Figure: 25 TAC §289.229(l)

Name of Record		Rule Cross-Reference	Time Interval Required for Record Keeping
Accelerators used for research and development and industrial operations			
(A)	Initial surveys	(f)(2)(C)	Until termination of registration
(B)	Tests and repairs	(f)(3)(A)(x)	5 years
(C)	Calibration, surveys	(f)(3)(F)	5 years
(D)	Contamination smear for units operating greater than 10 MeV	(f)(3)(G)	Until termination of registration
(E)	Receipt, transfers, and disposal	(f)(3)(H)	Until termination of registration
(F)	Training for operators	(f)(4)(B)	Until 2 years after the individual terminates employment
Therapeutic radiation machines, radiation therapy simulation systems, and EBT devices			
(G)	Credentials of operators EBT device operators	(h)(1)(C) (h)(1)(E)	Until 2 years after the individual terminates employment
(H)	Review of quality assurance program	(h)(1)(F)(vii)	5 years
(I)	Written OSP	(h)(1)(G)	Until transfer of machine or termination of registration
(J)	FDA variances	(h)(1)(H)	Until transfer of machine or termination of registration
(K)	Initial and Subsequent Surveys Therapy (below 1 MeV) Therapy (1 MeV and above) EBT device	 (h)(2)(D)(i)(II) (h)(3)(C)(i)(III) (h)(4)(B)(i)(II)	Until termination of registration

(L)	Calibration Therapy (below 1 MeV) Therapy (1 MeV and above) EBT device	 (h)(2)(D)(ii)(V) (h)(3)(C)(ii) (h)(4)(B)(ii)	5 years
(M)	Contamination smears for units operating greater than 10 MeV	(h)(1)(I)	Until termination of registration
(N)	QA checks and corrective actions Therapy (below 1 MeV) Therapy (1 MeV and above) EBT device	 (h)(2)(D)(iii)(VI) (h)(3)(C)(iii)(VII) (h)(4)(B)(iii)	5 years after the QA checks
(O)	Leakage measurements Therapy (1 MeV and above)	 (h)(3)(A)(i)	5 years
(P)	Protective devices forradiation therapy simulation systems	(h)(5)(A)(iii)(II)	3 years
(Q)	Film processing recordsfor simulators	(h)(5)(E)(i)(V), (VI), and (ii)	3 years
(R)	Digital imaging acquisition systems	(h)(5)(E)(iii)	3 years
(S)	CT dose measurements	(h)(5)(C)(iv)(III)	5 years
(T)	CT films resulting from quality control tests	(h)(5)(D)(ii)	1 year or until a new phantom image is performed
(U)	Reports of medical events	(j)(2)	Until termination of registration