

## K24 CHECKLIST: PA-20-193 (CLINICAL TRIAL REQUIRED)

Component	Page limit	Notes
Project Summary/Abstract	30 lines	
Project Narrative	3 sentences	
Bibliography & References Cited		Include references in Career Development Award supplemental form and Clinical trials information forms
Facilities & Other Resources		
Equipment		End with sentence: "Stanford equipment and resources listed above are available for use by the investigator at no direct cost to the sponsor unless specified in the budget and justification"
Biosketch	5 pages each	Key personnel: PI/PD, mentors, co-mentors, collaborators, consultants, advisory committee members
Current & Pending Support	3 pages	Must include Project Number, Source, Major Goals, Project / Proposal Start and End Dates, Total Award Amount.
Budget Justification		
Candidate Information and Goals for Career Development	6 pages (12 pages total with Research Strategy)	Sections: 1. Candidate's background 2. Career Goals and Objectives 3. Plan for Career Development
Specific Aims	1 page	
Research Strategy	6 pages (12 pages total with Candidate Information)	
Progress report publication list		For renewal applications: List titles and complete references of all publications resulting from the project since last competitive review
Training in Responsible Conduct of Research	1 page	
Candidate's Plan to Provide Mentoring	6 pages	Describe proposed and past mentoring activities
Description of Institutional Environment	1 page	Can include names of other faculty members willing to collaborate.
Institutional Commitment to Candidate's Research Career Development	1 page	Provided on institutional letterhead. Include statement that candidate will receive 3-6 person months (25-50% FTE) protected time.
Authentication of Key Biological and/or Chemical Resources	1 page	

### PDF Components for Human Subjects and Clinical Trials Information

Inclusion of Individuals  
Across the Lifespan

Inclusion of Women and  
Minorities

Recruitment and  
Retention Plan

Study Timeline

Protection of Human  
Subjects

Sections:

1. Risks to Human Subjects
2. Adequacy of Protection Against Risks
3. Potential Benefits of the Proposed  
Research to Research Participants and  
Others
4. Importance of the Knowledge to be Gained

Data and Safety  
Monitoring Plan

Statistical Design and  
Power

Dissemination Plan