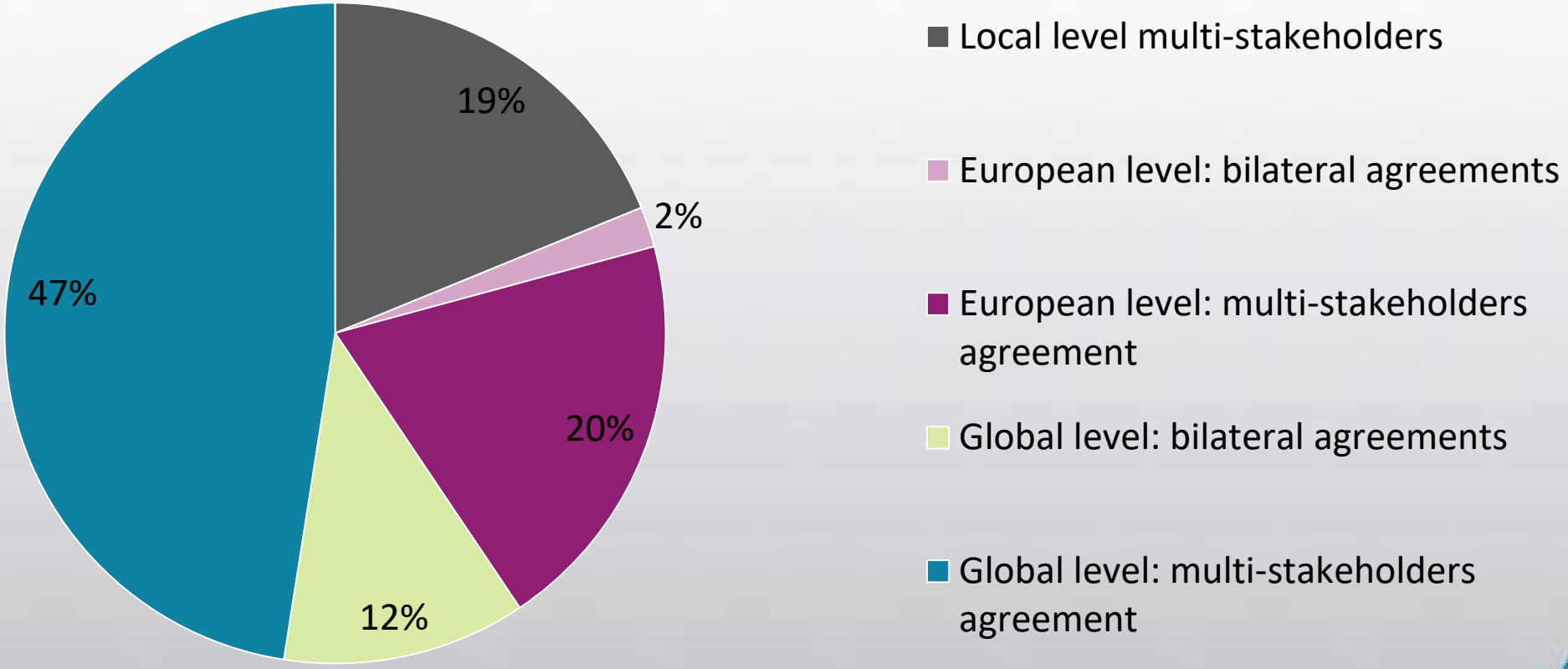
Results



The Code should be organised at:

1. Local level multi-stakeholders
2. European level bilateral agreements between interested stakeholders
3. European level multi-stakeholders agreement
4. Global level bilateral agreements between interested stakeholders
5. Global level multi-stakeholders agreement

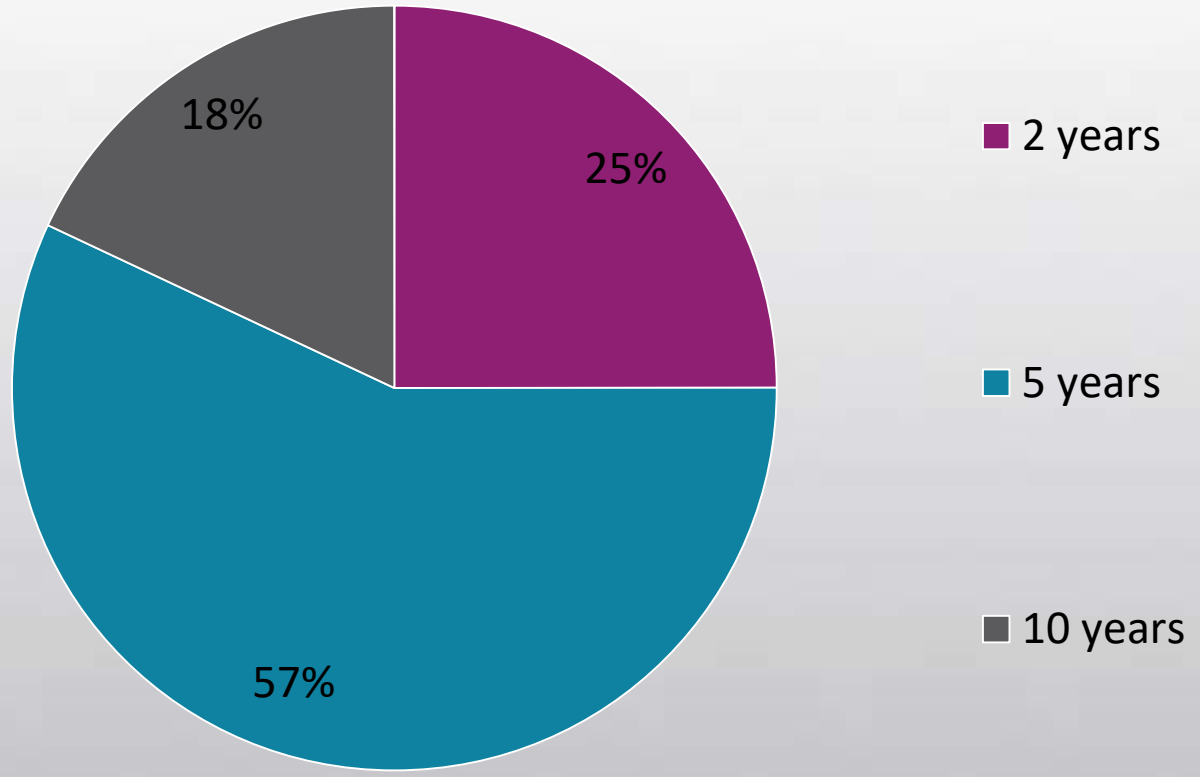
Results


19%
2%
20%
12%
47%
Local level multi-stakeholders
European level: bilateral agreements
European level: multi-stakeholders agreement
Global level: bilateral agreements
Global level: multi-stakeholders agreement

We should allocate resources to make the Code a reality within (?) years.

1. 2 years
2. 5 years
3. 10 years

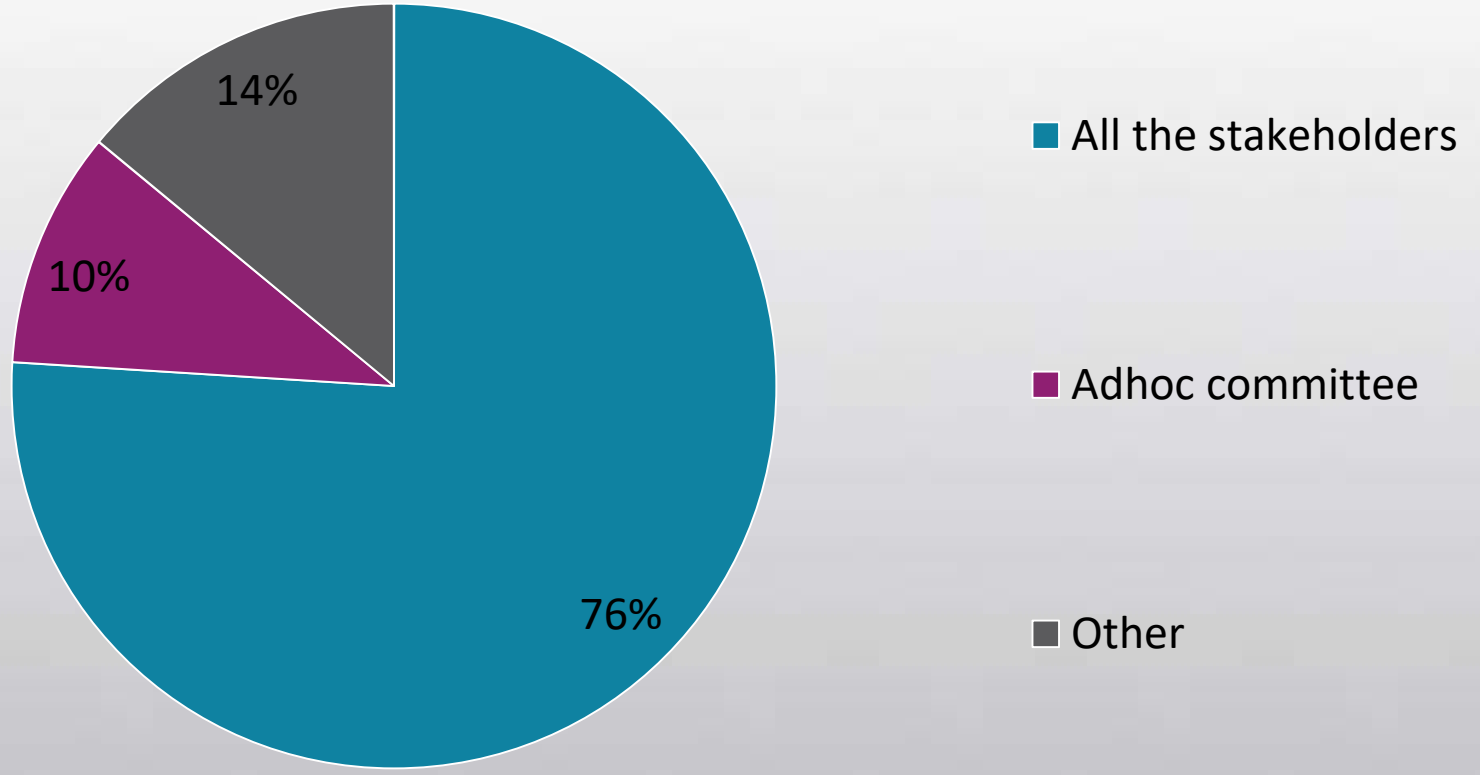
Results



Who should drive the development of a Stakeholders Code?

1. EFPIA, Medicines for Europe, MedTech Europe, AdvaMed, COCIR, Biomed Alliance and European Patients Forum
2. Ad hoc group of compliance officers from different stakeholder organisations
3. Other

Results



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