

## **Event Checking Chart**

## Cases of Cancer, both AIDS and Non-AIDS Defining Cancers

Name of centre and cohort Patient ID code: Year of birth (yyyy):		Gender: [ ] Male [ ] Female Date of first diagnosis (dd/mm/yy):		
1.	<ul> <li>Diagnosis         Please complete this form if the patient has been diagnosed with a malignant disease (excluding pre-cancers, relapses, basal and squamous cell skin cancers)     </li> <li>For the patients' cancer disease, please provide specific type:</li></ul>			
	Primary location (if known):	(e.g.	lung); unknown [ ]	
	If available, please include the: ICD-10	e: ICD-10, or ICD-9 code		
2. \$	<ul> <li>2. Stage (spread) at diagnosis (Tick one only):</li> <li>[] Localized (growth within the organ of origin)</li> <li>[] Disseminated (spread to tissue outside the organ of origin, incl to regional lymph nodes)</li> <li>[] Unknown</li> </ul>			
3. I	Histology/cytology Is a pathology report (or summary hereof) available [ ] Yes, full report [ ] Summary of report for 'unknown', please complete Question 4		[ ] Unknown	
	If yes, please include a copy of the full report	(And provide a brief sum	nmary in English):	
4. I	<ul> <li>4. If the diagnosis is not confirmed by histology/cytology, is the diagnosis based on (Tick all that apply, 1 at a minimum, and please provide source documentation):</li> <li>I. [] Radiology or other imaging technique (cancer suggestive findings)</li> <li>II. [] Biochemical assay (elevated markers of cancerous growth (e.g. prostate specific antigen, alpha-fetoprotein, cancer cell markers)</li> <li>III. [] Strong suspicion of cancer by clinical inspection (skin metastasis, suspected malignant melanoma, suspected cancerous growth visualized during endoscopy/anoscopy)</li> <li>IV. [] Other</li> <li>Of those marked above, please specify:</li> </ul>			
dis If y	Has the patient previously received chemo- and isease? [ ], Yes [ ], No yes, please tick off the appropriate box: chemother or fatal cases, please also complete a CoDe form.	[ ], Unkr	nown	
Sig	ignature:the Study Cod	ordinating Office, Date:_	(dd/mm/yyyy)	
Monitored at site by: Date: Date: dd/mm.			Date:	

Please return this form to the DAD study coordinating office incl. copies of other relevant documents from the medical record (made anonymous and labeled with the patients ID-code) by air- or email and provide the cohort coordinating office with a copy of the chart.