

# knkdns

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## **STATEMENT OF COMPLIANCE**

The trial will be carried out in accordance with International Conference on Harmonisation Good Clinical Practice (ICH GCP) and the following: • United States (US) Code of Federal Regulations (CFR) applicable to clinical studies (45 CFR Part 46, 21 CFR Part 50, 21 CFR Part 56, 21 CFR Part 312, and/or 21 CFR Part 812) • National Institutes of Health (NIH)-funded investigators and clinical trial site staff who are responsible for the conduct, management, or oversight of NIH-funded clinical trials have completed Human Subjects Protection and ICH GCP Training. • The protocol, informed consent form(s), recruitment materials, and all participant materials will be submitted to the Institutional Review Board (IRB) for review and approval.

## 1 PROTOCOL SUMMARY

### 1.3 Schedule of Activities (SoA)

[illegible]

3 OBJECTIVES AND ENDPOINTS



**Figure: nexus**  
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Type	dc dc vdc v	Endpoints	fdjivedhv
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deygdyegwfyegf	efvewgbeyyef	geyfre	hfuwehf
ewgfywefgweywey	hgcuyfwgfywe	ufweuf	wefgefefuew
gfeugewyfbewf	yefbewgew	yfgewfyge	hwebfweyfgew

