Evaluation sheet for the efficacy of Uridine ointment for the treatment of the hand-foot-syndrome induced by 5-Fluorouracil (5-FU) or the oral derivates Capecitabin (C) or Uracil/Tegafur (UFT).

Patient (Initials + Age)	Gender ☒		Causing agent (5-FU/C/UFT)	Tox. grade (CTC[1])	Time of treatment with 5-FU/C/UFT (cumulative dose)	Affected (H, F, H+F?[2])	Duration of treatment [weeks] until *distinct* onset of relief	Inpatient (**I**) or Outpatient (**O**) treatment	Frequency of use per day (min. q3d)
	Male (**M**)	Female (**F**)							

[1] **CTC Version 2, revised4/30/99:** adverse event → Dermatology/Skin → Hand-foot skin reaction

Grade

0 None
1 skin changes or dermatitis without pain (e.g. erythema, peeling)
2 skin changes with pain, not interfering with function
3 skin changes with pain, interfering with function
4 not defined

[2] H = Hand, F = Foot, H+F = both