

## Appendix C – Contextual Deployment Scenarios

The following hypothetical scenarios illustrate system behavior across representative physiological contexts. These do not constitute efficacy claims but serve to demonstrate the structural fidelity of Co-Aegis under variable constraints. All cases assume v2.5 architecture unless otherwise noted.

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### C1 Immunocompetent Solid Tumour (CT26 model)

#### **Context:**

Subcutaneous colorectal tumour with stable hypoxic core ( $O_2 \leq 0.5\%$ ) and intact immune function.

#### **Observed sequence:**

- Hypoxia threshold reached ( $O_2 \leq 0.5\%$  sustained  $\geq 6$  min)
- spo0A expression initiated; germination proceeds
- HSV-TKmut PET signal confirmed at d2 post-injection ( $^{18}\text{F}$ -FHBG)
- Cas9 abort payload remains dormant (no doxycycline administered)
- On d5, plasmid instability detected (partial dapA dropout)
- Pfault module initiates PSM exposure
- No systemic CFU detected; bystander tissue unaffected

#### **Conclusion:**

System remained within design constraints. Exposure logic triggered by deviation and resolved by host immune clearance. No intervention required.

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## **C2 Immunocompromised Mucosal Lesion**

### ***Context:***

Post-transplant patient with neutropenia and suppressed T-cell function. Localized necrotic ulcer in lower gut.

### ***Observed sequence:***

- Passive spore ingress into lesion site
- O<sub>2</sub> fluctuation between 0.7–1.1 %; germination not triggered
- On d3, transcript-level instability detected (Cas9 silencing event)
- Pfault module initiates PSM exposure
- No immune activity observed over 72 h
- PET remains negative (HSV-TKmut inactive)
- TMR-03 signature detected in urine at d4
- External biologic (anti-FLAG nanobody) administered on d4
- Clearance of exposed constructs confirmed within 48 h

### ***Conclusion:***

System signaled fault correctly. Lack of immune surveillance prevented autonomous resolution. Clearance required biologic intervention. No off-target activation occurred.

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### **C3 Non-Tumour Hypoxia: Chronic Ischemic Ulcer**

#### **Context:**

Peripheral chronic ulcer with localized ischemia; intermittent O<sub>2</sub> dips to 0.9 % but no sustained hypoxic descent.

#### **Observed sequence:**

- Multiple transient O<sub>2</sub> events (O<sub>2</sub> ≤ 1.2 % for < 4 min)
- Hypoxia persistence threshold not met
- spo0A remained suppressed; no germination
- Harmonizer peptide expressed; no inflammatory markers detected
- No exposure logic triggered
- Natural clearance observed by d6

#### **Conclusion:**

Refined germination gating prevented false activation in marginal hypoxic zone. Dormancy harmonizer preserved ecological neutrality. Passive presence resolved without immune event.