International Agency for Research on Cancer



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Type of evaluation	Agent	Evidence
Efficacy	Screening by conventional cytology	Sufficient
	Screening by liquid-based cytology	Sufficient
	Screening by automated cytology	Sufficient
	Testing for human papillomavirus infection as the primary screening modality	Sufficient
	Screening by visual inspection with application of acetic acid (VIA)	Limited
	Screening by visual inspection with application of Lugol's iodine (VILI)	Limited
Overall	Screening by conventional cytology, every 3–5 years between the ages of 35 and 64 years, in a high-quality programme	Sufficient
	HPV testing, using a validated system, as the primary screening modality	Sufficient
	Screening by visual inspection with application of acetic acid (VIA) or with Lugol's iodine (VILI)	Limited