A NEW INDEPENDENT PARTNERSHIP

Stronger together.



CREACT Life Sciences Consulting welcomes

Chantal Deschamps (Synerva Life Sciences Consulting) as an independent collaborating partner

WHY THIS MATTERS

More than expanded expertise.

01

Agility

More flexibility when your development priorities shift.

02

Responsiveness

Better capacity to distribute workload and meet your critical deadlines.

03

Continuity

Two senior experts, no single-person dependency.

04

Long-term teamwork

An extension of your team, built on trust over time.

WHAT WE DO

Regulatory Affairs & CMC consulting

Regulatory strategy

Regulatory landscape, Roadmaps, gap assessments, regulatory intelligence and risk management.

Agency interactions

Scientific advice, pre-IND & INTERACT meetings, orphan and expedited designations.

Submissions

CTA / CTIS, · MHRA · TGA · IND Module 1, 2 and 3, IMPD, IB · PIP - iPSP authored and delivered

Technical CMC

QTPP / CQA, control strategy, comparability, stability and Module 3 authoring, Cell Banking and virus seed banking testing strategy, mRNA testing

One objective: streamline your drug development.

OUR COMBINED TRACK RECORD

Proven with biotechs, in front of the agencies

39+

biotechs supported,
plus 1 CDMO

14

scientific advices
EMA and national agencies FAMHP · ANSM · AIFA · PEI

13 / 16

orphan designations granted
EMA and FDA

11 · 7

IMPDs · Investigator's Brochures
authored

10

CTA/CTIS · CTA MHRA · CTA TGA (CTN) . IVD performance study

6

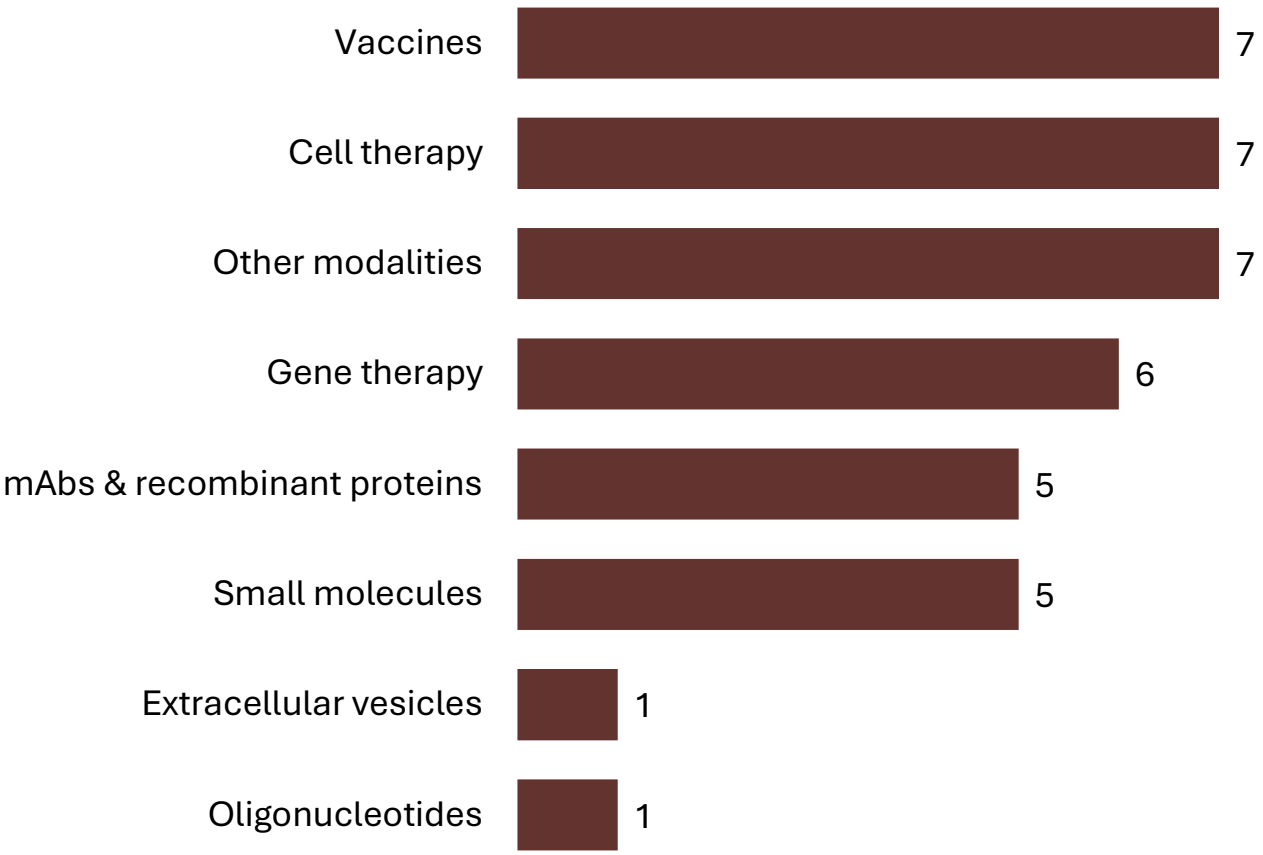
FDA Interact (2) · Pre-IND (3) · IND (1)

2

1 Module 3 BLA (US)
1 Module 3 MAA (MHRA)

ACROSS MODALITIES

Programmes supported, by modality



Deep CMC, hands-on: cell and virus banks, comparability, CQA / QTPP, BLA and MAA Module 3 authoring.

LET'S CONNECT

Let's discuss your development priorities

- Planning a clinical trial application?
- Facing a regulatory or CMC challenge?
- Need senior expertise, without the overhead of a large consultancy?

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